Environment Canada
Environmental Protection Service
Pacific Region
Yukon Branch

A BIOLOGICAL AND WATER QUALITY ASSESSMENT AT A PLACER MINE ON BARLOW CREEK, YUKON TERRITORY

Regional Program Report 84-16

bу

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#### ABSTRACT

An assessment of water quality and biological conditions was carried out at a placer gold mine on Barlow Creek from June to September 1982. The water quality, sediment and bottom fauna characteristics were investigated at four sample stations.

Water quality parameters were influenced by the placer mining activity. During periods of sluicing activity downstream stations showed elevated levels for suspended sediments (NFR), settleable solids, filterable residue (FR), turbidity, extractable heavy metals, phosphorous and nitrites. Dissolved oxygen was slightly lower at downstream stations compared to an upstream control station.

The sediment composition of the stream bottom was seen to vary considerably from station to station and at the control station during the season. At downstream stations the sediment composition was significantly higher in percentage of fines (fine sands and silt) on sluicing days. These fines tended to be re-suspended and moved downstream during periods of non-sluicing or higher flows so that the stream bottom composition appears to recover to a composition typical of the natural stream over time. In general, there was an increase in levels of sediment metals at downstream stations during the season which reflected the general increase of fine sediments, especially during periods of sluicing activity.

Bottom fauna types, numbers and diversity were determined for each station during the season. Diversity indices were variable during the season and comparison of the control station with downstream stations shows a general overall decrease in numbers and diversity index values at downstream stations. The dominant groups in terms of abundance were the Ephemeroptera, Diptera and Plecoptera whose numbers varied in response to seasonal change and habitat alterations.

#### RÉSUMÉ

De juin à septembre 1982, on a procédé à une analyse de la qualité de l'eau et des conditions biologiques de la rivière Barlow, à l'emplacement d'un gisement aurifère alluvial. On a analysé la qualité de l'eau, les caractéristiques des sédiments et de la faune profonde à quatre points différents d'échangillonnage.

L'activité minière a eu un effet sur les paramètres de la qualité de l'eau. Durant les périodes de lavage du minerai on a relevé une grande turbidité dans les stations situées en aval, une proportion élevée de sédiments en suspension (résidus non filtrables), de solides en voie de sédimentation, de résidus filtrables, de métaux lourds extractibles, de phosphore et de nitrites. Dans les stations situées en aval on a relevé une proportion d'oxygène dissous légèrement inférieure à celle relevée dans une station située en amont.

On a constaté que durant la saison la composition des sédiments prélevés au fond du cours d'eau variait considérablement d'une station à l'autre, y compris la station de contrôle, Dans les stations situées en aval, la composition des sédiments comprenait un pourcentage sensiblement plus élevé de fins (sable fin et vase) les jours de lavage du minerai. Durant les périodes sans lavage ou les périodes de hautes eaux, ces fins étaient remis en suspension et transportés vers l'aval, ce qui redonnait progressivement au fond du cours d'eau sa composition naturelle. En règles générale, pendant la saison, on a constaté un accroissement de la proportion des métaux dans les sédiments aux stations situées en aval, ce qui correspondait à une plus grande accumulation des sédiments fins, en particulier durant les périodes de lavage du minerai.

Durant la saison, on a évalué pour chaque station les types, nombre et diversité de la faune profonde. Pendant ce même temps on a relevé des variations dans l'indice de diversité; en comparant les résultats de la station de contrôle et des stations situées en aval, on a constaté une diminution générale en nombre et en diversité dans les stations situées en aval. En importance numérique les groupes dominants étaient les éphéméroptères, les diptères et les plécoptères dont le nombre variait selon les changements saisonniers et les modifications de l'habitat.

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#### 1 INTRODUCTION

A study of water quality, sediment composition and bottom fauna, was conducted on six different dates between June and September, 1982. This study was carried out, by the Environmental Protection Service - Yukon Branch, in the watershed of Barlow Creek in the Stewart River area (see Figure 1). The purpose of the study was to gather information on the quality of the stream and to enable the Environmental Protection Service to evaluate any impact to the receiving waters from placer mining operations.

## 1.1 Background

The Stewart River was one of the first areas in the Yukon to attract gold miners. Gold was discovered in paying quantities from its bars for many years beginning in 1883 with production reachings it peak between 1885 and 1887 (Bostock, 1957). In 1895 coarse gold was first discovered on tributary streams of the Stewart River, and in 1897 the first placer claims were staked on Clear Creek, the creek to which Barlow Creek is a tributary (Queenstake, 1982). From that time up to the 1940's, hand mining was the only form of mining that took place on Clear Creek and a number of its tributaries including Squaw Creek, Barlow Creek and Zinc Creek. According to records of the Mining Recorder for Barlow Creek, 104 claims were staked by Syndicate Lyonnais du Klondike from 1901 to 1903. In 1912 many five year placer mining grants were issued to individuals working single claims.

With the increase in gold prices in 1973, Clear Creek and the majority of its tributaries, including Barlow Creek, were staked by various individuals and companies. In 1981, two new operations were initiated on Barlow Creek and Zinc Creek. Dawson Mining Equipment, one of the larger companies on Zinc Creek, undertook an intensive mining program in 1981 but was not active in 1982 (Maxwell, J. Pers. Comm.). In 1982 there was some small scale testing activity approximately 6 km upstream of the study area on Zinc Creek. The operation on Barlow Creek where the study area is located was begun in 1981 and has the potential for a lengthy operation in that 66 claims were held by the operators.

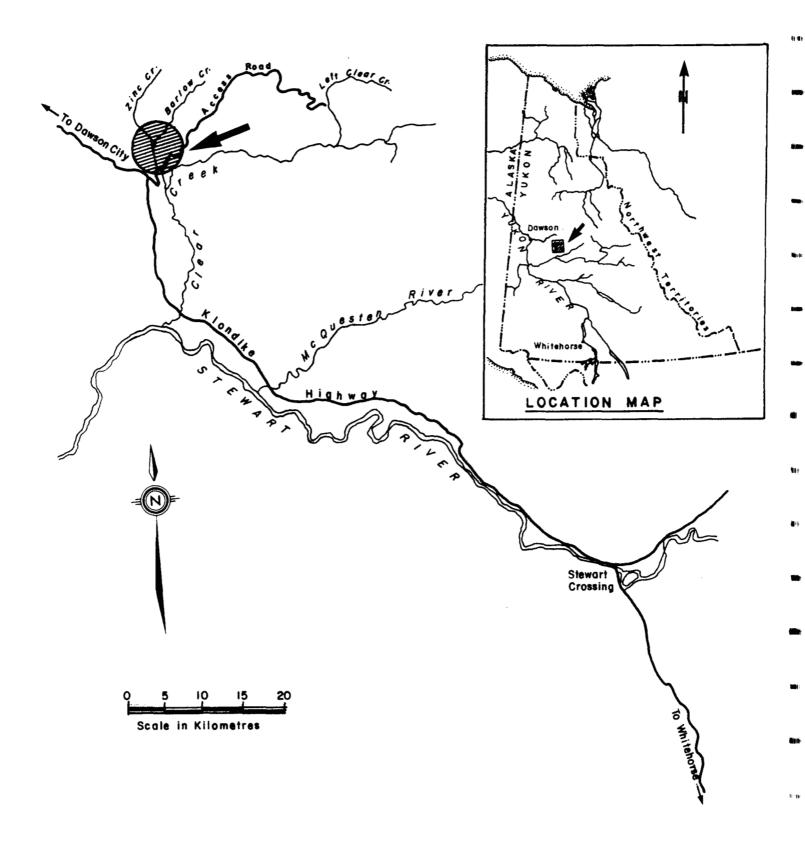


FIGURE I LOCATION OF BARLOW CREEK STUDY AREA

#### 2 STUDY AREA

The study area lies approximately 400 km northwest of Whitehorse and 100 km southeast of Dawson City (see Figure 1). The property has easy access via a 4 km secondary road which leaves the Klondike Highway at km 610. Barlow Creek is one of many small creeks in the area that flows into Clear Creek which in turn is one of the main tributaries of the Stewart River. This network of creek and river valleys is surrounded and separated by uplands which characterize the Stewart Plateau, a subdivision of the Yukon Plateau. This plateau rises steeply along the north-east side of the Tintina Valley south of the McQuesten River (Bostock, 1948).

The current study site is located on the lower reach of Barlow Creek with elevations ranging from 555 m to 580 m. Sample stations were selected to include an upstream control station which would be unaffected by mining activity; a station to measure mine effluent and two downstream stations (see Figure 2). The sample sites are described in Table 1 and illustrated in photographs, Figures 3 to 9.

## 2.1 Description

Mining at Barlow Creek during the study period involved mechanically stripping the overburden, stock piling the pay dirt and gradually feeding it through the mine's processing plant. The volume of dirt stripped for 1982 was 40,000 m³ with approximately 300 hours of sluicing between the months of July and September (DIAND, 1982). The processing plant on the site contained two trommell drums for classification of material, a stacker with a conveyor belt that carried away oversized waste rock, and a series of sluice boxes to gather gold. The concentrate from the process plant is further concentrated and cleaned in series of smaller riffles and specialized concentrator devices in a gold room. Other pieces of equipment on the site consist of a D8 cat and backhoe for digging and feeding the process plant. The water source for the mining operation was pumped directly from the Barlow Creek diversion through a 15 cm pipe at an approximate rate of 2300 L/min (500 IGPM). Barlow Creek was diverted to the northeast side

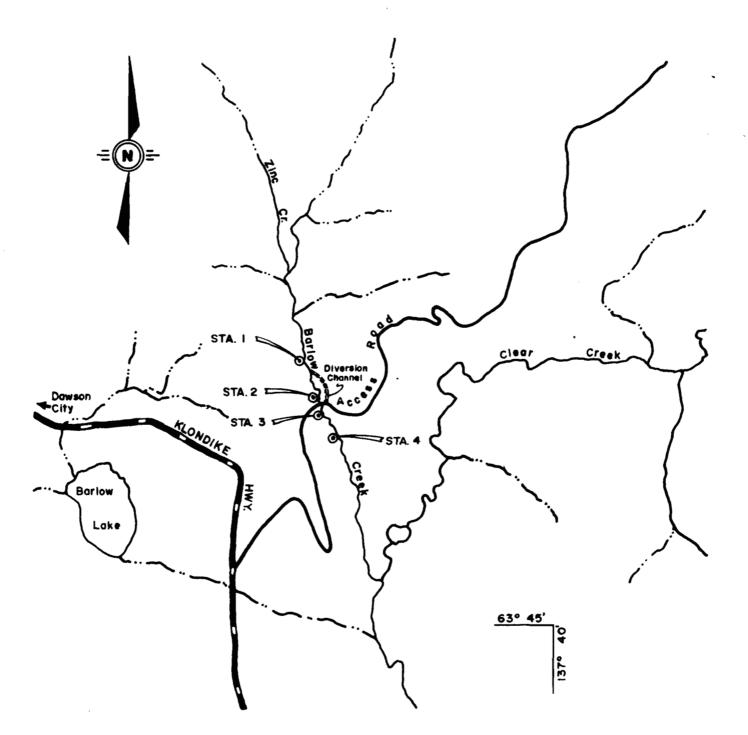


FIGURE 2 - BARLOW CREEK STUDY AREA SHOWING SAMPLE STATIONS

## LEGEND

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TABLE 1 DESCRIPTION OF SAMPLE SITES IN THE BARLOW CREEK STUDY AREA

	n con- kbrush, ssional m bed.	el from	ion corr grasses. sampling shallow	gh flow uicing.
REMARKS	Stable banks. Vegetation consists of black spruce, buckbrush, grasses and moss. Occassional gravel bar in the stream bed. Attempted electrofishing and found no fish. Of shade.	Constructed effluent channel from settling pond. O% shade.	banks. Vegetati of willows and bar near on. Water was too ectrofishing. 0%	Stable banks. Stream channel is narrow which results in high flow during rain periods and sluicing. Attempted electrofishing but found no fish. 30% shade.
	Stab sist gras grav Atte foun	Cons set1	Stable sists Gravel locatio	Stab narr duri Atte foun
STREAM BOTTOM	Large cobbles, coarse gravel and some sand. 90% gravel was discarded from sediment sample.	Mainly large boulders and cobbles.	Cobbles, small pebbles and fine silt.	Mainly small cobbles and gravel. Gravel cemented together and sediment. 90% gravel was discarded from sediment sample.
LOCATION	63°46'N 137°39'W on Barlow Creek 0.75 km upstream from the mining operation and 3 km upstream of its confluence with Clear Creek. Elevation 580 m (1900 ft.)	63°46'N 137°39'W. Mine tailings effluent near the operation, 5 m (33 ft.) above its confluence with Barlow Creek. Elevation 565 m (1850 ft).	63°46'N 137°39'W. 50 m (164 ft.) downstream of the confluence of the mine tailings effluent (Station 2) and Barlow Creek. Elevation 560 m (1837 ft.).	63°45'N 137°39' . 150m (492 ft.) downstream of station 3 on Barlow Creek. Elevation 555 m (1821 ft.).
STATION	-	0	n	4

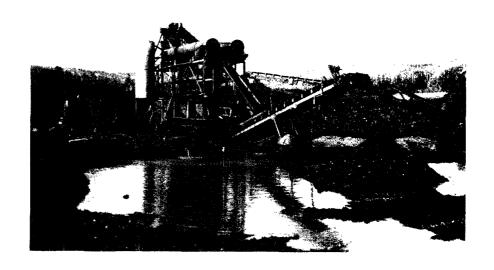


FIGURE 3 THE PROCESS PLANT AT BARLOW CREEK CONSISTS OF TWO TROMMELS,

A STACKER AND A SERIES OF SLUICES. NOTE THE PILE OF WASTE

ROCK BELOW THE STACKER.



FIGURE 4 BARLOW CREEK CONTROL STATION 1 LOCATED UPSTREAM OF THE MINING OPERATION.



FIGURE 5 STATION 2 IS LOCATED IN THE EFFLUENT CHANNEL 5 METRES ABOVE ITS CONFLUENCE WITH A DIVERTED SEGMENT OF BARLOW CREEK.

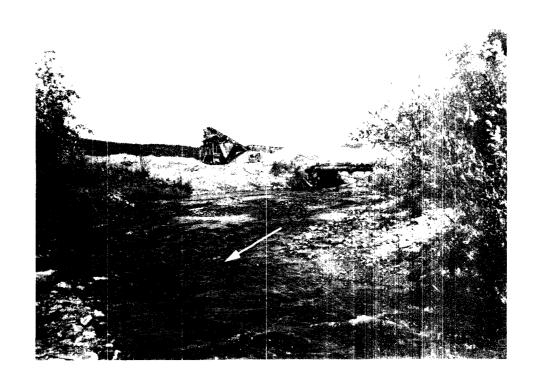


FIGURE 6 STATION 3 IS LOCATED 10 METRES DOWNSTREAM OF THE CLEAR CREEK ROAD BRIDGE.

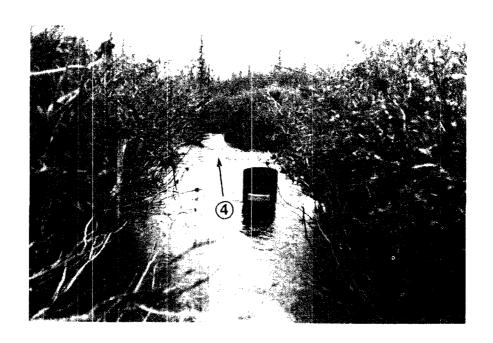


FIGURE 7 STATION 4 IS LOCATED 150 METRES DOWNSTREAM OF STATION 3.
HIGHER FLOWS ARE EVIDENT DURING ACTIVE SLUICING. CIRCULAR
BENTHIC SAMPLER IN THE FOREGROUND.

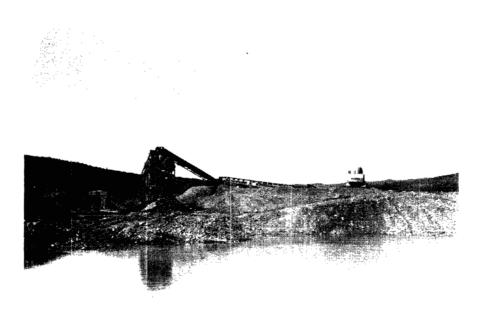


FIGURE 8 BARLOW CREEK PLACER MINING OPERATION. THE SETTLING POND IS LOCATED IN THE FOREGROUND.

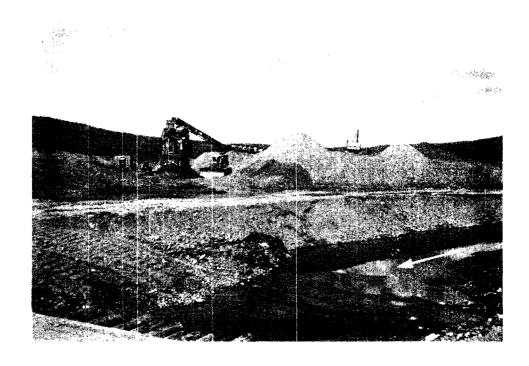


FIGURE 9 BARLOW CREEK MINING OPERATION, WITH SETTLING POND IN CENTRE
AND DIVERSION CHANNEL IN THE LOWER RIGHT HAND CORNER.

of the valley, re-entering the original channel below the settling pond. Water from the sluicing operation was discharged into settling ponds and tailings were pushed up to higher ground with future plans for restoration and revegetation.

The current camp and mining operation at Barlow Creek operated under a Water Use Authorization issued under the Northern Inland Waters Act. Water requirement for the camp was estimated at 2300 L/day (500 IGPD) and 5500 L/min (1200 IGPM) for the mining operation.

During the 1982 season, mining and testing activities were carried out at two small mining operations in the Barlow Creek drainage upstream of the study area. One operation with an estimated water requirement less than 2300 L/min. took place on the upper end of Barlow Creek on the Red Rooster tributary. This consisted of a relatively small testing program involving some stripping and trenching. Any waste water resulting from the testing program was discharged into a large hole rather than into the creek, thus preventing any disturbance. The second operation occurred 6 km upstream on Zinc Creek and basically involved some hand mining with an estimated water requirement of 700 L/min. for an approximate total of 15 hours of sluicing. Any significant influence arising downstream however, may be linked to mining activity and disturbance on Zinc Creek from the previous year (J. Maxwell, Pers. Comm.).

# 2.2 Mineralization Description

Yukon Group rocks of the Proterozoic age underlay the Barlow and Clear Creek watershed. This Group is primarily composed of quartzite grits, shales and bands of limestone. A major structural feature found in this area is the Tintina trench which is crossed by Clear Creek between Barlow Creek and the Stewart River (Queenstake, 1982). The terrain can be described as highly weathered with colluvial slopes and V-shaped, narrow, restricted valleys. Colluvial placer deposits are developed on terrain where active down-slope movement of superficial materials is occurring, generally over a weathered bedrock surface, and commonly associated with local primary lode sources.

Factors such as slope angle or gradient, thickness and type of slope material, size and specific gravity of heavy minerals, coefficient of friction and seasonal frost all affect extent and grade of colluvial placer deposits (Boyle, 1979). It is also important to note that gold associated with colluvial deposits tends to be coarser near the primary source.

#### 3 METHODS

Access to the area was by truck and each station was sampled six times. The sampling dates were June 26, July 8, July 19, August 9, September 1 and September 22, 1982.

# 3.1 Water Quality

Water samples were collected and preserved as described in Appendix I, Table 1 at each of Stations 1, 3 and 4. At Station 2, the tailings pond effluent stream, samples were collected for non-filterable residue (NFR) only.

Temperature, conductivity and pH were measured in the field using a Yellow Springs Instrument direct-reading salinity-conductivity-temperature meter and Model 296 Radiometer pH meter. Stream discharge (flow) was measured by using a Price-type current meter. Dissolved oxygen was analyzed using the Winkler's Azide modification method. Settleable solids were collected as grab samples and analyzed in the E.P.S. Whitehorse Lab within 2-4 days. Non-filterable residues were collected in duplicate. One sample was analyzed within 3 days in the Whitehorse Lab and the second sample was sent to the E.P.S. Vancouver Lab. All other analyses were performed by Laboratory Services, Environmental Protection Service, 4195 Marine Drive, West Vancouver, B.C. These included turbidity, colour, total hardness, filterable residue (FR), total alkalinity, total phosphates, nitrates, nitrites, ammonia, sulfate, chloride and the following extractable metals:

Aluminum (Al)	Copper (Cu)	Selenium (Se)
Antimony (Sb)	Iron (Fe)	Silicon (Si)
Arsenic (As)	Lead (Pb)	Silver (Ag)
Barium (Ba)	Magnesium (Mg)	Sodium (Na)
Beryllium (Be)	Manganese (Mn)	Strontium (Sr)
Cadmium (Cd)	Mercury (Hg)	Tin (Sn)
Calcium (Ca)	Molybdenum (Mo)	Titanium (Ti)
Chromium (Cr)	Nickel (Ni)	Vanadium (V)
Cobalt (Co)	Potassium (K)	Zinc (Zn)

The percent dissolved oxygen (% DO) saturation was calculated by first determining the dissolved oxygen saturation concentration (S') from the formula:

$$S' = S \frac{P}{760}$$
 (APHA et al 1975)

where S' = dissolved oxygen (DO) saturation concentration at the in situ temperature and atmospheric pressure

S = dissolved oxygen (DO) saturation concentration at sea level for in situ temperature

P = atmospheric pressure in mm of mercury (mm Hg) at site elevation

The percent dissolved oxygen saturation was obtained by using the ratio of field dissolved oxygen and S' in the following formula:

$$\frac{\text{Field DO}}{\text{S'}} \times 100 = \% \text{ DO Saturation}$$

where Field DO = Dissolved Oxygen concentration measured in the field

### 3.2 Sediments

Sediment samples were collected at the same time as water samples at Stations 1, 3 and 4. Sediment samples were collected on all dates except June 26 and at Station 4 on July 8 and 19. Sediments were not collected at Station 2 as this was strictly a settling pond effluent station in a constructed channel. Three replicate sediment samples were collected at each site using an aluminum shovel to scoop the samples into labelled Whirlpack bags. A description of sediment collection, preparation and analysis methods is given in Appendix I, Table 2. All sediment samples were shipped to Vancouver by air for analysis at Laboratory Services, Environmental Protection Service, West

Vancouver, British Columbia. Each of the three sediment samples collected at each station were analyzed for particle size and the following leachable metals:

Aluminum (Al)	Iron (Fe)	Silicon (Si)
Arsenic (As)	Lead (Pb)	Silver (Ag)
Barium (Ba)	Magnesium (Mg)	Sodium (Na)
Beryllium (Be)	Manganese (Mn)	Strontium (Sr)
Cadmium (Cd)	Mercury (Hg)	Tin (Sn)
Calcium (Ca)	Molybdenum (Mo)	Titanium (Ti)
Chromium (Cr)	Nickel (Ni)	Vanadium (V)
Cobalt (Co)	Phosphorus (P)	Zinc (Zn)
Copper (Cu)	Potassium (K)	

# 3.3 Bottom Fauna

Bottom fauna was sampled at three Barlow Creek sampling stations. Station 2 was located in the constructed effluent channel from the settling pond and was not suitable as a station for bottom fauna collection. Samples were collected at the same time as water and sediment samples on the six sampling dates. Three replicate samples were collected at each site using a 30 cm x 30 cm surber sampler (900 cm $^2$  or 1 ft $^2$ ) with a mesh size of 0.76 mm. Bottom fauna samples at Station 4 on July 8 and July 19, 1982, however, were not collected because high water conditions made it impossible to obtain a representative bottom sample. Bottom fauna collection, preservation and identification methods are given in Appendix 1, Table 3.

Diversity indices were calculated from the bottom fauna data collected, using the formula described by Pielou (1975) as follows:

Species Diversity (H') =  $-\Sigma$  (P<sub>i</sub> log<sub>10</sub> P<sub>i</sub>) i=1 where P<sub>i</sub> =  $n_i/N$   $n_i$  = total number of individuals in the ith genus in one sample N = total number of individuals identified to genus level in one sample g = total number of genera in one sample

The use of individuals identified to genus level instead of to species level results in slightly lower diversity indices (H') values (Hughes, 1978). The omission of bottom fauna that cannot be identified to genus or species will affect the validity of diversity on a comparative level, as the entire population failed to be represented.

## 3.4 Fish

Electrofishing of the creek at Stations 1, 3 and 4 was conducted on July 8, 1982. This was done to determine if fish were present and, if so, to obtain tissue samples for metal analysis. No fish were present. Notes were made on whether sample streams appeared suitable for fish habitat.

#### 4 RESULTS AND DISCUSSION

The results for the different parameters measured are generally seen to be greatly influenced by mining, particularly sluicing activity at the mine site. Sluicing was occurring during the sampling periods on July 19 and August 9, but had ceased for a few hours on September 22. There was no sluicing activity on any of the remaining sample dates.

## 4.1 Water Quality

Table 4 in Appendix I lists the accepted water quality criteria for drinking water and aquatic life with the appropriate reference sources. The water quality results are presented in their entirety in Appendix II whereas the following results and discussion with accompanying graphs concentrate on those parameters considered particularly important to water quality and affected by the placer mining activity.

# 4.1.1 Heavy Metals.

As illustrated in Figures 10 and 11, the following heavy metals, with the exception of mercury, are significantly affected downstream on Barlow Creek by placer mining activity:

Arsenic (As)

Copper (Cu)

Nickel (Ni)

Iron (Fe)

Mercury (Hg)

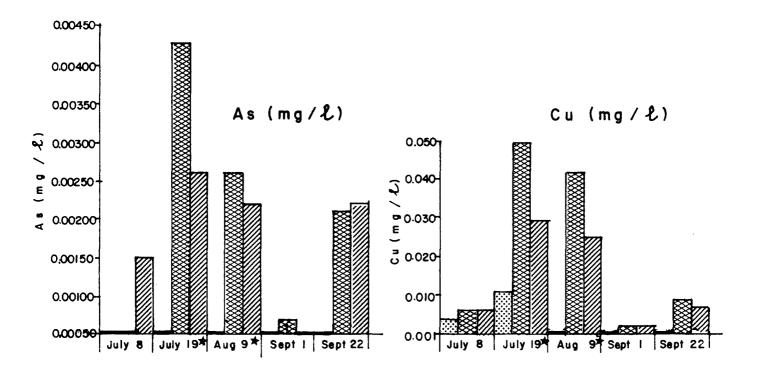
Manganese (Mn)

Nickel (Ni)

Lead (Pb)

Zinc (Zn)

Water analysis for extractable heavy metals reveals significant increases due to sluicing on July 19 and August 9. Downstream at Stations 3 and 4 arsenic levels ranged from <0.0005 mg/L to 0.0043 mg/L. Although values during sluicing periods increased 5 to 10 times from the upstream levels, they were still within the recommended limits of  $0.05 \, \text{mg/L/(total)}$  for drinking water and aquatic life.



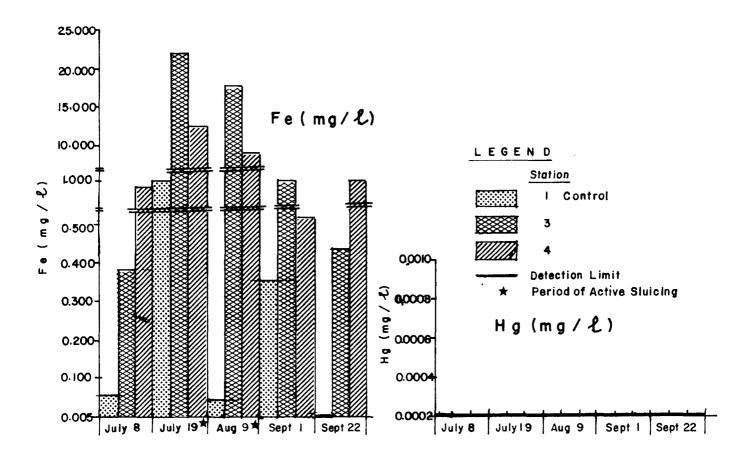


FIGURE IO HEAVY METALS IN WATER
(Origin of Y-axis equals Detection Limit) - 1982

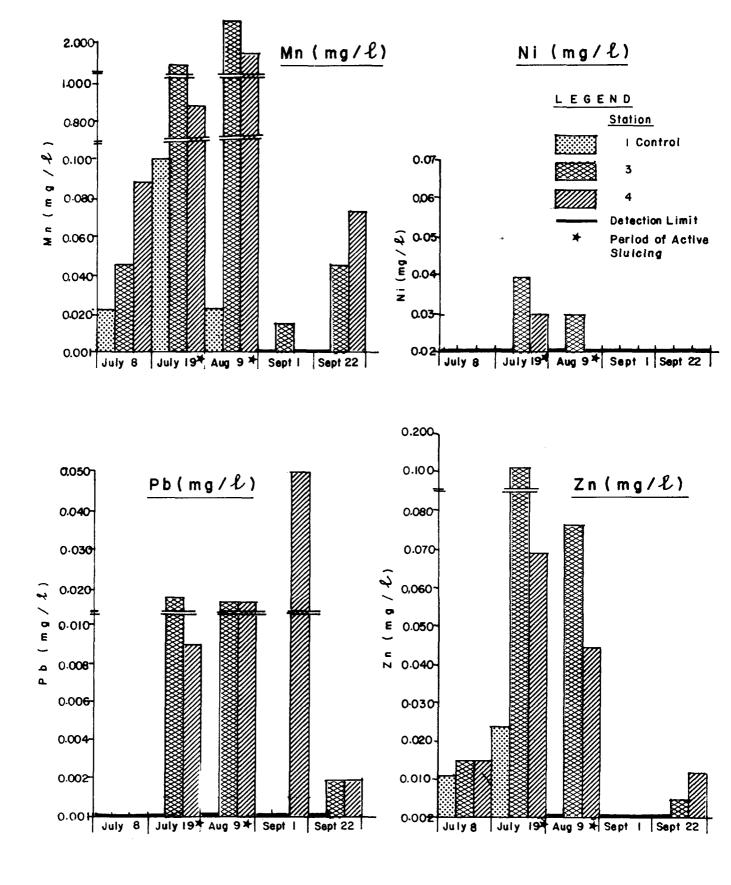


FIGURE II HEAVY METALS IN WATER

(Origin of Y-axis equals Detection Limit) - 1982

Copper levels increased as a result of mining and on both sluicing dates were above the recommended levels for aquatic life; 0.050 mg/L and 0.030 mg/L at Stations 3 and 4 on July 19, and 0.042 mg/L and 0.025 mg/L at Stations 3 and 4 on August 9. Copper also slightly exceeded recommended levels for aquatic life at downstream stations on July 8; 0.006 mg/L at Stations 3 and 4, and on September 22; 0.009 mg/L at Station 3 and 0.007 mg/L at Station 4. At Station 1 (control station), a level of 0.011 mg/L was found on July 19. This level may have been caused by upstream testing and sluicing activity.

Iron levels exceeded recommended limits for drinking water and aquatic life at all downstream stations throughout the sampling period and at the control (Station 1) on two occasions. The highest values found were 22.9 mg/L and 12.6 mg/L at Stations 3 and 4 on July 19 and 17.4 mg/L and 8.74 mg/L at Stations 3 and 4 on August 9. Elevated values for iron may be attributed to mining activity upstream on Barlow Creek or on the property itself.

Mercury levels were all below the detection limit of 0.0002 mg/L and therefore do not pose a problem.

Levels for manganese were found to be significantly increased at both downstream stations on July 19 and August 9. Values of 1.47 mg/L and 0.874 mg/L at Stations 3 and 4 exceeded the recommended limits for drinking water and aquatic life, as did values of 2.60 mg/L and 1.80 mg/L found at Stations 3 and 4 on August 9. Elevated levels of manganese that exceeded limits for aquatic life and drinking water were also noted at Station 1 on July 19.

Nickel levels only slightly exceeded recommended limits for aquatic life at Stations 3 and 4 on July 19 with values of 0.04 mg/L and 0.03 mg/L, and at Station 3 on August 9 with a value of 0.03 mg/L. All other values for nickel were below the 0.02 mg/L detection limit.

All levels for lead were within drinking water limits at all locations; however, some exceeded recommended limits for aquatic life. Values of  $0.017 \, \text{mg/L}$  and  $0.009 \, \text{mg/L}$  were found at Stations 3 and 4 on July 19 and  $0.016 \, \text{mg/L}$  for Stations 3 and 4 on August 9. These levels were noted to be approximately 20 times greater than at upstream

stations. An anomalous level of 0.050 mg/L also occurred at Station 4 on September 1. Possible error or contamination to the sample collection might explain this elevated level.

Zinc was also present at levels which exceeded recommended limits for both drinking water and aquatic life. These values were 0.111 mg/L and 0.069 mg/l at Stations 3 and 4 on July 19 and 0.076 mg/L and 0.044 mg/L at Stations 3 and 4 on August 9. Other heavy metals that appeared to increase as a result of mining activity included Aluminum (A1), Cobalt (C), Chromium (Cr) and Magnesium (Mg). These values may be found in Appendix II.

Heavy metals present in the water were clearly more abundant on dates when sluicing was occurring at the time of sampling as shown in Figures 10 and 11. These elevated levels corresponded with higher suspended sediment (NFR) values for the same dates which is expected because the metals are typically associated with fine grained material that makes up suspended sediments (Paski, Pers. Comm., 1982). It is seen in the above results and from Figures 10 and 11 that levels for copper, iron and manganese at Station 1 on July 19 in particular were higher than on other sample dates and exceeded the recommended levels for aquatic life (Cu, Fe, Mn) and drinking water (Fe, Mn). It is suggested that this increase in extractable metals at Station 1 on July 19 is due to placer testing activities on upstream tributaries to Barlow Creek or the prolonged rainfall which would increase natural erosion into the stream.

# 4.1.2 <u>Suspended Sediments (Non-Filterable Residue - NFR).</u>

All suspended sediments were collected by grab sample and analyzed at the EPS Whitehorse laboratory two to four days after collection. Duplicate samples were also analyzed at the EPS laboratory in Vancouver as a cross reference for quality control, and corresponded very closely with the results of the EPS Whitehorse laboratory. The results presented in Appendix II are those determined in the Whitehorse laboratory.

Suspended sediments can be defined as any particulate matter suspended in water and that is retained by a 0.45 micron filter or a

filter of defined pore size. The term is also synonymous with non-filterable residue (NFR). Suspended sediments are comprised of sand, silt, clay and detritus, and are usually the major cause of concern in water with regard to a placer mining operation. Griffiths and Walton (1978) summarize the direct and indirect effects of sediment on fish. The direct effects are: (1) mortality of adult fish as a result of mechanical and abrasive damage; (2) impairment of reproduction, growth and survival; and (3) increase of disease within a population. The indirect effects result from: (1) habitat modifications, (2) alterations in food sources and other biotic relationships, and (3) reduction of visibility.

Good discussions of the effects and potential effects of suspended sediments on fish, the aquatic habitat and bottom invertebrates have been presented by Pickral (1981), Griffiths and Walton (1978), Cordone and Kelley (1961), and Rosenberg and Snow (1975).

Water quality with regard to suspended sediments in the upper reaches of Barlow Creek appears relatively undisturbed and indicates low levels of less than 5 mg/L of suspended sediments on most sample On July 19, however, Station 1 exhibited a dates (see Figure 12). suspended sediment level of 47 mg/L. This could have been attributed to some testing work from upstream operations on Zinc Creek or the Little Rooster tributary of Barlow Creek or as a result of rainfall which increased natural erosion into the stream. Very high values for suspended sediments were found at downstream stations, Stations 3 and 4, on the two sluicing dates, July 19 and August 9. sediments on July 19 measured as high as 6990 mg/L, 1008 mg/L and 570 mg/L at Station 2 (tailings pond effluent), Station 3 and Station 4 respectively; and 4530 mg/L, 984 mg/L and 2065 mg/L at Stations 2, 3 and 4 respectively on August 9. During sampling times on September 1, active sluicing was not occurring and the suspended sediment levels at downstream stations all exhibited greatly reduced values as compared to sluicing times. The suspended sediment levels on September 22 at Stations 3 and 4 were 211 mg/L and 142 mg/L respectively. activity had ceased for several hours on September 22 when suspended sediments were collected at Stations 3 and 4 and found to be slightly

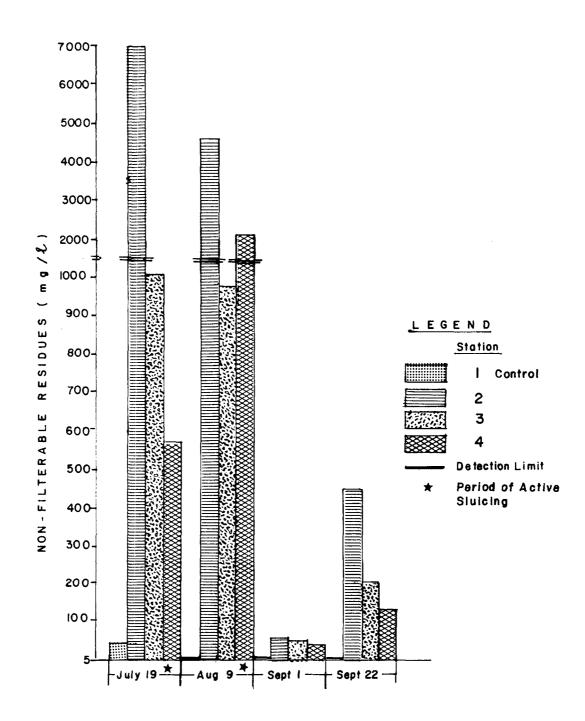


FIGURE 12 NON-FILTERABLE RESIDUE (Origin of Y-axis equals Detection Limit)

elevated (211 mg/L and 142 mg/L respectively). Effluent had time to settle in the tailings pond, which explains the observed results.

Suspended sediment levels influence or may influence other water quality parameters such as turbidity, colour, settleable solids and dissolved oxygen. These parameters and others are described in the following section.

## 4.1.3 Other Water Quality Parameters.

The results for other water quality parameters that were noticeably different between upstream and downstream stations and probably influenced by placer mining are provided below. The values for these parameters, in many cases, exceeded drinking water standards or recommended levels for aquatic life. The parameters include dissolved oxygen, settleable solids, turbidity, colour, total hardness, phosphorus, nitrite ( $NO_2-N$ ) and ammonia ( $NH_3-N$ ). Results for these and other parameters are provided in Appendix II.

- (a) Flow. Flow measurements provide only an approximation of flow in that there were variations in recorded flow rates between stations on the creek. The flow data in Appendix II indicates input to the stream between upstream and downstream stations based on average flows. The average flow rate (m3/s) at Station 3 is greater than at Station 4 suggesting that some of the flow is diverted from the stream channel to an underground aquifer or channel between Station 3 and 4 or that there are significant errors in the flow measurements. The range of flows at Station 1 varied from 0.1 m3/s to 0.7 m3/s with the peak of 0.7 m3/s occurring on July 19 which followed a period of prolonged rainfall. This period of highwater would probably contribute to the elevated levels of suspended sediments (NFR) and certain metals noted at Station 1 on July 19.
- (b) <u>Temperature</u>. The results show a general increase between Station 1 and Station 4. The general downstream increase from  $7^{\circ}\text{C}$  at Station 1 to  $9^{\circ}\text{C}$  at Stations 3 and 4 on July 19, and  $9^{\circ}\text{C}$  at

Station 1 to 11°C at Stations 3 and 4 on August 9 may be caused by the fact that the operation had a storage pond to provide adequate water for prolonged sluicing and because during sluicing there was a high percentage of sluicing water recirculated. The greatest temperature difference between upstream and downstream stations occurred in mid-season when the warming of the ponded water would be greatest, whereas at the beginning and end of the season the upstream and downstream temperature differences were less.

(c) Dissolved Oxygen (DO). Slight decreases in dissolved oxygen were noted at all downstream stations as compared to the Station 1 control. The DO levels were between 0.1 mg/L to 1.1 mg/L lower at downstream stations than respective upstream stations, which generally did not pose a significant decrease. However small, this reduction may have been caused by surface and organic material introduced into the water from sluicing and runoff which may create an oxygen demand. The DO levels remained well above minimum levels recommended for aquatic life. The percent saturation for DO, which is directly related to the actual concentration, is used as the reference standard for this parameter. Dissolved oxygen saturation did not meet the drinking water standard at any of the downstream stations except on July 8, which corresponds with the fact that mining activity did not influence those stations until after that date. Except for July 8, the percent DO saturation was less than 100% at the control station throughout the entire operating season. This could be due to placer activity on upstream tributaries to Barlow Creek or groundwater influencing the stream during the latter parts of the Under the conditions present in this stream reach, the decrease in percent DO saturation is not great although it was detectable on sluicing dates.

- (d) Turbidity. Increased turbidity generally exists in water when an increased surface area of silt and clay particles reflect more light than would larger sand particles (Taekema 1983). turbidity is directly related, though not necessarily linear, with suspended sediments it was found, as expected, to be greatly elevated at downstream stations on July 19 and August 9. Values as high as 800 FTU (Formazin Turbidity Units) and 600 FTU occurred at Stations 3 and 4 respectively on July 19, and 1280 FTU and 1200 FTU at Stations 3 and 4 on August 9. Significantly elevated turbidity levels of 180 FTU and 95.0 FTU were also demonstrated at Stations 3 and 4 during the last sampling date of the season, September 22, which occurred immediately after sluicing had ceased for the day. These values compare with levels less than 0.1 FTU to 2.9 FTU at the Station 1 control. Other downstream values during non-sluicing times range between 10.0 and 30.0 FTU, indicating turbidity resulting from non-point sources along the stream in the vicinity of the operation.
- (e) <u>Settleable Solids</u>. Settleable solids are commonly defined as "any particulate matter in suspension that will settle out of suspension in a standard unit of time" (Department of Environment, 1979). Settleable solids were elevated on one sampling date, July 19. Values ranged from 0.9 mL/L at Station 3 and 0.2 mL/L at Station 4. Values at Stations 3 and 4 on August 9 were both less than 0.1 mL/L. All other values for settleable solids at the control and downstream at Stations 3 and 4 were less than 0.1 mL/L.
- (f) Colour. Colour exceeded recommended levels for drinking water by demonstrating values greater than 100 colour units at Stations 3 and 4 on July 19. On this date, however, a value of 65 colour units was found at the control (Station 1). It is possible that this elevated level was caused from testing and mining activity upstream although naturally high flows existed at the time. A colour level of 20 colour units occurred at Station 3 on September

- 1. Colour measurements at Stations 3 and 4 were not analyzed for August 9 due to high turbidity interferences. High values for colour may be due to increased concentrations of organic matter introduced to water from the stripping process.
- (g) <u>Hardness</u>. Relatively high levels of total hardness were noted throughout the sampling period. Values ranged from 66.4 124 mg/L as CaCO<sub>3</sub> at Station 1 to 94.0 214 mg/L as CaCO<sub>3</sub> at downstream stations. Levels of 195 and 130 mg/L as CaCO<sub>3</sub> occurred at Stations 3 and 4 on July 19 and 214 and 150 mg/L as CaCO<sub>3</sub> on September 1. These results clearly exceeded the recommended levels of 80-100 mg/L for drinking water.
- (h) Phosphorus (as PO4). Excessive amounts of phosphates in streams may cause algal blooms and odours that are detrimental to fish. To avoid nuisance concentrations of algae, average total phosphorus concentrations should not exceed 0.020 mg/L (Ontario Ministry of Environment, 1978). This level however was exceeded at Stations 3 and 4 on three sampling dates. These were 0.920 mg/L and 0.510 mg/L on July 19, 1.05 mg/L and 1.00 mg/L on August 9 and 0.225 mg/L and 0.145 mg/L on September 22. One other slightly elevated level that may have been significant was 0.057 mg/L found at Station 1 on July 19. Other values at the control station ranged between <0.005 0.008 mg/L and between 0.020 0.033 mg/L at Stations 3 and 4 during nonsluicing times.
- (i) <u>Nitrites</u>. Nitrite measurements exceeded the recommended drinking water standard of 0.001 mg/L at most downstream stations and at the control station on July 19. These levels ranged from 0.007 mg/L to 0.037 mg/L. The detection limit for nitrite (0.005 mg/L) is greater than the recommended drinking water limit. All Station 1 samples except on July 19, Station 4 on September 1 and Stations 3 and 4 on September 22 were all less than the detection limit.

- (j) Ammonia. Levels for ammonia were all found to be within recommended limits for drinking water and aquatic life. Calculations taken from APHA et al (1975) were used on ammonia levels to determine the percentage of un-ionized ammonia. Levels of un-ionized ammonia that exceed the 0.02 mg/L recommended level for aquatic life indicates excess concentration of metabolic products in the water which may result in stress to aquatic organisms.
- (k) Conductivity, pH, filterable residues, sulphates, nitrates and alkalinity were all within recommended levels for drinking water and aquatic life.

### 4.2 Sediments

Sediment samples were collected at Stations 1, 3 and 4 on each of the sampling dates and subsequently analyzed for particle size composition and leachable metals. No sediment samples were collected at Station 2, or at Station 4 on July 8 and 19. When analyzing and discussing the results of sediment analysis it is necessary to be aware of the variability that can occur among replicate samples due to natural variations in stream bottom composition. Despite the limitation on interpretation the data shows the effects of sluicing on sediment composition and sediment chemistry at downstream stations compared to the control, and this can be related to the effectiveness of the settling ponds.

# 4.2.1 Sediment Particle Size Analysis.

Results of the sediment particle size analysis are given in Appendix III, Table 1. Illustrations of particle size in terms of mean percent composition can be found in Figures 13 and 14.

Increased levels of silts and clays occurred downstream of the mining activity throughout the operating season. Especially high concentrations of silts and fine sands were found at Station 3 on two sampling dates when sluicing occurred (July 19 and August 9). During

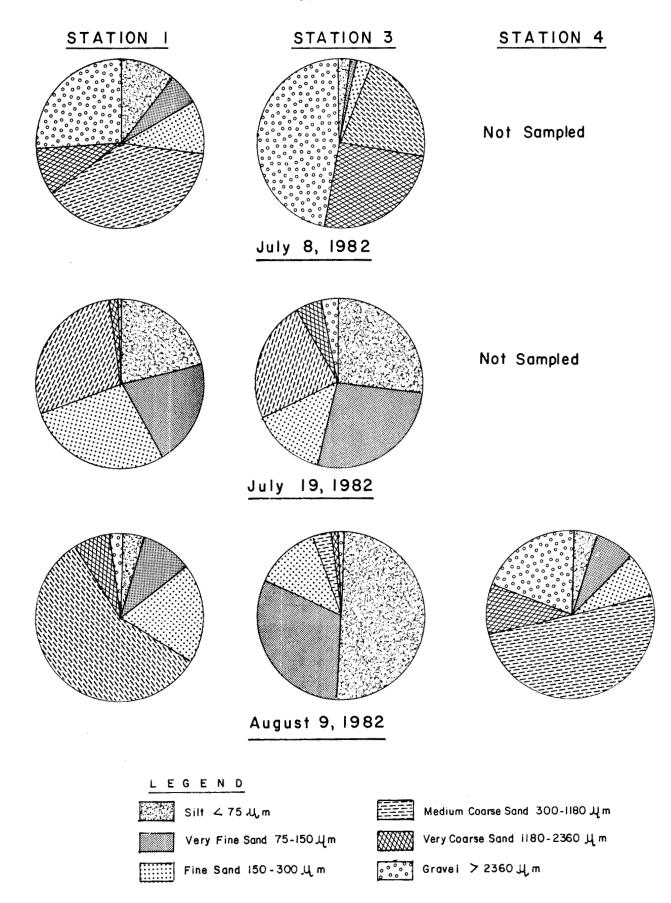
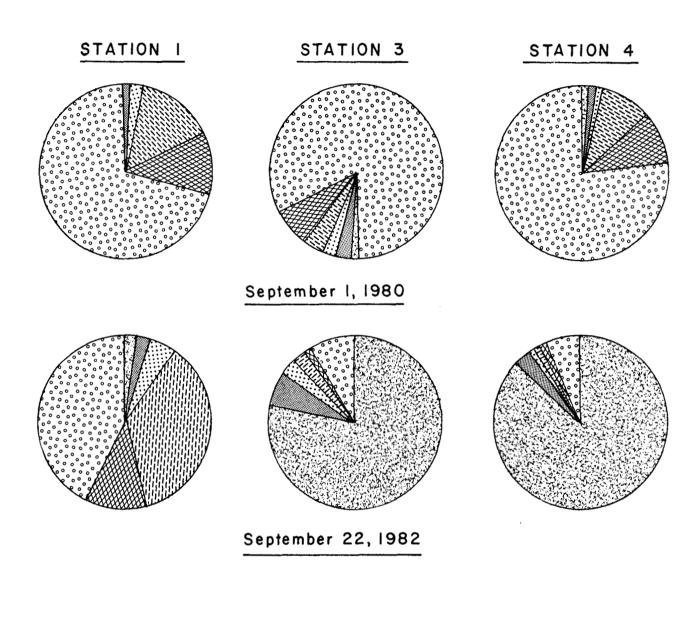


FIGURE 13 SEDIMENT PARTICLE SIZE ANALYSIS - MEAN PERCENT COMPOSITION



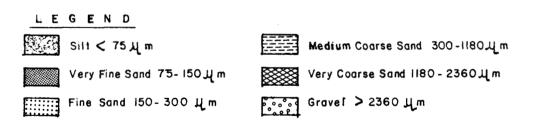


FIGURE 14 SEDIMENT PARTICLE SIZE ANALYSIS - MEAN PERCENT COMPOSITION

these two periods percent composition of silts and fine sands (75 um to 350 um) ranged from 67.4% to 97% downstream at Station 3, but only 19.4% at Station 4 on August 9. Eighty percent of the sediment collected at Station 4 on August 9 ranged in size from medium coarse sand to gravel (300 um to >2360 um). Due to the deep and narrow channel at Station 4 and an increase in discharge during sluicing, a "flushing out" process may have occurred to explain this state of recovery to the stream bed. Sediment at Stations 3 and 4 on September 1 consisted mostly of gravel which ranged in size >2360 um, 96.1% and 97.6% respectively. Sluicing had ceased for a period during this time and provided enough time for fine particles to be resuspended and transported further downstream allowing the stream bed in the study area to return to a more natural composition for the area.

Interesting sediment samples were collected at Stations 3 and 4 on September 22 where 89.6% to 93.5% of the sediment gathered was considered fine and ranged in size from 75 - 300 um. Sluicing was estimated at 300 hours between June and September and a thrust in sluicing activity at the end of the season may explain the large increase of fine sediments found downstream on this date. Also, sluicing had ceased just prior to sampling on this date so it is unlikely that there had been sufficient time for the stream to resuspend and move sediments further downstream. The flow rate was also declining later in the season and the stream was less able to resuspend the sediments.

## 4.2.2 <u>Sediment Metal Concentrations</u>.

The concentration of metals analyzed in the portion of the sediment that was smaller than 150 um are given in Appendix III, Table 2 for all stations sampled. Sediments at Station 4 on July 8 and 19 were not collected due to high water and the inability to obtain a representative sample. A list of mean values for sediment metal concentrations is found in Table 2.

Generally there appears to be some variability in the heavy metal results from the sediment sampled throughout the operating season. A mean value of 5.5 mg/kg was found at Station 3 on July 8 and

TABLE 2 BARLOW CREEK MEAN VALUES FOR SEDIMENT METAL CONCENTRATIONS
(All mean concentrations given in mg/kg dry weight unless otherwise noted)

982	4		\$	14267	16.6	-	134	0.3	3563	<0.3	10.4	24.3	29.6	25967	0.12	1563	4817	587	12,8	163	20	655	13	3637	<2	28.7	844	22	82.4
September 22, 1982	۳۱		\$	15200	16.2		132	0.3	3430	<0.3	11.9	25.3	34.2	28633	0.11	1670	5260	582	13.5	173	22	693	16	3830	\$	28.2	894	82	91.5
Septe	_		\$	12200	7.7	-	113	0.3	4507	<0,3	11.6	28.1	23.1	24200	0.10	1090	4137	753	10.7	133	20	756	٥	3543	<2	34.3	1307	35	57.1
286	4		\$	12967	8.4	9.7	131	<0.2	4123	<0.3	13.2	28.4	24.0	25633	60.0	1740	4397	688	5.1	173	24	670	=	3083	\$	30.2	1108	82	62,3
September 1, 1982	'n		\$	12600	8.5	11.1	128	<0.2	4040	<0.3	11.3	24.6	24.3	25700	0.03	1517	4333	650	5,3	147	23	678	01	3763	8	29.1	1048	37	62.4
Sep	-		ŝ	11400	7.6	6.4	6.76	<0.2	4147	<0.3	6.6	22,3	23.8	25667	0.14	1267	4200	602	4.7	137	21	745	80	3847	\$	29.3	1213	35	59.7
~!	4		\$	9757	5.8	8.2	106	<0.2	3440	<0.3	12.6	21.3	18.4	18733	0.15	755	3520	447	2.3	120	19	629	8	3227	<b>~</b>	23.7	989	28	49.0
Augus† 09, 1982	٣		\$	12267	0.9	10.0	151	<0.2	4080	<0.3	12.4	24.3	22.4	21267	0.15	876	4357	634	2.4	163	21	649	٥	3687	<b>4</b>	31.2	672	33	60.3
BnV	~		\$	11133	4.9	11.5	108	<0.2	3807	<0.3	9.01	25.1	22.7	21400	0.11	666	3977	618	3.0	137	50	629	80	3827	<2	28.2	883	32	55.2
~!	4																												
July 19, 1982	£		\$	7027	4.3	5.5	78.9	<0.2	2577	<0.3	9.9	16.0	16.1	14000	0.15	363	2943	373	9.1	93	41	652	ĸ	2300	<b>4</b> 2	15.7	239	81	43,3
<u>-</u>	-		\$	7213	5,5	3.1	76.4	<0.2	2730	<0.3	6.5	17.1	17.5	14733	0.15	385	3060	394	2.2	8	15	704	7	2400	\$	16.0	275	61	44.4
2	4	1																								22.0			
July 8, 1982	٣																									22.5			
	MS: 1		\$	7880	5.2	6.1	91.5	<0.2	2707	<0.3	7.9	9.61	20.4	15933	0.10	359	3347	514	40°8	8	8	809	^	2900	<2	18.5	215	61	49.8
DATE:	STATIONS:		٧6	<del>-</del>	٧s	8	83	8	<b>8</b>	8	ပိ	ن د	3	F.	θΉ	¥	Ð.	ž	9	10 2	ž	۵.	9	S	S	s	=	>	Zu

between 16.2 mg/kg and 16.6 mg/kg at Stations 3 and 4 on September 22; all of which were considered to be within normal limits. Manganese and zinc decreased slightly during the two sluicing periods as compared to the other sampling dates. Hean values for manganese ranged from 373 mg/kg to 976 mg/kg; a normal level for manganese is in the area of 1000 mg/kg (Paski, 1982). Mean values for zinc were also found to decrease during sluicing with a value of 43.3 mg/kg at Station 3 on July 19 and 60.3 mg/kg and 49.0 mg/kg at Stations 3 and 4 on August 9. After this period zinc levels gradually increased to mean values of 62.4 mg/kg and 62.3 mg/kg at Stations 3 and 4 on September 1 and 91.5 mg/kg and 82.4 mg/kg at Stations 3 and 4 on September 22. With other heavy metals such as copper, iron, mercury, nickel and lead, mean values increased over the operating season (see Figures 15 and 16).

A mean value of 16.1 mg/kg for copper was found at Station 3 on July 19, and 22.4 mg/kg and 18.4 mg/kg at Stations 3 and 4 on August By the end of the season values had increased slightly to 34.2 mg/kg and 29.6 mg/kg at Stations 3 and 4 on September 22. Mean values for copper found at the control sites varied little from the downstream stations. Mean values for iron were found to generally increase over the summer. A mean value of 14000 mg/kg was found at Station 3 on July 19 and 21267 mq/kq and 18733 mq/kq at Stations 3 and 4 on August 9. Values for mercury were most elevated during sluicing and declined in sediments toward the end of the summer. A level of 0.15 mg/kg was found at Station 3 on July 19 and 0.15 mg/kg each at Stations 3 and 4 on August 9. Mercury levels ranged between 0.03 mg/kg and 0.15 mg/kg. These concentrations are similar to other placer mining streams in Nickel and lead both rose Yukon Territory (Mathers et al 1981). gradually over the duration of the operating season although they remained within normal levels.

It was difficult to draw any firm conclusions due to the sample variability found for the heavy metals at Barlow Creek. Some metals decreased during sluicing periods and recovered by the end of the summer, while others gradually increased over the summer and suddenly escalated at the end of the season. The latter event was likely the result of a build up of sediment due to a concentrated

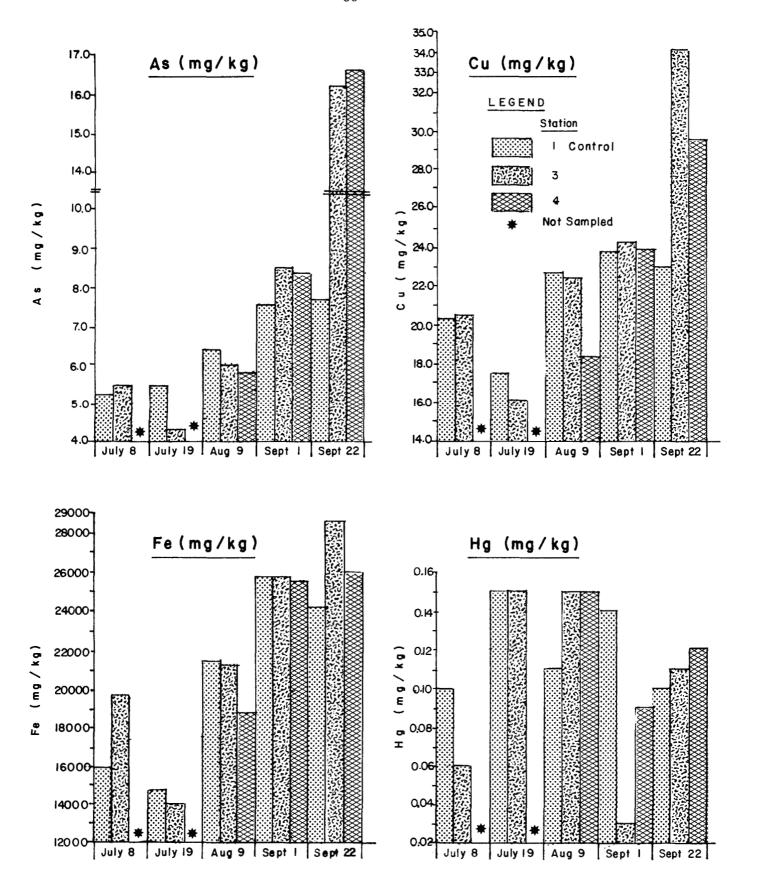
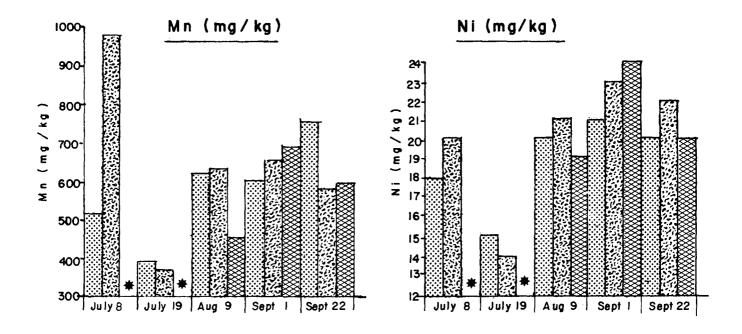


FIGURE 15 MEAN HEAVY METALS IN SEDIMENT



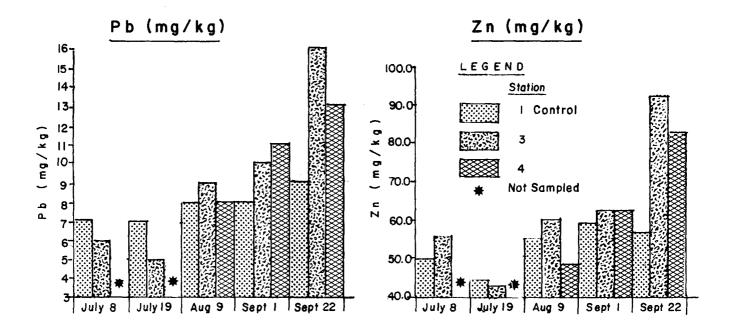


FIGURE 16 MEAN HEAVY METALS IN SEDIMENT

sluicing period which took place near the end of the season. Sluicing was estimated by the operator as 300 hours between June and September.

Although most of the heavy metals increased, all were considered to be within normal values for stream sediment (Paski, 1982). The build up of fine sediments, thus the increase in heavy metal concentrations, might suggest that the retention time of the settling pond was less effective at settling the heavier metal bearing sediments at the end of the operating season than at the beginning.

#### 4.3 Bottom Fauna

## 4.3.1 Bottom Fauna Numbers and Diversity Indices.

A summary of Barlow Creek bottom fauna numbers and diversity indices is given in Table 3. Stations 1, 3 and 4 were the only stations where bottom fauna were collected. Station 2 (settling pond effluent) was unsuitable for collection. Appendix IV, Table 1 lists the bottom fauna taxonomic groups as in Pennak (1978), found in Barlow Creek. Appendix IV, Table 2 lists the numbers of individuals in each taxonomic group and the diversity index for each sample.

Diversity index values were calculated (Table 3) to assist further assessment of the impact that a placer mining operation could have on Barlow Creek. The diversity index is used to express the "richness" of a community, represented by the number of different taxonomic groups (genera in this case). Communities of high diversity are characterized by large numbers of species with no single species overwhelmingly abundant. Communities of low diversity contain few species, some of which are represented in disproportionately high numbers. Generally diversity values greater than 0.90 (in  $log_{10}$  or 3.0 in log<sub>2</sub>) are found in unpolluted, productive waters while heavily polluted waters have values less than 0.30 (in log10 or 1.0 in log<sub>2</sub>)" (Wihlm and Dorris, 1968). The diversity indices in this study fall within a comparable range to those found in other Yukon streams subjected to placer mining such as Clear Creek (Burns, 1980). fauna populations (number per m<sup>2</sup>) generally have been found to be quite low, possibly due to cold climate of the study area and

SUMMARY OF BARLOW CREEK BOTTOM FAUNA AND DIVERSITY INDICES

TABLE 3

CALCULATED NUMBER PER M <sup>2</sup>	43	* * *	* * *	11 75 22	258 247 86	344 54 323
NUMBER PER FT.2	4 17 17	* * *	* * *	- 7 2	24 23 8	32 5 30
DIVERSITY (H1)	0.48	* * *	* * *	00.0	0.66	0.42
STATION	4 - 1 - 2 - 4 - 5 - 5 - 5	4 4 4 1 1 1 1 2 W	4 4 4 - 1 - 1 5 2 5	4 4 1 1 2 1 4 1 1 2 1 1 1 1 1 1 1 1 1 1	4 4 4 1 1 1 2 2 2	4 - 1 - 2 - 4 - 3
CALCULATED NUMBER PER M <sup>2</sup>	65	65 65 86	22 624 194	291 32 280	291 312 129	183 237 258
NUMBER PER FT.2 (0.093m <sup>2</sup> )	φ & ι	8 9 9	2 58 18	27 3	27 29 12	17 22 24
DIVERSITY (H <sup>1</sup> )	0.20	0.60	00.00	0.78 0.28 0.56	0.84	0.70
STATION	3 - 1	3 - 2 - 3 - 3	1 1 1 1 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	3 3 3 1 2 2 2 3 3 4 3 4 4 4 4 4 4 4 4 4 4 4 4 4	3 3 1 - 1 - 2 2 - 1 - 2 3 - 1 - 3 3 - 1 - 3 3 - 3 3 - 3 3 - 3 3 - 3 3 - 3 3 3 - 3	3 - 2 3 - 3
CALCULATED NUMBER PER M <sup>2</sup>	785 861	3228 4390 2077	97 54 215	258 452 667	441 291 377	280 452 742
NUMBER PER FT. <sup>2</sup> (0.093m <sup>2</sup> )	73	300 408 . 193	50 20	24 42 62	41 27 35	26 42 69
DIVERSITY (H <sup>1</sup> )	0.57	0.67	0.73 0.41 0.52	0.28	0.70	0.74
STATION	1 - 1 - 1 - 3	1 - 1 - 3 2 - 1	1 - 1 - 3 - 1 - 3	1 - 1 - 3 - 3	1 1 1 2 2 1	1 - 2 3 3
DATE 1982	June 26	Ju 1.7	Jul. 01	Aug.	Sep#	Sep † 22

\*- Bottom fauna not collected due to highwater conditions.

disturbance from an operation upstream. The bottom fauna standing crop appears low as a result, thus affecting the potential as a food source for fish in this type of habitat.

Diversity (H') values taken over six sampling periods for Station 1 (control) ranged from 0.28 to 0.89 and actual numbers of individuals collected by Surber sampler ranged from 5 to 408 (per 0.093 m² or 1.0 ft²). On July 19 diversity ranged between 0.41 and 0.73 at Station 1, and between 0 and 0.08 at Station 3. Benthic samples were not collected at Station 4 on July 19 due to high water conditions precluding obtaining benthic samples. Total numbers of individuals ranged from 5 to 20 at Station 1 and from 2 to 58 at Station 3. On August 9 diversity ranged between 0.28 and 0.56 at Station 1, 0.28 to 0.78 at Station 3 and zero at Station 4. Total numbers collected at this date ranged from 24 to 62 at Station 1, 3 to 27 at Station 3 and 1 to 7 at Station 4.

Significant variation in terms of diversities and total number of individuals were found at Station 1 on July 19 and Stations 1, 3 and 4 on August 9. Low numbers of individuals at Station 1 may have been the result of some mining disturbance six kilometers upstream on Zinc Creek. At Station 4 on August 9 benthic invertebrates were greatly reduced. These lower levels are considered to be due to sampling variability rather than impact from mining activity since there was not a corresponding reduction at Station 3. The three most abundant orders of bottom fauna found in terms of numbers of individuals were Ephemeroptera, Diptera and Plecoptera, which are further discussed in the following section.

## 4.3.2 Three Most Abundant Orders Found in Barlow Creek.

A list of the three most abundant orders and the component taxa found in Barlow Creek is given in Table 4. The numbers found and percent abundance based on the totals for these three orders is shown in Table 5. These values represent the numbers obtained in three replicate samples (0.279  $\rm m^2$  or 3  $\rm ft^2$ ) at each station. Graphs illustrating actual abundance and percent abundance for the three orders can be found in Figures 17, 18 and 19.

TABLE 4 THREE MOST ABUNDANT ORDERS AND THEIR TAXONOMIC GROUPS FOUND IN BARLOW CREEK

Ephemeroptera: Ameletus sp.

Baetis sp. Cinygmula sp. Epeorus sp.

Diptera:

Chironomidae adult
Chironomidae pupae
Brillia sp.
Cardiocladius sp.
Cricotopus sp.
Diplocladius sp.
Epoicocladius sp.
Eukiefferiella sp.
Heterotrissocladius sp.

Orthocladius sp.

Diamesa sp.

Monodiamesa sp.

Pseudodiamesa sp.

Culicidae

Chelifera sp.

Prosimulium sp.

Metacnephia sp.

Erioptera sp.

Tipula sp.

Plecoptera:

Subimago
Capnia sp.
Alloperla sp.
Utaperla sp.
Podmosta sp.

Arcynopteryx sp.

Cultus sp.

Megarcys sp.

Skwala sp.

Unid. dam.

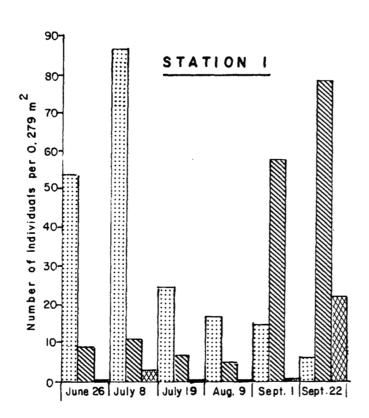
BARLOW CREEK BOTTOM FAUNA DATA - NUMBERS AND PERCENT CALCULATED FOR THE THREE MOST ABUNDANT ORDERS

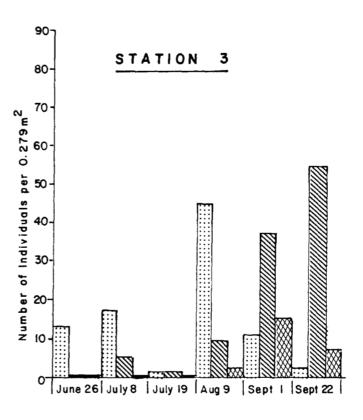
TABLE 5

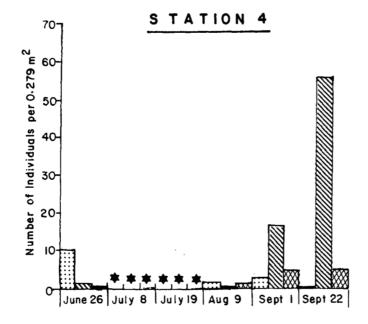
<del></del>	+	<b>+</b>		
September 22, 1982	Per- centage	6 21 21	3 85 12	0 8 8
Septembe	Total	6 79 23	52 7 7	55 5
September 1, 1982	Per- centage	20 79 1	17 59 24	12 72 16
September	Total Number	15 59 1	11 37 15	18 4 4 25
Augus† 9, 1982	Per- centage	80 20 0	78 18 4	50
August	Total Number	16 4 0 20	44 10 2 2 56	2 10
July 19, 1982	Per- centage	78 22 0	50	* * * *
July 19	Tota! Number	25 0 25 32	2 0 1	* * * *
July 8, 1982	Per- centage	3 11	18 0	* * * *
ا با سر	Tota! Number	87 11 3 101	17 4 4 0	* * * *
June 26, 1982	Per- centage	88 12 0	001	<u>10</u> 0
June 20	Tota! Number	53	0 0 51	0 0 1
		E D P Tot	To P	E P P Tot
SAMPLE	STATION	-	٣	4

E - Ephemeroptera D - Diptera P - Plecoptera

<sup>\* -</sup> Bottom fauna not sampled due to high water conditions.







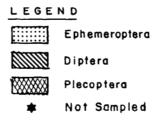


FIGURE 17 BOTTOM FAUNA DATA - ABUNDANCE

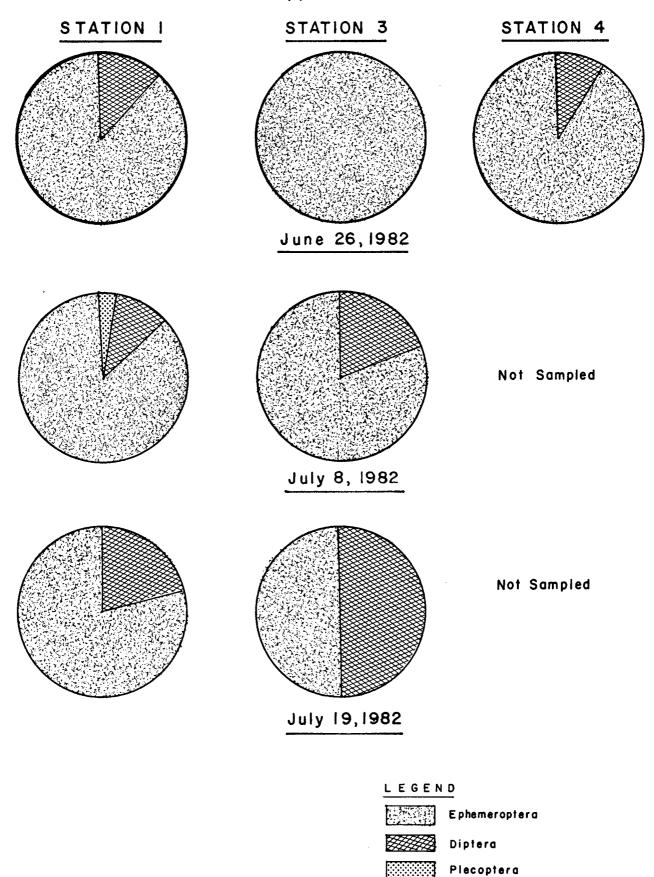


FIGURE 18 BOTTOM FAUNA DATA - PERCENT ABUNDANCE

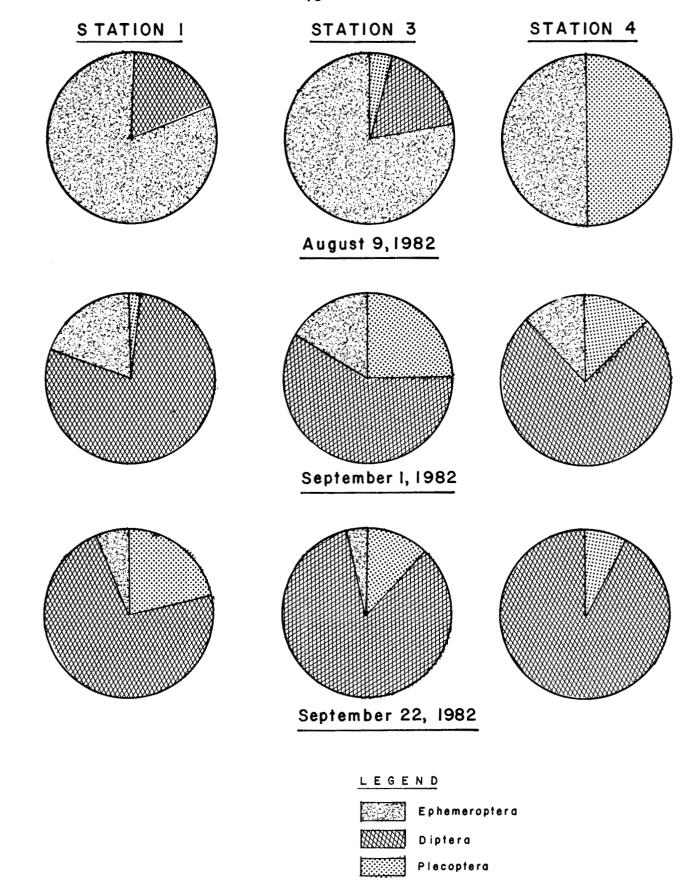


FIGURE 19 BOTTOM FAUNA DATA - PERCENT ABUNDANCE

Because of their sensitivity to low levels (10 - 15 mg/L) of suspended sediments (Rosenberg and Snow, 1977), the invertebrate populations are important as monitoring agents of environmental changes resulting from siltation. Increased levels of suspended sediments may fill interstices in the gravel substrate, thereby reducing the exchange of oxygenated water crucial to the survival of macro-invertebrates. In addition to this, increased sedimentation can change substrate composition, which may correspondingly change the community structure more suitable to the new habitat. A change in benthic habitat as a result of sedimentation usually results in a decline in important fish food organisms (Trichoptera, Ephemeroptera, Plecoptera) and the replacement with more tolerant species such as Chironomidae and Oligochaeta (Griffiths and Walton, 1978).

The three most abundant orders were chosen to determine if any general observations could be made with regard to their numbers as related to sluicing or non-sluicing conditions, or seasonal trends influenced by sluicing. Eighty-seven individuals of the order Ephemeroptera were found at Station 1 on July 8, representing 86% of the three orders. Numbers of individuals declined to 25 on July 19 and 16 on August 9, but still represented 78% and 80% respectively of the three most abundant orders. This reduction in population continued to the end of the summer. Seasonal succession of emergence could be a significant factor to this decline according to Usinger (1956). Conversely, individuals from the order Diptera increased in numbers at Station 1 on September 1 and 22 where mean sediment composition consisted of 91.7% and 97% medium coarse sand to gravel (300->2360 µm).

Abundance of Ephemeroptera was similar at Station 3 on June 26 and July 8 with individuals numbering 13 and 17 respectively. Mean sediment composition for July 8 only consisted of 7.8% silt and fine sand. Numbers, however, were reduced to 1 individual on July 19 where sediment was composed of 67.4% fine sand and silt, and increased to 44 on August 9 where substrate was comprised of 97% fine sand and silt. Numbers of Ephemeroptera on the two sluicing dates were quite inconsistant with each other. Ninety-seven percent fine sand and silt on August 9 did not provide a habitat where one would expect to see an

increase of Ephemeroptera. This cleanwater invertebrate group is known for its inability to survive high sediment levels (Langer, 1980). All of this might lead to the conclusion that a significant degree of variability existed in the collection of the sediment samples. The last two sampling periods in September show a reduction of Ephemeroptera which may be linked to a natural seasonal decline due to colder water temperatures. Ephemeroptera at Station 4 numbered ten or less throughout the sampling period.

Individuals of the order Diptera were less than 25% in abundance at the control station on all dates except on September 1 and September 23 where they represented 79% and 73% respectively. Numbers of individuals of Station 3 remained less than ten up to and including August 9. However, at this time individuals increased to 37 on September 1, where sediment was composed of 96.1% medium coarse sand to gravel (300 to >2360 um). The number of individuals at Station 3 on September 22 also increased to 52 where sediment was composed of approximately 89.6% silt to fine sand (<300 um). Both increases to the Diptera population occurred under two different sediment conditions. On August 9, zero individuals were collected at Station 4 where sediment was composed of 79.7% coarse sand to gravel. Eighteen (72%) and 55 (92%) individuals were collected at Station 4 on September 1 and 22 respectively. Sediment composition consisted of 97.6% medium sand to gravel (300 to >2360 um) on September 1 and 93.5% fine sand to silt (<75-300 um) on September 22.

Individuals of the order Plecoptera were extremely low in numbers throughout most of the operating season, but increased slightly at the end of the summer. Because numbers were so low at the control (Station 1), it was difficult to draw any meaningful conclusions. Perhaps the significant disturbance made upstream on Zinc Creek the year previous influenced the population, as this order is dependant on abundant oxygen supply for its survival (Pennak, 1978). In any case, low populations were present at the very end of the operating season at all three stations.

In spite of the two active sluicing periods, invertebrate results are not very conclusive. However, there appears to be a slight

decrease in total numbers of the three orders at Station 3 during sluicing periods (see Table 5). The order Diptera appears more tolerant as shown by the numbers at Stations 3 and 4 on September 1 and 22. These figures may suggest that Diptera individuals possess the ability to adapt more effectively to a stressed and changing environment created from sluicing activity than the more sensitive individuals belonging to the other two orders.

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#### **ACKNOWLEDGEMENTS**

The authors would like to thank R. Campeau for collating all the data and C. Ayris for preparing all the graphs. Also, special thanks to all the Environmental Protection Service regular and 1982 summer staff for their involvement in sample collection, lab analysis, drafting, typing and manuscript review.

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APPENDICES

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## APPENDIX I

COLLECTION, PRESERVATION, ANALYSIS OR IDENTIFICATION METHODS AND WATER QUALITY CRITERIA

WATER SAMPLE COLLECTION, PRESERVATION AND ANALYSIS METHODS TABLE 1 APPEND IX I

PARAMETER	DETECT! ON LIMIT	COLLECTION AND PRESERVATION PROCEDURE <sup>1</sup>	ME ANALYTICAL PROCEDURE	METHOD SECTION <sup>2</sup>
Temperature		In situ temperature reading.	Standard Centigrade Thermometer	
MO		In situ flow measurements using a Price-type current meter.	Cross section of the stream was measured and the velocity of flow was calculated using the standard Price-type Current Meter Method.	
Dissolved Oxygen	1.00 mg/L	Duplicate samples collected in 300 mL glass BOD bottles. The BOD bottles were rinsed 3 times with sample before filling. Preserved with 2 mL manganese sulphate and 2 mL alkali-iodide-azide solution and shaken 15 times. A water seal was maintained and DO analysis was done within 7 days.	lodometric Azide Modification Winkler Titration Method	0.48
Нd		Small aliquots of sample were taken and read soon after collection. No preservative.	Potentiometric	080
Conductivity	0.2 umhos/cm	In situ measurement. Laboratory measurement. No preservative. The measurement was taken from the same as NHz below.	YSI Conductivity Meter Model 33 Radiometer Conductivity Meter (CDMC) with radiometer conductivity cell.	044

WATER SAMPLE COLLECTION, PRESERVATION AND ANALYSIS METHODS (continued) TABLE 1 APPEND IX 1

PARAMETER	DETECTION LIMIT	COLLECTION AND PRESERVATION PROCEDURE <sup>1</sup>	ANALYTICAL PROCEDURE	METHOD SECTION <sup>2</sup>
Ammonia NH3-N	0.005 mg/L	Single samples collected in 2 litre linear polyethylene containers. Each container was rinsed 3 times with sample before it was filled. No preservatives. Stored at 4°C.	Phenol Hypochlorite-Colori- metric-Automated	058
Colour	5 (colour units)	Same sample as NH3.	Platinum-Cobalt Visual Compar- ison	040
Turbidity	0.1 (FTU)	Same sample as NH3.	Nephelometric Turbidity	130
Settleable Solids	0.1 mL/L	Same collection procedure as for NH3.	Settleable Matter	112
Non-Filterable Residue (NFR)	5 mg/L	Same sample as NH3.	Filtration, drying and weigh- ing of residue on filter	104
Filterable Residue (FR)	10 mg/L	Same sample as $NH_3oldsymbol{\mathfrak s}$	Filtration, drying and weigh- ing of filtrate	100
Total Alkalinity	1.0 mg/L as CaCO <sub>3</sub>	Same sample as NH3.	Potentiometric Titration	900
Total Phosphate T PO <sub>4</sub> -P	0.005 mg/L	Same sample as $NH_3ullet$	Acid-persulphate, Autoclave Digestion	980
Nitrite NO2-N	0.005 mg/L	Same sample as NH3.	Diazotization-Colorimetric- Automated	070

WATER SAMPLE COLLECTION, PRESERVATION AND ANALYSIS METHODS (continued) TABLE 1 APPEND IX |

PARAME TER	DETECTION LIMIT	COLLECTION AND PRESERVATION PROCEDURE <sup>1</sup>	ANALYTICAL PROCEDURE	METHOD SECTION <sup>2</sup>
Nitrate NO <sub>3</sub> -N	0.01 mg/L	Same sample as NH3.	Cadmium Copper Reduction Colorimetric Automated	072
Sulphate SO <sub>4</sub>	1.0 mg/L	Same sample as NH <sub>3</sub> .	Barium Chloranilate -UV Spectrophotometric	122
Chloride	0.5 mg/L	Same sample as NH3.	Thiocyanate-Combined Reagent- Colorimetric	024
Silica Total Si	0.5 mg/L	Same sample as NH <sub>3</sub> .	Ascorbic Acid Reduction - Colorimetric	118
Mercury Total Hg	0.0002 mg/L	Single samples were collected in a 200 ml linear polyethylene bottle. Preserved with a 10 ml 5% nitric dichromate solution.	Open Flameless System for Hg-AAS Determination	211 224 284 411
Extractable Metals A1 As Ba Be Ca Cd Co Cr Cu	mg/L 0.05 0.05 0.001 0.001 0.005 0.005	Single samples collected in 200 ml linear polyethylene bottles. Each bottle was rinsed 3 times with sample before filling. Preserved to a pH <1.5 using 2.0 ml concentrated HNO3.	Inductively Coupled Argon Plasma (ICAP) combined with Optical Emission Spectrometer (OES)	592

WATER SAMPLE COLLECTION, PRESERVATION AND ANALYSIS METHODS (continued) TABLE 1 APPEND IX I

PARAMETER	DE TECTION LIMIT	COLLECTION AND PRESERVATION PROCEDURE <sup>1</sup>	ANALYTICAL PROCEDURE	METHOD SECTION <sup>2</sup>
Extractable Metals (continued)	mg/Ł			
Fe	0.005			
D) Ε Σ W	0.001			
Mo	0.005			
Na	0.5			
: Z	0.02			
Pb	0.04			
Sb	0.05			
Sө	0.05			
Sn	0.01			
Sr	0.001			
1.	0.002			
>	0.01			
Zn	0.005			
AS.	0.0005	Same sample as metals.	Hydride Generation - ICAP	
P	0.0005	Same as sample metals.	Graphite Atomic Absorbtion	Δ+0m 1.
3 <del>1</del>	0.001	Same sample as metals.	Flameless Technique (AAS)	Absorption
Zn	0.002	Same sample as metals. Same sample as metals.		Jarrel I-Ash

WATER SAMPLE COLLECTION, PRESERVATION AND ANALYSIS METHODS (continued) TABLE 1 APPENDIX !

PARAMETER	DETECTION LIMIT	COLLECTION AND PRESERVATION PROCEDURE <sup>1</sup>	ANALYTICAL PROCEDURE	METHOD SECTION <sup>2</sup>
	0•0005 mg/L	Same sample as metals.	Graphite Furnace Atomic Absorption Spectrophotometry	210
	0.01 mg/L	Same sample as metals.	Flame Atomic Emission Spectro- photometry	210 423
Total Hardness	0.030 mg/L as CaCO <sub>3</sub>	Same sample as metals.	The sum of the ICAP results for Mg x 4.116 and Ca x 2.497 reported as mg/L CaCO <sub>3</sub>	
As described	As described in Environment Canada	da (1976).	÷	
As described	As described in Department of Envi	vironment (1979).		
As described	As described in the U.S. Dept. of	f the Interior (1975).		

Jarrel 1-Ash 850 Manual METHOD CODE 1 078 423 1 Department of Environment, Department of Fisheries and Oceans, Laboratory Manual, Environmental Protection Flame Atomic Emission Spectrophotometry Graphite Furnace Atomic Absorption Spectrophotometry SEDIMENT COLLECTION, PREPARATION AND ANALYSIS METHODS (continued) Standard Sieving Operation ANALYSIS (Unreliable in this type of digestion) Service, Fisheries and Marine Service (1979). Sample was oven-dried. Same as other metals. PREPARAT I ON TABLE 2 Metals (Leachable) (continued) Particle Size PARAMETER 0.08 0.08 2 0.2 0.3 2 0.3 APPEND IX I 0.2 0.2 8 Ag р C Р  $\checkmark$ 

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APPEND IX

IDENTIFICATION AND ENUMERATION	Bottom fauna was sent to Dr. C. Low, Consulting invertebrate Biologist, Nanaimo, B. C. for identification to genus, species if possible, and enumeration.
LABORATORY PROCEDURES	Bottom fauna was removed from other material in a labelled vial containing 70% methanol.
FIELD COLLECTION, SAMPLING PROCEDURES AND PRESERVATION	Surber Sampler: Creek and river samples were taken using a surber sampler with a 60 cm long net (mesh size 0.76 mm). Area sampled was 900 cm² (1 ft²). Surber samples were washed into a cup at the bottom of a plankton net (0.75 mm mesh size), put in separate labelled glass jars and preserved with 10% formalin. 3 samples were taken at each station.

WATER QUALITY CRITERIA FOR DRINKING WATER AND AQUATIC LIFE TABLE 4 APPEND IX I

SUBSTANCE	RECOMMENDED LEVEL(S) FOR DRINKING WATER	REFERENCE (S)	RECOMMENDED LEVEL(S) FOR AQUATIC LIFE	REF ERENCE (S)
Physical				
Colour Pt. Counts	15			
Odour and taste Turbidity J.T.U.	5			
Chemical				
Alkalinity mg/L	Not considered a public			
(Total)		4	>20	۶
Aluminum (AI) mg/L	Not considered a public			
	health problem	7	0.1	5
Ammonia (NH3-N) mg/L Antimony (Sb) mg/L	0.5 (Total)	4	0.02 (un-ionized)	٤
Arsenic (As) mg/L	0.05		0.05	2 (total)
Barium (Ba) mg/L	1.0	-	5.0	7
Boron (Bo) mg/L	1.0	_		
Cadmium (Cd) mg/L	0.005	-	0.0002	2 (total)
Calcium (Ca) mg/L	75-200	7		
Chloride (CI)mg/L	250	-		
Chromium (Cr) mg/L Cobal+ (Co) mg/L	0.05	-	0.04	2 (total)
Conductivity @ 25°C	Depends on dissolved			
(umhos/cm)	salts	7	150-500	v
Copper (Cu) mg/L	1.0		0.002	2 (total)
Cyanide (CN) mg/L	0.2	-	0.005	
Dissolved oxygen (% saturation)	Near 100%	4	>5.0 mg/L	· •
			ı	

WATER QUALITY CRITERIA FOR DRINKING WATER AND AQUATIC LIFE (continued) TABLE 4 APPEND IX I

SUBSTANCE	FOR DRINKING WATER	REF ERENCE (S)	FOR AQUATIC LIFE	REFERENCE(S)
Fluoride (F) mg/L	1.5		1.5	7
Hardness (Total)				•
as $mg/L$ CaCO $_3$	80-100	-		
Iron (Fe) mg/L	0.3	-	1.0	٢
ead (Pb) mg/L	0.05	-	0.005 (soft H <sub>2</sub> O*)	2 (total)
			0.01 (hard H <sub>2</sub> 0*)	2 (total)
Magnesium (Mg) mg/L	50	4	7	
Manganese (Mn) mg/L	0.05	-	1.0	7
Mercury (Hg) mg/L . Molybdenum (Mo)	0.002	-	0.0001-0.0002	2 (total)
Nickel (Ni) mg/L	0.25	2	0.025 (soft H <sub>2</sub> O*)	2 (total)
			0.25 (hard H2O*)	
Nitrate (NO <sub>3</sub> -N) mg/L	10	_	, Z.,	
Nitrite (NO2-N) mg/L	0.001			
pH units	6.5 - 8.5	_	6.5 - 9.0	~
Phosphorus (P) mg/L				•
(Total)			0.020 to prevent algae	5
Potassium (K) mg/L				
Residue: Filterable mg/L			70 - 400 with a maximum	
(Total dissolved solids)	1000	4	of 2000	9
Residue: Non-Filterable (mg/L)				
Selenium (Se) mg/L	0.01	•	0.01	(1e+0+) (
Silica (Si) mg/L				
Silver (Ag) mg/L	0.05	-	0.0001	2 (total)
Sodium (Na) mg/L	20	-		
Strontium (Sr) mg/L	10	_		
Suiphate (SO <sub>4</sub> ) mg/L	500	-		

WATER QUALITY CRITERIA FOR DRINKING WATER AND AQUATIC LIFE (continued) TABLE 4 APPEND IX I

SUBSTANCE	RECOMMENDED LEVEL(S) FOR DRINKING WATER	REF E RENCE (S)	RECOMMENDED LEVEL(S) FOR AQUATIC LIFE	REFERENCE(S)
Tin (sn) mg/L	Not present in natural waters	7		
Titanium (Ti) mg/L				
Total Inorganic Carbon (TIC)				
Total Organic Carbon (TOC)	5.0	Lo		
Vanadium (V) Zinc (Zn) mg/L	5.0	-	0.030	īV
* Soft water has a total hardness 95 mg/L as ${\sf CaCO}_3$ (Reference 6).		3003. Hard water ha	. Hard water has a total hardoness of more than	ə than
REFERENCE S:				
1. Health & Welfare Canada,	Health & Welfare Canada, Guidelines for Canadian Drinking Water Quality 1978,	ıking Water Quality	1978, Supply and Services, Canada (1979).	, Canada (1979).
2. Inland Waters Directorate, Guidelines Environment Canada, Ottawa	Inland Waters Directorate, Guidelines for Surface Water Quality, Vol. 1, Inorganic Chemical Substances. Environment Canada, Ottawa (1979, 1980).	er Quality, Vol. 1,	Inorganic Chemical Substa	ances.
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RECOMMENDED LEVEL(S)  SUBSTANCE FOR DRINKING WATER REFERENCE(S) FOR AQUATIC LIFE REFERENCE(S)	4. Anonymous, Guidelines for Establishing Water Quality Objectives for the Territorial Waters of the Yukon and Northwest Territories. Report of the Working Group on Water Quality Objectives to the Chairmen, Water Boards, Yukon and Northwest Territories, July (1977).	5. Ontario Ministry of the Environment, Water Management - Goals, Policies, Objectives and Implementation Procedures of the Ministry of the Environment. (1978).	6. Environment Canada, Pollution Sampling Handbook. Pacific Region Laboratory Services, Fisheries Operations and Environmental Profection Service, West Vancouver, B.C. (1976).	7. California State Water Resources Control Board, Water Quality Criteria. Publication No. 3-A Second Edition by McKee and Wolf. (1963).			
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APPENDIX II

WATER QUALITY DATA

APPENDIX II

## BARLOW PLACER STUDY WATER QUALITY DATA

PARAMETER		STATION 1	STATION 3	STATION 4
Fiow m <sup>3</sup> /s	a b c d e f	0.2 0.7 0.3 0.1	0 · 2 1 · 4 0 · 4 0 · 2 0 · 4	0.3 0.3 0.9 0.2 0.2
Temperature (°C)	a b c d e	8 10 7 9 6 4	8 10 9 11 7 4	7 11 9 11 7 4
Dissolved Oxygen (mg/L)	a b c d e f	 11.7 10.1 10.6 11.2 12.0	 10.8 9.6 10.1 10.8 11.9	 10.6 9.8 10.0 10.8
% D.O. Saturation (%)	a b c d e f	 111 89 98 97 98	96 88 97 94 97	 102 90 96 94 96

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

APPENDIX II

PARAMETER		STATION 1	STATION 2	STATION 3	STATION 4
Settleable Solids (mL/L)	a b c d e f	 <0.1 <0.1 <0.1 <0.1	 0.5 0.8 <0.1 0.4	<pre> &lt;0.1 0.9 &lt;0.1 &lt;0.1 &lt;0.1</pre>	 0.2 <0.1 <0.1
Non-Filterable Residue Whitehorse Lab (mg/L)	a b c d e f	 47 <5 <5 <5	 6990 4530 60 452	 1008 984 50 211	 570 2065 40 142
Non-Filterable Residue Vancouver Lab (mg/L)	a b c d e f	7 48 <5 <5	   	20 1160 110 75 211	 40 526 945 38 138
Filterable Residue (mg/L)	a b c d e f	148 103 131 183 187	   	 155 140 190 185 208	 163 146 185 182 189

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

APPENDIX II

PARAMETER		STATION	STATION	STATION	STATION
				31/11/01	1 2 I WILLOW
		1	2	3	4
рН	а				
In Situ	ь	7.8	7 • 4	7.8	7.7
	С			7.8	7.3
	d	8.4	7.9	8 • 2	7.9
	е		6.9	7 • 1	7.5
	f	8.0	7.9	7.8	
<del></del>					
- 11					
pH	a	7.7		7.8	7.8
Laboratory	b	1		6.7	6.7
	C	6.7		1	i .
	d	8.1		7.6	7.6
	e	7.6		7.6	7.6
	f	7.7	****	7.7	7.7
		405		100	100
Conductivity	a	105		108	102
In Situ	Ь	135		140	150
(umhos/cm)	С	80		82	80
	d	152		158	157
	е	158		160	163
	f	143		162	153
011					
Conductivity	a				
Laboratory	Ь	214		213	216
(umhos/cm)	С	125		124	121
	d	187		231	239
	e	260		255	260
	f	270	1	270	270

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

APPENDIX II

PARAMETER		STATION 1	STATION 3	STATION 4
Colour (colour units)	a b c d e	7 65 7 5 5	 15 >100 † 20 5	15 >100 t 10 5
Turbidity (FTU)	a b c d e	 <0.1 2.9 0.1 <0.1	10.0 800 1280 30.0 180	13.5 600 1200 10.8 95.0
Total Alkalinity (mg/L as CaCO <sub>3</sub> )	a b c d e	54.0 34.0 48.0 62.0	52.0 34.0 52.0 61.0 62.0	54.0 34.0 56.0 61.0 62.0
Total Hardness (mg/L as CaCO <sub>3</sub> )	a b c d e f	 100 66•4 110 124 76•6	99•1 195 214 129 94•0	106 130 150 126 133

# Date sampled:

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

\* Active Sluicing

<sup>†</sup> Could not remove all turbidity from sample; therefore all colour units will be falsely high

APPENDIX II

PARAMETER		STATION 1	STATION 3	STATION 4
Total	a			
P04-P	Ь	<0.005	0.020	0.030
(mg/L)	c	0.057	0.920	0.510
3	d	<0.005	1.05	1.00
	е	0.008	0.033	0.021
	f	0.005	0.225	0 • 1 4 5
NO -N				
NO <sub>2</sub> -N	a	<0.005	0.007	0.008
(mg/L)	Ь	0.008	0.007	0.031
	c	<0.005	0.037	0.031
	e	<0.005	0.034	<0.005
	f	<0.005	<0.005	<0.005
NO3-N	a			
(mg/L)	Ь	0.07	0.05	0.05
	c	0.01	0.03	0.05
	d	0.05	0.04	0.04
	е	0.04	0.03	0.04
	f	0.09	0.11	0.10
NH3-N †				
(mg/L)	a b	<0.005	0.005	0.006
(mg/L/	c	0.071	0.083	0.060
	d	<0.005	0.072	0.056
	e	<0.005	0.011	0.005
	f	<0.005	0.010	0.011
_				

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

- † All values for un-ionized  $NH_3$ -N are less than 0.02 mg/L
- \* Active Stuicing

APPENDIX II

PARAMETER		STATION 1	STATION 3	STATION 4
SO <sub>4</sub>	a			
4	Ь	62.5	55.5	54.5
	c	25.4	26.1	25.7
	d	42.0	54.0	55.0
	e	62.5	62.5	63.5
	f	66.5	71.5	72.0
CI				
(mg/L)	a b	<0.5	0.5	0.5
\mg/ \( \)	c	1.0	6.6	4.6
	d	<0.5	3.0	2.3
	e	0.6	0.9	0.7
	f	0.6	0.6	0.6
Ag	a		10.07	
(mg/L)	b	<0.03 <0.03	<0.03 <0.03	<0.03 <0.03
	c		<0.03	<0.03
	d	<0.03 <0.03	<0.03	<0.03
	e f	<0.03	<0.03	<0.03
			(0.03	<b>10.03</b>
Ai	a			
(mg/L)	Ь	<0.05	0.25	0.48
5	c	0.92	11.2	6.24
	d	<0.05	8.07	3.95
	е	0.24	0.60	0.36
	f	<0.05	0.42	0.96
			<u> </u>	

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Stuicing

PARAMETER		STATION 1	STATION 3	STATION 4
As	а			
(mg/L)	b	<0.0005	<0.0005	0.0015
<b>3</b> . –	c	<0.0005	0.0043	0.0026
	d	<0.0005	0.0026	0.0022
	e	<0.0005	0.00072	<0.0005
	f	<0.0005	0.0021	0.0022
Ва	a			
(mg/L)	Ь	0.033	0.041	0.063
g,	c	0.046	0.440	0.244
		0.034	0.497	0.339
	e	0.042	0.057	0.048
	f	0.018	0.052	0.072
Be	а			
(mg/L)	Ь	<0.001	<0.001	<0.001
( mg/L/	c	<0.001	<0.001	<0.001
	d	<0.001	<0.001	<0.001
	e	<0.001	<0.001	<0.001
	f	<0.001	<0.001	<0.001
Ca	a			
(mg/L)	Ь	25.3	24.6	25.9
	c	14.2	19.8	16.4
	d	28.0	32.0	27 • 2
	e	31.4	31.5	31.4
	-			

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

APPENDIX II

## BARLOW PLACER STUDY WATER QUALITY DATA (Continued)

PARAMETER		STATION 1	STATION 3	STATION 4
Cd	a			
(mg/L)	ь	<0.0005	<0.0005	<0.0005
g,	c	<0.0005	0.0005	<0.0005
	d	<0.0005	<0.0005	<0.0005
	e	<0.0005	<0.0005	<0.0005
	f	<0.0005	<0.0005	<0.0005
Co				
(mg/L)	a b	<0.005	<0.005	<0.005
( mg/L)	1 1	<0.005	0.038	0.022
	c d	<0.005	0.042	0.022
	e	<0.005	<0.005	<0.005
	f	<0.005	<0.005	<0.005
Cr	a			-~-
(mg/L)	Ь	<0.005	<0.005	<0.005
	c	0.060	0.019	0.010
	d	<0.005	0.009	<0.005
	е	<0.005	<0.005	<0.005
	f	<0.005	<0.005	<0.005
Cu				
	a	0.004	0.006	0.006
(mg/L)	Ь	0.004	0.000	0.006 0.030
	c d	<0.001	0.042	0.035
	e	0.001	0.002	0.025
	f	<0.001	0.002	0.002
	'	<b>\0.001</b>	0.009	0.007

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982
- Active Sluicing

APPENDIX II

PARAMETER		STATION 1	STATION 3	STATION 4
Fe	a			
(mg/L)	Ь	0.059	0.383	0.957
<b>3</b>	c	1.76	22.9	12.6
	d	0.044	17.4	8.74
	e	0.345	1.05	0.560
	f	<0.005	0.429	1.48
Hg	a			
(mg/L)	Ь	<0.0002	<0.0002	<0.0002
(mg/L)	c	<0.0002	<0.0002	<0.0002
	d	<0.0002	<0.0002	<0.0002
	e	<0.0002	<0.0002	<0.0002
	f	<0.0002	<0.0002	<0.0002
К	a			
(mg/L)	b	0.33	0.40	0.63
\mg/L/	c	0.36	0.98	0.69
		0.36	1.10	0.72
	e	0.41	0.74	0.68
	f	0.43	1.64	1.49
Mg				
mg (mg/L)	a b	8.9	8.6	9.0
(mg/L/	c	5.4	9.4	7.3
	d	9.6	12.9	9.9
	e	10.7	10.9	10.7
	1 1	6.7	7.9	11.1

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

PARAMETER		STATION 1	STATION 3	STATION 4
Mn	a			
(mg/L)	ь	0.022	0.046	0.089
<b>J</b>	c	0.139	1.47	0.874
	0	0.023	2.60	1.80
	e	<0.001	0.016	<0.001
	f	<0.001	0.046	0.074
Мо	a			
(mg/L)	Ь	<0.005	<0.005	<0.005
(mg/L)	c	<0.005	<0.005	<0.005
	d	<0.005	<0.005	<0.005
	e	<0.005	<0.005	<0.005
	f	<0.005	<0.005	<0.005
Na	a			
(mg/L)	Ь	3.0	2.9	3.0
	C	1.3	3.1	3.0
	d	3.0	3.2	2.8
	е	3.3	3.4	3.4
	f	2.3	2.6	3.4
Ni	a			
(mg/L)	Ь	<0.02	<0.02	<0.02
· · · g, = ·	c	<0.02	0.04	0.03
	d	<0.02	0.03	<0.02
	e	<0.02	<0.02	<0.02
	f	<0.02	<0.02	<0.02

## Date sampled:

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

\* Active Sluicing

PARAMETER		STATION 1	STATION 3	STATION 4
Pb	а			
(mg/L)	Ь	<0.001	<0.001	<0.001
<b>3</b>	c	0.001	0.017	0.009
	d	<0.001	0.016	0.016
	e	<0.001	<0.001	0.050
	f	<0.001	0.002	0.002
Sb	a			
(mg/L)	Ь	<0.05	<0.05	<0.05
,	c	<0.05	<0.05	<0.05
	d	<0.05	<0.05	<0.05
	e	<0.05	<0.05	<0.05
	f	<0.05	<0.05	<0.05
Se				
(mg/L)	a b	<0.05	<0.05	<0.05
( mg/L)	c	<0.05	<0.05	<0.05
	d	<0.05	<0.05	<0.05
	e	<0.05	<0.05	<0.05
	f	<0.05	<0.05	<0.05
Si	a			
(mg/L)	Ь	4.2	4.4	4.3
g / C /	c	3.2	3.9	3.7
	a	3.7	4.6	4.6
	e	4.4	4.5	4.4

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

<del> </del>			<del></del>	<del> </del>
PARAMETER		STATION 1	STATION 3	STATION 4
Sn	a			
(mg/L)	Ь	<0.01	<0.01	<0.01
,g, _ ,	c	<0.01	<0.01	<0.01
	d	<0.01	<0.01	<0.01
	e	<0.01	<0.01	<0.01
	f	<0.01	<0.01	<0.01
Sr ( )	a	0.140	0.140	0.150
(mg/L)	Ь	0.149	0.142	0.152
	C	0.084 0.167	0.133	0.106
	d	0.184	0.208 0.185	0.172
	e f	0.118	0.140	0.184
		0.116	0.140	0.191
Τi	a			
(mg/L)	Ь	<0.002	0.005	0.010
, and the second	c	0.004	0.036	0.024
	d	<0.002	0.026	0.017
	e	<0.002	0.004	<0.002
	f	<0.002	<0.002	0.006
V				
(mg/L)	a	<0.01	<0.01	<0.01
viiig/L/	C	<0.01	0.03	0.01
		<0.01	0.03	0.01
	e	<0.01	<0.01	<0.01
	f	<0.01	<0.01	<0.01

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982

<sup>\*</sup> Active Sluicing

PARAMETER		STATION 1	STATION 3	STATION 4
Zn (mg/L)	a b c d e f	0.011 0.024 <0.002 <0.002 <0.002	 0.015 0.111 0.076 <0.002 0.005	0.015 0.069 0.044 <0.002 0.012

- a) June 26, 1982 d) August 9, 1982\* b) July 8, 1982 e) September 1, 1982 c) July 19, 1982\* f) September 22, 1982
- \* Active Sluicing

APPENDIX III

SEDIMENT DATA

BARLOW CREEK SEDIMENT PARTICLE SIZE ANALYSIS - STATION 1 TABLE 1 APPEND IX 111

PERCENT COMPOSITION	> 2360 um 1180 - 2360 um 300 - 1180 um 150 - 300 um 75 - 150 um < 75 um	7		12.4 5.1 43.5 16.7 0.6	0.0 0.2 20.3	6.8 14.3 55.8 14.3 5.3	75.3	11.1 23.2 3.9 2.0		20.8 29.7 3.7	1.0 53.6	4.1 57.7 23.0 9.0	14.4 19.7 3.2 1.3			4.5	6.1	4.1 56.5 19.7 10.8	11.0	14.6 52.3 3.9 1.9	MEAN PERCENT COMPOSITION		37.9	1.0 31.0 25.7 23.8	7.5 56.6 19.0 8.4	14.0 1.8 0.7	13.4 34.1	
	> 2360 um	Gravel	!					58.3	-	39.9	0.2	0.7	60•3	48.3	-	22.9	0.5	9.0	75.8	26.1			25.2	0.2	2.7	70.5	44.2	
	_	SAMPLE DATE NUMBER 1982	June 26	July 08	1 - 1 July 19	Aug. 09	Sep + 01	Sept 22	June 26	July 38	1 - 2 July 19	Aug. 09		Sept 22	June 26		1 - 3 July 19			Sept 22		June 26	STN. July 08		Aug. 09		Sep† 22	

BARLOW CREEK SEDIMENT PARTICLE SIZE ANALYSIS - STATION 3 TABLE 1 APPEND IX 111

NO	150 - 300 um 75 - 150 um < 75 um	nd Fine Sand Very Fine Sand Silt	-	1.4	-	10.7 24.7 62.5		_		-	17.4 31.9 28.8			3.2		. —	10.8	12.6 29.5 53.7	1.6	12.5	
PERCENT COMPOSITION	300 - 1180 um 150 - 30	Med. Coarse Sand Fine Sa			10.0				-		19.5					19.8					
PERCE	1180 - 2360 um 30	Very Coarse Sand Med	-	24.5	1.5	0.1	5.7	7.0	-	22.6	6•1	0.2	3.8	0.2	!	21.3	9.11	0.2	7.8	0.3	
	> 2360 um	Gravel	-	45.8	0.5	0.2	87.2	24.6	* 1	43.5	9•0	0.0	88.1	0.1	;	52.5	6.1	1.6	85.8	0.1	
	_	DATE 1982	June 26	July 08	July 19	Aug. 09	Sept 01	Sept 22	 June 26	July 08	July 19	4ng 09	Sep+ 01	Sept 22	June 26	July 08	July 19	Aug. 09	Sept 01	Sept 22	
	(	SAMPLE			3 - 1						3 - 2						3 - 3				

MEAN PERCENT COMPOSITION

_						
June 26	-	1	!	!	1	!
July 08	47.2	22.7	22.3	2.3	1.7	3.8
	2.4	5.1	25.1	12.9	27.4	27.1
	09*0	0.1	2.3	13.5	31.1	52.4
	86•0	5.8	4.3	1.3	1.5	-:
	8,30	0.40	1.7	3.4	6.7	79.5

BARLOW CREEK SEDIMENT PARTICLE SIZE ANALYSIS - STATION 4 TABLE 1

APPEND IX 111

			Ψ1	PERCENT COMPOSITION			
	-	> 2360 um	1180 - 2360 um	300 - 1180 um	150 - 300 um	75 - 150 um	< 75 im
SAMPLE	DATE 1982	Gravel	Very Coarse Sand	Med. Coarse Sand	Fine Sand	Very Fine Sand	Silt
	June 26		1	-			
		!	;	-		!	!
4 - 1	July 19	:	!	1	;		
	Aug. 09	27.7	14.5	43.0	5.8	4.5	4-4
	Sep† 01	78.6	9.1	10.4	8.0	0.5	0.5
	Sept 22	0.0	0.1	0.4	6.0	3.9	94.7
	June 26	!.	i	-	1 1		
	July 08	}	;	-	!	!	
4 - 2	July 19	!	1	;	-	!	1
	Aug. 09	13.1	10.0	57.1	10.1	5.9	8.5
	Sept 01	82.8	8.1	6.8	0.5	9•0	2
	Sept 22	0.2	0.2	0.3	8.0	6.9	91.6
_							
	June 26	1	:	!	† !	:	!
	July 08		!!!!	!	1 1	!!!	!
4 1 3	July 19	1	;		:	;	;
	Aug. 09	13.3	9.4	54.1	11.5	9•9	5.2
		74.8		10.7	6.0	0.1	1.5
	Sept 22	17.8	0	0.4	-:-	2.9	77.6
			MEA	MEAN PERCENT COMPOSITION	NO		
	June 26	;	1	!		1	
STN.	July 08	:	;	:	;	!	
4	July 19	-	;	;	1 2 3	;	
	Aug. 09	18.0	11.3	50.4	9.1	5.7	4.6
	Sept 01	78•8	9.5	9.3	0.7	0.7	0.
	Sept 22	0•9	0.1	0.4	6.0	4.6	88.0
					the state of the s		

APPENDIX III TABLE 2 RARLOW CREEK SEDIMENT CHEMISTRY DATA - STATION 1
(All concentrations given in mg/kg dry weight unless otherwise noted)

982	- 3					-	106	0.3	4 160	<0.3	10.9	28.1	22.7	24300	0.11	1140	4120	813	10.7	140	61	708	10	3610	<b>&lt;</b> 5	31.8	1150	35	57.7
September 22, 1982	1 - 2		<5	1 2400.	7.7	-	119	0.3	4670	7.0	12.2	28.2	23.8	24000	0.10	1060	4 200	711	<0.8	130	21	168	6	88	\$	35.3	1320	<2	57.2
Septe	-																												56.3
382	- J		ŝ	11900	8.3	11.7	9.66	<0.2	4140	<0.3	8.6	23.1	24.6	27300	01.0	1390	4340	624	4.6	140	21	731	6	3070	\$	30.0	1370	36	62.2
September 1, 1982	1 - 2		Ş	11300	6.9	5.5	98.7	<0.2	4150	<0.3	10.0	22.9	23.7	24900	0.07	1210	4 200	585	5.1	150	22	744	ω	3930	\$	29.2	1110	34	59.2
des	-																												57.8
2	۲ -		\$	11400	0.9	13.3	112	<0.2	3750	<0.3	11.6	25.5	23.5	21500	0.08	1030	4050	619	2.9	140	21	597	6	3680	<b>4</b>	28.3	867	32	55.5
August 9, 1982	1 - 2		\$	10900	5.9	0.6	109	<0.2	4100	<0.3	10.0	24.8	21.3	20900	0.16	970	3810	583	3.0	140	18	695	0	3950	\$	7.62	917	33	52.7
- V			\$	11100	7.2	12.2	103	<0.2	3570	<0.3	10.3	25.1	23.3	21800	0.10	866	4070	653	3.0	130	20	296	9	3850	2	26.6	865	30	57.4
2	- 3		\$	7380	5.9	3.8	86.8	<0.2	2960	<0.3	5.7	17.2	15.6	14400	0.14	398	2960	349	1.7	100	14	741	ī	2480	<b>\$</b>	17:6	309	21	43.5
July 19, 1982	1 - 2		ŝ	7240	6.2	4.0	70.1	<0.2	2590	<0.3	9.4	17.7	19.5	15500	0.16	383	3230	418	2.5	80	17	700	01	2380	\$	14.7	257	18	46.3
ſ	-		Ĉ	7020	4.5	1.6	72.4	<0.2	2640	<0.3	4.5	16.4	17.5	14300	0.14	374	2990	414	2.4	8	14	670	9	2340	\$	15.6	258	18	43.3
2	1 - 3		£	7410	4.8	8.6	86.3	<0.2	2470	<0.3	7.3	18.7	19.9	15400	0.11	317	3250	515	0.1	70	92	592	7	2700	\$	16.9	172	17	48.3
July 8, 1982	1 - 2	1				5.1																							
	Ä:-		\$	7370	5.4	4.5	86.1	<0.2	2610	<0.3	7.2	18.4	18.7	14900	0.05	348	3110	451	8°0>	100	18	607	r.	2850	\$	17.3	216	18	47.3
DATE:	STAT I ON:		Αg	¥	As	<u>в</u>	Ba	Be	ပီ	PS	ပိ	ن د	n <sub>o</sub>	F.e	Нg	<u>~</u>	ω	ξ	ě.	e Z	ž	a	P Q	SI	-S	Sr	Ξ	>	uz

0.10 1570 4860 558 <0.8 17 643 15 5230 <2 22.8 638

26900

120 <0.2 2820

3 - 3

September 22, 1982 11.8 26.7 36.6 29900 0.12 1650 5480 600 14.0 180 26 717 18 3960 2 2 2 2 29.8 924 - 2 <0.3 0.3 3590 3880 <0.3 11.5 27.4 32.2 29100 0.10 1790 589 13.1 210 24 719 16 2300 3 32.1 1120 388 99.0 0.03 1580 4340 655 4070 4070 40.3 13.7 29.2 24.1 5.3 150 24 664 12 3040 <2 29.3 1170 38 3 - 3 September 1, 1982 0.03 1530 4460 687 7 <0.3 10.7 22.9 25.1 5.0 140 23 703 10 10 <2 30.4 954 37 26000 25000 3850 <0.3 9.5 21.7 23.7 3 - 1 14.2 119 <0.2 150 22 266 8 4120 <2 27.5 1020 35 59.8 1440 4200 609 5.7 3 - 3 August 09, 1982 (All concentrations given in mg/kg dry weight unless otherwise noted) 3540 <0.3 9.6 22.2 19.6 19400 3 - 2 11.7 850 3930 540 2.4 <2
26.1
640
30
53.2</pre> 140 18 623 8 45 12800 6.7 9.3 168 40.0 40.5 14.2 24.7 25.1 22100 0.15 983 4610 696 170 22 696 10 3770 BARLOW CREEK SEDIMENT CHEMISTRY DATA - STATION 3 2.2 88.1 <0.2 2920 <0.3 8.1 17.3 16.1 14900 0.15 410 3060 377 2.0 100 15 717 6 2550 <2 17.8 317 21 3 - 3 July 19, 1982 6.6 15.4 15.5 13400 0.15 100 14 592 52320 42 15.2 220 18 6.6 80.8 <0.2 2430 <0.3 3 - 2 363 2870 362 4.4 7.8 67.9 60.2 2380 60.3 5.1 15.4 16.6 13700 0.16 2900 80 13 646 4 2030 m 3 - 3 July 8, 1982 5.3 10.0 129 40.2 3510 40.3 6.2 22.7 20.10 0.05 3110 <2 25.7 475 28 56.4 648 3940 899 <0.8 3 - 2 160 648 TABLE 2 5.8 40.2 2810 40.3 9.2 20.1 20.8 90.0 -471
3680
1120
<0.8
90
22
22
7
7
2960
<22
20.3
210
210
21 APPEND 1X 111 m STATION: DATE: 

BARLOW CREEK SEDIMENT CHEMISTRY DATA - STATION 4 (All concentrations given in mg/kg dry weight unless otherwise noted) TABLE 2 APPENUIX III

		`																												
	786	4 - 3		4.5	15800	16.9	į	141	0.3	3300	2.0.3	9.3	26.3	30.6	28000	0.10	1810	5210	618	13.2	961	g	88	13	2240	<b>&lt;</b> 2	31.8	1020	88	87.1
	September 22, 1982	4 - 2		< 5	14700	14.2	1	145	0•3	3730	<b>4.0.3</b>	10.9	25.4	27.4	26000	0.16	1540	5030	\$	12,4	170	೫	919	14	4050	<b>&lt;</b> 2	30.2	784	88	80.2
	Septe	4 - 1		45	12300	18.8	1	116	7 0.2	3060	< 0.3	10.9	21.1	30.7	23900	0.10	1340	4210	498	\$ <b>0</b>	130	15	961	12	4620	<b>&lt;</b> 2	24.0	728	8	80.0
	1982	4 - 3					_						_					_												64.7
	September 01, 1982	4 - 2																			_									59.1
(na	S d	4 - 1																												63.2
ICI WISC IN	81	4 - 3											_	_							_		_			_	_			47.3
20 6531110	August 09, 1982	4 - 2																												49.2
אל מין ארושווי מוווכזי סמומו אוזכ ווסנכת	Aug	4 - 1	1																											9.05
8v/8ii ii>4:6	286	4 - 3		!	!	1	!	1	!	1	!	!	!	1	!	1	1	1	!	;	;	;	;	1	1	;	1	}	;	;
	July 19, 1982	4 - 2		1	1	1	!	1	1	;	;	1	;	1	1	!	1	1	1	1	1	1	1	1	1	1	1	1	1	1
	اب	4 - 1		;	!	1	 	1	;	;	!	1	;	!	¦ 	1	1	!	!	1	1	;	1	1	1	1	1	!	  -	1
	011	4 - 3		}	1	;	1	;	1	1	<b>!</b>	1	;	ł	1	ł	1	ł	ł	<b>¦</b>	ł	1	ł	ł	1	1	1	ł	1	1
	July 8, 1982	4 - 2		¦	1	!	1	;	1	1	}	}	;	ł	;	;	ł	}	ŀ	<b>!</b>	ł	ł	ł	ł	ł	ł	ł	ł	ł	!
	J.	N: 4 - 1		!	!	1	!	!	!	!	!	1	!	!	1	!	!	1	!	1	1	!	1	;	!	;	<b>!</b>	1	1	1
	DATE:	STATION:		₽ E	<b>8</b>	As.	2	ga ,	æ ,	J.	3	3	<u>ئ</u>	3	به	완	¥	Σ Σ	٤	<u>Q</u>	Sa.	E	۵	£	Si	ঠ	۲	ï	>	υZ

Note: Sediment samples were not collected on July 8 and 19 due to high water conditions. Unable to collect a representative sample.

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APPENDIX IV

BOTTOM FAUNA DATA

APPENDIX IV TABLE 1 BOTTOM FAUNA TAXONOMIC GROUPS FOUND IN BARLOW CREEK

	Phylum:	Annelida
	FNY LUM:	Annerida
1	Class:	Oligochaeta
	Order:	Lumbriculida
	Family:	Lumbriculidae
1.		Kincaidiana hexatheca
		<del></del>
	Order:	Haplotaxida
2.	Family:	Enchytraeidae
	5	T 1.161 1.1
7	Family:	Tubificidae
3 <b>.</b>		Telmatodrilus sp. Tubifex sp.
5.		llyodrilus sp.
		1170071103
6.	Class:	Hirudinae
	Phylum:	Anthropoda
	01	Annalisation
7.	Class: Order:	Arachnoidae Acari
′•	order:	Acari
8.	Order:	Arachnida
	0, 00, 0	
	Class:	Insecta
	Order:	Plecoptera
9.		Subimago
	Family:	Capniidae
10.		Capnia sp.
	C-mil	Chlanaranlina
11.	Family:	Chloroperlinae Alloperla sp.
12.		Utaperla sp.
}		oruper ra sp.
	Family:	Nemouridae
	,	
]		

APPENDIX IV TABLE 1 BOTTOM FAUNA TAXONOMIC GROUPS FOUND IN BARLOW CREEK (Continued)

Tamily:	Periodidae
amily:	Arcynopteryx sp.
	Cultus sp.
	Megarcys sp.
	Skwala sp.
	unid. dam.
Order:	Ephemeroptera
amily:	Baetidae
	Ameletus sp.
	Baetis sp.
amily:	Heptageniidae
	Cinygmula sp.
	Epeorus sp.
amily:	Siphlonuridae
	Siphlonurus sp.
Order:	Trichoptera
	Limnephilidae
	Clostoeca sp.
	Ecclisomya sp.
Order:	Diptera
family:	Chironomidae
	Chironomidae adult
	Chironomidae pupae
	Brillia sp. Cardiocladius sp.
	Cricotopus sp.
	Diplocladius sp.
	Epoicocladius sp.
	Eukiefferiella sp.
	Heterotrissocladius sp.
	Orthocladius sp.
Subfamily	Diamesinae
	Tamily: Tamily: Order:

APPENDIX IV TABLE 1 BOTTOM FAUNA TAXONOMIC GROUPS FOUND IN BARLOW CREEK (Continued)

<del></del>		
36.		Diamesa sp.
37.		Monodiamesa sp.
38•	Family:	Pseudodiamesa sp.
39.	Family:	Culicidae
,,,		04.10.00
	Family:	Empididae
40.		Chelifera sp.
	Family:	Simulidae
41.		Prosimulium sp.
42.		Metacnephia sp.
	Family:	Tipulidae
43.		Erioptera sp.
44.		Tipula sp.
	Order:	Homoptera
45.	Family:	Aphididae
	,	,

Sept. 22/82 1-2 -111411-11011111-01111 1-3 Sept. 1/82 1-1 1-2 1-1111111001-0110111 1-3 August 9/82 1-2 -1-3 July 19/82 1-1 1-2 July 8/82 1-2 -1-3 June 26/82 1-2 <u>-</u> Heterotrissociadius sp. Piecoptera subimago TAXONOMIC GROUP Siphlonurus sp. Chironomidae adult Chironomidae pupae Eukiefferiella sp. Cardiocladius sp. Epoicocladius sp. Arcynopteryx sp. Orthocladius sp. Hyodrilus sp. Cricotopus sp. Enchytraeidae Cinygmula sp. Podmosta sp. Ameletus sp. Tubifex sp. Epeorus sp. Brillia sp. Baetis sp. Cultus sp. Skwala sp. Capnia sp. 10 13. 21. 14. 22. 23. 26. 27. 15. 17. 19. 20. 30. 29.

BARLOW CREEK BOTTOM FAUNA DATA - STATION 1

TABLE 2

APPENDIX IV

0.89 1-3 13 69 57 0.78 1-2 34 Ξ 22 26 0.70 0.66 1-3 35 23 Sept. 1/82 1-2 24 27 Ξ 0.44 0.56 0.70 1 1 1 1 1 1 27 1-3 Augus† 09/82 62 1-2 42 **b.**28 -0.73 0.41 0.52 5-20 17 July 19/82 1-2 5 -0.60 0.61 1-3 9 July 8/82 1-2 79 0.67 300 99 Ξ 1-3 × × × June 26/82 0.70 Column totals recorded per ft<sup>2</sup> 1-2 8 41 0.57 -73 32 Column Totals Total Number TAXONOMIC GROUP (H') Diversity Pseudodiamesa sp. Chelifera sp.
Prosimulium sp.
Metacnephia sp. 36. Diamesa sp. Culicidae adult X = Not sampled.Erioptera sp. Tipula sp. Aphididae E NOTE: 41. 39. 40. 42. 43.

BARLOW CREEK BOTTOM FAUNA DATA - STATION 1

7

TABLE

APPEND IX IV

TABLE 2 BARLOW CREEK BOTTOM FAUNA DATA - STAFLON 3

APPENDIX 1V

	June 26/82	July 9/82	July 19/82	August 9/82	Sept. 1/82	Sept. 22/82
TAXONOMIC GROUP	3-1 3-2 3-3	3-1 3-2 3-3	3-1 3-2 3-3	3-1 3-2 3-3	3-1 3-2 3-3	3-1 3-2 3-3
:						
1. Kincaidiana hexatheca	× '		'	1	-	,
2. Enchytraeidae	× -	,	,	'	-	ı
3. Telmatrodrilus sp.	×	,		ı	- 1	
4. Tubifex sp.	×	1		ı	· ·	
8. Arachnida	× !		1	,		1 1
10. Capnia sp.	× 1	1	;	,		
	× 1	;	,	:	۰ ،	
13. Podmosta sp.	× 1	1	;	,		
	× 1	,	1	'		
	× 1	,	1	,	2 2 2	
18. Unid. dam.	× ,	,	1		. ,	- 1
	, ×	;	1	2 - 1		
	- 3 ×		;	1 - 2	3 -	,
	5 2 ×	1 5 5	-	ŧ	-	1
	× 1	-	1	8 2 15	1 2 1	
23. Siphionurus sp.	× 1	1	;		1	f I
	× '		'	1	· ·	,
•	× '	,	:	1	1	- '
	× ,	2	1	1	1	1
	×	,	,	1		- 2 2
	× ,	1	1	1	-	
	× 1	,	1			
	× 1	,	;	2 - 1	1	
31. Diplocladius sp.	× 1	,	1	-	٠	1 1
34. Heterotrissocladius sp.	× '	,	,	;	-	

	June 26/82	28/93	-n <sub>r</sub>	July 8/82	61	200	July 19/82	32	August	i	9/82	Sept	Sept. 1/82	2	Sep	Sept. 22/82	82
IAXONOMIC GROUP	3-1 3-2	2 3-3	3-1	3-2	3-3	3-1	3-2	3-3	3-1	3-2	3-3	3-1	3-2	3-3	3-1	3-2	3-3
35. Orthocladius sp. 36. Diamesa sp. 37. Monodiamesa sp. 38. Pseudodiamesa sp. 39. Culicidae adult 40. Chelifera sp. 42. Metacnephia sp. 44. Tipula sp.		* * * * * * * *	11-1-11						01101111	1 1 1 1 1 1 1 1	1110111	101-111-	101-1-11	161-111	10101111	1 = 1 4 1 1 1 1	10141111
(T) Column Totals	9	×	<b>6</b> 0	9	80	2	58	82	27	٣	26	27	8	12	17	22	24
(N) Total Number	6 7	×	4	9	æ	2	58	18	27	٣	56	24	28	6	17	20	21
(H¹) Diversity	0.20 0.47	×	0.60 0.20		0.70	00.0	0.08	0.00	0.78	0.28	0.56	0.84	0.94	0.73	0.70	0.55	0.70
NOTE: Column totals recorded per ft <sup>2</sup> x = not sampled.	ded per fi	7	_		-						<del>-</del>						

BARLOW CREEK BOTTOM FAUNA DATA - STATION 3

TABLE 2

APPENDIX 1V

4-3 Sept. 22/82 4-2 4-1 4-3 Sept. 1/82 4-2 4-1 9/82 4-3 4-2 August 4-1 Bottom fauna was not collected on July 8 and 19 due to high water conditions. 4-3 July 19/82 4-2 4-1 July 8/82 4-2 4-1 4-3 June 26/82 4-2 4-1 sp. Heterotrissociadius TAXONOMIC GROUP Chironomidae pupae Cardiocladius sp. Cricotopus sp. Pseudodiamesa sp. Monodiamesa sp. Siphlonurus sp. Enchytraeidae 11. Alloperia sp. 13. Podmosta sp. Ameletus sp. Baetis sp. Cinygmula sp. Epeorus sp. Diamesa sp. Tubifex sp. Brillia sp. Cultus sp. Skwala sp. 6. Hirudinae 15. 19. 27. 28. 29. 30.

APPENDIX IV TABLE 2

BARLOW CREEK BOTTOM FAUNA DATA - STATION 4

	June 26/82		July 8/82	3/82	3	July 19/82	/82	Aug	August 9/82	/82	Sept	Sept. 1/82		Sept	Sept. 22/82	82
TAXONOMIC GROUP	4-1 4-2 4-3		4-1 4-2	2 4-3	4-1	4-2	4-3	4-1	4-2	4-3	4-1	4-2	4-3	4-1	4-2	4-3
40. Chelifera sp. 42. Metacnephia sp. 44. Tipula sp. 45. Aphididae	* * * *		× × × × ×	× × × ×	× × × ×	× × × ×	× × × ×	1 1 1	1 1 1	1111	~ 1 1 1		1 1 1	1 1 1 1	1 1 1	1 1
(T) Column Totals	* 17 *		×	×	×	×	×	-	7	2	24	23	æ	32	2	30
(N) Total Numbers	* 8		×	×	×	×	×	0	0	-	6	80	9	24	4	25
(H¹) Diversity	0.48 0.29 *		×	×	×	×	×	00.0	00.00	99.0 00.0 00.0		0.75	0.48	0.42	0.30	0.50
NOTE: Bottom fauna was not collected on	collected on	July 8	and 19	July 8 and 19 due to high water conditions.	. high	water	cond!†	ions.					_			
* = not sampled																

APPENDIX IV

BARLOW CREEK BOTTOM FAUNA DATA - STATION 4