President’s Message

Paul Morris

Discussions with AAG members at our biennial symposia, by email, or at other conferences or workshops indicate that most members rate AAG’s publications (EXPLORE and GEEA) as significant reasons for belonging to the Association. Apart from being the means of spreading applied geochemical knowledge, both seem to draw together the applied geochemistry community, which is spread over all continents. The most recent editions of GEEA (volume 10 numbers 1 and 2) - released in February and May this year - include a collection of papers presented at the Exploration 07 meeting in Toronto in September, 2007. The papers reflect on both advances made in exploration geochemistry in the last decade and future directions, discuss advances in analysis, the efficacy of different digestion techniques, the use of groundwater in mineral exploration, and approaches to understanding large, regional geochemical datasets. These types of paper provide concise and often thought-provoking summaries, as well as including a comprehensive list of references. They are a valuable complement to the more deposit- or technique-focussed papers which span the breadth of applied geochemistry.

This edition of EXPLORE contains a discourse on sampling written, in his inimitable style, by Cliff Stanley. The focus of the paper - how much is enough (in terms of sampling) - has been a vexatious issue since exploration geochemistry was seen as a cost-effective approach to exploring for a variety of commodities, and I’m sure that Cliff’s exposition will provoke some worthwhile responses. Revisiting issues such as estimating ideal sample numbers highlights the importance of computer technology, which at the embryonic phase of exploration geochemistry was embryonic, precluded any rigorous statistical examination, and instead relied on consulting voluminous tables and making calculations using electronic equipment with limited capabilities. In hoping for some replies to Cliff’s paper, it should be pointed out that EXPLORE is a great vehicle for canvassing different points of view. Publishing in EXPLORE does not necessarily require writing a paper, as there is scope for letters-to-the-editor, which I hope will inspire some members to contribute.

In looking at a fundamental of exploration geochemistry such as sample size, a few members will cast their minds back to the late 1960s and early 1970s when these issues were often discussed. It’s timely to reflect now in that 2010 is AAG’s (AEG’s) 40th birthday. However, the formalisation of the Association was preceded by the inaugural applied geochemistry symposium held in late April 1966 in Toronto. After the next (1968) meeting, a group of applied geochemists, including Alan Coope, John Hansuld, and Bob Garrett, started up the Association of Exploration Geochemists (AEG). AEG/AAG has passed over the same hurdles and through the same hoops as most other professional societies (increasing costs, fluctuating membership, journal demises and rebirths, change in focus) but has maintained its relevancy to the applied geochemistry community, and expanded to embrace environmental as well as exploration geochemistry. These successful adaptations over the past 40 years are due to the commitment of AAG’s Council, and the ongoing support of its members.

I browsed a couple of websites today, AAG’s and that put together by Pertti Sarala and his group in Rovaniemi, Finland. Bob Eppinger, Andrew Ransom and Gemma Bonham-Carter have updated and expanded AAG’s website, www.appliedgeochemists.org, and I encourage you to not only visit it, but also send any contributions, upcoming event details, etc directly to Bob, eppinger@usgs.gov, for addition to the site. Judging by the ever-changing website for the 25th IAGS meeting in Rovaniemi, Finland (August 2011), Pertti Sarala and his local organising committee are continually working to provide AAG members with a memorable meeting. Already Actlabs and ESSA have indicated they will be taking up Trade Exhibitor opportunities, and ALS Minerals have signed up as a sponsor. The documentation available on the website for trade display and sponsorship options is easily available and comprehensive, and I urge companies to have a look and take the opportunity to reach a wide range of applied geochemistry professionals at this meeting, which is now less than 12 months away.

Paul Morris
President
How Many Samples Are Enough?

Introduction

Recently, I had the good fortune to be selected to go on a CIMM lecture tour across Canada to present a talk on quality control to geoscientists in the mining industry. After several of these talks, a number of geoscientists asked me the question: “how many samples are enough?”

Now, I have heard this question asked many times before, and am aware of the ‘conventional wisdom’ that a dataset of approximately 30 samples is generally considered to be ‘enough’ for geochemical applications. Consequently, although my immediate answer was “30”, the widespread interest in this question amongst geoscientists motivated me to investigate the origins of this ‘rule of thumb’.

In the geological literature, this concept apparently dates back to a publication by Bob Garrett (1979). In that paper, Bob noted that the standard error of the mean is a function of the inverse of the square root of the number of samples. Consequently, he plotted $n$ versus $1/\sqrt{n}$ and pointed out that the resulting curve basically ‘levelled out’ at approximately $n = 30$. As a result, Bob concluded that there is no real advantage to using more than 30 samples to estimate the mean, because no substantial further decrease in the estimation error of the mean can be achieved by such effort.

Clearly Bob’s analysis is highly practical. Unfortunately, the point at which Bob’s curve ‘levels out’ is a function of both the scale one plots the curve at, and the ‘eye of the beholder’, making use of such an approach rather arbitrary. Bob readily acknowledged this, and pointed out that the number of samples required “will depend on the particular situation” (Garrett 1979), a prudent caveat. Unfortunately, this hasn’t really prevented the widely quoted and employed “30 samples” from being used by probably the majority of geoscientists as anything but a hard and fast rule (i.e., immutable and inviolable) that should apply in all situations.

Until now, I can’t say that I have been anything but guilty of employing this erstwhile ‘constant’ in an array of geochemical applications where its use may have been inappropriate. After I understood the reason for the conventional wisdom of “30 samples”, I was justifiably chagrined, and decided to see if I could derive a more comprehensive way to determine “how many samples are enough” that doesn’t have the inherent subjectivity of the present ‘conventional wisdom’.

In undertaking this task, it became immediately apparent that Bob Garrett (1979) was on the right track, as the standard error of a statistic provides information about the confidence one can place in an estimate of that statistic. However, experience shows that geoscientists are many times concerned with the relative error of a statistical estimate instead of the absolute error. As a result, to determine “how many samples are enough”, one should probably also consider the relative standard error of that statistic.

Because Bob Garrett’s suggested ‘rule of thumb’ (1979) is based on the formula of the standard error of the mean, the ‘30 samples’ that Bob advocated applies only to the estimate of a mean of a set of observations. It is likely that the estimation of other statistics (e.g., the standard deviation, correlation coefficient, etc.) would require different numbers of samples in order to obtain an adequate estimate. Fortunately, the mean is a statistic that is commonly investigated by geoscientists, and the formula for the standard error of the mean ($se_x$) is:

$$se_x = \frac{s_x}{\sqrt{n}}$$

where $n$ is the number of samples and $s_x$ is the standard deviation. Because the relative standard error of the mean ($rse_x$) is merely the standard error of the mean divided by the mean, the relative standard error statistic is:

$$rse_x = \frac{se_x}{\bar{x}} = \frac{\frac{S_x}{\sqrt{n}}}{\bar{x}} = \frac{s_x}{\bar{x}\sqrt{n}} = \frac{CV}{\sqrt{n}},$$

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where CV is coefficient of variation, equal to the standard deviation divided by the mean ($= s_x / \bar{x}$).

Now, the standard approach to making decisions about the quality of an estimate of a statistic involves the use of formal inferential significance tests. Unfortunately, these commonly require assumptions about the distribution of the data (e.g., that the data are normally distributed). In my experience, geochemical data only very rarely exhibit distributions similar to ideal statistical distributions (e.g., Sinclair 1976). As a result, any method to determine “how many samples are enough” probably should not assume a statistical distribution for one’s data. This is particularly true given that when one asks such a question, one typically has, or can afford, only a small number of samples to begin with, and thus one has, and will continue to have, very little evidence to deduce what distribution the data actually derive from.

Consequently, a method to assess “how many samples are enough” should be ‘distribution-free’ in order to have maximum utility. Fortunately, derivation of the formula for the standard error of the mean, and thus the formula for the relative standard error of the mean, involves only first-principles calculus, whereby a first order Taylor polynomial expansion of the formula for the mean is used to propagate the errors in a group of measurements into the error in the mean of that group of measurements (Stanley 1990; Appendix). Thus, the absolute and relative standard errors of the mean formulae (Equations 1 and 2) are not derived using any assumption about the distribution of the data. Consequently, these formulae are ‘distribution free’, and can be used to determine “how many samples are enough” for data derived from any and all distributions (e.g., normal, Poisson, binomial, hypergeometric, exponential, multi-modal, etc.).

An assessment of the literature demonstrates that all standard error formulae are inverse functions of the number of samples used in calculating the standard error of a statistic (e.g., Arsham, undated). Aside from indicating that as n increases, estimates of the mean become more precise, this functional relation makes it possible to determine “how many samples are enough” to obtain a reliable estimate of the mean using the absolute or relative standard error of the mean formula.

This can be done by first determining the tolerance level one is willing to accept in one’s estimate of the mean (e.g., an absolute error of 5 ppm [1 std. dev.] on the mean estimate of a set of trace element concentrations, or say 5% relative error on that mean estimate). By assigning this tolerance level to the absolute or relative standard error of the mean ($s_e$ or $rse$) in Equations 1 or 2, and then algebraically manipulating these equations such that they are expressed in terms of the number of samples, we obtain:

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\[ n = \left( \frac{s}{se_x} \right)^2, \quad (3) \]

for the absolute standard error of the mean, and:

\[ n = \left( \frac{CV}{rse_x} \right)^2, \quad (4) \]

for the relative standard error of the mean.

Functional analysis of Equations 3 and 4 reveals that the number of samples required to obtain an adequate estimate of the mean is dependent on both the confidence level chosen (\(se_x\) or \(rse_x\)), and either the absolute (s) or relative (CV) variation in the data. This result is intuitive, as samples from a distribution that exhibits little absolute or relative variation (i.e., with a low value of s or CV) will each be relatively good estimates of the mean of that distribution, and so only a small number of such samples will be necessary to obtain a good estimate of the mean. In contrast, samples from a distribution that exhibits high absolute or relative variation (i.e., with a high value of s or CV) will mostly be poor estimates of the mean, and so a much larger number of these samples will be required to obtain a mean estimate of adequate quality.

Figures 1 and 2 graphically illustrate how many samples are necessary to estimate the mean of measurements from a suite of samples at a variety of absolute and relative precision levels (\(se_x\) or \(rse_x\)). Note that in both cases, the larger the absolute or relative variation in the data (s or CV), the more samples are necessary to obtain a given level of confidence in the estimate of the mean.

Unfortunately, determining “how many samples are enough” to estimate a mean requires prior knowledge of the absolute or relative variation in the data (s or CV). If no samples have been analyzed, this information will not be available. As a result, in order to estimate the mean of a suite of samples to a certain level of precision, one has to approach this problem iteratively. If one can make a ‘provisional’ estimate of the absolute or relative variation (s or CV), either using a guess, assumption, experience, samples that have already been analyzed, or results from historical datasets, and identify an acceptable tolerance level (\(se_x\) or \(rse_x\)), then one can make a provisional estimate of “how many samples are enough” using Equations 3 or 4, respectively. Collecting and analyzing this provisional number of samples will provide (additional) data to make an (improved) estimate of the

Figure 1. Number of samples (n) required to obtain an estimate of the mean with a desired level of absolute estimation error (\(se_x\)). Note that the number of samples required is a function of the standard deviation (s), a measure of the variation in the measurements, and the number of samples (n), and thus is not, universally, 30.

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Figure 2. Number of samples (n) required to obtain an estimate of the mean with a desired level of relative estimation error (rse). Note that the number of samples required is a function of the coefficient of variation (CV), a measure of the relative variation in the measurements, and the number of samples (n), and thus is also not, universally, 30.

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absolute or relative variation (s or CV), and these can then be used to make a better estimate of “how many samples are enough”, again using Equations 3 or 4. Obviously, a poor initial estimate of the amount of variation that exists will lead to a poor estimate of the number of samples required, but because this procedure can be undertaken iteratively, better estimates of the absolute or relative variation can be obtained as more and more samples are collected, such that an adequate estimate of the mean, at whatever confidence level is required, can eventually be obtained.

The following example illustrates how the above approach can be used to determine “how many samples are enough.”

Case History

The North American Soil Geochemical Landscapes Project (NASGLP) is a tri-national program begun in 2006 that seeks to provide a consistent set of C-horizon soil geochemical data across North America (Natural Resources Canada 2008). Collection of soil samples at a density of one sample per ~770 km² (Goodwin et al. 2009) on a random, stratified grid was conducted in Nova Scotia, Canada as part of a pilot study in 2007 and 2008. Analytical results (< 63 μm size fraction, nitric-perchloric-hydrochloric-hydrofluoric acid digestion, ICP-MS finish) confirm that As concentrations in Nova Scotia are generally elevated relative to the Canadian Soil Quality Guideline (Maximum Allowable Concentration; MAC) of 12 ppm (Canadian Council of Ministers of the Environment 2009; pers. comm., Terry Goodwin, Nova Scotia Department of Natural Resources), as concentrations range from 2.2 to 345.7 ppm (cf., the average As concentration in soils in Southern California is 1.5 ppm; n = 1086; Chernoff et al., un-dated).

The MAC’s comprising the Canadian Soil Quality Guidelines have been developed to determine whether a soil concentration is at or below a level where no appreciable risk to human and environmental health is expected. If the maximum soil concentration at a site exceeds the MAC, a potential problem exists, requiring further study and possible remediation (Willis 2006).

Results from the NASGLP indicate that south of the Cobequid-Chedabucto Fault Zone (CCFZ) within the Meguma Terrane, 75% of the samples have As concentrations that exceed the MAC (pers. comm. Goodwin 2010), a probable consequence of the abundance of arsenopyrite (up to 15% in Meguma Supergroup meta-sedimentary rocks comprising the basement to that terrane. In contrast, north of the CCFZ, only 25% of the samples have As concentrations that exceed the MAC (pers. comm. Goodwin 2010), a proportion that is still high, probably because erosion products from the Meguma Supergroup were deposited in Carboniferous basins north of the CCFZ. These observations indicate that many uncontaminated sites in Nova Scotia would be considered ‘contaminated’ using the national MAC for As because of the naturally high background As concentrations that exist in the province.

As a result, it is probably appropriate to identify an alternative ‘made in Nova Scotia’ As concentration (pers. comm. Terry Goodwin, NS DNR) for local use, in lieu of the Canadian MAC of 12 ppm. How this As concentration will be identified is not entirely clear, because MAC’s are not merely a series of threshold separating background from anomalous concentrations, as they have been developed using “conservative and protective assumptions, and (risk assessment) principles/methods to ensure protection of both sensitive human and ecological receptors under typical conditions” (Willis 2006). As a result, the natural frequency distribution of As concentrations in soil is only one factor contributing to the assignment of an MAC; other factors include analytical geochemistry methods and detection limits; industrial production and method-of-use factors; physical and chemical properties; mineral and water speciation; exposure mechanisms; environmental fate and behavior in soils, waters, air, microbes, plants and animals; bio-availability and bio-accumulation; metabolic behavior; acute and chronic toxicity levels (e.g., LD₅₀’s) in a variety of organisms (e.g., microbes, invertebrates, plants, livestock, wildlife, experimental animals, and humans); treatment and remediation ease, cost, and success probability; and benchmarks established by other jurisdictions (e.g., the US EPA). Unfortunately, the large number of factors that influence the assignment of an MAC make most MAC’s appear to have been chosen subjectively because the ‘influence weights’ each factor is assigned are not known, and many of these factors are not quantitative.

Nevertheless, in order establish an appropriate local MAC, the frequency distribution of As concentrations in soils needs to be understood so that the range of background concentrations can be identified using probability plot analysis (Sinclair 1976). Unfortunately, the only Nova Scotia-wide soil survey that has been undertaken is that of the NASGLP, and it provides only 72 samples (including 3 field duplicates; Goodwin et al. 2009). This is generally inadequate to characterize the distribution of As in Nova Scotia (with a landmass of 53,338 km²), and so very many more samples will need to be collected and analyzed.

As a result, in 2010, the Nova Scotia Department of Natural Resources (NS DNR) intends to collect additional soil samples using the same sampling and analysis protocols as the NASGLP, in order to obtain a better understanding of the As concentrations in the province. This will create a

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larger database with a higher sample density than the original NASGLP survey.

Because initial results indicate that soils developed over different lithologies in Nova Scotia exhibit different As concentrations, the NS DNR has divided the province into eight general but distinct lithozones (soils developed over different bedrock lithologies). As part of their effort to determine the frequency distribution of As concentrations in these lithozones, they intend to estimate a number of statistics that contribute to the understanding of the distribution of As concentrations in soils developed over each. These statistics will include the mean, standard deviation, and quantiles, among others, and it is hoped that at least a crude approximation of the frequency distribution of As concentrations in each lithozone can also be obtained from these data.

Two of the lithozones defined by the NS DNR are underlain by: (i) ‘granite’, including but not limited to the peraluminous South Mountain batholith, and (ii) ‘lower Paleozoic meta-greywacke and slate’ (the Meguma Supergroup). NASGLP individual soil sample As concentration results from these lithozones are presented in Table 1, along with their means, standard deviations and coefficients of variation (Goodwin et al. 2009). Although roughly similar numbers of NASGLP samples were collected and analyzed (17 and 15, respectively), the As concentration statistics for soils from these lithozones are significantly different.

Overall, As concentrations are high, with average concentrations of 13 and 47 ppm for granite and lower Paleozoic meta-greywacke and slate, respectively (Table 1). Furthermore, soils developed over lower Paleozoic meta-greywacke and slate exhibit more variable As concentrations than granite (standard deviations of 85 and 7 ppm, and coefficients of variation of 182 and 55 %, respectively; Table 1). Note that although these Meguma Supergroup rocks are known to contain As-bearing saddle reef gold deposits (Sangster & Smith 2007), all of the randomly located NASGLP samples ended up being collected away from known mineralization, so the higher As concentrations may be associated with yet-to-be discovered mineralization, or are merely samples from the high end of an elevated background As frequency distribution. In either case, these elevated As concentrations justify health and environmental concern. Additionally, the standard deviations and coefficients of variation of As concentrations over different lithozones indicate that a larger number of samples will be necessary to determine a reliable average As concentration in soils over lower Paleozoic meta-greywacke and slate, than in soils over granite.

**Example Calculation**

Fortunately, because at least some NASGLP As concentrations have already been measured from these two lithozones, we can use these data to provide ‘provisional’ estimates of the relative variations (CVs) of As concentrations in soils over these two lithozones (Table 1). These can then be used to obtain estimates of the number of samples (n) required to calculate mean As concentrations for each lithozone to a certain precision level. After collecting and analyzing n additional samples, better estimates of the relative variations (CVs) can then be made using this larger dataset, and these can be used to further refine/confirm the number of samples necessary to reliably estimate the mean As concentrations in the various lithozones. Obviously, this process can proceed iteratively until sufficiently reliable mean concentrations have been obtained.

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**Table 1. Arsenic concentrations (ppm) and statistics from soil samples collected as part of the NASGLP in Nova Scotia (Goodwin et al. 2009).**

<table>
<thead>
<tr>
<th>Samples &amp; statistics</th>
<th>Over granite lithozone</th>
<th>Over lower Paleozoic meta-greywacke &amp; slate lithozone</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>13.8</td>
<td>11.4</td>
</tr>
<tr>
<td>2</td>
<td>4.8</td>
<td>81.1</td>
</tr>
<tr>
<td>3</td>
<td>12.3</td>
<td>37.3</td>
</tr>
<tr>
<td>4</td>
<td>18.5</td>
<td>15.8</td>
</tr>
<tr>
<td>5</td>
<td>12.1</td>
<td>29.9</td>
</tr>
<tr>
<td>6</td>
<td>33.1</td>
<td>16.1</td>
</tr>
<tr>
<td>7</td>
<td>2.2</td>
<td>13.4</td>
</tr>
<tr>
<td>8</td>
<td>15.3</td>
<td>5.9</td>
</tr>
<tr>
<td>9</td>
<td>15.3</td>
<td>345.7</td>
</tr>
<tr>
<td>10</td>
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<td>26.2</td>
</tr>
<tr>
<td>11</td>
<td>15.7</td>
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</tr>
<tr>
<td>12</td>
<td>13.7</td>
<td>41.9</td>
</tr>
<tr>
<td>13</td>
<td>4.7</td>
<td>22.2</td>
</tr>
<tr>
<td>14</td>
<td>15.9</td>
<td>25.2</td>
</tr>
<tr>
<td>15</td>
<td>3.0</td>
<td>16.9</td>
</tr>
<tr>
<td>16</td>
<td>18.3</td>
<td></td>
</tr>
<tr>
<td>17</td>
<td>10.8</td>
<td></td>
</tr>
<tr>
<td>Mean</td>
<td>13.35</td>
<td>46.61</td>
</tr>
<tr>
<td>Standard Deviation</td>
<td>7.36</td>
<td>84.77</td>
</tr>
<tr>
<td>CV (%)</td>
<td>55.12</td>
<td>181.85</td>
</tr>
</tbody>
</table>

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Table 2 illustrates the number of samples required to estimate mean As concentrations in each lithozone at various tolerances, based on the results from the initial NASGLP samples. For example, for the soils overlying the granite lithozone, the calculated sample size necessary to achieve a 10% tolerance level ($rse$) is, coincidentally, $(55.12/10)^2 = 30.38$ samples. In contrast, the number of samples necessary to estimate the mean As concentration in soils over the lower Paleozoic meta-greywacke and slate lithozone is $(181.85/10)^2 = 330.69$. Obviously, the larger coefficient of variation for As concentrations in this lithozone necessitates the collection and analysis of far more samples.

Table 2. Calculated number of samples required to obtain estimates of the mean As concentration at various tolerance levels for the granite and lower Paleozoic meta-greywacke and slate lithozones of Nova Scotia.

<table>
<thead>
<tr>
<th>Tolerance level (%)</th>
<th># of samples over the granite lithozone</th>
<th># of samples the lower Paleozoic meta-greywacke &amp; slate lithozone</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>3038</td>
<td>33069</td>
</tr>
<tr>
<td>2</td>
<td>760</td>
<td>8267</td>
</tr>
<tr>
<td>5</td>
<td>122</td>
<td>1323</td>
</tr>
<tr>
<td>10</td>
<td>30</td>
<td>331</td>
</tr>
<tr>
<td>20</td>
<td>8</td>
<td>83</td>
</tr>
<tr>
<td>30</td>
<td>3</td>
<td>37</td>
</tr>
<tr>
<td>40</td>
<td>2</td>
<td>21</td>
</tr>
<tr>
<td>50</td>
<td>1</td>
<td>13</td>
</tr>
<tr>
<td>100</td>
<td>0</td>
<td>3</td>
</tr>
</tbody>
</table>

The results in Table 2 illustrate that significant differences exist between these two lithozones. The approximately 3 times larger CV, 181.85, observed in soil samples from the lower Paleozoic meta-greywacke and slate lithozone makes it necessary to collect and analyze approximately 10 times more samples to obtain the same relative standard error of the mean in each lithozone. Obviously, the relatively high variability in As concentrations in the lower Paleozoic meta-greywacke and slate lithozone makes estimating the mean more difficult. Nevertheless, using Equation 4 and the strategy described above to determine “how many samples are enough” will allow the NS DNR to estimate, in advance, how many samples should be collected to obtain adequate, or at least the best available, estimates of the average As soil concentration in all eight lithozones within Nova Scotia. Based on the results in Table 2, it is likely that the additional samples that they intend to collect will not be sufficient to obtain mean concentration estimates with highly acceptable errors for all eight lithozones. Nevertheless, this procedure will allow them to obtain the best estimates possible given their survey constraints.

Discussion

Given that elevated As concentrations represent a potential hazard in Nova Scotia, the concentrations of other elements measured in the NASGLP with crustal anomalous background geochemical concentrations should also be assessed to obtain other ‘made in Nova Scotia’ MAC’s, where appropriate (e.g., U, whose concentrations are known to also be elevated in Nova Scotia because of the abundance of peraluminous granite, and the presence of Carboniferous basins that host sediments shed by these eroding intrusions). Such an analysis will provide additional estimates of the numbers of samples necessary to estimate the mean concentrations of other elements for each lithozone, and these numbers of samples are likely to be different than those calculated for As, above. As a result, before collecting additional samples in the follow-up survey, the NS DNR will have to balance their tolerance for mean concentration estimation error for each element in each lithozone with a number of other factors (e.g., bio-accumulation and bio-availability; toxicity; treatment and remediation ease, cost, and success probability; etc.) to determine how many samples to collect from each lithozone, so that they maximize the quality of the resulting data.

Now that we understand all of the factors that control “how many samples are enough”, let us consider the estimation error we would obtain in terms of relative standard error on the mean if we were to use the ‘conventional wisdom’ of 30 samples (Garrett 1979) to estimate the mean As concentration in the granite and lower Paleozoic meta-greywacke and slate lithozones, above. Using Equation 2 and the observed coefficients of variation in each lithozone, we can see that in soils over the lower Paleozoic meta-greywacke and slate lithozone, there will be about 33% error on the mean As concentration estimate, and in soils over the granite lithozone, there will be only about 10% error on the mean As concentration estimate. These precision levels are significantly different, and may or may not be adequate for the purposes described above, likely because “how many samples are enough” is a function of more than just the number of samples (Garrett 1979), as the variability of the data (in absolute or relative terms) and the desired absolute or relative standard error on the mean also figure prominently.

Conclusions

Formulae have been derived that allow determination of how many samples are necessary to obtain an ‘adequate’ estimate of the mean of sample measurements. These derivations do not assume that any specific data distribution exists, making this approach perfectly general and applicable to geochemical (and other) data collected by the mineral exploration/mining (and other) industries and disciplines.

The formulae derived to determine “how many samples are enough” do not support what has become the ‘conventional wisdom’ that 30 samples are adequate to obtain a good estimate of the mean, as other factors such as confidence level and variation in the underlying dataset also exert control. Rather, these results indicate that a quantitative strategy
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can be used to more comprehensively address the unspecified “particular situations” that Bob Garrett alluded to in his 1979 paper when he first investigated the question: “how many samples are enough?”

Acknowledgements

Many thanks are extended to Terry Goodwin, geochemist for the Province of Nova Scotia Department of Natural Resources, for providing the data used in the example to illustrate how to determine “how many samples are enough”. This contribution also benefited from a helpful review by Dr. Robert Garrett of the Geological Survey of Canada.

References


of a single geological entity for which we are calculating a statistic, propagation of the errors in the $x_i$ into $f$ can be achieved using the following generative formula:

$$se_f^2 \equiv \sum_{i=1}^{n} \left( \frac{\delta f}{\delta x_i} \right)^2 se_{x_i}^2,$$  (5)

(Stanley 1990).

In order to derive the standard error of the mean (in this case, our $f$), we propagate the error in $x_i$ through the formula for the mean:

$$\bar{x} = \frac{1}{n} \sum_{i=1}^{n} x_i,$$  (6)

This requires the partial derivatives of the mean with respect to $x_i$:

$$\frac{\delta \bar{x}}{\delta x_i} = \frac{1}{n}.$$  (7)

Substituting these partial derivatives into Equation 5, using the variance of $x$ ($\sigma^2$) as the common variance in all $x_i$'s, and setting all covariances equal to zero, we obtain the well known formula for the standard error of the mean (Equation 1):

$$se_{\bar{x}} = \sqrt{\sum_{i=1}^{n} \left( \frac{1}{n} \right)^2 se_{x_i}^2} = \sqrt{n \left( \frac{1}{n^2} \right) \sigma^2} = \sigma \sqrt{n}.$$  (8)

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**CALENDAR OF EVENTS**

International, national, and regional meetings of interest to colleagues working in exploration, environmental and other areas of applied geochemistry. These events also appear on the AAG web page at: www.appliedgeochemists.org <http://www.appliedgeochemists.org/> Please let us know of your events by sending details to: Steve Amor, Geological Survey of Newfoundland and Labrador, P.O. Box 8700, St. John’s, NL, Canada A1B 4J6; Email: StephenAmor@gov.nl.ca Telephone: 709-729-1161.

**2010**

15-17 September 2010. 11th International Symposium on Environmental Radiochemical Analysis, Chester UK. Website: http://tinyurl.com/yghqp3o


5-9 November 2010. 36th International Symposium on Environmental Analytical Chemistry, Rome Italy. Website: www.iseac36.it <http://www.iseac36.it/>

19-20 November 2010. Eighth Swiss Geoscience Meeting, Fribourg Switzerland. Website: http://geoscience-meeting.scnatweb.ch <http://geoscience-meeting.scnatweb.ch/>

24-26 November 2010. International Conference on Geological Sciences and Engineering, Venice, Italy. Website: http://tinyurl.com/3a98o33 <http://tinyurl.com/3a98o33>


8-11 December 2010. 11th European meeting on Environmental Chemistry, Portorož Slovenia. Website: www.ung.si/~emec11 <http://www.ung.si/~emec11>
2011


2012

6-11 February 2012. 10th International Kimberlite Conference, Bangalore India. Website: http://10ikcbangalore.com <http://10ikcbangalore.com/>


5-15 August 2012. 34th International Geological Congress, Brisbane Australia. Website: www.34igc.org <http://www.34igc.org/>

**RECENT PAPERS**

This list comprises titles that have appeared in major publications since the compilation in EXPLORE Number 146. Journals routinely covered and abbreviations used are as follows: Economic Geology (EG); Geochemica 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.


Afzal, P., et al., 2010. Geochemical anomaly separation by multifractal modeling in Kahang (Gor Gor) porphyry system, central Iran. J. Geochem. Explor. 104(1/2): 34-47.


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Putnis, A. and John, T., 2010. Replacement Processes in the

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**RECENT PAPERS**

*continued from Page 18*

**Modular Course in Exploration Geochemistry**

**Date:** December 9-18, 2010  
**Location:** Laurentian University, Sudbury, Ontario, Canada  
**Information:** Dr. Michael Lesher, Mineral Exploration Research Centre, Department of Earth Sciences, Laurentian University, Willet Green Miller Centre, 935 Ramsey Lake Road, Sudbury, ON, Canada, P3E 2C6  
**Tel.** +1 (705) 675-1151 x2276  
**E-mail:** mlesher@laurentian.ca  
**website:** [http://earthsciences.laurentian.ca](http://earthsciences.laurentian.ca)
Earth's Crust. Elements 6(3): 159-164.


Tanaka, T., et al., 2010. Petrological and geochemical characteristics of intrusive rocks related to porphyry copper mineralization and the implications of the genesis of deposits in the Namosi area, Viti Levu, Republic of the Fiji Islands. Res. Geol. 60(1): 35-


Wang, Y., 2010. Physical and chemical characteristics of the host rocks in controlling the mineralization of the Chinkuashihi high-sulfidation gold-copper deposits, northeastern Taiwan. J. Geochem. Explor. 104(1/2): 61-


The Association of Applied Geochemists announces the 2010 AAG Student Paper Competition

The AAG announces the 8th biennial Student Paper Competition. The paper must address an aspect of exploration geochemistry or environmental geochemistry related to mineral exploration and be based on research performed as a student. The student must be the principal author and the paper must have been published in Geochemistry: Exploration, Environment, Analysis no more than three years after completion of the degree. All eligible papers in 2009 and 2010 volumes of GEEA will be reviewed by the selection panel.

The winner will receive:
A cash prize of $1000CAD generously donated by SGS Minerals Services.

A 2-year membership of AAG, including the society's journal (GEEA), EXPLORE newsletter, publication of an abstract and CV of the winner, a certificate of recognition and $500US towards expenses to attend an AAG-sponsored meeting, courtesy of AAG.

The results of the 2010 competition will be announced at the 25th IAGS in mid 2011. Details are available from the chair of the committee or the AAG Students' page (http://www.appliedgeochemists.org/).

David Cohen
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WELCOME TO ROVANIEMI

The 25th International Applied Geochemical Symposium of the Association of Applied Geochemistry (AAG) will take place in Rovaniemi, Northern Finland, from 22-26 August 2011. The meeting will focus on applied geochemistry under the theme Towards sustainable geochemical exploration, mining and the environment, which encompasses a variety of disciplines, including applied geochemistry, new ways of analysis, interpretation of data and the importance of taking care of the environment in mineral exploration and mining. A technical programme, special sessions, workshops, and pre and post-excursions are designed to support the theme.

IMPORTANT DATES AND DEADLINES

Second Circular and Call for Abstracts  30 September 2010
Third Circular                              31 January 2011
Deadline for abstracts                     31 March 2011
Notification of acceptance sent by         30 April 2011
Deadline for early-bird registration       31 May 2011
25th IAGS Conference                       22-26 August 2011
Deadline for Special Issue submission      15 November 2011

For further information check the conference website at: www.iags2011.fi
or contact the conference office (congress@ulapland.fi)
This volume is a companion to a drift exploration short course conducted at the Geological Association of Canada (GAC) 2007 Meeting in Yellowknife. The short course and this publication was sponsored by the Mineral Deposits Division of the GAC with additional sponsorship from the Alberta Geological Survey, Geological Survey of Canada, Overburden Drilling Management Limited, Apex Geoscience Limited and Shear Minerals Limited. The authors represent a wide range of specialties and possess many years of experience in their particular fields of interest. Federal and provincial geological surveys as well as academia and the exploration industry have all contributed to this volume.

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Full Reference:
EXPLORE Publication Schedule
Quarterly newsletters in March, June, September, December

Deadlines for submission of articles or advertisements:
- March newsletter: January 15
- June newsletter: April 15
- September newsletter: July 15
- December newsletter: October 15

Information for Contributors

Manuscripts should be double-spaced and submitted in digital format using WORD. Photos and figures (colour on black and white) should be submitted as separate digital files and as high resolution jpeg or PDF files. Tables should be submitted as separate digital files in EXCEL format. All scientific/technical articles will be reviewed. All contributions may be edited for clarity or brevity.

Formats for headings, scientific notations, references and figures must follow the Guide to Authors for Geochemistry: Exploration, Environment, Analysis (GEEA) that are posted on the GEEA website at: http://www.groslc.org.uk/template.cfm?name=gkea_instructions_for_authors

Submissions should be sent to:
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