

# **Radiochemistry at the Environmental Monitoring Section at Institute for Energy Technology (IFE), Kjeller, Norway**

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# Radiochemical activities

- Environmental monitoring programme
  - Low-level liquid radioactive effluents
  - Environmental samples from the surroundings
- Personnel monitoring (bioassay)
- Research projects
- Commercial services



Kjellerholen  
VA1  
SD1

IFE

VA2

Utslippsledning

SD3

Nybrua

Lillestrøm

VA4  
SD4

Rælingsbrua

VA5  
SD5

FI6

BATHAVNA

Svelle  
VA7  
SD7  
BP7

Glomma

# Samples

- NALFA
  - Sediment ( $\gamma$ -emitters and Pu)
  - Water ( $\gamma$ -emitters,  $^3\text{H}$  and  $^{90}\text{Sr}$ )
- Rain water ( $\gamma$ -emitters,  $^3\text{H}$ ,  $^{90}\text{Sr}$  and Pu)
- Farms
  - Milk ( $\gamma$ -emitters and  $^{90}\text{Sr}$ )
  - Agricultural products ( $\gamma$ -emitters,  $^{90}\text{Sr}$  and Pu)
- Gras ( $\gamma$ -emitters,  $^{90}\text{Sr}$  and Pu)
- River
  - Water ( $\gamma$ -emitters,  $^3\text{H}$ ,  $^{90}\text{Sr}$  and Pu)
  - Sediment ( $\gamma$ -emitters,  $^{90}\text{Sr}$ , U and Pu)
  - Fish ( $\gamma$ -emitters,  $^{90}\text{Sr}$  and Pu)
  - Water plants ( $\gamma$ -emitters,  $^{90}\text{Sr}$  and Pu)

# U, Th, Pu, Am, Cm, Sr and Ra

- Pretreatment of samples (urine, discharge water, precipitation, river water, produced water, sediment, soil, biological material)
- Separation and determination of U, Th, Pu, Am, Cm, Sr and Ra

TRU (Pu(III),  
Am(III))

Ra



UTEVA (U(VI),  
Th(IV))

Sr(II)

# Sr, U, Pu, Am and Cm in liquid effluents

- 200 ml sample added tracers is evaporated and dry ashed
- Dissolved in 3 M HNO<sub>3</sub> added sulfamic acid, ascorbic acid and Al(NO<sub>3</sub>)<sub>3</sub> (Fe(II), Pu(III))
- Loaded on UTEVA-TRU-Sr columns coupled together
- On-column oxidation of Pu(III) to Pu(IV) with NaNO<sub>2</sub>
- UTEVA: U(VI) eluted with 0.01 M HCl
- TRU: Am(III) and Cm(III) eluted with 4 M HCl  
Pu(III) eluted with 4 M HCl + TiCl<sub>3</sub>
- Sr: Sr eluted with 0.05 M HNO<sub>3</sub> (ingrowth of <sup>90</sup>Y)

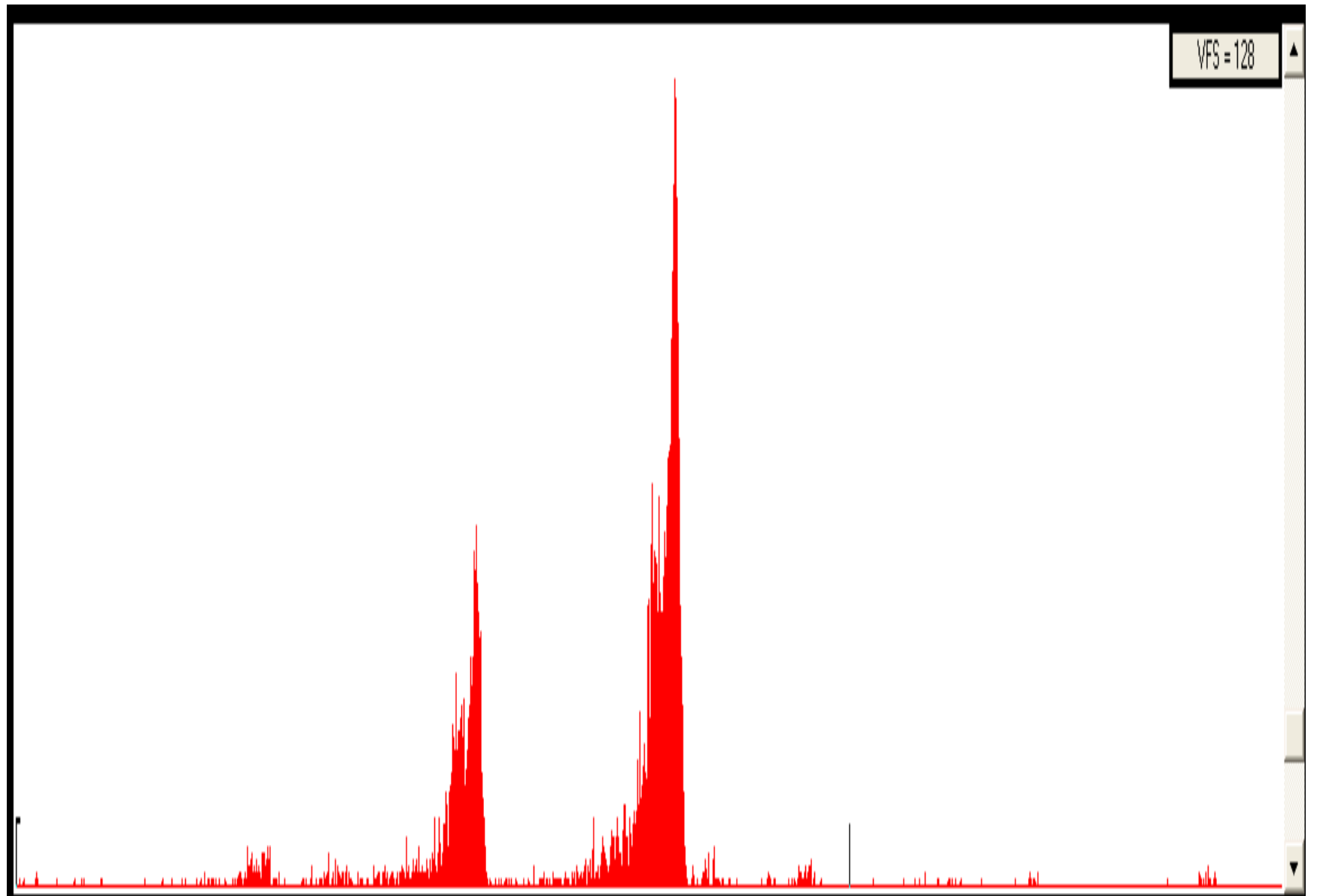
# Sr and Pu in sediment

- Drying and ashing
- Leaching with aqua regia
- Evaporation and dissolution in dilute HCl
- Oxalate precipitation followed by ashing
- Dissolution in 3 M HNO<sub>3</sub> added sulfamic acid, ascorbic acid and Al(NO<sub>3</sub>)<sub>3</sub> (Fe(II), Pu(III))
- Loaded on TRU-Sr columns coupled together
- TRU: Pu(III/IV) (source prep.: co-precipitation)
- Sr: Sr



# Sr, U, Pu and Am in urine

- Phosphate precipitation followed by ashing
- Dissolution in 3 M HNO<sub>3</sub> added sulfamic acid, ascorbic acid and Al(NO<sub>3</sub>)<sub>3</sub> (Fe(II), Pu(III))
- Load on UTEVA-TRU-Sr columns coupled together
- UTEVA: U(VI)
- TRU: Pu(III/IV), Am(III), Cm(III)
- Sr: Sr



# $^{210}\text{Pb}$ , $^{226}\text{Ra}$ and $^{228}\text{Ra}$ in produced water

- 2 litre produced water is heated with  $\text{HNO}_3$  and  $\text{MnO}_4^-$  after addition of  $^{133}\text{Ba}$  tracer
- pH adjustment to 1-2
- Reduction of  $\text{MnO}_4^-$  to  $\text{MnO}_2$
- Addition of sulphate ( $\text{H}_2\text{SO}_4$  and  $\text{Na}_2\text{SO}_4$ )
- Addition of Pb and Ba ( $\text{Pb}(\text{Ba})\text{SO}_4$ )
- Collection of  $\text{MnO}_2$  and  $\text{Pb}(\text{Ba})\text{SO}_4$
- Drying of the precipitate
- Gamma determination of  $^{210}\text{Pb}$ ,  $^{226}\text{Ra}$  and  $^{228}\text{Ra}$

# $^{210}\text{Pb}$ and $^{210}\text{Po}$ in produced water

- Addition of  $^{209}\text{Po}$  and stable lead
- Treatment of sample with acids and oxidizing agents
- Co-precipitation of Pb and Po with  $\text{MnO}_2$
- Dissolution with  $\text{HCl} + \text{H}_2\text{O}_2$
- Dissolve with 2 M  $\text{HCl}$  and load on Sr-Resin
- Wash with 2 M  $\text{HCl}$  and 6 M  $\text{HNO}_3$
- Po: 6 M  $\text{HNO}_3$
- Pb: 6 M  $\text{HCl}$

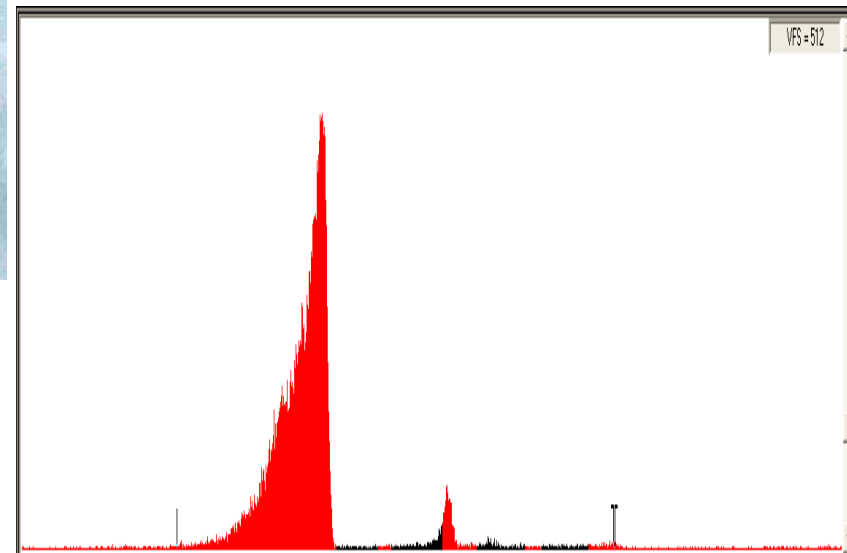
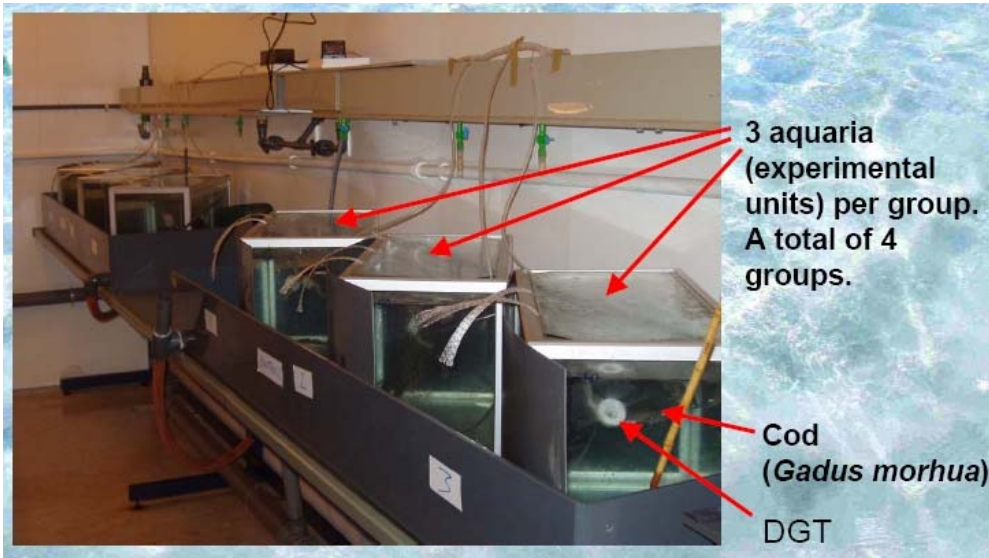
# U, Th and Ra in Si materials

- Dissolution of Si-materials with  $\text{HNO}_3$  – HF
- Treatment with  $\text{HNO}_3$
- Dissolution in 3 M  $\text{HNO}_3$
- Load on UTEVA
- UTEVA: Th(IV) eluted with 4 M HCl  
U(VI) eluted with 0.01 M HCl
- Ra sep. by sulfate precipitations
- Source prep.: co-precipitation

# $^{226}\text{Ra}$ and $^{228}\text{Ra}$ in seawater

- $\text{MnO}_2$  co-precipitation of Ra (pH 6-7!)
- Washing of the precipitate (wash out traces of Ca)
- Dissolution with  $\text{HCl} - \text{H}_2\text{O}_2$
- $\text{PbSO}_4$  co-precipitation of Ra
- Dissolution with DTPA
- $\text{BaSO}_4$  co-precipitation of Ra  $\rightarrow$   $\alpha$ -spec.
- Ingrowth of  $^{228}\text{Th}$  from  $^{228}\text{Ra}$   $\rightarrow$  Th separation and  $\alpha$ -spec

# $^{226}\text{Ra}$ with DGTs



# $^{90}\text{Sr}$ in seawater

- $^{90}\text{Sr}$  co-precipitation with  $\text{Ca}(\text{Sr})\text{CO}_3$
- Sr separation from Ca by repeated nitrate precipitations
- Sr separation from Y by repeated hydroxide precipitations
- Sr separation from Ba, Ra and Pb by repeated chromate precipitations
- Ingrowth of  $^{90}\text{Y}$
- $^{90}\text{Y}$  separation from  $^{90}\text{Sr}$  by hydroxide and sulphate precipitations



# $^{90}\text{Sr}$ in seawater – new

- Acidification and equilibrium between  $^{90}\text{Sr}$  and  $^{90}\text{Y}$
- Co-precipitation of Y carrier and  $^{90}\text{Y}$  with  $\text{Fe}(\text{OH})_3$
- Dissolution and evaporation
- Y extraction with TBP from 14 m  $\text{HNO}_3$
- Y strip with  $\text{H}_2\text{O}$
- Is further separation from Th required? Phosphate precipitation of Th?
- Source preparation by hydroxide and oxalate precipitation
- Risø beta counter and EDTA titration

**Thanks!**