









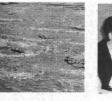








NORTHERN RIVER BASINS STUDY PROJECT REPORT NO. 59 **ECOTOXICOLOGY OF DEPOSITIONAL SEDIMENTS ATHABASCA RIVER MAY AND SEPTEMBER, 1993**







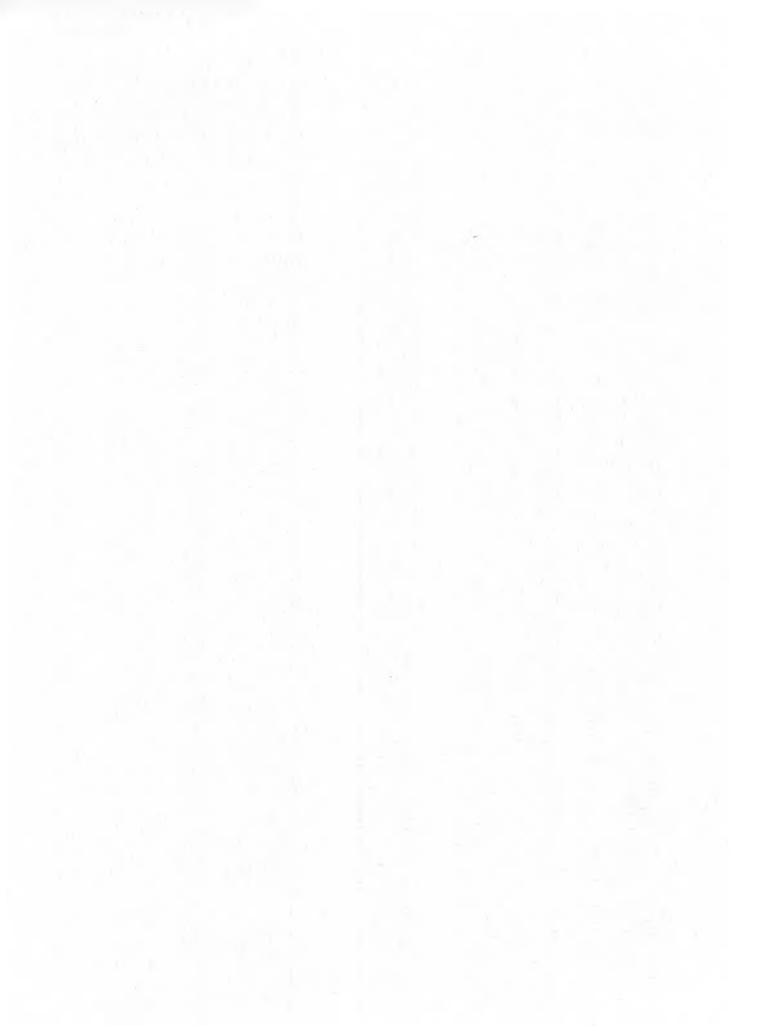








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by

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NORTHERN RIVER BASINS STUDY PROJECT REPORT NO. 59 ECOTOXICOLOGY OF DEPOSITIONAL SEDIMENTS ATHABASCA RIVER MAY AND SEPTEMBER, 1993

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PREFACE:

The Northern River Basins Study was initiated through the "Canada-Alberta-Northwest Territories Agreement Respecting the Peace-Athabasca-Slave River Basin Study, Phase II - Technical Studies" which was signed September 27, 1991. The purpose of the Study is to understand and characterize the cumulative effects of development on the water and aquatic environment of the Study Area by coordinating with existing programs and undertaking appropriate new technical studies.

This publication reports the method and findings of particular work conducted as part of the Northern River Basins Study. As such, the work was governed by a specific terms of reference and is expected to contribute information about the Study Area within the context of the overall study as described by the Study Final Report. This report has been reviewed by the Study Science Advisory Committee in regards to scientific content and has been approved by the Study Board of Directors for public release.

It is explicit in the objectives of the Study to report the results of technical work regularly to the public. This objective is served by distributing project reports to an extensive network of libraries, agencies, organizations and interested individuals and by granting universal permission to reproduce the material.

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ECOTOXICOLOGY OF DEPOSITIONAL SEDIMENTS ATHABASCA RIVER, MAY AND SEPTEMBER, 1993

STUDY PERSPECTIVE

Organic contaminants which enter aquatic ecosystems can become associated with particles of organic and inorganic materials in depositional zones. The presence and persistence of contaminants in these sediment depositional zones may constitute a source of toxicity to organisms which live on or near the substrate. As an example, benthic invertebrates are bottom-dwelling organisms that are very sensitive to environmental change. Toxicity from contaminants may have direct impacts on these species as well as indirect effects on other organisms which use them as food. Benthic invertebrates are considered good overall indicators of contaminants in sediments because, as a group, they are in direct contact with sediment solids. Sediment quality can be described through a three part study (TRIAD) examining the benthic invertebrate community structure, the toxicity of depositional sediments on selected life forms, followed by contaminant analyses of these sediments if results from toxicity testing are positive.

This project was designed to test the toxicity of depositional sediments from the upper Athabasca River using freshwater benthic invertebrates in

Related Study Questions

- 1a) How has the aquatic ecosystem, including fish and/or other aquatic organisms, been affected by exposure to organochlorines or other toxic compounds?
- 4a) What are the contents and nature of the contaminants entering the system and what is their distribution and toxicity in the aquatic ecosystem with particular reference to water, sediments and biota?
- 13b) What are the cumulative effects of manmade discharges on the water and aquatic environment?
- 14) What long term monitoring programs and predictive models are required to provide an ongoing assessment of the state of the aquatic ecosystems? These programs must ensure that all stakeholders have the opportunity for input.

chronic exposure studies under laboratory conditions. Additional analyses were performed on the *in situ* benthic invertebrate community structure to describe species distribution and abundance. Information from these tests will be used to determine cumulative effects of the Hinton combined effluent by comparing upstream and downstream sites.

In 1993, sediments and benthic invertebrates were collected from depositional areas on the upper Athabasca River upstream and downstream of Hinton; eight sites in May and seven sites in September. Sediments from these sites were subjected to chronic toxicity tests in the laboratory using four species of bottom-dwelling invertebrates; an amphipod, a chironomid, a mayfly and an oligochaete worm. The tests measured the effects of exposure to potentially contaminated sediments over a 10-day or 28-day period using the young of each species. The endpoints that were measured include survival, growth (amphipod, chironomid and mayfly) and reproduction (oligochaete worm). Sediments tested with the four species of invertebrates exhibited low toxicity in the laboratory with the exception of sediments collected downstream of Hinton, near the mouth of the Berland River and at the Windfall Bridge. At these two sites, reproduction of the oligochaete worm was reduced compared to the upstream control sites. Invertebrate species diversity was also reduced at Windfall Bridge for samples collected in the fall of 1993. Elevated levels of several metals at both sites, the high percentage of sand at the Berland River site, as well as other unmeasured contaminants may have contributed to the observed toxicity and reduced diversity.

Toxicity testing under laboratory conditions in this study indicated that further work is required. Nonetheless, this information will be incorporated into a multivariate statistical model to determine the environmental health for the reach of the river that was investigated. Results from this research will also assist with the task of cumulative effects assessment and development of biomonitoring guidelines for the rivers.

REPORT SUMMARY

The following report summarizes the results from the toxicological testing of depositional sediments in the upper Athabasca River using freshwater benthic invertebrates in chronic exposure studies under laboratory conditions as well as analyses of the *in situ* benthic invertebrate community structure. Sediments were collected at one or two sites upstream from Hinton (one control at site ARC in May 1993 and two controls at sites ARC and ARC2 in September 1993) and at five to six sites at varying distances downstream from Hinton on the same dates in the spring and fall. The four species of benthic invertebrates used in the tests were the amphipod, *Hyalella azteca*, the chironomid, *Chironomus riparius*, the mayfly, *Hexagenia spp.* and the oligochaete worm, *Tubifex tubifex*. The toxicity tests were conducted under controlled laboratory conditions and utilized young of each species. The tests measured the effects of exposure to potentially contaminated sediments over 10-d to 28-d depending on species. The endpoints determined were survival, growth (*H. azteca*, *C. riparius and Hexagenia*) and reproduction (*T. tubifex*). The structure of the benthic invertebrate communities at all sites were determined using percent abundance and three diversity indices (Shannon-Wiener, Simpson's and Margalef's).

All sediments tested with the four species of invertebrates exhibited low toxicity in the laboratory with the exception of sediments collected upstream of the Berland River (BR) and near Windfall Bridge (WB). At these two sites, reproduction by the tubificid worm, *T. tubifex*, was reduced compared to the control sites and this reduction was statistically significant at WB. Diversity was also reduced at Windfall Bridge for samples collected in the fall of 1993 for benthic invertebrate community structure. Levels of several metals, *i.e.*, arsenic, nickel, chromium and cadmium were slightly above the low effects level (LEL) set for another province (Ontario) at these two sites. In addition, the particle size distribution at BR-T consisted of a high percentage of sand. These two factors as well as other unmeasured contaminants may have contributed to the observed toxicity and reduced diversity.

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1.0 INTRODUCTION

Many organic contaminants which enter aquatic ecosystems become associated with particles of organic material in low-energy depositional zones in the benthic environments. The presence and persistence of contaminants which sorb to organic material in these zones may constitute a source of toxicity to organisms which live in or near the substrate such as epibenthic and burrowing invertebrates. Such toxicity may have direct detrimental effects on these species as well as indirect effects on the other organisms which use them as food (i.e., many species of fish, amphibians or shore-birds).

There have been very few studies in riverine environments in Canada which address the question of the toxicity of contaminants in sediments to organisms which live in or near this environment. This is mainly due to a shortage of standardized methodologies for assessing toxicity to benthic organisms as well as the lack of a multi-disciplinary approach to determining the effects of contaminants on ecosystem health. Benthic invertebrates have been traditionally considered the best overall indicators of contamination in sediments because, as a group, they are in direct contact with sediment solids as well as the interstitial waters and they have been effective in a wide range of studies (Burton et al. 1992; ASTM 1993; USEPA 1994). The Sediment Quality Triad (Chapman 1990) is an effects-based approach used to describe sediment quality which incorporates measures of sediment chemistry (grain size, metals, organic content, etc.), sediment toxicity (whole-sediment laboratory bioassays) and benthic infaunal community structure (diversity, richness, etc.).

Several species of invertebrates have been recommended as suitable organisms for the acute and chronic laboratory testing of sediments. The freshwater amphipod, *Hyalella azteca*, and the chironomids, *Chironomus tentans* or *C. riparius*, have received the most attention but other organisms have also been used such as the mayfly, *Hexagenia* spp.and the oligochaete worms, *Lumbriculus variegatus* or *Tubifex tubifex*. These organisms are found in lakes, ponds and streams throughout North America and have a variety of feeding habits which range from grazing on the surface of sediments for algae and organic detritus (e.g., *H. azteca)* to burrowing and ingesting sediment particles (e.g., *T. tubifex*). Thus, benthic invertebrates can be exposed to contaminanted sediments through a variety of mechanisms from passive diffusion of toxicants dissolved in the intersitial water to ingestion of particles of sediment to which contaminants have sorbed.

The objectives for this study were as follows: (1) to collect fine sediments from depositional areas upstream (two sites) and downstream (five sites) of Hinton in the upper Athabasca River; (2) to assess the toxicity of these sediments to four species of benthic invertebrates (the amphipod, *Hyalella azteca*, the chironomid, *C. riparius*, the mayfly, *Hexagenia* and the oligochaete worm, *T. tubifex*) in chronic laboratory toxicity tests; (3) to describe benthic invertebrate species distributions and abundances across stations; (4) characterize the sediments for their physical and chemical variables; (5) evaluate the toxicity of sediments using the Sediment Quality Triad.

2.0 METHODS

2.1 COLLECTION OF SEDIMENTS

2.1.1. SITES OF COLLECTION OF SEDIMENTS FOR BENTHIC INVERTEBRATE COMMUNITY ANALYSIS AND TOXICITY TESTS

Forty samples were collected in May 1993 (eight sites: ARC (Control), ATHA, HB, OB, EL, BR, BER, WB; five replicates each) and 35 samples (seven sites; ARC (Control), ARC2 (Control), HB, OB, EL, BR, WB; five replicates each) were collected in September 1993 using either a Ponar or Eckman grab for analysis of the benthic invertebrate community structure. The samples were preserved in formalin until analysis in the laboratory. Additional samples were collected in the fall of 1993 for use in whole sediment toxicity tests. For these tests, five replicates were collected at each site using an Ekman dredge and kept separate for use in bioassays (true field replicates). The samples were kept at 4°C following collection until they could be used in toxicity tests (approximately 6-8 weeks). The specific locations of the samples are outlined in more detail in Table 1.

2.2 PHYSICAL AND CHEMICAL ANALYSES OF SEDIMENTS

Particle size determination of each manipulated sediment was performed on lypholysized samples at the National Water Research Institute in Burlington, Ontario, Canada, following the procedure outlined by Duncan and LaHaie (1979). Large particles (>0.88 mm) were removed from the sediment sample prior to analysis. The sediment was then placed in a sodium metaphosphate solution, mixed for fifteen minutes and wet-sieved through a 0.063 μ m mesh. The material remaining on the sieve was dried, added to the large particles previously removed and the total was recorded as percent sand and gravel. The remaining suspension was analyzed using a sedigraph analyzer and results were expressed as percentage silt and clay.

Major elements, total phosphorous, total organic carbon (TOC), loss on ignition (LOI) and total Kjeldahl nitrogen were analyzed by Bondar Clegg & Co. Ltd, Ottawa, Canada using standard techniques outlined in USEPA (1981). Concentrations of metals were determined by acid digestion followed by ICP-AES analysis (Multi-channel Jarrell-ASH AtomComp 1100) using the methods of McLaren (1981).

2.3 WHOLE-SEDIMENT TOXICITY TESTS

2.3.1 Culture and Conduction of Tests

Chronic toxicity tests with four species of benthic invertebrates, the amphipod, Hyalella azteca; the chironomid, Chironomus riparius; the mayfly, Hexagenia spp. and the oligochaete tubificid worm, Tubifex tubifex, were conducted with all sediments collected from the Athabasca River as well as a clean control sediment from Long Point, Lake Erie, for biological quality assurance.

Complete details of the culture of organisms and conditions of each toxicity test for *C. riparius* and *T. tubifex* are described elsewhere (Reynoldson *et al.* 1991; Day *et al.* 1994; Reynoldson *et al.* 1995).

Culture of *H. azteca* was conducted according to the procedure described in Borgmann *et al.* (1989). Eggs of the mayfly *Hexagenia* spp. (both *H. limbata* and *H. rigida*) were collected during late June and July in 1991 according to the method of Hanes and Ciborowski (1992) and organisms were cultured using the procedure of Bedard et al. (1992). Tests with *H. azteca, C. riparius* and *T. tubifex* were conducted in replicates of 5 (true field replicates) in 250 mL glass beakers containing 60 to 100 mL of sediment with approximately 100 to 140 mL of overlying water (City of Burlington dechlorinated tap water). Tests with the mayfly, *Hexagenia* were conducted in replicates of 5 in 1 L glass jars with 150 mL of test sediment and 850 mL overlying water. The sediment was allowed to settle for 24 h prior to addition of the test organisms. Tests were initiated with the random addition of 15 organisms per beaker for *H. azteca* and *C. riparius*, 10 organisms per jar for *Hexagenia* spp. and 4 organisms per beaker for *T. tubifex*. Juveniles of *H. azteca* were 3 to 7 d old at test initiation; *C. riparius* larvae were first instars and were approximately 3 d postoviposition; *Hexagenia* nymphs were 1.5 to 2 months old (approximately 5 to 10 mg wet weight) and *T. tubifex* adults were 8-9 weeks old.

The organisms were fed during the course of exposure e.g., 8 mg of moistened Nutrafin^R fish food flakes was added twice weekly to beakers containing the midge, *C. riparius* and the amphipod, *H. azteca. Hexagenia* were fed 0.5 mL of YCT (yeast:cerophyll;trout chow) twice weekly and 8 mg Nutrafin^R was mixed in with each sediment in each container at the beginning of the bioassays with *T. tubifex*.

Tests were conducted at $23\pm1^{\circ}$ C with a 16L:8D photoperiod. Tests were static with the periodic addition of distilled water to replace water lost during evaporation. Each beaker was covered with a plastic petri dish with a central hole for aeration using a Pasteur pipette and air line. Dissolved oxygen concentrations and pH were measured at the beginning, middle and end of each exposure period. Tests were terminated after 10 d for *C. riparius*, 21 d for *Hexagenia* and 28 d for *H. azteca* and *T. tubifex* by passing the sediment samples through a 500 μ m mesh sieve. Sediment from the *T. tubifex* test was passed through an additional 250 μ m mesh sieve at test completion to obtain cocoons and young worms. Endpoints measured in the tests were: *H. azteca*, survival and growth (increase in mg dry wt/ind.); *C. riparius*, survival and growth (increase in mg dry wt/ind.), *Hexagenia* spp., survival and growth (increase in mg dry wt./ind. from day 0 to 21-d); and *T. tubifex*, survival and production of coccoons and live young. Mean dry weights of *H. azteca*, *C. riparius* and *Hexagenia* spp.were determined after drying the surviving animals from each replicate as a group to a constant weight in a drying oven (60 °C). Initial weight of *H. azteca* and *C. riparius* was considered to be zero. Initial weight of *Hexagenia* was determined from a subset of animals.

2.3.2 Statistical Analysis of Toxicity Data

The data for each species and measured endpoint were tested for normality and a statistical comparison of the responses of each species in sediment for a given site was conducted using analysis of variance (ANOVA). If significant effects were found and these data passed the tests for normality and homogeneity, comparison of means was performed using the Student-Newman-Keuls pairwise multiple comparison. All comparisons for toxicity were related to the results for the two upstream control sites, ARC and ARC2. All statistical analyses were performed using the microcomputer software package, SigmastatTM (Jandel, California) and significance is at a level of $P \le 0.05$ unless otherwise noted.

Responses in sediments were also compared to acceptability levels of survival, growth and/or production of young for the same four species used by Day *et al.* (1995) and obtained from a range of reference sediments (258 stations) in the Great Lakes with large differences in grain size and organic carbon. These acceptability criteria were set at the 5th percentile on the normal distribution curve for the range in responses for each endpoint and species in 258 reference sites and are as follows: *C. riparius* - % survival \geq 68.0, growth \geq 0.22 mg dry wt/ind.; *H. azteca* - % survival \geq 74.7, growth \geq 0.22 mg dry wt/ind.; *Hexagenia* spp. - % survival \geq 84.0, growth mg dry wt/ind. \leq 0.50; *T. tubifex* - \geq 31 cocoons, \geq 35 total young (Table 2).

2.4 PROCESSING OF SAMPLES FOR INVERTEBRATE COMMUNITY ANALYSIS

2.4.1 Sample Processing

The processing of samples collected for benthic invertebrate community analysis generally followed the procedures described by Alberta Environment (1990). Samples were first prepared by removing the formalin in which they were stored and rinsing them through a series of sieves. Mesh sizes of 1 mm, 229 μ m and 74 μ m were used. Organic material was separated from inorganic material (sand) in the two finest fractions by elutriation. The three fractions obtained were stored in 80% ethanol. The finest fraction (< 74 μ m) was kept but not sorted.

The coarse fraction (> 1 mm) of each sample was sorted in its entirety under a dissecting scope at a magnification of at least 7X. Because these depositional samples contained a large amount of organic debris, it was decided that the fine fraction would be subsampled according to the method of Wrona *et al.* (1982; cited in Saffran, 1994). Subsampling was standardized to at least one quarter (five 50 mL subsamples) of the fine fraction. Subsample counts were often low (< 100 organisms) but it was felt that the time taken to sort the entire fraction would have been excessive for the amount of data obtained in these cases.

The cone subsampler (Wrona *et al.* (1982; cited in Saffran, 1994) was also used to facilitate identification of Oligochaeta. When the number of worms was greater than 400, one quarter was removed for identification. The proportions of different families were then applied to the total number of oligochaetes in the sample, which had previously been counted.

2.4.2 Quality assurance/quality control

Sample cleaning, fractioning and subsampling were consistantly performed by one person who also supervised the sorting process. Four spring samples and four fall samples (10.7% of the total samples) were chosen at random to verify sorting efficiency. Re-sorting was undertaken by an individual not involved in the original sorting. A recovery of 95%, as suggested in Environment Canada (1993) was considered to be the minimum acceptable standard. The results of the QA/QC analysis are presented in Saffran et al. (1994).

2.4.3 Biotic indices used in data comparisons

Diversity indices are mathematical expressions which use three components of community structure; namely, richness (number of species present), eveness (uniformity in the distribution of individuals among the species) and abundance (total number of organisms present), to describe the response of a community to the quality of its environment (Metcalfe 1989). In this study, three of the most widely used measures of diversity were used to describe the data collected from the Athabasca River. These three indices are as follows:

Shannon-Wiener	$d = -\sum Ni / N \log 2 Ni / N$
----------------	----------------------------------

Simpson $d = 1 - \left[N_i(N_i-1)/N(N-1) \right]$

Margalef $d = (S - 1)/\log_e N$

Where: d = diversity

N = total number of individuals of all species collected

 N_i = number of individuals belonging to the ith species

S = number of species

3.0 RESULTS

3.1 TOXICITY TESTS

The results of the bioassays are summarized in Figures 1 to 4 and in Appendix A. In general, there was no indication of toxicity at any of the sites for three of four species and for seven of eight chronic endpoints. For example, survival of *C. riparius, Hexagenia* and *H. azteca* in sediments collected downstream of Hinton was equal to or greater than survival of these species in

sediments collected either upstream of Hinton (control sites), in the reference Lake Erie sediment used for QA/QC or the mean survival measured in 258 uncontaminated sediments from the nearshore areas of the Great Lakes (dotted line). Growth of all three species was similarly not reduced at any of the sites.

Only one toxicity test endpoint was lowered by exposure to sediments collected downstream of Hinton. Production of live young by the tubificid worm, *T. tubifex*, was significantly reduced in sediments collected above the Berland River (BR) and near Windfall Bridge (WB). Production of cocoons was not similary affected.

3.2 BENTHIC INVERTEBRATE COMMUNITY STRUCTURE

Tables 3 and 4 (and Appendix B) list the taxonomical groupings and species (where identified) of benthic invertebrates identified at all sites. Tables 3 and 4 also present the percent abundance of the total number of organisms counted in five replicate samples collected. The spring samples generally contained fewer organisms than the fall samples (see report by Saffran *et al.* 1994 in Appendix B).

Chironomids, especially the Chironomini and Orthocladiinae groups, dominated the spring samples collected at the control (ARC) site in terms of percent abundance. Chironomids were also very abundant in sediments collected downstream at Obed Bridge (OB). Samples collected at other downstream sites contained large percentages of tubificid worms (e.g., see data for Blue Ridge (BER), upstream of the Berland River (BR), and Windfall Bridge (WB)). The site upstream of the Emerson Lakes (ATHA) which was only sampled in the spring had a large percentage of ostracods.

The three diversity indices calculated for the spring samples (Fig. 5; Appendix A) were all relatively low (including the control samples). The greatest diversity in the spring was found at the Obed Bridge site (OB) and the site upstream of Emerson Lakes (ATHA).

Chironomids also dominated the two control sites sampled in the fall (i.e., ARC and ARC2). A high percentage of tubificid worms was found at most other sites located downstream of Hinton with the exception of the site at Weldwood Haul bridge which had a mixture of cladocerans, copepods and ostracods as well as several groups of chironomids (Appendix B). Diversity indices calculated for these sites indicated again that diversity at all sites was low (Fig.6). Diversity was the highest at the Weldwood Bridge site (HB). The lowest diversity was found at the Windfall Bridge site (WB) and this low diversity was statistically lower than all other sites, especially the control sites, for 2/3 indices (Shannon-Wiener and Simpson's).

3.3 PHYSICAL-CHEMICAL ANALYSES

The major nutrients (total N and P) and several metals of concern as well as the percent total organic carbon (TOC), loss on ignition (LOI), silt, sand and clay, are presented in Table 5.

Concentrations of metals at most sites were below the Low Effects Concentrations (LEL) set by the Province of Ontario (Persaud *et al.* 1992) for freshwater sediments with the exception of arsenic at BR and WB, chromium at all sites, cadmium at EL and nickel and copper at WB. WB and BR are the two sites where reproduction was lower in *T. tubifex* and diversity of the benthos was reduced (Windfall Bridge, only).

Examination of the physical parameters for sediments collected from all sites (percent sand, silt and clay) indicated that the sediments varied considerably in their particle size distribution with the Berland River site having the highest percentage of sand (74.3%). The percentage of clay was highest at the Windfall Bridge site.

4.0 DISCUSSION

Sediments located downstream of Hinton in the upper Athabasca River do not appear to be particularly toxic to benthic invertebrates living in or near the benthos or to invertebrates exposed to such sediments after removal from these habitats. Survival and growth of three benthic invertebrates in culture in the laboratory and exposed to field-collected sediments, i.e., H. azteca, C. riparius and Hexagenia, were all high and above the acceptability criteria for other reference sites and in comparison to control sites. However, sediments collected from the Athabasca River near the Windfall Bridge (WB) and upstream of the Berland River (BR) indicated sublethal effects to tubificid worms (*i.e.*, statistically reduced reproduction in comparison to control sediments, ARC and ARC2). Production of young at one site (i.e., upstream of the Berland River) was also lower than the range of production of young noted for 258 clean sediments collected from depositional areas in the nearshore areas of the Laurentian Great Lakes. The three components of community structure, i.e., richness, abundance and evenness measured using diversity indices were also reduced at the Windfall Bridge site. Tubificid worms were prevalent at this site although taxonomic determination to the level of genus or species was not conducted and therefore it is unknown if the species, T. tubifex, was present. Diversity at all sites was low in general but this is considered normal for depositional areas in rivers and streams located in the prairie provinces (T.B. Reynoldson, per. comm.).

The Berland River site had a high percentage of sand which would not be nutritionally acceptable to tubificid worms in comparison to sediment(s) with a greater component of organic material. However, the protocol for the tubificid bioassay includes a food component designed to 'even' out the responses of the organism with regard to nutrition. The Berland River (BR) and the Windfall Bridge (WB) sites also had levels of arsenic, chromium, coper and nickel higher than the criterea set for the Province of Ontario for low effects to invertebrates and thus the presence of such contaminants or other unreported contaminants in these sediments could be causing toxicity and resulting in reduced diversity.

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5.0 CONCLUSIONS

Sediments located downstream of Hinton in the upper Athabasca River are not particularly toxic to benthic invertebrates living in and near the sediments with the exception of sediments upstream of the Berland River and near Windfall bridge where toxicity to tubificid worms (reduced reproduction) was noted and where species diversity (WB, only) was lower. A slight elevation of arsenic and chromium was noted at these two sites and further information on the presence/absence of other contaminants in these sediments may help further interpretation of the data with regard to what is causing the observed toxicity and lower diversity.

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TABLES

Table 1. Description and coding for depositional sites sampled on the Upper Athabasca River in May and September/October, 1993.

Location	Description	Site Label
Athabasca River u/s Hinton Control 1	200 meters above Maskuta Creek right side in open bay. No bar to main current. Silt with organic layers	ARC B - Spring B,C,T - Fall
Athabasca River u/s Hinton (Control 2)	In protected bay just downstream of ARC, more sand and less organic debris than ARC	ARC2 B,C,T - Fall
Athabasca River at Weldwood Haul Bridge	Approx. 1 km below Weldwood Haul Bridge right bank opposite Fish Ck. Small bay sand overlain with brown silt-clay 1 cm thick	HB B - Spring B,C,T - Fall
Athabasca River at Obed	Small rock bay 1 km. below Obed bridge right side. Near Baseline Ck. More silt than at previous sites. Organic debris 3 cm below surface.	OB B - Spring B,C,T - Fall
Athabasca River at Emerson Lakes	Left side approx. 300 meters below May sampling Location. Open bay with sandy beach	EL B - Spring B,C,T - Fall
Athasbasca River upstream of EL	-	ATHA B - Spring
Athabasca River u/s of Berland River	Sampled in bay 250 meters below May sampling site. About 1.5 km above bridge. Shallow, fairly sandy, numberous emerging and in-place invertebrates	BR B - Spring B,C,T - Fall
Athabasca River at Windfall	Same site as sampled in May, 2 km below bridge, right side. Very good depositional zone. Fines in abundance. Numerous invertebrates.	WB B - Spring B,C,T - Fall
Blue Ridge	-	BER B - Spring

Table 2. MEAN VALUES FOR PERCENT SURVIVAL, INCREASE IN BIOMASS (MG DRYWEIGHT PER INDIVIDUAL) OR REPRODUCTION OF FOUR SPECIES OF BENTHIC INVERTEBRATES IN SEDIMENT COLLECTED FROM 258 REFERENCE SITES IN NEARSHORE AREAS OF THE LAURENTIAN GREAT LAKES. TOXICITY IS INDICATED WHEN VALUED ARE BELOW THE LEVEL OF THE 5TH PERCENTILE OF RESPONSES FOR EACH SPECIES.

Species (Endpoint)	Mean Values (±S.D.) for Reference Sites in the Great Lakes	Value at the 5th percentile on the normal distribution curve below which toxicity is indicated
Chironomus riparius % Survival Growth ¹	84.3 (11.0) 0.34 (0.07)	68 % 0.22
Hyalella azteca % Survival Growth ¹	88.6(7.1) 0.49 (0.14)	74.7 % 0.22
Hexagenia spp. % Survival Growth ² Unfed ³	96.0 (5.4) 4.81 (4.46)	84 % 0.38
Fed ⁴ Tubifex tubifex Cocoons	2.86 (1.06)	0.58
Unfed ⁵ Fed ⁶	34 (7) 38 (5)	24 31
Total Young Unfed ⁵ Fed ⁶	64 (30) 113 (36)	21 35

¹Biomass measured as mg dry weight/individual at test termination

²Change in biomass measured as mg dry weight/individual from test intiation to test termination

³Based on unpublished data from 50 reference sites in the nearshore areas of the Great Lakes where mayfly nymphs were not fed a supplemental diet ⁴Based on unpublished data from 208 reference sites in the nearshore areas of the

Great Lakes where mayfly nymphs were fed a supplemntal diet of 0.5 mL yeast-Cerophyll-trout chow 3X weekly

⁵Based on unpublished data from 50 reference sites in the nearshore areas of the Great Lakes were adult tubificids were not fed a supplemental diet ⁶Based on unpublished data from 208 reference sites in the nearshore areas of the Great Lakes where adult tubificids were fed a supplemental diet of 80 mg NutrafinTM at test initiation

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Table 3. CONTRIBUTION OF TAXA TO TOTAL ABUNDANCE FOR SAMPLES COLLECTEDIN MAY 1993 FROM EIGHT SITES ON THE UPPER ATHABASCA RIVER..

			% AB	UNDANCE OF	TOTAL NUM	BERS		
SPECIES/SITE	ARC	ATHA	НВ	OB	EL	BR	BER	WB
ANNELIDA OLIGOCHAETA HAPLOTAXIDA Naididae Tubificidae	7.8	0.3 2.4	47.1	3.2	0.04 57.9	90.5	0.2 57.4	81.9
ARTHROPODA ARACHNOIDA ACARI CRUSTACEA CLADOCERA COPEPODA OSTRACODA INSECTA DIPTERA Ceratopogonidae Chironomidae	2.0	1.7 12.9 65.6	0.4 5.3 3.2 3.4	5.5 0.1 2.8 0.3	0.2 0.2 0.5	0.5 0.5 0.3		0.1
Chironomini Tanytarasini Orthocladiinae Diamesinae Tanypodinae Chironomid adult Chironomid pupae Empididae Chelifera Hemerodromia	67.2 2.7 11.3 7.2 0.7 0.3	7.8 4.8 1.3 0.1	19.1 0.4 4.4 1.8 7.4 5.3	43.8 2.6 3.4 35.6 1.2 0.03 1.5 0.03 0.06	17.8 0.3 0.7 21.9	5.9 0.5 0.3 1.2 0.2	21.9 0.1 12.5 4.3	9.1 0.3 0.1 7.5 0.1
EPHEMEROPTERA Ephemerellidae <i>Serratella</i> Baetidae <i>Baetis</i> HEMIPTERA TRICHOPTERA MEGALOPTERA			0.2	0.09 0.1	0.04		0.1	0.1
MOLLUSCA GASTROPODA PULMONATA Lymnaeidae Planorbidae		0.2				0.02		
PELECYPODA Sphaeriidae Pisidium	0.7	0.4	0.2 0.1		0.04	0.06		0.1
NEMATODA		2.1	1.3		0.1	0.2	2.6	
TARDIGRADA			0.2					

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Table 4. CONTRIBUTION OF TAXA TO TOTAL ABUNDANCE FOR SAMPLESCOLLECTED IN SEPTEMBER 1993 FROM SEVEN SITES ON THE UPPERATHABASCA RIVER.

			% ABUNDAI	NCE OF TOTAL	L NUMBERS		
SPECIES/SITE	ARC	ARC2	HB	OB	EL	BR	WB
ANNELIDA							1
OLIGOCHAETA							
HAPLOTAXIDA							
Naididae	0.8	0.6	1.0	1.1	1.3	1.1	0.03
Tubificidae	2.1	13.9	24.7	66.3	65.6	58.0	84.2
ARTHROPODA							
ARACHNOIDA							
ACARI	0.3	0.6	0.1	0.1	0.4	0.06	
CRUSTACEA							
CLADOCERA	15.4	17.6	9.1	9.9	10.6	4.9	3.2
COPEPODA	5.0	0.8	12.2	0.5	0.2	5.2	0.2
OSTRACODA	10.4	5.5	13.7	1.0	0.7	0.4	1.1
INSECTA							
DIPTERA		1		1			
Ceratopogonidae	0.04	1		0.01			0.02
Chironomidae							
Chironomini	12.1	14.6	8.9	8.2	10.3	22.6	5.8
Tanytarasini	34.8	33.9	7.8	3.4	0.8	5.8	0.7
Orthocladiinae	6.1	5.9	8.7	1.8	0.8	0.3	
Diamesinae	10.4	5.4	6.8	6.0	7.9	0.7	0.1
Tanypodinae	1.1	0.4	0.2	0.6	1.0	0.1	3.9
Chironomid adult	0.7		4.1	0.0	1.0	0.1	
Chironomid pupae	0.7	0.3	1			0.01	
Empididae	0.09	0.5			0.03	0.01	
Chelifera	0.3			0.03	0.05		
Hemerodromia	0.5			0.05			
EPHEMEROPTERA							
Ephemerellidae				0.06	0.08	0.04	
Serratella					0.00	0.01	
Baetidae							0.2
Baetis	0.07		0.1				0.2
HEMIPTERA				0.01			0.01
TRICHOPTERA				0.03			0.01
MEGALOPTERA				0.05	0.01		0.01
					0.01		
MOLLUSCA							
GASTROPODA						ļ	
PULMONATA	0.04	[0.00		0.00	
Lymnaeidae	0.04		0.1	0.02	0.01	0.03	0.01
Planorbidae			0.1				
Physidae				0.01			0.2
PELECYPODA					1		
Sphaeriidae				0.7		0.4	0.7
Pisidium			0.7		0.1		
NEMATODA	1.1	0.6	1.8	0.3	0.1	0.3	0.2

CHEMICAL PARAMETERS FOR SEDIMENTS COLLECTED FROM THE ATHABASCA RIVER. Table 5.

EL-T 633 BR-T 779 WB-T 706		PPM	Pb PPM	Zn PPM	Ni PPM	PPM PPM	Cr PPM	As PPM	70C	% 10	% Sand	% Silt	% Clav
	1220	12	20	45	15	1.7	36	\$	1.16	22.6	23.3	51.7	25
	940	13	30	69	15	0.3	35	11	0.46	19.7	74.3	22.8	2.9
	1510	16	24	59	21	< 0.2	43	00	1.51	22.2	3.5	68.5	28.1
	852	13	20	47	15	< 0.2	36	< 5 2	0.73	24.8	58.7	34.5	6.8
	985	12	24	50	15	0.2	35	<5	0.72	23.0	18.7	64.7	16.5
	938	14	22	49	17	0.4	36	< V S	1.47	23.4	64.9	28.5	6.6
	1010	13	21	54	17	< 0.2	35	\$	1.37	21.6	62.9	34.9	2.2
	550	16	31	120	16	0.6	26	ور م	001	,			
SEL 2000	4800	110	250	820	75	10	110	33	10	ı	1	1	

1 Province of Ontario's sediment quality criteria for metals; LEL (lowest observed effects level) and SEL (severe effects level)

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APPENDIX A: RAW DATA AND DIVERSITY INDICES

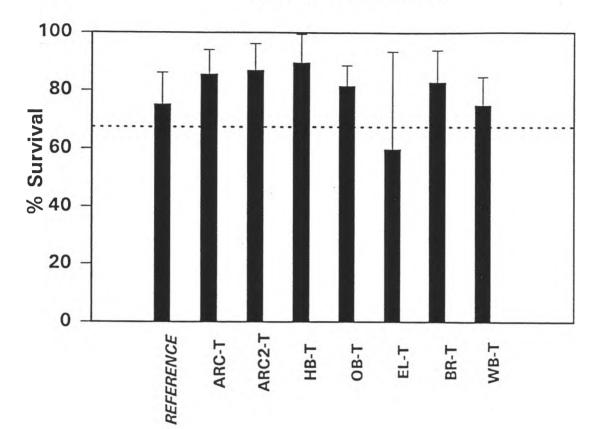
RAW DATA FROM THE UPPER ATHABASCA RIVER:

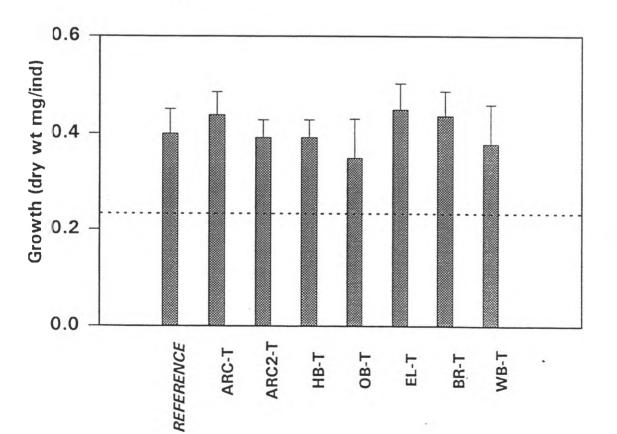
	SITE	SPECIES		SPECIES		SPECIES		SPECIES		
Athabasca	EL-T	CHSUR 66.67 93.33	0.5 0.371	100	5.863 5.42	100 100	0.725	4:	9 3	139 128
		93.33		100	5.786			4		142
		93.33		100				49		163
		66.67	0.446	100	5.210	100	0.227	5.	2	85
		59.5(33.9)	0.45(0.05)	98(4.5)	5.48(0.32)	96(5.9)	0.48(0.18)	48(3)	131(29)	
	BR-T	86.67						40		3
		100		100		100	0.4	50		2
		73.33			5.759			49		5 3
		73.33			2.95 5.037			41		3 20
		80	0.393	100	5.037	00.07	0.510	40	2	20
		82.7(11.2)	0.44(0.05)	100 (0)	5.12(1.30)	94.7(5.6)	0.52(0.19)	47(4)	7(8)	
	WB-T	60	0.394	100		93.33		40	6	32
		86.67		100				41		36
		73.33						3		1
		73.33						50		91
		80	0.488	100	6.165	93.33	0.419	4	/	25
		74.7(9.9)	0.38(0.08)	100(0)	7.18(0.87)	87.9(7.3)	0.75(0.22)	45(6)	37(33)	
	LONG POINT	80				100				123
		73.33						4		95
		73.33						41		75
		86.67						4:		83 92
		60	0.400	100	0.430	93.33	0.004	4	5	92
		75(11.)	0.40(0.05)	100(0)	5.23(0.74)	95.9(4.7)	0.48(0.10)	44(3)	113(26)	
	ARC-T	86.67	0.402	90	3.942	100	0.286	4	2	104
		80						4		149
		100					0.526	3	3	135
		80								181
		80	0.462	100	4.333	93,33	0.448	4	4	135
		85.3(8.7)	0.44(0.05)	92(4.5)	4.30(0.29)	98.7(2.9)	0.47(0.11)	42(6)	113(26)	
	ARC2-T	100	0.331	100	5.13	100	0.392	4	4	139
		86.67		100				4	4	79
		86.67			4.741			4		136
		73.33						4		152
		86.67	0.405	90	4.165	100	0.425	4	7	137
		86.6(9.4)	0.39(0.04)	96(5.5)	4.50(0.49)	94.7(8.7)	0.43(0.02)	45(1)	129(29)	
	HB-T	100		100				4	1	92
		86.67						4		142
		00	0.368	400	3.763	73.33	0.685	4		131
		80					0 5 (0	4		147
		80	0.379	100	3.448	100		4		142
			0.379	100	3.448	100		4 4		159
		80 100	0.379	100 100	3.448 3.75	100 100	0.309	4		
	OB-T	80 100 89.3(10.1) 86.67	0.379 0.38 0.392(0.04) 0.291	100 100 98(4.5) 100	3.448 3.75 3.73(0.17) 4.079	100 100 90.7(11.2) 100	0.309 0.47(0.15) 0.309	4 44(2) - 4	5 133(25) 3	159 137
	OB-T	80 100 89.3(10.1) 86.67 86.67	0.379 0.38 0.392(0.04) 0.291 0.364	100 100) 98(4.5) 100 100	3.448 3.75 3.73(0.17) 4.079 4.473	100 100 90.7(11.2) 100 93.33	0.309 0.47(0.15) 0.309 0.546	4 44(2) - 4 3	5 133(25) 3 7	159 137 81
	OB-T	80 100 89.3(10.1) 86.67 86.67 86.67	0.379 0.38 0.392(0.04) 0.291 0.364 0.422	100 100) 98(4.5) 100 100 100	3.448 3.75 3.73(0.17) 4.079 4.473 2.195	100 100 90.7(11.2) 100 93.33 100	0.309 0.47(0.15) 0.309 0.546 0.628	4 44(2) - 4 3 4	5 133(25) 3 7 3	159 137 81 88
	OB-T	80 100 89.3(10.1) 86.67 86.67 86.67 73.33	0.379 0.38 0.392(0.04) 0.291 0.364 0.422 0.43	100 100) 98(4.5) 100 100 100 100	3.448 3.75 3.73(0.17) 4.079 4.473 2.195 3.4	100 100 90.7(11.2) 100 93.33 100 100	0.309 0.47(0.15) 0.309 0.546 0.628 0.469	4 44(2) - 4 3 4 3	5 133(25) 3 7 3 2	159 137 81 88 155
	OB-T	80 100 89.3(10.1) 86.67 86.67 86.67	0.379 0.38 0.392(0.04) 0.291 0.364 0.422 0.43	100 100) 98(4.5) 100 100 100 100	3.448 3.75 3.73(0.17) 4.079 4.473 2.195 3.4	100 100 90.7(11.2) 100 93.33 100 100	0.309 0.47(0.15) 0.309 0.546 0.628 0.469	4 44(2) - 4 3 4 3	5 133(25) 3 7 3	159 137 81 88

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ATHABASCA RIVER - 1993

Chironomus riparius





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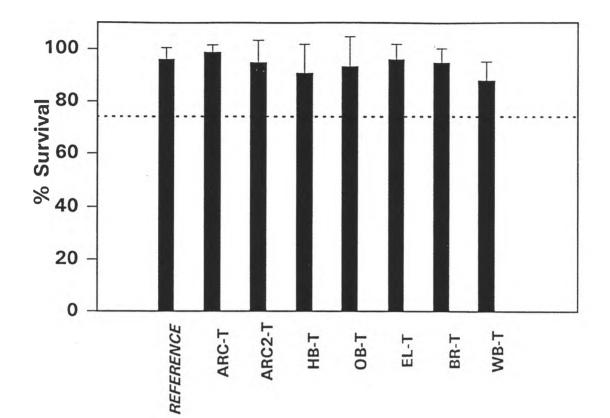
Fig. 1

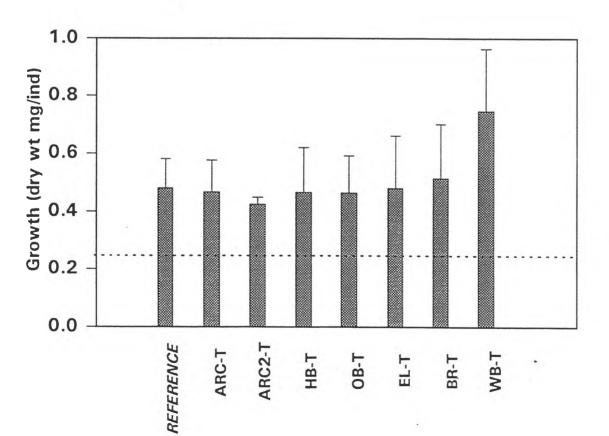
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ATHABASCA RIVER - 1993

Fig. 2

Hyalella azteca



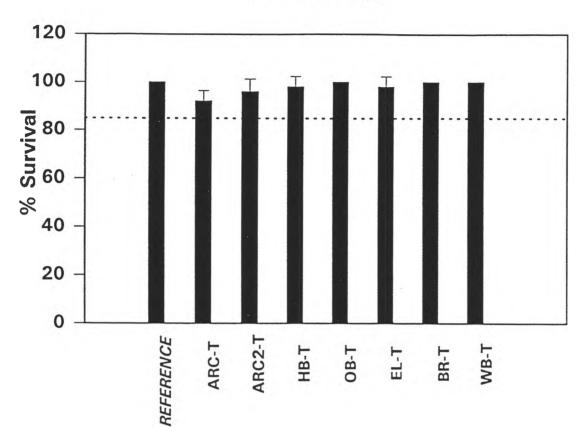


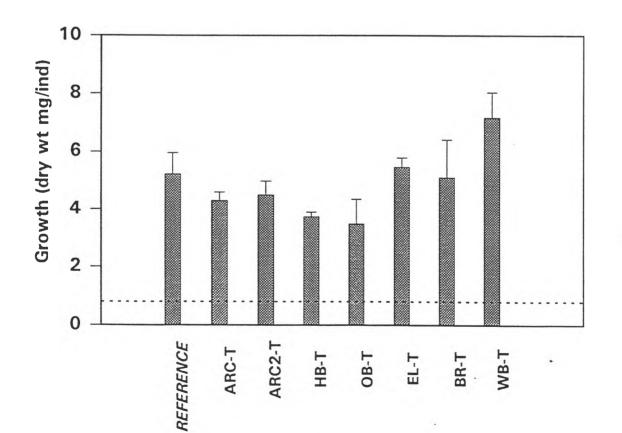
ATHABASCA RIVER - 1993

Fig. 3

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Hexagenia spp.

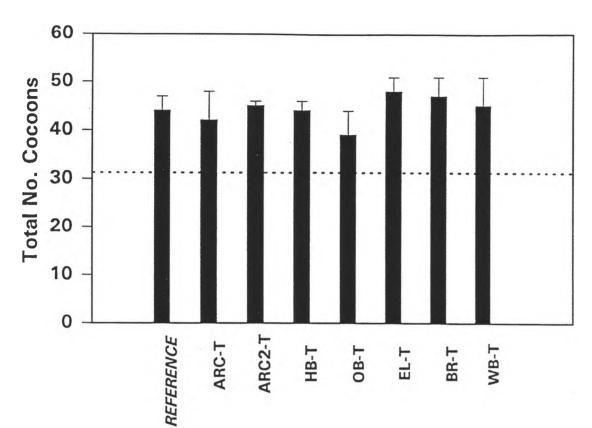


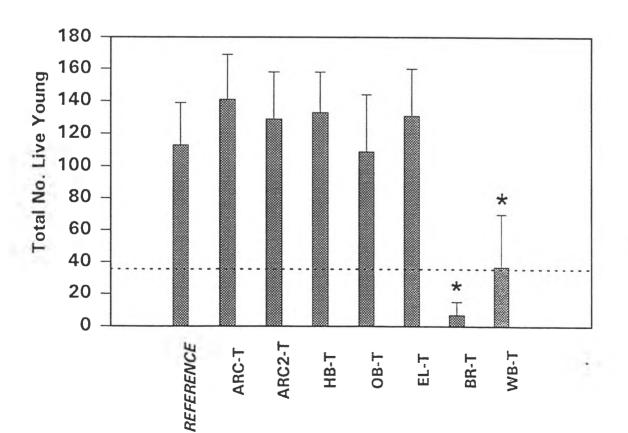


ATHABASCA RIVER - 1993



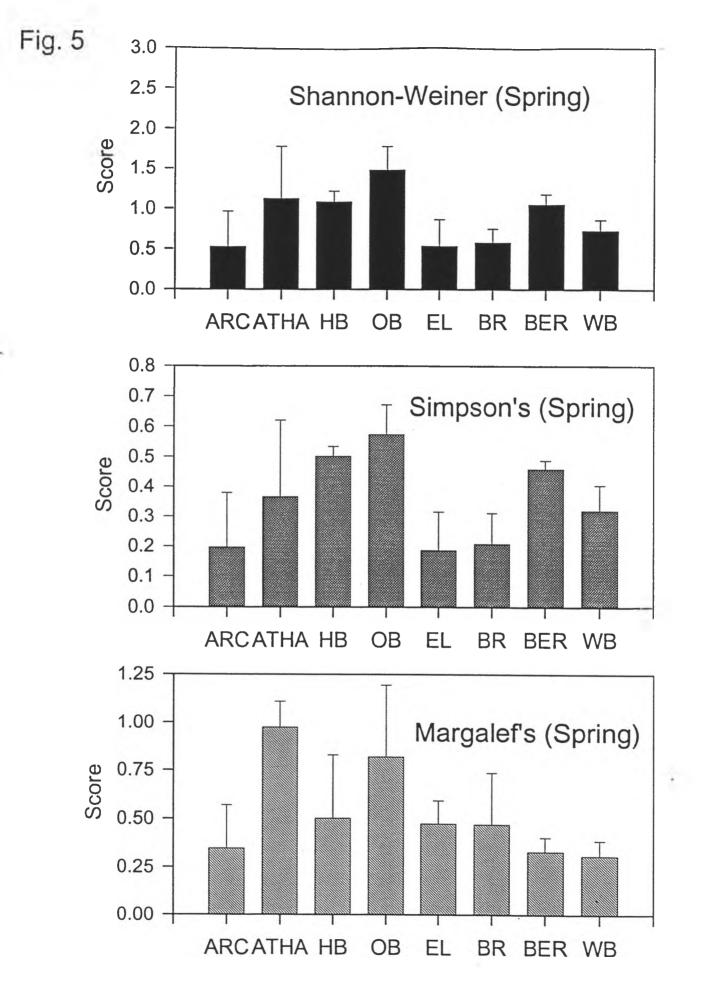
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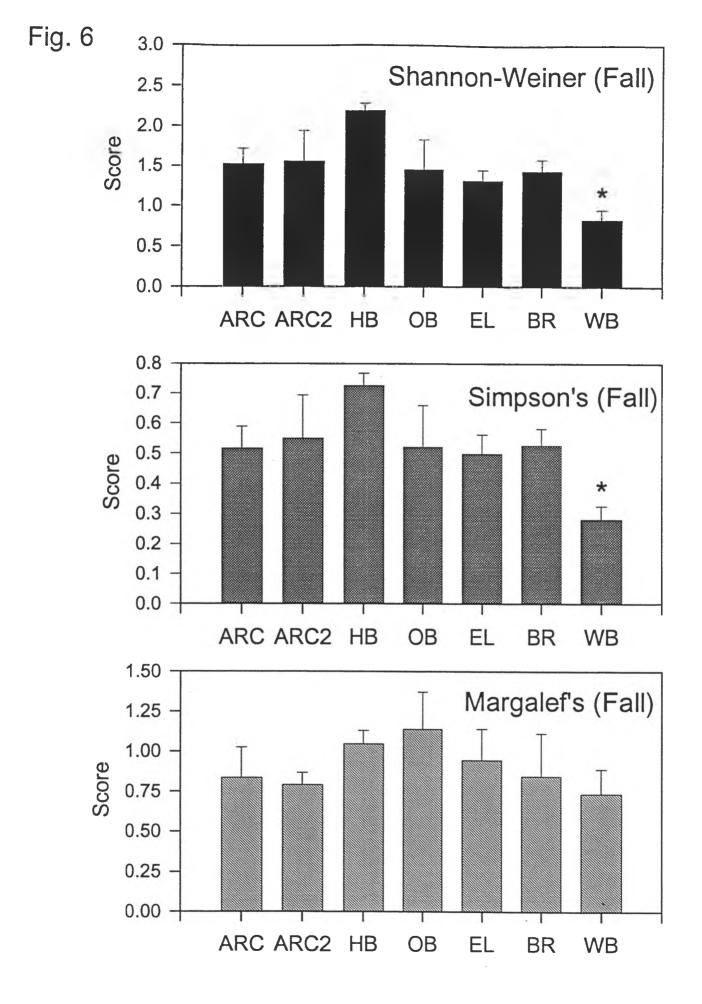
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APPENDIX B: SUMMARIES OF TOTAL INVERTEBRATE COUNTS FROM SAFFRAN

For ease of reference this appendix contains Tables 2 and 3 extracted from a previous report, namely;

Saffran, K. 1995. Northern River Basins Study Project Report No. 50, Aquatic Macroinvertebrate Identifications, Athabasca River, May and September, 1993. Northern River Basins Study, Edmonton, Alberta

AREL	93/05/05 Sorters*: P.H., D.P., K	.S.				
		1	2	3	4	5
ANNELIDA						
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae			1		
	Tubificidae	102	179	177	372	573
ARTHROPO	DA					
	ARACHNOIDA					
	ACARI		2			4
	CRUSTACEA					
	CLADOCERA		4			
	OSTRACODA	4	7			
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	31	114	54	106	126
	Tanytarsini		4			3
	Orthocladiinae		7		4	6
	Diamesinae	66	41	70	186	166
	Tanypodinae					1
	Chironomid Pupae		2	4		
	EPHEMEROPTERA					
	Ephemerellidae					
	Serratella					1
MOLLUSCA						
	PELECYPODA					
	Sphaeriidae					
	Pisidium	1				
NEMATODA	A	1	1			1
	TOTAL	205	361	306	668	881

Table 2.Summary of total invertebrate counts from eight sites on the upperAthabasca River, May 1993.

* Sorters: P.H. = Paul Hvengaard, D.L. = Darcy Lightle, D.P. = Dee Patriquin, K.S. = Karen Saffran, and N.W. = Nancy Westworth

	93/05/05 Sorters: D.P., K.S	1	2	3	4	5
ANNELIDA						
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Tubificidae	114	124	48	52	4
ARTHROPOI	DA					
	ARACHNOIDA					
	ACARI	2				
	CRUSTACEA					
	CLADOCERA					4
	COPEPODA	10	12			
	OSTRACODA			8	12	
1	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	55	12	24	33	3
	Tanytarsini	1			1	
	Orthocladiinae	25	2		9	
	Diamesinae		8		6	
	Tanypodinae	3	6	6		4
	Chironomid Pupae	10	2	5	3	2
	EPHEMEROPTERA					
	Baetidae					
	Baetis	2				
	PLECOPTERA (small)	2				
MOLLUSCA						
]	PELECYPODA					
	Sphaeriidae					
	Pisidium			1		
NEMATODA		2			8	
TARDIGRAD		2		e -		~ •
	TOTAL	228	166	92	124	203

ARATHAB 93/05/07 Sorters: D.	P., K.S.					
		1	2	3	4	5
ANNELIDA						
OLIGOCHAETA						
HAPLOTAXIDA						
Naididae			4		3	
Tubificidae		14	13	6	5	12
ARTHROPODA						
CRUSTACEA						
CLADOCERA		12		4	16	4
COPEPODA		40	176	4	20	32
OSTRACODA		240	192	256	2800	412
INSECTA						
DIPTERA						
Ceratopogo	nidae		1			
Chironomid	lae					
Chirono	mini	14	119	6	9	17
Tanytar	sini	31	17		34	18
Orthocla	adiinae	3	8	5	8	4
Diamesi	nae		4			
Tanypoo	dinae	1		1		
Chirono	mid Pupae		1	1		
MOLLUSCA						
GASTROPODA						
PULMONATA						
Lymnaeidae	۱ •	1		1	1	1
PELECYPODA						
Sphaeriidae						
Pisidium	1		5	1	1	1
NEMATODA		8	16		16	4
T	OTAL	364	556	285	2913	505

ARC	93/05/05	Sorters: P.H., D.P., K.S	•				
			1	2	3	4	5
ANNELID	A						
	OLIGOCH	AETA					
	HAPLO	DTAXIDA			-		
]	Fubificidae		10	4	5	4
ARTHRO	PODA						
	ARACHNO	DIDA					
	ACARI	[5		1	
	INSECTA						
	DIPTE	RA					
	(Chironomidae					
		Chironomini	18	26	35	74	44
		Tanytarsini	8				
		Orthocladiinae	9			15	9
		Diamesinae	4	8		8	1
		Chironomid Adult	1			1	
	E	Empididae					
		Chelifera			1		
	PLECC	PTERA (small)			2		
		TOTAL	40	49	42	104	58

OB	93/05/05 Sorters: D.P., K.S.					
		1	2	3	4	5
ANNEL	IDA					
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Tubificidae	18	5	4	41	29
ARTHR	OPODA					
	ARACHNOIDA					
	ACARI	2	12		152	
	CRUSTACEA					
	CLADOCERA				4	
	COPEPODA		72	4	8	
	INSECTA					
	DIPTERA					
	Ceratopogonidae				1	
	Chironomidae					
	Chironomini	46	417	207	262	388
	Tanytarsini	1	23	26	12	16
	Orthocladiinae	3	21	44	27	8
	Diamesinae	102	268	119	349	234
	Tanypodinae	1	5	1	24	2
	Chironomid Pupae		1			
	Chironomid Adult	1	13		16	14
	Empididae					
	Chelifera				1	
	Hemerodromia	1			1	
	EPHEMEROPTERA					
	Ephemerellidae					
	Serratella		1			2
NEMAT						4
	TOTAL	175	838	405	898	697

ARBR	93/05/06 Sorters: D.P, K.S.					
		1	2	3	4	
ANNELID	A					
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Tubificidae	653	1664	426	1316	324
ARTHROP	ODA					
	CRUSTACEA					
	CLADOCERA		8		16	
	COPEPODA	8		12	4	
	OSTRACODA	8			4	
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	39	90	47	49	61
	Tanytarsini		15			7
	Orthocladiinae	6	1	6	1	1
	Diamesinae		13	1	8	35
	Tanypodinae	1		1	5	2
MOLLUSC	CA					
	GASTROPODA					
	PULMONATA					
	Lymnaeidae	1				
	PELECYPODA					
	Sphaeriidae					
	Pisidium			2	1	
NEMATOI	DA				8	
	TOTAL	716	1791	495	1412	430

ARBER	93/05/06	Sorters: D.P., K.S.					
			1	2	3	4	5
ANNELIDA	1						
	OLIGOC	HAETA					
	HAPI	LOTAXIDA					
		Enchytraeidae					1
		Naididae	1				
		Tubificidae	145	209	137	47	154
ARTHROP	DDA						
INSECTA							
DIPTER	A						
		Chironomidae					
		Chironomini	41	59	54	52	58
		Tanytarsini		1			
		Orthocladiinae	31	11	19	47	43
		Diamesinae	10	15	19	1	7
		Chironomid Pupae	3	1	2		5
	EPHE	EMEROPTERA					
		Ephemerellidae	1				
NEMATOD	A			8	8		16
		TOTAL	232	304	239	147	284

ARW	93/05/06 Sorters: D.P., K.S.			-		
		1	2	3	4	5
ANNELII	DA					
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Tubificidae	464	733	231	185	143
ARTHRO	PODA					
	ARACHNOIDA					
	ACARI	1				1
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	48	49	27	42	28
	Tanytarsini		4		1	1
	Orthocladiinae	2	17		1	
	Diamesinae	28	55	17	43	18
	Tanypodinae		1	1		
	EPHEMEROPTERA					
	Ephemerellidae					
	Serratella					1
	MOLLUSCA					
	PELECYPODA					
	Sphaeriidae					
	Pisidium			1		
	TOTAL	543	859	277	272	192

ABR	93/09/17 Sorters: D.P., K.S.					
		1B	2B	3B	4B	5B
ANNELIDA						
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae	32	8	3	17	47
	Tubificidae	2004	2396	627	403	159
ARTHROPC	DDA					
	ARACHNOIDA					
	ACARI					6
	CRUSTACEA					
	CLADOCERA	108	244	20	48	48
	COPEPODA	164	120	64	57	100
	OSTRACODA		24	4		12
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	662	409	317	370	422
	Tanytarsini	64	54	51	219	173
	Orthocladiinae	1	7	4	8	9
	Diamesinae	11	12	15	11	20
	Tanypodinae	12				1
	Chironomid Pupae					1
	EPHEMEROPTERA (small)					4
MOLLUSCA	X					
	GASTROPODA					
	PULMONATA					
	Lymnaeidae			1	1	1
	PELECYPODA					
	Sphaeriidae					
	Pisidium	12	12	1	3	7
NEMATOD		8	16			4
	TOTAL	3078	3302	1107	1137	1014

Table 3.Summary of total invertebrate counts from eight sites on the upperAthabasca river, September 1993.

WB 93/09/17 Sorters: D.P., K.S.					
	B1	B2	B3	B4	B5
ANNELIDA					
OLIGOCHAETA					
HAPLOTAXIDA					
Naididae					4
Tubificidae	2936	2166	1242	2004	4188
ARTHROPODA					
CRUSTACEA					
CLADOCERA	336	48	8	8	76
COPEPODA	12	12			
OSTRACODA	40	28	21	17	52
INSECTA					
DIPTERA					
Ceratopogonidae	1			1	1
Chironomidae					
Chironomini	198	138	121	138	264
Tanytarsini	9	36	16	9	28
Diamesinae	17			1	à
Tanypodinae	3	88	153	123	215
Tabanidae					
Chrysops			1		
HEMIPTERA					
Corixidae		1	1		
TRICHOPTERA					
Limnephilidae					1
MOLLUSCA					
GASTROPODA					
PULMONATA					
Lymnaeidae			1		
PELECYPODA					
Sphaeriidae					
Pisidium	42	21	4	16	24
NEMATODA	1		4		20
TOTAL	3595	2538	1572	2317	4873

OB	93/09/17 Sorters: D.L., D.P., K	.S.				
		B1	B2	B3	B4	B5
ANNELIDA	L					
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae	48	31	36	65	43
	Tubificidae	364	6678	2179	1503	2972
ARTHROPO	DDA					
	ARACHNOIDA					
	ACARI	4	1	4	22	5
	CRUSTACEA					
	CLADOCERA	128	552	368	336	660
	COPEPODA	16	20	20	20	20
	OSTRACODA	68	20	32	60	24
	INSECTA					
	DIPTERA					
	Ceratopogonidae					1
	Chironomidae					
	Chironomini	255	317	354	340	418
	Tanytarsini	152	120	65	209	158
	Orthocladiinae	41	24	51	186	77
	Diamesinae	231	189	271	310	240
	Tanypodinae	13	15	6	57	25
	Empididae					
	Chelifera	4	4			
	EPHEMEROPTERA					
	Ephemerellidae				12	
	HEMIPTERA					
	Corixidae	1				1
	PLECOPTERA (small)	1				
	TRICHOPTERA					
	Brachycentridae					
	Brachycentrus	1	2	1		1

CONTINUED.../

OB	93/09/17	/CONTI	NUED				
			B 1	B2	B3	B4	B5
MOLLUSCA	X	· · · · · · · · · · · · · · · · · · ·					
	GASTROPOD	A					
	PULMONA	TA					
	Lym	naeidae			2	1	
	Phys	idae					
	F	°hysa	1				
	PELECYPODA	A Contraction of the second se					
	Spha	eriidae					
	P	isidium	23	47	10	29	29
NEMATODA	A		20	8	12	4	12
		TOTAL	1371	8028	3411	3154	4686

ARC 93/09/15 Sorters: D.L., D.P., K.S.	5.				
	<u>B1</u>	B2	B3	B4	B5
ANNELIDA					
OLIGOCHAETA					
HAPLOTAXIDA					
Naididae	1	20	2	6	16
Tubificidae	8	4	32	30	45
ARTHROPODA					
ARACHNOIDA					
ACARI		4	11	2	2
CRUSTACEA					
CLADOCERA	128	108	96	160	368
COPEPODA	56	36	40	52	96
OSTRACODA		84	44	164	288
INSECTA					
DIPTERA					
Ceratopogonidae				2	
Chironomidae					
Chironomini	89	172	166	135	111
Tanytarsini	238	384	264	342	714
Orthocladiinae	43	45	81	31	143
Diamesinae	60	124	137	147	110
Tanypodinae	2	23	4	11	21
Chironomid Pupae	1	2		1	1
Empididae	8				
EPHEMEROPTERA					
Baetidae					
Baetis	4				
MOLLUSCA					
GASTROPODA					
PULMONATA					
Lymnaeidae			1	1	
NEMATODA		21	30	12	
TOTAL	638	1027	908	1096	1915

ARC2	93/09/15 Sorters: D.L., D.P., K.	S.				
		B 1	B2	<u>B3</u>	B4	B5
ANNELIDA						
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae	9	4	6	8	6
	Tubificidae	268	261	22	22	251
ARTHROPC	DA					
	ARACHNOIDA					
	ACARI	9	4	4	4	12
	CRUSTACEA					
	CLADOCERA	272	192	120	204	252
	COPEPODA	16	16		4	12
	OSTRACODA	64	24	76	80	80
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	143	204	206	191	122
	Tanytarsini	338	221	515	658	274
	Orthocladiinae	62	73	37	101	77
	Diamesinae	123	69	27	41	57
	Tanypodinae	8	5		9	4
	Chironomid Pupae	5	3	3	3	2
	PLECOPTERA (small)		4			
NEMATODA	A	16		8		12
	TOTAL	1333	1080	1024	1325	1161

HB	93/09/15 Sorters: K.S., N.W.					
		B1	B2	B3	B4	B5
ANNELIDA						
(OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae	16	11	7	13	8
	Tubificidae	252	189	149	292	410
ARTHROPOD	DA					
4	ARACHNOIDA					
	ACARI				6	
(CRUSTACEA					
	CLADOCERA	36	44	108	56	232
	COPEPODA	36	40	256	204	104
	OSTRACODA	28	116	244	148	180
1	NSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	39	46	131	136	112
	Tanytarsini	20	36	136	113	102
	Orthocladiinae	24	45	185	90	109
	Diamesinae	23	42	133	67	90
	Tanypodinae	9	25	101	39	40
	Chironomid Adult					4
	HEMIPTERA					
	Corixidae			1		4
MOLLUSCA						
(GASTROPODA					
	PULMONATA					
	Lymnaeidae	4		1		
	Planorbidae	1				1
F	PELECYPODA					
	Sphaeriidae					
	Pisidium	9	6	10	9	1
NEMATODA		16	16	16	20	24
	TOTAL	513	616	1478	1193	1421

EL	93/09/15 Sorters: D.L, K.S., N	.W.				
		B1	B2	<u>B3</u>	B4	B5
ANNELID	A					
	OLIGOCHAETA					
	HAPLOTAXIDA					
	Naididae	76	70	26	4	3.
	Tubificidae	1653	2501	3532	1015	1740
ARTHROP	PODA					
	ARACHNOIDA					
	ACARI	15	13	13	14	13
	CRUSTACEA					
	CLADOCERA	480	600	348	64	190
	COPEPODA	4	8	12		
	OSTRACODA	17	24	44	12	12
	INSECTA					
	DIPTERA					
	Chironomidae					
	Chironomini	244	388	360	323	325
	Tanytarsini	13	23	20	32	46
	Orthocladiinae	20	18	26	33	29
	Diamesinae	210	290	280	227	251
	Tanypodinae	46	46	6	38	28
	Empididae					
	Chelifera				4	
	EPHEMEROPTERA					
	Ephemerellidae	4	4	4		
	MEGALOPTERA					
	Sialidae					
	Sialis	1				
	PLECOPTERA (small)	1				
MOLLUSC	A					
	GASTROPODA					
	PULMONATA					
	Lymnaeidae		1		1	
	PELECYPODA					
	Sphaeriidae					
	Pisidium	7	6	2	1	5
NEMATO	DA		12	4	4	
	TOTAL	2791	4004	4677	1772	2686

APPENDIX C: TERMS OF REFERENCE

NORTHERN RIVER BASINS STUDY

SCHEDULE A - TERMS OF REFERENCE

Project 2326-C1: Ecotoxicology of Depositional Sediments in the Upper Athabasca River

I. PROJECT BACKGROUND, RATIONALE AND RELEVANCE:

The assessment of water and sediment quality is essential to the execution of any proposed watershed management strategy. In addition, it is critical to determining the need for the success of any point and non-point source abatement programs. In any such strategy, it is also important to identify and protect desirable and self-sustaining aquatic communities but this is difficult using current techniques. Traditional environmental monitoring programs have emphasized the collection of biotic and abiotic samples for the analytical determination of residues of contaminants in a chemical-by-chemical approach and (more rarely) compared the biota of upstream sites (above a point-source) to downstream sites to determine the extent of declines in communities. Although this approach has proved valuable in establishing cause-and-effect in certain circumstances, it doesn't provide information on the types of biological communities which should be present if and when contamination is discontinued or removed.

Recent developments in multivariate statistical analyses have shown that the type of species assemblages (especially macroinvertebrate communities) at a reference site in a riverine or nearshore environment may be predicted using physical and chemical variables not affected by anthropogenic activities. This approach has potential in determining environmental health and in the setting of ecosystem objectives in a watershed. For example, a set of reference sites can be sampled within a watershed for their benthic community assemblages and selected chemical/physical variables measured. Community type at a potentially contaminated site with similar physical and chemical variables can then be predicted using a statistical model. A comparison between the existing community at a contaminated site and the predicted community can then be made to see if deterioration in community structure has occurred. This approach can also be used in determining whether remediation has allowed a return to the expected community.

Additional classification of reference and contaminated sites based on responses from toxicity tests conducted in the laboratory using samples from these sames sites as above can be added as a second set of guidelines for determining impairment. This approach has been shown to be successful in the development of sediment quality guidelines in the near-shore areas of the Great Lakes.

SCHEDULE A

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It is the intent of this research to develop biological guidelines for benthic biota in the northern river basins using predictive models based on community structure from a number of clean (reference) and contaminated sites and with sediment bioassays. The results from this study will answer a number of crucial Study Board Questions and advance science in the development of biological sediment criteria. It will also add to a National Programme aimed at developing National Sediment Guidelines.

II. REQUIREMENTS

The triad approach (use of benthic community structure, laboratory toxicity tests, and physicalchemical parameters incorporated into predictive multivariate statistical models) will be carried out in a 'feasibility' study in the fall of 1993. Depositional sediments from two sites upstream and five sites downstream from the bleached kraft mill effluent at Hinton will be sampled by the NRBS in mid-September. This project will provide information on the toxicity of the sediments to four species of benthic invertebrates using chronic toxicity tests conducted in the laboratory.

The following methodology is to be employed for chronic toxicity testing of the sediments:

- 1) Five field replicate samples of sediment were collected at the seven river sites. These were placed in plastic bags and held at 4° C before chronic toxicity testing.
- 2) Culture of *C. riparius* are to be conducted according to the ASTM (1992) procedure. Culture of *H. azteca* are to be conducted according to the procedure described in Borgmann *et al* (1989). Eggs of the mayfly *Hexagenia* spp. (a mixture of *H. limbata and H. rigida*) are to be collected and organisms are to be cultured using the procedure of Hanes and Ciborowski (1992) and Bedard *et al* (1992).
- 3) Tests with *H. azteca*, *C. riparius* and *T. tubifex* are to be conducted in 250 glass beakers containing 60 to 100 mL of sieved (500 um), homogenized sediment with approximately 100 to 140 mL of overlying carbon-filtered, dechlorinated and aerated Lake Ontario water. Tests with the mayfly, *Hexagenia*, are to be conducted in 1 L glass jars with 150mL of test sediment and 850 mL overlying water. The sediment is allowed to settle for 24 h prior to the addition of animals. Test are to be initiated with the random addition of 15 organisms per beaker for *H. azteca* and *C. riparius*, 10 organisms per beaker for *Hexagenia* spp. and 4 organisms per beaker for *T. tubifex*. Juveniles of *H. azteca* are to be 3 to 7 d old at test initiation; *C. riparius* larvae are first instars and approximately 3 d post-oviposition; *Hexagenia* nymphs are 1.5 to 2 months old (approximately 5 to 10 mg wet weight) and *T. tubifex* adults are 8-9 weeks old. Tests are to be conducted at 23±1·C with a 16L:8D photoperiod.

SCHEDULE A

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Tests are to be static with the periodic addition of distilled water to replace water lost during evaporation. Each beaker should be covered with a plastic petri dish with a central hole for aeration using a Pasteur pipette and air line. Tests are to be terminated after 10 d for *C. riparius*, 21 d for *Hexagenia* and 28 d for *H. azteca* and *T. tubifex*. Endpoints to be measured in the tests are: *H. azteca*, survival and growth (mean dry weight in mg); *C. riparius*, survival and growth; *Hexagenia*, survival and growth; and *T tubifex*, survival and production of coccoons and young.

III. REPORTING REQUIREMENTS

- 1) The Contractor is to provide ten copies of the draft report to the NRBS Component Coordinator by December 31st, 1993. The draft report is to summarize and interpret the toxictiy testing carried out by the Contractor in II, above.
- 2) Three weeks after the receipt of review comments on the draft report, the Contractor is to provide the Component Coordinator with two unbound, camera copies and ten cerlox bound copies of the final report. The final report is to include the following: an acknowledgement section that indicates any local or native involvement in the project, table of contents, list of tables, list of figures and an appendix with the Terms of Reference for this project. Text for the report should be in Times Roman 12 point font. If photographs are to be included in the report they should be high contrast black and white. All tables and figures in the report should be clearly reproducible by a black and white photocopier. Along with copies of the final report, the Contractor is to supply an electronic version of the report in Word Perfect 5.1 format. Electronic copies of tables, figures and data appendices in the report are also to be submitted to the Component Coordinator along with the final report. These should be submitted in a spreadsheet (Quattro Pro preferred, but also Excel or Lotus) or database (dBase IV) format. Where appropriate, data in tables, figures and appendices should be geo-referenced

IV. CONTACTS

The Component Coordinator for this project is:

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This project has been proposed by the Contaminants Component of the NRBS. The Contaminants Component Leader is:

Dr. Brian Brownlee National Water Research Institute 867 Lakeshore Road P.O. Box 5050 Burlington, Ontario phone: (416) 336-4706 L7R 4A6 fax: (416) 336-4972

V. LITERATURE CITED

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