

## Estimating soil properties with a proximal gamma-ray spectrometer using windows and full-spectrum analysis methods

2<sup>nd</sup> Global Workshop on Proximal Soil Sensing

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## General structure of this talk

General structure

- Introduction
- Materials and Methods
- Results
- Conclusions
- Future research

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## Background

Introduction

- Why soil sensors?
  - Conventional soil analysis is **expensive** and **laborious**
- Why proximal soil sensors?
  - Remote sensors – low **spatial** resolution
  - Interfering** factors
- What does a gamma sensor measure?
  - Gamma rays – **decay of radioisotopes** ( $^{40}\text{K}$ ,  $^{238}\text{U}$  and  $^{232}\text{Th}$ )
- What does abundance of radionuclides reflect?
  - regolith** features, for instance
  - Low count rate – **young** or **sandy** soil with leached profile
  - High count rate – **weathered** soils with **clay minerals**, **gravels** or **silica**

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## Spectral analysis methods

Introduction

- Full-spectrum analysis (FSA) method
- Conventional energy-windows (EWs) method

Spectrum is adapted from Wilford et al. (1997)

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## Objectives

Introduction

- To evaluate a proximal gamma-ray spectrometer **"the Mole"** to ascertain **quantitative relationships** between radiometric data and soil properties
- To compare two data analysis methods, **the EWs** and **the FSA**, to estimate soil properties using the data of the Mole

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## Gamma-ray spectrometer "the Mole"

Materials and methods

```

    graph TD
        Radioisotopes --> CsI_Tl_Crystal
        CsI_Tl_Crystal --> Photomultiplier
        Photomultiplier --> Amplifier
        Amplifier --> MultichannelAnalyser[Multichannel Analyser]
        MultichannelAnalyser --> DataLogging
        DataLogging --> OutputCounts[Output counts]
    
```

Picture is adapted from Van Egmond et al. (2010)

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## The Mole collecting the data

Materials and methods

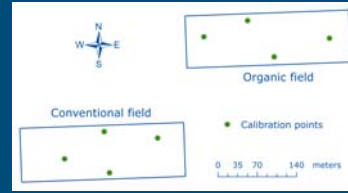


Received from Soil Conapny (2011)

## Data collection

Materials and methods

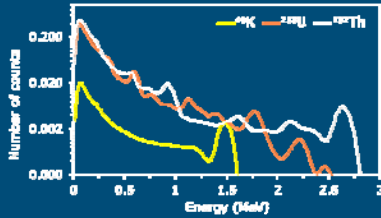
- Site – Lelystad, the Netherlands (total area = 5.5 ha)
- Calibration spectra were collected for 5 minutes
- Soil samples were taken from top 0-25 cm depth
- They were analysed for clay, OM, TN, TP, TK, Mg and pH



## Full-spectrum (FSA) method

Materials and methods

- Fits standard spectra to a measured spectrum
- A standard spectrum is the pure response of a detector having a fixed concentration (1 Bq/kg) of a radionuclide

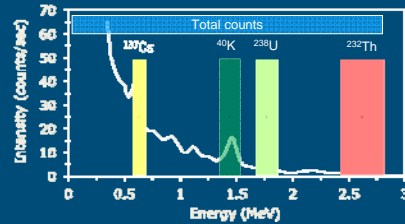


Hendricks et al. (2001)  
Reprinted from Van Edmond et al. (2010)

## Energy-windows (EWs) method

Materials and methods

- Sums counts in EWs of  $^{137}\text{Cs}$ ,  $^{40}\text{K}$ ,  $^{238}\text{U}$ ,  $^{232}\text{Th}$  and TC
- Calibration factors were determined from the standard spectra of the Mole



## Calibration – the FSA method

Results

Field	Soil property	R <sup>2</sup> <sub>adj</sub>	Radionuclide
Conventional	Clay content (%)	0.98**	$^{232}\text{Th}$
	OM (%)	0.96*	$^{40}\text{K}$
Organic	TN (mg/kg)	0.99**	$^{238}\text{U}$
	TP (mg/kg)	0.91*	$^{238}\text{U}$
Combined	Clay content (%)	0.54*	$^{232}\text{Th}$
	TN (mg/kg)	0.50*	$^{238}\text{U}$
	Mg (mg/kg)	0.50*	$^{238}\text{U}$

\*\* Relations are significant at 0.01 level; \* relations are significant at 0.05 level

- Correlations were **not consistent** with radionuclides
- Correlations were **variable** across fields
- R<sup>2</sup><sub>adj</sub> values were **higher** in individual fields

## Calibration – EWs method

Results

Field	Soil property	R <sup>2</sup> <sub>adj</sub>	Radionuclide
Conventional	OM (%)	0.88*	$^{238}\text{U}$
	TP (mg/kg)	0.94*	$^{137}\text{Cs}$
Organic	TN (mg/kg)	0.82	$^{238}\text{U}$
	TK (mg/kg)	0.82	$^{238}\text{U}$
Combined	Clay content (%)	0.46*	$^{232}\text{Th}$
	OM (%)	0.47*	$^{137}\text{Cs}$
	TN (mg/kg)	0.54*	$^{238}\text{U}$


\*\* Relations are significant at 0.01 level; \* relations are significant at 0.05 level

- Correlations were **not consistent** with radionuclides
- Correlations were **variable** across fields
- R<sup>2</sup><sub>adj</sub> values were **higher** in individual fields


## Calibration – comparison of FSA and EWs

**Results**

- Results in both methods were **comparable**; the **FSA** method, however, performed **better**
- Uncertainty in EWs – **background** and **limited information**
- Results were **variable** across fields and methods
- Relations were **not consistent**, but **site-specific**
- Both methods related **different properties** in the same field
- Correlations were **higher** in individual fields
- Clay content** and **TN** showed some **consistency** in relationships


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## Validation of models across fields

**Results**


- The FSA models predicted clay, TN and pH across fields

Field	Soil property	R <sup>2</sup>	RMSEP
Conventional	Clay content (%)	0.98	0.92
	TN (mg/kg)	0.69	54.37
	pH	0.75	0.05
Organic	Clay content (%)	0.83	0.66
	TN (mg/kg)	0.55	116.77
	pH	0.81	0.25


- The EWs predicted clay and TK with low correlation

Field	Soil property	R <sup>2</sup>	RMSEP
Conventional	Clay content (%)	0.51	0.57
	TK (mg/kg)	0.51	4.59
Organic	OM (%)	0.26	0.41
	TP (mg/kg)	0.46	12.69

- The FSA method outperformed the EWs method


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## Conclusions

**Conclusions**

- The Mole is **capable** to relate soil properties
  - Correlations are **variable** across fields and methods suggesting that they are **site-specific**
  - Clay content** shows some consistent correlation with <sup>232</sup>Th
  - TN** shows some consistent correlation with <sup>238</sup>U
  - Other soil properties show **mixed** correlations
- FSA versus EWs
  - Both methods show potential to relate soil properties
  - FSA method **outperforms** the EWs method during calibration
  - FSA **outperforms** EWs also during validation (**clay, TN and pH**)
  - Both methods related **different properties** in the same field


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## Future research needed


**Future research**

- Site-specific behaviour of correlations needs to be understood taking **more calibration** samples within a field
- Next step is to make **maps** and **compare** them with **existing maps** or **dense ground-truth** samples to bring things closer to practice of precision agriculture
- Calibration spectra should also be measured **on-the-go** to better **harmonise** them with the **field spectra**




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## Thank you for your attention




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