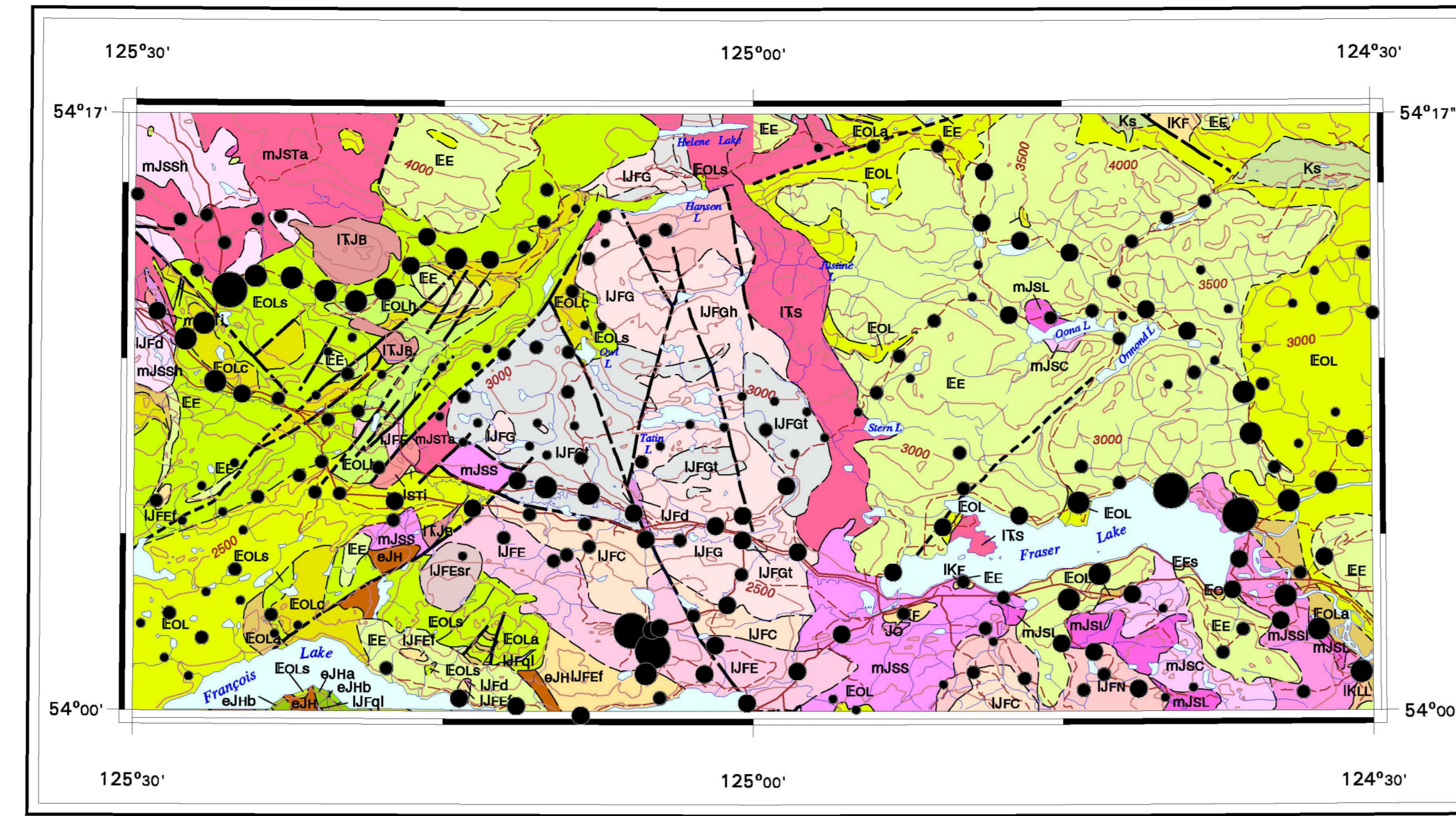


**THORIUM**

INAA

ppm	Th	Percentile
11	●	Maximum
5.0	●	98
3.6	●	95
3.0	●	90
2.4	●	75
1.0	●	50
< 0.1	●	Minimum

217 Samples  
Scalar Exponent = 2.0

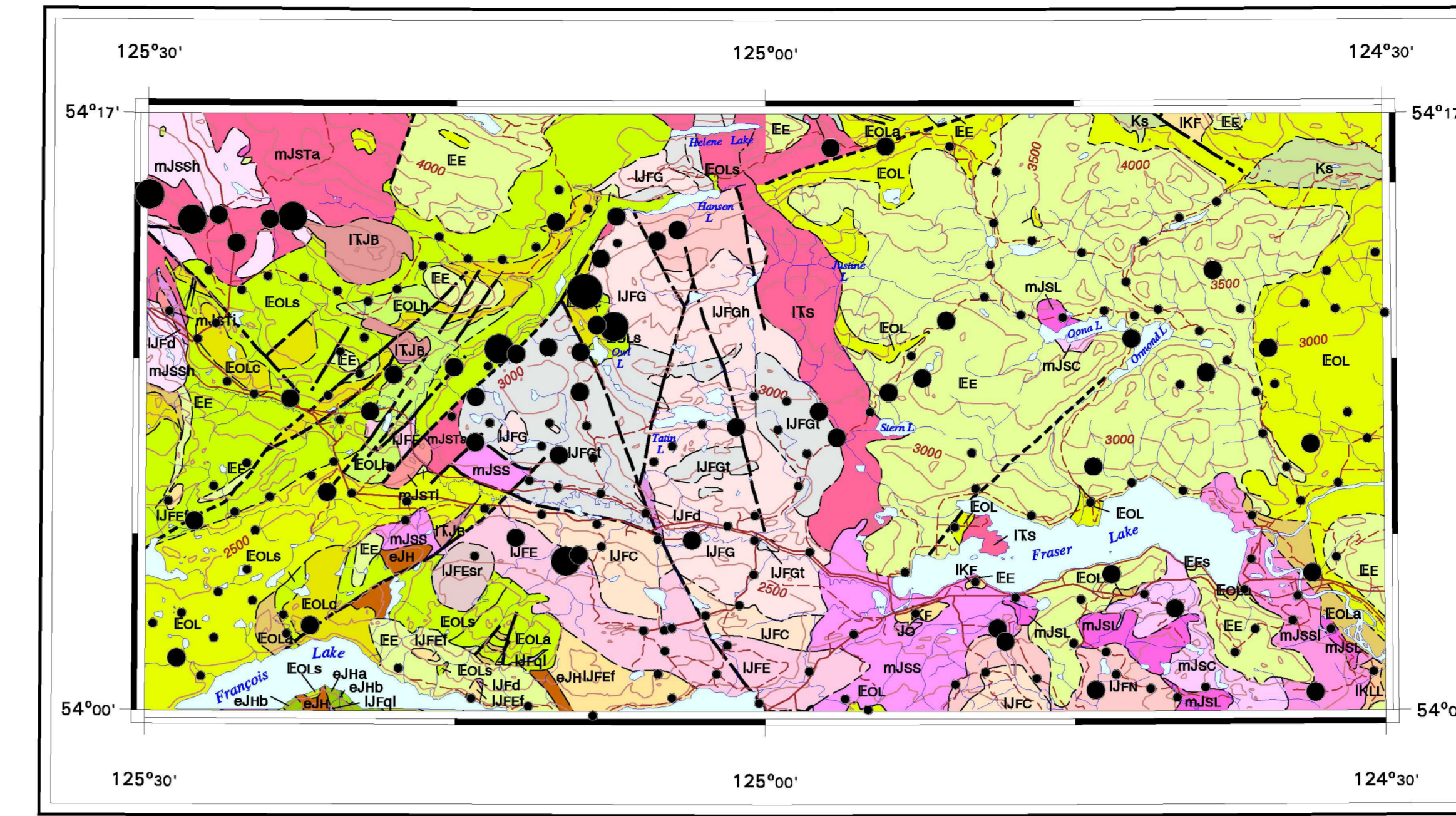


**IRON**

INAA

percent	Fe	Percentile
4.0	●	Maximum
3.5	●	98
3.2	●	95
2.7	●	90
1.9	●	75
0.8	●	50
0.13	●	Minimum

217 Samples  
Scalar Exponent = 1.0

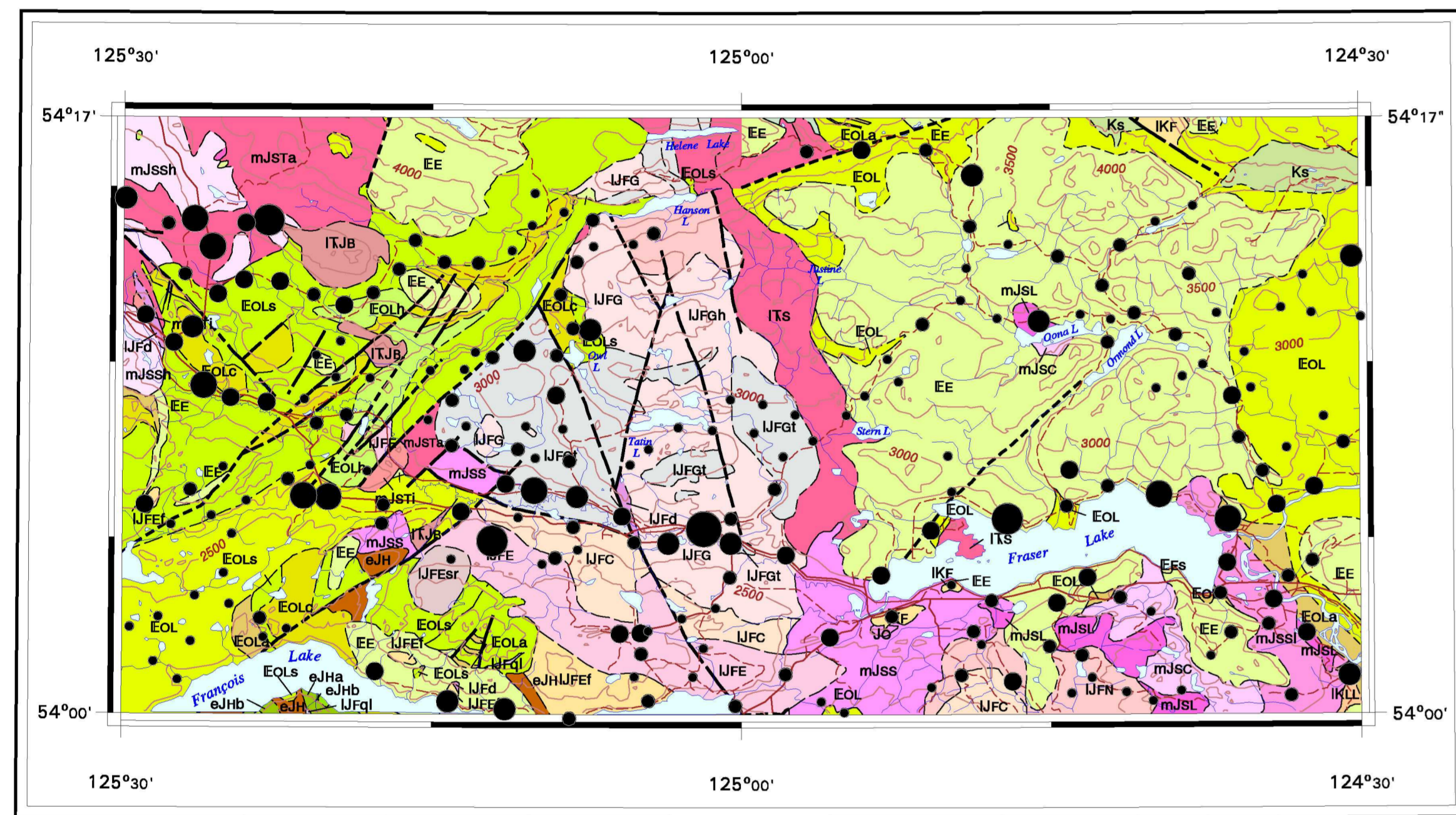


**MAGNESIUM**

ICP-ES

percent	Mg	Percentile
3.7	●	Maximum
2.7	●	98
2.3	●	95
2.0	●	90
1.5	●	75
1.2	●	50
0.4	●	Minimum

217 Samples  
Scalar Exponent = 1.5

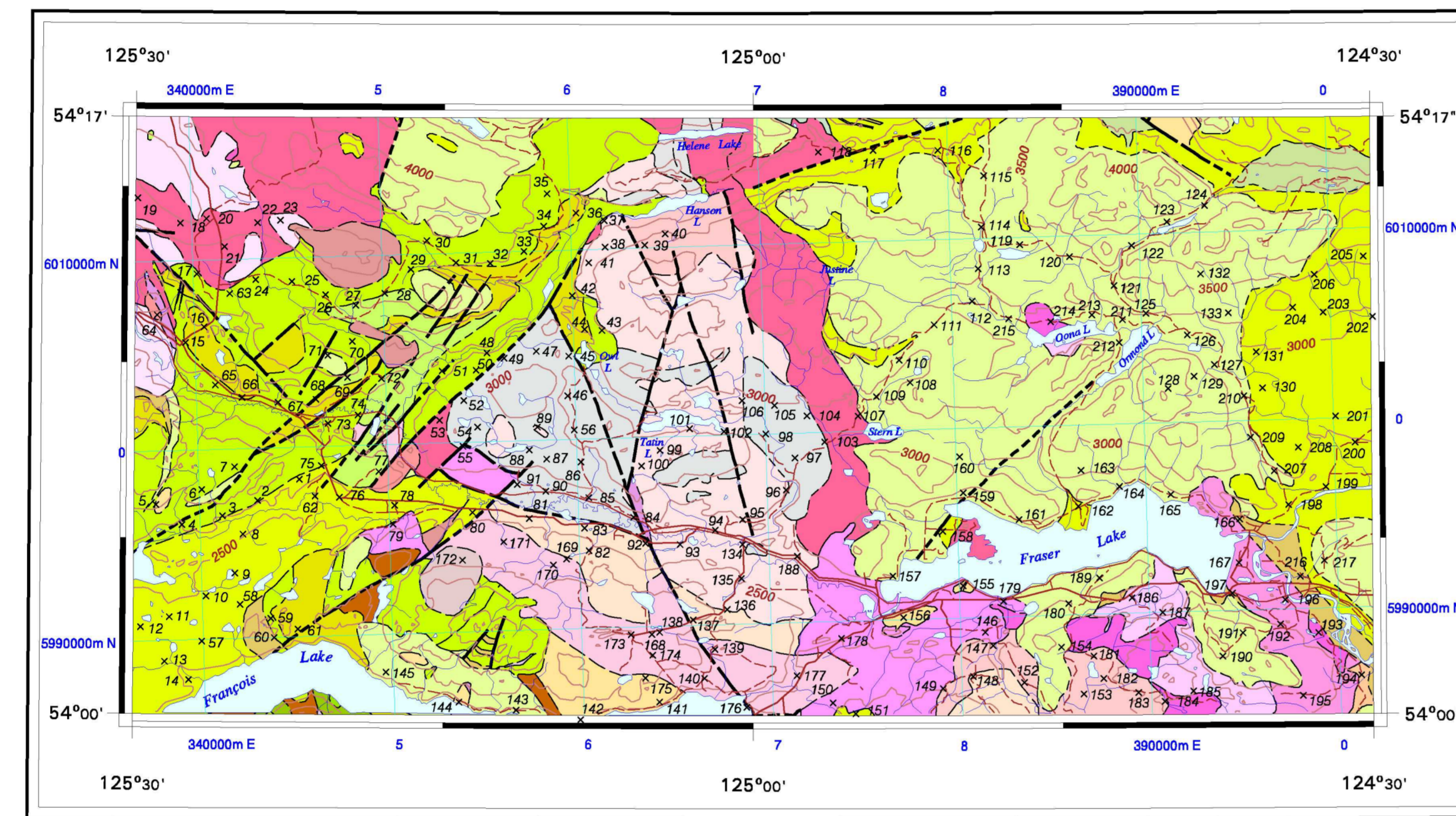


**NICKEL**

ICP-ES

ppm	Ni	Percentile
43	●	Maximum
36	●	98
31	●	95
25	●	90
19	●	75
13	●	50
< 2.0	●	Minimum

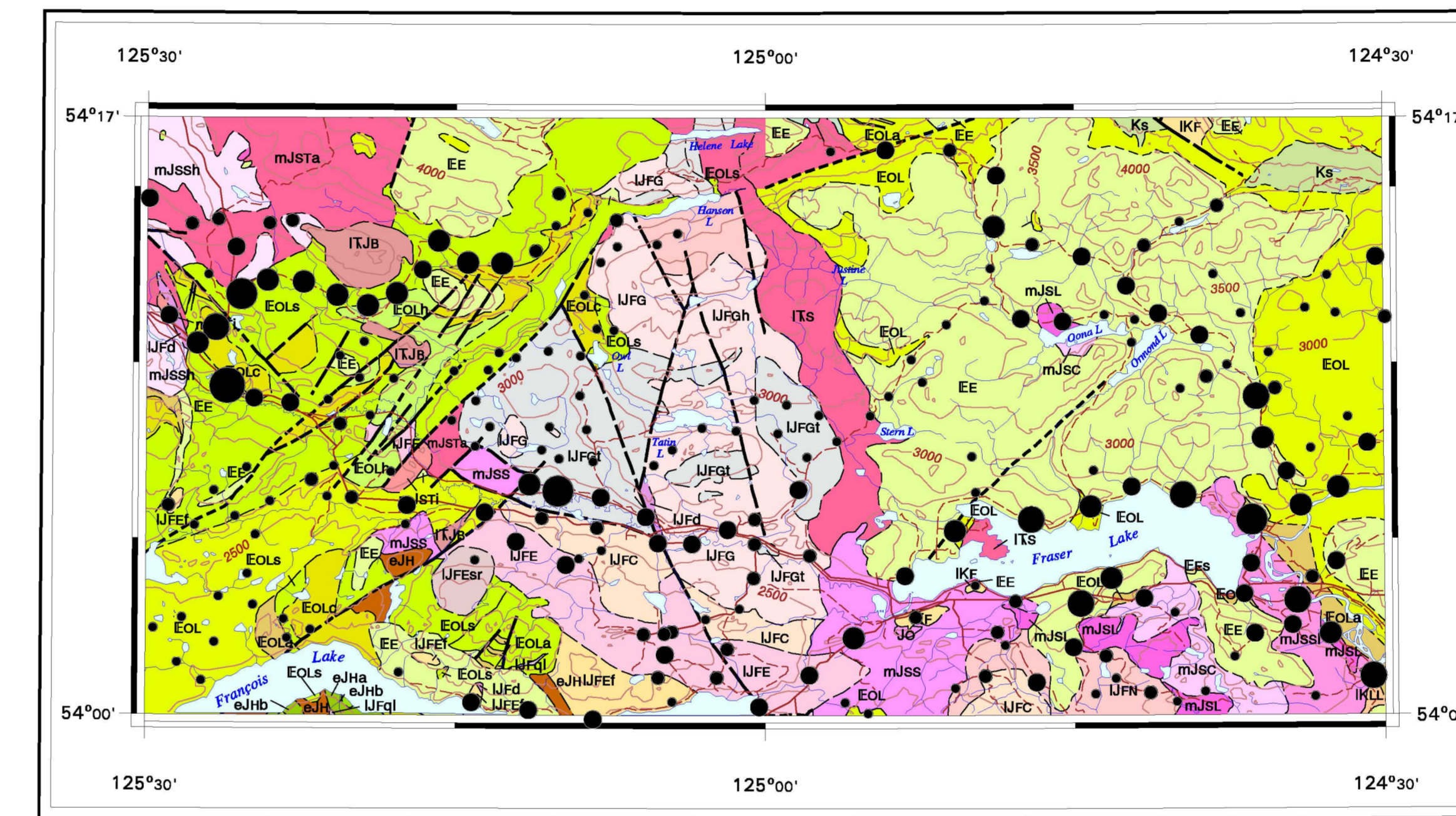
217 Samples  
Scalar Exponent = 1.0



**SAMPLE LOCATION MAP**

217 Sample Numbers

(All data for each sample are listed on separate sheets separately as Open File D36966)

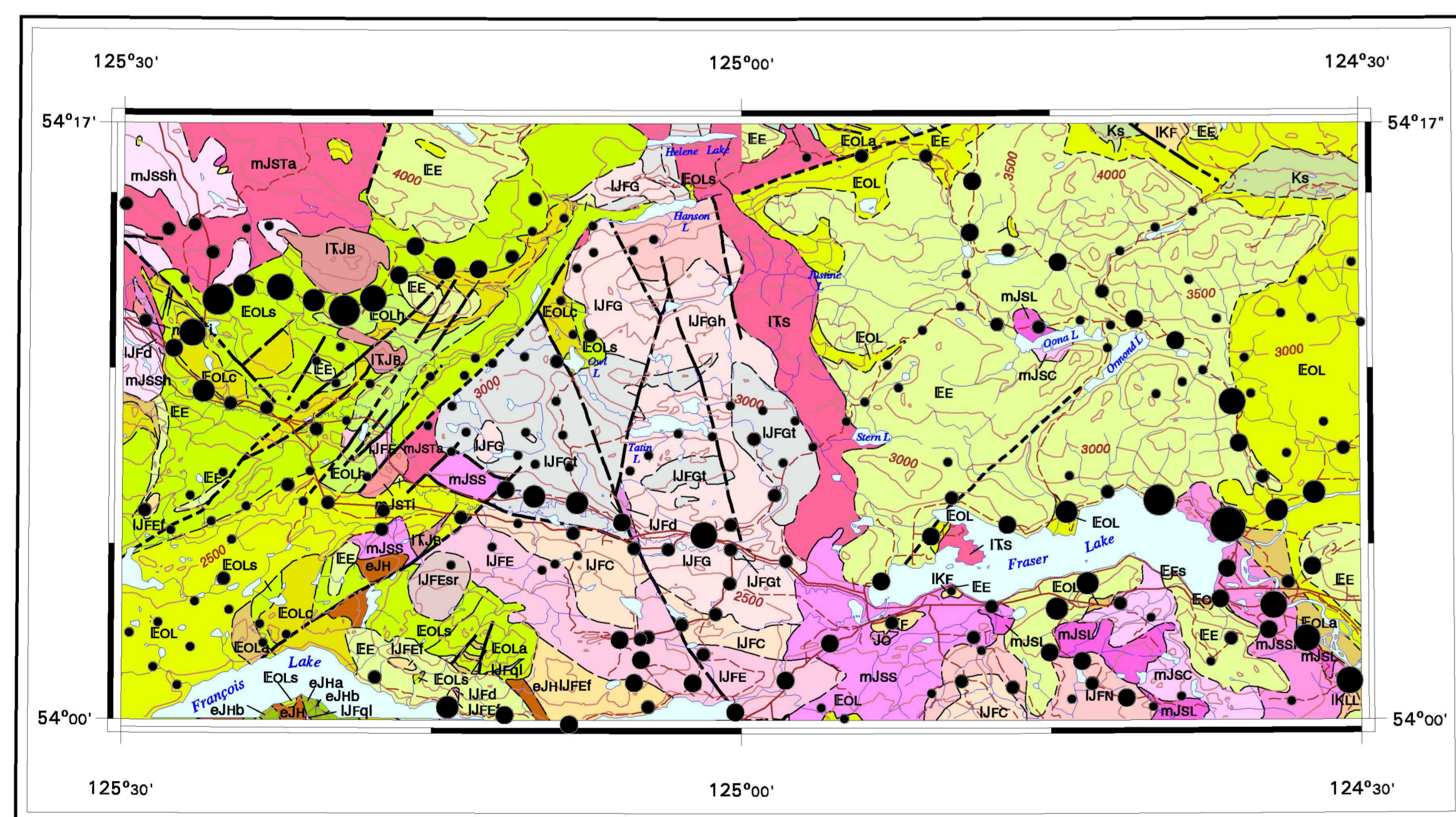


**COBALT**

INAA

ppm	Co	Percentile
18	●	Maximum
16	●	98
14	●	95
11	●	90
8	●	75
6	●	50
2.0	●	Minimum

217 Samples  
Scalar Exponent = 1.0

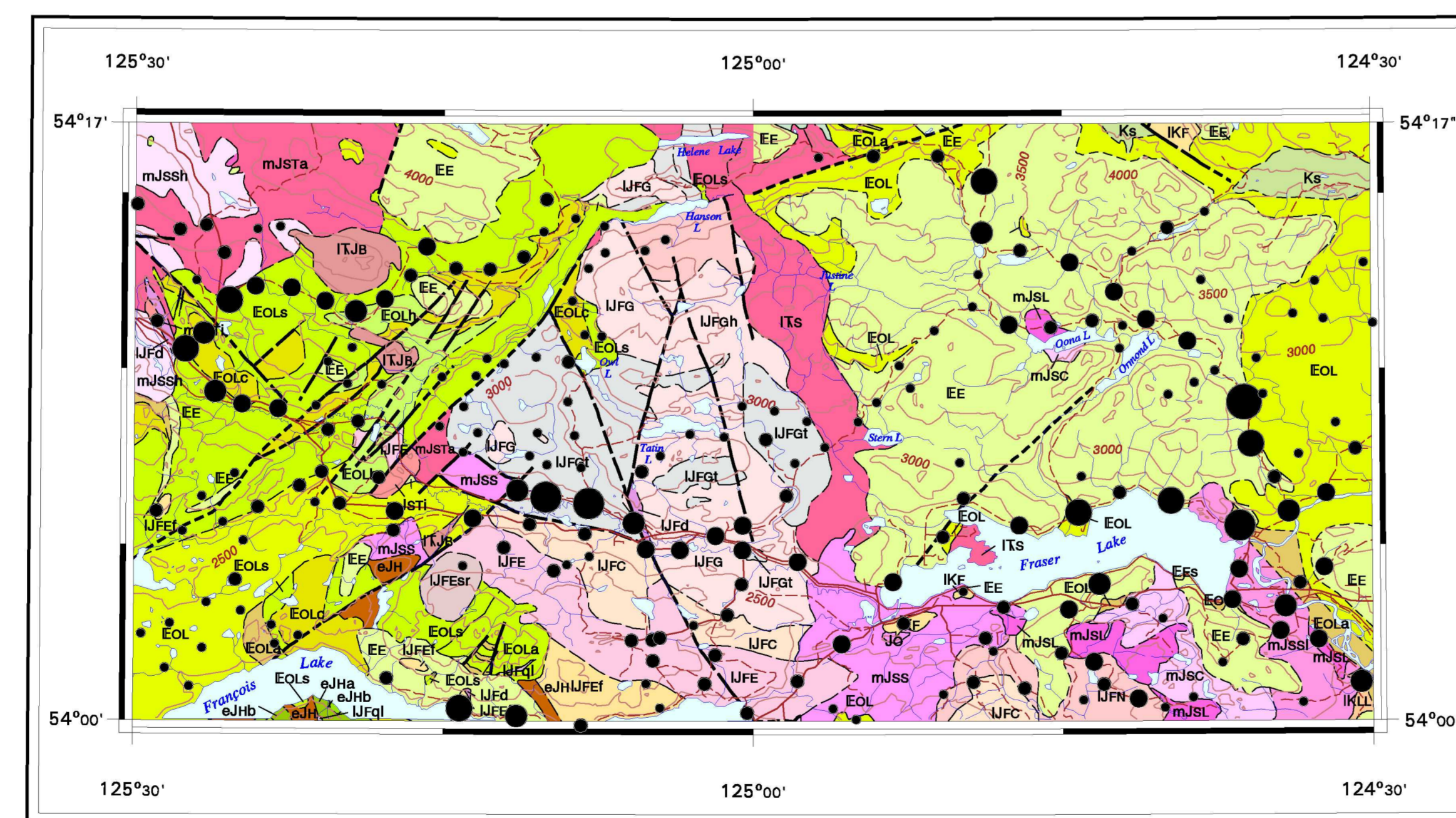


**VANADIUM**

ICP-ES

ppm	V	Percentile
89	●	Maximum
77	●	98
64	●	95
58	●	90
37	●	75
16	●	50
2.0	●	Minimum

217 Samples  
Scalar Exponent = 1.0

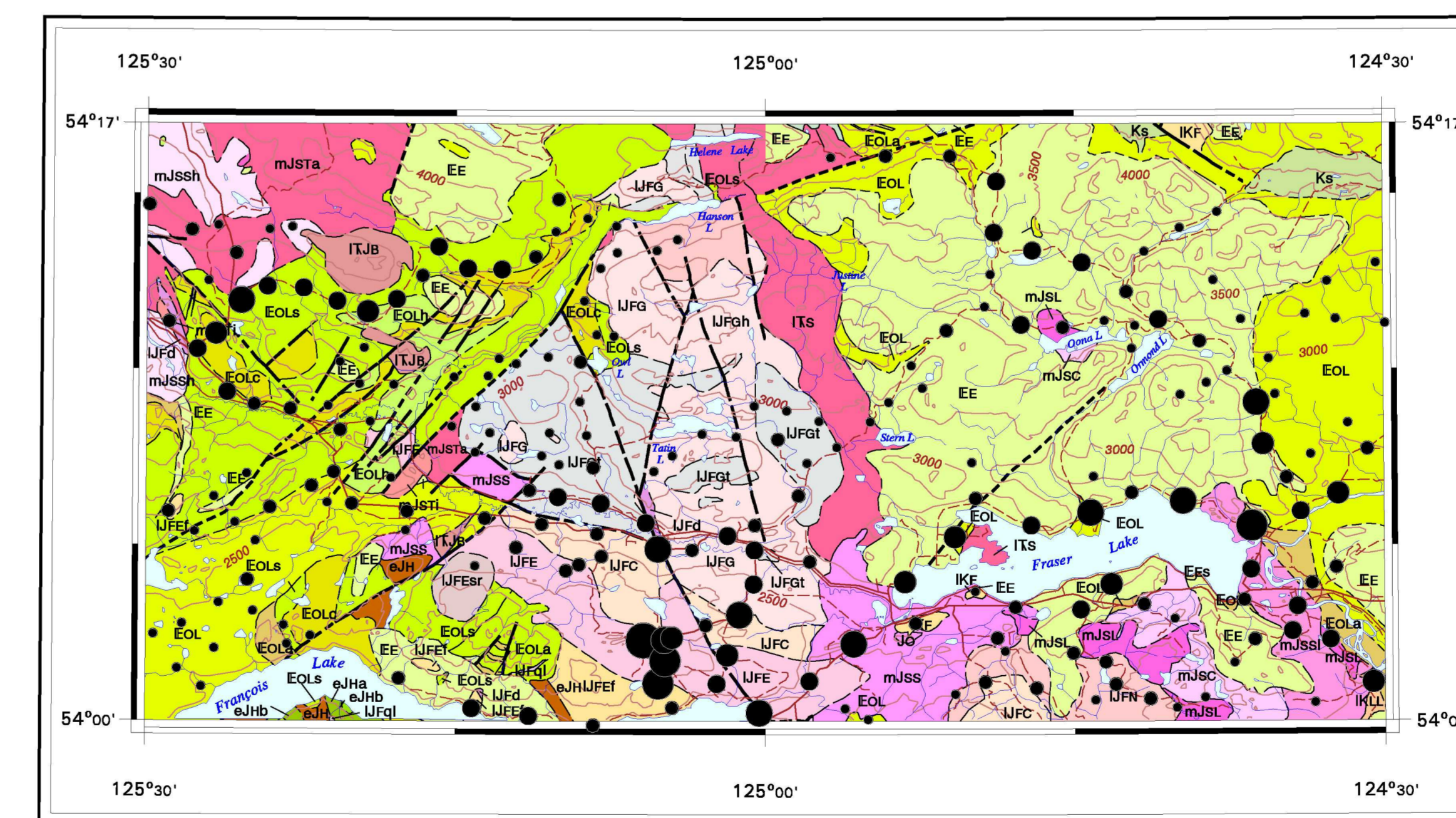


**CHROMIUM**

INAA

ppm	Cr	Percentile
62	●	Maximum
52	●	98
44	●	95
40	●	90
27	●	75
13	●	50
< 1	●	Minimum

217 Samples  
Scalar Exponent = 1.0



**LANTHANUM**

INAA

ppm	La	Percentile
80	●	Maximum
52	●	98
32	●	95
27	●	90
19	●	75
8	●	50
1.0	●	Minimum

217 Samples  
Scalar Exponent = 1.0

OPEN FILE 36966

## Biogeochemical Survey of the Fraser Lake Area Using Outer Bark of Lodgepole Pine (NTS 93K02/03)

### MAFIC SUITE OF ELEMENTS WITH THORIUM AND LANTHANUM

CENTRAL BRITISH COLUMBIA

Scale 1:400 000 - Echelle 1/400 000

Kilometers 0 10 20 30 40 Kilomètres

Transverse Mercator Projection  
Map Scale: 1:400 000  
© Her Majesty the Queen in Right of Canada, 1999

Projection: Transverse Mercator  
Map Scale: 1:400 000  
© Her Majesty the Queen in Right of Canada, 1999

Scene: view of the Fort Fraser Area Photo credit: C. Dunn  
Collection of outer bark Photo credit: C. Dunn  
Bark samples after reduction to ash Photo credit: C. Dunn

**NOTES**

This sheet of nine maps is one of three of similar format provided in this Open File. Grouped together are elements of certain affinities and/or similar distribution patterns. The three sheets are entitled:

- 1) Base Metals, Gold and Palladium Elements (Open File 36966)
- 2) Mafic Suite of Elements with Thorium and Lanthanum (Open File 36966)
- 3) Alkali metals, Alkaline Earths, Manganese and Aluminum (Open File 36966).

The elements depicted on these sheets are either those which show moderate geochemical data relief in the survey area, or they are representative of a closely associated suite of elements (e.g. of the rare earth elements, only La is shown because the distribution patterns of the other 7 determined (Ce, Eu, Lu, Nd, Sm, Tb, Yb) are almost identical). Full data listings of concentrations of these and other elements in each sample are given in the accompanying diskette. The diskette includes, also, a more detailed description of the methodology, analytical methods and analytical quality control.

**Rationale for Biogeochemical Surveys**

The roots of a single large tree extract elements from many cubic metres of soil, overburden, ground-water and sometimes bedrock. These elements are then transferred to aerial parts of the tree where they may become locally concentrated. In a multi-disciplinary survey program, data derived from the analysis of an appropriate vegetation sample medium permits geochemical mapping, with enhanced background to anomaly content of certain elements, which may assist both in mapping bedrock and in the search for concealed zones of mineralization.

Because each species of plant has a different requirement for, and tolerance to, a range of chemical elements, some partitioning of elements takes place and there is selective absorption and transference into the plants. For biogeochemical exploration, conifers provide suitable and effective sample media because they are primitive plants that have a wide tolerance to many trace elements. The outer bark is a repository for many elements that are not required for the metabolic function of the tree.

The geochemical information supplied by the vegetation is different from that of the soil. Just as two methods of geophysical survey may provide totally different information, so may two methods of geochemical survey. A high correlation between distribution patterns of two geochemical sample media is the exception rather than the rule. In geological environments where there is sufficient concentration of metals to form a mineral deposit, such a 'natural' metal of elements may be sufficient to generate biogeochemical anomalies above the mineral source (e.g. by upward diffusion) or close to it (e.g. by groundwater movement or movement in electrochemical cells). This, however, usually have geochemical anomalies displaced down-ice from the mineralized source. Such factors need to be taken into consideration when interpreting geochemical results.

**Reconnaissance Survey**

Lodgepole pine is the most common tree species in the Nechako area, and many metals concentrate in its outer bark. Hence, pine bark was selected as the sample medium for a reconnaissance-level biogeochemical survey in the Nechako project area. Samples were obtained by scraping approximately 100 g of outer bark from around the circumference of mature trees. The preferred sample interval was 2 km along roads, trails, and tracks. To minimize the effects of airborne contamination from roads, samples were collected at a minimum distance of 100 m from the forest. A few lesser used trails and tracks this distance was reduced to approximately 50 m. Within the survey area a two-person crew sampled trees at 217 sites within a 10-day period in early August 1998. Because of the lack of roads and trails in some areas, the sampling grid is not even. However, on average the sample coverage is approximately 1 site per 8 km<sup>2</sup>.

Bark samples were returned to the GSC laboratories in Ottawa where they were air-dried then reduced to ash by controlled ignition at 475°C for 24 hours. Ash samples were submitted for the analysis of 36 elements by instrumental neutron activation (INAA) at Activation Laboratories Ltd. (Ancaster, ON), and 30 elements by inductively coupled plasma emission spectrometry (ICP-ES) at Kamei Laboratories, Vancouver. The INAA analysis reports the total concentration of elements in the sample. The ICP-ES is performed on an acid digest of the ashes, and provides data on the total or near total concentrations of most elements that were determined. Data for some elements are obtained by both methods, but after review of the data from a quality control standpoint only the more accurate and precise set of values is recorded in the data listings.

**Map Production and Data Presentation**

The proportional dot maps are plotted using the Universal Transverse Mercator projection (NAD83 datum), with a central meridian of 120° (Zone 10). They were generated using AMI (ARC/INFO Macro Language). The macro, with its corresponding input menu, prompts the user to input percentile levels plotted, and an appropriate scaling exponent for each element to be mapped. Proportional dots are then generated, using the ARC/INFO SPOT/SIZE, POINT/PT and SPOT commands, with the user specifying an appropriate minimum and maximum dot size. Arguments for individual elements were chosen to provide the best view of the analytical data. Accordingly, care should be exercised when attempting to compare different elements plotted with different percentiles. Comments on distribution patterns are given in the digital file on the accompanying diskette.

**LEGEND**

<b>TERTIARY</b>	<b>Endako Group</b>	<b>LJFG</b>	Edmonton Phase: biotite granite and granodiorite, porphyritic
<b>EE</b>	andeste, basalt flows, breccia, vesicular amygdaloidal, tuff, hydrothermal, tuff	<b>LJFGI</b>	Fraser Lake Subphase: biotite granite, hornblende granodiorite
<b>EEs</b>	siltstone, sandstone, coal, tuff	<b>LJFGH</b>	Fraser Lake Subphase: biotite granite to quartz monzonite
<b>EOL</b>	Ootsa Lake Group (EOL - EOLh)		
<b>EOLa</b>	undifferentiated EOLa - EOLh		
<b>EOLa</b>	hydrothermal, rhyolite flows, breccia, tuff, locally flow banded; includes some felsic subvolcanic intrusions		
<b>EOLc</b>	andeste, basalt, diorite; includes some undifferentiated Ootsa Lake Group units		
<b>EOLs</b>	Savary Phase: diorite, mylonite and minor andeste and rhyolite		
<b>EOLh</b>	Hicks Hill Phase: diorite with phenocrystic biotite, hornblende, plagioclase		
<b>MESOZOIC</b>			
<b>Chocomaus</b>			
<b>Kasaska Formation</b>			
<b>JKK</b>	olive-green hornblende, plagioclase phytic diorite to diorite andeste; generally the phenocrysts have euhedral cores		
<b>Ks</b>	andeste plagioclase porphyry		
<b>IKF</b>	Fraser Phase: granodiorite, K-feldspar megacrystic		
<b>IKLL</b>	Lag Lake Phase: granite and granodiorite, pink, fine to medium grained		
<b>Late Jurassic</b>			
<b>LJLq</b>	Fraser Lake Suite (LJL - LJLh)		
<b>LJLqj</b>	quartz melt porphyry		
<b>LJLn</b>	NEW Phase: biotite granite to granodiorite, K-feldspar megacrystic		
<b>LJLd</b>	accolated porphyritic granite dykes		
<b>LJLp</b>	Casby Phase: granite, granodiorite		
<b>LJLr</b>	Endako Phase: biotite + hornblende granite to granodiorite		
<b>LJLrj</b>	Fraser Subphase: biotite granite to granodiorite		
<b>LJLrj</b>	Sam-Ross Creek Subphase: microcline granite		
<b>LA</b>	La		
<b>LAj</b>	La		
<b>LAk</b>	La		
<b>LAh</b>	La		
<b>LAi</b>	La		
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