

Newsletter for the Association of Exploration Geochemists

NUMBER 83 APRIL 1994

TECHNICAL NOTES

Hydrogeochemical surveys in Newfoundland - Geological mapping with REEs in lake waters.

A major component of the research carried out in the past several years by the Analytical Methods Development Section at the Geological Survey of Canada (GSC) has focused on the design and implementation of cost-effective methods to determine trace and ultra-trace elements at their "natural" (anthropogenic) levels in surface waters. Currently thousands of lake and stream sediments and their corresponding waters are collected across Canada under the federal-provincial National Geochemical Reconnaissance (NGR) Program in support of the exploration and mining industry. Unlike the extensive range of elements determined in the sediments, those measured in waters are generally limited to U, F, pH, Ca and Mg. This is a reflection of the priorities at the initiation of the program, during the "uranium boom days" of the 1970s, and at a time when analytical capability for other trace elements was poor. Since then, dramatic advances have been made in water analysis, mainly due to the establishment of the techniques of inductively coupled plasma emission spectrometry (ICP-ES) and mass spectrometry (ICP-MS). Both techniques enjoy such features as multi-element detection and a wide linear dynamic range (approximately 10⁵) but the sensitivity of ICP-MS is significantly superior to that of ICP-ES, generally by several orders of magnitude. Under a Mineral Development Agreement (MDA) with Newfoundland, the GSC has applied this new analytical methodology in a lake water survey in Bay d'Espoir in 1991 and a stream water survey in Bay Verte in 1992. This paper describes results from the former project for the rare earth elements (REEs).

Research to date on the distribution of the REEs in surface waters appears to have been concentrated almost exclusively in the area of marine geochemistry, more specifically on the degree to which river input and estuarine processes define the seawater composition of these elements (Hoyle et al., 1984; Goldstein and Jacobsen, 1988; Elderfield et al., 1990).

Ground water contents of the REEs have been measured at selected analogue sites in nuclear waste assessment studies to predict future mobility of both lanthanides and actinides. Miekeley et al. (1992) determined "dissolved" (empirically defined as <0.45 μ m) and particulate-held REEs in 42 near-surface and deeper ground water samples from the Osamu Utsumi uranium mine and the Morro do Ferro thorium and REE deposit in Brazil. The analytical methods employed by

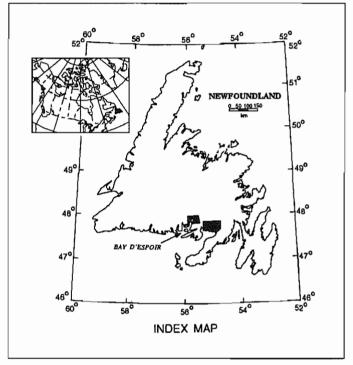


Figure 1. Bay d'Espoir location.

these workers are much too cumbersome and timeconsuming to apply on a large scale. Typically, from 2 to 20 litres of sample are collected and lengthy ion exchange procedures are invoked to achieve adequate detection levels.

The method developed at the GSC is based upon preconcentration and separation of the REEs, from 50 ml of sample, using chelation and analysis of the eluant by nebulisation ICP-MS. The resin used, marketed by Dionex Corp. (Sunnyvale, CA, U.S.A.), is known as the

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Information for Contributors to EXPLORE

Scope This Newsletter endeavors to become a forum for recent advances in exploration geochemistry and a key informational source. In addition to contributions on exploration geochemistry, we encourage material on multidisciplinary applications, environmental geochemistry, and analytical technology. Of particular interest are extended abstracts on new concepts for guides to ore, model improvements, exploration tools, unconventional case histories, and descriptions of recently discovered or developed deposits.

Format Manuscripts should be double-spaced and include camera-ready illustrations where possible. Meeting reports may have photographs, for example. Text is preferred on paper and 5-or 3-inch IBM-compatible computer diskettes with ASCII (DOS) format that can go directly to typesetting. Please use the metric system in technical material.

Length Extended abstracts may be up to approximately 1000 words or two newsletter pages including figures and tables.

Quality Submittals are copy-edited as necessary without reexamination by authors, who are asked to assure smooth writing style and accuracy of statement by thorough peer review. Contributions may be edited for clarity or space.

All contributions should be submitted to:

EXPLORE

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Information for Advertisers

EXPLORE is the newsletter of the Association of Exploration Geochemists (AEG). Distribution is quarterly to the membership consisting of 1200 geologists, geophysicists, and geochemists. Additionally, 100 copies are sent to geoscience libraries. Complimentary copies are mailed to selected addresses from the rosters of other geoscience organizations, and additional copies are distributed at key geoscience symposia. Approximately 20% of each issue is sent overseas.

EXPLORE is the most widely read newsletter in the world pertaining to exploration geochemistry. Geochemical laboratories, drilling, survey and sample collection, specialty geochemical services, consultants, environmental, field supply, and computer and geoscience data services are just a few of the areas available for advertisers. International as well as North American vendors will find markets through **EXPLORE**.

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NOTES FROM THE EDITOR

Despite claims of demise of the mineral exploration industry, technical notes continue to be submitted to EXPLORE. This issue features an article by Gwendy Hall and John McConnell on hydrogeochemical surveys in Newfoundland and an article by Reinhard Leinz and Don Hoover that updates us on developments with the CHIM method.

In the President's Message for this issue, Graham Taylor draws our attention to the changing times in the mineral exploration industry. This is a subject that the AEG Council has given considerable attention to recently. Most agree that the AEG must change in response to changing conditions in our industry in order to meet the needs of the membership, but the type and direction of change remains a topic of lively discussion. Some would have the AEG change its mandate and audience to the extent that a name change would be required, while others are basically happy with the organization as it was originally founded and is currently functioning. No AEG member should ignore this debate over fundamental issues. You are all urged to watch the pages of EXPLORE for continued discussion. Most importantly, be aware of any votes by the membership over the next several months and make sure your opinion is

Contributor's deadlines for the next four issues of EXPLORE are as follows:

<u>Issue</u>	Publication date	Contributor's deadline
84	July 1994	May 31, 1994
85 86	October 1994	August 31, 1994 November 30, 1994
87	January 1995 April 1995	February 28, 1995
07	April 1995	Tebruary 20, 1995

Owen Lavin
Editor EXPLORE



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PRESIDENT'S MESSAGE

During 1990 and 1991 a Committee chaired by Alan Coope worked very hard to produce a comprehensive "Report on the Identity of the Association of Exploration Geochemists." I refer you to EXPLORE No. 73 of October 1991 in which the nine recommendations of this Committee were detailed. Many of these recommendations have already been implemented and Council is presently discussing ways of initiating others such as Penrose-type lectures.



Geochemistry is still a major weapon in the armory of many companies as they expand their exploration activities into South America, the S.W. Pacific, S.E. Asia and West Africa, as is evidenced by the number of membership applications from geoscientists working in these areas. There is no doubt that this trend will continue as the alienation of prospective land and environmental considerations make exploration in Canada, USA and Australia less attractive. And yet, these three countries have, and still do, provide the majority of the membership of the Association.

What are these members now doing? Many of them are now involved in disciplines which are removed from our core business of exploration geochemistry. Some of them have ceased membership in the AEG because their career is no longer in the exploration industry. There are others, such as the petroleum geochemists, who are members of other scientific organizations but whose activities could be encompassed within an expanded AEG. It seems to me that a broadening of our charter would benefit both AEG and the many geochemists in the community.

A more immediate reason to reconsider our charter is occasioned by the present negotiations with Elsevier for the journal contract. There is potential to widen the scope of the Journal of Exploration Geochemistry to embrace disciplines other than exploration geochemistry and which may require a title change. I have already discussed these matters with the Negotiating Committee and there is general consensus within the committee that we should be at least considering some changes.

To aid this process I will prepare a discussion paper for the next issue of **EXPLORE**. In the meantime, I urge you to reread **EXPLORE** No. 73 and provide me with your vision for AEG in the 21st Century.

The ballot to change the By-laws to reflect coincidence of the term of office of the Executive and Councilors with the financial (calendar) year was overwhelmingly in the affirmative. The present Executive and Council will hold office until 31 December 1994. It is my desire that as we approach our 25th year, we can do so in the knowledge that the AEG is vibrant and assured of another quarter century of service to geochemistry.

Graham F. Taylor AEG President

ANNUAL GENERAL MEETING

As many of you know, the scheduled Annual General Meeting (AGM) of the Association of Exploration Geochemists had to be canceled because our President, Graham Taylor, had to return to unexpectedly to Australia, and the Association's finances had not been audited. We have rescheduled the AGM for April 27, 1994. Because of these unusual circumstances the AGM will be held via conference telephone in both Ottawa, Ontario, Canada and Denver, Colorado, USA. In Ottawa the conference call will be at 5PM EDT at the Geological Survey of Canada, 601 Booth Street and in Denver at 3 PM MDT at the Denver Federal Center, Building 20, Branch of Geochemistry, Room 1111. All available members are encouraged to attend the AGM if at all possible.

Sherman P. Marsh Secretary



NEWS RELEASE

Geologic Tours in the Las Vegas Area

The Nevada Bureau of Mines and Geology, in cooperation with the U.S. Bureau of Land Management and Department of Geosciences at the University of Nevada, Las Vegas, has published *Geologic Tours in the Las Vegas Area* (NBMG Special Publication 16), written by Becky Weimer Purkey, Earnest M. Duebendorfer, Eugene I. Smith, Jonathan G. Price, and Stephen B. Castor.

This 156-page book provides a geologic guide for interested local citizens and tourists as they drive the main roads, visiting the many sites of natural beauty and historical importance in the Las Vegas Valley area. Road logs for five trips are provided.

The guide is available for \$12.00 at the NBMG sales office (Room 310, Scrugham Engineering-Mines Building, University of Nevada, Reno) or for \$14.00 by mail from Nevada Bureau of Mines and Geology/178, University of Nevada, Reno NV 89557-0088. Mail orders must be prepaid by check or money order made out to "Board of Regents." Telephone orders (702-784-6691), using Visa or MasterCard, are also accepted.



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NO CHEMICALS AND FEW STEPS PRODUCES ACCURATE AND PRECISE RESULTS PAGE 4 NUMBER 83 EXPLORE

Technical Notes

Continued from Page 1

MetPac CC-1 column and is similar to Chelex-100 in that it is based on iminodiacetate functionality. In the pH range of 5-6, the resin selectivity is optimized for retention of the REEs and transition metals relative to the alkali and alkaline earth elements. By using an ammonium acetate eluant in this pH range, alkaline earth elements can be eluted while the REEs (and transition metals) remain strongly bound. The resin does not concentrate anions such as halides, nitrate, sulphate, phosphate or organic anions. It should be noted that it is the free REE cation being chelated; elements bound as charged or neutral inorganic or organic complexes, or in colloidal form, are not preconcentrated. Nitric acid is the preferred eluant (at 0.8 M) for analysis by ICP-MS as isobaric spectral interferences are minimized. The preconcentration procedure was designed to be fully automated, for operator-free functioning overnight, so that the following morning a tray of 40 samples is ready for nebulisation into the ICP mass spectrometer. A preconcentration factor of 10 was selected, but this can be increased or decreased easily to suit the needs of the particular project.

The ICP mass spectrometer employed was the Perkin-Elmer/SCIEX Elan model 250, upgraded to the 500. Corrections were made for the isobaric interferences of the light REE oxides on the heavy REE isotopes. The relative standard deviations (RSD) of synthetic water standards taken through the procedure were in the range of 1-2 % (n=5).

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FOR THE LAST 10 YEARS, OVER 3 MILLION SAMPLES WERE ANALYZED

Results for six replicate applications of this method to 50 ml aliquots of Ottawa River and Meech Lake control samples are given in Table 1. Note the excellent RSDs: for Ottawa River varying from ±3% for Pr at 31 ng l-1 (ppt, parts-per-trillion) to $\pm 17\%$ for Tb at only 1.8 ng l⁻¹; and for Meech Lake ranging from $\pm 2.3\%$ for La at 33 ng l⁻¹ to a maximum of $\pm 21\%$ for Tb at 0.7 ng l⁻¹. Not surprisingly, higher RSDs tend to be associated with those elements present at very low concentrations. With the exception of Gd in Ottawa River, RSDs greater than ±10% occur for those elements whose concentrations are below 5 ng l-1, namely Sm, Eu, Tb, and Lu. Also shown in Table 1 are control data for a bulk tap water control which has been analyzed 27 times (i.e., 27-50 ml amounts) over a period of approximately one year. Ottawa River is used for the supply of tap water, hence its similar concentrations. Note that precision has not degraded over time when these results are compared to those for the six replicates of Ottawa River carried out consecutively during the initial development of the method. Detection limits are in the range 0.2-1 ng l-1 using a definition based upon that concentration equal to three times the standard deviation obtained in five replicate analyses of a 5 ng l-1 synthetic standard solution. We have reported all survey data to lower limits of 1.0 ng l-1 as the noise encountered with the ICP-MS measurement fluctuated with its performance (aging instrument) from day to day.

Table 1. Reproducibility of results for the REEs in: Ottawa River (n=6); Meech Lake (n=6); and tap water (n=27). All in ng l⁻¹.

	Ottawa	River	Meech	Lake	Tap w	Tap water	
Element	mean	SD	mean	SD	mean	SD	
La	158	8	33	0.8	159	9	
Ce	154	9	42	1.7	156	8	
Pr	31	1	7	0.2	32	2	
Nd	119	6	29	2.4	120	8	
Sm	17	0.8	5	0.6	18	1	
Eu	3	0.4	7	0.3	3	0.4	
Gd	14	2	5	0.8	15	3.7	
ТЪ	1.8	0.3	0.7	0.15	2.0	0.5	
Dy	11	1	4	0.4	11	1	
Ho	2	0.1	1	0.1	3	0.1	
Er	8	0.3	3	0.2	8	0.7	
Tm	1	0.1	0.5	0.05	1	0.1	
Yb	6	0.6	4	0.3	7	0.8	
Lu	1	0.2	0.8	0.14	1	0.2	

SD: Standard Deviation

The lake and stream water survey was carried out in southern Newfoundland, north of Fortune Bay and east of Bay d'Espoir (Fig. 1). A total of 136 lake water samples, including 11 field (site) duplicates, were collected in one day using a float-equipped helicopter. Samples were collected about 20 cm below the surface after initial rinsing of the bottle with the lake water. Of these lakes, 108 had been sampled for the organic-rich centre lake bottom sediment in an NGR survey in 1980 (Friske and Hornbrook, 1991). Within 24 hr. of collection, samples were filtered through a 0.45 μ m filter paper into a 250 ml bottle, using Millipore glass

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Technical Notes

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Table 2. Field duplicate data for REEs in lakes and streams in Bay d'Espoir. All measurement in ng l-1

Sample	La	Ce	Pr	Nd	Sm	Eu	Tb	Gd	Dy	Ho	Er	Tm	Yb	Lu
91-54	186	440	46	183	36	7	5	31	23	5	16	1	7	1
91-55	196	468	48	192	36	8	5	36	23	5	16	2	7	1
91-61	131	303	34	133	25	6	4	26	21	5	15	2	9	1
91-62	130	306	34	136	29	6	4	27	21	5	15	2	10	1
91-80	31	46	7	26	5	2	1	7	5	1	3	<1	3	<1
91-81	34	52	7	30	7	2	1	7	5	1	3	<1	3	<1
91-94	118	122	24	87	15	3	2	16	8	2	4	<1	2	<1
91-95	97	104	21	78	14		2	13	7	1	4	<1	2	<1
91-105	135	116	26	94	18	3	2	16	10	2	5	1	4	<1
91-106	139	123	27	99	19	4	2	18	10	2	5	1	4	<1
91-131	70	134	15	57	10	1	2	12	11	3	10	1	12	2
91-132	46	91	10	39	7	1	1	7	8	2	7	1	7	1
91-141	33	48	9	31	8	<1	2	8	12	3	11	2	15	2 3
91-142	43	64	11	42	10	<1	2	13	16	4	14	2	19	
91-146	199	188	50	197	50	2	13	59	76	20	70	12	87	14
91-147	204	189	53	192	47	2	12	61	74	20	71	12	91	14
91-150	228	324	57	228	54	3	12	57	65	17	56	9	65	10
91-151	214	319	56	221	53	4	12	66	69	17	57	10	70	10
91-175	202	277	50	177	34	2	8	39	48	12	39	7	43	6
91-176	200	272	51	177	40	2	8	46	49	12	40	7	47	7
91-179	107	152	27	100	23	2	6	33	39	10	34	6	42	6
91-180	105	141	26	97	25	1	5	29	38	10	35	6	42	6

apparatus and acidified with 1 ml of 16 M HNO₃ (Ultrex). Earlier studies indicated that: (1) acidification to 0.1 or 0.4 % in HNO3 stabilized the REEs in solution at ng l-1 levels for at least two months, and (2) the Nalgene bottles used "as is" rather than acid-washed did not lead to contamination problems. About 50 elements, including the REEs, have been determined in these waters from Bay d'Espoir.

Data for the REE contents of 11 field duplicate lake water samples are given in Table 2. Precision appears to be excellent, even for the less abundant REEs, and is not noticeably inferior to the analytical precision shown previously by the control data (Table 1). A broader evaluation of precision is obtained by combining duplicate data from several concurrent hydrogeochemical projects and representing the enlarged dataset graphically as "Thompson and Howarth" plots, as has been done for La, Sm, Er and Yb in Figure 2. In this approach, the absolute difference between the duplicate pair is plotted against their mean on a log-log scale and a 95th percentile line for a selected precision (twice the RSD) is then drawn according to the pattern of the data (Thompson and Howarth, 1978). Thus, the 95th percentile for the selected precision on each graph is an upper confidence limit (i.e., the "worst" case precision value) above which no more than 5% of the points should lie. Field duplicate data (n=23 pairs) have been distinguished from blind (lab) duplicate data (n=27), the former being labelled as stars and

the latter as circles in Figure 2. Although greater variability would be expected in results for the field duplicates, rather than for sample splits, this was not apparent from the graphs and therefore data were viewed as one group. Note that numerous points fall on or just below the x-axis, indicating that there was no difference in values obtained for the duplicates. Hence, at concentrations in the range from several ng l-1 to several hundred ng l-1, a field and analytical precision of 25-30% (i.e. ± 15% RSD) or better can be

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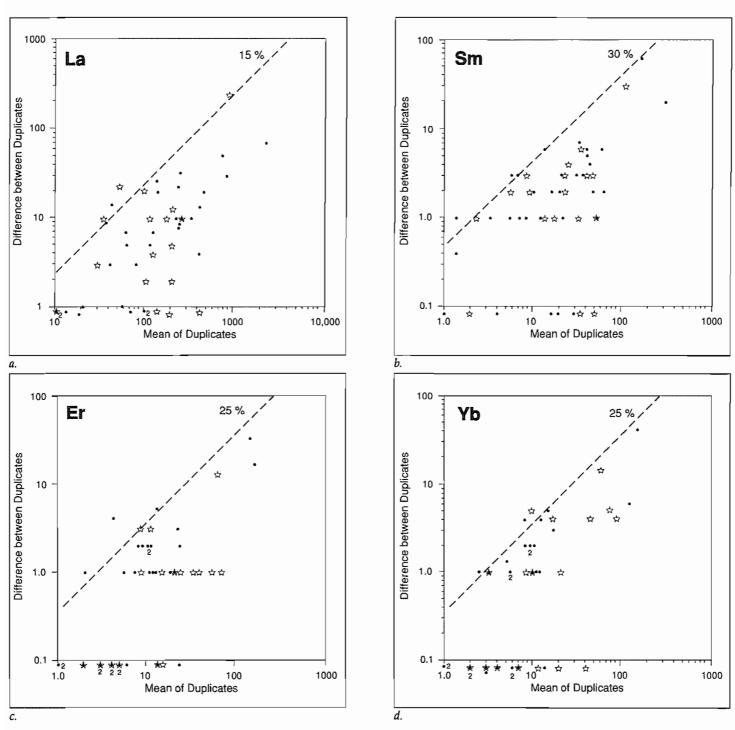


Figure 2. Thompson and Howarth plots representing precision for: (a) La; (b) Sm; (c) Er; and (d) Yb. Values in ng l⁻¹ (ppt). Stars represent field duplicates, and circles represent blind (lab) duplicates.

expected for Sm, Er and Yb. Superior precision of 15% is shown for the more abundant La, present in these 50 samples in the concentration range of 10-2400 ng l⁻¹.

Summary statistical data for the REEs in the 135 lake waters are given in Table 3. As mentioned previously, results were available for some of the REEs in the centre lake bottom

sediments of most of these lakes from the National Geochemical Reconnaissance (NGR) database. The elements La, Ce, Sm, Eu, Tb, Yb and Lu had been measured in the minus 180 μ m fraction by instrumental neutron activation analysis (INAA) in a multi-element commercial package.

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Technical Notes

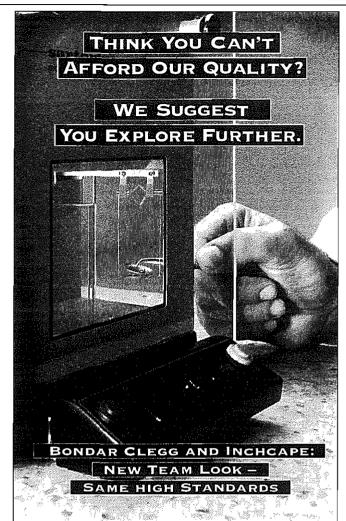
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lable 3. Summary statistics for REE concentrations in 136 lake vaters in Bay d'Espoir. All values in ng l⁻¹

lement	Range	Median	Mean	SD
Ja	10 - 883	130	148	119
Ce Ce	12 - 766	208	233	152
r	2 - 139	29	35	23
٧d	8 - 489	113	132	86
im	2 - 83	23	27	17
3u	<1 - 16	2	3.3	2.7
Лъ	<1 - 17	4	4.9	3.6
3d	2 - 96	23	29	20
Эу	2 - 112	21	27	22
-To	<1 - 29	4	6.4	5.6
3r	1 - 99	15	21	19
ľm	<1 - 17	2	3.1	3.1
ľb	1 - 113	10	20	22
.u	<1 - 17	1	2.9	3.4

There was a problem with the INAA data for Lu, mostly eported erroneously low at the method's detection limit of 1.02 μ g g⁻¹ and therefore comparison with the water data vas not possible. Spearman rank correlation coefficients between the two media for the 108 coincident sites are: 0.38 or La; 0.45 for Ce; 0.48 for Sm; 0.31 for Eu; 0.60 for Tb; and 1.76 for Yb. Data for Eu in both sediment and water are probably too close to the method's detection limits to have confidence in this correlation. This is not the case for the other five REEs. A correlation coefficient greater than 0.40 is ignificant at the 99.9% confidence level. Clearly, a strong correlation is present for Yb in the two media and that for Tb s high, despite the low and restricted concentration range, rom <1 to 17 ng l⁻¹ of Tb, in the lake waters.

Examples of the distribution patterns in both lake water ind sediment media are shown for Yb, Tb and La in Figure 1. The survey area is divided into two contrasting geological errains separated by the northeast trending Hermitage Bay ault. To the southeast, the older Avalon Zone comprises predominantly volcanic, sedimentary and intrusive rocks of Hadrynian age which are intruded by a Devonian high-silica granite. To the northwest, the Gander and Dunnage Zones ire composed of clastic-sedimentary and volcanic rocks, also ntruded by granites (Colman-Sadd et al., 1990). The earlier VGR sediment survey showed the area to the northwest of he fault to be higher in the elements As, Ba, Co, Cu, Cs, Ni and Sb, and lower in F, U and the REEs in comparison to the irea to the southeast. The similarities in both sediment and water maps for Yb, and to a lesser extent, Tb are striking Fig. 3). The north-western area of both media maps has considerably lower contents of Yb and Tb than the southeast, reflecting the contrasting geology - the west being iominantly fine-grained siliciclastic terrain and the east nainly granitic. The southern section of the area, east of the fault, elevated in REE concentrations (dark shading), is anderlain by a late marginal phase of the granite that is



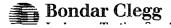
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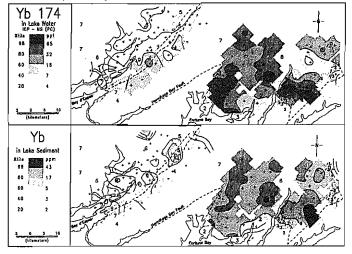
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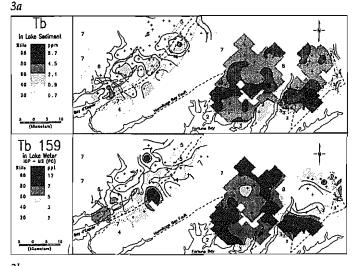


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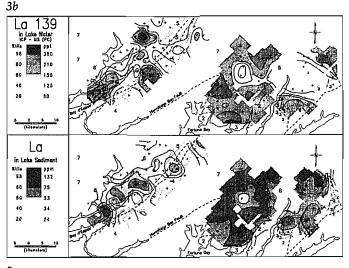


Figure 3. Elemental contour maps for (a) Yb; (b) Tb; and (c) La in lake water and lake sediment. Geology is as follows: 1: granite; 2: bimodal volcanic rocks; 3: siliciclastic sedimentary rocks; 4: clastic and volcanic rocks; 5: felsic and mafic volcanic rocks; 6: siliciclastic sedimentary rocks; 7: granite; 8: high-silica granite.

region north-west of the fault, there appears to be a ridge of high REEs extending north-east from Bay d'Espoir parallel to the stratigraphic and structural trends and is likely due to REE enrichment in one or more stratigraphic units. The single high in the extreme northwest is evident in Figures 3b (Tb in water) and 3c (La in water) should be disregarded as it is a small lake near a road and has anomalous concentrations of SO_4 (12.5 mg I^{-1}) and Cl (13.2 mg I^{-1}).

Another way to compare water and sediment information for the REEs is to examine their chondrite-normalized plots; these were drawn for all 108 coincident sites. For every lake, the similarity between the plots for REEs in water and sediment is remarkable. Forty seven sites are clearly located over the biotite-bearing Ackley granite and their chondritenormalized plots are typified by those for samples 126 and 150 in Figure 4a. All show distinct Eu negative anomalies, flat heavy REEs patterns and various degrees of enrichment in the light REEs (e.g., greater for sample 126 in Fig. 4a). Weak negative Ce anomalies in the waters usually correspond to slight Ce enrichments in the sediment, probably indicating oxidation of Ce3+ to Ce4+ and subsequent precipitation of the less soluble species. There is no apparent dependence of the shape of the plot on the textural classification of the granite, that is, as fine or coarse grained, equigranular or porphyritic. Plots for waters collected within siliciclastic sedimentary drainage basins resemble those shown in Figure 4b for samples 47 and 94, overlying the Riches Island Formation of green-grey pelite and siltstone with graphitic slate, quartzite and psammite. These patterns exhibit a very gradual decline from the light to heavy REEs and are devoid of features such as significant Eu or Ce anomalies. Rather similar to these are patterns for waters collected over a semi-pelitic and psammitic paragneiss (samples 69 and 70 in Fig. 4c), but here more distinct Eu anomalies are shown by the sediments and not by the waters. Signatures of the lake waters and sediments overlying felsic and mafic volcanics are shown by samples 158 and 159 in Figure 4d. Again, light REE enrichment is evident, together with Eu negative anomalies, though of much less intensity than shown earlier by the granitic lithology. These findings suggest that: (1) there is little fractionation of the REEs as they become fixed by the lake sediment; and (2) the elemental patterns in the waters may be quite stable over time, though overall concentrations undoubtedly do change (e.g., with climatic conditions).

In summary, application to 135 lakes in Bay d'Espoir has demonstrated the potential of REE signatures in lakes, streams and ground waters to be used to trace fluxes and identify their origins. There appears to be negligible fractionation of the REEs during processes of dissolution and precipitation, as evidenced by the similarity between chondrite-normalized plots of lake water and matching sediment and by the distinct features of these plots within lithologies. The excellent productivity of this method, at about 100 samples a day, and the sample volume requirement of only 50 ml or less will ensure a rapid growth in information in the future on the geochemical cycling of the REEs. A later report on this area will describe the application of this and other new analytical methodologies to examine the potential of stream chemistries in exploration for base metal and Au mineralisation in the Bay Verte Peninsula, Newfoundland. Continued on Page 9 XPLORE NUMBER 83 PAGE 9

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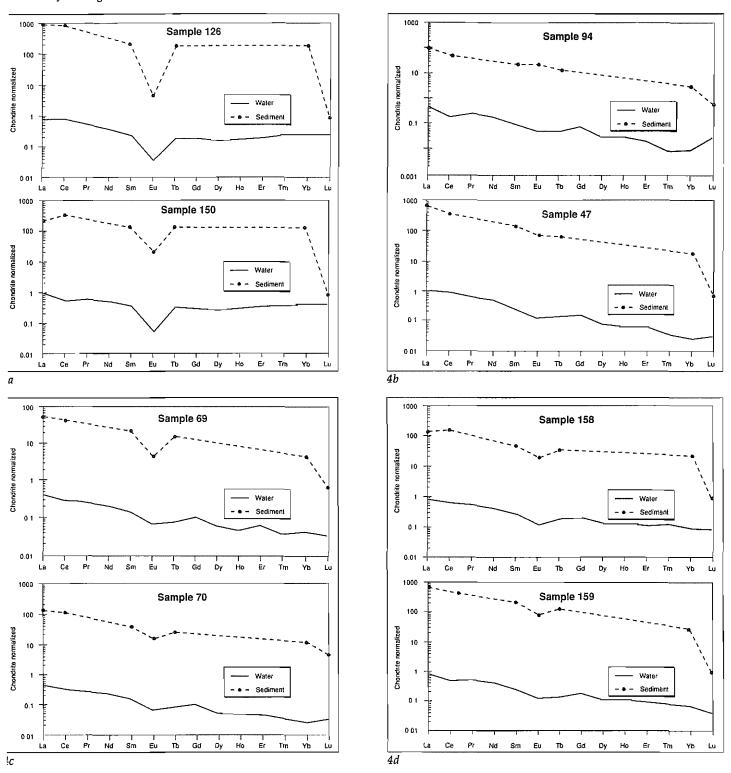


Figure 4. Chondrite-normalized plots for (a) samples 126 and 150 over Ackley granite; (b) samples 47 and 94 over pelite and siltstone; (c) samples 69 and 70 over semi-pelitic and psammitic gneiss; and (d) samples 158 and 159 over felsic and mafic volcanics. [Lu in sediment is rroneously low]

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Acknowledgements

This work was funded under the Canada-Newfoundland Cooperation Agreement on Mineral Development 1990-1994. The authors are grateful to J.E. Vaive and J.C. Pelchat of the Analytical Method Development Section of the GSC for superb analyses, and to Chris Finch for his participation in the fieldwork.

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Ideal CHIM with the Newly-Developed, NEOCHIM Electrode

Review

CHIM is an electrogeochemical exploration method developed more than twenty years ago in the former Soviet Union where it has been used successfully for locating buried mineral deposits (Goldberg et al., 1990). In principle, CHIM uses an electric field to cause ions at depth to migrate into special collector electrodes. Russian investigators stated that CHIM effectively transported cations from a buried deposit through 100 meters or more of overburden to the electrodes placed on the soil surface. They hypothesized that the cations were transported mainly by the strong electric field applied in CHIM. There is little mention in the Russian literature regarding the transport and collection of anions.

We recently reported results in this newsletter (Leinz and Hoover, 1993) that indicate that ions collected by conventional CHIM are, for the most part, generated in situ in chemical reactions proximal to the electrode and that they are transported into the electrode predominantly by diffusion. We attribute these results to the fact that conventional CHIM electrodes, as described by Goldberg et al. (1990) and Smith et al. (1993), permit the loss of acid from the electrode to the soil by diffusion. Acid is used in the conventional electrode to neutralize hydroxyl (OH⁻) ions produced in the electrode by electrolysis during operation $(2H_2O + 2e^- \rightarrow 2OH^- + H_2\uparrow)$. Generally, nitric acid is used. Goldberg et al. (1990) call for the addition of concentrated HNO₃ to the electrode, as necessary, to prevent alkalization of the soil by OH ions. Such alkalization would impede the mobility of cations.

Problems caused by acid loss

Ideally, CHIM should affect the transport of only those ions soluble in the soil moisture. The presence of a large quantity of H⁺ ions proximal to the electrode promulgates conditions that are not permissive for conducting ideal extractions because: 1) the H⁺ ions react with soil matter, releasing cations which are otherwise not available for electrical extraction. The acid-soluble ions may or may not be associated with buried ore; 2) the H⁺ ions are transported back into the electrode by the electric current in preference over other cations. Our tests with the conventional electrode indicated that cations other than H⁺ account for a maximum of only 0.1% of the charge transfer delivered by the current. Most of this amount is likely due to diffusion; and 3) the large number and high mobility of the H⁺ ions also constrain the volume of soil sampled. Of the mobile cations drawn towards the electrode by the current, few enter the electrode. Most of them accumulate in the zone flooded with H+ from which the H⁺ is preferentially transported.

We recognized that the loss of acid from the conventional electrode was not permissive for conducting the ideal extractions possible of CHIM, at least in principle.

Subsequently, we developed an electrode that overcomes the problem. We have tested the new electrode and are encouraged by results that we feel are closer to those expected of ideal extractions.

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The new electrode briefly described

Our new electrode (patent pending, 1993, Fig. 1) is comprised of two chambers, an outer chamber containing an electrolyte fluid, which receives the collected ions, and an nner chamber containing an electrolyte fluid in which a graphite or metal working electrode is immersed. The two chambers are separated by a conductive salt bridge. As with the convential CHIM electrode, electrical contact is made through a disk of synthetic parchment sealed into the bottom of the outer chamber. The salt bridge is a gel salt solution. It is held in place at the bottom of the inner chamber by a econd parchment disk. The salt bridge has a two-fold unction: 1) to allow the passage of current from the inner to the outer chamber; and 2) to prevent undesirable ions in the nner chamber from entering the outer chamber. This design an function as either an anion or cation collector.

Besides the requirement of conductivity, there are few onstraints on selecting the composition of the fluids and the alt bridge. The parchment membrane is permeable to ions, o not only can the electric field drive ions through but liffusion of ions across the membrane can occur as well. During operation of the electrode as a cation collector, ations from the soil are brought into the outer chamber fluid and anions present in the fluid migrate out into the soil. The everse happens when the electrode is used as an anode ollector. Because of this, consideration of the reactivity of

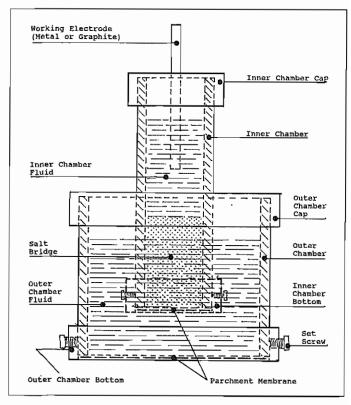


Figure 1. The NEOCHIM electrode, patent pending, by Hoover and Leinz, 1993. The prototype is constructed from chemical-resistant plastic.

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he soil with the ions in the fluid is necessary. In an ideal HIM run, the ions moving out of the electrode are passive owards soil matter and serve only to conduct current. The luids in the two chambers and the salt bridge normally hould be unreactive towards each other and to the products f electrolysis in the inner chamber. Finally, the ion oncentration of the fluids and the salt bridge must be ufficient to balance the charge transfer of the applied urrent.

lesults of field tests

We tested our new electrode, which we have named the JEOCHIM electrode, in four separate experiments at the Cokomo mine in the Central City district, west of Denver, Colorado, where we conducted some of our previous CHIM xperiments (Smith et al., 1993; Leinz and Hoover, 1993). The four experiments were conducted using both cation and nion collector electrodes in each run. Electric power was rovided by a 10kW generator with 800 volts applied to the lectrodes. Line and site placement were the same as for the revious CHIM experiments (Smith et al., 1993). Site 00 is ocated over a gold-bearing, base-metal vein concealed by 3 neters of transported overburden. Cathodes and anodes were separated by a distance of about 3 meters. The arameters for each of the experimental runs are summarized a Table 1.

The quantity of ions collected is dependent on the total harge transfer supplied to the electrodes by the current. 'he amount of charge transfer depends on the duration of ne run and the current at each electrode. For the most part, 'able 2 shows that the quantity of Cu, Pb, and Zn collected acreases with time as expected. Normalizing the data to mpere-hours reveals that strikingly similar amounts of zinc re collected per unit charge in each of the four runs (Fig. 2), vivid demonstration of comparable efficiencies of the lectrical extractions. This is near the result expected of ideal HIM. The data for copper and lead depicted in Figure 2, hough not as striking as the zinc data, reflect reproducibility reviously not achievable with conventional CHIM elecrodes. Deviations of Cu from the norm in run IV (Fig. 2) are ttributed to the effects of chloride entering the soil from the athode fluid or, in some cases, to analytical uncertainty at

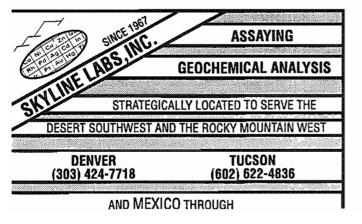


Table 1. Parameters of four experimental CHIM runs conducted in 1993 at the Kokomo mine, Central City district, Colorado, to test the newly developed NEOCHIM electrode. The working electrode for each run was a 5mm-diameter graphite rod. Electrical power was supplied by a 10kW generator.

	RUN	11	RUN	1 11	RUN	Ш	RUN IV	
	Cathode	Anode	Cathode	Anode	Cathode	Anode	Cathode	Anode
Outer Chamber								
Diameter, mm	40	40	40	40	40	40	100	100
Volume, ml	So	50	50	50	50	50	400	400
Composition	.01N	.01N	.1N	.1N	.1N	.1N	.7N	.7N
-	KNO ₃	KCI	KCI					
Inner Chamber								
Diameter, mm	25	25	25	25	25	25	40	40
Volume, ml	25	25	25	25	25	25	70	70
Composition	1N	1N	1N	1N	1N	1N	4N	4N
_	HNO ₃	HCI	KOH					
Salt Bridge								
Diameter, mm	25	25	25	25	25	25	40	40
Volume, ml	20	20	20	20	20	20	70	70
Composition (in	1N	1N	1N	1N	1N	1N	4N	4N
5% agar gel)	KNO ₃	KCI	KCl					
Date of Run (1993)	July	27	July	27	Aug. 12		Sept. 2	1,22
Duration of Run	0.5	Hrs	1.5	Hrs	3.5 Hrs 20 Hr		drs	

Table 2. Total micrograms (μg) of copper, lead, and zinc collected in the cathode solution during four experimental CHIM runs conducted at the Kokomo mine, Central City district, to test the NEOCHIM electrode.

		Cop	per			Lea	ad				Zinc	
RUN	ī	II	III	IV	1	11	111	١٧	1	11	III	١٧
SITE												
50N	0	0	2.5	12	0	0	30	520	3.5	9.0	35	144
40N	8.5	49	185	680	35	230	75 0	2160	40	165	500	1440
30N	1.0	.50	3.5	640	5.0	0	25	1120	9.5	24	35	640
20N	.50	1.5	8.5	64	0	5.0	45	0	10	45	75	560
10N	6.5	26	75	1640	5.0	15	70	400	80	550	900	4800
000	7.0	16	14	2480	10	15	55	1480	42	230	175	2680
105	1.0	1.0	3.0	٥	0	5.0	35	0	2	16	40	204
205	1.0	1.0	7.0	0	0	0	25	0	3.5	14	125	800
305	.50	0	3.0	36	0	0	35	0	2.0	7.0	35	600
405	.50	.50	8.5	0	0	0	50	0	1.5	6.5	35	180
50S	1.0	1.0	4.5	0	5.0	5.0	50	200	3.0	13	30	168

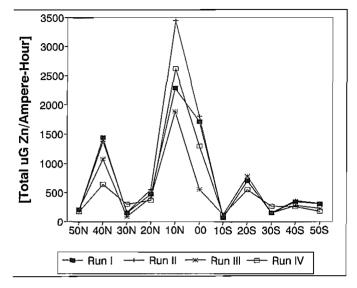
the limit of detection. Overall, our results indicate the NEOCHIM electrode yields extraction efficiencies more than two orders of magnitude better than the conventional CHIM electrode. The efficiencies were determined by ratioing the total equivalents of cations collected by each electrode to the amount of electrical charge supplied to the electrode.

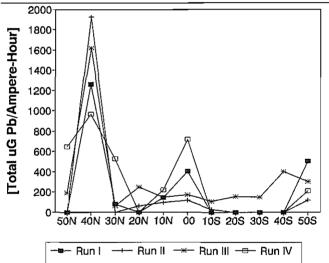
The collection of gold and other anionic species

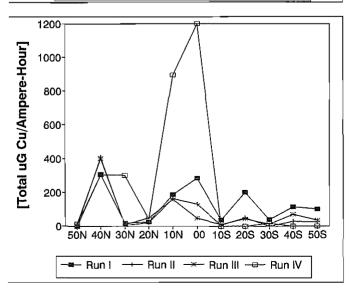
Most notable of our anode results (Table 3) is the occurrence of gold in some of the solutions of run IV. It is important to note that gold was not recovered in detectable amounts in any of the cathode solutions. Run IV was planned with the specific intent to test gold collection. The larger electrode and more concentrated fluids allowed us to conduct the extended run we felt necessary for collecting detectable amounts of gold that might be present in ionic form in soil moisture. Several months prior to run IV, we collected soil samples at each electrode site. An additional

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in the cathode solutions of four separate CHIM runs at the Kokomo mine, Central City district, Colorado, using the newly developed NEOCHIM electrode. The data are normalized to the amount of current (ampere-hours) delivered to each electrode, placed along a line at 10-meter intervals. A gold-bearing base-metal vein is located 3 meters beneath the surface at site 00.

bulk sample was collected near site 00 over the vein. Six splits of the bulk sample were analyzed by graphite furnace atomic absorption (Meier, 1980) for water-soluble gold, gold associated with organic matter (Gregoire, 1985), and total gold. We were unable to detect any water-soluble gold but found 95 to 97 percent of the gold associated with the organic matter in each of the splits. The data in Figure 3 shows a strong correlation between the gold associated with the organic matter in the soils from each site and the gold collected in the anodes during run IV. The mobility of gold as anionic organic complexes has been suggested by some investigators (Curtin et al., 1970; Baker, 1978; Varshal et al., 1990; Vlassopoulos et al., 1990). Ong and Swanson (1969) have suggested that gold migrates in association with complex organic acids as colloids of negative charge.

Anode data for zinc, sulfur, and arsenic (Table 3) implies the occurrence of zinc as both anions and cations with the cationic forms predominating in the surface environment at the Kokomo mine. There is a strong correlation between the sulfur determined in the anode solution and sulfate ion determined in water extracts of the soil samples from each

Table 3. Total quantities of zinc, sulfur, arsenic, and gold collected in the anode solution of experimental CHIM run IV conducted at the Kokomo mine, Central City district, to test the NEOCHIM electrode. (---) means not determined.

SITE	Zn Total μg	S Total μg	As Total μg	Au Total ng
50N	12	1200		0
40N				
30N	90	2200		110
20N	40	1000	30	0
10N	9 0	3800		144
00	110	180	96	92
105	55	0		20
20S	55	6700	52	12
30S	110	0	26	0
40S	40	0		0
50S	60	0		0



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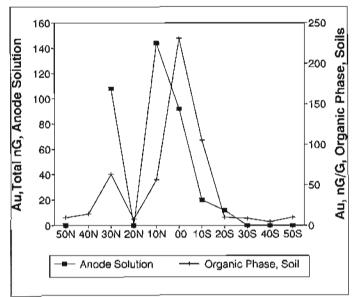


Figure 3. Total nanograms of gold (ng) collected in the anode solution of CHIM run IV at the Kokomo mine using the NEOCHIM electrode, and total nanograms of gold associated with organic matter in soil samples collected from the sites of electrode placement. Site 00 lies over the mineralized vein. The anode solution at site 40S was not recovered.

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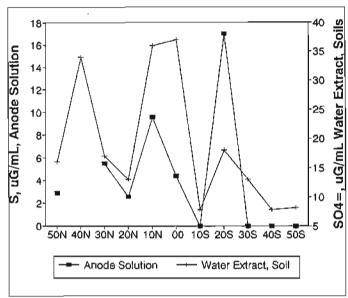


Figure 4. Concentrations, in $\mu g/ml$ (ppm), of sulfur collected in the anode solution of CHIM run IV at the Kokomo mine using the NEOCHIM electrode, and concentrations of sulfate in water extracts of soil samples, in $\mu g/g$ of soil (ppm), collected from the sites (50N-50S) of electrode placement. Site 00 lies over the mineralized vein. The anode solution at site 40S was not recovered.

site (Fig. 4). Arsenic was found in all four of the anode solutions selected for the analysis and was more abundant in the anode solution from site 00, 96 micrograms (μ g), than in the cathode solution from the same site (30 μ g). Analytical constraints prevented analysis of the anode solutions for other naturally occurring anions.

Conclusion

The NEOCHIM electrode overcomes the problems resulting from acid loss suffered by the conventional electrode. With the new electrode, we can more confidently use CHIM for its intended purpose: to provide representative samplings of electrically mobile, naturally occurring ions in the soil. It is now possible to more definitively determine the utility of CHIM for mineral exploration or other geochemical purposes. This new technology may also benefit the understanding of the mobility and transport of chemical elements under natural conditions, especially those of low abundance, such as gold, in the earth's surface environment.

Reinhard W. Leinz Donald B. Hoover U.S. Geological Survey M.S. 973, Federal Center Denver, CO 80225 (303) 236-2449

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CORRECTION

An astute reader noted an error in Table 1 of the article The Use of Silica Sand Cleaners in Pulverizing Müls to Reduce Cross Contamination by Lynda Bloom in EXPLORE No. 82. The corrected table is reproduced below.

Table 1. Determination of gold in silica sand cleaners

Wt. of	Sample	Gold Assays	Weighted	%
Gold(g)/	Weight	(ppm)	Average	Carry
Wt. of	(g)		(ppm)	~over
Sand(g)				ľ
0.0034/200	166.06	20.5, 13.4, 16.6, 18.5, 17.2,	17.2	
		16.9		1
Cleaner 1	134.42	0.18, 0.19, 0.43, 0.11, 0.10	0.21	1
Cleaner 2	68.70	Tr, Nil, Nil	Nil	
0.0034/100	99.16	37.7, 33.2, 27.9, 28.4	32.5	
Cleaner 1	69.46	0.44, 0.32, 1.02	0.47	1
Cleaner 2	69.60	0.06, 0.05, 0.06	0.06	
0.0069/100	98.93	72.7, 51.7, 78.0, 56.7	66.5	_
Cleaner 1	69.05	0.33, 0.19, 0.18	0.25	0.3
Cleaner 2	68.90	0.04, 0.04, 0.06	0.04	



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Special Issue

Mineral Deposits of Indonesia - Discoveries of the Past 25 Years

T.M. van Leeuwen, J.W. Hedenquist, L.P. James and J.A.S. Dow (Editors)

Preface

Indonesian mineral deposits - introductory comments, comparisons and speculations
R.H. Sillitoe

25 Years of mineral exploration and discovery in Indonesia T.M. van Leeuwen

Magmatic arcs and associated gold and copper mineralization in Indonesia
J.C. Carlile and A.H.G. Mitchell

Geologic and geochemical zoning of the Grasberg Igneous Complex, Irian Jaya, Indonesia

G.D. MacDonald and L.C. Arnold

Skarn Cu-Au orebodies of the Gunung Bijih (Ertsberg) district, Irian Jaya, Indonesia

H.J. Mertig, J.N. Rubin and J.R. Kyle

The Batu Hijau porphyry copper-gold deposit, Sumbawa Island, Indonesia

S.J. Meldrum, R.S. Aquino, R.I. Gonzales, R.J. Burke, A. Suyadi, B. Irianto and D.S. Clarke

Geology, porphyry Cu-Au, and epithermal Cu-Au-Ag mineralization of the Tombulilato district, North Sulawesi, Indonesia

J.A. Perello

Buligidun prospect: a copper, gold and tourmaline bearing porphyry and breccia system in northern Sulawesi,
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Upcoming JGE

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H. Lubis, S. Prihatmoko and L.P. James

Porphyry molybdenum mineralization in a continental collision setting at Malala, northwest Sulawesi, Indonesia T.M. van Leeuwen, R. Taylor, A. Coote, and F.J. Longstaffe

Sediment-hosted gold mineralization in the Ratatotok District, North Sulawesi, Indonesia

S.J. Turner, P.A. Flindell, D. Hendri, I. Harjana, P.F. Lauricella, R.P. Lindsay, B. Marpaung and G.P. White

Integrated exploration success for gold at Wetar, Indonesia D.M. Sewell and C.J.V. Wheatley

The Lerokis and Kali Kuning submarine exhalative goldsilver-barite deposits, Wetar Island, Maluku, Indonesia D.M. Sewell and C.J.V. Wheatley

The Gunung Pongkor gold-silver deposit, West Java, Indonesia

A. Basuki, D. Aditya Sumanagara and D. Sinambela Epithermal gold deposits in West Java, Indonesia: geology, age and crustal source

E. Marcoux and J.-P. Milesi

Structural controls and genesis of epithermal gold-bearing breccias at the Lebong Tandai mine, Western Sumatra, Indonesia

D.H. Jobson, C.A. Boutier and R.P. Foster

Alteration and epithermal mineralization in the Masupa Ria volcanic center, Central Kalimantan, Indonesia J.F.H. Thompson, H.Z. Abidin, R. Both, S. Martosuroyo, W.J. Rafferty and A.J.B. Thompson

Alluvial gold in Kalimantan, Indonesia: a colloidal origin? J.B. Seeley and T.J. Senden

High Au, Ag, Mo, Pb, V and W content of fumarolic deposits at Merapi volcano, central Java, Indonesia I. Kavalieris

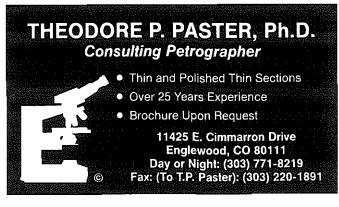
The mercury "tromol" mill: an innovative gold recovery technique, and a possible environmental concern L.P. James



1994 SME ANNUAL MEETING

Geochemistry Session

The 1994 SME Annual Meeting convened in Albuquerque, New Mexico, on February 14-17. Of the 77



sessions scheduled, the geochemistry session entitled Geochemical Exploration for Gold: "Using Old Techniques for New Discoveries" and the six geology sessions, including an excellent session entitled Ore Deposits of Chile, were of greatest interest to explorationists. Between 40 and 60 explorationists attended the geochemistry session.

Several presentations proved very interesting. Alan Morris suggested, in his presentation of the surficial geochemistry over the Meikle deposit, Carlin Trend, Nevada, that deep (+800 ft.), high-grade gold mineralization beneath barren bedrock does, in fact, yield recognizable geochemical patterns on the surface. Charlie Tapper, in his presentation on the Twin Creeks Mine, Getchell Trend, Nevada, told of new discoveries and improved geologic understanding as a result of combining the Chimney and Rabbit Creek mines. John Roberts' presentation on the use of spontaneous potential electro-geochemistry for detecting faults and/or intense argillic alteration beneath shallow to moderate cover should spark increased experimentation and use of this technique in the Great Basin and elsewhere. Finally, Joyanto Routh's presentation on the impact of the Van Stone Pb-Zn mine, Washington, on the Columbia River and surrounding environment was an excellent example of the application of geology-based environmental geochemistry.



TAUSON INTERNATIONAL SYMPOSIUM

IV Joint International Symposium on Exploration Geochemistry, A Tribute to Academician L.V. Tauson, September 7-10, 1994. Irkutsk, Russia

I. TECHNICAL SESSIONS

- Geochemical mapping for solving fundamental problems of geology, environment and exploration
- Geochemical methods of exploration and environment assessment
- Memorial Session, dedicated to L.V. Tauson (geochemical classification of geological formations, geochemistry of ore-magmatic systems)
- 4. Reference samples of different geochemical media and analytical means of geochemical investigations
- 5. Recent methods of geochemical data processing and geochemical map compilation

II. WORKSHOPS

- International Geochemical Mapping (including programmes "Geochemical map of Russia" and "Geoecology of Russia")
- 2. Geochemical prospecting for precious metals in the northern hemisphere
- 3. Baikal International Geochemical Polygon (programmes "Baikal Drilling", "Global change of the Environment and Climate", "Baikal Polygon, Russia")
- 4. Geochemical mapping and monitoring aimed at medical and biological problems

Tauson International Symposium

Continued from Page 16

- III. SESSION OF INTERNATIONAL AND NATIONAL GEOCHEMICAL ORGANIZATIONS AND **PROJECTS**
- Council of International Association of Geochemistry and Cosmochemistry
- Council of Association of Exploration Geochemists
- Baikal Drilling Project
- Projects IGCP no. 360 "Global Geochemical Baselines"
- Session of Coordination Council on multi-purpose geochemical mapping and geoecology (Project "Geochemical map of Russia" and "Geoecology of Russia")
- Working Group "Global Geochemical Mapping" IAGC

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I۷. **EXCURSIONS**

- 1. Environmental geochemistry of Irkutsk and its surroundings
- 2. Geochemical monitoring of the landscapes of the Western Baikal region
- 3. Murun charoite deposit
- 4. Gold deposits, Trans-Baikal region (Darasun, Balei)
- Gold deposits, South-east Siberia (Zun-Kholba, Sukhoi
- 6. Phlogopite and lazurite deposits of the South Baikal
- 7. Geochemistry of the bottom sediments of Baikal lake and
- 8. Geology and mineralogy of the Baikal region

OFFICIAL LANGUAGES:

English and Russian

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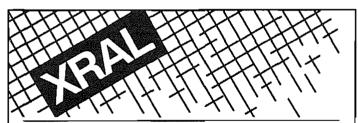
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CALENDAR OF EVENTS

International, National and Regional Meetings of Interest to Colleagues Working in Exploration and Other Areas of Applied Geochemistry.

- ■May 10-14, '94, Mining Latin America, mtg., Santiago de Chile, by Institution of Mining and Metallurgy and others (IMM, 44 Portland Place, London, W1N 4BR, England; TEL: 44 71 580 3802; FAX: 44 71 436 5388)
- ■May 13-18, '94, GAC-MAC, ann. mtg., and MDD-GAC Short Course on Alteration Processes, Waterloo (G. Roberts, Waterloo '94, Dept. of Earth. Sci., Univ. of Waterloo, Waterloo, Ontario N2L 3G1, Canada; TEL: (519) 885-1211; FAX: (519) 746-7484; short course contact D.R. Lentz, GSC, PO Box 50, Bathurst, New Brunswick E2A 3Z1, Canada; TEL: (506) 546-2070; FAX: (506) 546-3994)
- ■May 16-18, '94, Peru, First International Gold Symposium and Exhibition, Lima, Peru, by Comite Aurifero, Sociedad Nacional de Minero y Petroleo and Randol International Ltd. (Sondra Wells, Randol International Ltd., Golden, CO, USA, FAX: (303) 526-1650 or Dr. Guillermo Albareda, Comite Aurifero Sociedad Nacional de Minero y Petroleo, Las Flores 346, Lima 27, Peru. TEL: (51-14) 70-4260, FAX: (51-14) 70-4245)
- ■May 17-19, '94, GAC-MAC, ann. mtg., Edmonton, Alberta (J. W. Kramers, Alberta Geological Survey, Box 8330, Station F, Edmonton, T6H 5X2, Canada; TEL: (403) 438-7644; FAX: (403) 438-3364)



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- ■June 6-10, '94, Mining History, mtg., Golden, CO, by Colorado School of Mines (Mining History Association, Box 15030, Denver, CO 80215)
- ■July 17-19, '94, Conf. on Lead and Arsenic Exposure in the Rocky Mountains, Salt Lake City, Utah (Society of Environmental Geochemisty and Health, Rocky Mountain Region, P.O. Box 70915, Salt Lake City, UT 84170, FAX: (801) 322-8398 or (801) 261-2194)
- Aug. 28-Sept. 3, '94, European Association of Geochemistry Meeting and 4th Goldschmidt Conference, Edinburgh (Dr. B. Harte, Department of Geology and Geophysics, Grant Institute, University of Edinburgh, West Mains Road, Edinburgh, EH9 3JW, UK)
- ■Sept. 5-7, '94, Prospecting in Areas of Glaciated Terrain, mtg., St. Petersburg, Russia (Institution of Mining and Metallurgy, 44 Portland Place, London W1N 4BR, UK, TEL: (071) 580 3802; FAX: (071) 436 5388)
- ■Sept. 8-12, '94, Joint International Symposium on Exploration Geochemistry, Irkutsk: a tribute to Academician L.V. Tauson (Pavel Koval, Institute of Geochemistry, P.O.Box 4019, 664033 Irkutsk-33, Russia; TELEX: 133 163 Taiga SU; TEL: 395(2) 46-59-78)
- ■Sept. 12-15, '94, 3rd Symposium on Environmental Geochemistry, Kraków, Poland (Dr hab. Edeltrauda Helios Rybicka, Faculty of Geology, Geophysics and Environmental Protection, University of Mining and Metallurgy, Al. Mickiewicza 30, 30-059, Poland; TEL: (48) 12-333290; FAX: (48) 12-332936)
- ■Sept. 18-22, '94, Geoanalysis '94, International Symposium on Analysis of Geological and Environmental Materials, UK (Doug Miles, British Geological Survey, Keyworth Nottingham NG12 5GG, UK; TEL: 44-36-602-3100; FAX: 44-36-602-3200)
- ■Sept. 18-22, '94, Mineral Exploration '94 (including a 1-day symposium on Latin American Mineral Deposits), Lake Tahoe, NV, U.S.A. (Dr. Phil Newall & Dr. Alan Butcher, CSM Associates Ltd., Pool, Redruth, Cornwall TR15 3SE, UK; TEL: 44 (0)209 717724; FAX: 44 (0)209 716977)
- ■Sept. 19-21, '94, Exploration and Mining Geology of World Class Deposits, Sudbury, Ontario, Sudbury Geological Discussion Group, and Canadian Institute of Mining, Metallurgy and Petroleum (Ruth Debicki, MNDM, A3-933 Ramsey Lake Road, Sudbury, Ontario P3E 6B5 Canada; TEL: (705) 670-5627; FAX: (705) 670-5622)
- ■Sept. 26-27, '94, Manganese Mineralization: Geochemistry and Mineralogy of Terrestrial and Marine Deposits, London, England (Dr. Keith Nicholson, School of Applied Sciences, Robert Gordon University, Aberdeen AB1 1HG, Scotland. TEL: 0224-262821/262802, FAX: 0224-262828)
- ■Oct. 2-5, '94, 1994 Annual Meeting of the Geothermal Resource Council (GRC), theme: Restructuring the Geothermal Resource Council, Salt Lake City, UT, USA (Philip M. Wright, Geothermal Resource Council, PO Box 1350, Davis, CA 95617-1350, TEL: (916) 758-2360, FAX: (916) 758-2839)
- ■Oct. 5-7, '94, Porphyry Copper Deposits from Alaska to Chile Symposium, Tucson, AZ, by Arizona Geological Society, Society for Mining, Metallurgy and Exploration, and U.S. Geological Survey (Jim Laukes, The University of Arizona Extended

Calendar of Events

Continued from Page 18

University, 1955 E. Sixth St., Tucson, AZ 85719-5224; TEL: (800) 955-8632; FAX: (602) 621-3269; E-mail (Internet): jlaukes.ccit.arizona.edu)

- ■Oct. 25-27, '94, Geological Society of America, ann. mtg., Seattle, WA (Vanessa George, G.S.A., Box 9140, Boulder, CO 80301; TEL: (303) 447-2020)
- ■Oct. 26-29, '94, 4th Asia Pacific Mining Conference and Exhibition, Jakarta, Indonesia (Warjono Soemodinoto or Joe Widartoyo, IMA Secretariat, Jl. Prof. Dr. Supomo SH No.10, Jakarta 12870, Indonesia; TEL/FAX: 62 (021) 828-0763 / 830-3632)
- ■Oct. 30-Nov. 1, '94, RANDOL Latin American Mining Opportunities, conference, Vancouver, British Columbia (RANDOL International LTD., 21578 Mountsfield Drive, Golden, CO 80401, TEL: (303) 526-1626; FAX: (303) 526-1650) ■Mar. 6-9, '95, SME Annual Meeting and Exhibit, Denver, CO (Meetings Dept., SME Inc., PO Box 625002, Littleton, CO 80162-5002; TEL: (303) 973-9550; FAX: (303) 979-3461)
- ■Apr. 3-7, '95, Centennial Geocongress 1995, Johannesburg, The Geological Society of South Africa (The Congress Secretariat, Centennial Geocongress, PO Box 36815, Johannesburg 0102, South Africa; TEL/FAX: 27 12 47 3398)
- ■Apr. 10-13, '95, Geology and Ore Deposits of the American Cordillera, Geological Society of Nevada Symposium III (Bob Hatch, Geological Society of Nevada, P.O. Box 12021, Reno NV 89510; TEL: (702) 323-4569; FAX: (702) 323-3599)
- May 15-19, '95, 17th International Geochemical Exploration Symposium, "Exploring the Tropics", Townsville, Queensland, Australia (Russell Myers, 17 IGES, National Key Centre in Economic Geology, James Cook University, Townsville, Queensland 4814, Australia; TEL: (077) 814486; FAX: (61) 77-815522)
- ■June 3-6, '95, International Field Conference on Carbonate-hosted Lead Zinc Deposits, SEG Anniversary Field Conference (David Leach or Martin Goldhaber, USGS, Branch of Geochemistry, MS 973, PO Box 25046, Federal Center, Denver, CO 80225, USA, FAX: (303) 236-3200, e-mail: dleach@helios.cr.usgs.gov)
- June 7-9, '95, African Mining '95, Windhoek, Namibia (IMM, 44 Portland Place, London W1N 4BR, UK; TEL: (071) 580-3802; FAX: (071) 436-5388)
- ■Nov. 6-9, '95 Geological Society of America, ann. mtg., New Orleans, LA (Vanessa George, 3300 Penrose Place, Boulder, CO 80301; TEL: (303) 447-2020; FAX: (303) 447-1133

Please check this calendar before scheduling a meeting to avoid overlap problems. Let this column know of your events.

Fred Siegel

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NEW MEMBERS

To All Voting Members:

Pursuant to Article Two of the Association's By-Law No.1, names of the following candidates, who have been recommended for membership by the Admissions Committee, are submitted for your consideration. If you have any comments, favorable or unfavorable, on any candidate, you should send them in writing to the Secretary within 60 days of this notice. If no objections are received by that date, these candidates will be declared elected to membership. Please address comments to Sherman P. Marsh, Secretary AEG, U.S. Geological Survey, Mail Stop 973, Box 25046, Federal Center, Denver, Colorado 80225, U.S.A.

Editors note: Council has decided that all new applicants will receive the journal and newsletter upon application for membership. The process of application to the Nepean, Ontario office, recommendation by the Admissions Committee, review by the Council, and publication of applicant's names in the newsletter remains unchanged.

FELLOW

Cook, Stephen J. Geochemist B.C. Geological Survey Victoria, BC, Canada

MEMBERS

Bloomer, Anthony G. Rio Tinto Exploration Sandton, South Africa

Bolster, Simon J.S. Geomorphologist Normandy Poseidon Townsville, Qld, Australia

Bolter, Ernst University of Missouri Rolla, MO, U.S.A.

Bond, Roger Geologist Newmont Exploration Elko, NV, U.S.A

Brewer, Adrian
Principal Geologist
RGC Exploration
Fyshwick, ACT, Australia

Goldsworthy, Julian
Exploration Geologist
Newcrest Mining
Belmont, WA, Australia

Harding, Tony CRA Exploration Karratha, WA, Australia

Higgins, M.
Professor
University of Quebec
Chicoutimi, PQ, Canada

Hudson, Geoffrey R.T.

Operations Manager

Australian Mineral Foundation
Glenside, SA, Australia

Lindhorst, Jeffrey W. Geologist Geopeko Parkes, NSW, Australia

Ouedraogo, Mourou F. Bureau Mines and Geology Ouagadougou, Burkina Faso

Powell, Matthew B. Big Creek Exploration Services Sun Valley, NV, U.S.A.

Piekenbrock, Joe R. Placer Dome Anchorage, AK, U.S.A.

Silakul, Tawisak Senior Geologist Padaeng Industry Chiang Mai, Thailand

Turner, Dean D. Senior Project Geologist FMC Gold Reno, NV, U.S.A.

Viladevall, Manuel Professor Zona Universitaria de Pedrales Barcelona, Spain

STUDENT

Laurus, Kathy University of British Columbia Vancouver, BC, Canada

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BOOK REVIEW

Randol International recently published the 1994 edition of their Mexican Mining Directory. This 248 page reference volume begins with tables of mining statistics for Mexico, then provides a listing of mining and exploration companies operating in Mexico. This is followed by listings of mines, government associations, a buyers guide, and a calendar of events. Anyone exploring, or considering exploring in Mexico would be well advised to obtain a copy of this \$US85 volume. For more information contact Randol International Ltd, Golden Colorado at (303) 526-1626.

EXPLORE staff



RECENT PAPERS

This list comprises titles that have appeared in major publications since the compilation in EXPLORE Number 82. Journals routinely covered and abbreviations used are as follows: Economic Geology (EG); Geochimica et Cosmochimica Acta (GCA); the USGS Circular (USGS Cir); and Open File Report (USGS OFR); Geological Survey of Canada Papers (GSC Paper) and Open File Report (GSC OFR); Bulletin of the Canadian Institute of Mining and Metallurgy (CIM Bull); Transactions of Institute of Mining and Metallurgy, Section B: Applied Earth Sciences (Trans IMM). Publications less frequently cited are identified in full. Compiled by L. Graham Closs, Department of Geology and Geological Engineering, Colorado School of Mines, Golden, CO 80401-1887, Chairman AEG Bibliography Committee. Please send new references to Dr. Closs, not to EXPLORE.

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