

Tobia Howell's Ceramic Materials

THE ACCIDENTAL GLAZE PROJECT

"Name the greatest of all inventors. Accident."

Mark Twain

Has this happened to you? Out of the kiln comes a new set of glaze tests. One of the tests is incredibly beautiful!! It's a blood red at cone 6 or a green glaze with only cobalt as a colorant or maybe it's just the best glaze you've seen. You want to share it with the world! But then you realize:

- a) That was the test you tossed in 10 grams of... (oh gosh what was that) just to see what would happen.
- b) It's the garbage (scrap) glaze
- c) You know what's in it then try to reproduce it and it doesn't work. It could be the firing cycle but you can't be sure.
- d) You were sure you would remember that formulation

Usually when we design glazes we do just that. There is a huge amount of information out there to help you guide your testing process. What we don't do often enough is go outside those guidelines. Over 30% of great discoveries are accidental. Scrap or garbage glazes are a perfect example, rarely are they just 5 or 6 ingredients in intelligent well thought out proportions. Likely you will have upwards of 15 different ingredients. This large number of variables can have the most amazing results. Remember how hard it was to reproduce "Albany Slip"?

Enter "The Accidental Glaze Project"

Project goals:

The goal of this project is to gather a very large number of accidental glazes (any glaze with an accidental component to them). Through the efficiencies of numbers and the donations of volunteers (testing and administration) we reduce the cost of a full spectrum mineralogical analysis from \$100 USD to less than \$15 USD (The goal is \$10 USD)

From this we make a publication and you or your studio can get full credit for its discovery.

PS (This is a green project encouraging potters to keep their cast away glazes instead of discarding them.)

There are many roles to take in this project. You can be a partner, participant or something in between. If you are interested in finding out what your accidental glaze is, or you want to be published or you just can't wait to see what everyone's accidental glaze contains, or you're a glaze intellectual and want to see what new discoveries this project can uncover.

Let me know if you would like to participate and what role you would like to play. This is definitely a group project.

Tobia's Samples as Received



When and how to use xrf to measure ceramics?

The answers to this question are simple and based on fundamental physics, which we have not yet found a way to violate:

- •One can never determine the weight percent of a non-uniform material because it is non-uniform UNLESS you covert it to a uniform material.
- •Nor can one ever use xrf to report content it cannot measure like C and O and H. So the content listed in this calibration is only the weight percent of the elements that are measured. It does not add up to 100 percent.
- •The Tracer spot size is 3 by 4 mm, if your substance has a mixture of particles smaller than 0.2 mm and they are well mixed then you will get a reasonable answer relative to wt %
- •A detailed calibration with the ceramic materials that were provided was done after using the raw Spectra from the Tracer to verify the given values validity.
- •If you material is a conglomerate then it is <u>not</u> uniform and the only way to get a reasonable estimate of its average elemental content is to take a large quantity of it grind it up and then press it into a pellet, there is no other method that will work.
- •How did Bruker check / corrected for any matrix effect due to material containing heterogeneous particles with size larger than mud (silt/clay)? Actually you have to do this by grinding your material up and pressing it to pellets. See item 5 $_{\odot}$
- •A calibration that is very accurate on uniform ceramics without preparation and non-uniform material ground and pressed in to pellets.
- •Note ceramics or other material with paint, slip, and glaze or any covering like dirt on them are VERY non-uniform!
- •Your sample also has to be infinitely thick relative to the range of the emitted photon from each element of interest. See table on next page, you sample must be thicker than the values list for each element noted.

| Element | Photon Emitted energy | Analysis depth in |
|---------|-----------------------|-------------------|
| | (keV) | Ceramic(cm) |
| | | |
| 0 | 0.53 | 0.00001 |
| Na | 1.04 | 0.0007 |
| Mg | 1.2 | 0.00096 |
| Al | 1.47 | 0.0017 |
| Si | 1.74 | 0.0027 |
| Р | 2.01 | 0.0013 |
| Ca | 3.69 | 0.0064 |
| Cr | 5.41 | 0.0192 |
| Fe | 6.4 | 0.03 |
| Cu | 8.01 | 0.058 |
| Zn | 8.64 | 0.077 |
| Pb | 10.55 | 0.113 |
| Zr | 15.78 | 0.384 |

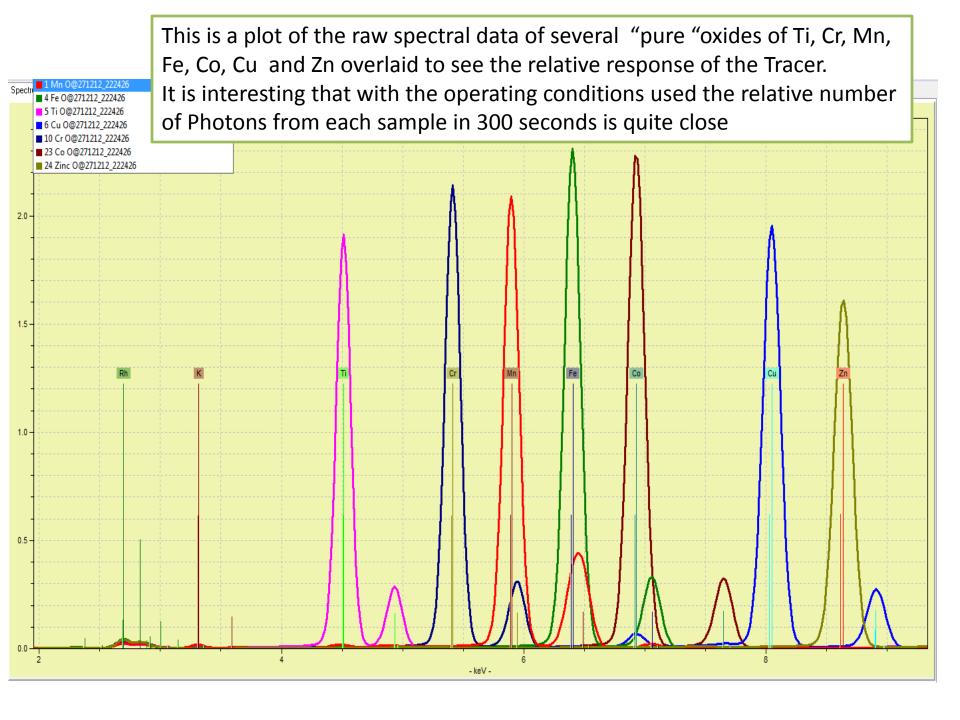
System and settings

A Bruker Tracer IV SD system was used

- 1. No beam filter
- 2. Tube voltage was 15 kV
- 3. Tube current was 44 micro amps
- 4. Vacuum at 4 Torr
- 5. Each sample was placed 3 mm thick in a 1.25" sample cup
- 6. Each analysis was done for 300 sec

THE SAMPLES AS ANLYZED IN XRF ANALYSIS CUPS WITH 4 MICRON POLY WINDOWS

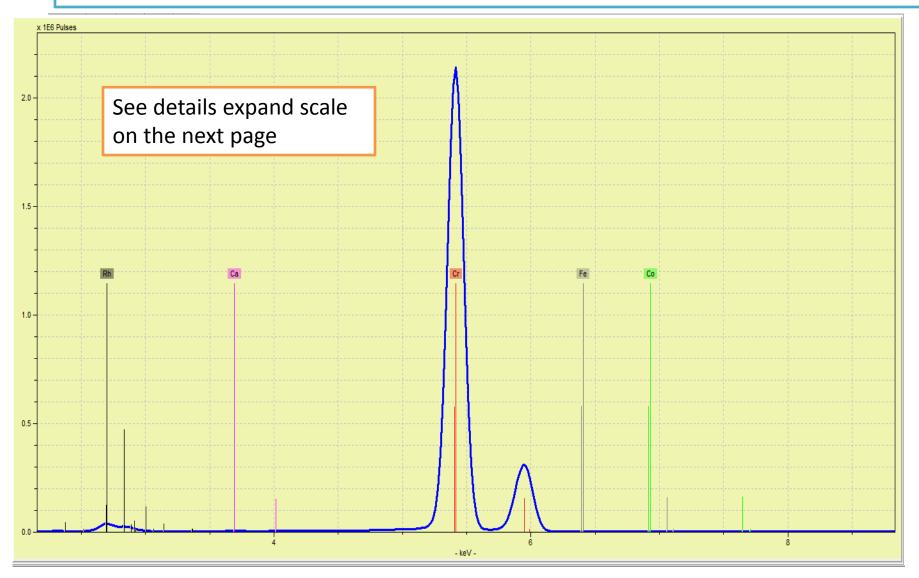




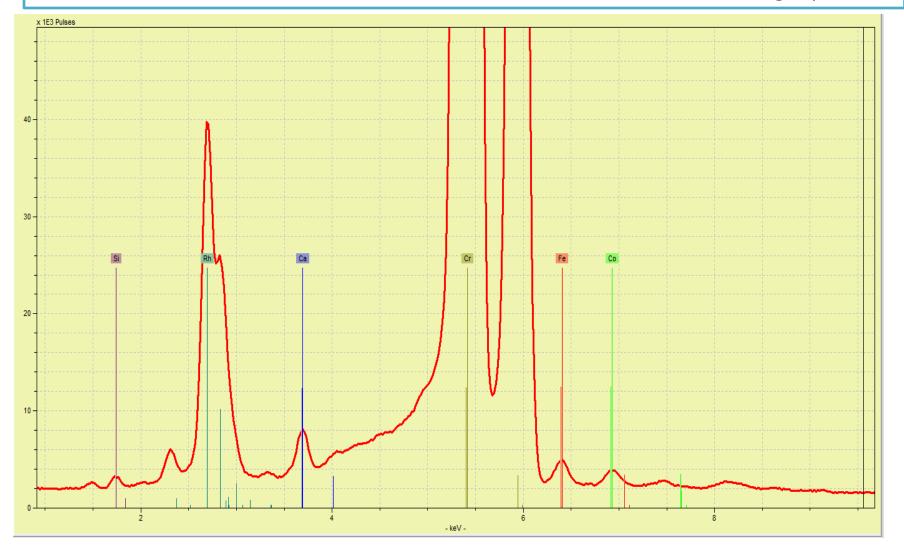
The following is a quick look at some of the "pure" oxides.

Note the Tracer detected trace elements as noted, these were not indicated in the given material content!

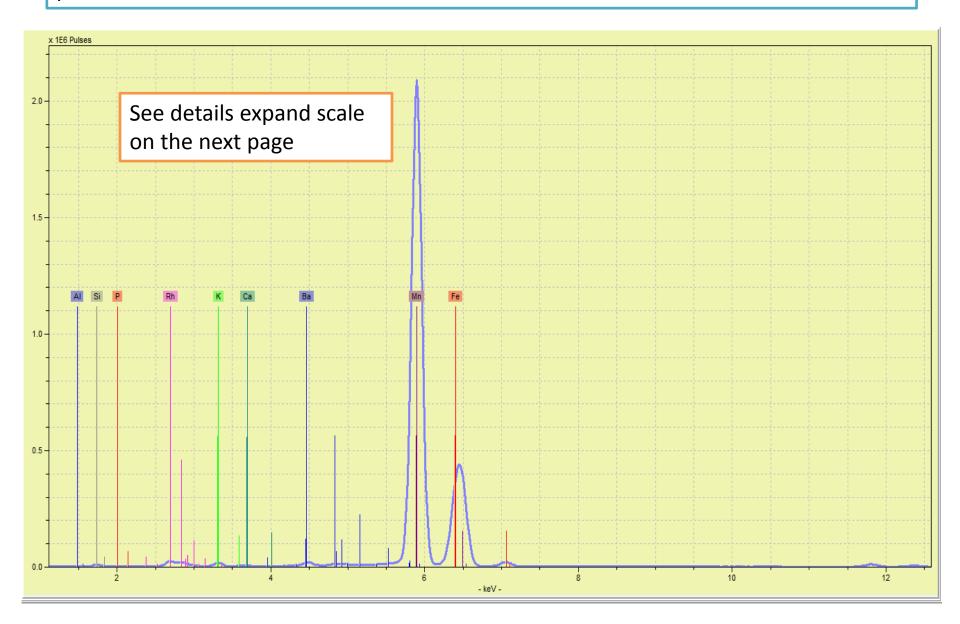
<u>Cr oxide</u>: trace elements are Si , Ca, Fe and Co these are at around 0.01 weight percent



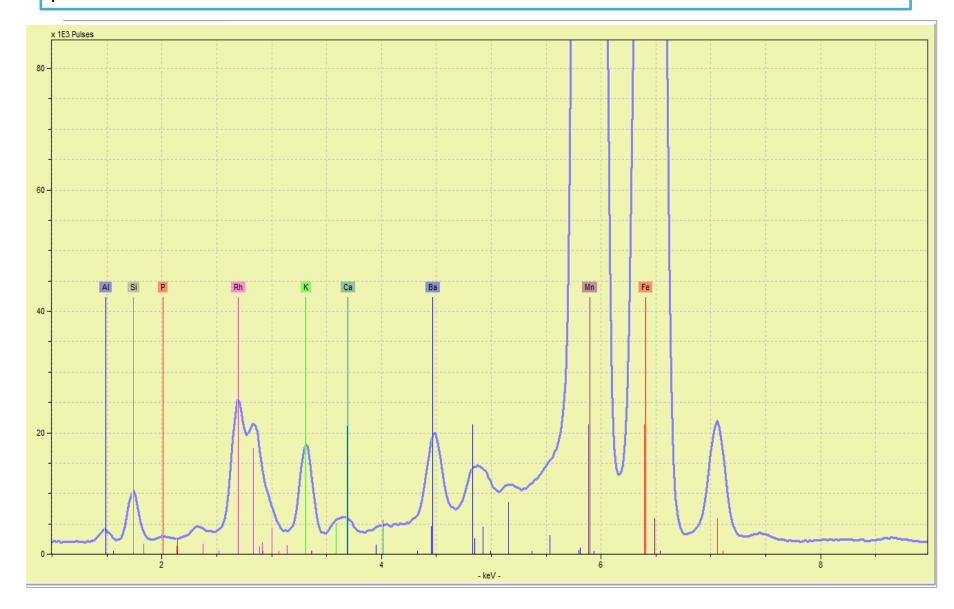
Cr oxide: trace elements are Si , Ca, Fe and Co these are at around 0.01 weight percent



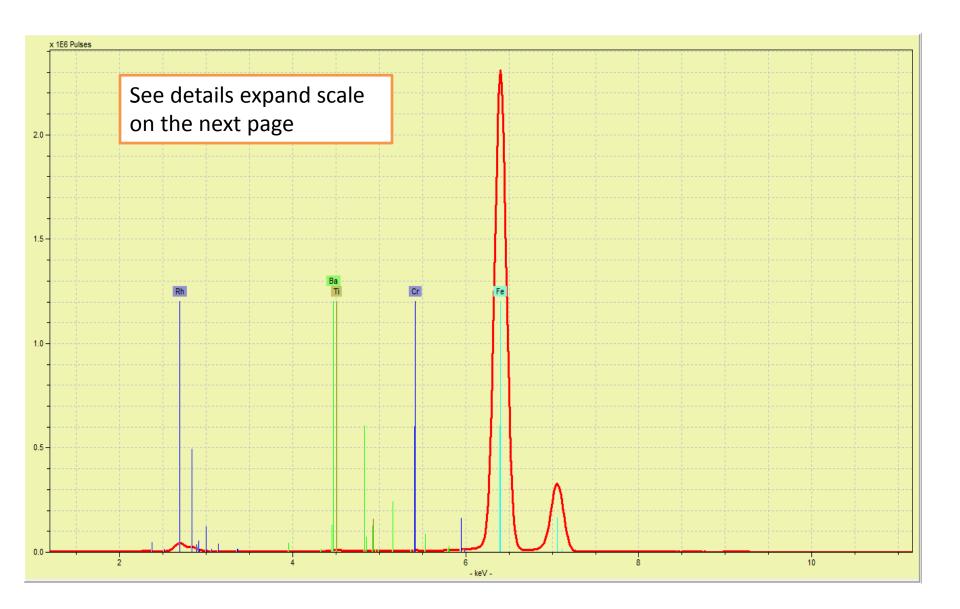
Mn oxide: trace elements are Al, Si, P, K, Ca, and Fe these are at around 0.01 weight percent or less



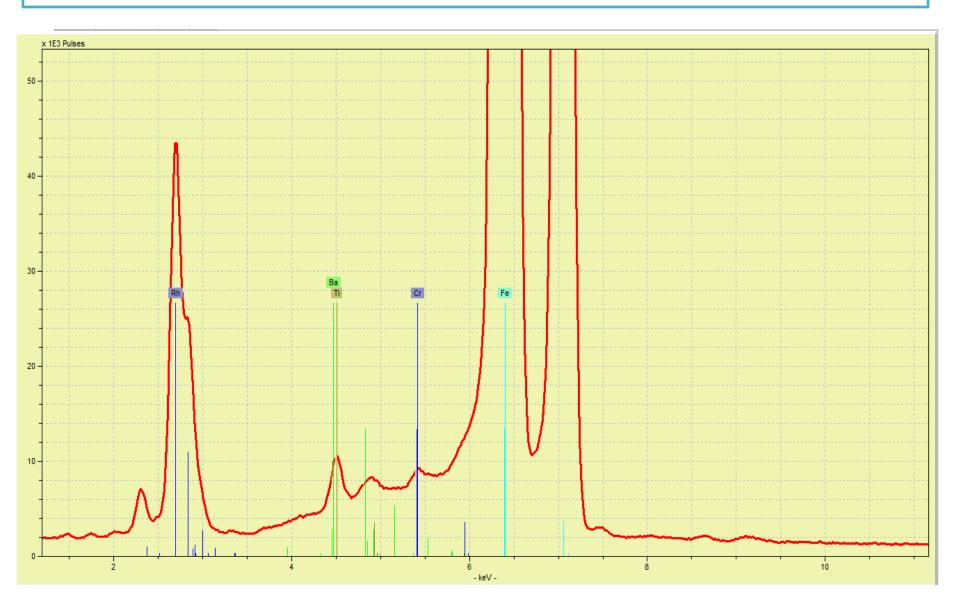
<u>Mn oxide</u>: trace elements are Al, Si , P, K, Ca, and Fe these are at around 0.01 weight percent or less



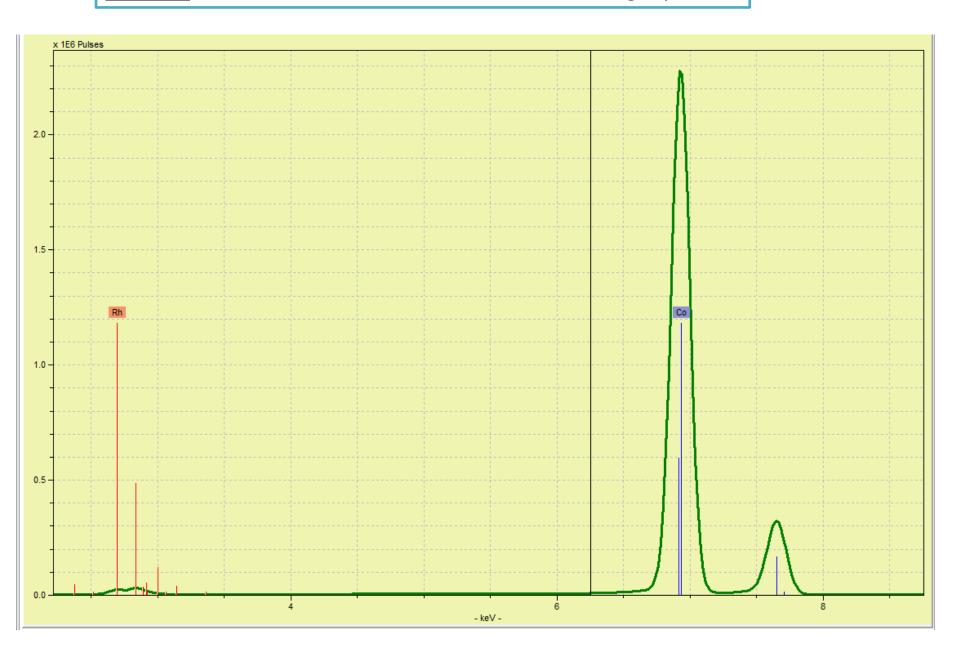
Fe oxide: trace elements are Ba, Ti and Cr these are at around 0.01 weight percent or less



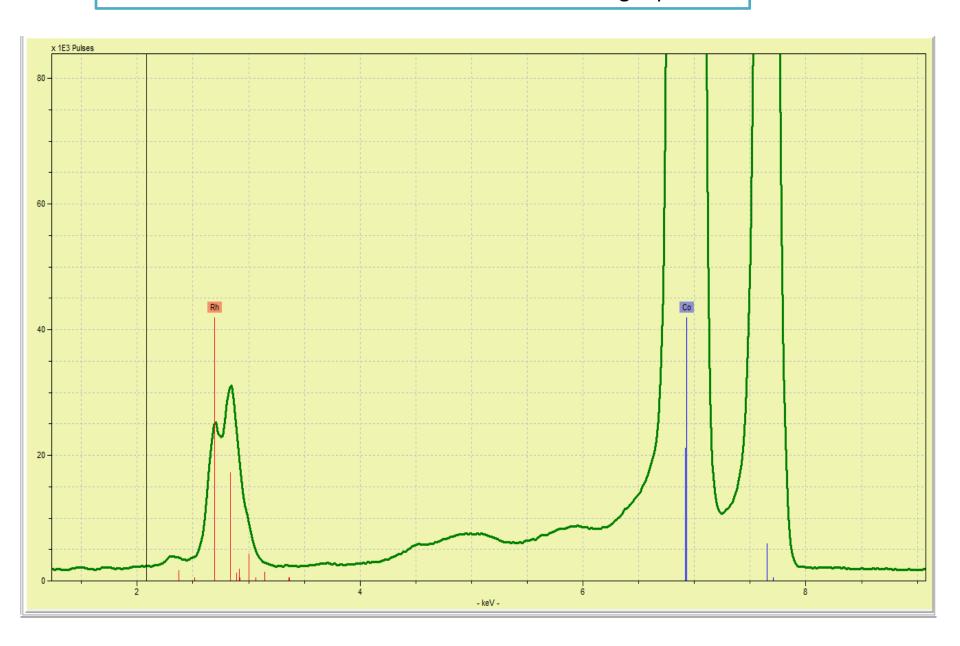
Fe oxide: trace elements are Ba, Ti and Cr these are at around 0.01 weight percent or less



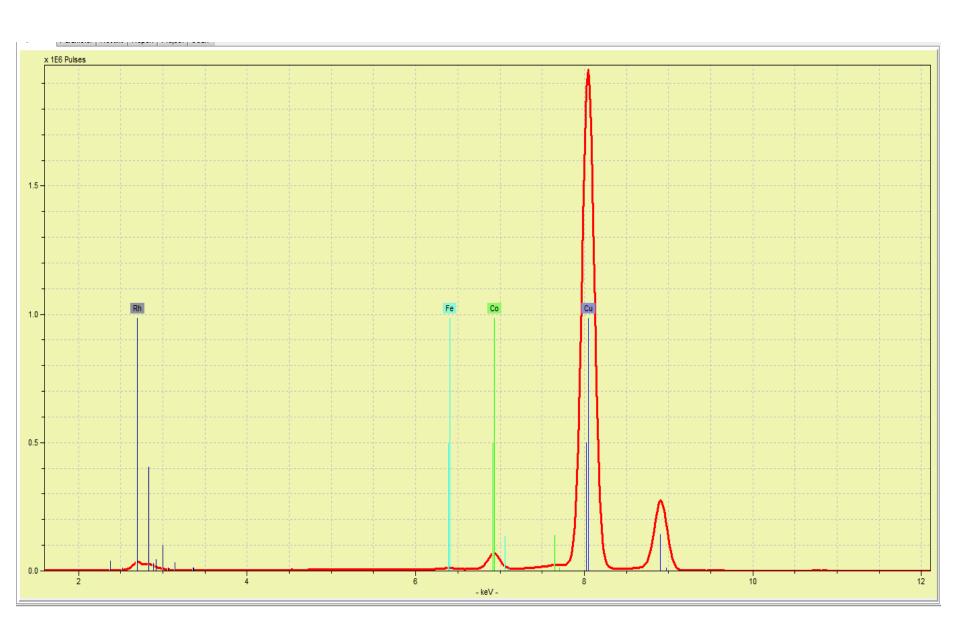
Co oxide: there are no trace elements above 0.01 weight percent



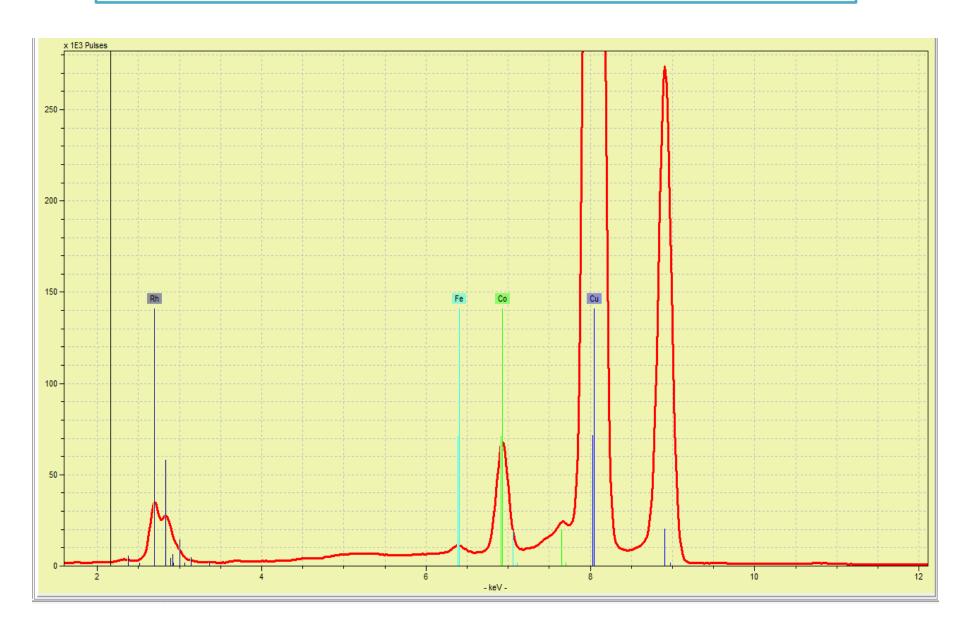
Co oxide there are no trace elements above 0.01 weight percent



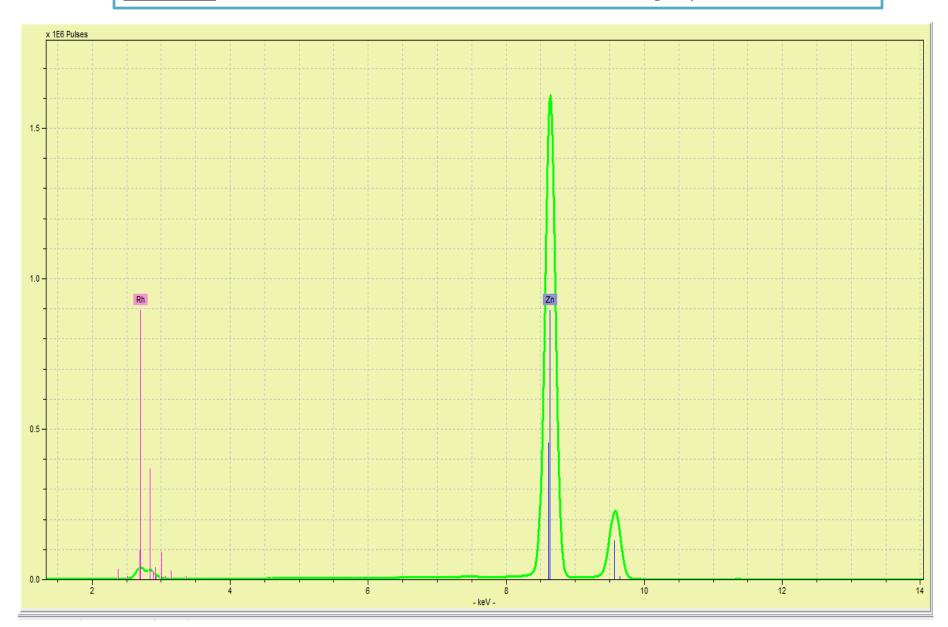
Cu oxide with Co above at about 2.0 weight percent and a trace of Fe.

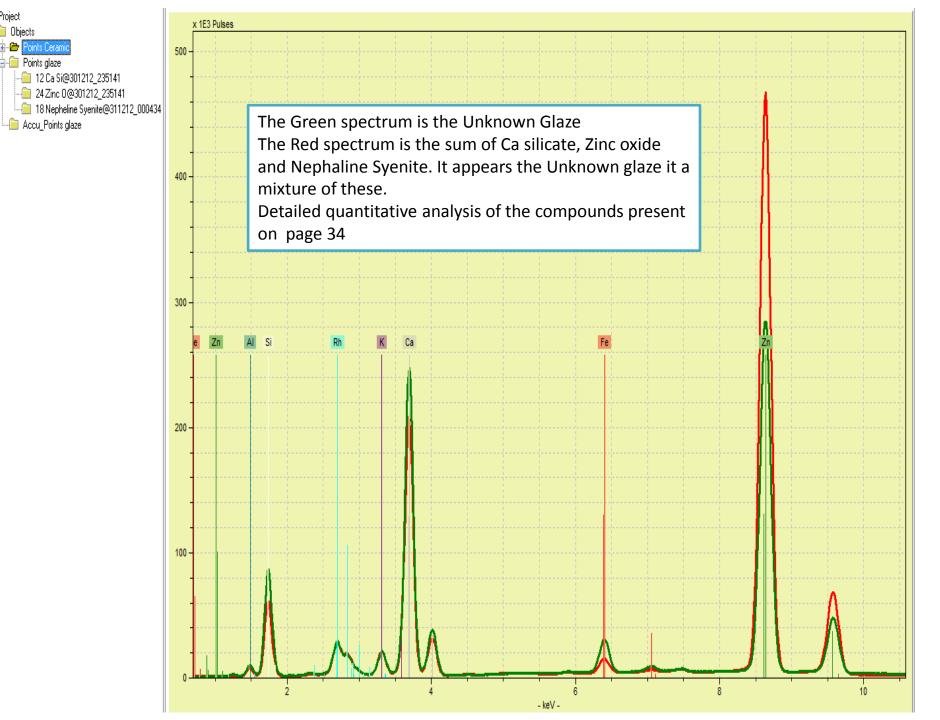


Cu oxide with Co above at about 2.0 weight percent and a trace of Fe.



Zn oxide: there are no trace elements above 0.01 weight percent





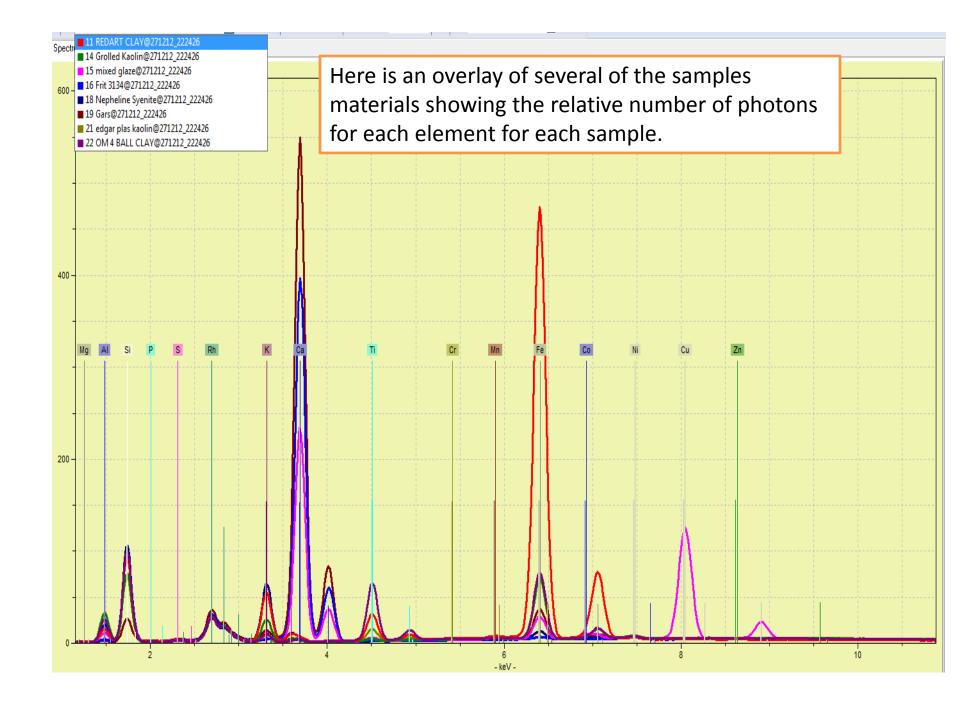
Project

Objects 🖫 🗁 Points Ceramic

🗀 🧰 Points glaze



Bruker has software that can quickly give one the net number of photons from each element in a given spectra. This allows one to quickly compare the relative concentration of any element in any material, uniform or non-uniform. One can do up to 50000 spectra at once. The software is called ARTAX. This was done to all the data that was taken on the samples sent. Direction and the result of this analysis follows.



STEPS REQUIRED TO DO SPECTRA ANALYSIS USING ARTAX SOFTWARE provided with the Tracer

This gives the net number of photons form each element for each spectrum analyzed

Net area analysis in Artax 7

- 1. In ARTAX, click on 'File', click on "open spectrum", file the folder that has all your txt spectrum files, highlight them all and click open
- 2. Click on "project", click on "new project", right click on "object", click on "add node", enter "Points" in name box. Highlight this folder.
- 3. Go back to "Project" tab. click on "Add spectrum".
- 4. Click on "File", click on "save project", give a name (.rtx), click on "save"
- 5. Click on "spectrum" tab
- 6. Go to method list and pick your method! See below
- 7. Highlight the Points folder
- 8. Click on "Analyze" and then on "Evaluate Results", a progress bar should appear as all the spectra are being evaluated with "your named method"
- 9. Then click on "Export" and then on" Results to Excel" Then a box will appear so you name the excel file and put it in a folder that you want the results to be in!
- 10. Now immediately resave your project file because it now contains your spectra and your results. Use the same name you did before and save on top of the old version of the rtx file.
- 11. Now go to the folder you saved your results in and open the file and got to the Points tab to see all your net area data. You then edit out the area that gives you no information.

Method Creation

- •To create a method open a spectrum that is typical of the spectra you want to analysis, get the periodic table and LABEL ALL THE ELEMENTS THAT ARE IN THE SPECTRUM, YOU CAN NOT SKIP ONE JUST BECAUSE YOU ARE NOT INTERESTED IN IT, YOU MUST LABEL ALL THE ELEMENTS THAT THE SYSTEM HAS DETECTED IN YOUR SAMPLE.
- •Then click on the Method editor that is to the left of the method name.
- •Click on Identification and make sure the dot is in the Preset list option to the left of the periodic table
- Then click Get elements
- •Then go to the Name box and type in whatever name you would like for your method
- •Click on Corrections then set cycles at 9 and then pick your energy range for fitting, typically the range of analysis.
- •Then click on add

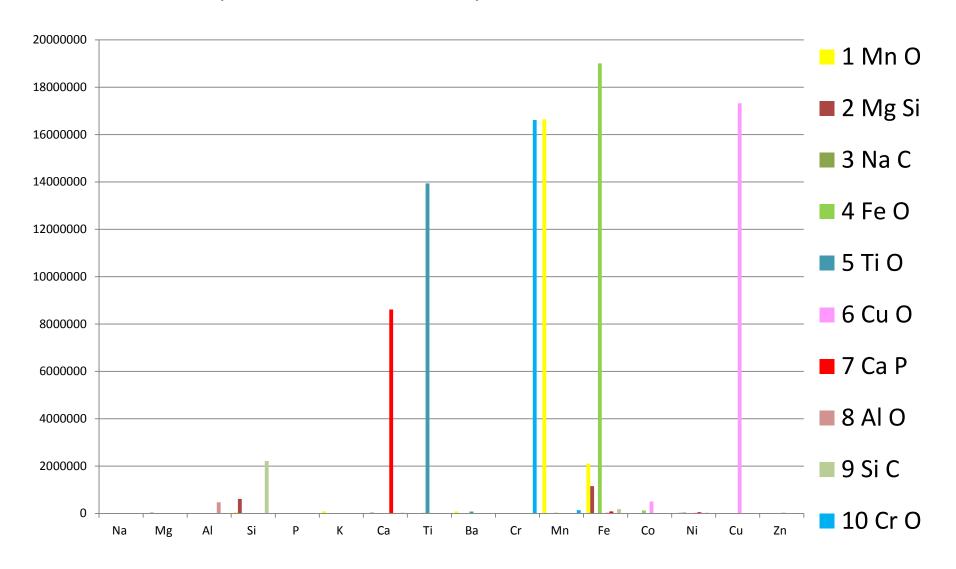
Then click on ok at the bottom of the popup method editor window. Your method should now be in the method window at the top of the Artax screen

Net number of photons for each element in each sample in 60 seconds This is proportional to the concentration of each element in each sample

| | Na | Mg | Al | Si | Р | К | Ca | Ti | Ba | Cr | Mn | Fe | Co | Ni | Cu | Zn |
|----------------------|------|-------|--------|---------|--------|--------|---------|----------|--------|----------|----------|----------|----------|--------|----------|----------|
| 1 Mn O | 1 | 177 | 10578 | 46353 | 1879 | 92855 | 13107 | 19216 | 87534 | 7888 | 16643255 | 2110582 | 2 | 8867 | 2387 | 7963 |
| 2 Mg Si | 1 | 31508 | 3732 | 613291 | 179 | 1 | 32112 | 530 | 2725 | 9108 | 7534 | 1154827 | 21423 | 26041 | 2727 | 3163 |
| 3 Na C | 4153 | 198 | 943 | 1788 | 1238 | 1 | 2529 | 2772 | 3729 | 1 | 3210 | 13474 | 10450 | 41865 | 863 | 1206 |
| 4 Fe O | 1 | 96 | 2301 | 1800 | 1246 | 1 | 862 | 28687 | 12852 | 9565 | 35148 | 19005647 | 124631 | 3112 | 2449 | 3900 |
| 5 Ti O | 1 | 428 | 3011 | 955 | 5159 | 14014 | 2019 | 13937706 | 78256 | 288 | 1703 | 2372 | 75 | 15729 | 4945 | 29 |
| 6 Cu O | 1 | 1 | 1388 | 1352 | 1038 | 0 | 6256 | 1 | 4851 | 13052 | 1 | 37749 | 504050 | 31395 | 17324804 | 40118 |
| 7 Ca P | 272 | 1469 | 2639 | 14025 | 2699 | 12276 | 8614015 | 2155 | 834 | 1 | 13711 | 89004 | 6252 | 52747 | 264 | 2581 |
| 8 AI O | 66 | 3387 | 474538 | 1002 | 1245 | 1 | 4535 | 1531 | 11821 | 5447 | 326 | 16987 | 10223 | 18353 | 10577 | 15868 |
| 9 Si C | 1 | 261 | 18005 | 2216848 | 62 | 0 | 6235 | 14153 | 12355 | 17018 | 3615 | 180016 | 13228 | 35668 | 9012 | 5024 |
| 10 Cr O | 1 | 142 | 3273 | 5460 | 602 | 402 | 29834 | 0 | 3395 | 16611361 | 143817 | 11645 | 12403 | 6006 | 6025 | 2183 |
| 11 REDART CLAY | 1 | 755 | 82892 | 546175 | 538 | 347937 | 28250 | 203553 | 7048 | 5874 | 21966 | 3846650 | 31933 | 15781 | 4332 | 6149 |
| 12 Ca Si | 1 | 256 | 11691 | 605750 | 1561 | 4255 | 4929616 | 17040 | 2792 | 3019 | 38754 | 248444 | 4381 | 22179 | 780 | 490 |
| 13 Si O | 23 | 357 | 11218 | 1382275 | 1 | 1 | 1703 | 274 | 4175 | 5285 | 10533 | 23601 | 3883 | 24898 | 191 | 2634 |
| 14 Grolled Kaolin | 1 | 527 | 176041 | 441419 | 1034 | 153298 | 8899 | 6504 | 8707 | 3498 | 11148 | 545805 | 5193 | 26102 | 6952 | 5190 |
| 15 mixed glaze | 1 | 1 | 57710 | 569915 | 1042 | 37902 | 1609432 | 23813 | 3479 | 3386 | 20083 | 204692 | 39443 | 19971 | 1069684 | 9120 |
| 16 Frit 3134 | 647 | 49 | 16605 | 555550 | 965 | 2746 | 2726678 | 10229 | 417 | 1723 | 3475 | 27214 | 3158 | 22506 | 1139 | 1646 |
| 17 custer fledspar | 38 | 1 | 80264 | 808351 | 1150 | 631258 | 24335 | 1 | 1317 | 2444 | 9152 | 99903 | 2184 | 20199 | 1981 | 1009 |
| 18 Nepheline Syenite | 620 | 1 | 110665 | 622034 | 257 | 416283 | 23369 | 0 | 767 | 3460 | 5830 | 67885 | 2586 | 25517 | 1542 | 1999 |
| 19 Gars | 1 | 3326 | 6032 | 150224 | 1089 | 54798 | 3779967 | 12835 | 3019 | 719 | 14766 | 268801 | 5786 | 26566 | 1146 | 1796 |
| 20 Ca Mg | 1 | 19155 | 1138 | 15556 | 1255 | 3204 | 5418443 | 0 | 766 | 710 | 11872 | 303621 | 5016 | 40708 | 784 | 624 |
| 21 edgar plas kaolin | 1 | 105 | 188014 | 434889 | 2405 | 22898 | 35339 | 79169 | 12720 | 7574 | 8894 | 523936 | 4731 | 29791 | 4031 | 4967 |
| 22 OM 4 BALL CLAY | 1 | 166 | 145542 | 575809 | 922 | 50793 | 17736 | 446150 | 14085 | 8403 | 6537 | 594267 | 6660 | 26246 | 2697 | 5378 |
| 23 Co O | 1 | 31 | 2132 | 1008 | 598 | 0 | 2761 | 4757 | 13864 | 1 | 14415 | 21644 | 19379759 | 117669 | 248 | 1055 |
| 24 Zinc O | 1782 | 1 | 1601 | 941 | 678 | 1 | 2438 | 1 | 8589 | 14575 | -92 | 16432 | 25072 | 38794 | 4955 | 14725926 |
| 25 Sr C | 1 | 1 | 8347 | 536313 | 73729 | 1 | 6825 | 1 | 123630 | 1690 | 11134 | 4716 | 5228 | 42816 | 193 | 5843 |
| 26 rutile | 1 | 196 | 3901 | 9889 | 20622 | 2650 | 8278 | 14509075 | 81718 | 13926 | 4681 | 345027 | 5866 | 8214 | 9662 | 2611 |
| 27 Li C | 1 | 176 | 1333 | 1447 | 1343 | 7284 | 12681 | 12058 | 9146 | 1117 | 3 | 9714 | 12784 | 67175 | 6943 | 792 |
| 28 Zr Si | 678 | 135 | 1561 | 309379 | 806030 | 1 | 1497 | 9738 | 16846 | 3419 | 507 | 34875 | 24060 | 19282 | 148283 | 7638 |
| Unknown glaze | 1 | 7062 | 46846 | 506829 | 650 | 116217 | 1696162 | 1 | 2439 | 3885 | 6369 | 216702 | 4631 | 25123 | 558 | 2561620 |
| Unknown | 1 | 14515 | 30779 | 262413 | 2685 | 88351 | 631435 | 487784 | 5304 | 55900 | 339637 | 1074097 | 218790 | 15184 | 1273513 | 1895135 |

Note the raw spectral data is never wrong.

This is a plot of the net number of photons for the Oxides indicated

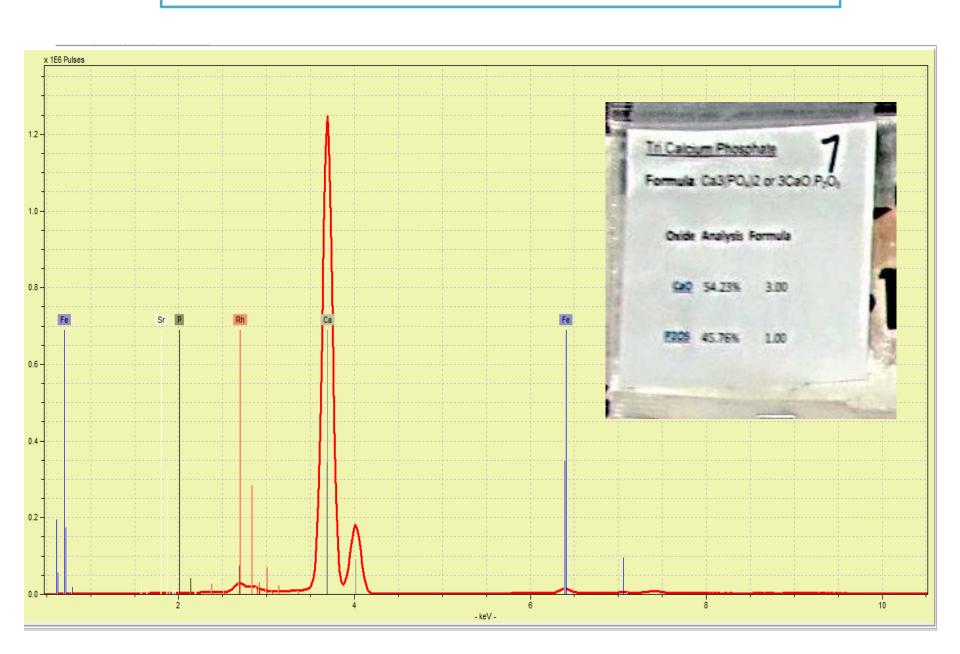


Note the raw spectral data is never wrong.

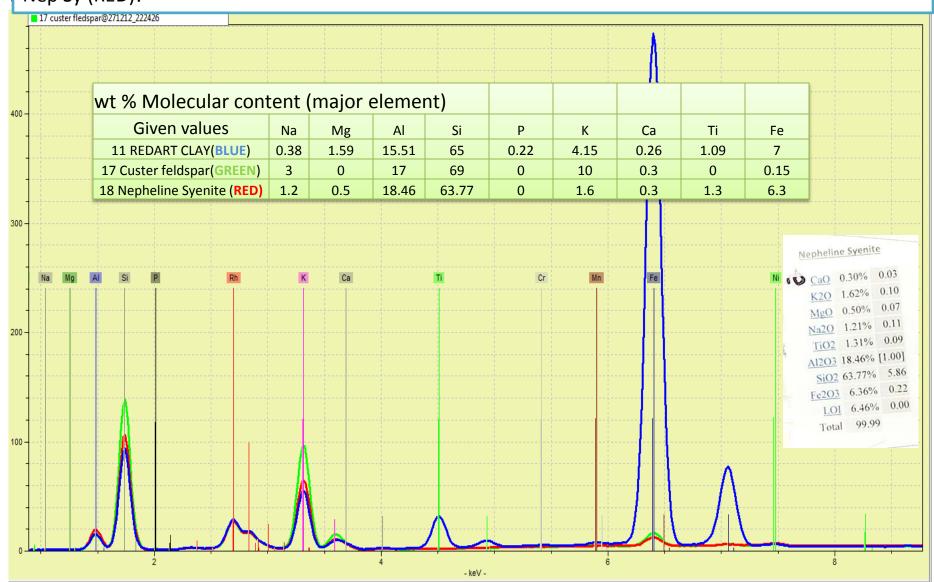
The samples shown in the following slides have raw spectra that are inconsistent with the given content.

Note the raw spectral data is never wrong.

There a problem with the Ca Phosphate: There is no Phosphorous!



<u>There is a problem with the Nepheline Syenite (RED):</u> The raw spectrum shows too much K and way to little Ti and Fe relative to the given concentrations in the table. Note the raw elemental spectrum comparing it to Redart (BLUE) and Custer(GREEN) and the table of given values. I suspect this is not Nep Sy (RED)!

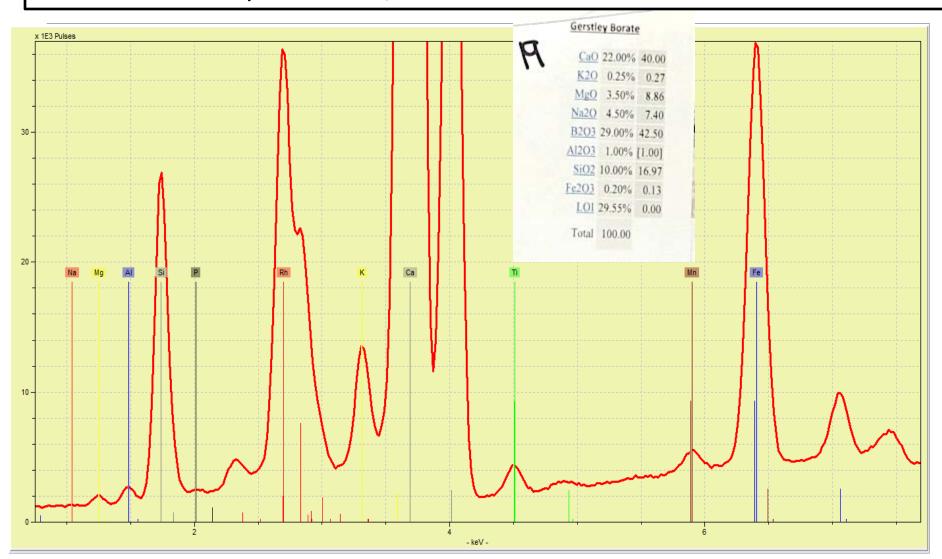


The samples shown in the following slides have raw spectra that the major element contents appear correct but the trace elements apparent in the raw spectra are not in the given content.

Note the raw spectral data is never wrong.

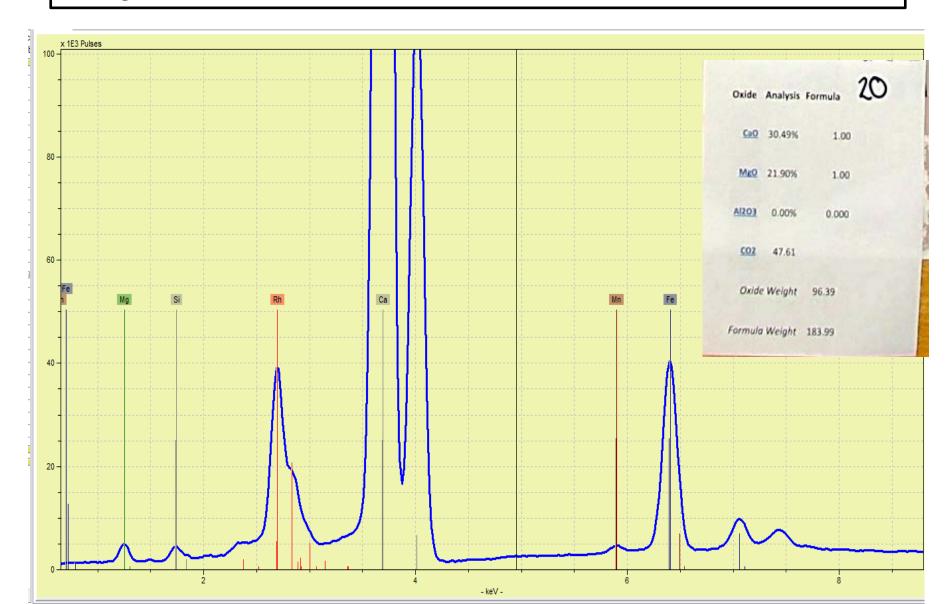
Gerstley Borate

This is and example of the major element content looking correct but the Trace elements in Gerstley Borate of P, Ti and Mn are not listed.



Ca /Mg

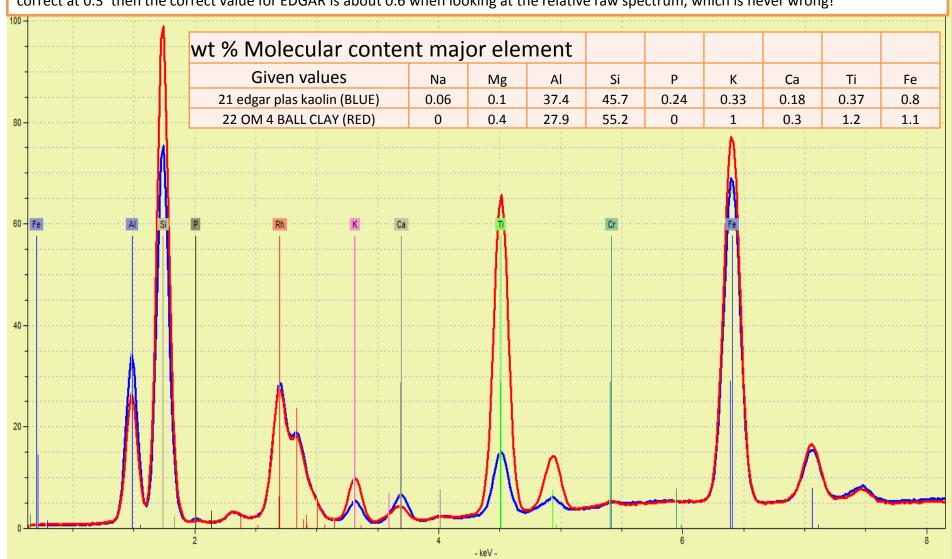
This is and example of the major elements being correct but the Tracer elements in Ca /Mg of Si, Mn and Fe are not listed. Note Fe is about 0.5% and the Si is about 3%.



The next 2 slides shows you how you can use the raw spectra between 2 samples to determine the accuracy of the given values.

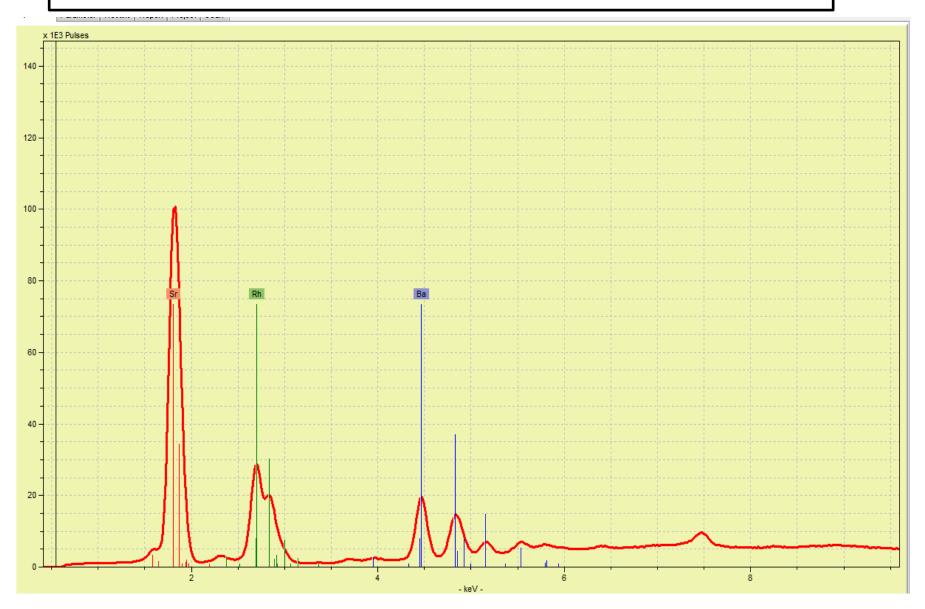
Raw spectra are never wrong. If an element has twice the number of photons from one spectra to an other then there are twice as many atoms for that element present in the sample with twice the photons.

This is and example of the some of major elements between two samples not agreeing. The Al, Si, K and Fe ratios look ok when comparing the given concentration and the raw spectra. The Ca is NOT; the OM 4 Ball (RED) is suppose to be roughly twice the EDGAR (BLUE) according to the given numbers. It looks like the given value of the Ca concentration for EDGAR (blue) is low. If the OM BALL 4 given value for Ca is correct at 0.3 then the correct value for EDGAR is about 0.6 when looking at the relative raw spectrum, which is never wrong!



Sr C

It is apparent from the Raw spectrum that the Sr C has a lot of Ba in it! This is not listed on the concentration sheet.



Quantitative analysis

The following weight percent analysis was done by calibrating the Tracer with the know samples given. Only those references that had spectra that appeared to be consistent with the given values were used (see table below) to do the calibration. The calibration was done by analyzing each sample 300 SECONDS, reading the raw spectra into the calibration software and typing in the given values and then creating a calibration response matrix. That was then used to "calibrate" the Tracer response to a given elemental concentration. The calibration matrix was then used to determine the content shown in the next charts. Note the calibration is only as good as the references values used to do the calibration.

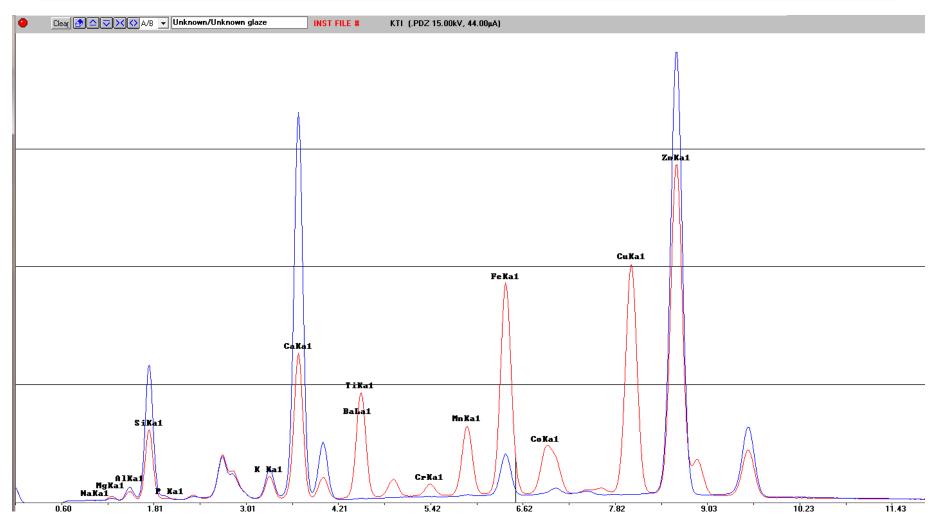
Given wt % Molecular content (major element) values and Samples used in the calibration

Only those references that had spectra consistent with the given values were used (see table below)

| | NaKa1 | MgKa1 | AlKa1 | SiKa1 | P Ka1 | K Ka1 | CaKa1 | BaLa1 | TiKa1 | CrKa1 | MnKa1 | FeKa1 | CoKa1 | CuKa1 | ZnKa1 |
|----------------------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|-------|
| 1 Mn O | 0 | 0 | 0.2 | 0.4 | 0 | 0.1 | 0.05 | 0.01 | 0.03 | 0 | 100 | 0.08 | 0 | 0 | 0 |
| 2 Mg Si | 0 | 31.67 | 0 | 63.38 | 0 | 0 | 0.08 | 0 | 0 | 0 | 0 | 0.8 | 0 | 0 | 0 |
| 3 Na C | 49 | 0 | 2 | 0.5 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 4 Fe O | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.2 | 0.05 | 0 | 99.75 | 0 | 0 | 0 |
| 5 Ti O | 0 | 0 | 0 | 0 | 0 | 0.1 | 0 | 0 | 99.9 | 0 | 0 | 0 | 0 | 0 | 0 |
| 6 Cu O | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0.05 | 2 | 97.95 | 0 |
| 8 AI O | 0 | 0 | 99.7 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 9 Si C | 0 | 0 | 0 | 99 | 0 | 0 | 0.03 | 0 | 0.03 | 0 | 0 | 0.1 | 0 | 0 | 0 |
| 10 Cr O | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 99.9 | 0 | 0.05 | 0.05 | 0 | 0 |
| 11 REDART CLAY | 0.38 | 1.59 | 15.51 | 65 | 0.22 | 4.15 | 0.26 | 0 | 1.09 | 0 | 0 | 7 | 0 | 0 | 0 |
| 12 Ca Si | 0 | 0 | 0.5 | 50 | 0 | 0 | 47 | 0 | 0.3 | 0 | 0.3 | 1.5 | 0 | 0 | 0 |
| 13 Si O | 0 | 0 | 0 | 99.9 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 14 Grolled Kaolin | 0.1 | 0.3 | 37 | 48 | 0 | 1.9 | 0 | 0 | 0 | 0 | 0 | 0.7 | 0 | 0 | 0 |
| 16 Frit 3134 | 10 | 0 | 2 | 46 | 0 | 0 | 20 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 |
| 17 custer fledspar | 3 | 0 | 17 | 69 | 0 | 10 | 0.3 | 0 | 0 | 0 | 0 | 0.15 | 0 | 0 | 0 |
| 19 Gars | 4.5 | 3.5 | 1 | 10 | 0.02 | 0.25 | 22 | 0 | 0.02 | 0 | 0.02 | 0.2 | 0 | 0 | 0 |
| 20 Ca Mg | 0 | 21.9 | 0 | 3 | 0 | 0 | 30.5 | 0 | 0 | 0 | 0.05 | 0.5 | 0 | 0 | 0 |
| 21 edgar plas kaolin | 0.06 | 0.1 | 37.4 | 45.7 | 0.24 | 0.33 | 0.6 | 0 | 0.37 | 0 | 0 | 0.8 | 0 | 0 | 0 |
| 22 OM 4 BALL CLAY | 0 | 0.4 | 27.9 | 55.2 | 0 | 1 | 0.3 | 0 | 1.2 | 0 | 0 | 1.1 | 0 | 0 | 0 |
| 23 Co O | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 99.9 | 0 | 0 |
| 24 Zinc O | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 99.9 |
| 26 rutile | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 0 | 90 | 0.5 | 0.5 | 10 | 0 | 0 | 0 |

Calculated wt % Molecular content (major element) indicated Using Tracer IV SD calibration done with references provided

| Samples | Na | Mg | Al | Si | Р | K | Ca | Ti | Cr | Mn | Fe | Co | Cu | Zn |
|----------------------|-------|--------|--------|--------|-------|-------|--------|-------|-------|-------|-------|-------|-------|--------|
| Unknown glaze (BLUE) | 2.954 | 9.558 | 11.164 | 47.285 | 0.020 | 0.736 | 12.951 | 0.000 | 0.000 | 0.000 | 0.792 | 0.078 | 0.049 | 25.779 |
| Unknown (RED) | 3.452 | 13.844 | 6.661 | 24.698 | 0.020 | 0.675 | 3.403 | 5.061 | 0.061 | 0.166 | 0.791 | 0.906 | 9.018 | 18.184 |



Calculated wt % Molecular content (major element) indicated Using Tracer IV SD calibration done with references provided

| wt % Molecular | conten | ıt (maj | or ele | ment) | | | | | | | | | | |
|----------------------|--------|---------|--------|--------|-------|-------|--------|--------|-------|-------|-------|-------|-------|--------|
| | Na | Mg | Al | Si | Р | К | Ca | Ti | Cr | Mn | Fe | Со | Cu | Zn |
| 11 REDART CLAY | 3.431 | 1.892 | 16.240 | 52.446 | 0.027 | 4.163 | 0.360 | 1.955 | 0.013 | 0.016 | 7.585 | 0.151 | 0.000 | 0.000 |
| 14 Grolled Kaolin | 5.430 | 0.819 | 35.931 | 46.520 | 0.027 | 1.722 | 0.083 | 0.000 | 0.000 | 0.000 | 0.741 | 0.083 | 0.000 | 0.000 |
| 15 mixed glaze | 3.577 | 2.069 | 10.789 | 53.757 | 0.022 | 0.257 | 13.203 | 0.210 | 0.000 | 0.000 | 0.789 | 0.214 | 0.692 | 0.000 |
| 16 Frit 3134 | 3.254 | 1.782 | 1.963 | 47.145 | 0.018 | 0.019 | 20.854 | 0.000 | 0.000 | 0.009 | 0.889 | 0.018 | 0.000 | 0.000 |
| 17 custer fledspar | 3.984 | 0.536 | 15.867 | 75.460 | 0.029 | 9.995 | 0.667 | 0.000 | 0.000 | 0.000 | 0.830 | 0.065 | 0.000 | 0.000 |
| 18 Nepheline Syenite | 4.076 | 0.000 | 22.191 | 62.089 | 0.029 | 6.078 | 0.414 | 0.000 | 0.000 | 0.000 | 0.848 | 0.079 | 0.000 | 0.000 |
| 21 edgar plas kaolin | 5.596 | 0.335 | 37.755 | 45.332 | 0.026 | 0.243 | 0.166 | 0.685 | 0.002 | 0.005 | 0.739 | 0.084 | 0.000 | 0.000 |
| 22 OM 4 BALL CLAY | 5.011 | 0.859 | 28.819 | 57.802 | 0.028 | 0.554 | 0.115 | 4.524 | 0.030 | 0.054 | 0.747 | 0.090 | 0.000 | 0.000 |
| 26 rutile | 1.380 | 0.908 | 0.324 | 0.900 | 0.010 | 0.050 | 0.052 | 89.673 | 0.321 | 0.492 | 0.808 | 0.000 | 0.000 | 0.000 |
| wt % Molecular | conten | ıt (maj | or ele | ment) | | | | | | | | | | |
| Samples | Na | Mg | Al | Si | Р | К | Ca | Ti | Cr | Mn | Fe | Со | Cu | Zn |
| Unknown glaze | 2.954 | 9.558 | 11.164 | 47.285 | 0.020 | 0.736 | 12.951 | 0.000 | 0.000 | 0.000 | 0.792 | 0.078 | 0.049 | 25.779 |
| Unknown | 3.452 | 13.844 | 6.661 | 24.698 | 0.020 | 0.675 | 3.403 | 5.061 | 0.061 | 0.166 | 0.791 | 0.906 | 9.018 | 18.184 |

The Accidental Glaze is perhaps the unknown glaze and not accidental but because the material used was not what it was said to be. To remove all accidents one must actually do a detailed elemental analysis of all the materials used. Glazes are like special artistic glasses, the devil is very much in the details, Including the traces elements!

It is clear from the Tracer elemental analysis of this material; what was said of these materials does not always match what is actually there!

Much more could be determined concerning THE materials using the Tracer..... But the above should give one a glimpse of it capabilities

THE END