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Special Report 76

Geochemical, Mineralogical and Kimberlite Indicator Mineral Electron Microprobe Data from Silts, Heavy Mineral Concentrates and Waters from a National Geochemical Reconnaissance Stream Sediment and Water Survey

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Alberta Energy and Utilities Board

Alberta Geological Survey

Geochemical, Mineralogical and Kimberlite Indicator Mineral Electron Microprobe Data from Silts, Heavy Mineral Concentrates and Waters from a National Geochemical Reconnaissance Stream Sediment and Water Survey in the Northern and Southwestern Buffalo Head Hills, Northern Alberta (Parts of NTS 84B, 84C, 84F and 84G)

Introduction

A regional stream sediment and water geochemical survey was undertaken in the northern and southwestern part of the Buffalo Head Hills in 2004. The 2004 data are part of an ongoing survey of the Buffalo Head Hills area initiated in 2001 (Friske et al., 2003; McCurdy et al., 2004). This National Geochemical Reconnaissance (NGR) project contributes to Alberta's plan for a multi-year, multi-disciplinary study in the northern part of the province.

Analytical data accompany this document for 49 elements in stream sediments, 50 elements in heavy mineral concentrates, and 60 variables in waters from 122 sites sampled in 2004. Selected results from the 2004 NGR survey were released in Prior et al. (2005a; 2005b). The Geological Survey of Canada, under the Targeted Geoscience Initiative II (TGI II) and Northern Resources Development Program, and the Alberta Energy and Utilities Board/Alberta Geological Survey (EUB/AGS), funded the 2004 survey.

Analytical results and field observations contribute to building a national geochemical database for resource assessment, mineral exploration, geological mapping, and environmental studies. Sample collection, preparation procedures and analytical methods are strictly specified and carefully monitored to ensure consistent and reliable results regardless of the area, the year of collection or the analytical laboratory undertaking the analyses (Friske and Hornbrook, 1991).

The northern Buffalo Head Hills survey area lies south and southeast of La Crete, a community located 550 km north-northwest of Edmonton (Fig. 1). Access to sample sites was mainly by helicopter. Seismic cut lines and pipeline right-of-ways provide some access for all-terrain vehicles. Services available in La Crete include lodging, gas stations, grocery and hardware stores, and restaurants. There is a small airport with no services.

Samples were collected in the southwestern part of the survey area by helicopter based at Red Earth Creek, Alberta, a community located 350 km north-northwest of Edmonton (Fig. 1). Services available in Red Earth Creek include lodging, gas stations, grocery and hardware stores, and restaurants. There is a small airport with several companies providing helicopter and charter services.

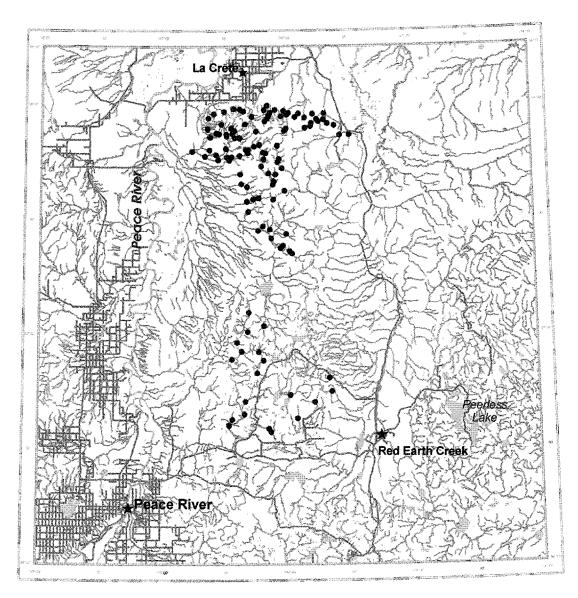


Figure 1. Location map shows area of survey south and southeast of the village of La Crete, and west of Red Earth Creek. Black circles indicate the location of sample sites.

Sample Collection

Stream Sediments (Silts)

At each site a pre-labelled Kraft paper bag (12.5 cm x 28 cm with side gusset) (Fig. 2) was two-thirds filled with silt or fine sand collected from the active stream channel. In practice, the silt sample was collected after water samples were collected but before a bulk sediment sample. Commonly, the sampler collected handfuls of silt from various

points in the active stream channel while moving gradually upstream, normally over a distance of 5 to 15 m. If the stream channel consists mainly of clay, coarse material or organic sediment from which suitable sample material is scarce or absent, moss mat from the stream channel, which commonly contains trapped silt, may be added to the sample. A field duplicate pair of silt samples, assigned sequential sample numbers, is collected within each block of 20 samples. The first sample of the pair is assigned a replicate status value of 10 and the second is assigned a replicate status value of 20. Routine (non-duplicate) field samples are assigned replicate status values of 0.

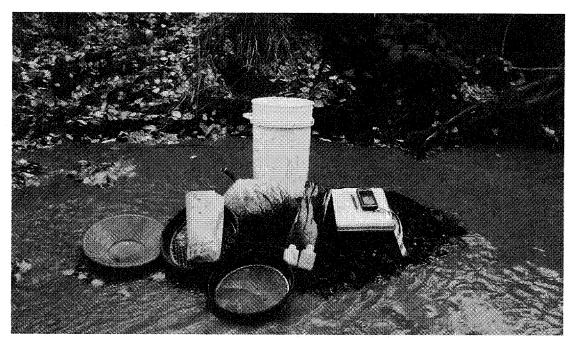


Figure 2. Pre-labelled Kraft paper bags and plastic bottles are used to collect samples of stream silts and stream waters. A bulk sample, for heavy mineral processing, is collected by wet-sieving coarse-grained stream sediment using a US Sieve Series 12-mesh (1.68 mm) sieve and collecting <12 mesh grains in a plastic pail lined with a polyethylene sample bag. The gold pan is used for adding water for wet sieving, not for heavy mineral concentrate panning. A sample composed of granules and pebbles, for archive, is collected at bulk sample sites by sieving >12 mesh material through a US Sieve Series 2-mesh (10 mm) sieve and collecting the <10 mm material in a labelled Kraft paper bag. Flagging tape with a sample site number is used to mark sample sites. Field observations are noted on pre-printed water-resistant paper (see Appendix).

Stream Waters

Waters were sampled in mid-channel, from flowing water where possible. At each site two water samples were collected in 125-ml Nalgene HDPE bottles (Fig. 2). Samples were collected after first rinsing bottles at least two times in flowing water before a final fill. Field duplicate pairs of water samples, assigned sequential sample numbers, are collected within each block of 20 samples. The first sample of each pair is assigned a replicate status value of 10 and t he second is assigned a replicate status value of 20. Routine (non-duplicate) field samples are assigned replicate status values of 0.

Heavy Mineral Concentrates

Ideal sites for the collection of sediments for the heavy mineral concentrate fraction are located at the upstream points of mid-channel bars. Material was collected from a single location where possible, or within close proximity otherwise. A 5-gallon (22.7 litre) plastic pail was lined with a heavy-duty polyethylene bag (18x24 inches, 4 Mil). Material was wet-sieved through a 12-mesh (1.68 mm) stainless steel sieve, placed on top of the pail, until a sample weight of 10-15 kg was attained. The bag lining the pail was taped shut with black plastic (electrical) tape and placed into a second bag with a sample number and taped. Samples were shipped directly to a commercial lab for preparation and analysis.

Sample Preparation

Stream Sediments (Silts)

The Kraft paper bags containing the silt samples were placed into plastic bags, taped with electrical tape and shipped directly to a commercial lab, where they were air-dried at temperatures below 40° C and sieved through a minus 80-mesh (177 µm) screen. Control reference and duplicate samples were inserted into each block of twenty sample numbers.

Waters

One set of water samples remained unaltered and was packed and shipped to a laboratory for analysis. The second set of water samples was filtered within 24 hours of collection through single-use Millipore Sterivex-HV 0.45-µm filter units attached to 50-ml sterile plastic syringes. After 50 ml of water from the second set of samples was filtered into new 60-ml bottles, the remainder was used for the determination of pH and conductivity before being discarded. Using an Eppendorf pipettor with disposable plastic tips, 0.5 ml 8M HNO₃ was added to filtered water samples. Syringes were reused after rinsing with distilled, de-ionized water, but replaced daily. Control reference samples (filter, acid and travel blanks*) were added to each batch of samples. Filtered and acidified waters were kept in a cool dark place until shipment to the lab. Control reference samples were inserted into each block of 20 water samples. No duplicate water samples were introduced.

Heavy Mineral Concentrates

Bulk sediment samples were progressively reduced by different laboratory procedures to concentrate heavy minerals. Initially a 500-g character sample was taken and stored

^{*} Filter (sample) blanks are 60-ml bottles filled with deionized water used in the field that has been filtered and acidified at the same time as routine samples; acid blanks are samples of the deionized water used in the field and acidified (but not filtered) at the same time as routine samples; travel blanks are bottles of deionized water pre-filled at the GSC lab in advance of field sampling and acidified in the field with the survey samples.

before a low-grade table concentrate was prepared from the remainder. Gold grains were observed at this stage and counted, measured and classified as to degree of wear (reflecting distance of transport). The table reject was re-tabled to scavenge possible unrecovered kimberlite indicator minerals and magmatic massive sulphide indicator minerals. The concentrate from both tabling runs was separated in methylene iodide diluted with acetone to S.G. 3.20 to recover heavy minerals including Cr-diopside and forsterite olivine. Magnetite was removed after the heavy liquid separation and the remaining concentrate cleaned with oxalic acid to remove limonite stains. The dried concentrate was sieved to separate it into several size fractions, (<0.25 mm, 0.25 to <0.5 mm, 0.5 mm to <1.0 mm, \geq 1.0 mm to 2.0 mm). The <0.25 mm fraction was kept for chemical analysis and the 0.25 to 0.50 mm fraction was sorted with a Carpco® drum magnetic separator into strongly, moderately, weakly and non-paramagnetic fractions.

Analytical Procedures

Stream Sediment (Silt) Analyses

Instrumental Neutron Activation Analysis (INAA)

Weighed and encapsulated samples, normally 30 g, were packaged for irradiation along with internal standards and international reference materials. Samples and standards were irradiated together with neutron flux monitors in a two-megawatt pool type reactor. After a seven-day decay period, samples were measured with a high-resolution germanium detector. Typical counting times were 500 seconds. Elements determined by INAA are listed in Table 1.

Table 1 Elements in stream sediments and the <0.25 mm fraction of heavy mineral concentrates determined by Instrumental Neutron Activation analysis.

ELEMENT		DETEC	TION LEVEL
As	Arsenic	0.5	ppm
Au	Gold	2	ppb
Ba	Barium	50	ppm
Br	Bromine	0.5	ppm
Ce	Cerium	5	ppm
Co	Cobalt	5	ppm
Cr	Chromium	20	ppm
Cs	Cesium	0.5	ppm
Eu	Europium	1	ppm
Fe	Iron	0.2	pct
Hf	Hafnium	1	ppm
La	Lanthanum	2	ppm
Lu	Lutetium	0.2	ppm
Мо	Molybdenum	1	ppm
Na	Sodium	0.02	pct
Ni	Nickel*	10	ppm
Rb	Rubidium	5	ppm
Sb	Antimony	0.1	ppm
Sc	Scandium	0.2	ppm
Sm	Samarium	0.1	ppm
Та	Tantalum	0.5	ppm
Tb	Terbium	0.5	ppm
Th	Thorium	0.2	ppm

Ti	Titanium*	500	ppm
U	Uranium	0.2	ppm
W	Tungsten	1	ppm
Yb	Ytterbium	2	ppm
* HMC samples only			

INAA data for silver, cadmium, iridium, selenium, tin, tellurium, zinc, and zirconium are not published because of inadequate detection limits and/or precision.

Inductively Coupled Plasma Mass Spectrometry (ICP-MS) and Other Analyses

For determination of the 38 elements listed in Table 2, a one-gram sample was leached with 6 ml of a mixture of HCl, HNO₃, and distilled, deionized water (2:2:2 v/v) at 95° C for one hour. The sample solution was diluted with de-ionized water to 20 ml and analysed by inductively coupled plasma emission spectroscopy on a Jarell-Ash instrument and inductively coupled plasma mass spectroscopy on a Perkin-Elmer Elan instrument.

Loss-on-ignition was determined using a one-gram sample. The sample, weighed into a Leco® crucible, was placed into a 100°C muffle furnace and brought up to 500° C for one hour. The oven was cooled to 100°C and crucibles transferred to a desiccator for cooling to room temperature. The crucibles were re-weighed, and the difference was reported as loss-on-ignition.

Tin in heavy mineral concentrates was prepared for analysis by heating a 0.2 g sample mixed with 1.5 g of LiBO₂ flux in a muffle furnace for 15 minutes at 1050°C in a graphite crucible. The molten mixture was removed and immediately poured into 100 mL of 5% HNO₃. The solution was shaken for two hours and then an aliquot was poured into a test tube. Analytical values were determined using a Perkin-Elmer Elan 6000 ICP-MS (Gravel, 2003).

Table 2 Variables in stream silts and the <0.25 mm fraction of heavy mineral concentrates determined by Inductively Coupled Plasma - Mass Spectrometry and other analytical methods

VARIA	BLE	DETEC	CTION LEVEL
Ag	Silver	2	ppb
ΑĬ	Aluminum	0.01	pct
As	Arsenic	0.1	ppm
B†	Boron	1	ppm
Ва	Barium	0.5	ppm
Bi	Bismuth	0.02	ppm
Ca	Calcium	0.01	pct
Cd	Cadmium	0.01	ppm
Co	Cobalt	0.1	ppm
Cr	Chromium	0.5	ppm
Cu	Copper	0.01	ppm
F	Fluorine	10	ppm
Fe	Iron	0.01	pct
Ga	Gallium	0.2	ppm
Hg	Mercury	5	ppb
K	Potassium	0.01	pct

La	Lanthanum	0.5	ppm
LOI*	Loss-on-ignition	1	pct
Mg	Magnesium	0.01	pct
Mn	Manganese	1	ppm
Mo	Molybdenum	0.01	ppm
Na	Sodium	0.001	pct
Ni	Nickel	0.1	ppm
P	Phosphorus	0.001	pct
Pb	Lead	0.01	ppm
S	Sulphur	0.01	pct
Sb	Antimony	0.02	ppm
Sc	Scandium	0.1	ppm
Se	Selenium	0.1	ppm
Sn	Tin (LiBO ₂ Fusion)	1	ppm
Sr	Strontium	0.5	ppm
Те	Tellurium	0.02	ppm
Th	Thorium	0.1	ppm
Ti	Titanium	0.001	pct
TI	Thallium	0.02	ppm
U	Uranium	0.1	ppm
V	Vanadium	2	ppm
W	Tungsten	0.1	ppm
Zn	Zinc	0.1	ppm
*Silt san	nples only		

Fluorine was determined using 0.25-gram sample splits weighed into Ni crucibles. One gram of sodium hydroxide was added and the mixture was fused at 600°C in a muffle furnace. The fusion product was dissolved into a mixture of 7 ml of de-ionized water and 5 ml of 30% sulphuric acid. The solutions were transferred to plastic beakers and 5 ml of 30% ammonium acetate added. The volume was made up to 90 ml with de-ionized water. The pH was tested and adjusted to 7.8 with either sodium hydroxide or sulphuric acid. Fluorine content was determined using a fluorine selective ion electrode.

Water Analyses

The pH of stream waters was determined at the field base location using a Hanna Instruments Combo® waterproof tester with automatic temperature compensation, a range of 0.00 to ± 14.00 pH, resolution of 0.01 pH and an accuracy of ± 0.05 pH. Meters were calibrated using commercial buffer solutions with pH values of 4.0, 7.0 and 10.0.

Conductivity of stream waters was determined at the field base location using a Hanna Instruments Combo® waterproof tester with a range of 0 to 3999 μ S/cm, a resolution of 1 μ S/cm and a full-scale accuracy of ±2%. Meters were calibrated using commercial conductivity standards with values of 1413 μ S/cm and 84 μ S/cm.

Chloride (Cl⁻), fluoride (F⁻), nitrate (NO₃⁻) and sulphate (SO₄²⁻) in untreated waters (water samples not filtered or acidified in the field) were determined by ion chromatography (IC) at Geological Survey of Canada laboratories in Ottawa. Untreated water is identified as UU-Water (unfiltered, unacidified) in the data table.

Uranium in untreated waters was determined by inductively coupled plasma-mass spectrometry (ICP-MS) at Geological Survey of Canada laboratories in Ottawa.

Trace and Major Elements

Acidified and filtered stream water samples were analyzed for trace metal and major elements at Geological Survey of Canada laboratories in Ottawa. A complete list of elements and stated detection limits are given in Table 3.

Trace metal analysis was performed using a VG PQII ICP-MS with a Meinhard concentric glass nebulizer, Type K (solution uptake rate 1 ml min⁻¹), a quartz Scott-type double-pass chilled spray chamber (2°C) and a 27 MHz standard quartz torch. The argon flow-rates are: Cool 12.5 l min⁻¹, Auxiliary 0.85 l min⁻¹, and Nebulizer 0.9 l min⁻¹. The RF power is 1350 watts. Isotopes measured and corrections for spectral interferences are detailed in Hall et al. (1995) and Hall et al. (1996). Data for hafnium and zirconium are not published because these elements are not sufficiently stabilized in waters by the addition of nitric acid. Data for indium, selenium, silver, tantalum and thulium are not published because of inadequate detection limits and/or precision.

 Table 3
 Major and trace elements determined in stream waters.

ELEM	ENT	DETEC LEVEL	TION	LABORATORY METHOD
Water	s -Filtered, Acidified (FA-Water)			
Al As	Aluminum Arsenic	2 0.1	ppb ppb	ICP-MS ICP-MS
В	Boron	0.5	ppb	ICP-MS
Ba Be	Barium	0.2	ppb	ICP-MS
Cd	Beryllium Cadmium	0.005 0.02	ppb	ICP-MS ICP-MS
Ce	Cerium	0.02	ppb ppb	ICP-MS
Cs	Cesium	0.01	ppb	ICP-MS
Co	Cobalt	0.05	ppb	ICP-MS
Cr	Chromium	0.03	ppb	ICP-MS
Cu	Copper	0.1	ppb	ICP-MS
Dy	Dysprosium	0.005	ppb	ICP-MS
Er	Erbium	0.005	ppb	ICP-MS
Eu	Europium	0.005	ppb	ICP-MS
Ga	Gallium	0.01	ppb	ICP-MS
Gd	Gadolinium	0.005	ppb	ICP-MS
Ge	Germanium	0.02	ppb	ICP-MS
Но	Holmium	0.005	ppb	ICP-MS
La	Lanthanum	0.01	ppb	ICP-MS
Li	Lithium	0.02	ppb	ICP-MS
Lu	Lutetium	0.005	ppb	ICP-MS
Mn	Manganese	0.1	ppb	ICP-MS
Мо	Molybdenum	0.05	ppb	ICP-MS
Nb	Niobium	0.01	ppb	ICP-MS
Nd	Neodymium	0.005	ppb	ICP-MS
Ni	Nickel	0.2	ppb	ICP-MS
Pb	Lead	0.01	ppb	ICP-MS
Pr	Praseodymium	0.005	bbp	ICP-MS
Rb	Rubidium	0.05	ppb	ICP-MS
Re Sb	Rhenium	0.005 0.01	ppb	ICP-MS ICP-MS
Sm	Antimony Samarium	0.005	ppb	ICP-MS
Sn	Tin	0.005	ppb ppb	ICP-MS
Sr	Strontium	0.5	ppb	ICP-MS
Tb	Terbium	0.005	ppb	ICP-MS
Te	Tellurium	0.003	ppb	ICP-MS
Ti	Titanium	0.52	ppb	ICP-MS
π	Thallium	0.005	ppb	ICP-MS
Ü	Uranium	0.005	ppb	ICP-MS
v	Vanadium	0.1	ppb	ICP-MS

بحسسون				
W	Tungsten	0.02	ppb	ICP-MS
Y	Yttrium	0.01	ppb	ICP-MS
Yb	Ytterbium	0.005	ppb	ICP-MS
Zn	Zinc	0.5	ppb	ICP-MS
Ca	Calcium	0.02	ppm	ICP-ES
Fe	Iron	0.005	ppm	ICP-ES
K	Potassium	0.05	ppm	ICP-ES
Mg	Magnesium	0.005	ppm	ICP-ES
Na	Sodium	0.05	ppm	ICP-ES
P	Phosphorus	0.05	ppm	ICP-ES
s	Sulphur	0.05	ppm	ICP-ES
Si	Sílicon	0.02	ppm	ICP-ES
Waters -	-Unfiltered, not Acidified (UU-Water)			,
CI.	Chloride	100	ppb	ic ·
F	Fluoride	50	ppb	l ič
NO ₃	Nitrate	50	ppb	lic
SO ₄	Sulphate	50	ppb	ic
Cond.	Conductivity	1 1	μS/cm	EC (electrolytic conductivity)
pH	-	0.1	μω/GII	GCE (glass-calomel electrode)
ΰ	Uranium	0.05	ppb	ICP-MS

Major element analysis was performed using a Perkin-Elmer 3000DV Inductively Coupled Plasma – Emission Spectrometer (ICP-ES) with a cross-flow nebulizer (solution uptake rate 1 ml min⁻¹), a Ryton Scott-type double-pass spray chamber and a custom demountable quartz ICP-ES torch. The argon flow-rates are: Plasma 15.0 I min⁻¹, Auxiliary 0.5 I min⁻¹, and Nebulizer 0.7 I min⁻¹. The RF power is 1350 watts. All elements were analyzed in axial mode except for sodium and potassium. These elements were run in radial mode. Inter-element correction factors were applied as required to correct for various spectral interferences. Data for scandium are not published because of inadequate detection limits and/or precision.

Heavy Mineral Concentrate Analyses

Kimberlite indicator minerals (KIMs) were picked and identified from each of three size fractions (0.25-0.5 mm, 0.5 mm-1.0 mm, 1.0-2.0 mm). Fractions exceeding a 100 g threshold were characterized by a 100 g split and normalized to represent the total sample weight. Following removal of the kimberlite indicator minerals, 100 grains were randomly selected from each 0.25-0.5 mm fraction and identified. After 100 grains were identified they were recombined with the source sample fraction. The 0.25-0.5 mm, 0.5-1.0 mm and 1.0-2.0 mm fractions (minus KIMs) were archived. The <0.25 mm fraction of the heavy mineral concentrate was sent to a commercial lab where it was ground in a ceramic mill and analyzed by a combination of ICP-MS, INAA and specific methods. Kimberlite indicator mineral grains underwent electron microprobe analysis for chemical characterization. A mineralogical consultant evaluated results of electron microprobe data.

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Ottawa, Ontario

Analysis: Acme Analytical Laboratories

Vancouver, British Columbia

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Mississauga, Ontario

GSC Analytical Chemistry Laboratory

Ottawa, Ontario

GSC Analytical Method Development Laboratory

Ottawa, Ontario

GSC Mineralogical Laboratory (electron microprobe)

Ottawa, Ontario

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Ottawa, Ontario

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NATIONAL GEOCHEMICAL RECONNAISSANCE (Silt Sediment, Bulk Sediment and Water Field Card - 2005)

	Yak	i i de la companya d	STATE OF THE	CHALL SALE OF A		THE TORS
GENERAL PHYSIOGRAPHY Mountainous Hilly	STREAM SOURCE(S) Ground SpringMelt Glacier	WATER COLOUR	CONTAMINATION(S) None Possible Probable	STREAM SEDIMENT SAMPLE COLOUR(S)	BULK SITE Longitudinal Bar Transverse Bar	BULK SITE COMPOSITION Cobbles % Pebbles %
Plateau Plain Swamp	Recent Rain Unknown	WATER CLARITY Transparent	Definite Mining Industry	STREAM SEDIMENT	Point Bar Diagonal Bar Boulder Trap	Sand % Silt %
SURFACE EXPRESSION	STREAM CLASS Primary	Partially Cloudy Cloudy	Agriculture Domestic Forestry	COMPOSITION Sand % Silt & Clay %	Log Trap Vegetation Trap Bedrock Step	Clay % Organics %
Hummocky Inclined Level	Secondary Tertiary Quaternary	VEGETATION Coniferous Deciduous	Burn Other	Organics %	Pool Gravel Veneer Stream Bed	PEBBLE LITHOLOGY(IES)
DRAINAGE PATTER Dendritic	Undefined STREAM TYPE Permanent	Mixed Grass Bog	BANK PRECIPITATE	SAMPLE TYPE(S) Silt Sediment (SS)	Beaver Dam	%
Herringbone Rectilinear Trellis	Intermittent Re-emergent Undefined	Other BANK TYPE(S) Alluvium		Bulk Sediment (BS) Pebble (PB) Water (SW)	SITE RATING Good Good to Moderate Moderate	BEDROCK EXPOSED No Yes
Poor Discontinuous Closed	STREAM FLOW Stagnant	Colluvium Till Outwash	Colour(s) BOTTOM PRECIPITATE No Yes	Ofns/	Moderate to Poor Poor	Į.
SITE DRAINAGE Well Moderate	Slow Moderate Fast Torrential	Bare Rock Talus/Scree Organic Other	Į.	IN-SITU WATER	CLAST SHAPE Rounded % Sub-Angular/ %	Type(s) BOULDERS PRESENT No Yes
Poor	A D	Longitude (DI	Colour(s) D)	Latitude (DD)	Sub-Rounded 76 Angular % Platy/Flat %	Type(s)
COMMENTS:						
kre Täll	YEAR	SANGU	RP WETH	DEFICH DATE	THAE	cdulacions
GENERAL PHYSIOGRAPHY	STREAM SOURCE(S) Ground	WATER COLOUR	CONTAMINATION(S) None	STREAM SEDIMENT SAMPLE COLOUR(S)	BULK SITE Longitudinal Bar	BULK SITE COMPOSITION
Mountainous Hilly Plateau Plain	SpringMelt Glacier Recent Rain	WATER CLARITY	Possible Probable Definite	<u>-</u>	Transverse Bar Point Bar Diagonal Bar	Cobbles % Pebbles % Sand %
Swamp SURFACE	Unknown STREAM CLASS Primary	Transparent Partially Cloudy Cloudy	Mining Industry Agriculture Domestic	STREAM SEDIMENT COMPOSITION Sand ——— % Silt & Clay ——— %	Boulder Trap Log Trap Vegetation Trap	Silt % Clay % Organics %
EXPRESSION Hummocky Inclined Level	Secondary Tertiary Quaternary	VEGETATION Coniferous Deciduous	Forestry Burn Other	Organics %	Bedrock Step Pool Gravel Veneer	PEBBLE LITHOLOGY(IES)
DRAINAGE PATTER	Undefined N STREAM TYPE Permanent	Mixed Grass Bog	BANK PRECIPITATE	SAMPLE TYPE(S) Silt Sediment (SS)	Stream Bed Beaver Dam SITE RATING	% %
Herringbone Rectilinear Trellis Poor Discontinuous	Intermittent Re-emergent Undefined STREAM FLOW	Other BANK TYPE(S) Alluvium Colluvium Till	Colour(s)	Bulk Sediment (BS) Pebble (PB) Water (SW)	Good Good to Moderate Moderate Moderate to Poor Poor	BEDROCK EXPOSED No Yes
SITE DRAINAGE Well Moderate Poor	Stagnant Slow Moderate Fast Torrential	Outwash Bare Rock Talus/Scree Organic Other	□ No Yes □ Colour(s)	IN-SITU WATER	CLAST SHAPE Rounded —— % Sub-Angular/ Sub-Rounded —— %	Type(s) BOULDERS PRESENT No Yes
	A D	Longitude (DD		Latitude (DD)	Angular % Platy/Flat %	Type(s)
COMMENTS:						

NTS SPEET	e.g. 0 8 4 F I 3
NTS Sheet: National Topographic System 1:250,000 index refer and occupy the first four boxes. The final two boxes are used for	ence, consisting of three numbers and one letter e.g. 085E the 1:50,000 sheet identification e.g. 13. if applicable.
1,	
YEAR	e.g. 2 0 0 5
Year: The four-digit year, e.g. 2005.	
- HAMALE	e a 19/mpal a
NUMBER 1	e.g. 13050 2
Sample Number: A four-digit sample number e.g. 1002. The first other three digits are a sequential series from 001 to 999, for example 1000 to 999	st digit refers to the collection party crew number, while the nole:
Crew 1 samples range from 1002 to 1999,	- -
Crew 2 samples range from 2002 to 2999, Crew 3 samples range from 3002 to 3999,	
REP	a±0
STAT	e.g. [970]
Rep Stat (Replicate Status): A two digit number e.g. 00, definir 00 routine sample	ng the relationship of the current sample to others in the survey
10 first sample of a field duplicate pair	
20 second sample of a field duplicate pair 80 blind duplicate number (empty bag) for a blind duplicate	e cut of one of previous 18 field samples
90 control reference number (empty bag) for cut of a control	of reference sample
WHITH	e.g. VIII
Stream Width: width of the stream estimated in metres to the close	st 1/10 th of a metre e.g. 1.1metres wide.
	[marken]
DEPTH	e.g. [10][11]
Stream Depth: depth of the stream estimated in metres to the closes	st 1/10 th of a metre e.g. 0.1 metres deep.
	<u></u>
DATE NO	e.g. 1 T TO 7
Date: date of collection, DD MM format, e.g. July 11 = 11 07	
TIME	e.g. 1 442 6
me: time of day (24 hour clock), e.g. 2:26pm = 14:26	S.6. [-].]-[-]

Collectors: initials of the collection crew, first three boxes for the navigator (A.B.C.), and the last three for the second sampler (D.E.F.), e.g. ABC DEF

GENERAL PHYSIOGRAPHY	STREAM TYPE
Mountainous - rugged area of uplift having at least 300m gain	Permanent - year-round flow
from base to peak	Intermittent - seasonal flow during wet season or spring runoff
Hilly - natural elevation change, of less than 300m, while having a well defined outline	Re-emergent - discontinuous stream course
	Undefined -
Plateau - an elevated area of fairly level ground	STREAM FLOW
Plain - any area of level or near-level open land	Stagnant - little or no flow
Swamp - low waterlogged area having shrubs and/or trees	Slow - speed of a slow walker
	Moderate - speed of someone briskly walking
SURFACE	Fast - speed of a jogger
EXPRESSION	Torrential - speed of a quick jogger
Hummocky - series of rounded knobs and kettles	WATER COLOUR
Inclined - constant sloping surface	dominant colour (if any)
Level - flat or gently sloping	of the stream <u>water</u>
DRAINAGE PATTERN	
Dendritic - "tree-like" network of streams	WATER OLABITY
Herringbone - V-shaped pattern of streams	WATER CLARITY
Rectilinear/Trellis - series of parallel streams with near right-angle	Transparent - clear (any colour) Partially Cloudy - semi opaque (any colour)
turns and perpendicular intersections Parallel - streams flowing parallel before joining at small angles	
Poor/Deranged - no clear geometry in the drainage and no	Cloudy - opaque or nearly opaque (any colour)
true stream valley pattern	VEGETATION
Discontinuous - stream disappears for a short distance then re-appears down slope	Coniferous - having needle-like leaves, e.g. spruce, pine, incl. tamarack.
Closed	Deciduous - trees that shed their leaves annually, e.g. maple, poplar
SITE	Mixed - roughly equal mixture of coniferous and deciduous trees
DRAINAGE	Grass - grasslands surrounding site
Well - stream channel well developed and well drained	Bog - waterlogged spongy ground, sphagnum moss dominate
Moderate	Other
Poor - stream channel poorly developed and poorly drained	
STREAM SOURCE	BANK TYPE
Ground - stream flow originates from natural springs or seeps	Alluvium - clay, silt, sand or gravel recently deposited by stream action
SpringMelt - stream flow greater due to melting of winter's snow	Colluvium - accumulation of material through the action of gravity
Glacier - stream originates from melting glacier	Till - glacial till (unsorted)
Recent Rain - stream flow greater due to recent rain	Outwash - stratified sand or gravel deposited by glaciofluvial melt water
Unknown	Bare Rock - bedrock
	Talus/Scree - loose rock fragments derived from an adjacent steep rocky slope
STREAM CLASS (determined from 1:250,000 NTS)	Organic - peaty organic soil or sediment Other
Primary - smallest stream, originates from springs and seeps	
Secondary - stream below confluence of two primary streams	
Tertiary - stream below confluence of two secondary streams	
Quaternary - stream below confluence of two tertiary streams	
Undefined	

CONTAMINATION	
None - no sign of any hum	an activity
Possible - some human ac of contamination Probable - site, area distu	ctivity in area, no obvious sign
· · · · · · · · · · · · · · · · · · ·	ination due to human activity
Mining	
Industry	
Agriculture	
(Specify) Domestic	
Forestry	
Other	
	Syntax
BANK PRECIPITATE	single colour
☐ No Yes ☐	dominant-subordinant
\	multiple distinct colours dominant ; subordinate
Colour(s)	
BOTTOM PRECIPITATE	Syntax
☐ No Yes ☐	single colour dominant-subordinant
$\int_{\mathbb{R}}$	multiple distinct colours
V	dominant ; subordinate
Colour(s)	
STREAM SEDIMENT	Syntax single colour
SAMPLE COLOUR(S)	dominant-subordinant
₿	multiple distinct colours dominant ; subordinate
	aominant y suportantate
STREAM SEDIMENT COMPOSITION	
Sand — % - particles bet	ween 0.0625 and 2 mm in size
	when squeezed into a ball
- holds togethe	r when squeezed, silt is fine
grained and h	gritty feel, clay is very fine as slippery feel
Organics % - muck-like light of organic m	ght weight sediment composed

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SAMPLE TYPE(S)	Check <u>ALL</u> applicable boxe
Silt Sediment (SS)	·
Bulk Sediment (BS)	
Pebble (PB)	
Water (FA)	
IN-SITU WATER	
PHI donb	If applicable
N A D Datum,	either NAD27 or NAD83
	Longitude
Cascapal Degrees	(decimal degrees to at least five decimal places)
	Latitude
Dedimai Degrees	(decimal degrees to at least five decimal places)
	iivo decidai piaces)

BULK SEDIMENT SITE	
Longitudinal Bar - elongated bodies of sediment parallel to stream flow	
Transverse Bar - lobate bodies of sediment oriented roughly perpendicular to stream flow	
Point Bar - elongated bodies of sediment that form on the inside of stream bend, often attached to the inside bank	PEBBLE SHAPE
Diagonal Bar - elongated bodies of sediment orientated obliquely to the stream flow	Rounded — % - smooth and rounded clasts
Boulder Trap - sediment on the down-stream side of a boulder	Sub-Angular/ Sub-Rounded ——— % - rough and semi-rounded clasts
Log Trap - sediment on the down-stream side of a log	Angular % - sharp edged angular clasts
Vegetation Trap - sediment on the down-stream side of vegation	Platy/Flat % - disclike clasts, one dimension much shorter than other two
Bedrock Step - sediment collects down-stream of break in bedrock slope	
Pool - sediment collects down-stream of waterfall or set of rapids	
Gravel Veneer - thin layer of gravel atop finer sediment	
Stream Bed - sediment taken from main stream channel	
Beaver Dam - course sediment exposed by flushing action adjacent or below dam	
	BULK SEDIMENT SITE COMPOSITION
·	Cobbles % - particles between 64 and 256 mm in size
	Pebbles % - particles between 2 and 64 mm in size
SITE RATING*	Sand ——— % - particles between 0.0625 and 2 mm in size - will fall apart when squeezed into a ball
Good - Clast supported, tightly packed, poorly sorted gravel in well	Silt % - particles between 0.02 and 0.0625 mm - holds together when squeezed, silt is fine
formed bedrock depression, pothole or crevice. Clast sizes: boulders,	grained with gritty feel
cobbles, pcbbles, granules. Matrix contains sand and silt. Excavation to bedrock and/or presence of abundant well-rounded clasts enhances	Clay % - particles smaller than 0.02 mm - holds together when squeezed, clay is very fine
site rating. Lack of boulders diminishes rating.	grained and has slippery feel Organics % - muck-like light weight sediment composed
Good to Moderate - Clast supported, tightly packed, poorly sorted gravel upstream or downstream of prominent rock bar or large	of organic materials
boulder and preferably at a level well below the obstruction. Clast	DEDDI E LETTO COMETO DE LA CARACTERIA (CARACTERIA)
sizes: boulders, cobbles, pebbles, granules. Matrix contains sand and silt. Excavation to bedrock and/or presence of abundant well-	PEBBLE LITHOLOGY(IES) - Rough estimate of pebble lithology(ies)
rounded clasts enhances site rating.	%
Moderate - Clast supported, poorly sorted gravel amongst boulders.	%
Packing moderate to tight. Clast sizes boulders (mainly small),	%
cobbles, pebbles, granules. Matrix contains sand and silt. Excavation to bedrock and/or presence of many well-rounded clasts and/or	%
association with some kind of obstruction enhances site rating. Lack of boulders diminishes rating.	
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Moderate to Poor - Matrix supported, generally losely packed gravel strewn on river bed and not associated with any prominent obstruction.	
Sorting is moderate to poor. Boulders are rare or absent. Main clast	BEDROCK EXPOSED
sizes: cobbles, pebbles, granules. Matrix contains sand and silt.	∐ No Yes ☐
Poor - Matrix supported, very loosely packed, fine gravel. Clasts are	\
relatively rare and often form a thin surface veneer on sand or are confined to isolated lenses within a sand mass. Clast sizes: cobbles	Type(s)
(rare), pebbles, granules. Matrix of sand and/or silt. No associated obstruction.	
	BOULDERS* PRESENT
*Muggeridge, M.T., 1986. Pathfinder sampling techniques for locating	No Yes
primary sources of diamond: Recovery of indicator minerals, diamonds and geochemical signatures. Journal of Geochemical Exploration	↓
53 183-204.	Type(s)

* > 256 mm