Patented Medicine Prices Review Board

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Conseil d'examen du prix des médicaments brevetés

Year-over-Year Changes in the Patented Medicine Price Index 1988-1997

Tenth Annual Report

For the Year Ended December 31, 1997

1988 1989 1990 1991 1992 1993 1994 1995 1996 1997

Visit us at our Web site: http://www.pmprb-cepmb.gc.ca or call us at our toll-free number: 1-877-861-2350



The Patented Medicine Prices Review Board Standard Life Centre Box L40 333 Laurier Avenue West Suite 1400 Ottawa, Ontario K1P 1C1

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Conseil d'examen du prix des médicaments brevetés

May 29, 1998

The Honourable Allan Rock, P.C., Q.C., M.P. Minister of Health House of Commons Ottawa, Ontario KIA 0A6

Dear Minister:

I have the honour to present to you, in accordance with the provisions of sections 89 and 100 of the *Patent Act*, the Annual Report of the Patented Medicine Prices Review Board for the year ended December 31, 1997.

Yours very truly,

-Is Elqu Robert G. Elgie

Robert G. Elg Chairperson

Mission and Values of the PMPRB



The mission of the Patented Medicine Prices Review Board (PMPRB) is to contribute to Canadian health care by ensuring that prices of patented medicines are not excessive. The PMPRB achieves this by:

- promoting voluntary compliance with Guidelines established by the Board
- reviewing prices and taking remedial action when necessary
- analysing and reporting to Canadians on price trends of all medicines and on research and development conducted by patentees
- consulting with interested parties on Guidelines and other matters of policy
- fostering awareness of the Board's mandate, activities and achievements through communication, dissemination of information and public education.

In fulfilling its mission the PMPRB is committed to innovative leadership based on the following values:

- effectiveness and efficiency
- fairness
- integrity
- mutual respect
- transparency of process
- a supportive and challenging work environment.

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Chairperson's Message

As it moved into its second decade of operations in 1997, the Patented Medicine Prices Review Board focussed on identifying ways which would enable it to become more effective, responsive and accessible to Canadians while functioning in an increasingly dynamic environment. New drugs, new medical technologies and an aging population are just some of the forces which are influencing the evolution of our health care system and the work of the PMPRB.

Consistently, health care ranks as the number one issue of importance to Canadians. In this context, Canada, along with many other countries, continues to look for ways to contain rising costs. In 1996, according to the latest figures published by Health Canada, drugs alone accounted for 14.4% of total health care expenditures, representing the fastest growing cost component within our health care system. This trend underscores the need for more and better information on key elements such as hospital, physician and drug costs and on cost drivers such as utilization, prescribing habits of physicians, pricing and other factors. The Federal/Provincial/Territorial Ministers of Health have assigned task forces to look into these matters.

Many of these concerns were voiced by witnesses during the hearings of the Standing Committee on Industry reviewing Bill C-91. Echoing these concerns in its April 1997 Report, the Standing Committee noted that "... many witnesses wanted [the PMPRB] to fulfill a wider role in a more publicly responsive way... [prompting the recommendation to the Government that] the mandate of the [PMPRB] be reviewed and strengthened." In addition, the Standing Committee recommended that "... the PMPRB consult with consumers, health care professionals, experts, and the provinces to assess its current statistical reporting and find out what other information it could provide to the public."

In 1997, the Board embarked on a comprehensive review and renewal process examining a range of issues, including those raised in the Standing Committee report. The Board initiated a broad consultation with its stakeholders to examine the role, functions and methods of the PMPRB with a view to being more relevant to the needs of those it serves. While the Board has always endeavored to consult widely, this expanded consultation process represented a major, new undertaking for the PMPRB.

In November 1997, the Board released a detailed Discussion Paper, followed in February and March 1998 by a series of public information sessions held in every province and territory. The purpose of the consultation was to solicit constructive input and suggestions regarding issues of concern that are within the Board's mandate and scope of activities. The main areas covered in the Discussion Paper include: drug prices and cost issues; consultation, information and transparency issues as they pertain to the Board's public accountability; and issues regarding pricing methods and guidelines.

To date, the consultation process has proven to be worthwhile on several levels. It has enabled the Board members to hear directly from consumers and others regarding their concerns as well as encouraging stakeholders to express their views on how these concerns could and should be addressed. Similarly, the consultations have furnished valuable feedback on how the Board might improve the "way we do business." In effect, the consultation process has already served to influence the way the Board functions in terms of an increasing emphasis on availability, access and two-way communications. The impact and benefits of the consultation process can be measured in other ways as well. Results thus far have served to validate the Board's direction toward strengthening public accountability. From an operational perspective, the Board is currently working on the development and implementation of a formal consultation policy supporting the broader objective of further strengthening the PMPRB's public accountability. Work is also moving ahead on a review



ioto courtesy of Health Cana

of the Board's pricing methods and Guidelines to ensure their appropriateness and, where possible, identify opportunities for improvement. In addition, the Board will be conducting further analyses and will be reporting on issues surrounding drug prices and cost drivers.

In 1997, the Board worked to better identify and respond to the needs of consumers and other stakeholders, to promote a process of continuous dialogue and exchange of ideas, and to strengthen public accountability and operational transparency. The Board is always seeking to enhance the transparency of its operations by, for instance, developing mechanisms to report publicly on the Board's analyses of new medicines. In addition, the Board is also exploring new approaches to the way it operates as reflected by its recent invitation for comments relating to the drug Humalog prior to deciding whether to accept a proposed Voluntary **Compliance Undertaking** (VCU) to lower the price of this medicine.

In last year's Annual Report, the PMPRB signaled its intention to place greater emphasis on consultations with its stakeholders, to keep them better informed and to promote public debate on the pricing, usage and cost of drugs in Canada. In 1997, furthering these goals has been the focal point of our efforts as an organization. The Board has worked to better identify and respond to the needs of consumers and other stakeholders, to promote a process of continuous dialogue and exchange of ideas, and to strengthen public accountability and operational transparency. To further these goals, the Board continues to collaborate with the provinces in a variety of areas. In particular, the Board provides expertise to the Federal/Provincial/Territorial Task Force on Pharmaceutical Pricing Issues as well as being a member of the Pharmaceutical Issues Committee. The Board wants to ensure the highest possible degree of efficiency, integrity and fairness in Canada's patented medicine price review system.

As Chairperson of the PMPRB, I am especially proud of the Board members and PMPRB staff, particularly with respect to the professionalism they have demonstrated in supporting the consultation and renewal process which has involved so much additional effort during the past year.

In embarking on our second decade, yet another milestone in the life of our agency has been marked with the departure from the Board of Dr. Harry C. Eastman. Dr. Eastman served as Chairperson and Chief Executive Officer from the Board's founding in 1987 to 1995, as well as serving as a Board member from 1987 to 1997. On behalf of all Board members and PMPRB staff, I wish to thank Dr. Eastman for his invaluable contribution to the PMPRB over the past decade. In renewing itself, the PMPRB is building upon the experience and accomplishments of its first ten years, while initiating the changes and improvements necessary to ensure its continued effectiveness in protecting the interests of Canadian consumers.

Colut Is Elqu

Robert G. Elgie Chairperson

Mandate

The PMPRB is an independent quasi-judicial body created by Parliament in 1987 under the *Patent Act*. The PMPRB protects consumer interests and contributes to Canadian health care by regulating the prices charged by manufacturers of patented medicines to ensure that they are not excessive.

The PMPRB reports to Parliament through the Minister of Health. The Annual Report, which covers each calendar year, includes a review of the PMPRB's major activities, analyses of the prices of patented medicines and of the price trends of all drugs, and reports on the R&D expenditures by patent-holding drug manufacturers.

Jurisdiction

The PMPRB is responsible for regulating the maximum prices that patentees may charge for prescription and non-prescription patented drugs sold in Canada for human and veterinary use. In most cases that price is the "factorygate" price at which the manufacturer sells the product to wholesalers, hospitals or pharmacies. The PMPRB's jurisdiction includes patented medicines marketed or distributed under voluntary licences. The Board has no authority to regulate the prices of non-patented drugs, including generic drugs sold under compulsory licences, and does not have jurisdiction over prices charged by wholesalers or retailers nor over pharmacists' dispensing fees.

In Canada, Health Canada assesses new medicines to ensure that they conform with the *Food and Drugs Act* and *Regulations*. Formal authorization to market or distribute a medicine is granted through a Notice of Compliance (NOC). A medicine may be temporarily distributed with specified restrictions before receiving a NOC, as an Investigational New Drug or under the Special Access Program.

The PMPRB regulates the price of each patented drug product, including each strength of each dosage form of a patented medicine. This is normally the level at which Health Canada assigns a Drug Identification Number (DIN) or General Public (GP) number.

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Membership

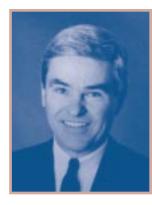
The Board consists of no more than five part-time members appointed by the Governor in Council for a term of five years. There is currently one vacancy on the Board.



Chairperson: **Robert G. Elgie,** LL.B., M.D., F.R.C.S. (C)

Dr. Elgie, a lawyer and neurosurgeon, Fellow of the Royal College of Surgeons (Neurosurgery), was the founder and first Director of Dalhousie University's Health Law Institute from 1991 to 1996. He was also the part-time Chair of the

Workers' Compensation Board of Nova Scotia from 1992 to 1996. Dr. Elgie has taught at the Medical Schools of Queen's University and the University of Toronto, and has held several positions with the Scarborough General Hospital, including Chief of Medical Staff. In 1977, he was elected to the Ontario Legislative Assembly and subsequently served in several Cabinet positions. He resigned from the Ontario Legislature in September 1985 to become Chair of the Workers' Compensation Board of Ontario where he served until 1991. Dr. Elgie was appointed Member and Chairperson of the PMPRB in March 1995.



Vice-Chairperson: **Réal Sureau**, FCA

Mr. Sureau, a chartered accountant, is President of Sureau Management Limited and Director, Business Development, Montreal Baseball Club Inc.

From June 1995 to June 1996, he was President of the Order of Chartered

Accountants of Québec. Through the years, he was a member of several committees of the Order, including the Disciplinary Committee, the Professional Practice Committee, the Professional Development Committee and the Committee on Government Finances. He was Vice-President, Finance, at Forex and CanamManac. Mr. Sureau sits on the board of directors of many organizations, including Gaz Métropolitain, the *Institut de réadaptation de Montréal and la Fondation des paraplégiques du Québec*. Mr. Sureau was appointed Member and Vice-Chairperson of the PMPRB in October 1995.

Members:

Judith L. Glennie, Pharm., D., FCSHP

Dr. Glennie is a clinical pharmacist specializing in pharmacoeconomics. She is currently President of J. L. Glennie Consulting Inc. and works with the Ottawa General Hospital Pharmaceutical Outcomes Research Unit and various governmental and non-governmental health agencies. She is an affiliate researcher with the Loeb Research Institute and adjunct professor with the Faculty of Medicine, University of Ottawa. Dr. Glennie is completing a Masters of Science in Community Health Science, Faculty of Medicine, University of Manitoba. Dr. Glennie was appointed Member of the PMPRB in March 1995.



Ysolde Gendreau, B.C.L., LL.B., LL.M., Ph.D.

Dr. Gendreau is a professor in the Faculty of Law of the Université de Montréal, where she teaches intellectual property law and competition law. She is also sessional lecturer at McGill University, where she teaches intellectual property law. Dr. Gendreau was appointed Member of the PMPRB in October 1995.



Consultations: Examining the Role, Functions and Methods of the Patented Medicine Prices Review Board

In April 1997, the Standing Committee on Industry reviewed the drug patent legislation, Bill C-91. Although the main issues during the review were the regulatory framework for drug patent policy, Linkage Regulations, patent terms, pharmacare and related matters, the mandate and activities of the Patented Medicine Prices Review Board were also examined and debated.

The Board's mandate flows from policy established by Parliament through the legislation and its regulations and is focussed on protecting consumer interests by ensuring that prices for patented drugs are not excessive. In its report, the Standing Committee noted that witnesses tended to give only limited recognition of the Board's success to date in controlling drug prices.

Consequently, the Committee made two recommendations directly related to the Board. The first such recommendation was addressed to the government and advocated:

- strengthening the mandate of the PMPRB;
- an audit by the Auditor General; and
- greater access to non-proprietary information filed with the PMPRB.

The second recommendation on the PMPRB was addressed directly to the Board. This recommendation urged the Board to consult with consumers, health care professionals, other experts, and the provinces to assess its current statistical reporting and to determine what other information might be gathered and shared with the public. From its perspective, the Board recognized the value and appropriateness of looking at opportunities to improve the transparency of the regulatory and price review process. In addition, the Board acknowledged the need to further the discussion with respect to concerns that the Committee heard as to the growth of expenditures on drugs and the growing cost to the health care system.

Responding to the Challenge

The Board initiated an internal review of its activities to examine existing functions and to relate them to potential new requirements arising out of the Committee's recommendations.

A major element of this review was to seek advice, on an informal basis, from some of our stakeholders who, traditionally, had not been closely involved in the work of the Board. The purpose of this advice gathering was to determine how the Board might enhance its consultations. The results of this initiative were used to help shape the consultation process that followed.

In November 1997, the Board launched a public consultation process through the publication of a Discussion Paper. The first step in the consultations centered on the role, functions and methods of the Board and on questions relating to the Board's role in disseminating information, strengthening its public accountability, as well as its price guidelines and price review methodologies.

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Process

We sought to ensure that the consultations would be as broad and inclusive as possible. Towards this end, the Discussion Paper was circulated to the people on our regular mailing list, approximately 900 recipients. As other interested parties were identified or came forward to request a copy of the Discussion Paper, we eventually reached a point of having more than 2,000 copies in circulation.

We also recognized the importance of ensuring the opportunity for face-to-face contact to encourage two-way communications. During February and early March, the Board held a series of public information sessions across Canada. These sessions were preceded by notifying people on our mailing list, issuing media advisories, and publishing notices in major regional newspapers — all designed to encourage public involvement in the review. A session was held in every province and territory. In total, thirteen meetings took place, conducted by the Board members and staff and were attended by close to 300 Canadians.

To further support the consultation process, we invited written submissions to be filed by the end of March. The Board received 60 substantive submissions from interested organizations and individuals. In total, these submissions run to more than several hundred pages of text.

At the end of April, the Board held a two-day policy hearing at which 24 organizations or individuals made presentations in support of their submissions. The purpose of this hearing was to ensure that interested stakeholders were afforded every opportunity to clarify and, as appropriate, elaborate upon the information contained in their submissions.

Results and Next Steps

The Board is currently evaluating the submissions and plans to make a report in late summer which may set out some changes and proposals for further consultation.

We will be communicating these results through distribution of printed copies of the report, our web site and our NEWSLETTER. We are also encouraging stakeholders to communicate with us through our toll-free number: 1-877-861-2350 or via e-mail (pmprb@pmprb-cepmb.gc.ca).

We have concluded that the release of the Discussion Paper and the holding of the consultation sessions across Canada were worthwhile initiatives for the Board. They have afforded us direct contact with consumers and other stakeholders while the submissions, oral and in writing, represent significant sources of new information and feedback on issues and how they might be addressed.

This expanded consultation process represents a new stage in the evolution of the Board with an increased emphasis on ensuring closer links with consumers and other stakeholders. The Board is committed to strengthening its public accountability including the transparency of its operations.

Future Direction of the Board

Where we can identify ways of improving our effectiveness, we are doing so and will continue to do so. We recognize the need to remain relevant to the needs of consumers and our other stakeholders by promoting a constant exchange of information, improving the timeliness and quality of information we provide and ensuring that our pricing methods and guidelines are appropriate in helping us fulfill our mandate.

Sales of Drugs in Canada in 1997

The Pharmaceutical Industry in Canada

The global pharmaceutical industry is dominated by a number of large multinational enterprises based in several countries. Most of these companies have Canadian subsidiaries which, along with a few domestic pharmaceutical firms, account for the manufacture, sale and distribution of drugs in Canada. It has been reported that the top ten pharmaceutical companies accounted for approximately 50% of total sales in 1997, up from 45% in 1996.¹ Of the top



ten firms, one was a Canadian company supplying generic products.²

It has been reported that sales of drugs worldwide increased in 1997 by 8.6% over 1996 to over \$400 billion. In Canada, total sales of drugs increased at about the same rate. 7.0%, to an estimated total of \$7.0 billion. The Canadian market for drugs has traditionally represented less than 2% of the world market.

In Canada, the PMPRB protects consumer interests by regulating the prices charged by pharmaceutical patentees during the time they benefit from patent protection. Ordinarily, a patentee is the exclusive supplier of a patented product, but there may also be other medicines available. Although a drug that is no longer protected by a patent may become the subject of competition, there are also instances where the manufacturer of a non-patented drug may nevertheless remain the sole supplier.

In 1997, 76 companies reported sales of patented medicines in Canada to the PMPRB, an increase from 72 in 1996.³

The pharmaceutical industry continued to account for less than 2% of all sales and employment in the manufacturing sector of the Canadian economy in 1997. Because of its research and development (R&D) activities, however, the industry accounted for approximately 10% of total R&D in that sector.⁴ This is consistent with this industry's relative performance since 1987. (For a report on patentees' R&D expenditures in 1997, please refer to page 28).

Sales of Pharmaceutical Drugs in Canada

Table 1 shows manufacturers' sales of all drugs and of patented drugs in Canada since 1990. Total sales by manufacturers of pharmaceuticals in 1997 in Canada are estimated

¹ See IMS. Canadian Pharmaceutical Industry Review. 1997. Table 1.

² In 1996, two Canadian companies were in the top ten, Apotex and Novopharm. In 1997, Novopharm moved to eleventh position. See IMS, Canadian Pharmaceutical Industry Review, 1997, Table 1.

³ A list of all reporting patentees and patented drug products appears in Table 13.

Statistics Canada Catalogues, 31-2003 and 88-202.

| Year | Tot | al | Pate | nted | Patented as |
|------|-----------------------|----------------|-----------------------|----------------|------------------------|
| | Sales (\$billions) | Change* (%) | Sales (\$billions) | Change* (%) | Percentage of Total |
| 1997 | 7.0 | 7.0 | 3.7 | 22.6 | 52.3 |
| 1996 | 6.6 | 10.0 | 3.0 | 12.8 | 45.0 |
| 1995 | 6.0 | 1.7 | 2.6 | 10.8 | 43.9 |
| 1994 | 5.9 | 9.3 | 2.4 | -2.1 | 40.7 |
| 1993 | 5.4 | 12.5 | 2.4 | 9.4 | 44.4 |
| 1992 | 4.8 | 9.1 | 2.2 | 14.0 | 43.8 |
| 1991 | 4.4 | 18.9 | 2.0 | 13.1 | 43.2 |
| 1990 | 3.7 | - | 1.7 | - | 43.2 |

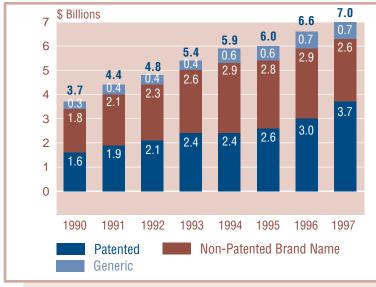
Table 1 Manufacturers' Sales of All Drugs and Patented Drugs, 1990 - 1997

* Percentage changes reflect exact values and not rounded values. Source: PMPRB, Statistics Canada and IMS Canada.

at \$7.0 billion, an increase of 7.0% from 1996.⁵ In 1997, patentees reported total factory-gate sales of patented drugs of \$3.7 billion, an increase of 22.6% from 1996. This increase

represents the largest annual growth in patented drug sales during the 1990's. Furthermore, for the first time patented drug sales accounted for over 50% of the total sales

Figure 1 Manufacturers' Sales of Patented and Non-Patented Drugs 1990-1997



Sources: PMPRB, Statistics Canada and IMS Canada

by manufacturers. Since prices and sales have been tracked by the PMPRB patented drugs had never accounted for more than 45% of the total sales. For further comments on this issue, please refer to page 20, Trends in Quantities of Sales of Patented Drug Products.

Figure 1 shows the growth in annual sales of patented and non-patented drugs from 1990 to 1997. Sales of non-patented drugs can be estimated as the difference between the total sales of pharmaceuticals and the sales of patented drugs as reported to the PMPRB.

⁵ See Statistics Canada Cansim #'s D667757, D315488, D401624 and D451712.

The most recent figures on annual total sales from Statistics Canada are for 1995. The PMPRB relies on the most current annual report of sales by Statistics Canada when it becomes available. Beginning with the 1995 Annual Report the PMPRB has adopted a method of estimating total sales for the years that Statistics Canada's data are not available.

Patentees are required, under the Patented Medicines Regulations, to submit to the PMPRB annual total pharmaceutical sales information for both patented and non-patented drug products sold in Canada. IMS Canada Ltd. publishes reports of sales by individual firms. Total sales by all manufacturers can be estimated by adding the total sales of patentees and the estimated IMS sales for non-patentees. These estimates will be revised when the annual sales estimates published by Statistics Canada become available.

Non-patented medicines include products that were previously subject to patent protection, those that are not yet or never have been protected by a patent, and generic copies of existing patented drugs. Information filed by patentees with the PMPRB indicates that most non-patented drugs are sold by companies that also sell patented drugs. During the 1990's sales of non-patented drugs typically accounted for about 50% of the total drug sales of patentees. In 1997, the proportion of non-patented drug sales dropped substantially to about 40% while patented drug sales increased to about 60% of the total drug sales of patentees.

Non-patented sales also include sales by generic companies. IMS Canada estimates sales by non-patentees to be \$710 million or 11% of the total pharmaceutical market in 1997, a slight decline from 1996.⁶

The PMPRB regulates the maximum prices of individual patented drug products, including each strength and dosage form of them. A total of 981 patented drug products were sold in Canada during all or part of 1997. This number represents an increase from the 917 products sold in 1996. The number of patented drugs reported to the PMPRB is smaller than the 1,431 patented drugs with a Notice of Compliance (NOC) because not all products with a NOC were actually sold in Canada during the year. In addition, patented drugs that do not have a NOC but are sold as Investigational New Drugs or under the Special Access Program administered by Health Canada are subject to review by the PMPRB. In 1997, there were over 21,000 drug products sold in Canada. Of these drug products approximately 6,000 were prescription drugs. Although the number of patented drug products with a NOC in 1997 represented only 6% of the total number of drug products approved for sale, sales of patented drugs accounted for 52.3% of total sales as shown in Table 1.



^{photo} courtesy of Health Canada

⁶ See IMS, *Canadian Pharmaceutical Market: Drug Store and Hospital Purchases*, December 1997. Generic companies include those that belong to the Canadian Drug Manufacturers Association (CDMA).

Trends in Drug Prices and Expenditures

Prices of Patented Drugs in 1997

The PMPRB maintains the Patented Medicine Price Index (PMPI), an index of manufacturers' prices for patented drugs as reported annually by the PMPRB. The PMPI measures the average change from the previous year in the actual prices of all patented drugs sold in Canada. Because the PMPI is derived from the actual prices charged by manufacturers for all patented medicines, it provides a precise measure of price changes for drugs reported to the PMPRB.⁷

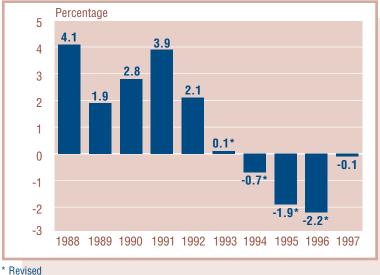
In 1997, manufacturers' prices of patented drugs were virtually unchanged from 1996. The prices of patented drugs, as measured by the PMPI, fell by 0.1% from the level in 1996. As shown in Figure 2, prices for patented drugs had declined by about 2% per year in the previous two years; the PMPI went down 2.2% in 1996 and 1.9% in 1995.

Price Trends of All Drugs — Patented and Non-Patented

The *Patent Act* provides that the PMPRB shall consider changes in the Consumer Price Index (CPI) in determining if the price of a patented medicine is excessive. As shown in Figure 3, consumer prices, as measured by the CPI, increased in every year since 1988 by an amount greater than the PMPI with the exception of 1992.⁸ In 1997, consumer prices increased by 1.6%.

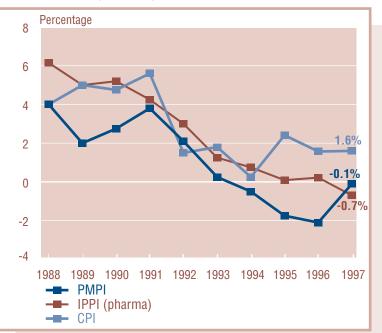
It is not unexpected that the overall increases in patented drug prices have been less than the increases in the CPI. The PMPRB limits apply on a product-by-product basis; in other words, no patented drug product can have its price increase by more than the CPI, and some will increase by less. The policies of some provincial governments also influence the annual rate of change in drug prices.

Figure 2 Year-over-Year Changes in the PMPI 1988-1997



Source: PMPRB

Figure 3 Year-over-Year Changes in PMPI, IPPI (pharma) and CPI 1988-1997



Sources: PMPRB and Statistics Canada

⁷ See the PMPRB's A description of the Laspeyres methodology used to construct the Patented Medicine Price Index (PMPI), April, 1997, for an explanation of the PMPI.

⁸ To facilitate and encourage compliance by patentees, the PMPRB's CPI-adjusted methodology uses the forecast rate of CPI inflation published by the Department of Finance. The methodology is self-correcting over time. The forecast CPI inflation rate for 1992 had been 3.2% but the actual rate was 1.5%. For a full explanation of the CPI-adjusted methodology please refer to Schedule 4 of the PMPRB's *Compendium of Guidelines, Policies and Procedures*.

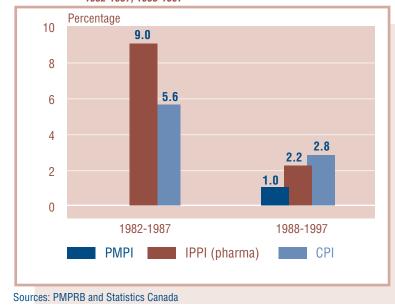
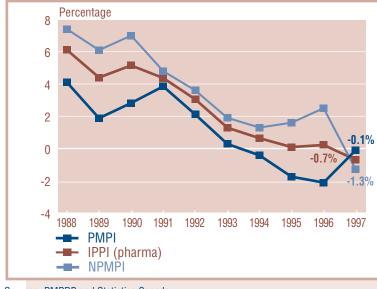


Figure 4 Summary of Price Trends Average Annual Percentage Changes 1982-1987; 1988-1997

Figure 5 Year-over-Year Changes in the IPPI (pharma), NPMPI and PMPI 1988-1997



Sources: PMPRB and Statistics Canada

Industrial Product Price Index (IPPI)

The pharmaceutical component of the Industrial Product Price Index [IPPI (pharma)], published by Statistics Canada, provides an index of manufacturers' prices for all pharmaceuticals, including both patented and non-patented drugs. In 1997, the IPPI (pharma) decreased by 0.7%. This is the first year that the prices of all drugs, as measured by Statistics Canada, went down. Figure 3 also shows that, with the exception of 1997, the IPPI (pharma) has increased every year since the creation of the PMPRB by an amount greater than the PMPI, and closer to changes in the CPI. In 1997, the IPPI was slightly below the PMPI.

As summarized in Figure 4, from 1988 to 1997, the IPPI (pharma) has increased on average by 2.2%, about 20% less than the average annual increase in the CPI of 2.8%. In contrast, prices for patented drugs have increased at a significantly lower rate over that period, by 1.0% per year on average.

Figure 4 also shows information on pharmaceutical price trends prior to the creation of the PMPRB in 1987. From 1982 to 1987, when there was no direct regulation of drug prices, price increases of all drugs, as measured by the IPPI (pharma), averaged 9.0% per year as compared with increases in the CPI of 5.6% per year. The decline in the rate of increase in the prices of all drugs relative to the CPI coincided with the introduction of federal price regulation of patented drugs, which represented between 40.7% and 52.3% of the sales of all drugs since 1988.

These trends imply that limiting price increases for all patented drugs has contributed to keeping the rate of increase in prices for all drugs lower than would otherwise have been the case.

Non-Patented Medicine Price Index (NPMPI)

There is no source of detailed price information for non-patented drugs that is comparable to the information for patented drugs. Therefore, it is necessary to estimate trends in prices of non-patented drugs from other sources. A nonpatented medicine price index (NPMPI) can be derived from the IPPI (pharma) and the PMPI. The IPPI (pharma) and the PMPI track the factory-gate prices of drug products; the IPPI (pharma) includes all drugs regardless of patent status, while the PMPI captures the complete basket of patented drug products. Given these two indices, it is reasonable to derive the NPMPI.⁹

Figure 5 shows how the NPMPI increased at a higher annual rate relative to the IPPI (pharma) and the PMPI from 1987 to 1996. In 1997, this trend was reversed. The change in non-patented drug prices was about one percentage point below the change in patented drug prices; non-patented drug prices fell by 1.3%, while patented drug prices fell by 0.1%.

Price Trends in Canada and the United States

The trends in drug prices in Canada can be compared with those in the United States. Figures 6 and 7 compare the annual changes in the pharmaceutical component of the U.S. Product Price Index [PPI (pharma)] to the annual changes in the IPPI (pharma) both before and after the introduction of federal price regulation in Canada. The U.S. PPI (pharma) measures price increases of all pharmaceuticals at the factory-gate. It is similar in construction to the Canadian IPPI.

Figure 6 shows the year-over-year changes in the U.S. PPI (pharma) and the Canadian IPPI (pharma) from 1982 and in the PMPI from 1988. Prior to the introduction of federal price regulation, the Canadian IPPI (pharma) increased in every year at a rate above the U.S. PPI (pharma). That trend reversed in 1987. In 1997, the U.S. PPI (pharma) increased by



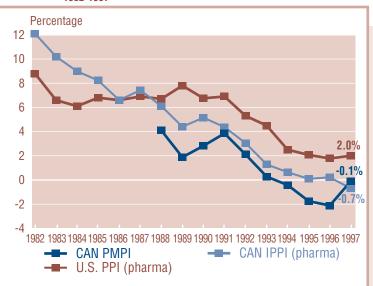
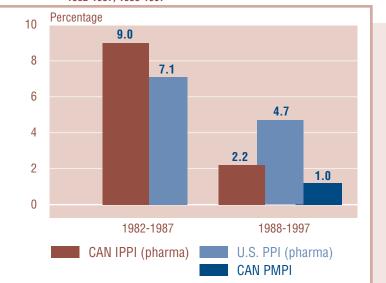




Figure 7 Summary of Price Trends Average Annual Percentage Changes 1982-1987; 1988-1997



Sources: Statistics Canada, the U.S. Bureau of Labor Statistics and the PMPRB

2.0% from 1996 compared to the decrease of 0.7% for the Canadian IPPI (pharma).

Figure 6 also shows that, since 1987, the PMPI grew every year at a much lower rate than the U.S. PPI (pharma).

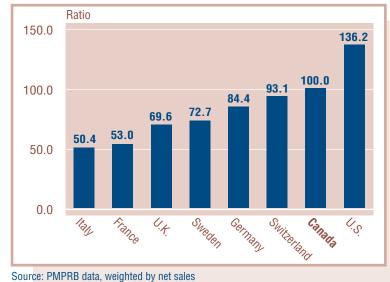
9 See the PMPRB's *The Impact of Federal Regulation of Patented Drug Prices*, February 1997.

Figure 7 shows these trends in summary form during the periods prior to and after the introduction of federal price regulation. From 1982 to 1987, Canadian drug prices, as measured by the IPPI (pharma), increased by almost 9% per year, more than the average annual rate of increase of 7.1% in the U.S. By contrast, the average annual rate of increase in



Figure 8 Ratio of Canadian Prices to Median International Prices 1987-1997

Figure 9 Average Foreign to Canadian Price Ratios All Patented Drug Products in 1987



10 France, Germany, Italy, Sweden, Switzerland, U.K., U.S.

the IPPI (pharma) declined to 2.2% from 1988 to 1997, well below the rate of 4.7% for the U.S. PPI (pharma). The average annual rate of increase of patented drug prices in Canada was only 1.0% from 1988 to 1997.

Relationship of Canadian Prices to Foreign Prices: Past and Present

The above price indices demonstrate how prices of drugs in Canada have changed over time. Another important role of price controls is to limit the introductory prices of new drug products in the Canadian market. One way of examining the combined effect of controls on introductory prices and price increases is to examine the trend in the relationship of prices in Canada to those in other countries. The next three figures show the relationship between Canadian prices of patented drugs and foreign prices over time.

Figure 8 shows the relationship between Canadian prices and median prices in the seven countries used for price comparison purposes, as listed in the *Patented Medicines Regulations*, over the period from 1987 to 1997.¹⁰ It shows that the average ratio of Canadian prices to median foreign prices has declined from 1.23 in 1987 to 0.89 in 1997. In other words, Canadian prices for patented drugs have declined from 23% above foreign prices to approximately 11% below. This represents about a 30% fall in the level of Canadian prices over this time period. This calculation is based on a revenue-weighted average of the ratio of the Canadian price to median international price for each patented drug product sold in that year.

Figures 9 and 10 show the relationship between Canadian prices and prices in each of the countries listed in the Regulations in 1987 and 1997. Figure 9 shows that in 1987, Canadian prices were, on average, below the U.S. but above the prices in all other countries. While Canadian prices were, on average, 36% below those in the U.S., they were almost twice as By 1997, Canada ranked third lowest, just below the U.K. high as prices in Italy and France. As shown in Figure 10, by 1997 Canadian prices relative to prices in each of the countries had changed dramatically. Canada ranked third lowest, just below the U.K. Prices

in the U.S., Switzerland, Germany and Sweden were higher, on average, while Italy and France were lower.

Increased Expenditures on Drugs

The above price indices show that since 1987, manufacturers' prices of all drugs, and patented drugs in particular, have increased at a more modest rate than before or have declined. Despite this moderation in price increases, total expenditures (including the manufacturer's price, markup and dispensing fees) on drugs have increased more rapidly. In 1996, according to the latest figures published by Health Canada, total expenditures on drugs have increased faster than other major components of health care, and reached 14.4% of total health expenditures (see Figure 11). Overall drug expenditures increased by 2.7% in 1996, 4.0% in 1995 and 3.6% in 1994 (see Figure 12).

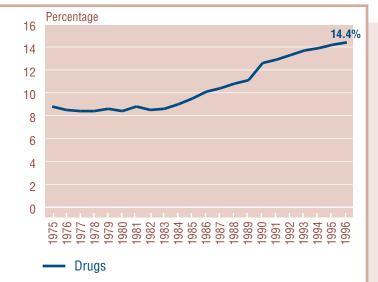
Several factors may account for this phenomenon of increasing total drug expenditures and moderating manufacturers' drug prices. The first issue which must be recognized in trying to explain any price-cost discrepancy is that total expenditures include wholesale or retail mark-ups as well as pharmacists' dispensing fees. Statistics Canada measures retail price changes of prescription drugs, including mark-ups and dispensing fees, with the Consumer Price Index for prescribed medicines, CPI (Rx). Since 1994, prices of prescription medicines at the retail level have been virtually constant as shown in Figure 13, even though total expenditures on drugs have been increasing.

Figure 10 Average Foreign to Canadian Price Ratios All Patented Drug Products in 1997



Source: PMPRB data, weighted by net sales





Source: Health Canada

The second issue which contributes to the confusion of rising drug costs in the face of moderating prices is the fact that the total drug "bill" (i.e. expenditures) is a function of a number of factors as outlined in Figure 14.¹¹ These factors often have an independent impact on total expenditures of drugs.

11 This figure is reproduced from the PMPRB's Discussion Paper, *Examining the Role, Functions and Methods of the Patented Medicine Prices Review Board*, November 1997.

Therefore, the control of one factor (e.g. drug prices at the factory or retail level) does not necessarily mean control of total expenditures. As an example, even if drug prices go down (as has more recently been the case), any change in the other factors affecting total expenditures may drive up total expenditures.



Figure 12 Year-over-Year Changes in Drug Expenditures 1976-1996



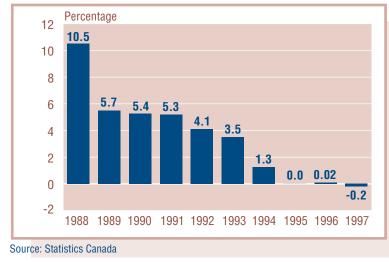


Figure 14 Factors Affecting Total Drug Expenditures

- changes in the total population
- changes in the demographics and health status of the population (i.e. towards those with increased medication needs)
- changes in the unit prices of drugs (both patented and non-patented)
- changes in retail and wholesale mark-ups and dispensing fees
- changes in the prescribing habits of physicians (i.e. from older, less expensive medications to newer, relatively more expensive medications [± improved therapeutic effect] to treat the same underlying diagnosis)
- changes in utilization of drugs on a per patient basis (i.e. more medications per patient per year)
- trends towards using drug therapy instead of other treatments (e.g. as alternatives to surgery in some cases)
- new diseases to be treated
- old diseases to be treated, where there existed no treatment before; old diseases better treated

Trends in Quantities of Sales of Patented Drug Products

Data available to the PMPRB allow it to measure changes in the quantities as well as the prices of patented drugs sold from year to year. This analysis reveals that the quantities of patented drugs sold have consistently increased at a much faster rate than prices. As shown in Figure 15, this trend has escalated in 1997.

In 1997, prices for patented medicines declined by 0.1%, on average, but the quantities sold increased by about 20%. In other words, while total sales of patented drugs increased by 22.6% in 1997, 88% of that increase is attributable to an increase in the amount of existing patented drugs consumed. About 12% of the increase in patented drug expenditures is attributable to the introduction of new patented drugs. Finally, the overall change in patented drug prices had virtually no impact in driving up the annual change in patented drug expenditures.

From 1988 to 1997, the average annual increase in quantities of patented drugs sold was approximately 10% as compared with an average annual increase of 1.0% in their prices.

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The index for the quantities of patented drugs sold may not be representative of total sales of all pharmaceuticals because patented drugs have represented between 41% and 52% of total sales since 1990. Among other things, this analysis does not take into account shifts in utilization between patented drugs and nonpatented drugs, nor does it account for changes in patent status. For example, drugs continue to be consumed even though their patents expire and their prices are no longer subject to the PMPRB's jurisdiction.

Sales by Major Therapeutic Group

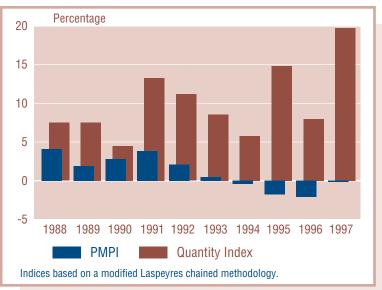
For price review purposes, the PMPRB classifies all drugs sold in Canada according to the Anatomical Therapeutic Chemical (ATC) classification system.

Table 2 breaks out the 981 patented drug products sold during 1997 according to major

therapeutic groups, showing the total expenditures, proportion of total expenditures and growth in expenditures from 1996. (The breakdown for patented medicines may differ from that for all drugs, including non-patented medicines.)

As shown in Table 2, the largest growth in patented drug expenditures came from the Blood and blood forming organs, Sensory organs, Antineoplastics and immunomodulating agents and Cardiovascular system groups. These groups experienced an increase in expenditures above 30% from the previous year. The average increase in expenditure over all therapeutic groups was 22.6%.

Figure 15 Year-over-Year Changes in the PMPI and in the Quantity Index 1988-1997



Source: PMPRB

Table 2 Patented Drug Expenditures by Major Therapeutic Group, 1997

| Major Therapeutic Group | 1997 Expenditures (\$Millions) | Proportion of total Expenditures in 1997 | Exper | Growth in Expenditures from 1996 | |
|--|--------------------------------------|---|-------|--|--|
| | | 11 1337 | \$M | % | |
| Alimentary tract and metabolism | 526.1 | 14.4 | 78 | 17.4 | |
| Blood and blood forming organs | 91.4 | 2.5 | 54 | 147.0 | |
| Cardiovascular system | 1,125.7 | 30.8 | 278 | 32.7 | |
| Dermatologicals | 89.0 | 2.4 | 7 | 8.5 | |
| Genito-urinary system and sex hormones | 90.4 | 2.5 | -33 | -33.1 | |
| Systemic hormonal preparations, excluding sex hormones | 28.1 | 0.8 | 25 | 0.9 | |
| General antiinfectives for systemic use | 518.9 | 14.2 | 81 | 18.5 | |
| Antineoplastics and immunomodulating agents | 213.8 | 5.9 | 60 | 38.8 | |
| Musculo-skeletal system | 107.4 | 2.9 | 18 | 20.7 | |
| Nervous system | 469.9 | 12.9 | 80 | 20.5 | |
| Antiparasitic products | 0.6 | 0.0 | 0 | 0.0 | |
| Respiratory system | 226.3 | 6.2 | 31 | 16.1 | |
| Sensory organs | 32.3 | 0.9 | 10 | 46.8 | |
| Various | 30.9 | 0.9 | 2 | 6.6 | |
| Veterinary products | 104.1 | 2.9 | 13 | 14.4 | |
| Totals | 3,655.1 | 100 | 683 | 22.6 | |

Rows and columns may not add to totals due to rounding. Source: PMPRB

Compliance and Excessive Price Guidelines

Under the *Patented Medicines Regulations* (*Regulations*), patentees are required to report information on the sales and prices of new patented medicines and to continue to file detailed information on sales and prices of each patented drug for the first and last sixmonth period of each year. The PMPRB reviews this pricing information on an ongoing basis to ensure that the prices charged by patentees comply with the Guidelines established by the Board. The Guidelines are published in the PMPRB's *Compendium of Guidelines, Policies and Procedures*.

The Guidelines are based on the price determination factors in section 85 of the *Patent Act*. In summary, the Guidelines provide that:

- prices for most new drugs are limited such that the cost of therapy for the new drug is in the range of the cost of therapy for existing drugs used to treat the same disease in Canada;
- prices of breakthrough drugs and those which bring a substantial improvement are limited to the median of the prices charged for those drugs in other industrialized countries listed in the *Regulations* (France, Germany, Italy, Sweden, Switzerland, U.K. and U.S.);
- price increases for existing medicines are limited to changes in the Consumer Price Index (CPI); and
- the price of a patented drug in Canada may, at no time, exceed the range of the prices for the same drug in foreign countries.

New Patented Drug Products in 1997

For purposes of the review of prices by the PMPRB, new patented drug products in 1997 include those introduced on the market in Canada or those previously marketed but first patented between December 1, 1996 and November 30, 1997. Because of the timing of the filing requirements under the *Regulations*

and the manner of calculating benchmark prices, drug products introduced or patented in December are considered to be new patented products in the following year.

There were 98 new patented drug products (DINs) representing 61 medicines sold in 1997. This is slightly higher than the average number of patented drug products introduced annually in Canada over the last several years. All but five of the new patented DINs in 1997 are for human use. Drug products for veterinary use have always been a relatively small percentage of the total number of new patented drug products.

The Board's Guidelines establish three categories of new patented drug products for purposes of conducting introductory price reviews.¹²

- Category 1 a new DIN of an existing or comparable dosage form of an existing medicine, usually a new strength of an existing drug (line extension).
- Category 2 the first drug product to treat effectively a particular illness or which provides a substantial improvement over existing drug products, often referred to as "breakthrough" or "substantial improvement."
- Category 3 a new drug or new dosage form of an existing medicine that provides moderate, little or no improvement over existing medicines.

Figure 16 provides a breakdown by category of the new patented DINs for human use between 1991 and 1997. The proportion of DINs in each category has been relatively constant over the years. The number of category 2 drugs has usually been less than 10%. The number of category 2 drugs in 1997, 13, appears to be higher than usual and warrants closer inspection. These 13 DINs represent 6 medicines; 7 of these DINs (3 medicines) were introduced on the market

12 For complete definitions of the categories, refer to the Compendium of Guidelines, Policies and Procedures, Chapter 3, section 3.

There were 28 new active substances among the new patented DINs for human use in 1997. in Canada well before 1997 but only came under the Board's jurisdiction when they received their first patent(s) in 1997. For example, Eprex has been sold in Canada since 1990 while 3TC has been on the market since 1995. Table 12, on page 39,

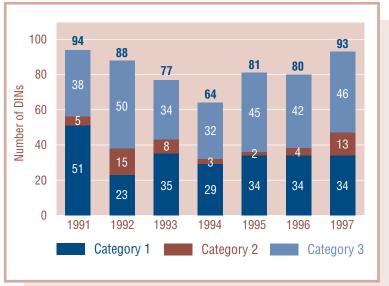
lists the drug products included in each category in 1997.

New Active Substances in 1997

Among the new patented DINs for human use sold in Canada in data-year 1997, there were 28 new active substances (NASs) compared to 21 NASs in 1996 and 20 NASs in 1995. Of the 28 NASs, four were introduced on the market in Canada prior to 1997 but only received their first patent(s) in 1997. A listing of the NASs and their assigned categories appears in Table 3. A NAS may represent more than one drug product if it is sold in more than one strength or dosage form. The 28 NASs listed were marketed as 51 presentations (DINs) in 1997. Manufacturers filed category 2 submissions for 10 of the patented NASs (20 DINs) listed in Table 3; five of the 10 were classified by the Human Drug Advisory Panel (HDAP) as category 2 new medicines following review of the patentee's submission. One other new medicine, Indinavir, (trade name: Crixivan) was classified as a category 2 drug by the HDAP without a submission from the patentee. All three of the protease inhibitor drug products approved and introduced in Canada for HIV therapy in 1996 were considered to be category 2 new drug products (i.e. Norvir, Crixivan and Invirase).

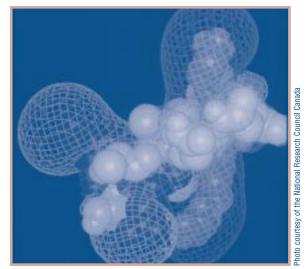
The PMPRB's list of patented NASs in any year may differ from the list of NASs approved by the Health Protection Branch (HPB). In some circumstances, a patented drug may be sold under the Special Access Program (SAP) before it receives a Notice of

Figure 16 Categorization of New Patented Drug Products (Human)



Source: PMPRB

Compliance (NOC); or it may be approved but not be introduced until some time later. For example, 18 of the 28 patented NASs were granted a NOC by HPB in 1997; eight in 1996; one in 1995 and one in 1990. The HPB list of New Active Substances, which reports on the NASs that received a NOC in 1997, lists 42 NASs; not all were introduced to the market in that year. In addition, not all NASs approved by HPB are subject to the PMPRB's jurisdiction. Twenty-four of the 42 NASs approved by HPB in 1997 are patented medicines and four more are known to the PMPRB as having patents pending.



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Price Review of New Patented Drugs in 1997

All 98 of the 1997 new patented drug products (DINs) were categorized as listed in Table 12. The majority (94%) were found to be priced within the Guidelines.

Of the 98 products, six are under review pending the review of additional information and two have been found to be outside the Guidelines and are the subject of an investigation.

Table 3New Patented Medicines in 1997
New Active Substances (Human)

| Chemical Name | Brand Name | Company | Category | ATC CLASS |
|-------------------------------------|------------|---------------------------|----------|-----------|
| New Active Substances Introduced in | 1997 (NAS) | | | |
| ACELLULAR PERTUSSIS VACCINE | ACEL-P | WYETH-AYERST CANADA | 3 | J07AJ |
| ANISTREPLASE | EMINASE | ROBERTS PHARMACEUTICAL | 3 | B01AD |
| ATORVASTATIN | LIPITOR | WARNER-LAMBERT CANADA | 3 | C10AA |
| COAGULATION FACTOR IX | BENEFIX | GENETICS INSTITUTE | 2 | B02BD |
| DAPT VACCINE | TRIPACEL | CONNAUGHT LABORATORIES | 3 | J07AJ |
| DAPT-IPV VACCINE | QUADRACEL | CONNAUGHT LABORATORIES | 3 | J07AJ |
| DAPT-IPV-HIB VACCINE | PENTACEL | CONNAUGHT LABORATORIES | 3 | J07AG |
| DESFLURANE | SUPRANE | ZENECA PHARMA INC. | 3 | N01AB |
| DOLASETRON MESYLATE | ANZEMET | HOECHST MARION ROUSSEL | 3 | A04AA |
| DONEPEZIL HYDROCHLORIDE | ARICEPT | PFIZER CANADA INC. | 2 | N07AA |
| EPOPROSTENOL | FLOLAN | GLAXO WELLCOME INC. | 3 | B01AC |
| FEXOFENADINE HYDROCHLORIDE | ALLEGRA | HOECHST MARION ROUSSEL | 3 | R06AX |
| FORMOTEROL FUMARATE | FORADIL | SANDOZ CANADA | 3 | R03AC |
| IRINOTECAN HYDROCHLORIDE | CAMPTOSAR | PHARMACIA & UPJOHN INC. | 2 | L01XX |
| LATANOPROST | XALATAN | PHARMACIA & UPJOHN INC. | 3 | S01EX |
| LETROZOLE | FEMARA | SANDOZ CANADA | 3 | L02BG |
| PANTOPRAZOLE SODIUM | PANTOLOC | SOLVAY KINGSWOOD INC. | 3 | A02BC |
| ROPINIROLE HYDROCHLORIDE | REQUIP | SMITHKLINE BEECHAM PHARMA | 3 | N04BC |
| ROPIVACAINE HYDROCHLORIDE | NAROPIN | ASTRA PHARMA INC. | 3 | N01BB |
| TAZAROTENE | TAZORAC | ALLERGAN INC. | 3 | D05AX |
| TECHNETIUM TC-99M TETROFOSMIN | MYOVIEW | AMERSHAM CANADA LTD. | 3 | V09AG |
| TOLCAPONE | TASMAR | HOFFMAN-LA ROCHE LTD. | 3 | N04BX |
| TOPIRAMATE | TOPAMAX | JANSSEN PHARMACEUTICAL | 3 | N03AX |
| TOPOTECAN HYDROCHLORIDE | HYCAMTIN | SMITHKLINE BEECHAM | 3 | L01XX |
| First Patent Granted in 1997 (FPG) | | | | |
| EPOETIN ALFA | EPREX | JANSSEN PHARMACEUTICAL | 2 | B03XA |
| INDINAVIR SULFATE | CRIXIVAN | MERCK FROSST CANADA | 2 | J05AE |
| LAMIVUDINE | 3TC | GLAXO WELLCOME INC. | 3 | J05AB |
| SAQUINAVIR MESYLATE | INVIRASE | HOFFMAN-LA ROCHE LTD. | 2 | J05AE |

Photo courtesy of Health Canada

Patent Pending

When a medicine, subject to a pending patent, is being sold, the Board will, when the patent is issued, review the price at which the medicine was sold during the pre-grant infringement period. This is normally from the date the patent is first laid open for inspection. The Board encourages patentees to seek advisory assistance during this patent pending period to ensure that the medicine is not being sold at an excessive price before the patent issues.

On April 29, 1998, the Board accepted a Voluntary Compliance Undertaking (VCU) in respect of Humalog, a medicine for which the patent is pending. Details of the circumstances and the VCU can be found at page 26.

Price Review of Existing Patented Drug Products in 1997

For the purposes of this report, existing medicines include all patented drug products that were on the market before 1997. The PMPRB's Guidelines limit the price changes for existing patented drugs to changes in the Consumer Price Index (CPI). In addition, the price of a patented drug product cannot exceed the highest price of the same drug product in the countries listed in the *Regulations*.¹³

A total of 893 existing patented drug products were sold during 1997. The prices of 880 drug products (over 98%) were reviewed and found to be within the Guidelines. A drug product is considered to be within the Guidelines if its price does not exceed the maximum allowable price and does not meet the criteria established for the commencement of an investigation.¹⁴



At the time of this report, the prices of 13 drug products (nine medicines) are under investigation. Of these 13 drug products, six involve similar issues and are being reviewed simultaneously. In the seven remaining cases, the manufacturers have been notified and the Board is awaiting their responses.

Update of the Ninth Annual Report

In the 1996 Annual Report, the PMPRB had reported that eight drug products (new and existing) were under review. Since then, six cases have been resolved through receipt of additional evidence which showed that the prices had been within the Guidelines. The remaining two products continue to be under review and are included in the 13 existing drug products under investigation mentioned previously.

¹³ France, Germany, Italy, Sweden, Switzerland, U.K. and U.S.

¹⁴ For a full explanation of the criteria for commencing an investigation please refer to Schedule 5 of the PMPRB's *Compendium of Guidelines, Policies and Procedures.*

Enforcement Activities

Voluntary Compliance Undertakings

Under the Compliance and Enforcement Policy, patentees are given an opportunity to make a Voluntary Compliance Undertaking (VCU) when Board staff conclude, following an investigation, that a price appears to have exceeded the Guidelines. Approval of a VCU by the Chairperson or Board is an alternative to the commencement of formal proceedings through the issuance of a Notice of Hearing.

The Board did not approve any VCUs in 1997, but a major investigation during 1997 resulted in a VCU in 1998, discussed below.

HUMALOG (insulin lispro)

Eli Lilly Canada Inc.

On April 29, 1998, the Chairperson of the Board approved a VCU by Eli Lilly Canada Inc. (Lilly) to reduce the manufacturer's price of Humalog by 23%, from \$30 to \$23.

On March 14, 1998, the Board had issued a public notice (Notice) in the Canada Gazette that it had received a VCU from Lilly in respect of the price of the medicine Humalog. The Notice observed that the Staff of the Board recommended the acceptance of the VCU by the Board and provided the provincial and territorial Ministers of Health and other interested persons with an opportunity to make submissions on the appropriateness of the VCU and its terms and provisions.

Background

Humalog is a rapid-acting analog of human insulin that has been approved by Health Canada for the treatment of patients with diabetes mellitus who require insulin for the maintenance of glucose homeostasis. An application for a Canadian patent pertaining to Humalog, filed in 1990, is still pending but Lilly expects that a patent will be granted by the end of 1998. In brief, the VCU provided that Lilly would lower the manufacturer's list price for Humalog by approximately 23%, such that the relationship between the prices of Humalog and regular insulin in Canada would reflect the relationship between the prices of those products in other countries.

In addition, the VCU proposed to offset all excess revenues which had been received by Lilly as a result of charging the higher price through a mechanism intended to provide Humalog at no charge to patients who had previously been treated with it.

Submissions

The Board received submissions from or on behalf of Ministers of Health of six provinces: Alberta, British Columbia, Nova Scotia, Prince Edward Island, Québec and Saskatchewan. Submissions were also received from the Canadian Diabetes Association and Novo Nordisk Canada Inc.

Most of the parties agreed with the proposed price reduction and with the methodology for determining the maximum non-excessive (MNE) price.

However, all of the submissions objected to the proposed method of achieving the offset of excess revenues, being the proposed distribution of Humalog at no charge to patients who had used the medicine in the past.

The Amended VCU

Following further discussions, on April 24, 1998, Lilly submitted an amended VCU that changed the proposed method of offsetting excess revenues. Pursuant to the amended VCU, Lilly undertook to make a payment to the Government of Canada to offset excess revenues of \$666,824. To the extent that any excess revenues may have been received in 1998 prior to the price reduction, they will be offset through further price reductions to ensure that the average price for Humalog in 1998 does not exceed the MNE price of \$22.1072.

Decision

Having considered all of the submissions from interested parties and the amendments to the VCU, the Chairperson accepted the amended VCU on April 29, 1998.

While the Board was sensitive to the submissions of some parties who argued for a larger reduction in the price of Humalog, it was considered on balance that the amended VCU is consistent with the provisions of the *Patent Act* and the Board's Compliance and Enforcement Policy. The amended VCU establishes a reasonable basis for determining the MNE price for Humalog consistent with the factors set out in subsection 85(1) of the *Patent Act*. Furthermore, the amended VCU will result in an immediate price reduction for Humalog, together with the offset of past excess revenues in a manner consistent with the submissions of interested parties.

The acceptance of the amended VCU is based on the particular circumstances of the medicine Humalog and its context in the international market for analogs of human insulin. This exceptional case does not warrant a review or amendment of the Excessive Price Guidelines, which remain applicable to medicines subject to the Board's jurisdiction.

Use of the Funds Paid to the Government of Canada

Under the *Patent Act*, the Board has no authority to order that funds paid to the Government of Canada to offset excess revenues be used for certain purposes. The Board invited comments relating to Humalog prior to deciding whether to accept a proposed VCU to lower the price of this medicine.

Pursuant to section 103 of the *Act*, the Ministers of Health in Canada have agreed in principle that funds collected as a result of Board Orders and VCUs be distributed to the provinces on a per capita basis. They further agreed that such funds should be used for pharmaceutical objectives.

In this case, the Canadian Diabetes Association recommended that any payment to governments should be directed towards a national diabetes related initiative. The Board brought that recommendation to the attention of governments.

Copies of the VCU, Public Notice and Reasons are available from the Secretary to the Board or on the Board's web site at http://www.pmprb-cepmb.gc.ca.

Analysis of Research-and-Development (R&D) Expenditures

The ratio of R&D expenditures to sales revenues for the patented pharmaceutical industry was 11.5% in 1997. With the adoption of the 1987 amendments to the *Patent Act*, the Pharmaceutical Manufacturers Association of Canada (PMAC) made a public commitment that the brand name pharmaceutical industry would increase its annual R&D expenditures as a percentage of sales to 10% by 1996.

Under the *Patent Act*, the PMPRB monitors and reports on R&D spending by patentees but it has no regulatory authority to influence the type of research or amount of R&D spending by patentees. The *Act* requires the PMPRB to report on how much each patentee spends annually on pharmaceutical R&D in relation to revenues and on how much the patented industry as a whole spends on R&D. For individual patentees, this calculation includes all revenues from Canadian sales of medicines, including revenues from licensing agreements.

Data Sources

Companies that reported sales of patented medicines in 1997 were also required to file R&D data for that calendar year as per the Patented Medicines Regulations (Regulations). Only companies with active Canadian patents pertaining to a medicine sold in Canada are required by the *Patent Act* to report on R&D expenditures. As new patents are granted and others expire, the group of companies required to file R&D data may change from year to year.

For 1997, 75 companies, including 35 PMAC members, filed reports on R&D. The data from these firms are the basis of this report. The total R&D expenditures for the 35 PMAC members totalled \$657.4 million in 1997, or 90.7% of the total R&D expenditures for the patented pharmaceutical industry as a whole; and their sales revenues totalled \$5,098.2 million, accounting for 81.1% of the total sales revenues.

R&D Expenditures

Pursuant to the *Regulations*, patentees must report those R&D expenditures that would have been eligible for an Investment Tax Credit for scientific research and experimental development under the provisions of the *Income Tax Act* in effect on December 1, 1987. Market research, sales promotions, quality control or routine testing of materials, devices or products and routine data collection are

Table 4 Total R&D Expenditures* and R&D-to-Sales Ratios, 1988 - 1997

| Year | Companies | Total R&D | Change from | Total Sales | Change from | R&D-to-S | ales Ratio |
|------|-----------|------------------------|----------------------|-------------------|----------------------|----------------------|-----------------------|
| | Reporting | Expenditures* (\$M) | Previous Year (%) | Revenues (\$M) | Previous Year (%) | All Patentees (%) | PMAC Patentees (%) |
| 1997 | 75 | 725.1 | 9.0 | 6,288.4 | 7.4 | 11.5 | 12.9 |
| 1996 | 72 | 665.3 | 6.4 | 5,857.4 | 9.9 | 11.4 | 12.3 |
| 1995 | 71 | 625.5 | 11.5 | 5,330.2 | 7.5 | 11.7 | 12.5 |
| 1994 | 73 | 561.1 | 11.4 | 4,957.4 | 4.4 | 11.3 | 11.6 |
| 1993 | 70 | 503.5 | 22.1 | 4,747.6 | 14.0 | 10.6 | 10.7 |
| 1992 | 71 | 412.4 | 9.6 | 4,164.4 | 6.9 | 9.9 | 9.8 |
| 1991 | 65 | 376.4 | 23.2 | 3,894.8 | 18.1 | 9.7 | 9.6 |
| 1990 | 65 | 305.5 | 24.8 | 3,298.8 | 11.0 | 9.3 | 9.2 |
| 1989 | 66 | 244.8 | 47.4 | 2,973.0 | 9.4 | 8.2 | 8.1 |
| 1988 | 66 | 165.7 | - | 2,718.0 | - | 6.1 | 6.5 |

Source: PMPRB

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PMPRB 1997 among the expenditures that are not eligible for an Investment Tax Credit and therefore are *not* included in the patentees' filings. Total R&D expenditures include current expenditures, capital equipment costs and allowable depreciation expenses.

Revenues from Sales

The 75 patentees reported total revenues of \$6.3 billion from Canadian sales of patented and non-patented drugs in 1997, up 7.4% over 1996. Of total sales revenues, less than 1% was generated by licensing agreements.

R&D-to-Sales Ratios

The ratio of R&D expenditures to sales revenues for the patented pharmaceutical industry was 11.5% in 1997, up from 11.4% in 1996. The ratio for the 35 companies that were members of the PMAC was 12.9% in 1997, up from 12.3% in 1996.

As shown in Table 5, of the 75 reporting companies, 18 companies (24.0%) reported having performed no R&D in 1997 compared to 11 in 1996. Sales revenues for companies with no R&D totalled \$335.2 million in 1997 accounting for 5.3% of total sales revenues for the patented pharmaceutical companies. The category of R&D-to-sales ratios between 0% and 10% experienced both a decrease in the number of companies and in the total sales revenues in 1997. There were 32 companies with total sales of \$1,879.8 million in 1997 as compared to 34 companies with total sales of \$2,000.4 million in 1996. The sales revenues



for companies with ratios of 10% or more experienced an increase of 8.8% in the total sales from \$3,744.6 million in 1996 to \$4,073.4 million in 1997 while the number of companies included in this category fell by 7.4%.

As shown in Table 4, patentees reported total R&D expenditures of \$725.1 million in 1997, an increase of 9.0% over 1996. Current expenditures accounted for \$679.2 million or 93.7% of total R&D expenditures. Capital equipment costs and allowable depreciation expenses amounted to 4.9% and 1.4% respectively.

Table 6 shows how current expenditures on R&D in 1997 were allocated among basic, applied, and other qualifying R&D. Total current expenditures on R&D rose by 7.8% in 1997.

| Range of R&D-to-Sales | | 1997 | | | 1996 | | % Change in Number of | % Change in Total Sales |
|--------------------------|-------------------------------------|----------------------------------|------|-------------------------------------|----------------------------------|------|-------------------------------------|----------------------------|
| Ratio | Number of Reporting Companies | Total Sales Revenues (\$M) | % | Number of Reporting Companies | Total Sales Revenues (\$M) | % | Reporting Companies 1997/1996 | Revenues 1997/1996 |
| 0% | 18 | 335.2 | 5.3 | 11 | 112.4 | 1.9 | 63.6 | 198.2 |
| 0% to 10% | 32 | 1,879.8 | 29.9 | 34 | 2,000.4 | 34.2 | -5.9 | -6.0 |
| >10% | 25 | 4,073.4 | 64.8 | 27 | 3,744.6 | 63.9 | -7.4 | 8.8 |
| Total | 75 | 6,288.4 | 100 | 72 | 5,857.4 | 100 | 4.2 | 7.4 |

Table 5 Range of R&D-to-Sales Ratios by Number of Reporting Companies and Total Sales Revenues

Source: PMPRB

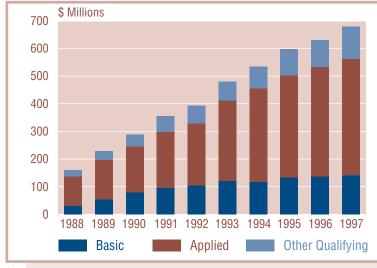
29

| Type of Research | 19 | 997 | 19 | 96 | Change in |
|------------------|-------|------|-------|------|--------------------------|
| | \$M | % | \$M | % | Expenditures 1997/1996 % |
| Basic | 140.4 | 20.7 | 136.6 | 21.7 | 2.8 |
| Applied | 421.3 | 62.0 | 396.4 | 62.9 | 6.3 |
| Other Qualifying | 117.5 | 17.3 | 97.1 | 15.4 | 21.0 |
| Total | 679.2 | 100 | 630.1 | 100 | 7.8 |

Table 6 Current R&D Expenditures* by Type of Research, 1996 and 1997

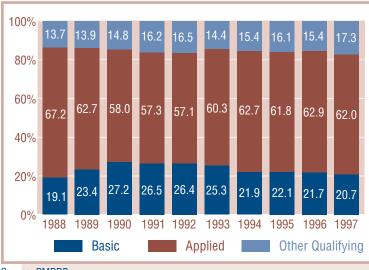
Source: PMPRB





Source: PMPRB

Figure 18 Share of Current R&D Expenditures by Type of Research 1988-1997



Source: PMPRB

Spending on basic research was \$140.4 million or 20.7% of the total. Basic research is defined as work that advances scientific knowledge without a specific application in view. The lion's share of R&D spending continued to be on applied research, \$421.3 million or 62.0% of the total. Applied research is directed towards some practical application, comprising the

manufacturing process, pre-clinical trials and clinical trials. Clinical trials accounted for 79.2% of total applied research expenditures, \$333.7 million, while manufacturing process accounted for \$51.0 million, or 12.1% of the total, and pre-clinical trials accounted for \$36.6 million or 8.7% of the total. Other qualifying research expenditures are for drug regulation submissions, bioavailability studies and Phase IV clinical trials.

Figure 17 shows current expenditures on R&D by type of research from 1988 to 1997 and Figure 18 shows their shares of expenditures during those years.

While spending on basic research went up by 2.8% in 1997, its share of total R&D continued to decline from 21.7% in 1996 to 20.7% in 1997.

Pharmaceutical patentees may report their expenditures on research they conduct themselves (intra-mural) and research performed by others, including universities and hospitals and other manufacturers (extramural). Table 7 presents the current R&D expenditures by R&D performer and identifies the intra-mural and extra-mural expenditures. Most R&D was carried out by patentees. In 1997, 55.3% of R&D expenditures was directed to R&D performed by the patentee, compared with 55.6% in 1996. Expenditures on R&D performed by universities and hospitals decreased by 2.8% to \$142.7 million from 1996. Expenditures on R&D performed by other companies on behalf of patentees increased, from \$70.3 million in 1996 to \$79.4 million in 1997, a 12.9% increase. The largest increase occurred in the category "others."

The category "others" includes individuals, organizations such as private clinics, and federal and provincial governments. This category incurred an increase of 30.3%.

In 1997, as in previous years, most of the R&D expenditures of pharmaceutical patentees were funded internally. Table 8 shows that in 1997, more than 98% of all patentees' R&D was funded by internal funds and funds provided by associated companies. The share of funding by governments increased to 1.0% and the share of others slightly decreased to 0.8%.

In 1997, R&D spending increased in all regions. The largest increases occurred in the Atlantic and Western provinces, by 45.5% and 11.3% respectively. There was no significant change in the regional distribution of R&D spending in 1997. As shown in Table 9, almost 90% of total expenditures continued to be made in Ontario and Québec. Table 11, on page 34, lists the current R&D expenditures by province and by R&D performer for 1997.

Table 7 Current R&D Expenditures* by R&D Performer, 1996 and 1997

| R&D Performer | 1 | 997 | | 1996 | | |
|-------------------------------|-------|------|-------|------|----------------|--|
| | \$M | % | \$M | % | 1997/1996 % | |
| Intra-mural | | | | | | |
| Patentees | 375.8 | 55.3 | 350.6 | 55.6 | 7.2 | |
| Extra-mural | | | | | | |
| Universities and Hospitals | 142.7 | 21.0 | 146.8 | 23.3 | -2.8 | |
| Other Companies | 79.4 | 11.7 | 70.3 | 11.2 | 12.9 | |
| Others | 81.3 | 12.0 | 62.4 | 9.9 | 30.3 | |
| Total | 679.2 | 100 | 630.1 | 100 | 7.8 | |

Source: PMPRB

Table 8 Total R&D Expenditures* by Source of Funds, 1996 and 1997

| Source of Funds | 1 | 997 | 1 | 1996 | | |
|-----------------------------------|-------|------|-------|------|----------------|--|
| | \$M | % | \$M | % | 1997/1996 % | |
| Company Funds | 712.1 | 98.2 | 654.4 | 98.4 | 8.8 | |
| Federal/Provincial Governments | 7.3 | 1.0 | 4.7 | 0.7 | 55.3 | |
| Others | 5.7 | 0.8 | 6.2 | 0.9 | -8.1 | |
| Total | 725.1 | 100 | 665.3 | 100 | 9.0 | |

* Total expenditures include capital equipment and allowable depreciation.

Source: PMPRB

Table 9 Current R&D Expenditures* by Location, 1996 and 1997

| Location of R&D | 19 | 97 | | 1996 | | |
|--------------------|---------------------|------|-------|------|----------------|--|
| | \$M | % | \$M | % | 1997/1996 % | |
| Atlantic Provinces | 16.0 | 2.4 | 11.0 | 1.8 | 45.5 | |
| Québec | 290.6 | 42.8 | 264.7 | 42.0 | 9.8 | |
| Ontario | 296.6 | 43.7 | 286.1 | 45.4 | 3.7 | |
| Western Provinces | 76.0 | 11.2 | 68.3 | 10.8 | 11.3 | |
| Yukon and N.W.T. | 0.17 | 0.0 | 0.0 | 0.0 | n/a | |
| Total | 679.41 ¹ | 100 | 630.1 | 100 | 7.8 | |

* Current expenditures exclude capital equipment and depreciation expenditures.

1 Table may not balance due to roundings. For more details on provincial breakdown, refer to Table 11.

Source: PMPRB

| Company | 1997 | R&D-to-Sales Ratio (%) | 1996 |
|--|--------------------|------------------------|--------------------|
| 3M Pharmaceuticals, 3M Canada Inc. | 2.3 | | 2.0 |
| Abbott Laboratories, Limited | 1.7 | | 1.9 |
| Alcon Canada Inc. | 0.0 | | 0.02 |
| Allergan Inc. | 7.2 | | 15.6 |
| Alpha Therapeutic Corporation | 0.0 | | 0.0 |
| AltiMed Pharmaceutical Company | 0.0 | | 0.0 |
| Amersham Canada Limited | 2.9 | | 1.2 |
| Amgen Canada Inc. | 79.9 ² | | 99.9 ² |
| Astra Pharma Inc. | 12.1 | | 13.9 |
| Ayerst Veterinary Laboratories | 0.0 | | 0.8 |
| Baxter Corporation | 0.2 | | 0.2 |
| Bayer Inc. | 8.3 | | 8.7 |
| Bayer Inc., Agriculture Division | 4.5 | | 4.7 |
| Berlex Canada Inc. | 10.3 | | 8.0 |
| Block Drug Company (Canada) Ltd. | 0.0 | | 0.0 |
| Boehringer Ingelheim (Canada) Ltd. | 46.4 | | 37.3 |
| Boehringer Mannheim Canada Ltd. | 5.7 | | 8.0 |
| Bracco Diagnostics Canada Inc. | 0.0 | | 0.0 |
| Bristol-Myers Squibb Pharmaceutical Group | 10.5 | | 11.9 |
| Canderm Pharma Inc. | 3.0 | | 2.5 |
| Cangene Corporation | 183.6 ² | | 271.5 ² |
| Carter-Horner Inc. | 6.2 | | 6.0 |
| CIBA Vision Canada Inc. | 1.8 | | 3 |
| Colgate Oral Pharmaceuticals (not a patentee in 1996) | 0.0 | | |
| Dahi Animal Health Inc. | 0.0 | | 0.0 |
| Dermik Laboratories Canada Inc. | 0.0 | | 0.0 |
| Draxis Health Inc. | 5.3 | | 10.9 |
| Du Pont Merck Pharma Inc. | 8.3 | | 9.4 |
| Eli Lilly Canada Inc. (includes Elanco Animal Health Division) | 15.2 | | 12.2 |
| Fabrigen Inc. | 0.0 | | 0.0 |
| Ferring Inc. | 3.9 | | 2.5 |
| Fournier Pharma Inc. | 10.3 | | 11.1 |
| Fujisawa Canada Inc. | 14.0 | | 17.9 |
| Galderma Canada Inc. | 0.0 | | 0.0 |
| GenDerm Canada Inc. | 1.2 | | 0.9 |
| Genetics Institute Inc. (not a patentee in 1996) | 0.0 | | |
| Glaxo Wellcome Inc. | 12.9 | | 11.5 |
| Hoechst Marion Roussel Canada Inc. | 20.9 | | 15.1 |
| Hoffmann-La Roche Limited | 9.4 | | 10.2 |
| ICN Canada Limited | 4.8 | | 7.0 |
| Immuno (Canada) Ltd. | 0.0 | | 0.8 |
| Janssen-Ortho Inc. | 13.6 | | 10.6 |
| Johnson & Johnson Merck Consumer Pharmaceuticals of Canada | 0.0 | | 0.05 continue |

Table 10Ratios of R&D Expenditures to Sales Revenues
by Reporting Patentee 1, 1996 and 1997

Table 10 continued

| Company | | R&D-to-Sales Ratio (%) | |
|--|-------------------|------------------------|-------------------|
| | 1997 | | 1996 |
| Leo Laboratories Canada Ltd. | 8.5 | | 8.7 |
| Ligand Pharmaceuticals (Canada) | 48.0 ² | | 98.8 ² |
| Mallinckrodt Medical Inc. | 1.1 | | 1.3 |
| Mallinckrodt Veterinary Inc. | 3.5 | | 1.3 |
| McNeil Consumer Products Company | 1.9 | | 2.5 |
| Merck Frosst Canada Inc. | 13.1 | | 14.4 |
| Nexstar Pharmaceuticals Inc. | 0.0 | | 0.0 |
| Novartis Animal Health Canada Inc. | 0.3 | | 4 |
| Novartis Consumer Health Canada Inc. | 2.9 | | 5 |
| Novartis Pharmaceuticals Canada Inc. | 10.7 | | 6 |
| Novo Nordisk Canada Inc. | 0.5 | | 1.4 |
| Nycomed (Canada) Inc. (not a patentee in 1996) | 0.0 | | |
| Organon Canada Ltd. | 2.0 | | 2.2 |
| Pasteur Mérieux Connaught Canada ⁷ | 70.0 | | 68.8 |
| Pfizer Canada Inc. | 12.5 | | 11.3 |
| Pharmacia & Upjohn Inc. | 11.0 | | 11.6 |
| Procter & Gamble Pharmaceuticals Canada, Inc. | 10.2 | | 14.1 |
| Purdue Frederick | 6.4 | | 7.9 |
| Rhône-Poulenc Rorer Canada Inc. | 11.0 | | 10.1 |
| Roberts Pharmaceutical Canada Inc. (not a patentee in 1996 |) 0.0 | | |
| Sanofi Winthrop | 13.6 | | 11.8 |
| Schering Canada Inc. | 13.2 | | 7.1 |
| Searle Canada Inc. | 11.4 | | 6.2 |
| SmithKline Beecham Pharma Inc. | 10.0 | | 9.0 |
| Solvay Pharma Inc. | 9.3 | | 14.2 |
| The Liposome Company Inc. | 5.0 | | 0.0 |
| Warner-Lambert Canada Inc. (Parke-Davis) | 12.0 ⁸ | | 17.6 |
| Warner-Wellcome Consumer Health Products | 2.0 | | 2.4 |
| Westwood-Squibb | 0.7 | | 0.5 |
| Wyeth-Ayerst Canada Inc. | 13.1 | | 12.6 |
| Yamanouchi Pharmaceutical Co., Ltd. | 0.0 | | 0.0 |
| Zeneca Pharma Inc. | 7.6 | | 7.6 |

1 The revenues from royalties are included in calculating each company's ratio, but are deducted, when appropriate, for the industry-wide aggregation to avoid double-counting. Federal and provincial government grants have been netted from the expenditures used to calculate the individual R&D-to-sales ratios but are included in the aggregate statistics. Differences between the list of firms filing data on prices and those filing R&D data are due to differences in reporting practices between patentees and their affiliates or licencees.

2 These ratios have been verified with the firm and are due to the fact that funding for R&D expenditures was provided by associated companies.

3 Formerly under CIBA-Geigy Canada Ltd. The R&D-to-sales ratio for CIBA-Geigy Canada Ltd. was 7.6% in 1996.

4 Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. merged in 1997. The R&D-to-sales ratio for Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. were, respectively, 12.0% and 7.6% in 1996. Novartis Animal Health Canada Inc. includes veterinary products that were previously sold by CIBA-Geigy Canada Ltd.

5 Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. merged in 1997. The R&D-to-sales ratio for Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. were, respectively, 12.0% and 7.6% in 1996. Novartis Consumer Health Canada Inc. includes over counter products that were previously sold by CIBA-Geigy Canada Ltd. and Sandoz Canada Inc.

6 Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. merged in 1997. The R&D-to-sales ratio for Sandoz Canada Inc. and CIBA-Geigy Canada Ltd. were, respectively, 12.0% and 7.6% in 1996. Novartis Pharmaceuticals Canada Inc. includes prescription drug products for human use previously sold by CIBA-Geigy Canada Ltd. and Sandoz Canada Inc.

7 Formerly known as Connaught Laboratories Limited.

8 Warner-Lambert Canada Inc. (Parke-Davis) purchased Jouvenial Inc. in 1997. The R&D-to-sales ratio for Jouvenial Inc. was 2.8% in 1996.

Source: PMPRB

Table 11 Current R&D Expenditures by Province and by R&D Performer, 1997

| Province | | | | R&D Performer | | | Percentage of Expenditures | |
|----------------------|--------------|--------------------|--------------------|------------------|-------------------|-------------------|-------------------------------|-------|
| | | Patentees | Other Companies | University | Hospitals | Others | Total | |
| Newfoundland | \$(000) % | 831.94 29.1 | 211.24 7.4 | 681.19 23.8 | 761.33 26.6 | 373.64 13.1 | 2,859.32 100 | 0.42 |
| Prince Edward Island | \$(000) % | 84.14 24.6 | 22.30 6.5 | 121.15 35.4 | 5.04 1.5 | 109.93 32.1 | 342.56 100 | 0.05 |
| Nova Scotia | \$(000) % | 1,325.43 12.8 | 947.36 9.1 | 2,703.19 26.1 | 3,700.90 35.7 | 1,699.88 16.4 | 10,376.76 100 | 1.53 |
| New Brunswick | \$(000) % | 127.10 5.3 | 661.99 27.7 | 189.55 7.9 | 724.76 30.4 | 683.75 28.6 | 2,387.15 100 | 0.35 |
| Quebec | \$(000) % | 185,502.21 63.8 | 41,489.24 14.3 | 8,411.60 2.9 | 23,905.16 8.2 | 31,247.54 10.8 | 290,555.75 100 | 42.78 |
| Ontario | \$(000) % | 174,089.80 58.7 | 25,798.85 8.7 | 15,945.84 5.4 | 45,264.76 15.3 | 35,476.6 12.0 | 296,575.84 100 | 43.66 |
| Manitoba | \$(000) % | 5,705.95 45.2 | 663.17 5.3 | 1,519.92 12.1 | 3,218.54 25.5 | 1,504.55 11.9 | 12,612.14 100 | 1.86 |
| Saskatchewan | \$(000) % | 544.12 8.5 | 703.44 11.0 | 2,175.99 34.1 | 1,606.29 25.2 | 1,345.82 21.1 | 6,375.66 100 | 0.94 |
| Alberta | \$(000) % | 3,714.49 11.8 | 6,122.33 19.4 | 9,935.44 31.5 | 5,721.53 18.1 | 6,041.68 19.2 | 31,535.46 100 | 4.64 |
| British Columbia | \$(000) % | 3,870.69 15.2 | 2,609.01 10.2 | 7,845.53 30.8 | 8,282.76 32.5 | 2,849.45 11.2 | 25,457.44 100 | 3.75 |
| Yukon & N.W.T. | \$(000) % | 1.96 1.2 | 163.65 97.4 | 2.21 1.3 | 0.00 0.0 | 0.22 0.1 | 168.04 100 | 0.02 |
| Canada | \$(000) % | 375,797.82 55.3 | 79,392.57 11.7 | 49,531.61 7.3 | 93,191.07 13.7 | 81,333.06 12.0 | 679,246.12 100 | 100 |

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The percentage under each R&D category gives the percentage of all money spent in that category in that province. Expenditures as a percentage of total means percentage of R&D expenditures in that province compared to total R&D in Canada. Rows and columns may not equal totals due to rounding. Current expenditures plus capital expenditures (equipment + depreciation) = total R&D expenditures. •

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Source: PMPRB

Glossary

Note To Reader: This glossary is included for the convenience of the reader. For more detailed information and definitions please refer to the *Patent Act*, the *Patented Medicines Regulations* and the *Compendium of Guidelines*, *Policies and Procedures*, or contact the PMPRB.

Active Ingredient: Chemical responsible for the claimed pharmacologic effect of a drug product.

Advance Ruling Certificate (ARC): A nonbinding certificate may be issued pursuant to subsection 98(4) of the *Act* at the request of a patentee when the Board is satisfied that the price or proposed price of the medicine would not exceed the maximum non-excessive price under the Board's Guidelines.

ATC: Anatomical Therapeutic Chemical classification system that divides drugs into different groups according to their site of action and therapeutic and chemical characteristics. This system is used as a guide for selecting comparable medicines for purposes of price review.

Dedication of Patent: A practice whereby a patentee notifies the Commissioner of Patents that it has surrendered its rights and entitlements flowing from the patent for the benefit of the public to use and enjoy (PMPRB Bulletin 15, January 1995, p.4).

Drug Identification Number (DIN):

A registration number that the Health Protection Branch of Health Canada assigns to each prescription and non-prescription drug product marketed under the *Food and Drug Regulations*. The DIN is assigned using information in the following areas: manufacturer of the product; active ingredient(s); strength of active ingredient(s); pharmaceutical dosage form; brand/trade name; and route of administration.

Drug Product: A particular presentation of a medicine characterized by its pharmaceutical dosage form and the strength of the active ingredient(s).

Drug Product, Existing: An existing drug product is a DIN for which a benchmark price has been established in accordance with the Board's Guidelines. (See Chapter 1, subsection 3.3 of the *Compendium of Guidelines, Policies and Procedures.*)

Drug Product, New: A new drug product is one for which the introductory price is under review. Patented drug products are considered new in the year during which they are first introduced on the market in Canada or the year they receive their first patent(s) if previously marketed. For price review purposes, new drug products for a given year are those introduced between December 1, of the previous year and November 30, of the reporting year. Because of the filing requirements under the Patented Medicines Regulations and the manner of calculating benchmark prices, drug products introduced in December are considered to have been introduced in the following year. (See Chapter 1, subsection 3.2 of the Compendium of Guidelines, Policies and Procedures.)

Emergency Drug Release (EDR) Program: See Special Access Program.

Generic Product: A drug product with the same active ingredient, strength and dosage form of a brand name drug product.

General Public (GP) Number: A number that the Health Protection Branch of Health Canada assigns to proprietary medicines that are registered according to the requirements of Division 10 of the *Food and Drug Regulations*. These products may be sold in non-pharmacy outlets in certain provinces.

Investigational New Drug (IND): A drug that has been authorized for clinical evaluation (i.e., testing on humans) by Health Canada but that is not yet approved for sale for the indication under study.

Licence, Compulsory: A licence granted by the Commissioner of Patents in accordance with subsection 39(4) of the *Patent Act* that has been continued pursuant to subsection 11(1) of the *Patent Act Amendment Act, 1992*, which permits the licencee to import, make, use or sell a patented invention pertaining to a medicine. Royalties payable are determined by the Commissioner of Patents who sets the terms of licences pursuant to subsection 39(5) of the *Patent Act*. Except for those compulsory licences issued prior to December 20, 1991, which are continued pursuant to subsection 11(1) of the *Patent Act*, licences issued after December 20, 1991 have no effect.

Licence, Voluntary: A contractual agreement between a patent holder and a licensee under which the licensee is entitled to enjoy the benefit of the patent or to exercise any rights in relation to the patent for some consideration (i.e., royalties in the form of a share of the licensee's sales).

Medicine: Any substance or mixture of substances made by any means, whether produced biologically, chemically, or otherwise, that is applied or administered *in vivo* in humans or in animals to aid in the diagnosis, treatment, mitigation or prevention of disease, symptoms, disorders, abnormal physical states, or modifying organic functions in humans and or animals, however administered.

For greater certainty, this definition includes vaccines, topical preparations, anaesthetics and diagnostic products used *in vivo*, regardless of delivery mechanism (e.g., transdermal, capsule form, injectable, inhaler, etc.). This definition excludes medical devices, *in vitro* diagnostic products and disinfectants that are not used *in vivo* (*Compendium of Guidelines, Policies and Procedures*, Introduction, subsection 1.5).

Notice of Compliance (NOC): A notice in respect of a medicine issued by the Health Protection Branch of Health Canada under section C.08.004 of the *Food and Drug Regulations*. The issuance of an NOC indicates that a drug product meets the required Health Canada standards for use in humans or animals and that the product is approved for sale in Canada.

Patent: An instrument issued by the Commissioner of Patents in the form of letters patent for an invention that provides its holder with a monopoly limited in time, for the claims made within the patent. A patent gives the patentee the exclusive right to make, sell or otherwise exploit the invention for the term of the patent.

Patentee: As defined by subsection 79(1) of the *Patent Act*, "the person for the time being entitled to the benefit of the patent for that invention (pertaining to a medicine) and includes, where any other person is entitled to exercise any rights in relation to that patent other than under a licence continued by subsection 11(1) of the *Patent Act Amendment Act, 1992*, that other person in respect of those rights;"

Research and Development (R&D): Basic or applied research for the purpose of creating new, or improving existing, materials, devices, products or processes (e.g., manufacturing processes).

Research and Development — **Applied Research:** Work that advances scientific knowledge with a specific practical application in view such as creating new or improved products or processes through manufacturing processes or through preclinical or clinical studies.

Research and Development — **Basic Research:** Work that advances scientific knowledge without a specific application in view.

Research and Development — **Clinical Research:** The assessment of the effect of a new medicine on humans. It typically consists of three successive phases, beginning with limited testing for safety in healthy humans then proceeding to further safety and efficacy studies in patients suffering from the target disease.

Research and Development — **Preclinical Research:** Tests on animals to evaluate the pharmacological and toxicological effects of medicines.

Research and Development Expenditures:

For the purposes of the Patented Medicines Regulations, 1994, in particular sections 5 and 6, research and development includes activities for which expenditures would have qualified for the investment tax credit for scientific research and experimental development under the *Income Tax Act* as it read on December 1, 1987.

Special Access Program (SAP): A program operated by Health Canada to give practitioners access to drugs that are not approved or otherwise available for sale in Canada. (Formerly the EDR Program.)

Voluntary Compliance Undertaking (VCU):

A written undertaking by a patentee to adjust its price to conform with the PMPRB's Excessive Price Guidelines (see Chapter 1 of the *Compendium of Guidelines, Policies and Procedures*). Pursuant to the Board's Compliance and Enforcement Policy (see Chapter 2, section 7) the Chairperson or the Board may approve a VCU in lieu of issuing a Notice of Hearing if it is consistent with the *Patent Act* and the policies of the Board and in the public interest. The Board reports publicly on all VCUs approved by the Chairperson or the Board.

List of Publications

- Patent Act
- Patented Medicines Regulations
- Compendium of Guidelines, Policies and Procedures
- Patentees' Guide to Reporting (1995)

ANNUAL REPORT Series (1989 to 1997)

NEWSletter Series (1997 to 1998)

BULLETIN Series (1988 to 1996)

ARTICLE Series

- Dr. H. C. Eastman Article "Pharmaceutical Price Review in Canada," 1992
- Dr. R. G. Elgie Article "Regulating Prices of Patented Pharmaceuticals in Canada: The Patented Medicine Prices Review Board," 1996

VCUs and DECISIONS of the Board (1993 to 1998)

• 1998: Humalog

SPEECH Series (1989 to 1998)

Toll-Free Number: 1-877-861-2350

If you have any questions or comments, please contact: Sylvie Dupont-Kirby Box L40 Standard Life Centre 333 Laurier Avenue West Suite 1400 Ottawa, Ontario K1P 1C1

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TO ORDER PMPRB PUBLICATIONS, contact Céline Carré at (613) 952-3303

- Chairperson's presentation to the Standing Committee on Industry, March 4, 1997
- CPIC'97 Chairperson's presentation to the Canadian Pharmaceutical Industry Conference 1997, October 20, 1997
- Chairperson's Introductory Remarks to the Standing Committee on Health, April 28, 1998

STUDY Series

- S-9301: International Price Comparison
- S-9302: International Price Comparison of the Top 200 Selling Patented Products Sold in Canada
- S-9303: Further Analyses on Introductory Medicines and Top-Selling Medicines
- S-9404: The Top 200 Selling Patented Drug Products in Canada (1993)
- S-9405: Interprovincial Price Comparisons in Canada (1988-1993)
- S-9506: Estimated Savings from Compliance and Enforcement Activities
- S-9607: The Top 200 Selling Patented Drug Products in Canada (1994)
- S-9708: The Impact of Federal Regulation of Patented Drug Prices
- S-9709: A Comparison of Pharmaceutical Research and Development Spending in Canada and Selected Countries
- S-9710: A Description of the Laspeyres Methodology Used to Construct the Patented Medicine Price Index (PMPI)

DISCUSSION PAPER Series

• D-9401: Measurement of Cost Savings to the Canadian Health Care System

Table 12Patented Drug Products Introduced in Canada in 1997

HUMAN

Category 1

| Brand Name | DIN | Company |
|---|----------|---|
| ACTHREL NF - 0.1MG/VIAL | | FERRING INC. |
| CERVIDIL 0.3 - 10MG/POUCH | 02231047 | FERRING INC. |
| EPREX - 10000UNIT/SYRINGE | 02231587 | JANSSEN-ORTHO INC. |
| EPREX - 1000UNIT/SYRINGE | 02231583 | JANSSEN-ORTHO INC. |
| EPREX - 2000UNIT/SYRINGE | 02231584 | JANSSEN-ORTHO INC. |
| EPREX - 3000UNIT/SYRINGE | 02231585 | JANSSEN-ORTHO INC. |
| EPREX - 4000UNIT/SYRINGE | 02231586 | JANSSEN-ORTHO INC. |
| FOSAMAX - 5MG/TAB | 02233055 | MERCK FROSST CANADA INC. |
| FRAGMIN - 25000UNIT/ML | 02231171 | PHARMACIA & UPJOHN INC. |
| HAVRIX 720 JUNIOR - 1440UNIT/ML | 02231056 | SMITHKLINE BEECHAM PHARMA INC. |
| HUMATROPE - 13.3MG/CART | 02229693 | ELI LILLY CANADA INC. |
| HUMATROPE - 6.7MG/CART | 02229692 | ELI LILLY CANADA INC. |
| IMODIUM LINGUAL - 2MG/TAB | 02230542 | MCNEIL CONSUMER PRODUCTS COMPANY |
| INNOHEP - 10000UNIT/ML | 02229755 | LEO LABORATORIES CANADA LTD. |
| INNOHEP - 20000UNIT/ML | 02229515 | LEO LABORATORIES CANADA LTD. |
| INTRON-A - 18000000UNIT/VIAL | 02231651 | SCHERING CANADA INC. |
| LOSEC - 10MG/TAB | 02230737 | ASTRA PHARMA INC. |
| LUPRON DEPOT - 22.5MG/VIAL | 02230248 | ABBOTT LABORATORIES LIMITED |
| MIRALUMA | | DU PONT MERCK PHARMA INC. |
| NAROPIN - 7.5MG/ML | 02229416 | ASTRA PHARMA INC. |
| NUTROPIN - 5MG/VIAL | 02216183 | HOFFMANN-LA ROCHE LIMITED |
| NUTROPIN AQ - 5MG/ML | 02229722 | HOFFMANN-LA ROCHE LIMITED |
| PAXIL - 10MG/TAB | 02027887 | SMITHKLINE BEECHAM PHARMA INC. |
| PEPCID AC - 10MG/TAB | 02185911 | JOHNSON & JOHNSON-MERCK |
| PNU-IMUNE 23 | 02230175 | WYETH-AYERST CANADA INC. |
| PULMICORT NEBUAMP - 0.125MG/ML | 02229099 | ASTRA PHARMA INC. |
| REACTINE - 5MG/TAB | 02223546 | PFIZER CANADA INC. |
| RHINOCORT AQ NASAL AEROSOL - 0.064MG/DOSE | 02231923 | ASTRA PHARMA INC. |
| ROFERON-A - 4500000UNIT/VIAL | 02217023 | HOFFMANN-LA ROCHE LIMITED |
| ROFERON-A - 600000UNIT/VIAL | 02217031 | HOFFMANN-LA ROCHE LIMITED |
| SENSODYNE-F 5%/0.24% | 02083310 | BLOCK DRUG COMPANY (CANADA) LTD. |
| SERZONE - 50MG/TAB | 02087294 | BRISTOL-MYERS SQUIBB PHARMÁCEUTICAL GROUP |
| SUPREFACT DEPOT - 6.6MG/VIAL | 02228955 | HOECHST MARION ROUSSEL CANADA INC. |
| ZITHROMAX - 600MG/TAB | 02231143 | PFIZER CANADA INC. |

Category 2

| Brand Name | DIN | Company | New Active Substance |
|-------------------------|----------|---------------------------|----------------------|
| ARICEPT - 10MG/TAB | 02232044 | PFIZER CANADA INC. | NAS |
| ARICEPT - 5MG/TAB | 02232043 | PFIZER CANADA INC. | NAS |
| BENEFIX - 1000UNIT/VIAL | 02231020 | GENETICS INSTITUTE, INC. | NAS |
| BENEFIX - 250UNIT/VIAL | 02231018 | GENETICS INSTITUTE, INC. | NAS |
| BENEFIX - 500UNIT/VIAL | 02231019 | GENETICS INSTITUTE, INC. | NAS |
| CAMPTOSAR - 20MG/ML | 02231622 | PHARMACIA & UPJOHN INC. | NAS |
| CRIXIVAN - 200MG/CAP | 02229161 | MERCK FROSST CANADA INC. | FPG ¹ |
| CRIXIVAN - 400MG/CAP | 02229196 | MERCK FROSST CANADA INC. | FPG |
| EPREX - 10000UNIT/ML | 02126591 | JANSSEN-ORTHO INC. | FPG |
| EPREX - 20000UNIT/ML | 02206072 | JANSSEN-ORTHO INC. | FPG |
| EPREX - 2000UNIT/ML | 02126575 | JANSSEN-ORTHO INC. | FPG |
| EPREX - 4000UNIT/ML | 02126583 | JANSSEN-ORTHO INC. | FPG |
| INVIRASE - 200MG/CAP | 02216965 | HOFFMANN-LA ROCHE LIMITED | FPG |

Category 3

| Brand Name | DIN | Company | New Active Substance |
|-----------------|----------|--------------------------|----------------------|
| 3TC - 10MG/ML | 02192691 | GLAXO WELLCOME INC. | FPG |
| 3TC - 150MG/TAB | 02192683 | GLAXO WELLCOME INC. | FPG |
| ACEL-P | 02215071 | WYETH-AYERST CANADA INC. | NAS |

| ALLEGRA - 60MG/TAB ANZEMET - 100MG/TAB ANZEMET - 20MG/ML ANZEMET - 50MG/TAB | 02231462 02231379 02231380 02231378 | HOECHST MARION ROUSSEL CANADA INC. HOECHST MARION ROUSSEL CANADA INC. HOECHST MARION ROUSSEL CANADA INC. HOECHST MARION ROUSSEL CANADA INC. NEXSTAR PHARMACEUTICALS INC. ROBERTS PHARMACEUTICAL CANADA INC. | NAS NAS NAS NAS |
|--|--|---|--|
| DAUNOXOME - 50MG/VIAL EMINASE - 30UNIT/VIAL | 02218046 02229390 | ROBERTS PHARMACEUTICALS INC. | NAS |
| ALLEGRA - 60MG/TAB ANZEMET - 100MG/TAB ANZEMET - 20MG/ML ANZEMET - 50MG/VIAL EMINASE - 30UNIT/VIAL EMINASE - 30UNIT/VIAL EMINASE - 30UNIT/VIAL ENTOCORT - 3MG/CAP FEMARA - 2.5MG/TAB FLOLAN - 0.5MG/VIAL FLOLAN - 0.5MG/VIAL FLOLAN - 1.5MG/VIAL FLOLAN STERILE DILUENT FORADIL - 0.012MG/DOSE HYCAMTIN - 4MG/VIAL IMITREX - 20MG/DOSE LIPITOR - 10MG/TAB LIPITOR - 20MG/TAB LIPITOR - 40MG/TAB MIREZE - 20MG/ML MYOVIEW NAROPIN - 10MG/ML NAROPIN - 10MG/ML NAROPIN - 5MG/ML PANTOLOC - 40MG/TAB PENTACEL PYLORID - 400MG/TAB REQUIP - 0.25MG/TAB REQUIP - 10MG/TAB REQUIP - 5MG/TAB REQUIP - 5MG/TAB REQUIP - 5MG/TAB REQUIP - 5MG/TAB TASMAR - 100MG/TAB TASMAR - 100MG/TAB TAZORAC - 0.5MG/G TAZORAC - 0.5MG/G TAZORAC - 0.5MG/G TAZORAC - 0.5MG/TAB TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 0.5MG/G TAZORAC - 0.5MG/TAB TAZORAC - 25MG/TAB TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 0.5MG/G TAZORAC - 25MG/TAB TAZORAC - 25MG/TAB TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 25MG/TAB TAZORAC - 0.5MG/G TAZORAC - 25MG/TAB TOPAMAX - 25MG/TAB TOPAMAX - 25MG/TAB TOPAMAX - 25MG/TAB TRIPACEL TWINRIX 720/20 XALATAN - 0.05MG/ML | 02229293 02231384 02230845 02230848 02230857 02230898 02231116 02230420 | ASTRA FRAMMA INC. NOVARTIS PHARMA CANADA INC. GLAXO WELLCOME INC. GLAXO WELLCOME INC. NOVARTIS PHARMA CANADA INC. SMITHKLINE BEECHAM PHARMA INC. GLAXO WELLCOME INC. | NAS NAS NAS NAS NAS NAS |
| IMITREX - 5MG/DOSE LIPITOR - 10MG/TAB LIPITOR - 20MG/TAB LIPITOR - 40MG/TAB | 02230418 02230711 02230713 02230714 | GLAXO WELLCOME INC. WARNER-LAMBERT CANADA INC. (PARKE-DAVIS) WARNER-LAMBERT CANADA INC. (PARKE-DAVIS) WARNER-LAMBERT CANADA INC. (PARKE-DAVIS) ALLERGAN INC. | NAS NAS NAS |
| MIREZE - 20MG/ML MYOVIEW NAROPIN - 10MG/ML | 02230446 | ALLERGAN INC. AMERSHAM CANADA LIMITED | NAS |
| NAROPIN - 2MG/ML NAROPIN - 2MG/ML NAROPIN - 5MG/ML PANTOLOC - 40MG/TAB | 02229410 02229411 02229415 02229453 | ASTRA PHARMA INC. ASTRA PHARMA INC. ASTRA PHARMA INC. SOLVAY PHARMA INC. | NAS NAS NAS |
| PENTACEL PYLORID - 400MG/TAB QUADRACEL | 02231343 02231831 02230946 | GLAXO WELLCOME INC. | NAS NAS |
| REQUIP - 0.25MG/TAB REQUIP - 1MG/TAB REQUIP - 2MG/TAB REQUIP - 5MG/TAB | 02230946 02232565 02232567 02232568 02232569 | SMITHKLINE BEECHAM PHARMA INC. SMITHKLINE BEECHAM PHARMA INC. SMITHKLINE BEECHAM PHARMA INC. SMITHKLINE BEECHAM PHARMA INC. | NAS NAS NAS NAS NAS |
| SPORANOX - 10MG/ML SUPRANE TASMAR - 100MG/TAB TASMAR - 200MG/TAB TAZORAC - 0.5MG/G | 02231347 02227428 02235914 02235921 02230784 | JANSSEN-ORTHO INC. ZENECA PHARMA INC. HOFFMANN-LA ROCHE LIMITED HOFFMANN-LA ROCHE LIMITED | NAS NAS NAS NAS |
| TAZORAC - 0.5MG/G TAZORAC - 1MG/G TOPAMAX - 100MG/TAB TOPAMAX - 200MG/TAB TOPAMAX - 25MG/TAB TRIPACEL THINK ZOO 000 | 02230784 02230785 02230894 02230896 02230893 02224143 | WARNER-LAMBERT CANADA INC. (PARKE-DAVIS) ALLERGAN INC. AMERSHAM CANADA LIMITED ASTRA PHARMA INC. ASTRA PHARMA INC. ASTRA PHARMA INC. SOLVAY PHARMA INC. CONNAUGHT LABORATORIES LTD. GLAXO WELLCOME INC. CONNAUGHT LABORATORIES LTD. SMITHKLINE BEECHAM PHARMA INC. SMITHKLINE BEECHAM PHARMA INC. ZENECA PHARMA INC. HOFFMANN-LA ROCHE LIMITED HOFFMANN-LA ROCHE LIMITED HOFFMANN-LA ROCHE LIMITED ALLERGAN INC. JANSSEN-ORTHO INC. JANSSEN-ORTHO INC. JANSSEN-ORTHO INC. SMITHKLINE BEECHAM PHARMA INC. PHARMACIA & UPJOHN INC. | NAS NAS NAS NAS NAS |
| TWINRIX 720/20 XALATAN - 0.05MG/ML | 02230578 02231493 | Smithkline Beecham Pharma Inc. Pharmacia & UPJOHN INC. | NAS |

VETERINARY

Category 3

| Brand Name | DIN | Company | New Active Substance |
|--------------------|----------|---------------------------------------|----------------------|
| AVIAX - 50000MG/KG | 02229543 | PFIZER CANADA INC.,ANIMAL HEALTH GROU | IP NAS |
| SENTINEL 115/5.75 | 02229547 | NOVARTIS ANIMAL HEALTH CANADA INC. | |
| SENTINEL 230/11.5 | 02229548 | NOVARTIS ANIMAL HEALTH CANADA INC. | |
| SENTINEL 46/2.3 | 02229551 | NOVARTIS ANIMAL HEALTH CANADA INC. | |
| SENTINEL 460/23 | 02229549 | NOVARTIS ANIMAL HEALTH CANADA INC. | |

1 FPG: First patent granted in 1997.

Table 13Patented Drug Products and Canadian Patentees,January 1 - December 31, 1997

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|---|--|----------------|--|------------|
| 3M PHAR | MACEUTICALS, 3M CANADA | INC. | | | |
| Human: 02162806 | MINITRAN 0.2 - 18MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 02163527 | MINITRAN 0.4 - 36MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 02163535 | MINITRAN 0.6 - 54MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 02164337 | MINITRAN 0.8 - 72MG/PATCH | nitroglycerin | C01DA | transdermal patch | not sold |
| 01966197 01966200 | TAMBOCOR - 50MG/TAB TAMBOCOR - 100MG/TAB | flecainide acetate flecainide acetate | CO1BC CO1BC | tablet | |
| 01900200 | TAMBOCOR - 150MG/TAB | flecainide acetate | CO1BC | tablet tablet | not sold |
| 00628239 | TAMBOCOR - 200MG/TAB | flecainide acetate | C01BC | tablet | not sold |
| ABBOTT I | ABORATORIES LIMITED | | | | |
| Human: 02146908 | BIAXIN - 25MG/ML | clarithromycin | J01FA | nowder for oral suspension | |
| 02140908 | BIAXIN - 250MG/ML BIAXIN - 250MG/TAB | clarithromycin clarithromycin | J01FA | powder for oral suspension tablet | |
| 02126710 | BIAXIN - 500MG/TAB | clarithromycin | J01FA | tablet | |
| 00891738 | CALCIJEX - 0.001MG/ML | calcitriol | A11CC | injectable solution | |
| 00891746 | CALCIJEX - 0.002MG/ML | calcitriol | A11CC | injectable solution | |
| 00596418 | EPIVAL - 125MG/TAB | divalproex sodium | N03AG | enteric-coated tablet | |
| 00596426 00596434 | EPIVAL - 250MG/TAB EPIVAL - 500MG/TAB | divalproex sodium divalproex sodium | N03AG N03AG | enteric-coated tablet enteric-coated tablet | |
| 00390434 | ERYBID - 500MG/TAB | erythromycin | J01FA | tablet | |
| 00682268 | ERYTHROCIN ADD-VANTAGE - | erythromycin lactobionate | J01FA | powder for injectable solution | |
| 00682276 | 500MG/VIAL ERYTHROCIN ADD-VANTAGE - 1000MG/VIAL | erythromycin lactobionate | J01FA | powder for injectable solution | |
| 00818658 | HYTRIN - 1MG/TAB | terazosin hydrochloride | C02CA | tablet | |
| 00818682 | HYTRIN - 2MG/TAB | terazosin hydrochloride | C02CA | tablet | |
| 00818666 | HYTRIN - 5MG/TAB | terazosin hydrochloride | CO2CA | tablet | |
| 00818674 02187876 | HYTRIN - 10MG/TAB HYTRIN-STARTER PACK | terazosin hydrochloride terazosin hydrochloride | CO2CA CO2CA | tablet tablet | expired |
| 02107070 | LUPRON DEPOT - 3.75MG/VIAL | leuprolide acetate | L02AE | powder for injectable solution | expireu |
| 00836273 | LUPRON DEPOT - 7.5MG/VIAL | leuprolide acetate | LOZAE | powder for injectable solution | |
| 02148722 | LUPRON DEPOT - 11.25MG/VIAL | leuprolide acetate | L02AE | powder for injectable solution | not sold |
| 02148730 | LUPRON DEPOT - 15MG/VIAL | leuprolide acetate | L02AE | powder for injectable solution | not sold |
| 02230248 | LUPRON DEPOT - 22.5MG/VIAL | leuprolide acetate | L02AE | powder for injectable solution | introduced |
| 02229137 02229145 | NORVIR - 100MG/CAP NORVIR - 80MG/ML | ritonavir ritonavir | J05AE J05AE | capsule oral solution | |
| 00769991 | PCE DISPERTAB - 333MG/TAB | erythromycin | J01FA | tablet | |
| 02165503 | PREVACID - 15MG/CAP | lansoprazole | A02BC | sustained-release capsule | |
| 02165511 | PREVACID - 30MG/CAP | lansoprazole | A02BC | sustained-release capsule | |
| 02172763 | SEVORANE | sevoflurane | N01AB | inhalation anesthetic | |
| 02016109 | SURVANTA - 25MG/ML | beractant | R07AA | endo-tracheal suspension | |
| ALCON CA | ANADA INC. | | | | |
| 00695688 | BETOPTIC - 5MG/ML | betaxolol hydrochloride | S01ED | ophthalmic solution | expired |
| 01908448 | BETOPTIC S - 2.5MG/ML | betaxolol hydrochloride | S01ED | ophthalmic suspension | 1 |
| 00568082 | BSS PLUS | sodium bicarbonate/ | 00.000 | | |
| 01045070 | | dextrose/glutathione | S01XA | ophthalmic solution | |
| 01945270 02076306 | CILOXAN - 3MG/ML IOPIDINE - 5MG/ML | ciprofloxacin hydrochloride apraclonidine hydrochloride | S01AX S01XA | ophthalmic solution ophthalmic solution | |
| 02070300 | IOPIDINE - 10MG/ML | apracionidine hydrochloride | S01XA | ophthalmic solution | |
| 02233143 | PATANOL - 1MG/ML | olopatadine hydrochloride | | ophthalmic solution | not sold |
| 00575240 | PILOPINE-HS - 40MG/G | pilocarpine hydrochloride | S01EB | ophthalmic gel | |
| 02132710 | PROFENAL - 10MG/ML | suprofen | M01AE | ophthalmic solution | not sold |
| 00390291 | TEARS NATURALE 1/3 | dextran/hydroxypropyl methylcellulose | S01XA | ophthalmic solution | expired |
| 00743445 | TEARS NATURALE II 1/3 | dextran/hydroxypropyl | | | |
| 00770045 | | methylcellulose | S01XA | ophthalmic drops | expired |
| 00778915 00778907 | TOBRADEX 3/1 TOBRADEX 3/1 | tobramycin/dexamethasone tobramycin/dexamethasone | S01CA S01CA | ophthalmic ointment ophthalmic suspension | |
| 00110301 | | | JUTUR | opininanino suspension | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|--|---|---|--|--|
| ALLERGA | | | AIU | | oominicitia |
| Human: | N ING. | | | | |
| 01968300 02230446 02011255 02011263 | ACULAR - 5MG/ML ILOTYCIN - 5MG/G MIREZE - 20MG/ML NAFTIN - 10MG/G NAFTIN - 10MG/G | ketorolac tromethamine erythromycin nedocromil sodium naftifine hydrochloride naftifine hydrochloride | S01BC S01AA S01GX D01AE D01AE | ophthalmic solution ointment ophthalmic drops cream gel | introduced expired expired |
| 02143291 02230784 02230785 | OCUFLOX - 3MG/ML TAZORAC - 0.5MG/G TAZORAC - 1MG/G | ofloxacin tazarotene tazarotene | S01AA D05AX D05AX | ophthalmic solution gel gel | introduced (nas) introduced (nas) |
| ALPHA TH | IERAPEUTIC CORPORATION | | | | |
| Human: 01924745 | ALPHANINE - 1UNIT/DOSE ALPHANINE SD - 1UNIT/DOSE | factor IX(human) factor IX(human) | B02BD B02BD | injectable solution injectable solution | not sold |
| 00740780 | PROFILATE - 1UNIT/DOSE | antihemophilic fáctor | B02BD | injectable solution | not sold |
| | PHARMACEUTICALS INC. | | | | |
| Human: 01912070 | DOMPERIDONE MALEATE - 10MG/TAB | domperidone maleate | A03FA | tablet | expired |
| 02128918 02128926 02128977 02128985 02128985 02128853 02128861 02128861 02128861 02128861 02128896 00756814 00615307 00615315 00615323 00615331 02039478 02039567 00828688 00878715 00828688 00878715 00851639 00851883 00851647 00851655 00878790 00675369 | KENRAL-CEFACLOR - 250MG/CAP KENRAL-CEFACLOR - 500MG/CAP KENRAL-CEFACLOR - 25MG/ML KENRAL-CEFACLOR - 50MG/ML KENRAL-CEFACLOR - 75MG/ML KENRAL-FLUOXETINE - 10MG/CAP KENRAL-FLUOXETINE - 20MG/CAP KENRAL-FLUOXETINE - 4MG/ML KENRAL-NIZATIDINE - 150MG/CAP | cefaclor cefaclor cefaclor cefaclor cefaclor fluoxetine hydrochloride fluoxetine hydrochloride fluoxetine hydrochloride nizatidine naproxen naproxen naproxen naproxen naproxen naproxen naproxen naproxen naproxen naproxen naproxen naproxen salbutamol sulfate captopril captopril captopril flunisolide naproxen sodium | J01DA J01DA J01DA J01DA J01DA N06AB N06AB N06AB A02BA M01AE | capsule capsule powder for oral suspension powder for oral suspension capsule capsule oral solution capsule capsule capsule capsule suppository tablet | not sold not sold |
| 01900897 | SYNFLEX DS - 550MG/TAB | naproxen sodium | M01AE | tablet | |
| _ | AM CANADA LIMITED | | | | |
| Human: | CERETEC | technetium tc-99m exametazime | V09AA | powder for injectable solutio | n |
| | HEPATATE II MEDRONATE II | technetium tc-99m tin colloid technetium tc-99m medronate sodium | V09DB V09BA | powder for injectable solutio powder for injectable solutio | n n |
| | MYOVIEW | technetium tc-99m tetrofosmin | V09AG | powder for injectable solution | introduced (nas) |
| | ANADA INC. | | | | |
| Human: 01968017 | NEUPOGEN - 0.3MG/ML | filgrastim | L03AA | injectable solution | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|---|--|----------------|---|------------------------------|
| | HARMA INC. | | | | |
| Human: 02148641 | BAMBEC - 10MG/TAB | bambuterol hydrochloride | R03CC | tablet | not sold |
| 02148668 | BAMBEC - 20MG/TAB | bambuterol hydrochloride | R03CC | tablet | not sold |
| 00719846 | BETALOC - 1MG/ML | metoprolol tartrate | C07AB | injectable solution | |
| 00402605 | BETALOC - 50MG/TAB | metoprolol tartrate | C07AB | tablet | |
| 00402540 | BETALOC - 100MG/TAB | metoprolol tartrate | C07AB | tablet | |
| 02028727 | BETALOC CR - 47.5MG/TAB | metoprolol succinate | C07AB | sustained-release tablet | not sold |
| 02028735 | BETALOC CR - 95MG/TAB | metoprolol succinate | C07AB | sustained-release tablet | not sold |
| 02028743 | BETALOC CR - 190MG/TAB | metoprolol succinate | C07AB | sustained-release tablet | not sold |
| 00497827 | BETALOC DURULES - 200MG/TAB | metoprolol tartrate | C07AB | sustained-release tablet | |
| 00444774 | BRICANYL SPACER - 0.25MG/DOSE | | R03AC | aerosol for inhalation | expired |
| 00786616 | BRICANYL TURBUHALER - 0.5MG/DOSE | terbutaline sulfate | R03AC | powder for inhalation | |
| 01958100 | CARDURA-1 - 1MG/TAB | doxazosin mesylate | CO2CA | tablet | expired |
| 01958097 | CARDURA-2 - 2MG/TAB | doxazosin mesylate | CO2CA | tablet | expired |
| 01958119 | CARDURA-4 - 4MG/TAB | doxazosin mesylate | CO2CA | tablet | expired |
| 00886858 | EMLA 25/25 | lidocaine/prilocaine | D04AB | cream | |
| 02057794 02166062 | EMLA 25/25 | lidocaine/prilocaine | D04AB D04AB | transdermal patch cream | not sold |
| 02229293 | EMLA STERILE 25/25 ENTOCORT - 3MG/CAP | lidocaine/prilocaine budesonide | A07EA | sustained-release capsule | introduced |
| 02052431 | ENTOCORT - 0.02MG/ML | budesonide | A07EA | enema | Introduced |
| 02134810 | FOSCAVIR - 24MG/ML | foscarnet sodium | J05AD | injectable solution | not sold |
| 02119579 | LOSEC - 10MG/CAP | omeprazole | A02BC | capsule | not sold |
| 00846503 | LOSEC - 20MG/CAP | omeprazole | A02BC | capsule | not sold |
| 02016788 | LOSEC - 40MG/CAP | omeprazole | A02BC | capsule | not sold |
| 02230737 | LOSEC - 10MG/TAB | omeprazole magnesium | A02BC | sustained-release tablet | introduced |
| 02190915 | LOSEC - 20MG/TAB | omeprazole magnesium | A02BC | sustained-release tablet | |
| 02229411 | NAROPIN - 2MG/ML | ropivacaine hydrochloride | N01BB | injectable solution | introduced (nas) |
| 02229415 | NAROPIN - 5MG/ML | ropivacaine hydrochloride | N01BB | injectable solution | introduced (nas) |
| 02229416 02229418 | NAROPIN - 7.5MG/ML NAROPIN - 10MG/ML | ropivacaine hydrochloride | N01BB | injectable solution | introduced |
| 02229418 | NITROGARD-SR - 1MG/TAB | ropivacaine hydrochloride nitroglycerin | N01BB C01DA | injectable solution sustained-release tablet | introduced (nas) not sold |
| 00749397 | NITROGARD-SR - 2MG/TAB | nitroglycerin | CO1DA | sustained-release tablet | not sold |
| 00749389 | NITROGARD-SR - 3MG/TAB | nitroglycerin | CO1DA | sustained-release tablet | not sold |
| 00749370 | NITROGARD-SR - 5MG/TAB | nitroglycerin | C01DA | sustained-release tablet | not sold |
| 00627127 | PENGLOBE - 400MG/TAB | bacampicillin hydrochloride | J01CA | tablet | |
| 00627135 | PENGLOBE - 800MG/TAB | bacampicillin hydrochloride | J01CA | tablet | |
| 02057778 | PLENDIL - 2.5MG/TAB | felodipine | C08CA | sustained-release tablet | |
| 00851779 | PLENDIL - 5MG/TAB | felodipine | C08CA | sustained-release tablet | |
| 00851787 | PLENDIL - 10MG/TAB | felodipine | CO8CA | sustained-release tablet | |
| 00817228 | PULMICORT INHALER - 0.05MG/DOSE | budesonide | R03BA | aerosol for inhalation | not sold |
| 00634549 | PULMICORT INHALER - 0.2MG/DOSE | budesonide | R03BA | aerosol for inhalation | not sold |
| 02229099 | PULMICORT NEBUAMP - | budesonide | R03BA | suspension for inhalation | introduced |
| 01978918 | 0.125MG/ML PULMICORT NEBUAMP - | budesonide | R03BA | suspension for inhalation | |
| 01978926 | 0.25MG/ML PULMICORT NEBUAMP - | budesonide | R03BA | suspension for inhalation | |
| 00634530 | 0.5MG/ML PULMICORT SPACER - 0.05MG/DOSE | budesonide | R03BA | aerosol for inhalation | not sold |
| 00814091 | PULMICORT SPACER - 0.2MG/DOSE | budesonide | R03BA | aerosol for inhalation | not sold |
| 00852074 | PULMICORT TURBUHALER - 0.1MG/DOSE | budesonide | R03BA | powder for inhalation | |
| 00851752 | PULMICORT TURBUHALER - 0.2MG/DOSE | budesonide | R03BA | powder for inhalation | |
| 00851760 | PULMICORT TURBUHALER - 0.4MG/DOSE | budesonide | R03BA | powder for inhalation | |
| 02051788 | RAMACE - 1.25MG/CAP | ramipril | C09AA | capsule | not sold |
| 02051796 | RAMACE - 2.5MG/CAP | ramipril | C09AA | capsule | not sold |
| 02051818 | RAMACE - 5MG/CAP | ramipril | C09AA | capsule | not sold |
| 00636460 | RHINOCORT - 0.05MG/DOSE | budesonide | R01AD | nasal aerosol | not sold |
| | | | | | |

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|---|---|---|----------------------------------|--|--|
| 02231922 | RHINOCORT AQUA - 0.032MG/DOSE | budesonide | R01AD | nasal aerosol | not sold |
| 01974424 02231923 | RHINOCORT AQUA - 0.05MG/DOSE RHINOCORT AQUA - 0.064MG/DOSE | budesonide budesonide | R01AD R01AD | nasal aerosol nasal aerosol | not sold introduced |
| 01974432 02035324 | RHINOCORT AQUA - 0.1MG/DOSE RHINOCORT TURBUHALER - | budesonide budesonide | R01AD R01AD | nasal aerosol powder for nasal inhalation | |
| 02035340 | 0.1MG/DOSE RHINOCORT TURBUHALER - 0.2MG/DOSE | budesonide | R01AD | powder for nasal inhalation | not sold |
| 02036363 02036371 02036398 02036401 | ROXIAM - 75MG/CAP ROXIAM - 150MG/CAP ROXIAM - 300MG/CAP ROXIAM - 300MG/CAP ROXIAM - 100MG/ML | remoxipride hydrochloride remoxipride hydrochloride remoxipride hydrochloride remoxipride hydrochloride | N05AL N05AL N05AL N05AL | capsule capsule capsule injectable solution | not sold not sold not sold not sold |
| AYERST V | ETERINARY LABORATORIES | | | | |
| Veterinary: 00844306 00844292 00849413 00844284 00844241 00844233 00844225 | AMIGLYDE - V - 50MG/ML AMIGLYDE - V - 250MG/ML CEFA - 50MG/ML CEFA - 50MG/TAB CEFA - 100MG/TAB CEFA - 200MG/TAB CEFA - 1000MG/TAB | amikacin sulfate amikacin sulfate cefadroxil cefadroxil cefadroxil cefadroxil cefadroxil | | injectable solution injectable solution powder for oral solution tablet tablet tablet tablet | not sold |
| 02215713 02228386 | CYDECTIN - 10MG/ML CYDECTIN POUR-ON - 5MG/ML ECOLAN ECOLAN-RC | moxidectin moxidectin escherichia coli vaccine escherichia coli rota corona vaccine | | injectable solution topical solution injectable suspension injectable suspension | not sold not sold not sold |
| 02148676 02148684 02148692 02231660 00845000 00844977 00844985 00844993 | GUARDIAN - 0.03MG/TAB GUARDIAN - 0.068MG/TAB GUARDIAN - 0.136MG/TAB HEVLAN TC (VACCINE) QUEST - 20MG/ML TORBUGESIC - 10MG/ML TORBUTROL - 1MG/TAB TORBUTROL - 5MG/TAB TORBUTROL - 10MG/TAB | moxidectin moxidectin hemorrhagic enteritis vaccine moxidectin butorphanol tartrate butorphanol tartrate butorphanol tartrate butorphanol tartrate | 1 | tablet tablet tablet injectable suspension oral gel injectable solution tablet tablet tablet | not sold not sold not sold not sold not sold |
| | ORPORATION | | | | |
| Human: 00781339 00781347 | ANTITHROMBIN III IMMUNO BEBULIN VH BEBULIN VH CMV IVEEGAM IMMUNO - | antithrombin III factor IX complex (human) factor IX complex (human) cmv immune globulin | B01AB B02BD B02BD J06BB | powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution | not sold not sold not sold |
| 02230772 | 1000MG/VIAL CRITILIP 20% | intravenous (human) medium and long chain triglycerides | B05BA | injectable suspension | not sold |
| 00609137 | FACTOR VII IMMUNO VH FEIBA VH IMMUNO GAMMAGARD S/D - 50MG/ML | factor VII concentrate factor VIII anti inhibitor immune globulin | BO2BD B02BD J06BA | powder for injectable solution powder for injectable solution powder for injectable solution | |
| 00808709 02206021 01959379 | HEMOFIL-M IMMUMINE VH IVEEGAM IMMUNO - 500MG/VIAL | intravenous (human) factor VIII factor IX (human) immune globulin intravenous (human) | B02BD B02BD J06BA | powder for injectable solution powder for injectable solution powder for injectable solution | not sold |
| 01959336 | IVEEGAM IMMUNO - 1000MG/VIAL | immune globulin intravenous (human) | J06BA | powder for injectable solution | not sold |
| 01959328 | IVEEGAM IMMUNO - 2500MG/VIAL | | J06BA | powder for injectable solution | not sold |
| 01959301 | IVEEGAM IMMUNO - 5000MG/VIAL | immune globulin intravenous (human) | J06BA | powder for injectable solution | |
| 01959298 | IVEEGAM IMMUNO - 7500MG/VIAL | immune globulin intravenous (human) | J06BA | powder for injectable solution | not sold |
| 01959344 | IVEEGAM IMMUNO - 10000MG/VIAL | immune globulin intravenous (human) | J06BA | powder for injectable solution | not sold |
| 00719603 00719811 | KRYOBULIN VH KRYOBULIN VH | factor VIII complex (human) factor VIII complex (human) | B02BD B02BD | powder for injectable solution powder for injectable solution | not sold not sold |

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|--|---|--|--|--|---|
| 00719838 01958968 01949020 01949012 01974327 | KRYOBULIN VH TISSEEL KIT VH 0.5 TISSEEL KIT VH 1.0 TISSEEL KIT VH 2.0 TISSEEL KIT VH 5.0 | factor VIII complex (human) fibrin sealant fibrin sealant fibrin sealant fibrin sealant fibrin sealant | B02BD B02BC B02BC B02BC B02BC | powder for injectable solution powder for topical solution powder for topical solution powder for topical solution powder for topical solution | not sold |
| BAYER IN | C . | | | | |
| Human: 00852082 02155907 02155990 | ADALAT FT - 10MG/TAB ADALAT XL - 30MG/TAB ADALAT XL - 60MG/TAB | nifedipine nifedipine nifedipine | C08CA C08CA C08CA | tablet sustained-release tablet sustained-release tablet | not sold |
| 00629243 02150948 02155931 02155958 02155966 02155974 02155982 02178842 | CANESTEN 1 - 500MG/TAB CANESTEN 1 COMBI-PAK CIPRO - 100MG/TAB CIPRO - 250MG/TAB CIPRO - 500MG/TAB CIPRO - 750MG/TAB CIPRO IV - 10MG/ML GAMIMUNE N - 50MG/ML | clotrimazole clotrimazole ciprofloxacin hydrochloride ciprofloxacin hydrochloride ciprofloxacin hydrochloride ciprofloxacin hydrochloride ciprofloxacin lactate immune globulin intravenous | G01AF G01AF J01MA J01MA J01MA J01MA J01MA J06BA | vaginal insert vaginal insert and cream tablet tablet tablet tablet injectable solution injectable solution | not sold |
| 01902695 | GAMIMUNE N - 100MG/ML | (human) immune globulin intravenous | J06BA | injectable solution | not sold |
| | GAMIMUNE N HT - 50MG/ML | (human) immune globulin intravenous | J06BA | injectable solution | |
| 02155923 02155915 02190885 02190893 02204592 02204606 02189135 | NIMOTOP - 30MG/CAP NIMOTOP IV - 0.2MG/ML PRANDASE - 50MG/TAB PRANDASE - 100MG/TAB PROLASTIN - 25MG/ML PROLASTIN - 25MG/ML THROMBATE III - 500UNIT/VIAL | (human) nimodipine nimodipine acarbose acarbose alpha1-proteinase alpha1-proteinase antithrombin III | N07XC N07XC A10BF A10BF R07AX R07AX B01AB | capsule injectable solution tablet tablet powder for injectable solution powder for injectable solution powder for injectable solution | not sold |
| 02189143 | THROMBATE III - 1000UNIT/VIAL C., AGRICULTURAL DIVISION | antithrombin III | B01AB | powder for injectable solution | |
| Veterinary: | - | | | | |
| 00812285 00719757 01923781 | BAYTRIL - 22.7MG/ML BAYTRIL - 32.3MG/ML BAYTRIL - 50MG/ML BAYTRIL - 100MG/ML | enrofloxacin enrofloxacin enrofloxacin enrofloxacin | | injectable solution egg dip concentrate injectable solution oral suspension | not sold |
| 00719765 00719773 00719781 | BAYTRIL - 100MG/ML BAYTRIL - 5.7MG/TAB BAYTRIL - 22.7MG/TAB BAYTRIL - 68MG/TAB | enrofloxacin enrofloxacin enrofloxacin enrofloxacin enrofloxacin | | injectable solution tablet tablet tablet | not sold |
| 00469319 02076241 02076268 00573795 00597856 | CUTTER PASTE - 455MG/G DRONTAL PLUS 113.4/22.7/22.7 DRONTAL PLUS 340.2/68/68 NEGABOT PLUS 67.5/400 VERCOM 34/3.4 | febantel febantel/praziquantel/pyrantel febantel/praziquantel/pyrantel febantel/metrifonate febantel/praziquantel | | oral paste tablet tablet oral paste oral paste | expired expired expired expired expired |
| | ANADA INC. | | | | |
| Human: 02169649 02231509 02231510 02227347 02229102 01989987 02202603 02187051 02202611 02078597 02078600 02078619 | BETASERON - 0.3MG/VIAL CLIMARA 0.05 - 3.9MG/PATCH CLIMARA 0.1 - 7.8MG/PATCH LEVOVIST 999/1 LEVOVIST 999/1 MAGNEVIST - 469MG/ML OSMOVIST 240 - 512.59MG/ML OSMOVIST 280 - 598MG/ML OSMOVIST 280 - 598MG/ML ULTRAVIST 240 - 499MG/ML ULTRAVIST 300 - 623MG/ML ULTRAVIST 370 - 769MG/ML | interferon beta-1b estradiol 17b estradiol 17b galactose/palmitic acid gadopentetate dimeglumine iotrolan iotrolan iopromide iopromide iopromide | L03AA G03CA G03CA V08DA V08DA V08CA V08AB V08AB V08AB V08AB V08AB V08AB | powder for injectable solution transdermal patch transdermal patch powder for injectable solution powder for injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution | not sold not sold not sold |

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|--|---|---|--|--|--|
| BLOCK DF | RUG COMPANY (CANADA) LT | D. | | | |
| Human: 02178796 01945092 | COLPERMIN - 187MG/CAP SENSODYNE-F 5%/0.24% | peppermint oil potassium nitrate/sodium | A02DA A01AA | enteric-coated capsule toothpaste | |
| 02083310 | SENSODYNE-F 5%/0.24% | fluoride potassium nitrate/sodium | A01AA | toothpaste | introduced |
| 00624098 | SENSODYNE-F 5%/0.8% | fluoride w/baking soda potassium nitrate/sodium monofluorophosphate | A01AA | toothpaste | |
| | GER INGELHEIM (CANADA) L | .TD. | | | |
| Human: 00790486 02057735 02057743 | BRONALIDE - 0.25MG/DOSE PROSTEP 11 - 15MG/PATCH PROSTEP 22 - 30MG/PATCH | flunisolide nicotine nicotine | R03BA N07BA N07BA | aerosol for inhalation transdermal patch transdermal patch | expired |
| Veterinary: 02080001 00698229 02126214 | SEDIVET - 10MG/ML SPUTOLYSIN - 5MG/G SPUTOLYSIN - 5MG/ML SUPER-OV - 7.5UNIT/ML | romifidine dembrexin dembrexin follicle stimulating hormone (porcine origin) | | injectable solution oral powder injectable solution powder for injectable solution | expired expired expired |
| BOEHRIN | GER MANNHEIM CANADA LT | D. | | | |
| Human: 02232770 02129094 02129009 02129019 02129027 02129035 02231763 02231764 02231765 02231766 01927078 02233013 | BONDRONAT - 1MG/ML DEMADEX - 10MG/ML DEMADEX - 5MG/TAB DEMADEX - 10MG/TAB DEMADEX - 20MG/TAB DEMADEX - 100MG/TAB EUCARDIC - 3.125MG/TAB EUCARDIC - 6.25MG/TAB EUCARDIC - 12.5MG/TAB EUCARDIC - 25MG/TAB OSTAC - 400MG/CAP RETAVASE - 10.8UNIT/VIAL | ibandronate torsemide torsemide torsemide torsemide carvedilol carvedilol carvedilol carvedilol carvedilol clodronate disodium reteplase | M05BA C03CA C03CA C03CA C03CA C07AG C07AG C07AG C07AG C07AG M05BA B01AD | injectable solution injectable solution tablet tablet tablet tablet tablet tablet tablet tablet tablet capsule powder for injectable solution | not sold not sold not sold not sold not sold not sold not sold not sold not sold |
| | DIAGNOSTICS CANADA INC. | | | | |
| Human: | CHOLETEC - 45MG/VIAL | technetium tc-99m | V09DA | powder for injectable solution | |
| 02229056 | PROHANCE - 279.3MG/ML | mebrofenin gadoteridol | V08CA | injectable solution | |
| | MYERS SQUIBB PHARMACE | UTICAL GROUP | | | |
| Human: 01911570 01911562 01911554 00695661 00546283 00546291 00546305 00891843 02015099 00620998 00621005 00621013 | AZACTAM - 500MG/VIAL AZACTAM - 1000MG/VIAL AZACTAM - 2000MG/VIAL CAPOTEN - 12.5MG/TAB CAPOTEN - 25MG/TAB CAPOTEN - 50MG/TAB CAPOTEN - 100MG/TAB CAPOZIDE 25/12.5 CAPOZIDE 25/15 CAPOZIDE 25/25 CAPOZIDE 50/25 | aztreonam aztreonam aztreonam captopril captopril captopril captopril/hydrochlorothiazide captopril/hydrochlorothiazide captopril/hydrochlorothiazide captopril/hydrochlorothiazide captopril/hydrochlorothiazide | C09BA C09BA C09BA | powder for injectable solution powder for injectable solution powder for injectable solution tablet tablet tablet tablet tablet tablet tablet tablet tablet | not sold not sold not sold not sold not sold not sold not sold not sold |
| 02163675 02163683 02163659 02163667 00702277 00824135 02230986 | CARDIOTEC CEFZIL - 25MG/ML CEFZIL - 50MG/ML CEFZIL - 50MG/TAB CEFZIL - 500MG/TAB DESYREL - 150MG/TAB DESYREL - 300MG/TAB ETOPOPHOS - 100MG/VIAL | technetium tc-99m teboroxime cefprozil cefprozil cefprozil cefprozil trazodone hydrochloride trazodone hydrochloride etoposide phosphate | V09GA J01DA J01DA J01DA J01DA J01DA N06AX N06AX L01CB | powder for injectable solution powder for oral suspension powder for oral suspension tablet tablet tablet tablet powder for injectable solution | not sold not sold not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|--|---|--|--|
| 02168529 02168863 02163624 02163632 | MAXIPIME - 1000MG/BOTTLE MAXIPIME - 2000MG/BOTTLE MAXIPIME - 500MG/VIAL MAXIPIME - 1000MG/VIAL | cefepime hydrochloride cefepime hydrochloride cefepime hydrochloride cefepime hydrochloride | JO1DA J01DA J01DA J01DA J01DA | powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution | not sold not sold not sold |
| 02163640 01907093 01907107 01907115 | MAXIPIME - 2000MG/VIAL MONOPRIL - 5MG/TAB MONOPRIL - 10MG/TAB MONOPRIL - 20MG/TAB | cefepime hydrochloride fosinopril fosinopril fosinopril | J01DA C09AA C09AA C09AA | powder for injectable solution tablet tablet tablet | not sold |
| 00893749 00893757 02222051 02087294 | PRAVACHOL - 10MG/TAB PRAVACHOL - 20MG/TAB PRAVACHOL - 40MG/TAB SERZONE - 50MG/TAB | pravastatin sodium pravastatin sodium pravastatin sodium nefazodone hydrochloride | C10AA C10AA C10AA N06AX | tablet tablet tablet tablet | introduced |
| 02087375 02087383 02087391 | SERZONE - 100MG/TAB SERZONE - 150MG/TAB SERZONE - 200MG/TAB | nefazodone hýdrochloride nefazodone hydrochloride nefazodone hydrochloride | N06AX N06AX N06AX | tablet tablet tablet | |
| 02087405 02113031 01940511 01940538 | SERZONE - 300MG/TAB STADOL NS - 10MG/ML VIDEX - 25MG/TAB VIDEX - 50MG/TAB | nefazodone hydrochloride butorphanol tartrate didanosine didanosine | N06AX N02AF J05AB J05AB | tablet injectable solution tablet tablet | not sold |
| 01940546 01940554 01940589 | VIDEX - 100MG/TAB VIDEX - 150MG/TAB VIDEX - 100MG/VIAL | didanosine didanosine didanosine | J05AB J05AB J05AB | tablet tablet powder for oral solution | not sold |
| 01940597 01940600 01940619 01940627 | VIDEX - 167MG/VIAL VIDEX - 250MG/VIAL VIDEX - 375MG/VIAL VIDEX - 2000MG/VIAL | didanosine didanosine didanosine didanosine | J05AB J05AB J05AB J05AB | powder for oral solution powder for oral solution powder for oral solution powder for oral solution | not sold not sold not sold not sold |
| 01940635 02216078 02216086 02216094 02216108 | VIDEX - 4000MG/VIAL ZERIT - 5MG/CAP ZERIT - 15MG/CAP ZERIT - 20MG/CAP ZERIT - 30MG/CAP | didanosine stavudine stavudine stavudine stavudine | J05AB J05AX J05AX J05AX J05AX | powder for oral solution capsule capsule capsule capsule | not sold not sold |
| 02216116 Veterinary: 00673064 | ZERIT - 40MG/CAP TORBUTROL - 2MG/ML | stavudine butorphanol tartrate | J05AX | capsule injectable solution | not sold |
| | I PHARMA INC. | | | | 101 3010 |
| Human: 01945149 | CONDYLINE - 5MG/ML | podofilox | D11AF | topical solution | |
| | CORPORATION | | | | |
| Human: 01930605 | WINRHO - 0.12MG/VIAL | RHo(D) immune globulin (human) | J06BB | powder for injectable solution | not sold |
| 01919326 | WINRHO - 0.3MG/VIAL | RHo(D) immune globulin (human) | J06BB | powder for injectable solution | not sold |
| 02022842 | WINRHO SDF - 0.12MG/VIAL | RHo(D) immune globulin (human) | J06BB | powder for injectable solution | |
| 02022834 | WINRHO SDF - 0.3MG/VIAL | RHo(D) immune globulin (human) | J06BB | powder for injectable solution | |
| 02230397 | WINRHO SDF - 1MG/VIAL | RHo(D) immune globulin (human) | J06BB | powder for injectable solution | not sold |
| CARTER-I | IORNER INC. | | | | |
| Human: 00511641 | DEPEN - 250MG/TAB | penicillamine | M01CC | tablet | |
| CIBA-VISI | ON | | | | |
| Human: 02029901 | AQUASITE 2/1 | polyethylene glycol | S01XA | ophthalmic drops | |
| 02131625 01940414 | LIVOSTIN - 0.5MG/ML VOLTAREN OPHTHA - 1MG/ML | 400/dextran 70 levocabastine hydrochloride diclofenac sodium | S01GX S01BC | ophthalmic suspension ophthalmic drops | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|---|--|--|---|---|--|
| | ORAL PHARMACEUTICALS | | | | |
| Human: 02048213 02048221 02048248 | VIADENT - 0.3MG/ML VIADENT - 1.2MG/ML VIADENT FLUORIDE - 14MG/ML | sanguinarine sanguinarine sanguinarine | A01AD A01AD A01AD | oral rinse toothpaste toothpaste | not sold |
| CONNAUC | HT LABORATORIES LTD. | | | | |
| Human: 01905023 02231343 00764221 02230946 02224143 | MULTITEST CMI PENTACEL PROHIBIT QUADRACEL TRIPACEL | skin test antigens DaPT-IPV-Hib vaccine Hib vaccine DaPT-Hib vaccine DaPT vaccine | V04CX J07AG J07AJ J07AJ J07AJ | injectable suspension injectable suspension injectable suspension injectable suspension injectable suspension | introduced (nas) not sold introduced (nas) introduced (nas) |
| DERMIK L | ABORATORIES CANADA INC. | | | | |
| Human: 02225271 | BENZAMYCIN 50/30 | benzoyl peroxide/ erythromycin | D10AF | topical gel | |
| DISTA PR | ODUCTS LIMITED | | | | |
| Human: 01968041 01968033 | TALIDAN - 10MG/CAP TALIDAN - 20MG/CAP | fluoxetine hydrochloride fluoxetine hydrochloride | N06AB N06AB | capsule capsule | not sold not sold |
| | EALTH INC. | | | | |
| Human: 02123312 02123320 02123339 02123347 01927272 | ELDEPRYL - 5MG/TAB PERMAX - 0.05MG/TAB PERMAX - 0.25MG/TAB PERMAX - 1MG/TAB SD DEPRENYL - 5MG/TAB | selegiline hydrochloride pergolide mesylate pergolide mesylate pergolide mesylate selegiline hydrochloride | N04BD N04BC N04BC N04BC N04BC N04BD | tablet tablet tablet tablet tablet | not sold |
| DU PONT | MERCK PHARMA INC. | | | | |
| Human: 02211114 02028786 00870935 | BIANDA - 25MG/VIAL CARDIOLITE MIRALUMA NEUROLITE - 20MCI/TEST SINEMET CR 25/100 SINEMET CR 50/200 | losoxantrone hydrochloride technetium tc-99m sestamibi technetium tc-99m sestamibi technetium tc-99m bicisate carbidopa/levodopa carbidopa/levodopa | | powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution sustained-release tablet sustained-release tablet | n introduced |
| ELI LILLY | CANADA INC. | | | | |
| Human: 00778338 00778346 00465194 00465194 00465208 00832790 00465216 00832804 02161761 02161788 00548375 0229863 00555665 02218054 02230308 02230308 02230309 00015377 02229692 02229693 02229694 00745626 00889113 | AXID - 150MG/CAP AXID - 300MG/CAP CECLOR - 250MG/CAP CECLOR - 500MG/CAP CECLOR - 500MG/CAP CECLOR - 37.4MG/ML CECLOR - 37.4MG/ML CECLOR BID - 75MG/ML CECLOR CD - 375MG/TAB CECLOR CD - 375MG/TAB CECLOR CD - 500MG/TAB CECLOR CD - 500MG/TAB CECLOR CD - 500MG/TAB ELDISINE - 5MG/VIAL GEMZAR - 200MG/VIAL GEMZAR - 200MG/VIAL GEMZAR - 1000MG/VIAL GEMZAR - 1000MG/VIAL GLUCAGON - 1MG/ML HUMATROPE - 6.7MG/CARTRIDGE HUMATROPE - 5MG/VIAL HUMATROPE - 5MG/VIAL HUMATROPE - 5MG/VIAL | nizatidine nizatidine cefaclor cefaclor cefaclor cefaclor cefaclor cefaclor cefaclor cefaclor cefaclor dirithromycin vindesine sulfate amifostine gemcitabine hydrochloride gemcitabine hydrochloride glucagon hydrochloride somatropin somatropin somatropin insulin (regular/isophane) human biosynthetic | A02BA A02BA J01DA J01DA J01DA J01DA J01DA J01DA J01DA J01DA J01DA J01DA J01FA L01CA V03AF L01BC L01BC L01BC H04AA H01AC H01AC H01AC H01AC | capsule capsule capsule capsule powder for oral suspension powder for oral suspension powder for oral suspension powder for oral suspension sustained-release tablet sustained-release tablet capsule tablet powder for injectable solution powder for injectable solution | n n n expired n introduced n introduced n not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|---|---|----------------|--|----------------------|
| 01962639 | HUMULIN-10/90 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 00889105 | HUMULIN-20/80 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 01962655 | HUMULIN-20/80 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 00795879 | HUMULIN-30/70 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 01959212 | HUMULIN-30/70 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 00889091 | HUMULIN-40/60 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 01962647 | HUMULIN-40/60 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 00889121 | HUMULIN-50/50 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 01962663 | HUMULIN-50/50 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 00646148 | HUMULIN-L - 100UNIT/ML | insulin (lente) human biosynthetic | A10AC | injectable suspension | |
| 00587737 | HUMULIN-N - 100UNIT/ML | insulin (isophane) human biosynthetic | A10AC | injectable suspension | |
| 01959239 | HUMULIN-N - 100UNIT/ML | insulin (isophane) human biosynthetic | A10AC | injectable suspension | |
| 00586714 | HUMULIN-R - 100UNIT/ML | insulin (regular) human biosynthetic | A10AB | injectable solution | |
| 01959220 | HUMULIN-R - 100UNIT/ML | insulin (regular) human biosynthetic | A10AB | injectable solution | |
| 00733075 | HUMULIN-U - 100UNIT/ML | insulin (ultralente) human biosynthetic | A10AD | injectable suspension | |
| 00514535 | ILETIN-II LENTE PORK - 100UNIT/ML | pork insulin/zinc | A10AC | injectable suspension | |
| 00514551 00513644 | ILETIN-II NPH PORK - 100UNIT/ML ILETIN-II REGULAR PORK - 100UNIT/ML | pork insulin/zinc/protamine pork insulin/zinc | A10AC A10AB | injectable suspension injectable solution | |
| 00446580 00446572 | ILETIN-LENTE - 100UNIT/ML ILETIN-NPH - 100UNIT/ML | pork/bovine insulin/zinc pork/bovine insulin/zinc/ | A10AC A10AC | injectable suspension injectable suspension | |
| 00446610 | ILETIN-PROTAMINE ZINC - 100UNIT/ML | protamine pork/bovine insulin/zinc/protamine | A10AD | injectable suspension | not sold |
| 00446564 00446602 | ILETIN-REGULAR - 100UNIT/ML ILETIN-SEMILENTE - 100UNIT/ML | pork/bovine insulin/zinc pork/bovine insulin/zinc | A10AB A10AC | injectable solution injectable suspension | not sold |
| 00446599 00015202 | ILETIN-ULTRALENTE - 100UNIT/ML ILOSONE - 250MG/CAP | pork/bovine insulin/zinc | A10AD J01FA | injectable suspension capsule | not sold |
| 00015474 | ILOSONE - 25MG/ML | erythromycin estolate erythromycin estolate | J01FA | oral suspension | |
| 00210641 00244384 | ILOSONE - 50MG/ML ILOSONE - 500MG/TAB | erythromycin estolate | J01FA J01FA | oral suspension | not sold |
| 00244304 | KEFLIN - 1000MG/VIAL | erythromycin estolate cephalothin sodium | J01DA | tablet powder for injectable solution | expired |
| 00244406 | KEFLIN - 2000MG/VIAL | cephalothin sodium | J01DA | powder for injectable solution | expired |
| 00298484 | KEFLIN - 4000MG/VIAL | cephalothin sodium | J01DA | powder for injectable solution | expired |
| 00659150 | KEFLIN - 20000MG/VIAL | cephalothin sodium | J01DA | powder for injectable solution | expired |
| 00752525 | KEFLIN ADD-VANTAGE - 1000MG/VIAL | cephalothin sodium | J01DA | powder for injectable solution | not sold |
| 00752533 00322288 | KEFLIN ADD-VANTAGE - 2000MG/VIAL KEFZOL - 500MG/VIAL | cephalothin sodium cefazolin sodium | J01DA J01DA | powder for injectable solution powder for injectable solution | not sold |
| 00322200 | KEFZOL - 1000MG/VIAL | cefazolin sodium | J01DA | powder for injectable solution | |
| 00411450 | KEFZOL - 10000MG/VIAL | cefazolin sodium | J01DA | powder for injectable solution | |
| 00411434 | KEFZOL ADD-VANTAGE - 500MG/VIAL | cefazolin sodium | J01DA | powder for injectable solution | not sold |
| 00411442 | KEFZOL ADD-VANTAGE - 1000MG/VIAL | cefazolin sodium | J01DA | powder for injectable solution | |
| 02044757 02044749 | LORABID - 200MG/CAP LORABID - 400MG/CAP | loracarbef loracarbef | J01DA J01DA | capsule capsule | not sold not sold |
| 02161206 | LORABID - 20MG/ML | loracarbef | J01DA | powder for oral suspension | not sold |
| 02161214 | LORABID - 40MG/ML | loracarbef | J01DA | powder for oral suspension | not sold |
| 00439304 | MANDOL - 500MG/VIAL | cefamandole nafate | J01DA | powder for injectable solution | not sold |
| 00439320 00439312 | MANDOL - 1000MG/VIAL MANDOL - 2000MG/VIAL | cefamandole nafate cefamandole nafate | J01DA J01DA | powder for injectable solution powder for injectable solution | |
| 50 1000 TZ | | solumination manato | 50101 | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|---|---|---|--|
| 00648930 | MANDOL ADD-VANTAGE - 1000MG/VIAL | cefamandole nafate | J01DA | powder for injectable solution | not sold |
| 00842621 | MANDOL ADD-VANTAGE - 2000MG/VIAL | cefamandole nafate | J01DA | powder for injectable solution | not sold |
| 00569755 00569763 00851825 00611182 02018985 00636622 01917021 | MOXAM - 1000MG/VIAL MOXAM - 2000MG/VIAL NEBCIN ADD-VANTAGE - 10MG/ML ONCOVIN - 1MG/ML PROZAC - 10MG/CAP PROZAC - 20MG/CAP PROZAC - 4MG/ML | moxalactam disodium moxalactam disodium tobramycin sulfate vincristine sulfate fluoxetine hydrochloride fluoxetine hydrochloride fluoxetine hydrochloride | J01DA J01DA J01GB L01CA N06AB N06AB N06AB | powder for injectable solution powder for injectable solution injectable solution injectable solution capsule capsule oral solution | expired expired not sold |
| 02216973 00888338 00886971 00886955 00886963 00887129 | REOPRO - 2MG/ML TAZIDIME - 500MG/VIAL TAZIDIME - 1000MG/VIAL TAZIDIME - 2000MG/VIAL TAZIDIME - 6000MG/VIAL TAZIDIME ADD-VANTAGE - 1000MG/VIAL | abciximab ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate | B01AC J01DA J01DA J01DA J01DA J01DA | injectable solution powder for injectable solution | not sold |
| 01980645 00800430 00788716 02015110 00015423 00722146 00803510 00803537 | TAZIDIME ADD-VANTAGE - 2000MG/VIAL VANCOCIN - 125MG/CAP VANCOCIN - 250MG/CAP VANCOCIN - 10000MG/VIAL VANCOCIN C.P 500MG/VIAL VANCOCIN C.P 1000MG/VIAL VANCOCIN C.P. ADD-VANTAGE - 500MG/VIAL VANCOCIN C.P. ADD-VANTAGE - | ceftazidime pentahydrate vancomycin hydrochloride vancomycin hydrochloride vancomycin hydrochloride vancomycin hydrochloride vancomycin hydrochloride vancomycin hydrochloride | J01DA J01XA J01XA J01XA J01XA J01XA J01XA J01XA | powder for injectable solution capsule powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution | |
| 02229250 02229269 02229277 02229285 | 1000MG/VIAL ZYPREXA - 2.5MG/TAB ZYPREXA - 5MG/TAB ZYPREXA - 7.5MG/TAB ZYPREXA - 10MG/TAB | olanzapine olanzapine olanzapine olanzapine | N05AH N05AH N05AH N05AH N05AH | tablet tablet tablet tablet | expired expired expired expired |
| FABRIGEN | I INC. | | | | |
| Human: 02078465 02078473 02078503 02078643 02078627 02078635 02078651 02078481 | AVIRAX - 200MG/CAP AVIRAX - 50MG/G AVIRAX - 50MG/G AVIRAX - 40MG/ML AVIRAX - 200MG/TAB AVIRAX - 400MG/TAB AVIRAX - 800MG/TAB AVIRAX - 1000MG/VIAL | acyclovir acyclovir acyclovir acyclovir acyclovir acyclovir acyclovir acyclovir sodium | J05AB D06BB J05AB J05AB J05AB J05AB J05AB J05AB | capsule ointment cream oral suspension tablet tablet tablet powder for injectable solution | not sold not sold not sold not sold not sold |
| FAULDING | G (CANADA) INC. | | | | |
| Human: | ONCOSCINT CR/OV - 1MG/KIT | satumomab pendetide | V09IB | injectable solution | not sold |
| FERRING Human: | | · | | | |
| 02231047 00836362 00873993 00402516 00824305 00824143 02125226 | ACTHREL - 0.1MG/VIAL ACTHREL NF - 0.1MG/VIAL CERVIDIL 0.3 - 10MG/POUCH DDAVP - 0.01MG/DOSE DDAVP - 0.04MG/ML DDAVP - 0.1MG/ML DDAVP - 0.1MG/TAB DDAVP - 0.2MG/TAB NIDAGEL - 7.5MG/G | corticorelin ovine triflutate corticorelin ovine triflutate dinoprostone desmopressin acetate desmopressin acetate desmopressin acetate desmopressin acetate desmopressin acetate metronidazole | V04CD V04CD G02AD H01BA H01BA H01BA H01BA H01BA G01AF | powder for injectable solution powder for injectable solution vaginal insert nasal aerosol injectable solution nasal solution tablet tablet vaginal gel | not sold introduced introduced |
| 02024179 00780197 02153521 02153548 02153556 02153564 | OCTOSTIM - 0.15MG/DOSE OCTOSTIM - 0.015MG/ML OCTOSTIM - 1.5MG/ML QUINTASA - 1000MG/DOSE QUINTASA - 2000MG/DOSE QUINTASA - 4000MG/DOSE QUINTASA - 1000MG/SUP | desmopressin acetate desmopressin acetate desmopressin acetate 5-aminosalicylic acid 5-aminosalicylic acid 5-aminosalicylic acid 5-aminosalicylic acid | H01BA H01BA H01BA A07EC A07EC A07EC A07EC | nasal aerosol injectable solution nasal solution rectal suspension rectal suspension rectal suspension suppository | not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form Comments |
|--|---|---|---|--|
| FOURNIE | R PHARMA INC. | | | |
| Human: 02230283 02146959 | LIPIDIL MICRO - 67MG/CAP LIPIDIL MICRO - 200MG/CAP | fenofibrate fenofibrate | C10AB C10AB | capsule not sold capsule |
| FUJISAW | A CANADA INC. | | | |
| Human: 02139294 02175991 02175983 02176009 | ADENOSCAN - 3MG/ML AMBISOME - 50MG/VIAL PROGRAF - 1MG/CAP PROGRAF - 5MG/CAP PROGRAF - 5MG/ML | adenosine amphotericin b tacrolimus tacrolimus tacrolimus | C01EB J02AA L04AA L04AA L04AA | injectable solution powder for injectable solution capsule capsule injectable solution |
| GALDERM | IA CANADA INC. | | | |
| Human: 02148749 02231592 02148757 02092832 | DIFFERIN - 1MG/G DIFFERIN - 1MG/G DIFFERIN - 1MG/ML METROGEL - 7.5MG/G | adapalene adapalene adapalene metronidazole | D10AD D10AD D10AD D06BX | topical gel topical cream not sold topical solution not sold topical gel |
| GENDERN | I CANADA INC. | | | |
| Human: 02142147 00740306 02004240 | ZONALON - 50MG/G ZOSTRIX - 0.25MG/G ZOSTRIX-HP - 0.75MG/G | doxepin hydrochloride capsaicin capsaicin | D04AA M02AB M02AB | cream cream cream |
| | INSTITUTE, INC. | | | |
| Human: 02231018 | BENEFIX - 250UNIT/VIAL | coagulation factor IX | B02BD | powder for injectable solution introduced (nas) |
| 02231019 | BENEFIX - 500UNIT/VIAL | (recombinant) coagulation factor IX | B02BD | powder for injectable solution introduced (nas) |
| 02231020 | BENEFIX - 1000UNIT/VIAL | (recombinant) coagulation factor IX (recombinant) | B02BD | powder for injectable solution introduced (nas) |
| GLAXO W | ELLCOME INC. | | | |
| Human: 02192691 02192683 00828521 00828548 02215039 02215047 01943049 02145286 00886882 | 3TC - 10MG/ML 3TC - 150MG/TAB BECLODISK - 0.1MG/DOSE BECLOVENT - 0.2MG/DOSE BECLOVENT - 0.1MG/CAP BECLOVENT - 0.2MG/CAP CEFTIN - 25MG/ML CEFTIN - 250MG/POUCH CEFTIN - 125MG/TAB | lamivudine lamivudine beclomethasone dipropionate beclomethasone dipropionate beclomethasone dipropionate beclomethasone dipropionate cefuroxime axetil cefuroxime axetil cefuroxime axetil | R03BA R03BA J01DA J01DA J01DA J01DA | oral solution introduced (nas) tablet introduced (nas) powder for inhalation powder for inhalation powder for inhalation powder for oral suspension powder for oral suspension tablet not sold |
| 00886890 00886904 01974394 01974408 01974416 01968092 | CEFTIN - 250MG/TAB CEFTIN - 500MG/TAB CEPTAZ - 500MG/VIAL CEPTAZ - 1000MG/VIAL CEPTAZ - 2000MG/VIAL CEPTAZ - 10000MG/VIAL | cefuroxime axetil cefuroxime axetil ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate | J01DA J01DA J01DA J01DA J01DA J01DA | tablet tablet powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution |
| 01927175 02038269 01927183 02230845 02230848 02230857 02048043 02213583 02213591 02213621 02213648 02213605 02213613 | EXOSURF - 108MG/KIT EXOSURF - 67.5MG/VIAL EXOSURF - 108MG/VIAL FLOLAN - 0.5MG/VIAL FLOLAN - 1.5MG/VIAL FLOLAN STERILE DILUENT FLONASE - 0.05MG/DOSE FLOVENT - 0.05MG/DOSE FLOVENT - 0.05MG/DOSE FLOVENT - 0.1MG/DOSE FLOVENT - 0.125MG/DOSE FLOVENT - 0.25MG/DOSE FLOVENT - 0.25MG/DOSE | colfosceril palmitate colfosceril palmitate colfosceril palmitate epoprostenol sodium epoprostenol sodium fluticasone propionate fluticasone propionate fluticasone propionate fluticasone propionate fluticasone propionate fluticasone propionate fluticasone propionate fluticasone propionate | R07AA R07AA B01AC B01AC B01AC R01AD R03BA R03BA R03BA R03BA R03BA R03BA R03BA | endo-tracheal suspension kit endo-tracheal suspension vial not sold endo-tracheal suspension vial not sold powder for injectable solution powder for injectable solution powder for injectable solution powder for injectable solution nasal spray aerosol for inhalation powder for inhalation powder for inhalation not sold powder for inhalation powder for inhalation powder for inhalation aerosol for inhalation aerosol for inhalation |
| 02213656 02213664 | FLOVENT - 0.25MG/DOSE FLOVENT - 0.5MG/DOSE | fluticasone propionate fluticasone propionate | R03BA R03BA | powder for inhalation not sold powder for inhalation not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|--|--|----------------|---|------------|
| 00640026 | FORTAZ - 500MG/VIAL | ceftazidime pentahydrate | J01DA | powder for injectable solution | |
| 00640034 | FORTAZ - 1000MG/VIAL | ceftazidime pentahydrate | J01DA | powder for injectable solution | |
| 00640042 | FORTAZ - 2000MG/VIAL | ceftazidime pentahydrate | J01DA | powder for injectable solution | |
| 00791679 | FORTAZ - 6000MG/VIAL | ceftazidime pentahydrate | J01DA | powder for injectable solution | |
| 02230418 | IMITREX - 5MG/DOSE | sumatriptan hemisulphate | N02CC | nasal spray | introduced |
| 02230419 | IMITREX - 10MG/DOSE | sumatriptan hemisulphate | N02CC | nasal spray | not sold |
| 02230420 02212188 | IMITREX - 20MG/DOSE IMITREX - 12MG/ML | sumatriptan hemisulphate sumatriptan succinate | N02CC N02CC | nasal spray injectable solution | introduced |
| 02212100 | IMITREX - 50MG/TAB | sumatriptan succinate | N02CC | tablet | |
| 02212161 | IMITREX - 100MG/TAB | sumatriptan succinate | N02CC | tablet | |
| 02142082 | LAMICTAL - 25MG/TAB | lamotrigine | N03AX | tablet | |
| 02142090 | LAMICTAL - 50MG/TAB | lamotrigine | N03AX | tablet | not sold |
| 02142104 | LAMICTAL - 100MG/TAB | lamotrigine | N03AX | tablet | |
| 02142112 | LAMICTAL - 150MG/TAB | lamotrigine | N03AX | tablet | |
| 02142120 | LAMICTAL - 200MG/TAB | lamotrigine | NO3AX | tablet | not sold |
| 02142139 02217422 | LAMICTAL - 250MG/TAB MEPRON - 150MG/ML | lamotrigine atovaguone | N03AX P01XA | tablet oral suspension | not sold |
| 02009358 | MEPRON - 250MG/TAB | atovaquone | P01XA | tablet | |
| 02087308 | MIVACRON - 2MG/ML | mivacurium chloride | M03AC | injectable solution | |
| 02091283 | NAVELBINE - 10MG/ML | vinorelbine tartrate | L01CA | injectable solution | |
| 02229422 | NIMBEX - 2MG/ML | cisatracurium besylate | M03AC | injectable solution | expired |
| 02229423 | NIMBEX - 10MG/ML | cisatracurium besylate | M03AC | injectable solution | expired |
| 01924656 | NUROMAX - 1MG/ML | doxacurium chloride | M03AC | injectable solution | |
| 02231831 | PYLORID - 400MG/TAB | ranitidine bismuth citrate | A02BA | tablet | introduced |
| 01902660 01902644 | RETROVIR - 100MG/CAP RETROVIR - 10MG/ML | zidovudine zidovudine | J05AB J05AB | capsule injectable solution | |
| 01902652 | RETROVIR - 10MG/ML | zidovudine | JOSAB | Syrup | |
| 02136139 | SEREVENT - 0.025MG/DOSE | salmeterol xinafoate | R03AC | aerosol for inhalation | |
| 02136147 | SEREVENT - 0.05MG/DOSE | salmeterol xinafoate | R03AC | powder for inhalation | |
| 02231129 | SEREVENT DISKUS - 0.05MG/DOSE | salmeterol xinafoate | R03AC | powder for inhalation | not sold |
| 00675210 | TRACRIUM - 10MG/ML | atracurium besylate | M03AC | injectable solution | expired |
| 02230409 | ULTIVA - 1MG/VIAL | remifentanil hydrochloride | N01AH | powder for injectable solution | |
| 02230410 | ULTIVA - 2MG/VIAL | remifentanil hydrochloride | N01AH | powder for injectable solution | |
| 02230411 02219492 | ULTIVA - 5MG/VIAL VALTREX - 500MG/TAB | remifentanil hydrochloride | N01AH J05AB | powder for injectable solution tablet | |
| 02219492 | VENTODISK - 0.2MG/DOSE | valacyclovir hydrochloride salbutamol sulfate | R03AC | powder for inhalation | |
| 02215004 | VENTODISK - 0.4MG/DOSE | salbutamol sulfate | R03AC | powder for inhalation | |
| 02213215 | VENTOLIN - 0.2MG/CAP | salbutamol sulfate | R03AC | powder for inhalation | |
| 02212323 | VENTOLIN - 0.4MG/CAP | salbutamol sulfate | R03AC | powder for inhalation | |
| 02035421 | VENTOLIN - 0.4MG/ML | salbutamol sulfate | R03CC | oral solution | |
| 00782351 | VOLMAX - 4MG/TAB | salbutamol sulfate | R03CC | sustained-release tablet | expired |
| 00782378 01959077 | VOLMAX - 8MG/TAB WELLFERON - 3000000UNIT/ML | salbutamol sulfate | R03CC L03AA | sustained-release tablet | expired |
| 02161176 | WELLFERON - 5000000UNIT/ML | interferon alpha-n1 interferon alpha-n1 | LOSAA | injectable solution injectable solution | not sold |
| 01959069 | WELLFERON - 10000000UNIT/ML | interferon alpha-n1 | LOSAA | injectable solution | 1101 3010 |
| 00782386 | ZANTAC - 15MG/ML | ranitidine hydrochloride | A02BA | oral solution | |
| 00603791 | ZANTAC - 25MG/ML | ranitidine hydrochloride | A02BA | injectable solution | |
| 00553379 | ZANTAC - 150MG/TAB | ranitidine hydrochloride | A02BA | tablet | |
| 00641790 | ZANTAC - 300MG/TAB | ranitidine hydrochloride | A02BA | tablet | |
| 00849421 | ZANTAC C - 150MG/CAP | ranitidine hydrochloride | A02BA | capsule | |
| 00849448 01951831 | ZANTAC C - 300MG/CAP ZANTAC EFFERVESCENT - | ranitidine hydrochloride ranitidine hydrochloride | A02BA A02BA | capsule effervescent granules | not sold |
| 01901001 | 150MG/POUCH | | AUZDA | enervescent granules | 1101 5010 |
| 01951823 | ZANTAC EFFERVESCENT - 300MG/POUCH | ranitidine hydrochloride | A02BA | effervescent granules | not sold |
| 02076284 | ZANTAC EFFERVESCENT - 150MG/TAB | ranitidine hydrochloride | A02BA | effervescent tablet | not sold |
| 02076292 | ZANTAC EFFERVESCENT - 300MG/TAB | ranitidine hydrochloride | A02BA | effervescent tablet | not sold |
| 00577227 00497843 | ZINACEF - 250MG/VIAL ZINACEF - 750MG/VIAL | cefuroxime sodium cefuroxime sodium | J01DA J01DA | powder for injectable solution powder for injectable solution | not sold |
| 00497843 | ZINACEF - 750MG/VIAL ZINACEF - 1500MG/VIAL | cefuroxime sodium | J01DA J01DA | powder for injectable solution | |
| 00890936 | ZINACEF - 7500MG/VIAL | cefuroxime sodium | J01DA | powder for injectable solution | |
| 02229639 | ZOFRAN - 0.8MG/ML | ondansetron hydrochloride | A04AA | oral solution | |
| 01911821 | ZOFRAN - 2MG/ML | ondansetron hydrochloride | A04AA | injectable solution | |
| 02213567 | ZOFRAN - 4MG/TAB | ondansetron hydrochloride | A04AA | tablet | |
| 02213575 | ZOFRAN - 8MG/TAB | ondansetron hydrochloride | A04AA | tablet | |
| 00590924 00569771 | ZOVIRAX - 200MG/CAP ZOVIRAX - 50MG/G | acyclovir | J05AB D06BB | capsule ointment | |
| 00008111 | | acyclovir | סמטטט | Untillent | |
| | | | | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|---|--|----------------|--|--------------------|
| 02039524 | ZOVIRAX - 50MG/G | acyclovir | D06BB | cream | |
| 00886157 | ZOVIRAX - 40MG/ML | acyclovir | J05AB | oral suspension | |
| 00634506 | ZOVIRAX - 200MG/TAB | acyclovir | JOSAB | tablet | |
| 01911627 | ZOVIRAX - 400MG/TAB | acyclovir | J05AB | tablet | |
| 01911635 00605336 | ZOVIRAX - 800MG/TAB ZOVIRAX - 500MG/VIAL | acyclovir acyclovir sodium | J05AB J05AB | tablet powder for injectable solution | |
| 00899321 | ZOVIRAX - 1000MG/VIAL | acyclovir sodium | JOSAB | powder for injectable solution | |
| ПОЕСПОТ | MARION ROUSSEL CANADA | - | | | |
| Human: | MANION NOUSSEL GANADA | INC. | | | |
| 02231463 | ALLEGRA - 60MG/CAP | fexofenadine hydrochloride | R06AX | capsule | not sold |
| 02231462 | ALLEGRA - 60MG/TAB | fexofenadine hydrochloride | R06AX | | introduced (nas) |
| 02050943 02050951 | ALTACE - 1.25MG/CAP | ramipril | C09AA C09AA | capsule capsule | |
| 02050951 | ALTACE - 2.5MG/CAP ALTACE - 5MG/CAP | ramipril ramipril | CO9AA CO9AA | capsule | |
| 02050986 | ALTACE - 10MG/CAP | ramipril | C09AA | capsule | |
| 00863890 | ANANDRON - 50MG/TAB | nilutamide | L02BB | tablet | expired |
| 00863904 | ANANDRON - 100MG/TAB | nilutamide | L02BB | tablet | expired |
| 02231380 | ANZEMET - 20MG/ML | dolasetron mesylate | A04AA | | introduced (nas) |
| 02231378 | ANZEMET - 50MG/TAB | dolasetron mesylate | A04AA | | introduced (nas) |
| 02231379 | ANZEMET - 100MG/TAB | dolasetron mesylate | A04AA | | introduced (nas) |
| 00760439 | CITRUCEL - 166.66666MG/G | methylcellulose | A06AC | powder | not sold |
| 00546208 00546216 | CLAFORAN - 500MG/VIAL CLAFORAN - 1000MG/VIAL | cefotaxime sodium cefotaxime sodium | J01DA J01DA | powder for injectable solution powder for injectable solution | |
| 00546224 | CLAFORAN - 2000MG/VIAL | cefotaxime sodium | J01DA | powder for injectable solution | |
| 00839248 | CLAFORAN ADD-VANTAGE - | cefotaxime sodium | J01DA | powder for injectable solution | |
| 00839256 | 1000MG/VIAL CLAFORAN ADD-VANTAGE - | cefotaxime sodium | J01DA | powder for injectable solution | not sold |
| 02059754 | 2000MG/VIAL DERMATOP - 1MG/G | prednicarbate | D07AC | ointment | not sold |
| 02093111 | NICODERM 7 - 36MG/PATCH | nicotine | N07BA | transdermal patch | |
| 02093138 | NICODERM 14 - 78MG/PATCH | nicotine | N07BA | transdermal patch | |
| 02093146 | NICODERM 21 - 114MG/PATCH | nicotine | N07BA | transdermal patch | |
| 02179725 | ORELOX - 100MG/TAB | cefpodoxime proxetil | J01DA | tablet | not sold |
| 00894869 00894877 | PENTASA - 250MG/CAP PENTASA - 250MG/TAB | 5-aminosalicylic acid 5-aminosalicylic acid | A07EC A07EC | capsule tablet | not sold |
| 01940384 | PENTASA - 500MG/TAB | 5-aminosalicylic acid | A07EC | tablet | |
| 02069539 | RENEDIL - 2.5MG/TAB | felodipine | CO8CA | sustained-release tablet | |
| 01989618 | RENEDIL - 5MG/TAB | felodipine | C08CA | sustained-release tablet | |
| 01989596 | RENEDIL - 10MG/TAB | felodipine | C08CA | sustained-release tablet | |
| 00619760 | RYTHMODAN LA - 250MG/TAB | disopyramide | C01BA | sustained-release tablet | |
| 02068036 | SABRIL - 500MG/POUCH | vigabatrin | N03AG | powder for oral solution | expired |
| 02068028 | SABRIL - 1000MG/POUCH | vigabatrin | N03AG | powder for oral solution | expired |
| 02068001 02067994 | SABRIL - 2000MG/POUCH SABRIL - 3000MG/POUCH | vigabatrin vigabatrin | N03AG N03AG | powder for oral solution powder for oral solution | expired expired |
| 02065819 | SABRIL - 500MG/TAB | vigabatrin | N03AG | tablet | expired |
| 00590908 | SELDANE - 60MG/TAB | terfenadine | R06AX | tablet | onphiou |
| 00786624 | SELDANE - 120MG/TAB | terfenadine | R06AX | tablet | |
| 00680028 | SUPREFACT - 1MG/ML | buserelin acetate | L02AE | injectable solution | expired |
| 00680036 | SUPREFACT - 1MG/ML | buserelin acetate | L02AE | nasal solution | expired |
| 02228955 00870927 | SUPREFACT DEPOT - 6.6MG/VIAL | buserelin acetate | L02AE | injectable implant tablet | introduced |
| 00070927 | TERFENADINE - 60MG/TAB TERFENADINE - 120MG/TAB | terfenadine terfenadine | R06AX R06AX | tablet | |
| 00586625 | TRENTAL - 400MG/TAB | pentoxifylline | C04AD | sustained-release tablet | expired |
| | NN-LA ROCHE LIMITED | | | | |
| Human: 02162725 | ANAPROX - 275MG/TAB | nanrovan addium | M01AE | tablat | |
| 02162725 | ANAPROX - 275MG/TAB ANAPROX DS - 550MG/TAB | naproxen sodium naproxen sodium | M01AE | tablet tablet | |
| 02102717 | ANAFROX DS - 550MG/TAB ANEXATE - 0.1MG/ML | flumazenil | V03AB | injectable solution | |
| 00550078 | BACTRIM 16/80 | trimethoprim/ | JOIEE | injectable solution | |
| 50000070 | 2.1011111110/00 | sulfamethoxazole | 00122 | jootable contion | |
| 00272477 | BACTRIM 20/100 | trimethoprim/ | J01EE | tablet | not sold |
| 00070405 | | sulfamethoxazole | | | |
| 00272485 | BACTRIM 8/40 | trimethoprim/ sulfamethoxazole | J01EE | oral suspension | |
| 00272469 | BACTRIM 80/400 | trimethoprim/ | J01EE | tablet | |
| | | sulfamethoxazole | | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|--|--|----------------|--|------------------------|
| 00371823 | BACTRIM DS 160/800 | trimethoprim/ sulfamethoxazole | J01EE | tablet | |
| 02162741 | CARDENE - 20MG/CAP | nicardipine hydrochloride | C08CA | capsule | |
| 02162733 | CARDENE - 30MG/CAP | nicardipine hydrochloride | C08CA | capsule | |
| 02143348 | CARDENE - 2.5MG/ML | nicardipine hydrochloride | C08CA | injectable solution | not sold |
| 02034514 | CARDENE SR - 30MG/CAP | nicardipine hydrochloride | C08CA | sustained-release capsule | not sold |
| 01950657 | CARDENE SR - 45MG/CAP | nicardipine hydrochloride | CO8CA | sustained-release capsule | not sold |
| 01950649 | CARDENE SR - 60MG/CAP | nicardipine hydrochloride | C08CA | sustained-release capsule | not sold |
| 02192748 02186802 | CELLCEPT - 250MG/CAP | mycophenolate mofetil | LO4AA | capsule | |
| 02160602 | CYTOVENE - 250MG/CAP CYTOVENE - 500MG/VIAL | ganciclovir ganciclovir sodium | J05AB J05AB | capsule powder for injectable solution | |
| 00692719 | FANSIDAR 500/25 | sulfadoxine/pyrimethamine | P01BD | tablet | |
| 00776173 | FEMSTAT - 20MG/G | butoconazole nitrate | G01AF | cream | expired |
| 00890987 | FEMSTAT - 20MG/G | butoconazole nitrate | G01AF | cream | expired |
| 00890979 | FEMSTAT - 100MG/SUP | butoconazole nitrate | G01AF | suppository | expired |
| | GARDRIN - 0.035MG/CAP | enprostil | A02BB | capsule | not sold |
| 01990918 | HIVID - 0.375MG/TAB | zalcitabine | J05AB | tablet | |
| 01990896 | HIVID - 0.75MG/TAB | zalcitabine | J05AB G01AF | tablet | ownired |
| 01951874 | IN-EX DUAL PACK | sulconazole and butoconazole nitrate | GUTAF | topical and vaginal cream | expired |
| 01955535 | IN-EX DUAL PACK | sulconazole and | G01AF | topical cream and vaginal ovule | expired |
| 01000000 | | butoconazole nitrate | GOIN | topical creatil and vaginal ovulo | CAPITOU |
| 01909959 | INHIBACE - 0.5MG/TAB | cilazapril | C09AA | tablet | not sold |
| 01911465 | INHIBACE - 1MG/TAB | cilazapril | C09AA | tablet | |
| 01911473 | INHIBACE - 2.5MG/TAB | cilazapril | C09AA | tablet | |
| 01911481 | INHIBACE - 5MG/TAB | cilazapril | C09AA | tablet | |
| 02181479 | INHIBACE PLUS 5/12.5 | cilazapril/hydrochlorothiazide | | tablet | not sold |
| 02216965 | INVIRASE - 200MG/CAP | saquinavir mesylate | J05AE | | roduced (nas) |
| 00899348 00899356 | MANERIX - 100MG/TAB MANERIX - 150MG/TAB | moclobemide | N06AG N06AG | tablet tablet | |
| 02166747 | MANERIX - 300MG/TAB | moclobernide | N06AG | tablet | |
| 02227371 | MEGALONE - 4MG/ML | fleroxacin | J01MA | injectable solution | not sold |
| 02227398 | MEGALONE - 200MG/TAB | fleroxacin | J01MA | tablet | not sold |
| 02227401 | MEGALONE - 400MG/TAB | fleroxacin | J01MA | tablet | not sold |
| 02162431 | NAPROSYN - 25MG/ML | naproxen | M01AE | oral suspension | |
| 02162458 | NAPROSYN - 500MG/SUP | naproxen | M01AE | suppository | not cold |
| 00299413 02162474 | NAPROSYN - 125MG/TAB NAPROSYN - 250MG/TAB | naproxen | M01AE M01AE | tablet tablet | not sold |
| 02162482 | NAPROSYN - 375MG/TAB | naproxen naproxen | M01AE | tablet | |
| 02162490 | NAPROSYN - 500MG/TAB | naproxen | M01AE | tablet | |
| 02162792 | NAPROSYN E - 250MG/TAB | naproxen | M01AE | tablet | |
| 02162415 | NAPROSYN E - 375MG/TAB | naproxen | M01AE | tablet | |
| 02162423 | NAPROSYN E - 500MG/TAB | naproxen | M01AE | tablet | |
| 02162466 | NAPROSYN SR - 750MG/TAB | naproxen | M01AE | sustained-release tablet | أدا محاجر |
| 02168871 02216183 | NAPROSYN SR - 1000MG/TAB NUTROPIN - 5MG/VIAL | naproxen | M01AE H01AC | sustained-release tablet powder for injectable solution | not sold introduced |
| 02216191 | NUTROPIN - 10MG/VIAL | somatropin somatropin | H01AC | powder for injectable solution | not sold |
| 02229722 | NUTROPIN AQ - 5MG/ML | somatropin | H01AC | injectable solution | introduced |
| 02204584 | PROTROPIN - 5MG/VIAL | somatrem | H01AC | powder for injectable solution | |
| 02204576 | PROTROPIN - 10MG/VIAL | somatrem | H01AC | powder for injectable solution | |
| 02162687 | RHINALAR - 0.25MG/ML | flunisolide | R01AD | nasal aerosol | |
| 00481823 | ROCALTROL - 0.00025MG/CAP | calcitriol | A11CC | capsule | |
| 00481815 00824291 | ROCALTROL - 0.0005MG/CAP ROCALTROL - 0.001MG/ML | calcitriol calcitriol | A11CC A11CC | capsule oral solution | |
| 00624291 | ROCEPHIN - 250MG/VIAL | ceftriaxone disodium | J01DA | powder for injectable solution | |
| 00657425 | ROCEPHIN - 500MG/VIAL | ceftriaxone disodium | J01DA | powder for injectable solution | not sold |
| 00657417 | ROCEPHIN - 1000MG/VIAL | ceftriaxone disodium | J01DA | powder for injectable solution | not oolu |
| 00657409 | ROCEPHIN - 2000MG/VIAL | ceftriaxone disodium | J01DA | powder for injectable solution | |
| 00851957 | ROCEPHIN - 10000MG/VIAL | ceftriaxone disodium | J01DA | powder for injectable solution | |
| 00894699 | ROCEPHIN ADD-VANTAGE - | ceftriaxone disodium | J01DA | powder for injectable solution | |
| 00894702 | 1000MG/VIAL ROCEPHIN ADD-VANTAGE - | ceftriaxone disodium | J01DA | powder for injectable solution | not sold |
| | 2000MG/VIAL | | | | |
| 01911988 | ROFERON-A - 3000000UNIT/VIAL | interferon alfa-2a | LO3AA | powder for injectable solution | not sold |
| 02217015 02217023 | ROFERON-A - 3000000UNIT/VIAL ROFERON-A - 4500000UNIT/VIAL | interferon alfa-2a interferon alfa-2a | L03AA L03AA | injectable solution injectable solution | introduced |
| 02217023 | ROFERON-A - 6000000UNIT/VIAL | interferon alfa-2a | LOSAA | injectable solution | introduced |
| 01911996 | ROFERON-A - 9000000UNIT/VIAL | interferon alfa-2a | LOSAA | powder for injectable solution | not sold |
| 02217058 | ROFERON-A - 9000000UNIT/VIAL | interferon alfa-2a | LO3AA | injectable solution | |
| | | | | | |

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|--|---|--|---|---|--|
| 00812471 01912003 02217066 00891002 | ROFERON-A - 1800000UNIT/VIAL ROFERON-A - 1800000UNIT/VIAL ROFERON-A - 1800000UNIT/VIAL ROFERON-A - 3600000UNIT/VIAL | interferon alfa-2a interferon alfa-2a | L03AA L03AA L03AA L03AA | powder for injectable solution powder for injectable solution injectable solution injectable solution | n not sold n not sold |
| 02235914 | TASMAR - 100MG/TAB | tolcapone | N04BX | tablet | introduced (nas) |
| 02235921 01911767 | TASMAR - 200MG/TAB TICLID - 125MG/TAB | tolcapone ticlopidine hydrochloride | N04BX B01AC | tablet tablet | introduced (nas) not sold |
| 02162776 | TICLID - 250MG/TAB | ticlopidine hydrochloride | B01AC | tablet | not solu |
| 00884499 02162644 | TORADOL - 5MG/ML TORADOL - 10MG/ML | ketorolac tromethamine ketorolac tromethamine | S01BC M01AB | ophthalmic solution injectable solution | not sold |
| 01908499 | TORADOL - 15MG/ML | ketorolac tromethamine | M01AB | injectable solution | not sold |
| 02162652 02162660 | TORADOL - 30MG/ML TORADOL - 10MG/TAB | ketorolac tromethamine ketorolac tromethamine | M01AB M01AB | injectable solution tablet | |
| 00784516 | VERSED - 1MG/ML | midazolam hydrochloride | N05CD | injectable solution | |
| 00766011 | VERSED - 5MG/ML | midazolam hydrochloride | N05CD | injectable solution | |
| Veterinary: | | ablartatragualina | | oral powdor | not cold |
| | AUREO S-700 - 154000MG/KG | chlortetracycline hydrochloride/sulfamethazin | е | oral powder | not sold |
| 02230809 | AUREO S-700G - 154000MG/KG | chlortetracycline hydrochloride/sulfamethazin | e | feed premix | |
| 02230721 | AUREO SP250 G | chlortetracycline hydrochl./sulfamethazine/ penicillin g proc. | | feed premix | |
| | AUREOMIX 500G | chlortetracycline hydrochl./ sulfamethazine/ penicillin g proc. | | feed premix | not sold |
| 02230722 | AUREOMIX 625G | chlortetracycline hydrochl./ sulfamethazine/ | | feed premix | |
| 02230724 | AUREOMYCIN 100G - 220000MG/KG | penicillin g proc. chlortetracycline hydrochloride | | feed premix | |
| 02230723 02230778 | AUREOMYCIN 50G - 110000MG/KG CYGRO - 10000MG/KG | chlortetracycline hydrochlorid maduramicin ammonium | le | feed premix oral powder | |
| ICN CANA | DA LTD. | | | | |
| Human: 00704008 | VIRAZOLE - 6000MG/VIAL | ribavirin | J05AB | powder for inhalation | |
| JANSSEN- | ORTHO INC. | | | | |
| Human: 00755818 01968440 01992872 01937383 01937391 01937405 01937413 | ALFENTA - 0.5MG/ML CYCLEN 0.25/0.035 CYCLEN 0.25/0.035 DURAGESIC - 2.5MG/PATCH DURAGESIC - 5MG/PATCH DURAGESIC - 7.5MG/PATCH DURAGESIC - 10MG/PATCH | alfentanil hydrochloride norgestimate/ethinyl estradiol norgestimate/ethinyl estradiol fentanyl fentanyl fentanyl fentanyl | | injectable solution tablet transdermal patch transdermal patch transdermal patch transdermal patch | expired expired |
| 00590665 02126575 02126583 02126591 02206072 02231583 | DURALITH - 300MG/TAB EPREX - 2000UNIT/ML EPREX - 4000UNIT/ML EPREX - 10000UNIT/ML EPREX - 20000UNIT/ML EPREX - 1000UNIT/SYRINGE | lithium carbonate epoetin alfa epoetin alfa epoetin alfa epoetin alfa epoetin alfa | N05AN B03XA B03XA B03XA B03XA B03XA | sustained-release tablet injectable solution injectable solution injectable solution injectable solution injectable solution | introduced (nas) introduced (nas) introduced (nas) introduced (nas) introduced |
| 02231584 02231585 02231586 02231587 02049430 02049414 02049422 | EPREX - 2000UNIT/SYRINGE EPREX - 3000UNIT/SYRINGE EPREX - 4000UNIT/SYRINGE EPREX - 10000UNIT/SYRINGE FLOXIN - 4MG/ML FLOXIN - 200MG/ML FLOXIN - 40MG/ML | epoetin alfa epoetin alfa epoetin alfa epoetin alfa ofloxacin ofloxacin ofloxacin | B03XA B03XA B03XA B03XA J01MA J01MA | injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution | introduced introduced introduced introduced not sold not sold not sold |
| 01968424 01968416 01968408 02022117 02236839 02236840 02236841 | FLOXIN - 200MG/TAB FLOXIN - 300MG/TAB FLOXIN - 400MG/TAB LEUSTATIN - 1MG/ML LEVAQUIN - 5MG/ML LEVAQUIN - 25MG/ML LEVAQUIN - 250MG/TAB | ofloxacin ofloxacin cladribine levofloxacin levofloxacin levofloxacin | J01MA J01MA J01MA L01BB J01MA J01MA J01MA | tablet tablet injectable solution injectable solution injectable solution tablet | not sold not sold not sold |

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|----------------------|---|---|----------------|----------------------------|------------------|
| 02236842 02020017 | LEVAQUIN - 500MG/TAB LIVOSTIN - 0.5MG/ML | levofloxacin levocabastine hydrochloride | J01MA R01AC | tablet nasal suspension | not sold |
| 00642851 | MOTILIUM - 10MG/TAB | domperidone | A03FA | tablet | expired |
| 00855820 | MOTILIUM - 10MG/TAB | domperidone maleate | A03FA | tablet | expired |
| 00703974 | NIZORAL - 20MG/G | ketoconazole | D01AC | cream | onpriou |
| 00788813 | NIZORAL - 20MG/ML | ketoconazole | J02AB | oral suspension | |
| 00633836 | NIZORAL - 200MG/TAB | ketoconazole | J02AB | tablet | |
| 00602957 | ORTHO 7/7/7 .575-1/.035 | norethindrone/ethinyl estradiol | G03AB | tablet | |
| 00602965 | ORTHO 7/7/7 .575-1/.035 | norethindrone/ethinyl estradiol | G03AB | tablet | |
| 02042541 | ORTHO-CEPT 0.15/0.03 | desogestrel/ethinyl estradiol | G03AA | tablet | expired |
| 02042533 | ORTHO-CEPT 0.15/0.03 | desogestrel/ethinyl estradiol | G03AA | tablet | expired |
| 02015978 | ORTHOCLONE-OKT3 - 1MG/ML | muromonab-cd3 | L04AA | injectable solution | |
| 00836354 | PREPULSID - 1MG/ML | cisapride monohydrate | A03FA | oral suspension | |
| 00836311 | PREPULSID - 5MG/TAB | cisapride monohydrate | A03FA | tablet | |
| 00836338 | PREPULSID - 10MG/TAB | cisapride monohydrate | A03FA | tablet | |
| 02054817 | PREPULSID - 20MG/TAB | cisapride monohydrate | A03FA | tablet | |
| 02236435 | PREPULSID QS - 5MG/TAB | cisapride monohydrate | A03FA | tablet | not sold |
| 02236441 | PREPULSID QS - 10MG/TAB | cisapride monohydrate | A03FA | tablet | not sold |
| 02236443 | PREPULSID QS - 20MG/TAB | cisapride monohydrate | A03FA | tablet | not sold |
| 02052407 | RENOVA - 0.5MG/G | tretinoin | D10AD | cream | not cold |
| 02236950 | RISPERDAL - 1MG/ML | risperidone | N05AX | oral solution | not sold |
| 02025280 | RISPERDAL - 1MG/TAB | risperidone | N05AX | tablet | |
| 02025299 02025302 | RISPERDAL - 2MG/TAB | risperidone | N05AX N05AX | tablet tablet | |
| 02025302 | RISPERDAL - 3MG/TAB RISPERDAL - 4MG/TAB | risperidone risperidone | N05AX N05AX | tablet | |
| 02025310 | SPORANOX - 100MG/CAP | itraconazole | J02AC | capsule | |
| 02047454 | SPORANOX - 100MG/CAP | itraconazole | JOZAC JOZAC | oral solution | introduced |
| 01951319 | SUFENTA - 0.05MG/ML | sufentanil citrate | N01AH | injectable solution | expired |
| 01934155 | TERAZOL 3 - 8MG/G | terconazole | G01AG | vaginal cream | expireu |
| 00894710 | TERAZOL 3 - 80MG/SUP | terconazole | G01AG | vaginal suppository | |
| 02130874 | TERAZOL 3 DUALPAK | terconazole | G01AG | vaginal cream and supposit | orv |
| 00894729 | TERAZOL 7 - 4MG/G | terconazole | G01AG | vaginal cream | ory |
| 00484938 | TOLECTIN - 400MG/CAP | tolmetin sodium | M01AB | capsule | |
| 00364126 | TOLECTIN - 200MG/TAB | tolmetin sodium | M01AB | tablet | |
| 00632740 | TOLECTIN - 600MG/TAB | tolmetin sodium | M01AB | tablet | |
| 02230893 | TOPAMAX - 25MG/TAB | topiramate | N03AX | tablet | introduced (nas) |
| 02230894 | TOPAMAX - 100MG/TAB | topiramate | N03AX | tablet | introduced (nas) |
| 02230896 | TOPAMAX - 200MG/TAB | topiramate | N03AX | tablet | introduced (nas) |
| 02028700 | TRI-CYCLEN .1821525/.035 | norgestimate/ethinyl estradiol | | tablet | |
| 02029421 | TRI-CYCLEN .1821525/.035 | norgestimate/ethinyl estradiol | | tablet | |
| Veterinary: 00788724 | APPERTEX - 2.5MG/TAB | clazuril | | tablet | not sold |
| 00731374 | WILDNIL - 3MG/ML | carfentanil citrate | | injectable solution | expired |
| JOHNSON Human: | & JOHNSON-MERCK | | | | |
| 02182912 | HISMANAL - 10MG/TAB | astemizole | R06AX | tablet | |
| 02185911 | PEPCID AC - 10MG/TAB | famotidine | A02BA | chewable tablet | introduced |
| 02185938 | PEPCID AC - 10MG/TAB | famotidine | A02BA | tablet | |
| JOUVEINA | AL INC. | | | | |
| Human: | | | | | |
| muniun. | CERICLAMINE - 100MG/CAP | cericlamine | N06AB | capsule | not sold |
| | CERICLAMINE - 150MG/CAP | cericlamine | N06AB | capsule | not sold |
| KNOLL PH | IARMA INC. | | | | |
| Human: | | | | | |
| 02231457 | MAVIK - 0.5MG/CAP | trandolapril | C09AA | capsule | not sold |
| 02231459 | MAVIK - 1MG/CAP | trandolapril | C09AA | capsule | not sold |
| 02231460 | MAVIK - 2MG/CAP | trandolapril | CO9AA | capsule | not sold |
| | | | | - stere er e | |
| | | | | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|--|---|--|--|
| LEO LABO | ORATORIES CANADA LTD. | | | | |
| Human: 01976133 02150956 02194341 02167840 02229755 02167859 02229515 02231478 00474517 00474525 00759546 | DOVONEX - 0.05MG/G DOVONEX - 0.05MG/G DOVONEX - 0.05MG/ML INNOHEP - 10000UNIT/ML INNOHEP - 10000UNIT/ML INNOHEP - 20000UNIT/ML INNOHEP - 20000UNIT/ML ONE-ALPHA - 0.00025MG/CAP ONE-ALPHA - 0.001MG/CAP ONE-ALPHA - 0.0002MG/ML | calcipotriol calcipotriol calcipotriol tinzaparin sodium tinzaparin sodium tinzaparin sodium tinzaparin sodium alfacalcidol alfacalcidol alfacalcidol | D05AX D05AX B01AB B01AB B01AB B01AB B01AB B01AB A11CC A11CC A11CC | ointment cream scalp lotion injectable suspension injectable suspension injectable suspension injectable suspension capsule capsule oral solution | introduced introduced not sold |
| LIGAND F | PHARMACEUTICALS INC. | | | | |
| Human: 02020033 02019876 02130181 | PHOTOFRIN - 15MG/VIAL PHOTOFRIN - 75MG/VIAL PROLEUKIN - 22000000UNIT/VIAL | porfimer sodium porfimer sodium aldesleukin | L01XX L01XX L03AA | powder for injectable solutior powder for injectable solutior powder for injectable solutior | 1 |
| LIPOSOM Human: | E COMPANY, INC. (THE) | | | | |
| 02231590 | ABELCET - 5MG/ML | amphotericin b lipid complex | J02AA | powder for injectable solution | ı |
| | KRODT MEDICAL INC. | | | | |
| Human: | 99MTC TECHNESCAN | technetium tc-99m mertiatide | V09CA | powder for injectable solutior | n not sold |
| | MAG3 UD PED. 99MTC TECHNESCAN | technetium tc-99m mertiatide | V09CA | powder for injectable solutior | 1 |
| 00727725 | MAG3 UD REG. HEXABRIX 160 - 294.5MG/ML | meglumine and sodium | V08AB | injectable solution | expired |
| 00788805 | HEXABRIX 200 - 369MG/ML | ioxaglate meglumine and sodium | V08AB | injectable solution | expired |
| 00603740 | HEXABRIX 320 - 589MG/ML | ioxaglate meglumine and sodium | V08AB | injectable solution | expired |
| 01900838 01900846 02034492 01900854 02034484 | OPTIRAY 160 - 339MG/ML OPTIRAY 240 - 509MG/ML OPTIRAY 300 - 636MG/ML OPTIRAY 320 - 678MG/ML OPTIRAY 350 - 742MG/ML TECHNESCAN MAG3 | ioxaglate ioversol ioversol ioversol ioversol ioversol technetium tc-99m mertiatide | V08AB V08AB V08AB V08AB V08AB V08AB | injectable solution injectable solution injectable solution injectable solution injectable solution powder for injectable solutior | not sold not sold |
| _ | KRODT VETERINARY INC. | | | | |
| Veterinary: 01950576 00670898 00673056 | CLINACOX - 5000MG/KG ESTRUMATE - 0.25MG/ML PLANATE - 0.0875MG/ML | diclazuril cloprostenol sodium cloprostenol sodium | | feed premix injectable solution injectable solution | not sold expired expired |
| | ROSST CANADA INC. | | | | |
| Human: 02182815 02182874 02182882 02229153 02229161 02229188 02229196 00587699 00576131 0223055 02201011 0223055 02201011 02201038 00568368 02230047 | COZAAR - 25MG/TAB COZAAR - 50MG/TAB COZAAR - 100MG/TAB CRIXIVAN - 100MG/CAP CRIXIVAN - 200MG/CAP CRIXIVAN - 300MG/CAP DOLOBID - 250MG/TAB DOLOBID - 500MG/TAB FOSAMAX - 5MG/TAB FOSAMAX - 10MG/TAB FOSAMAX - 40MG/TAB HEPTAVAX-B HYZAAR 50/12.5 | losartan potassium losartan potassium losartan potassium indinavir sulfate indinavir sulfate indinavir sulfate diflunisal diflunisal alendronate sodium alendronate sodium alendronate sodium vaccine - hepatitis B losartan potassium/ hydrochlorothiazide | C09CA C09CA J05AE J05AE J05AE J05AE N02BA N02BA M05BA M05BA M05BA J07BC C09DA | tablet tablet tablet capsule capsule capsule tablet tablet tablet tablet tablet tablet tablet tablet tablet tablet | not sold not sold introduced (nas) not sold introduced (nas) introduced not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|---|--|----------------|--|----------------------|
| 00463248 00466085 | INDOCID-SR - 75MG/CAP M-M-R II | indomethacin vaccine - measles/mumps/ rubella | M01AB J07BD | sustained-release capsule injectable suspension | |
| 00893668 | MEFOXIN ADD-VANTAGE - 1000MG/VIAL | cefoxitin sodium | J01DA | powder for injectable solution | |
| 00893676 | MEFOXIN ADD-VANTAGE - 2000MG/VIAL | cefoxitin sodium | J01DA | powder for injectable solution | |
| 00795844 00795860 | MEVACOR - 10MG/TAB MEVACOR - 20MG/TAB | lovastatin lovastatin | C10AA C10AA | tablet tablet | not sold |
| 00795852 | MEVACOR - 40MG/TAB | lovastatin | C10AA | tablet | |
| 01908294 | NOROXIN - 3MG/ML | norfloxacin | S01AX | ophthalmic solution | |
| 00643025 01942948 | NOROXIN - 400MG/TAB PEDVAXHIB | norfloxacin vaccine - hemophilus | J01MA J07AG | tablet injectable suspension | |
| 01342340 | | influenzae b | 307 AU | injectable suspension | |
| 00728101 | PEPCID - 8MG/ML | famotidine | A02BA | oral suspension | not sold |
| 00728128 00710121 | PEPCID - 10MG/ML PEPCID - 20MG/TAB | famotidine famotidine | A02BA A02BA | injectable solution tablet | |
| 00710121 | PEPCID - 40MG/TAB | famotidine | A02BA | tablet | |
| 00431648 | PNEUMOVAX 23 | vaccine - polyvalent | J07AL | injectable suspension | |
| 00717274 | | pneumoccocal | J01DH | powder for injectable colution | |
| 00717274 | PRIMAXIN 250/250 PRIMAXIN 250/250 ADD-VANTAGE | imipenem/cilastatin sodium | J01DH J01DH | powder for injectable solution powder for injectable solution | |
| 00717282 | PRIMAXIN 500/500 | imipenem/cilastatin sodium | J01DH | powder for injectable solution | |
| 01911449 | PRIMAXIN 500/500 ADD-VANTAGE | imipenem/cilastatin sodium | J01DH | powder for injectable solution | |
| 00890952 00893684 | PRIMAXIN IM 500/500 PRIMAXIN IM 750/750 | imipenem/cilastatin sodium imipenem/cilastatin sodium | J01DH J01DH | powder for injectable solution powder for injectable solution | not sold not sold |
| 00839388 | PRINIVIL - 5MG/TAB | lisinopril | C09AA | tablet | 1101 3010 |
| 00839396 | PRINIVIL - 10MG/TAB | lisinopril | C09AA | tablet | |
| 00839418 | PRINIVIL - 20MG/TAB | lisinopril | CO9AA | tablet | |
| 00839426 00839434 | PRINIVIL - 40MG/TAB PRINIVIL - 80MG/TAB | lisinopril lisinopril | CO9AA CO9AA | tablet tablet | not sold not sold |
| 02108194 | PRINZIDE 10/12.5 | lisinopril/hydrochlorothiazide | | tablet | 1101 3010 |
| 00884413 | PRINZIDE 20/12.5 | lisinopril/hydrochlorothiazide | C09BA | tablet | |
| 00884421 02010909 | PRINZIDE 20/25 | lisinopril/hydrochlorothiazide finasteride | CO9BA GO4BX | tablet tablet | |
| 02010909 | PROSCAR - 5MG/TAB RECOMBIVAX HB - 0.01MG/ML | vaccine - hepatitis B(rDNA) | J07BC | injectable suspension | |
| 01934023 | RECOMBIVAX HB - 0.04MG/ML | vaccine - hepatitis B(rDNA) | J07BC | injectable suspension | |
| 02171880 | TIMOPTIC XE - 2.5MG/ML | timolol maleate | S01ED | ophthalmic gel | |
| 02171899 02216205 | TIMOPTIC XE - 5MG/ML TRUSOPT - 20MG/ML | timolol maleate dorzolamide hydrochloride | S01ED S01EC | ophthalmic gel ophthalmic solution | |
| 02229702 | VAQTA - 50UNIT/ML | hepatitis A vaccine | J07BC | injectable suspension | |
| | | (inactivated) | | | |
| 00657298 | VASERETIC 10/25 | enalapril maleate/ hydrochlorothiazide | C09BA | tablet | |
| 00851795 | VASOTEC - 2.5MG/TAB | enalapril maleate | C09AA | tablet | |
| 00708879 | VASOTEC - 5MG/TAB | enalapril maleate | CO9AA | tablet | |
| 00670901 00670928 | VASOTEC - 10MG/TAB VASOTEC - 20MG/TAB | enalapril maleate enalapril maleate | C09AA C09AA | tablet tablet | |
| 00708887 | VASOTEC - 40MG/TAB | enalapril maleate | C09AA | tablet | not sold |
| 01923846 | VASOTEC I.V 1.25MG/ML | enalaprilat | C09AA | injectable solution | |
| 00884324 00884332 | ZOCOR - 5MG/TAB ZOCOR - 10MG/TAB | simvastatin simvastatin | C10AA C10AA | tablet tablet | |
| 00884340 | ZOCOR - 20MG/TAB | simvastatin | C10AA C10AA | tablet | |
| 00884359 | ZOCOR - 40MG/TAB | simvastatin | C10AA | tablet | |
| 02186810 | ZYMAXIN - 200MG/TAB | ibuprofen lysine | M01AE | tablet | expired |
| | ROSST-MSD AGVET | | | | |
| Veterinary: 02060531 | ENACARD - 1MG/TAB | enalapril maleate | | tablet | |
| 02060558 | ENACARD - 1MG/1AB ENACARD - 2.5MG/TAB | enalapril maleate | | tablet | |
| 02060566 | ENACARD - 5MG/TAB | enalapril maleate | | tablet | |
| 02060574 | ENACARD - 10MG/TAB | enalapril maleate | | tablet | |
| 02060582 00717150 | ENACARD - 20MG/TAB EQVALAN LIQUID FOR HORSES - | enalapril maleate ivermectin | | tablet oral liquid | |
| 00717130 | 10MG/ML | ויטוווסטווו | | orar liquiu | |
| 00594431 | EQVALAN PASTE FOR HORSES - 18.6916MG/G | ivermectin | | oral paste | |
| 00651966 | HEARTGARD 30 - 0.068MG/TAB | ivermectin | | tablet | |
| 00651982 | HEARTGARD 30 - 0.136MG/TAB | ivermectin | | tablet | |
| | | | | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|--|--|--|--|
| 00651990 00812315 | HEARTGARD 30 - 0.272MG/TAB HEARTGARD 30 CHEW - 0.068MG/TAB | ivermectin ivermectin | | tablet tablet | |
| 00812323 | HEARTGARD 30 CHEW - | ivermectin | | tablet | |
| 00870064 | 0.136MG/TAB HEARTGARD 30 CHEW - 0.272MG/TAB | ivermectin | | tablet | |
| 02216604 02216612 01909975 01909983 01909991 00622125 | HEARTGARD 30 FX - 0.055MG/TAB HEARTGARD 30 FX - 0.165MG/TAB HEARTGARD 30 PLUS 0.068/57 HEARTGARD 30 PLUS 0.136/114 HEARTGARD 30 PLUS 0.272/227 IVOMEC DRENCH FOR SHEEP - 0.83MG/ML | ivermectin ivermectin/pyrantel pamoate ivermectin/pyrantel pamoate ivermectin/pyrantel pamoate ivermectin | ` | tablet tablet tablet tablet tablet injectable suspension | not sold not sold |
| 00630470 | IVOMEC INJ. FOR CATTLE - 10MG/ML | ivermectin | | injectable suspension | |
| 00583340 | IVOMEC INJ. FOR SWINE - 10MG/ML | ivermectin | | injectable suspension | not sold |
| 00761842 01926950 | IVOMEC POUR-ON - 5MG/ML IVOMEC SR BOLUS FOR CATTLE - 1720MG/BOLUS | ivermectin ivermectin | | topical solution sustained-release bolus | |
| 01913085 | IVOMEC SWINE PREMIX - 6000MG/KG | ivermectin | | oral powder | |
| McNEIL C | ONSUMER PRODUCTS COMPA | ANY | | | |
| Human: 02236894 02184656 02230542 02132575 02182920 | CHILDREN'S MOTRIN - 20MG/ML HISMANAL - 1MG/ML HISMANAL - 2MG/ML IMODIUM LINGUAL - 2MG/TAB MEDIPREN - 20MG/ML NIZORAL - 20MG/ML | ibuprofen astemizole astemizole loperamide hydrochloride ibuprofen ketoconazole | M01AE R06AX R06AX A07DA M01AE D11AC | oral suspension oral suspension oral suspension tablet oral suspension shampoo | not sold not sold introduced not sold |
| NEXSTAR | PHARMACEUTICALS, INC. | | | | |
| Human: 02218046 | DAUNOXOME - 50MG/VIAL | daunorubicin, liposomal | L01DB | powder for injectable solution | introduced |
| NOVARTIS | S ANIMAL HEALTH CANADA IN | C. | | | |
| Veterinary: | | | | | |
| 02220563 02220555 00846422 00846430 00846449 00846457 02016710 02016699 02016702 02016729 02016729 02141434 | INTERCEPTOR CHEW - 2.3MG/TAB INTERCEPTOR CHEW - 5.75MG/TAB INTERCEPTOR CHEW - 11.5MG/TAB | milbemycin oxime | | tablet tablet tablet tablet tablet tablet tablet tablet tablet tablet tablet | not sold not sold not sold not sold not sold not sold not sold |
| 02141426 | INTERCEPTOR FLAVOUR - | milbemycin oxime | | tablet | |
| 02141418 | 5.75MG/TAB INTERCEPTOR FLAVOUR - | milbemycin oxime | | tablet | |
| 02141396 | 11.5MG/TAB INTERCEPTOR FLAVOUR - | milbemycin oxime | | tablet | |
| 02087332 02087340 02087359 02087367 02057832 02192632 0229547 02229548 02229551 02229549 | 23MG/TAB PROGRAM - 45MG/TAB PROGRAM - 90MG/TAB PROGRAM - 204.9MG/TAB PROGRAM - 409.8MG/TAB PROGRAM - 133MG/UNIT PROGRAM - 266MG/UNIT SENTINEL 115/5.75 SENTINEL 115/5.75 SENTINEL 230/11.5 SENTINEL 46/2.3 SENTINEL 460/23 | lufenuron lufenuron lufenuron lufenuron lufenuron/milbemycin oxime lufenuron/milbemycin oxime lufenuron/milbemycin oxime lufenuron/milbemycin oxime | | tablet tablet tablet oral suspension oral suspension tablet tablet tablet tablet | introduced introduced introduced introduced |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|--|--|----------------------------------|---|-------------|
| | S CONSUMER HEALTH | | | | |
| Human: 02018411 | CORSYM 7.5/0.8 | phenylpropanolamine/ chlorpheniramine polistirex | R01BA | sustained-release oral suspension | |
| 02018403 02018381 00667285 | DELSYM - 6MG/ML DURAFEDRIN - 12MG/ML LIPACTIN - 5MG/G | dextromethorphan polistirex pseudoephedrine polistirex heparin sodium/zinc sulfate | R05DA R01BA D11AX | sustained-release oral suspension sustained-release oral suspension gel | |
| 02162318 02162326 00651184 02162245 | MAALOX HRF MAALOX HRF PROBAX - 20MG/G PRODIEM PLAIN - 672MG/G | antacid-alginate compound antacid-alginate compound propolis psyllium | A02EA A02EA D02AX A06AC | oral suspension tablet ointment oral granules | not sold |
| 02162253 | PRODIEM PLUS 542/124 | psyllium/senna | A06AC | oral granules | |
| NOVARTIS | S PHARMA CANADA INC. | | | | |
| Human: 02059762 | AREDIA - 30MG/VIAL | pamidronate disodium | M05BA | powder for injectable solution | |
| 02059770 | AREDIA - 60MG/VIAL | pamidronate disodium | M05BA | powder for injectable solution | |
| 02059789 01968130 | AREDIA - 90MG/VIAL DYNACIRC - 1.25MG/CAP | pamidronate disodium isradipine | M05BA C08CA | powder for injectable solution capsule | not sold |
| 00872385 | DYNACIRC - 2.5MG/CAP | isradipine | C08CA | capsule | not sold |
| 00872393 | DYNACIRC - 5MG/CAP | isradipine | CO8CA | capsule | not sold |
| 02108186 | ESTRACOMB .05/.0525 | estradiol 17b/estradiol 17b & norethindrone acetate | G03FA | transdermal patch | |
| 00756792 | ESTRADERM 100 - 8MG/PATCH | estradiol 17b | G03CA | transdermal patch | |
| 00756849 | ESTRADERM 25 - 2MG/PATCH | estradiol 17b | G03CA | transdermal patch | |
| 00756857 02231384 | ESTRADERM 50 - 4MG/PATCH FEMARA - 2.5MG/TAB | estradiol 17b letrozole | G03CA L02BG | transdermal patch tablet intro | duced (nas) |
| 02230898 | FORADIL - 0.012MG/DOSE | formoterol fumarate | R03AC | | duced (nas) |
| 01943065 | HABITROL 14 - 35MG/PATCH | nicotine | N07BA | transdermal patch | |
| 01943073 | HABITROL 21 - 52.5MG/PATCH | nicotine | N07BA | transdermal patch | |
| 01943057 | HABITROL 7 - 17.5MG/PATCH | nicotine | N07BA | transdermal patch | |
| 02031094 | LAMISIL - 10MG/G | terbinafine hydrochloride | D01AE D01BA | cream | not cold |
| 02031108 02031116 | LAMISIL - 125MG/TAB LAMISIL - 250MG/TAB | terbinafine hydrochloride terbinafine hydrochloride | D01BA | tablet tablet | not sold |
| 02128209 | LENTARON - 250MG/VIAL | formestane | L02BG | powder for injectable solution | |
| 02061562 | LESCOL - 20MG/CAP | fluvastatin sodium | C10AA | capsule | |
| 02061570 | LESCOL - 40MG/CAP | fluvastatin sodium | C10AA | capsule | |
| 00885835 00885843 | LOTENSIN - 5MG/TAB LOTENSIN - 10MG/TAB | benazepril hydrochloride | CO9AA | tablet | |
| 00885851 | LOTENSIN - 20MG/TAB | benazepril hydrochloride benazepril hydrochloride | C09AA C09AA | tablet tablet | |
| 02162873 | LOTENSIN-HCT 10/12.5 | benazepril hydrochloride/ | C09BA | tablet | not sold |
| 02162881 | LOTENSIN-HCT 20/25 | hydrochlorothiazide benazepril hydrochloride/ | C09BA | tablet | not sold |
| | | hydrochlorothiazide | | | |
| 02162865 | LOTENSIN-HCT 5/6.25 | benazepril hydrochloride/ hydrochlorothiazide | C09BA | tablet | not sold |
| 01990926 | MIACALCIN - 50UNIT/ML | calcitonin salmon | V03AG | injectable solution | not sold |
| 01992880 | MIACALCIN - 100UNIT/ML | calcitonin salmon | V03AG | injectable solution | not sold |
| 02228947 02223740 | MIGRANAL - 4MG/ML NORPROLAC - 0.025MG/TAB | dihydroergotamine mesylate quinagolide hydrochloride | N02CA G02CB | nasal aerosol tablet | not sold |
| 02223759 | NORPROLAC - 0.05MG/TAB | quinagolide hydrochloride | G02CB | tablet | not sold |
| 02223767 | NORPROLAC - 0.075MG/TAB | quinagolide hydrochloride | G02CB | tablet | not sold |
| 02223775 | NORPROLAC - 0.15MG/TAB | quinagolide hydrochloride | G02CB | tablet | not sold |
| 02229854 | RESTORIL - 7.5MG/CAP | temazepam | N05CD | capsule | not sold |
| 00755591 01907182 | SANDIMMUNE - 25MG/CAP SANDIMMUNE - 50MG/CAP | cyclosporine | L04AA L04AA | capsule | |
| 00755605 | SANDIMMONE - JOUNG/CAP | cyclosporine cyclosporine | L04AA | capsule capsule | |
| 00593257 | SANDIMMUNE - 50MG/ML | cyclosporine | L04AA | injectable solution | |
| 00593249 | SANDIMMUNE - 100MG/ML | cyclosporine | L04AA | oral solution | |
| 02150689 | NEORAL - 25MG/CAP | cyclosporine | L04AA | capsule | |
| 02150662 | NEORAL - 50MG/CAP | cyclosporine | L04AA | capsule | |
| 02150670 02150697 | NEORAL - 100MG/CAP NEORAL - 100MG/ML | cyclosporine cyclosporine | L04AA L04AA | capsule oral solution | |
| 00839191 | SANDOSTATIN - 0.05MG/ML | octreotide | H01CB | injectable solution | |
| 00839205 | SANDOSTATIN - 0.1MG/ML | octreotide | H01CB | injectable solution | |
| 02049392 | SANDOSTATIN - 0.2MG/ML | octreotide | H01CB | injectable solution | |
| 00839213 | SANDOSTATIN - 0.5MG/ML | octreotide | H01CB | injectable solution | |
| 00584223 | TRANSDERM-NITRO 0.2 - 25MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| | | | | | |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|--|--|--|----------------------------------|
| 00852384 | TRANSDERM-NITRO 0.4 - | nitroglycerin | C01DA | transdermal patch | |
| 02046156 | 50MG/PATCH TRANSDERM-NITRO 0.6 - | nitroglycerin | C01DA | transdermal patch | |
| 02046164 | 75MG/PATCH TRANSDERM-NITRO 0.8 - 100MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 00568627 00568635 02204444 02204401 02204428 02204436 | VISKAZIDE 10/25 VISKAZIDE 10/50 VIVELLE 100 - 8.66MG/PATCH VIVELLE 37.5 - 3.28MG/PATCH VIVELLE 50 - 4.33MG/PATCH VIVELLE 50 - 6.56MG/PATCH | pindolol/hydrochlorothiazide pindolol/hydrochlorothiazide estradiol 17b estradiol 17b estradiol 17b estradiol 17b | C07BA C07BA G03CA G03CA G03CA G03CA | tablet tablet transdermal patch transdermal patch transdermal patch transdermal patch | |
| | RDISK CANADA INC. | | | | |
| Human: 01986813 | INSULATARD NPH - 100UNIT/ML | insulin (isophane) human semi-synthetic | A10AC | injectable suspension | not sold |
| 02056682 02056690 01985973 | LOGIPARIN - 10000UNIT/ML LOGIPARIN - 11700UNIT/ML MIXTARD 15/85 - 100UNIT/ML | tinzaparin sodium tinzaparin sodium insulin (regular/isophane) | B01AB B01AB A10AE | injectable suspension injectable suspension injectable suspension | not sold not sold not sold |
| 01986821 | MIXTARD 30/70 - 100UNIT/ML | human semi-synthetic insulin (regular/isophane) | A10AE | injectable suspension | not sold |
| 01985965 | MIXTARD 50/50 - 100UNIT/ML | human semi-synthetic insulin (regular/isophane) | A10AE | injectable suspension | not sold |
| 02024292 | NOVOLIN GE 10/90 - 100UNIT/ML | human semi-synthetic insulin (regular/isophane) | A10AE | injectable suspension | |
| 02024306 | NOVOLIN GE 20/80 - 100UNIT/ML | human biosynthetic insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 02024217 | NOVOLIN GE 30/70 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 02025248 | NOVOLIN GE 30/70 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 02024314 | NOVOLIN GE 40/60 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 02024322 | NOVOLIN GE 50/50 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | |
| 02024241 | NOVOLIN GE LENTE - 100UNIT/ML | | A10AC | injectable suspension | |
| 02024225 | NOVOLIN GE NPH - 100UNIT/ML | insulin (isophane) human biosynthetic | A10AC | injectable suspension | |
| 02024268 | NOVOLIN GE NPH - 100UNIT/ML | insulin (isophane) human biosynthetic | A10AC | injectable suspension | |
| 02024233 | NOVOLIN GE TORONTO - 100UNIT/ML | insulin (regular) human biosynthetic | A10AB | injectable suspension | |
| 02024284 | NOVOLIN GE TORONTO - 100UNIT/ML | insulin (regular) human biosynthetic | A10AB | injectable suspension | |
| 02024276 | NOVOLIN GE ULTRALENTE - 100UNIT/ML | insulin (ultralente) human biosynthetic | A10AD | injectable suspension | |
| 01988700 | NOVOLINSET 30/70 - 100UNIT/ML | insulin (regular/isophane) human semi-synthetic | A10AE | injectable suspension | not sold |
| 02024411 | NOVOLINSET GE 10/90 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | not sold |
| 02024438 | NOVOLINSET GE 20/80 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | not sold |
| 02024446 | NOVOLINSET GE 30/70 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | not sold |
| 02024454 | NOVOLINSET GE 40/60 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | not sold |
| 02025264 | NOVOLINSET GE 50/50 - 100UNIT/ML | insulin (regular/isophane) human biosynthetic | A10AE | injectable suspension | not sold |
| 02024403 | NOVOLINSET GE NPH - 100UNIT/ML | insulin (isophane) human biosynthetic | A10AC | injectable suspension | not sold |
| 02025256 | NOVOLINSET GE TORONTO - 100UNIT/ML | insulin (regular) human biosynthetic | A10AB | injectable suspension | not sold |
| 01986090 | NOVOLINSET NPH - 100UNIT/ML | insulin (isophane) human semi-synthetic | A10AC | injectable suspension | not sold |
| 01927477 | NOVOLINSET TORONTO - 100UNIT/ML | insulin (regular) human semi-synthetic | A10AB | injectable suspension | not sold |
| 01986805 | VELOSULIN HUMAN - 100UNIT/ML | insulin (regular) human semi-synthetic | A10AB | injectable suspension | not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|---|--|--|---|
| NYCOMED |) CANADA INC. | | | | |
| Human: 01962973 02172720 02172739 02172747 02172755 02172771 02145766 02145774 | OMNIPAQUE 140 - 302MG/ML OMNIPAQUE 180 - 389MG/ML OMNIPAQUE 240 - 518MG/ML OMNIPAQUE 300 - 647MG/ML OMNIPAQUE 350 - 755MG/ML OMNISCAN - 287MG/ML VISIPAQUE 270 - 550MG/ML VISIPAQUE 320 - 652MG/ML | iohexol iohexol iohexol iohexol gadodiamide iodixanol iodixanol | V08AB V08AB V08AB V08AB V08AB V08CA V08AB V08AB | injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution injectable solution | not sold |
| OHMEDA, | DIVISION OF CANADIAN OX | YGEN LIMITED | | | |
| Human: 00418994 | STADOL - 2MG/ML | butorphanol tartrate | N02AF | injectable solution | not sold |
| ORGANON | I CANADA LTD. (AKZO) | | | | |
| Human: 02042487 02042479 00687405 02129043 02108208 | MARVELON 0.15/0.03 MARVELON 0.15/0.03 NORCURON - 10MG/VIAL ORGARAN - 1250UNIT/ML ZEMURON - 10MG/ML | desogestrel/ethinyl estradiol desogestrel/ethinyl estradiol vecuronium bromide danaparoid sodium rocuronium bromide | G03AA G03AA M03AC B01AB M03AC | tablet tablet powder for injectable solutior injectable solution injectable solution | expired expired |
| PFIZER C | ANADA INC. | | | | |
| Human: 02232043 02232044 00871052 00871060 00871079 02141442 00891835 02024152 02024152 02024160 00891830 00891819 00891827 02230443 02230443 02230444 02093170 02093189 00878901 00878928 00878936 01942506 | ARICEPT - 5MG/TAB ARICEPT - 10MG/TAB CARDURA - 1MG/TAB CARDURA - 2MG/TAB CARDURA - 4MG/TAB CARDURA - 8MG/TAB DIFLUCAN - 150MG/CAP DIFLUCAN - 10MG/ML DIFLUCAN - 10MG/ML DIFLUCAN - 00MG/TAB DIFLUCAN - 00MG/TAB DIFLUCAN - 200MG/TAB GLUCOTROL XL - 5MG/TAB GLUCOTROL XL - 10MG/TAB MINIPRESS XL - 2.5MG/TAB MINIPRESS XL - 2.5MG/TAB NORVASC - 2.5MG/TAB NORVASC - 10MG/TAB PLAX PEPPERMINT 20/2/2.5 | donepezil hydrochloride donepezil hydrochloride doxazosin mesylate doxazosin mesylate doxazosin mesylate doxazosin mesylate fluconazole fluconazole fluconazole fluconazole fluconazole fluconazole fluconazole glipizide glipizide glipizide prazosin hydrochloride amlodipine besylate amlodipine besylate amlodipine besylate amlodipine besylate sodium benzoate/sodium salicylate/sodium lauryl sulfate | N07AA N07AA C02CA C02CA J02AC J02AC J02AC J02AC J02AC J02AC J02AC J02AC J02AC J02AC A10BB A10BB C02CA C08CA C08CA C08CA C08CA A01AD | tablet tablet tablet tablet tablet tablet tablet capsule injectable solution oral suspension oral suspension tablet tablet tablet sustained-release tablet sustained-release tablet sustained-release tablet sustained-release tablet tablet tablet tablet tablet tablet oral rinse | introduced (nas) introduced (nas) expired expired expired not sold not sold not sold not sold not sold not sold not sold |
| 01942409 | PLAX SOFT MINT 20/2/2.5 | sodium benzoate/sodium salicylate/sodium lauryl sulfate | A01AD | oral rinse | |
| 02223546 02223554 01900978 00874116 | REACTINE - 5MG/TAB REACTINE - 10MG/TAB REACTINE - 20MG/TAB UNASYN 1000/2000 | cetirizine hydrochloride cetirizine hydrochloride cetirizine hydrochloride sulbactam sodium/ampicillin sodium | R06AE R06AE R06AE J01CR | tablet tablet tablet powder for injectable solutior | introduced not sold not sold |
| 01955527 | UNASYN 1000/2000 | sulbactam sodium/ampicillin | J01CR | powder for injectable solution | not sold |
| 00874108 | UNASYN 500/1000 | sodium sulbactam sodium/ampicillin | J01CR | powder for injectable solution | not sold |
| 01955519 | UNASYN 500/1000 | sodium sulbactam sodium/ampicillin | J01CR | powder for injectable solution | not sold |
| 02091291 02223716 02223724 02229971 | ZITHROMAX - 250MG/CAP ZITHROMAX - 20MG/ML ZITHROMAX - 40MG/ML ZITHROMAX - 1000MG/POUCH | sodium azithromycin azithromycin azithromycin azithromycin | J01FA J01FA J01FA J01FA | capsule powder for oral suspension powder for oral suspension powder for oral solution | not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|--|--|----------------|--|------------------|
| 02212021 | ZITHROMAX - 250MG/TAB | azithromycin | J01FA | tablet | not sold |
| 02231143 | ZITHROMAX - 600MG/TAB | azithromycin | J01FA | tablet | introduced |
| 02132702 01962817 | ZOLOFT - 25MG/CAP ZOLOFT - 50MG/CAP | sertraline hydrochloride sertraline hydrochloride | N06AB N06AB | capsule capsule | |
| 01962779 | ZOLOFT - 100MG/CAP | sertraline hydrochloride | N06AB | capsule | |
| 01962787 | ZOLOFT - 150MG/CAP | sertraline hydrochloride | N06AB | capsule | not sold |
| 01962795 | ZOLOFT - 200MG/CAP | sertraline hydrochloride | N06AB | capsule | not sold |
| | ANADA INC.,ANIMAL HEALTH | GRUUP | | | |
| Veterinary: 02229543 | AVIAX - 50000MG/KG | semduramicin sodium | | oral powder | introduced (nas) |
| 01913034 | CESTEX - 12.5MG/TAB | epsiprantel | | tablet | |
| 01913026 | CESTEX - 25MG/TAB | epsiprantel | | tablet | |
| 01913018 01912992 | CESTEX - 50MG/TAB CESTEX - 100MG/TAB | epsiprantel epsiprantel | | tablet tablet | |
| 02038331 | CLAVAMOX 100/25 | amoxicillin trihydrate/ | | tablet | |
| 02000001 | 02/07/00/20 | clavulanate potassium | | labiot | |
| 02038315 | CLAVAMOX 200/50 | amoxicillin trihydrate/ | | tablet | |
| 02038358 | CLAVAMOX 300/75 | clavulanate potassium amoxicillin trihydrate/ | | tablet | |
| 02030330 | GLAVAIVION 300/75 | clavulanate potassium | | lablet | |
| 02027879 | CLAVAMOX 50/12.5 | amoxicillin trihydrate/ | | oral suspension | |
| 00000000 | | clavulanate potassium | | - | |
| 02038323 | CLAVAMOX 50/12.5 | amoxicillin trihydrate/ clavulanate potassium | | tablet | |
| 01950673 | DOMITOR - 1MG/ML | medetomidine hydrochloride | | injectable solution | not sold |
| 00755559 | DORMOSEDAN - 10MG/ML | detomidine hydrochloride | | sólution | |
| | ENDURALL - K | inactivated rabies vaccine, | | injectable solution | not sold |
| | FELOCELL CVR | porcine cell line origin modified live feline | | injectable solution | expired |
| | | rhinotracheitis- | | injectable solution | expireu |
| | | panleukopenia virus | | | |
| | FELOCELL CVR-C | modified live feline | | injectable solution | expired |
| | FELOMUNE CVR | rhinotracheitis modified live feline | | | |
| | | rhinotracheitis-calicivirus | | injectable solution | expired |
| | | vaccine | | inicatable colution | |
| | FIRSTDOSE CPV | modified live canine parvovirus vaccine | | injectable solution | |
| | LEUKOCELL | inactivated feline leukaemia | | injectable solution | not sold |
| | | vaccine | | | |
| 00725064 | LEUKOCELL 2 LIFE-GUARD-H.E. | feline leukemia vaccine nutrients/electrolytes | | injectable solution oral powder | |
| 00606081 | PARATECT BOLUS - 22700MG/TAB | morantel tartrate | | sustained-release oral device | not sold |
| 00778923 | PARATECT FLEX - 19800MG/TAB | morantel tartrate | | sustained-release oral device | not sold |
| 00606103 | POSISTAC - 12500MG/KG | salinomycin | | oral powder | not sold |
| 00633828 00603724 | POSISTAC - 50000MG/KG POSISTAC - 60000MG/KG | salinomycin salinomycin | | oral powder oral powder | not sold |
| 00003724 | RABGUARD-TC | inactivated rabies vaccine | | injectable solution | not sold |
| 02006936 | RESORB | electrolyte product | | pówder for oral suspension | expired |
| 00698083 | SYNERGISTIN 60/120 | sulbactam/ampicillin | | injectable suspension | |
| | VANGUARD 5 | modified live canine distemper-adenovirus type | 2 | injectable solution | |
| | VANGUARD 5 CV | canine combination vaccine | 2 | injectable solution | |
| | VANGUARD 5B | modified live canine | _ | injectable solution | |
| | VANGUARD 5L | distemper-adenovirus type modified live canine | 2 | injectable solution | |
| | VANGUARD SL | distemper-adenovirus type | 2 | injectable solution | |
| | VANGUARD CPV (KILLED) | inactivated cannine parvovirus vaccine | _ | injectable solution | |
| | VANGUARD CPV (MLV) | modified live canine parvovirus vaccine | | injectable solution | not sold |
| PHARMA | CIA & UPJOHN INC. | | | | |
| Human: | | | | | |
| 02071002 | ADRIAMYCIN PFS - 2MG/ML | doxorubicin hydrochloride | L01DB | injectable solution | |
| 02065657 | ADRIAMYCIN RDF - 10MG/VIAL | doxorubicin hydrochloride | L01DB | powder for injectable solution | |
| 00698040 02069555 | ADRIAMYCIN RDF - 20MG/VIAL ADRIAMYCIN RDF - 50MG/VIAL | doxorubicin hydrochloride doxorubicin hydrochloride | L01DB L01DB | powder for injectable solution powder for injectable solution | |
| 02009000 | | | LUIDD | powder for injectable solution | 1 |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|---|-------|--|---------------------------|
| 02069563 | ADRIAMYCIN RDF - 150MG/VIAL AMPEROZIDE - 0.5MG/TAB AMPEROZIDE - 2.5MG/TAB | doxorubicin hydrochloride amperozide amperozide | L01DB | powder for injectable solution tablet tablet | n not sold not sold |
| | AMPEROZIDE - 5MG/TAB | amperozide | | tablet | not sold |
| 02231622 | CAMPTOSAR - 20MG/ML | irinotecan hydrochloride | L01XX | injectable solution | introduced (nas) |
| 02132680 | COLESTID - 1000MG/TAB | colestipol hydrochloride | C10AC | tablet | () |
| 02132699 | COLESTID ORANGE - | colestipol hydrochloride | C10AC | oral granules | |
| 00030910 | 5000MG/DOSE CORTEF - 10MG/TAB | hydrocortisone | H02AB | tablet | |
| 00030929 | CORTEF - 20MG/TAB | hydrocortisone | H02AB | tablet | |
| 02063808 | DIPENTUM - 250MG/CAP | olsalazine sodium | A07EC | capsule | |
| 00875856 | DIPENTUM - 500MG/TAB | olsalazine sodium | A07EC | tablet | not sold |
| 02063794 | EMCYT - 140MG/CAP | estramustine phosphate disodium | L01XX | capsule | expired |
| 02168898 | ESTRING - 2MG/RING | estradiol | G03CA | vaginal ring | |
| 02132656 | FRAGMIN - 2500UNIT/ML | dalteparin sodium | B01AB | injectable solution | |
| 02132664 | FRAGMIN - 10000UNIT/ML | dalteparin sodium | B01AB | injectable solution | |
| 02132621 | FRAGMIN - 12500UNIT/ML | dalteparin sodium | B01AB | injectable solution | |
| 02132648 | FRAGMIN - 25000UNIT/ML | dalteparin sodium | B01AB | injectable solution | |
| 02231171 | FRAGMIN - 25000UNIT/ML | dalteparin sodium | B01AB | injectable solution | introduced |
| 02166100 | IDAMYCIN - 5MG/CAP | idarubicin hydrochloride | L01DB | capsule | |
| 02166119 | IDAMYCIN - 10MG/CAP | idarubicin hydrochloride | L01DB | capsule | not sold |
| 02166127 | IDAMYCIN - 25MG/CAP | idarubicin hydrochloride | L01DB | capsule | not sold |
| 02065630 | IDAMYCIN - 5MG/VIAL | idarubicin hydrochloride | L01DB | powder for injectable solution | 1 |
| 02068095 | IDAMYCIN - 10MG/VIAL | idarubicin hydrochloride | L01DB | powder for injectable solution | |
| | LINOMIDE - 2.5MG/TAB | roquinimex | L03AX | tablet | not sold |
| | LINOMIDE - 5MG/TAB | roquinimex | L03AX | tablet | not sold |
| 00000700 | LINOMIDE - 10MG/TAB | roquinimex | L03AX | tablet | not sold |
| 02063786 | MYCOBUTIN - 150MG/CAP | rifabutin | J04AB | capsule | expired |
| 00194913 | NEO-CORTEF 10/5 | hydrocortisone acetate/ neomycin sulfate | D07CA | ointment | |
| 00194921 | NEO-CORTEF 15/5 | hydrocortisone acetate/ | S01CA | ophthalmic ointment | not sold |
| 00194948 | NEO-CORTEF 15/5 | neomycin sulfate hydrocortisone acetate/ neomycin sulfate | S01CA | ophthalmic suspension | |
| 00194883 | NEO-CORTEF 5/5 | hydrocortisone acetate/ neomycin sulfate | S01CA | ophthalmic ointment | not sold |
| 00194891 | NEO-CORTEF 5/5 | hydrocortisone acetate/ neomycin sulfate | D07CA | ointment | |
| 02065703 | PHARMORUBICIN PFS - 2MG/ML | epirubicin hydrochloride | L01DB | injectable solution | |
| 02065746 | PHARMORUBICIN RDF - 10MG/VIAL | epirubicin hydrochloride | L01DB | powder for injectable solution | า |
| 00698202 | PHARMORUBICIN RDF - 20MG/VIAL | epirubicin hydrochloride | L01DB | powder for injectable solution | not sold |
| 02069512 | PHARMORUBICIN RDF - 50MG/VIAL | epirubicin hydrochloride | LOIDB | powder for injectable solution | 1 |
| 01951882 | PHARMORUBICIN RDF - | epirubicin hydrochloride | L01DB | powder for injectable solution | n not sold |
| 00860786 | 150MG/VIAL PREPIDIL - 0.5MG/SYRINGE | dinoprostone | G02AD | intra-uterine gel | |
| 02028689 | PROMIT - 150MG/ML | dextran 1 | B05AA | injectable solution | not sold |
| 01919679 | PROSTIN E2 - 1MG/SYRINGE | dinoprostone | G02AD | intra-uterine gel | 1101 3010 |
| 01919687 | PROSTIN E2 - 2MG/SYRINGE | dinoprostone | G02AD | intra-uterine gel | |
| 00030600 | SOLU-CORTEF - 100MG/VIAL | hydrocortisone sodium | HOZAB | powder for injectable solution | ı |
| 00030619 | SOLU-CORTEF - 250MG/VIAL | succinate hydrocortisone sodium | H02AB | powder for injectable solution | ı |
| 00030627 | SOLU-CORTEF - 500MG/VIAL | succinate hydrocortisone sodium | H02AB | powder for injectable solution | ı |
| 00030635 | SOLU-CORTEF - 1000MG/VIAL | succinate hydrocortisone sodium | H02AB | powder for injectable solution | ı |
| | STERECYT - 20MG/TAB | succinate prednimustine | L01AA | tablet | not sold |
| | STERECYT - 100MG/TAB | prednimustine | L01AA | tablet | not sold |
| 02231493 | XALATAN - 0.05MG/ML | latanoprost | S01EX | ophthalmic solution | introduced (nas) |
| 02153432 | ZINECARD - 250MG/VIAL | dexrazoxane | V03AF | powder for injectable solution | |
| 02153440 | ZINECARD - 500MG/VIAL | dexrazoxane | V03AF | powder for injectable solution | 1 |
| Veterinary: 00813567 02207753 | EXCENEL - 50MG/ML PIRSUE AQUEOUS GEL - 50MG/SYRINGE | ceftiofur sodium pirlimycin hydrochloride | | powder for injectable solution intramammary gel | 1 |

| PROCTER & GAMBLE PHARMACEUTICALS CANADA INC. Human (2019) CACDL - 400WG/TAB (20176017) Saminoss flucture athermoss flucture athermoss flucture eliciture athonate (active athonate) A07EC total totalpaste (able) tablet totalpaste (able) not sold 0205862 MACROBID OF ELI LILLY CANADA INC. | DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|--|---|---|----------------------------------|--|--------------|
| 01997580 ASACDL - 400MG/TAB 5-aminosalicylic acid A07EC tablet tablet tablet 221062 CREST GUM CARE - 454MG/G etidronate disodium and M05BB tablet tablet 22058662 MACROBID - 100MG/CAP antitrofurnation G04AC capsule activity activity of tablet table | PROCTER | & GAMBLE PHARMACEUTIC | ALS CANADA INC. | | | |
| 20203682 MACROBID - 100MG/CAP nitrofurantoin G04AC capsule PROVEL, DIVISION OF ELI LILLY CANADA INC. . | 01997580 02231062 | CREST GUM CARE - 4.54MG/G | stannous fluoride etidronate disodium and | A01AA | toothpaste | not sold |
| Veteringer 005/16333 Competitions Competitions Real Brank - 250000MG/KG estration Inicitable injectable solution not sold 01902867 MAXIBAN PREMIX narasin/nicarbazin abamectin injectable solution injectable solution 01902867 MAXIBAN PREMIX narasin feed premix regetable injectable solution injectable solution 00057604 TVLAN 50 SULFA G tylineitable timeitable istamethazine feed premix not sold 20180756 GASTROBID - 20MG/TAB metoclopramide A03FA sustained-release tablet not sold 02125331 HYDROMORPH CONTIN - hydrochioride N02AA sustained-release capsule not sold 02125334 HYDROMORPH CONTIN - hydrochioride N02AA sustained-release capsule not sold 02125354 HYDROMORPH CONTIN - hydrochioride N02AA sustained-release capsule not sold 02125354 HYDROMORPH CONTIN - hydrochioride N02AA sustained-release capsule not sold 02125354 HYDROMORPH CONTIN - hydrochioride N02AA su | 02063662 | MACROBID - 100MG/CAP | | G04AC | capsule | |
| 00616333 CARBIGRAN - 250000MC/KG nicrbazin feed premix not sold 00590932 COMPUDDSE - 24MG/MP abarrectin injectable solution injectable solution 002163355 ENDECTO - 10MG/ML abarrectin injectable solution injectable solution 00216355 ENDECTO - 10MG/ML atarsin/fucathazin feed premix injectable solution 00057648 MCOTL - 300MG/ML tilmicosin sulfate injectable solution injectable solution 00057648 TVLAN 50 SULFA G tylosin phosphate/ feed premix rot sold 00057538 MONTEBAN 70 PREMIX - narasin feed premix not sold 02125356 GASTROBID - 20MG/TAB metoclopramide A03FA sustained-release tablet not sold 02125331 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride <td>PROVEL,</td> <td>DIVISION OF ELI LILLY CANA</td> <td>DA INC.</td> <td></td> <td></td> <td></td> | PROVEL, | DIVISION OF ELI LILLY CANA | DA INC. | | | |
| Sulfamethazine PURDUE FREDERICK Human: 02163756 GASTROBID - 20MG/TAB metoclopramide hydrochloride A03FA sustained-release tablet not sold 0212532 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212533 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212537 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212539 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 0212539 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release tablet oxycodone hydrochloride N02AA sustained-release tablet oxycodone hydrochloride N02AA sustained-release tablet oxycodone hydrochloride N02AA sustained-release tablet oxycodo | 00616338 00590932 02123355 01902687 00857602 | COMPUDOSE - 24MG/IMP ENDECTO - 10MG/ML MAXIBAN PREMIX MICOTIL - 300MG/ML MONTEBAN 70 PREMIX - | estradiol abamectin narasin/nicarbazin tilmicosin sulfate | | injectable implant injectable solution feed premix injectable solution | not sold |
| Human: 02163756 GASTROBID - 20MG/TAB hydrochloride hydrochlo | 00637645 | TYLAN 50 SULFA G | | | feed premix | |
| 02163756 GASTROBID - 20MG/TAB metoclopramide A03FA sustained-release tablet not sold 02125323 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule 0212533 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule 0212533 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212538 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release capsule not sold 0212539 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release tablet cold 0212530 HYDROMORPH CONTIN - hydrochloride N02AA sustained-release tablet cold 02202441 OXYCONTIN - 10MG/TAB oxycodone hydrochloride N02AA sustained-release tablet | PURDUE | FREDERICK | | | | |
| 02125323 HYDROMORPH CONTIN - Mydrochoride hydrochoride N02AA sustained-release capsule 02125331 HYDROMORPH CONTIN - hydrochoride hydrochoride N02AA sustained-release capsule 02125335 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 0212536 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 0212537 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 0212538 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 0212539 HYDROMORPH CONTIN - hydrochoride hydrochoride N02AA sustained-release capsule not sold 02020241 0XYCONTIN - 20MG/TAB oxycodone hydrochoride N02AA sustained-release tablet capsule 02202441 0XYCONTIN - 20MG/TAB oxycodone hydrochoride N02AA sustained-release tablet capsule 0220246 0XYCONTIN - 20MG/TAB oxycodone hydrochoride N02AA sustained-release tablet cap | | GASTROBID - 20MG/TAB | metoclopramide | A03FA | sustained-release tablet | not sold |
| 02125331 HYDROMORPH CONTIN - hydromorphone 10MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule 02125358 HYDROMORPH CONTIN - 10MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule not sold 02125354 HYDROMORPH CONTIN - 12MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule not sold 02125374 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 02125382 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 02125382 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 02125380 HYDROMORPH CONTIN - hydrochloride hydrochloride N02AA sustained-release capsule not sold 020202441 OXYCONTIN - 10MG/TAB oxycodone hydrochloride N02AA sustained-release tablet ozycodone 02202456 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet ozycodone 02202464 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet ozycodone 022024563 <t< td=""><td>02125323</td><td></td><td>hydromorphone</td><td>N02AA</td><td>sustained-release capsule</td><td></td></t<> | 02125323 | | hydromorphone | N02AA | sustained-release capsule | |
| 02125358 HYDROMORPH CONTIN - 10MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125366 HYDROMORPH CONTIN - 12MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125374 HYDROMORPH CONTIN - 12MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125382 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 02125390 HYDROMORPH CONTIN - hydromorphone N02AA sustained-release capsule not sold 02202464 OXYCONTIN - 10MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202464 OXYCONTIN - 20MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202465 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202464 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202465 INTAL INHALER - 1MG/DOSE sodium cromoglycate R03BC aerosol for inhalation expired 00555649 INTAL SYNCRONER - 1MG/DOSE sodium cromoglycate | 02125331 | HYDROMORPH CONTIN - | hydromorphone | N02AA | sustained-release capsule | |
| 02125366 HYDROMORPH CONTIN - 12MG/CAP hydromorphone hydrochloride hydrochloride N02AA sustained-release capsule 02125374 HYDROMORPH CONTIN - 20MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125382 HYDROMORPH CONTIN - 24MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule not sold 02125390 HYDROMORPH CONTIN - 4VOROMORPH CONTIN - 20MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule not sold 02202441 OXYCONTIN - 10MG/TAB oxycodone hydrochloride N0220445 N02AA sustained-release tablet expired 02202476 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet expired 02202476 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet expired 02449636 TRILISATE 293/362 choline salicylate/ magnesium salicylate N02BA tablet expired 00555649 INTAL INHALER - 1MG/DOSE NO555649 sodium cromoglycate R03BC aerosol for inhalation powder for inhalation expired 00538641 INTAL SPINCAPS - 20MG/ CARTRIDE sodium cromoglycate R03BA aerosol for inhalation R05BA </td <td>02125358</td> <td>HYDROMORPH CONTIN -</td> <td>hydromorphone</td> <td>N02AA</td> <td>sustained-release capsule</td> <td>not sold</td> | 02125358 | HYDROMORPH CONTIN - | hydromorphone | N02AA | sustained-release capsule | not sold |
| 02125374 HYDROMORPH CONTIN - 20MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125382 HYDROMORPH CONTIN - 24MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02125390 HYDROMORPH CONTIN - 24MG/CAP hydrochloride hydrochloride N02AA sustained-release capsule not sold 02202441 OXYCONTIN - 10MG/TAB oxycodone hydrochloride 0xycodone hydrochloride N02AA sustained-release tablet 02202446 OXYCONTIN - 40MG/TAB oxycodone hydrochloride 0xycodone hydrochloride N02AA sustained-release tablet 02202484 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet expired 00449636 TRILISATE 293/362 choline salicylate N02BA tablet expired RHÔNE-POULENC RORER CANADA INC. Human: NOZAS sodium cromoglycate R03BC aerosol for inhalation expired 00261238 INTAL SPINCAPS - 20MG/ CARTRIDGE sodium cromoglycate R03BC aerosol for inhalation expired 00888397 MYKROX - 0.5MG/TAB cefixime J01DA sustained-release oral suspens | 02125366 | HYDROMORPH CONTIN - | hydromorphone | N02AA | sustained-release capsule | |
| 02125382 HYDROMORPH CONTIN - 24MG/CAP hydrocnloride hydrochloride N02AA sustained-release capsule 02125390 HYDROMORPH CONTIN - 30MG/CAP hydrochloride N02AA sustained-release capsule not sold 02202441 OXYCONTIN - 10MG/TAB oxycodone hydrochloride N02AA sustained-release tablet not sold 02202446 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet not sold 02202446 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet not sold 02202445 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet not sold 02202446 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet expired 02449636 TRILISATE 293/362 choline salicylate/ N02BA tablet expired M0555649 INTAL INTAL SPINCAPS - 20MG/ sodium cromoglycate R03BC aerosol for inhalation expired 00638641 INTAL SPINCAPS - 20MG/ sodium cromoglycate R03BC aerosol for inhalation expired 00838657 | 02125374 | HYDROMORPH CONTIN - | hydromorphone | N02AA | sustained-release capsule | not sold |
| 02125390 HYDROMORPH CONTIN - 30MG/CAP hydromorphone hydrochloride N02AA sustained-release capsule not sold 02202441 0XYCONTIN - 10MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202468 0XYCONTIN - 20MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202468 0XYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202476 0XYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202441 0XYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202441 0XYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 0220441 0XYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 00449636 TRILISATE 293/362 choline salicylate N02BA tablet expired RMÔNE-POULENC RORER CANADA INC. Human: 00261238 INTAL INHALER - 1MG/DOSE sodium cromoglycate R03BC aerosol for inhalation expired 00638641 INTAL SPINCAPS - | 02125382 | HYDROMORPH CONTIN - | hydromorphone | N02AA | sustained-release capsule | |
| 02202468 OXYCONTIN - 20MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202476 OXYCONTIN - 40MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202486 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202486 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 02202486 OXYCONTIN - 80MG/TAB oxycodone hydrochloride N02AA sustained-release tablet 00449636 TRILISATE 293/362 choline salicylate/ N02BA tablet expired RHÔNE-POULENC RORER CANADA INC. | 02125390 | HYDROMORPH CONTIN - | hydromorphone hydrochloride | N02AA | sustained-release capsule | not sold |
| Human: 00555649 00261238INTAL INHALER - 1MG/DOSE INTAL SPINCAPS - 20MG/ CARTRIDGEsodium cromoglycate sodium cromoglycateR03BC R03BCaerosol for inhalation powder for inhalationexpired00638641 00888397INTAL SYNCRONER - 1MG/DOSE MYKROX - 0.5MG/TABsodium cromoglycate metolazone codeine/chlorpheniramine polistirexR03BC CO3BA tabletaerosol for inhalation tabletexpired02195992 02195992 02195976SUPRAX - 20MG/ML SUPRAX - 200MG/TAB SUPRAX - 400MG/TAB 02177099cefixime cefiximeJ01DA tabletoral suspension tabletexpired02195984 02195984 02195984 02177099 TAXOTERE - 20MG/VIAL 00766038cefixime nedocromil sodiumJ01DA tabletcral suspension tabletexpired00766038 TILADE - 2MG/DOSEnot sold cefixineL01CD nedocromil sodiuminjectable solution aerosol for inhalationexpiredR05DAR05DASuspension tabletcefixime J01DA tabletJ01DA 02177090TAXOTERE - 20MG/VIAL docetaxelL01CD injectable solution aerosol for inhalationR05BC codeine/chlorpheniramine polistirexJ01DA tabletColspan="4">SupremationColspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan="4">Colspan= Colspan="4">Colspan= Colspan= Colspan=Colspan= Colspan= Colspan=Colspa | 02202468 02202476 02202484 | OXYCONTIN - 20MG/TAB OXYCONTIN - 40MG/TAB OXYCONTIN - 80MG/TAB | oxycodone hydrochloride oxycodone hydrochloride oxycodone hydrochloride oxycodone hydrochloride choline salicylate/ | NO2AA NO2AA NO2AA | sustained-release tablet sustained-release tablet sustained-release tablet | expired |
| 00555649 00261238INTAL INHALER - 1MG/DOSE INTAL SPINCAPS - 20MG/ CARTRIDGEsodium cromoglycate sodium cromoglycateR03BC R03BCaerosol for inhalation powder for inhalationexpired00638641INTAL SYNCRONER - 1MG/DOSE CARTRIDGEsodium cromoglycate metolazoneR03BC CO3BA tabletaerosol for inhalation tabletexpired006386397MYKROX - 0.5MG/TAB O0848397sodium cromoglycate metolazone polistirexR05DAaerosol for inhalation tabletnot sold02195992SUPRAX - 20MG/ML SUPRAX - 200MG/TAB 02195976cefixime codeine/chlorpheniramine polistirexJ01DA tabletoral suspension tabletexpired02195984SUPRAX - 200MG/TAB C2195984cefixime coetaxelJ01DA tablettablet tabletoral suspension tablet02177099TAXOTERE - 20MG/VIAL O766038cefixime nedocromil sodiumJ01DA R03BC aerosol for inhalationaerosol for inhalationR05DA sustained-release oral suspension02177099TAXOTERE - 20MG/VIAL docetaxelcefixime LO1CD injectable solution aerosol for inhalationR05DA sustained-release oral suspension02177080TAXOTERE - 20MG/VIAL docetaxeldocetaxel LO1CD LO1CDL01CD injectable solution aerosol for inhalationR0BERTS PHARMACEUTICAL CANADA INC.Human: | RHÔNE-P | OULENC RORER CANADA INC | | | | |
| 00638641INTAL SYNCRONER - 1MG/DOSE MYKROX - 0.5MG/TAB 00842702sodium cromoglycate metolazoneR03BC C03BA R05DAaerosol for inhalation tabletnot sold expired02195992SUPRAX - 20MG/ML C2195976cefixime cefiximeJ01DA J01DAoral suspension tabletexpired02195976SUPRAX - 200MG/TAB C2195984cefixime cefiximeJ01DA J01DAoral suspension tabletexpired02195976SUPRAX - 400MG/TAB C2195984cefixime cefiximeJ01DA J01DAtablet02177099TAXOTERE - 20MG/VIAL O0766038docetaxel TILADE - 2MG/DOSEL01CD nedocromil sodiumL01CD R03BC aerosol for inhalationROBERTS PHARMACEUTICAL CANADA INC.Human: | 00555649 | INTAL SPINCAPS - 20MG/ | | | | expired |
| 02195992SUPRAX - 20MG/MLcefiximeJ01DAoral suspension02195976SUPRAX - 200MG/TABcefiximeJ01DAtablet02195984SUPRAX - 400MG/TABcefiximeJ01DAtablet02177099TAXOTERE - 20MG/VIALdocetaxelL01CDinjectable solution02177080TAXOTERE - 80MG/VIALdocetaxelL01CDinjectable solution00766038TILADE - 2MG/DOSEnedocromil sodiumR03BCaerosol for inhalationROBERTS PHARMACEUTICAL CANADA INC.Human: | 00888397 | INTAL SYNCRONER - 1MG/DOSE MYKROX - 0.5MG/TAB | metolazone codeine/chlorpheniramine | C03BA | tablet | |
| Human: | 02195976 02195984 02177099 02177080 | SUPRAX - 200MG/TAB SUPRAX - 400MG/TAB TAXOTERE - 20MG/VIAL TAXOTERE - 80MG/VIAL | cefixime cefixime cefixime docetaxel docetaxel | J01DA J01DA L01CD L01CD | tablet tablet injectable solution injectable solution | |
| | | PHARMACEUTICAL CANADA | INC. | | | |
| | | EMINASE - 30UNIT/VIAL | anistreplase | B01AD | powder for injectable solution intro | oduced (nas) |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------------------|--|--|----------------|---|----------------------|
| SANOFI W | /INTHROP INC. | | | | |
| Human: | COROTROPE - 5MG/CAP | milrinone | C01CE | caplet | not sold |
| | COROTROPE - 7.5MG/CAP | milrinone | C01CE | caplet | not sold |
| | COROTROPE - 10MG/CAP | milrinone | C01CE | caplet | not sold |
| 02236913 | FRAXIPARINE - 9500UNIT/ML | nadroparin calcium | B01AB | injectable solution | not sold |
| 02017644 02006391 | INOCOR - 5MG/ML PRIMACOR - 1MG/ML | amrinone lactate milrinone lactate | C01CE C01CE | injectable solution injectable solution | expired |
| 02230643 | SKELID - 200MG/TAB | tiludronate disodium | M05BA | tablet | not sold |
| SCHERING | G CANADA INC. | | | | |
| Human: | | | | - inter- ent | اما معام |
| 02070855 02070871 | ALDERM - 0.5MG/G ALDERM - 0.5MG/G | alclometasone dipropionate alclometasone dipropionate | D07AB D07AB | ointment cream | not sold not sold |
| 02165465 | CEDAX - 200MG/CAP | ceftibuten | J01DA | capsule | not sold |
| 02165473 | CEDAX - 400MG/CAP | ceftibuten | J01DA | capsule | not sold |
| 02165449 | CEDAX - 18MG/ML | ceftibuten | J01DA | powder for oral suspension | not sold |
| 02165457 01970399 | CEDAX - 36MG/ML CHLOR-TRIPOLON N.D. 5/120 | ceftibuten | J01DA R06AX | powder for oral suspension | not sold |
| 01970399 | CHEUR-TRIPOLON N.D. 5/120 | loratadine/pseudoephedrine sulfate | RUOAA | tablet | |
| 02019973 | CLARITIN - 1MG/ML | loratadine | R06AX | syrup | |
| 00782696 | CLARITIN - 10MG/TAB | loratadine | R06AX | tablet | |
| 01945157 | CLARITIN EXTRA 5/120 | loratadine/pseudoephedrine sulfate | R06AX | tablet | |
| 00851736 | ELOCOM - 1MG/G | mometasone furoate | D07AC | ointment | |
| 00851744 | ELOCOM - 1MG/G | mometasone furoate | D07AC | cream | |
| 00871095 | ELOCOM - 1MG/ML | mometasone furoate | D07AC L03AA | lotion iniectable solution | |
| 00889067 00705896 | INTRON-A - 5000000UNIT/ML INTRON-A - 3000000UNIT/VIAL | interferon alfa-2b interferon alfa-2b | LOSAA | powder for injectable suspension | |
| 00705918 | INTRON-A - 5000000UNIT/VIAL | interferon alfa-2b | LO3AA | powder for injectable suspension | |
| 00705926 | INTRON-A - 10000000UNIT/VIAL | interferon alfa-2b | L03AA | powder for injectable suspension | |
| 02231651 | INTRON-A - 18000000UNIT/VIAL | interferon alfa-2b | L03AA | powder for injectable suspension | introduced |
| 02231652 02231653 | INTRON-A - 30000000UNIT/VIAL INTRON-A - 50000000UNIT/VIAL | interferon alfa-2b interferon alfa-2b | L03AA L03AA | powder for injectable suspension powder for injectable suspension | not sold not sold |
| 00713376 | K-DUR - 1500MG/TAB | potassium chloride | A12BA | sustained-release tablet | 1101 3010 |
| 02230921 | LEUCOMAX - 0.15MG/VIAL | molgramostim | L03AA | powder for injectable solution | not sold |
| 02230922 | LEUCOMAX - 0.3MG/VIAL | molgramostim | LO3AA | powder for injectable solution | not sold |
| 02230923 02230924 | LEUCOMAX - 0.4MG/VIAL LEUCOMAX - 0.7MG/VIAL | molgramostim | L03AA L03AA | powder for injectable solution | not sold |
| 02230924 | LOTRIDERM 0.5/10 | molgramostim betamethasone dipropionate/ | | powder for injectable solution cream | not sold |
| | | clotrimazole | | oroann | |
| 00503363 | NETROMYCIN - 25MG/ML | netilmicin sulfate | J01GB | injectable solution | not sold |
| 00503371 00503398 | NETROMYCIN - 50MG/ML NETROMYCIN - 100MG/ML | netilmicin sulfate netilmicin sulfate | J01GB J01GB | injectable solution injectable solution | |
| 01911910 | NITRO-DUR 0.2 - 40MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 02213370 | NITRO-DUR 0.3 - 60MG/PATCH | nitroglycerin | C01DA | transdermal patch | |
| 01911902 | NITRO-DUR 0.4 - 80MG/PATCH | nitroglycerin | CO1DA | transdermal patch | |
| 01911929 02011271 | NITRO-DUR 0.6 - 120MG/PATCH NITRO-DUR 0.8 - 160MG/PATCH | nitroglycerin nitroglycerin | C01DA C01DA | transdermal patch transdermal patch | |
| | | IIIIIOgiyceIIII | GUIDA | liansuerniai palon | |
| Veterinary: 02231742 | AQUAFLOR - 500000MG/KG | florfenicol | | powder | |
| 00882887 | DURA SE-120 - 360MG/BOLUS | selenite sodium | | bolus | not sold |
| 02216558 | NUFLOR - 300MG/ML | florfenicol | | injectable solution | |
| 02142155 | OPTIMMUNE - 2MG/G | cyclosporine | | ophthalmic ointment | |
| | ANADA INC. | | | | |
| Human: 02205971 | AMBIEN - 5MG/TAB | zolpidem tartrate | N05CG | tablet | not sold |
| 02205998 | AMBIEN - 10MG/TAB | zolpidem tartrate | N05CG | tablet | not sold |
| 02231676 | CHRONOVERA - 180MG/TAB | verapamil hydrochloride | C08DA | sustained-release tablet | not sold |
| 02231677 01946188 | CHRONOVERA - 240MG/TAB | verapamil hydrochloride | C08DA C07AB | sustained-release tablet tablet | not sold |
| 01946188 | KERLONE - 10MG/TAB KERLONE - 20MG/TAB | betaxolol hydrochloride betaxolol hydrochloride | CO7AB CO7AB | tablet | expired expired |
| 02010925 | MAXAQUIN - 400MG/TAB | lomefloxacin | J01MA | tablet | not sold |
| 00874213 | NITRODISC 0.2 - 16MG/PATCH | nitroglycerin | C01DA | transdermal patch | not sold |
| 00874221 | NITRODISC 0.3 - 24MG/PATCH | nitroglycerin | CO1DA | transdermal patch | not sold |
| 00874248 02188783 | NITRODISC 0.4 - 32MG/PATCH SYNAREL - 2MG/ML | nitroglycerin nafarelin acetate | C01DA H01CA | transdermal patch nasal solution | not sold |
| 02187108 | SYNPHASIC 1.0-0.5/0.035 | norethindrone/ethinyl estradio | | tablet | |
| 02187116 | SYNPHASIC 1.0-0.5/0.035 | norethindrone/ethinyl estradio | | tablet | |
| 66 | PMPRB | | | | |
| 66 | 1997 | | | | |

| SMITHKLINE BEECHAM PHARMA INC. Human: 01911937 AUGMENTIN 25/6.25 amoxicillin trihydrate/ clavulanate potassium JOTCR oral suspension 01911937 AUGMENTIN 250/125 amoxicillin trihydrate/ clavulanate potassium JOTCR tablet 01911945 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium JOTCR tablet 01911946 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium JOTCR tablet 01911947 BACTROBAN - 20M6/G mupirocin mupirocin DO6AX ointment 01919490 CEFIZOX - 1000MG/VIAL cefizoxime sodium JOTCR tablet 01919682 CLAVULIN 256/25 amoxicillin trihydrate/ clavulanate potassium JOTCR tablet 01916866 CLAVULIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium JOTCR tablet 02229650 COREG - 3 125MG/TAB carvediol COTAG tablet 02229651 COREG - 6 25MG/TAB carvediol COTAG tablet 02229652 COREG - 6 25MG/TAB carvediol COTAG tablet | |
|--|------------------|
| 01911937 AUGMENTIN 25/6.25 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911953 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911964 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911964 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911964 AUGMENTIN 50/12.5 amoxicillin trihydrate/ amoxicillin trihydrate/ clavulanate potassium J01CR tablet 0191687 BACTROBAN - 20MG/G TPICO mupirocin mupirocin J06AX ointment 0191686 CLAVULIN 25/6.25 amoxicillin trihydrate/ amoxicillin trihydrate/ J010FR J01CR tablet 01916866 CLAVULIN 25/12.5 amoxicillin trihydrate/ amoxicillin trihydrate/ J010FR J01CR tablet 02229650 COREG - 3.125MG/TAB clavulanate potassium J01CR tablet 02229651 COREG - 6.25MG/TAB carvediol C07AG tablet 02229652 COREG - 1.26MG/TAB carvediol C07AG tablet 02229105 COREG - 2.5MG/TAB carvediol C07AG tablet 02229110 <t< td=""><td></td></t<> | |
| 01911953 AUGMENTIN 250/125 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911945 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01911961 AUGMENTIN 500/125 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01916947 BACTROBAN - 20MG/G mupirocin advinate potassium J01DA powder for injectable soluti 01919490 CEFIZOX - 1000MG/VIAL cefitzoxime sodium J01DA powder for injectable soluti 01916866 CLAVULIN 250/125 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01916876 CLAVULIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 02229650 COREG - 3.125MG/TAB carvediol COTAG tablet 02229651 COREG - 2.5MG/TAB carvediol COTAG tablet 02229105 | not sold |
| 01911945 AUGMENTIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium oral suspension 01911961 AUGMENTIN 500/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01916947 BACTROBAN - 20MG/G mupirocin clavulanate potassium D06AX nasal ointment 01916940 CEFIZOX - 1000MG/IAL ceftizoxime sodium J01DA powder for injectable soluti 01916862 CLAVULIN 256/2.5 amoxicillin trihydrate/ J01CR ratsupension 01916866 CLAVULIN 250/12.5 amoxicillin trihydrate/ J01CR roral suspension 01916874 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR roral suspension 01916878 CAVULIN 500/125 amoxicillin trihydrate/ J01CR roral suspension 01916878 CAVULIN 500/125 amoxicillin trihydrate/ J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol CO7AG tablet 02229650 COREG - 3.125MG/TAB carvedilol CO7AG tablet 02229650 COREG - 3.25MG/TAB carvedilol CO7AG tablet 02229610 COREG - 2.5MG/TAB | not sold |
| 01911961 AUGMENTIN 500/125 amoxicilin trihydrate/ J016R tablet 01916947 BACTROBAN - 20MG/G mupirocin D06AX nasal ointment 01919490 CEFIZOX - 1000MG/VIAL ceftizoxime sodium J01DA powder for injectable soluti 01919804 CEFIZOX - 2000MG/VIAL ceftizoxime sodium J01CR oral suspension 01916862 CLAVULIN 256/25 amoxicilin trihydrate/ J01CR oral suspension 01916864 CLAVULIN 250/125 amoxicilin trihydrate/ J01CR tablet 01916874 CLAVULIN 50/12.5 amoxicilin trihydrate/ J01CR tablet 01916874 CLAVULIN 50/12.5 amoxicilin trihydrate/ J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol C07AG tablet 02229651 COREG - 12.5MG/TAB carvedilol C07AG tablet 02229652 COREG - 12.5MG/TAB carvedilol C07AG tablet 02229651 COREG - 12.5MG/TAB famciclovir J05AB tablet 02229105 FAMVR - 25MG/TAB famciclovir J05AB tablet < | not sold |
| 01916947 BACTROBAN NASAL - 20MG/G mupirocin D06AX nasal ointment 01919490 CEFIZOX - 1000MG/VIAL ceftizoxime sodium J01DA powder for injectable soluti 01919490 CEFIZOX - 2000MG/VIAL ceftizoxime sodium J01DA powder for injectable soluti 01916882 CLAVULIN 256/25 amoxicillin trihydrate/ J01CR oral suspension 01916884 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 01916885 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 01916858 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol C07AG tablet 02229651 COREG - 5.25MG/TAB carvedilol C07AG tablet 02229652 COREG - 12.5MG/TAB carvedilol C07AG tablet 02229650 COREG - 5.25MG/TAB carvedilol C07AG tablet 02229651 COREG - 12.5MG/TAB carvedilol C07AG tablet 02229105 FAMVR - 25MG/TAB farciclovir J05AB tablet < | not sold |
| 01919490 CEFIZOX - 1000MG/VIAL ceffizoxime sodium J01DA powder for injectable soluti 01919504 CEFIZOX - 2000MG/VIAL ceffizoxime sodium J01DA powder for injectable soluti 01916882 CLAVULIN 25/6.25 amoxicillin trihydrate/ J01CR clavulanate potassium 01916886 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR oral suspension 01916886 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol C07AG tablet 02229652 COREG - 12.5MG/TAB carvedilol C07AG tablet 0222915 FAMVIR - 125MG/TAB famciclovir J05AB tablet 0222911 FAMVIR - 125MG/TAB famciclovir J05AB tablet 0222912 FAMVIR - 125MG/TAB famciclovir J05AB tablet | not cold |
| 01919504 CEFIZOX - 2000MG/VIAL ceftizoxime sodium J01DA jowder for injectable soluti 01916882 CLAVULIN 25/6.25 amoxicillin trihydrate/ J01CR oral suspension 01916866 CLAVULIN 250/125 amoxicillin trihydrate/ J01CR tablet 01916874 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 01916878 CLAVULIN 50/12.5 amoxicillin trihydrate/ J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol C07AG tablet 02229651 COREG - 6.25MG/TAB carvedilol C07AG tablet 02229652 COREG - 1.25MG/TAB carvedilol C07AG tablet 02229653 COREG - 2.5MG/TAB carvedilol C07AG tablet 02229105 FAMVIR - 125MG/TAB famciclovir J05AB tablet 02229110 FAMVIR - 125MG/TAB famciclovir J05AB tablet 02229121 FAMVIR - 125MG/TAB famciclovir J05AB tablet 02217702 FAMVIR - 125MG/TAB famciclovir J05AB tablet 02229110 FAM | not sold |
| 01916882 CLAVULIN 25/6.25 amoxicillin trihydrate/ clavulanate potassium J01CR for al suspension 01916866 CLAVULIN 250/125 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01916874 CLAVULIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 01916876 CLAVULIN 50/12.5 amoxicillin trihydrate/ clavulanate potassium J01CR tablet 02229650 COREG - 3.125MG/TAB carvedilol C07AG tablet 02229651 COREG - 6.25MG/TAB carvedilol C07AG tablet 02229652 COREG - 12.5MG/TAB carvedilol C07AG tablet 02229610 FMCKR-B hepattiis B vaccine J07BC injectable solution 01919431 ENGERIX-B hepattiis A vaccine J07BC injectable suspension 02229110 FAMVIR - 125MG/TAB famciclovir J05AB tablet 0229110 FAMVIR - 120MG/TAB famciclovir J05AB tablet 0217102 FAMVIR - 120MG/TAB famciclovir J05AB tablet 0218707 HAVRIX 720 JUNIOR - hepatitis A vaccine <td< td=""><td></td></td<> | |
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| 1440EUNIT/ML(inactivated)02231116HYCAMTIN - 4MG/VIALtopotecan hydrochlorideL01XXpowder for injectable solution02230307INFANRIXDaPT vaccineJ07AJinjectable suspension02154471KREDEX - 25MG/TABcarvedilolC07AGtablet02154498KREDEX - 50MG/TABcarvedilolC07AGtablet02185881KYTRIL - 1MG/TABgranisetron hydrochlorideA04AAtablet02088371KYTRIL INJECTION - 1MG/MLgranisetron hydrochlorideA04AAinjectable solution01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABnabumetoneM01AXtablet02027835RELAFEN - 500MG/TABnabumetoneM01AXtablet02231352RELAFEN - 750MG/TABnabumetoneM01AXtablet02232567REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02230307INFANRIXDaPT vaccineJ07AJinjectable suspension02154471KREDEX - 25MG/TABcarvedilolC07AGtablet02154498KREDEX - 50MG/TABcarvedilolC07AGtablet02185881KYTRIL - 1MG/TABgranisetron hydrochlorideA04AAtablet0208371KYTRIL INJECTION - 1MG/MLgranisetron hydrochlorideA04AAinjectable solution01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970437PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABnabumetoneM01AXtablet02083538RELAFEN - 500MG/TABnabumetoneM01AXtablet02232567REQUIP - 0.25MG/TABnabumetoneM01AXtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | introduced |
| 02154471KREDEX - 25MG/TABcarvedilolC07AGtablet02154498KREDEX - 50MG/TABcarvedilolC07AGtablet02185881KYTRIL - 1MG/TABgranisetron hydrochlorideA04AAtablet02088371KYTRIL INJECTION - 1MG/MLgranisetron hydrochlorideA04AAinjectable solution01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 100MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02231352RELAFEN - 750MG/TABnabumetoneM01AXtablet02232567REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02154498KREDEX - 50MG/TABcarvedilolC07AGtablet02185881KYTRIL - 1MG/TABgranisetron hydrochlorideA04AAtablet02088371KYTRIL INJECTION - 1MG/MLgranisetron hydrochlorideA04AAinjectable solution01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet0223152RELAFEN - 750MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABnabumetoneM01AXtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | not sold |
| 02185881KYTRIL - 1MG/TABgranisetron hydrochlorideA04AAtablet02088371KYTRIL INJECTION - 1MG/MLgranisetron hydrochlorideA04AAinjectable solution01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet0203558RELAFEN - 750MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | not sold |
| 02088371KYTRIL INJECTION - 1MG/ML 01970410granisetron hydrochloride vefonicid sodiumA04AA J01DAinjectable solution powder for injectable soluti powder for injectable soluti01970410MONOCID - 500MG/VIAL 01970429cefonicid sodiumJ01DA powder for injectable soluti powder for injectable soluti02027887PAXIL - 10MG/TAB 01940481paroxetine hydrochlorideN06AB paroxetine hydrochloridetablet01940473PAXIL - 20MG/TAB 02027895paroxetine hydrochlorideN06AB paroxetine hydrochloridetablet02027895PAXIL - 50MG/TAB 02083531paroxetine hydrochlorideN06AB paroxetine hydrochloridetablet02083538RELAFEN - 500MG/TAB 02033538nabumetone nabumetoneM01AX M01AX M01AXtablet02232565REQUIP - 0.25MG/TAB 02232566nabumetone REQUIP - 0.5MG/TAB M01ABmotione ropinirole hydrochlorideN04BC N04BCtablet02232567REQUIP - 1MG/TAB REQUIP - 1MG/TABropinirole hydrochloride ropinirole hydrochlorideN04BC N04BCtablet | not sold |
| 01970410MONOCID - 500MG/VIALcefonicid sodiumJ01DApowder for injectable soluti01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABnabumetoneM01AXtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 01970429MONOCID - 1000MG/VIALcefonicid sodiumJ01DApowder for injectable soluti02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABparoxetine hydrochlorideN06ABtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet0223352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02027887PAXIL - 10MG/TABparoxetine hydrochlorideN06ABtablet01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 01940481PAXIL - 20MG/TABparoxetine hydrochlorideN06ABtablet01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | n expired |
| 01940473PAXIL - 30MG/TABparoxetine hydrochlorideN06ABtablet02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | introduced |
| 02027895PAXIL - 50MG/TABparoxetine hydrochlorideN06ABtablet02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02083531RELAFEN - 500MG/TABnabumetoneM01AXtablet02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02083558RELAFEN - 750MG/TABnabumetoneM01AXtablet02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | not sold |
| 02231352RELAFEN - 1000MG/TABnabumetoneM01AXtablet02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | |
| 02232565REQUIP - 0.25MG/TABropinirole hydrochlorideN04BCtablet02232566REQUIP - 0.5MG/TABropinirole hydrochlorideN04BCtablet02232567REQUIP - 1MG/TABropinirole hydrochlorideN04BCtablet | not sold |
| 02232566 REQUIP - 0.5MG/TAB ropinirole hydrochloride N04BC tablet 02232567 REQUIP - 1MG/TAB ropinirole hydrochloride N04BC tablet | not sold |
| 02232567 REQUIP - 1MG/TAB ropinirole hydrochloride N04BC tablet | introduced (nas) |
| | not sold |
| 02232568 REQUIP - 2MG/IAB ropinirole hydrochloride N04BC tablet | introduced (nas) |
| | introduced (nas) |
| 02232569 REQUIP - 5MG/TAB ropinirole hydrochloride N04BC tablet | introduced (nas) |
| 01927280 TAGAMET - 6MG/ML cimetidine hydrochloride A02BA injectable solution | |
| 01916750 TAGAMET - 60MG/ML cimetidine hydrochloride A02BA oral solution | |
| 01916807 TAGAMET - 150MG/ML cimetidine hydrochloride A02BA injectable solution | |
| 01916793 TAGAMET - 200MG/TAB cimetidine A02BA tablet | not sold |
| 01916815 TAGAMET - 300MG/TAB cimetidine A02BA tablet | |
| 01916785 TAGAMET - 400MG/TAB cimetidine A02BA tablet | |
| 01916777 TAGAMET - 600MG/TAB cimetidine A02BA tablet | |
| 01916769 TAGAMET - 800MG/TAB cimetidine A02BA tablet | not sold |
| 01916920 TICAR - 1000MG/VIAL ticarcillin disodium J01CA powder for injectable soluti | |
| 01916912 TICAR - 3000MG/VIAL ticarcillin disodium J01CA powder for injectable soluti | |
| 01916904 TICAR - 6000MG/VIAL ticarcillin disodium J01CA powder for injectable soluti | |
| 01916890 TICAR - 20000MG/VIAL ticarcillin disodium J01CA powder for injectable soluti | n not sold |

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|--|---|--|--|---|--|
| 00803480 01916939 | TICAR - 30000MG/VIAL TIMENTIN 3000/100 | ticarcillin disodium ticarcillin disodium/ | J01CA J01CR | powder for injectable solutior powder for injectable solutior | |
| 02230578 | TWINRIX 720/20 | clavulanate potassium combined hepatitis A & B vaccine | J07BC | injectable suspension | introduced |
| | PHARMA INC. | | | | |
| Human: 01919334 01919342 01919350 01919369 02224151 02229453 | LUVOX - 25MG/TAB LUVOX - 50MG/TAB LUVOX - 75MG/TAB LUVOX - 100MG/TAB PANTO-BYK - 40MG/TAB PANTOLOC - 40MG/TAB | fluvoxamine maleate fluvoxamine maleate fluvoxamine maleate fluvoxamine maleate pantoprazole sodium pantoprazole sodium | N06AB N06AB N06AB N06AB A02BC A02BC | tablet tablet tablet tablet tablet tablet | expired expired expired expired not sold introduced (nas) |
| WARNER | LAMBERT CANADA INC. (PAF | RKE-DAVIS) | | | |
| Human: 01947664 01947672 01947680 01947699 | ACCUPRIL - 5MG/TAB ACCUPRIL - 10MG/TAB ACCUPRIL - 20MG/TAB ACCUPRIL - 40MG/TAB COGNEX - 10MG/CAP COGNEX - 20MG/CAP COGNEX - 30MG/CAP COGNEX - 40MG/CAP | quinapril hydrochloride quinapril hydrochloride quinapril hydrochloride quinapril hydrochloride tacrine hydrochloride tacrine hydrochloride tacrine hydrochloride tacrine hydrochloride | C09AA C09AA C09AA C09AA N07AA N07AA N07AA N07AA | tablet tablet tablet capsule capsule capsule capsule capsule | introduced (peo) |
| 02230711 02230713 02230714 02053136 | LIPITOR - 10MG/TAB LIPITOR - 20MG/TAB LIPITOR - 40MG/TAB MAXAIR - 0.25MG/DOSE | atorvastatin calcium atorvastatin calcium atorvastatin calcium pirbuterol acetate | C10AA C10AA C10AA R03AC | tablet tablet tablet aerosol for inhalation | introduced (nas) introduced (nas) introduced (nas) |
| 02033130 02231095 02231096 | REZULIN - 200MG/TAB REZULIN - 400MG/TAB | troglitazone troglitazone | A10BG A10BG | tablet tablet | not sold not sold |
| | OD-SQUIBB PHARMACEUTIC | LS INC. | | | |
| Human: 02010291 01909150 01962701 01962728 | HALOG - 1MG/ML LACHYDRIN - 120MG/ML ULTRAVATE - 0.5MG/G ULTRAVATE - 0.5MG/G | halcinonide lactic acid halobetasol propionate halobetasol propionate | D07AD D02AX D07AD D07AD | topical solution lotion cream ointment | expired |
| WYETH-A | YERST CANADA INC. | | | | |
| Human: 02215063 02215071 02036274 02036428 02103664 02103664 02103680 02103699 02103702 02103710 02126206 | ACEL-IMUNE ACEL-P ALREDASE - 200MG/TAB CEFOTAN - 1000MG/VIAL CEFOTAN - 2000MG/VIAL EFFEXOR - 12.5MG/TAB EFFEXOR - 25MG/TAB EFFEXOR - 37.5MG/TAB EFFEXOR - 50MG/TAB EFFEXOR - 75MG/TAB EFFEXOR - 100MG/TAB HIBTITER | DaPT vaccine acellular pertussis vaccine tolrestat cefotetan disodium venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride venlafaxine hydrochloride vaccine - hemophilus influenzae b | J07AJ J07AJ A10XA J01DA J01DA N06AA N06AA N06AA N06AA N06AA N06AA N06AA | injectable suspension injectable suspension tablet powder for injectable solution powder for injectable solution tablet tablet tablet tablet tablet tablet | |
| 02042231 02042258 02042266 02042274 00673013 00673021 00673048 02042304 02042312 01919644 00665117 00665126 | INDERAL-L.A 60MG/CAP INDERAL-L.A 80MG/CAP INDERAL-L.A 120MG/CAP INDERAL-L.A 160MG/CAP MAGNACEF - 500MG/VIAL MAGNACEF - 1000MG/VIAL MICRO-K EXTENCAPS - 600MG/CAP MICRO-K EXTENCAPS - 750MG/CAP MICRO-K LS - 1875MG/PCK MINOCIN - 50MG/TAB MINOCIN - 100MG/TAB | propranolol hydrochloride propranolol hydrochloride propranolol hydrochloride propranolol hydrochloride ceftazidime pentahydrate ceftazidime pentahydrate ceftazidime pentahydrate potassium chloride potassium chloride potassium chloride minocycline hydrochloride | C07AA C07AA C07AA J01DA J01DA J01DA A12BA A12BA A12BA J01AA J01AA | sustained-release capsule sustained-release capsule sustained-release capsule sustained-release capsule powder for injectable solutior powder for injectable solutior sustained-release capsule sustained-release capsule oral suspension tablet tablet | not sold |

PMPRB

| DIN/GP | Brand Name | Generic Name | ATC* | Dosage Form | Comments |
|----------|------------------------------------|---|--------|--|-------------------|
| 02170728 | MONOCOR - 5MG/TAB | bisoprolol fumarate | C07AB | tablet | not sold |
| 02170701 | MONOCOR - 10MG/TAB | bisoprolol fumarate | C07AB | tablet | not sold |
| 02173468 | NOVANTRONE - 2MG/ML | mitoxantrone hydrochloride | L01DB | injectable solution | |
| 00564974 | PIPRACIL - 2000MG/VIAL | piperacillin sodium | J01CA | powder for injectable solution | |
| 00564982 | PIPRACIL - 3000MG/VIAL | piperacillin sodium | J01CA | powder for injectable solution | |
| 00564990 | PIPRACIL - 4000MG/VIAL | piperacillin sodium | J01CA | powder for injectable solution | |
| 00857548 | PIPRACIL - 40000MG/VIAL | piperacillin sodium | J01CA | powder for injectable solution | |
| 02230175 | PNU-IMUNE 23 | vaccine - polyvalent | J07AL | injectable suspension | introduced |
| 01970348 | REV-EYES - 25MG/VIAL | pneumoccocal | S01EX | anthalmia colution | not cold |
| 01970346 | TAZOCIN 2000/250 | dapiprazole hydrochloride piperacillin sodium/ | J01CR | ophthalmic solution powder for injectable solution | not sold |
| 02170017 | TA2001N 2000/230 | tazobactam sodium | J010h | powder for injectable solution | |
| 02170795 | TAZOCIN 3000/375 | piperacillin sodium/ | J01CR | powder for injectable solution | |
| 02110100 | 1120011 0000/010 | tazobactam sodium | 001011 | | |
| 02170809 | TAZOCIN 4000/500 | piperacillin sodium/ | J01CR | powder for injectable solution | |
| | | tazobactam sodium | | , · · · · · · · · · · · · · · · · · · · | |
| 02093405 | TETRAMUNE | DPT-hemophilus b conjugate | J07CA | injectable suspension | not sold |
| | | vaccine | | | |
| ZENECA F | PHARMA INC. | | | | |
| Human: | | | | | |
| 02224135 | ARIMIDEX - 1MG/TAB | anastrozole | L02BG | tablet | |
| 02188880 | BREVIBLOC - 10MG/ML | esmolol hydrochloride | C07AB | injectable solution | |
| 02188872 | BREVIBLOC - 100MG/ML | esmolol hydrochloride | C07AB | injectable solution | not sold |
| 02188864 | BREVIBLOC - 250MG/ML | esmolol hydrochloride | C07AB | injectable solution | |
| 02184478 | CASODEX - 50MG/TAB | bicalutamide | L02BB | tablet | |
| 02218488 | MERREM - 500MG/VIAL | meropenem | J01DH | powder for injectable solution | |
| 02218496 | MERREM - 1000MG/VIAL | meropenem | J01DH | powder for injectable solution | |
| 02219018 | MERREM ADD-VANTAGE - 500MG/VIAL | meropenem | J01DH | powder for injectable solution | not sold |
| 02219026 | MERREM ADD-VANTAGE - | meropenem | J01DH | powder for injectable solution | not sold |
| | 1000MG/VIAL | | | President and a second s | |
| 02236951 | SEROQUEL - 25MG/TAB | quetiapine fumarate | N05AX | tablet | not sold |
| 02236952 | SEROQUEL - 100MG/TAB | quetiapine fumarate | N05AX | tablet | not sold |
| 02236953 | SEROQUEL - 200MG/TAB | quetiapine fumarate | N05AX | tablet | not sold |
| 00007400 | | al a a fluina na a | | independence and a start of the | tue dure ed (mee) |

lisinopril/hydrochlorothiazide C09BA lisinopril/hydrochlorothiazide C09BA

lisinopril/hydrochlorothiazide C09BA

desflurane

raltitrexed

lisinopril

lisinopril

lisinopril

lisinopril

goserelin acetate goserelin acetate

N01AB

L01BA

C09AA

C09AA

C09AA

C09AA

L02AE

L02AE

tablet

tablet

tablet

tablet

tablet

tablet

tablet

inhalation anesthetic

injectable implant injectable implant

powder for injectable solution

02227428 02229566

02103729

02045737

02045729

02049333

02049376

02049384

00839345

02049325

02225905

SUPRANE

TOMUDEX - 2MG/VIAL

ZESTORETIC 10/12.5 ZESTORETIC 20/12.5 ZESTORETIC 20/25

ZESTRIL - 5MG/TAB ZESTRIL - 10MG/TAB ZESTRIL - 20MG/TAB

ZESTRIL - 40MG/TAB ZOLADEX - 3.6MG/VIAL ZOLADEX LA - 10.8MG/VIAL

not sold

introduced (nas)