
FIELD PORTABLE XRF ANALYSERS

GEOS Mining
“The Consultants Experience”
2007 – 2009

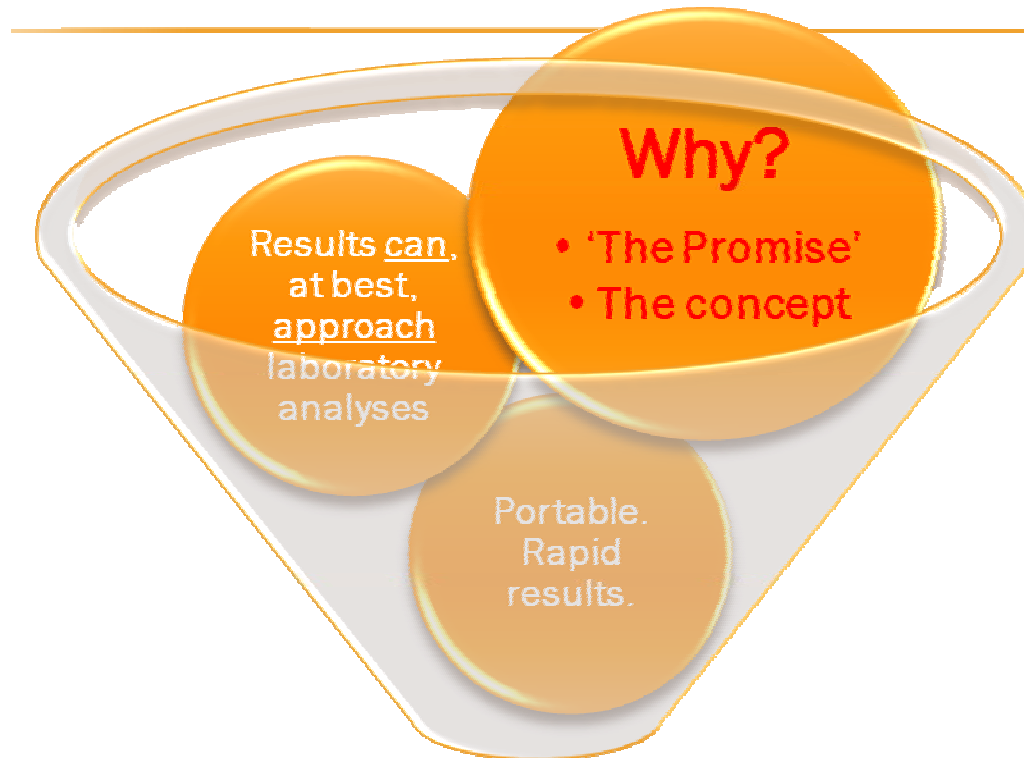


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JUSTIFICATION



Efficiency Cost Saving

- Quicker and larger areal coverage
- Enabling modification/adaption of the ongoing work program

Niton XRF analyser (Niton Mining Brochure)



A quoted ... "Element Range of Detection: Potassium (K) through to Uranium (U)" ... "detection limits down to ten parts-per-million (ppm)"

The image shows a standard periodic table of elements. The elements are arranged in rows and columns, with their atomic numbers and chemical symbols. The table includes elements from Hydrogen (H) to Oganesson (Og). The elements are color-coded by groups: Group 1 (purple), Group 2 (blue), Groups 3-10 (various colors), Groups 11-18 (various colors), and the f-block (various colors).

Typical 'experienced' target elements & 'practical' detection limits:

Element	Detection Limit Range (ppm) *
Cu, Pb, Zn	10-100
Ag	50-150
As	10-100
Co	10-100
Cr	20-100
Mo	10-100
Ni	10-100
Sn	50-150
U	20-100
W	50-100

(* ..Soils.. powders, fine grained homogenous materials)

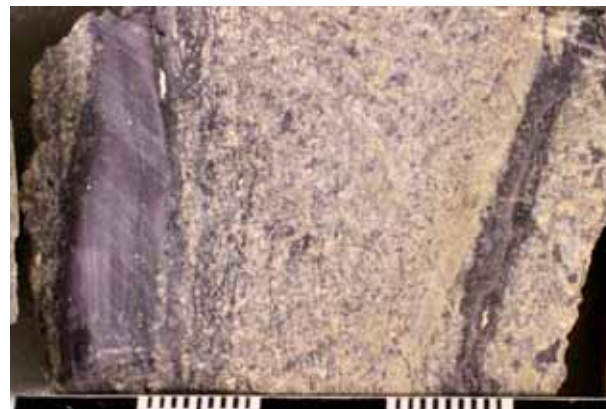
APPLICATION

- **In situ Geochemical surveys**
 - **Soil**
 - **“Termitus”**
- **Mineralisation discrimination**
 - **Drill core**
 - **Rock samples**
 - **Drill chip**
 - **Vein sets**



Termite mound survey

F-U-Mo veins



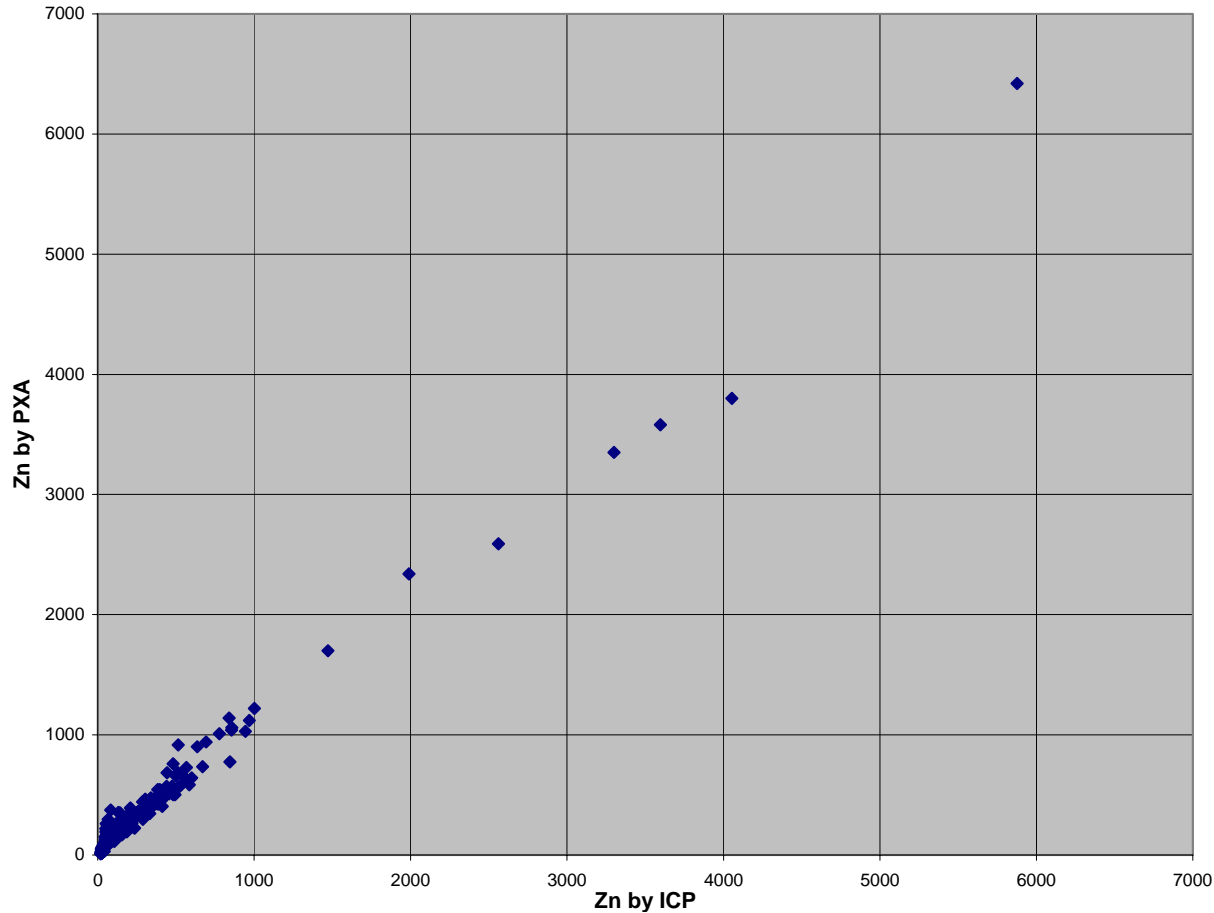
Vein gossan



"THE EXPERIENCE"

Soil survey FP XRF vs. laboratory ICP assay (ppm)

Zinc pxa vs lab ICP



Good Correlation due to:

- Fairly homogenous fine grained soil
- Relative low detection capability for Zn by FP XRF
- Low error for XRF Zn

Soils

Pb, Zn, As, Fe, Ti



U, Mo, Cu



Sn, Bi, W



Rock/Core/Chip

Massive Mineralisation



Vein / Disseminated Mineralisation



TYPICAL ISSUES

- Generally high detection limits
- No direct analysis for gold
- Spot readings
 - Often unrepresentative, rock/core/chip
 - Ground homogenised materials ideal
- Permit users to become analysts
 - Without training OR understanding
- Great risk of errors



Great need for Pre survey planning

- WHAT ,WHAT, WHAT AND WHAT??

.... CONTINUED.....

- ✘ Masking of level soil anomalies
 - + e.g. Cu error consistently >120 ppm masking low level Cu anomalism
 - + Recommend at least 10% of survey stations/points sampled chemical analysis

- ✘ Contamination
 - + Cover instrument window with thin plastic (cling film)
 - ✘ Beware !!! - Anomalous Zn – due to green plastic cover
 - + Plastic bagged samples analysed through plastic
 - + Daily calibration of instrument



- ✘ “Spike” readings
 - + High values not reproduced in readings taken close to or at the original site
 - + Probably caused by
 - ✘ Small rock/mineral fragments in soil surveys
 - ✘ Imperfect PF XRF window fit to sample
 - ✘ High moisture
 - + Take repeat readings, record them as such



- ✘ High nugget effect from core or rock samples
 - + Operator bias
 - + One 'spot' reading on core not representative of one metre
 - + Procedures to randomize readings
 - ✘ multiple/repeat readings

- ✘ Erroneous results
 - + Unusual matrices
 - + Weathering effects on rock surfaces
 - + Instrument introduced errors
 - ✘ Bad fit (instrument vs. operator ?)
 - ✘ Low battery
 - ✘ Over heating

- ✘ Data not validated by laboratory analysis
 - + MUST be regarded as indicative ONLY
 - + MUST be clearly recorded as such



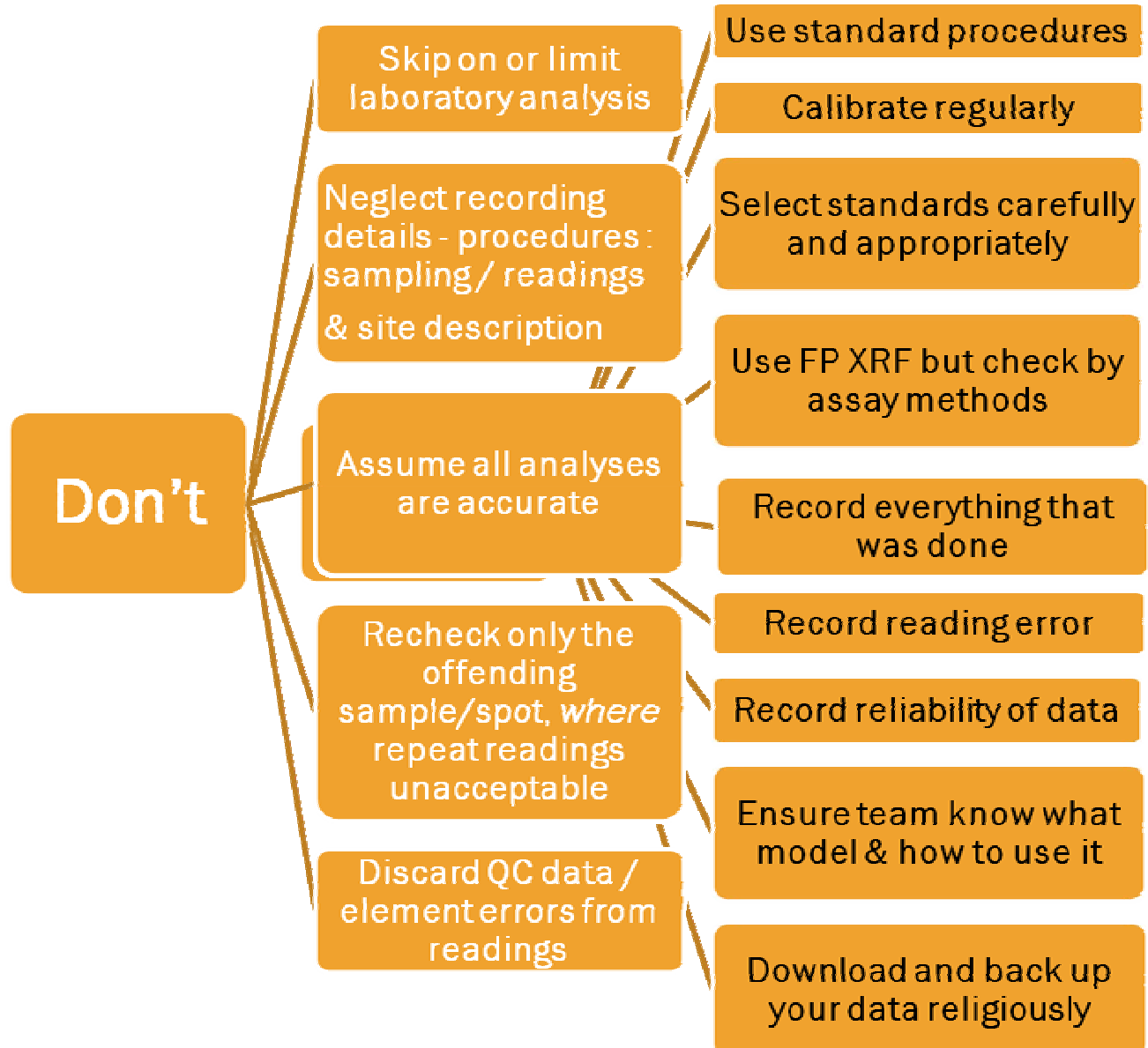
CONCLUSIONS

Poor methodology
and hence data will
lead to poor choices

Quality control of the
data is critical

Pre-program setup
of FP XRF is crucial

- Includes understanding limits of methodology and instrument
- Style and type of target mineralisation (including pathfinder elements)
- Materials to be tested
- Likely errors that will be encountered / introduced





FIELD GEOLOGIST QUOTE

“....TO GET THE MOST OUT OF (FP XRF), IT IS ALL ABOUT CONSISTENCY, CONSISTENCY, CONSISTENCY!”



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