

## Comparison of leaching methods for the estimation of bioavailability in two CCA-contaminated soils

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## Background - Risk Assessment of polluted sites

□ Total concentrations of contaminants compared to permissible levels (e.g. PNECS, guideline limit values from national EPA's) → estimation of risk

□ Lack of site-specific information, soil conditions, combination effects, and bioavailability may lead to over- and underestimation of risk

□ Other tools to better estimate risk is necessary



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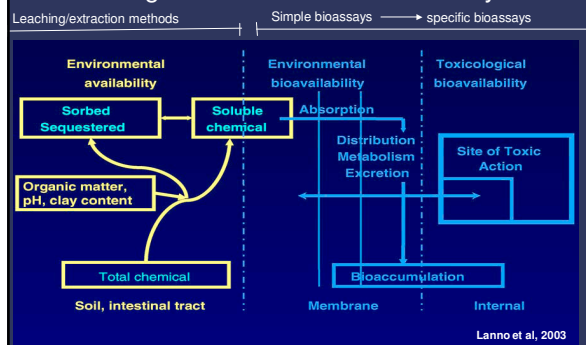
## Background - Risk Assessment of polluted sites

- Combination of toxicological methods and chemical analysis is beneficial
- Standardised bioassays ( USEPA, ASTM, OECD,etc.) have been proposed  
Plants, soil living organisms e.g. earthworms, springtails, plants etc.
- Ecologically relevant, but often laborious → high costs!!
- Spatial distribution is necessary at large contaminated sites
- Fast och cheap surrogate methods for screening purposes are needed



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## Background - What is bioavailability?



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### Aims of the study

- Compare the relative bioavailability of arsenic and arsenic species between methods
  - Leaching/extraction efficiency – yield of metals/metalloid in the leachates
  - Cytotoxicity and genotoxicity in the leachates



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### Leaching/extraction methods for estimation of bioavailability

- **ASE** - Accelerated Solvent Extraction ( or Pressurized liquid extraction (PLE))  
10MPa and **150 °C**, **100 °C**, **50 °C**
- **Batch** leaching EN- 124 57-2, L/S=10, 24h.
- Batch leaching 0.01 M **CaCl<sub>2</sub>**, L/S 10, 2h (Houba et al. 1996).
- **IVG** anaerobic, 37 °C, 2 x 1h (Rodriguez et al., 1999)



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### ASE - Accelerated Solvent Extraction

Environmental sample in



H<sub>2</sub>O

Biocompatible Extract out

- High through put, ca 20 min/sample
- High temp. & pressure
- Solvent is water
- Medium effective extraction
- Worst case scenario of mobile and water soluble fraction of contaminants



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### Bioassays

✓ **L-929**-cytotoxicity: General toxicity, disturbed basal cellular functions affecting growth.

✓ **Genotoxicity using CHO-cells:** mutagenic properties e.g. metal/metalloid (Helleday et al.2001)

Control	Day1	Day2	Day3	Day4	Day5	Day6	Day7	Day8	Day9	Blank
AA8	Day1	Day2	Day3	Day4	Day5	Day6	Day7	Day8	Day9	Blank
UV4	Day1	Day2	Day3	Day4	Day5	Day6	Day7	Day8	Day9	Blank
UV5	Day1	Day2	Day3	Day4	Day5	Day6	Day7	Day8	Day9	Blank
EM9	Day1	Day2	Day3	Day4	Day5	Day6	Day7	Day8	Day9	Blank

- **AA8: Wild-type**
- **UV4: Deficient in DNA-repair**
- **UV5: Deficient in DNA-repair**
- **EM9: Deficient in DNA-repair**



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### Test soils

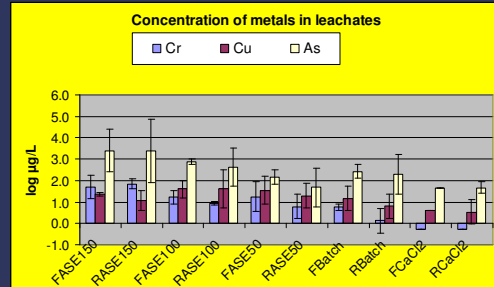
Wood impregnation sites using CCA-based preservatives

	Forsmo mg/kg	Robertsfors mg/kg
Arsenic	276	311
Cadmium	3	4
Chromium	45	54
Cobalt	6	4
Copper	18	10
Lead	20	13
Nickel	11	6
Zinc	135	105
TOC	2.6%	8.7%



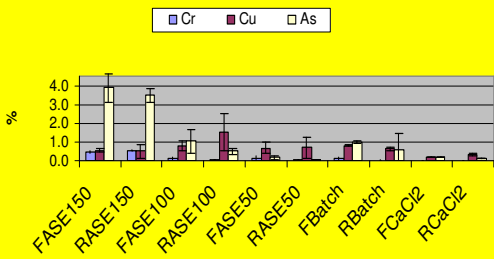
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### Results



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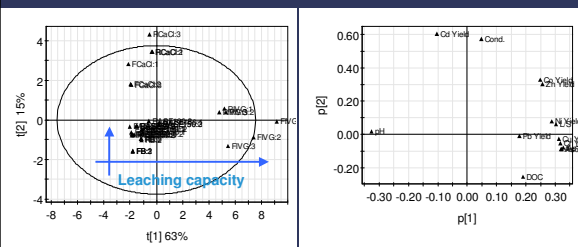
### Metal yield



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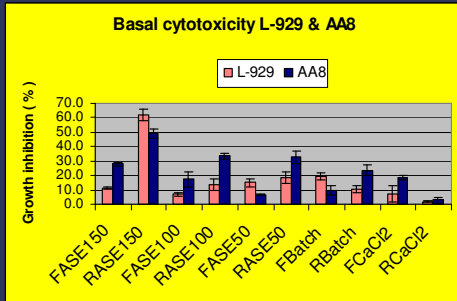
### Principal component analysis (PCA)

Chemical characteristics



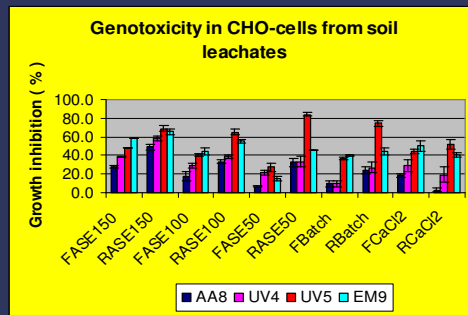
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### Leachate toxicity



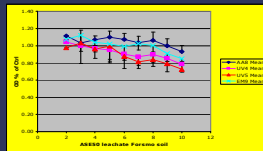
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### Leachate Genotoxicity

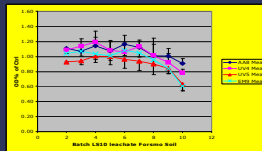


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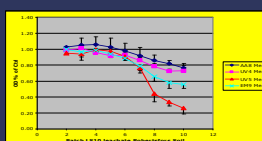
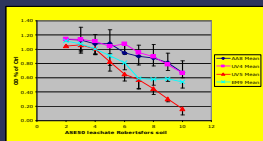
### ASE50



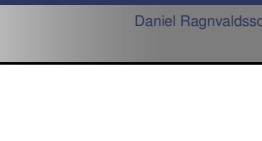
### Batch



### Forsmo leachates

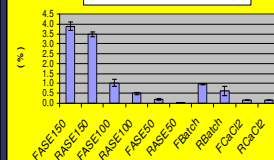


### Robertsfors leachates

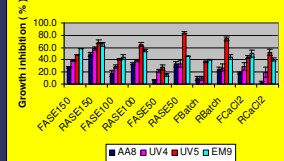


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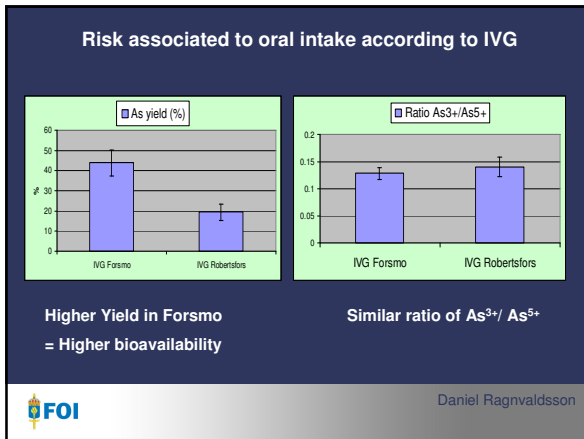
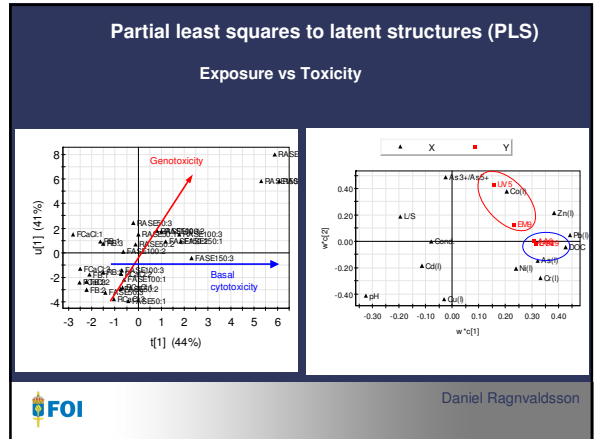
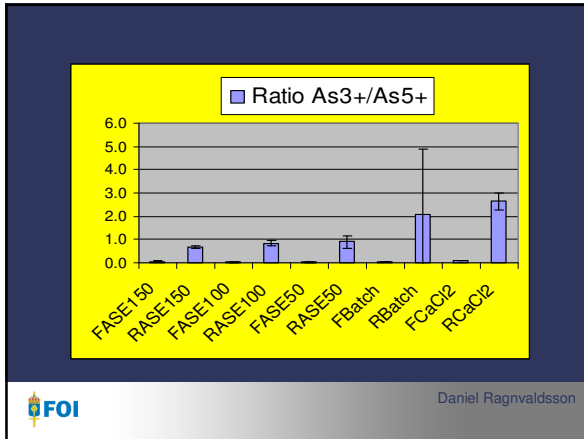
### Extraction of total arsenic (%)



### Genotoxicity in CHO-cells from soil leachates



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### Conclusion 1

- Higher yield and toxicity in PLE/ASE at 150 C – suitable for screening purposes to assess the toxic potential in soil samples
- Comparable results from ASE at lower temperatures (100 °C & 50 °C) and the batch leaching methods, EN12457-2 and 0.01M CaCl<sub>2</sub>

## Conclusion 2

- ❑ Speciation of Arsenic is essential for risk assessment of CCA - contaminated sites
- ❑ Both analysis of chemical speciation and bioassays for genotoxicity reveal these differences compared to total concentrations!



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## Conclusion 3

- Drink Forsmo soil leachates - Eat Robertsfors soil!

Thank you for your attention!!



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