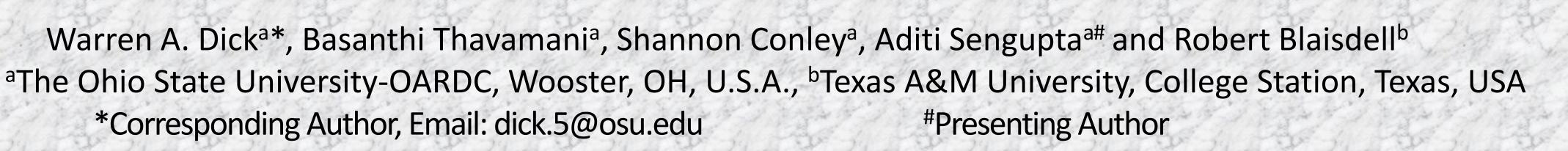
Prediction of Selected Soil Carbon Enzymes and Properties in a Diverse Population of Ohio Soils Using Near Infrared Reflectance Spectroscopy Analysis



Abstract

A Near Infrared Spectroscopy (NIRS) method was used to predict soil organic and amino sugar concentrations, and β -glucosidase and β carbon glucosaminidase activity in 184 diverse soils of Ohio. The measured variable values were calibrated to NIR spectral data with partial least squares regression analysis. The multivariate models developed were validated using the full cross validation method and the test set method with a test set size of 50 samples. Statistical analysis of the spectral data was done using the multivariate analysis software Unscrambler 8.0 (CAMO Inc). The first differential transformation of the NIR region (1100-2498 nm) gave good results. The NIRS method predicted well the organic carbon (OC) and amino sugar concentrations in soil (R² was 0.91 for OC and 0.90 for amino sugar). The enzyme activity values were also well predicted (R^2 was 0.82 for both soil β -glucosidase and β -glucosaminidase enzyme activity).

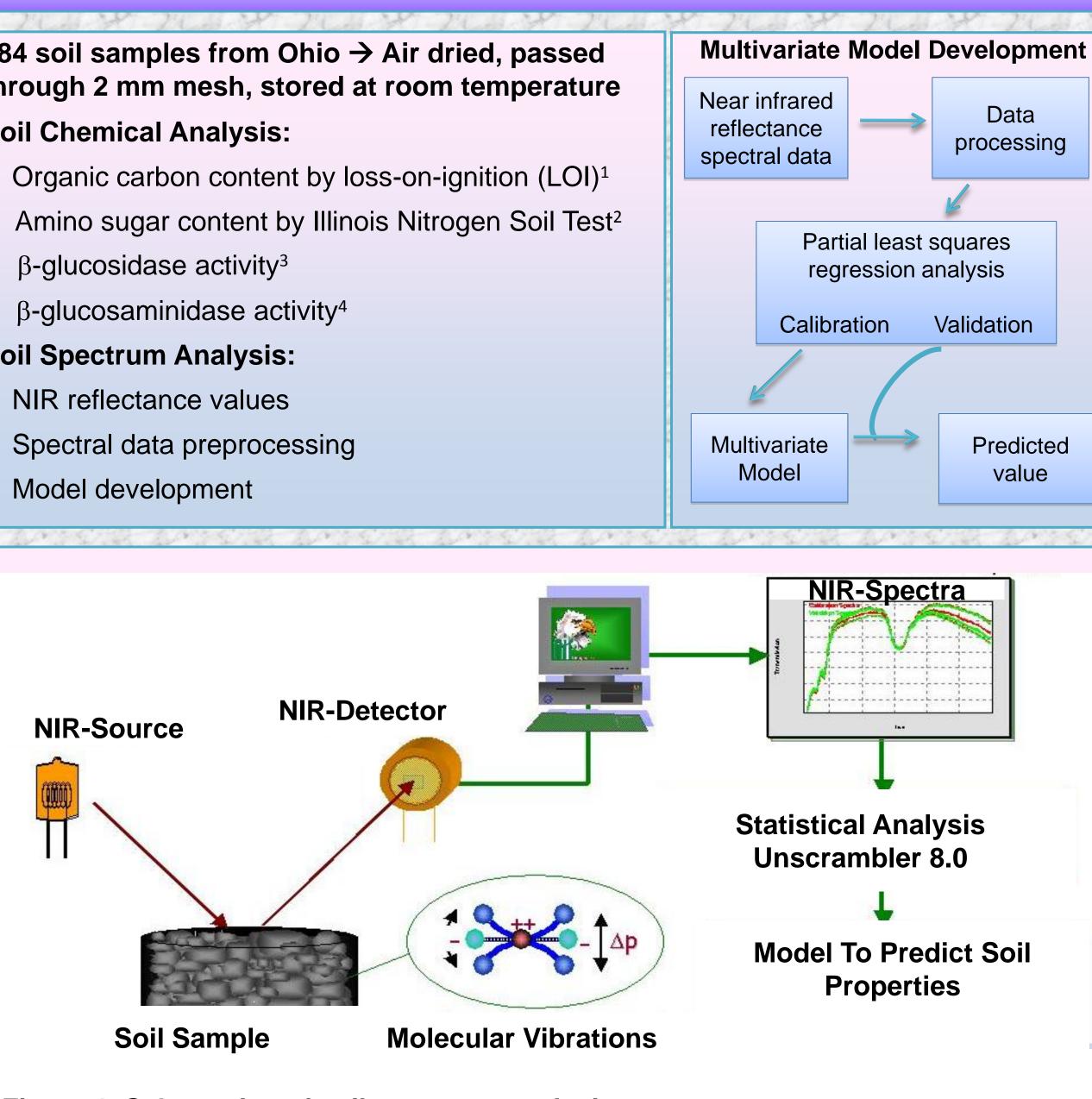
Methods

184 soil samples from Ohio \rightarrow Air dried, passed through 2 mm mesh, stored at room temperature

Soil Chemical Analysis:

Organic carbon content by loss-on-ignition (LOI)¹ Amino sugar content by Illinois Nitrogen Soil Test² β -glucosidase activity³ β -glucosaminidase activity⁴

Soil Spectrum Analysis:





Advantages of NIRS Analysis

Rapid assessment of soil properties	Simultaneous measurement of variables
Nondestructive to samples	Precision of analysis
No use of hazardous chemicals	Simple sample preparation

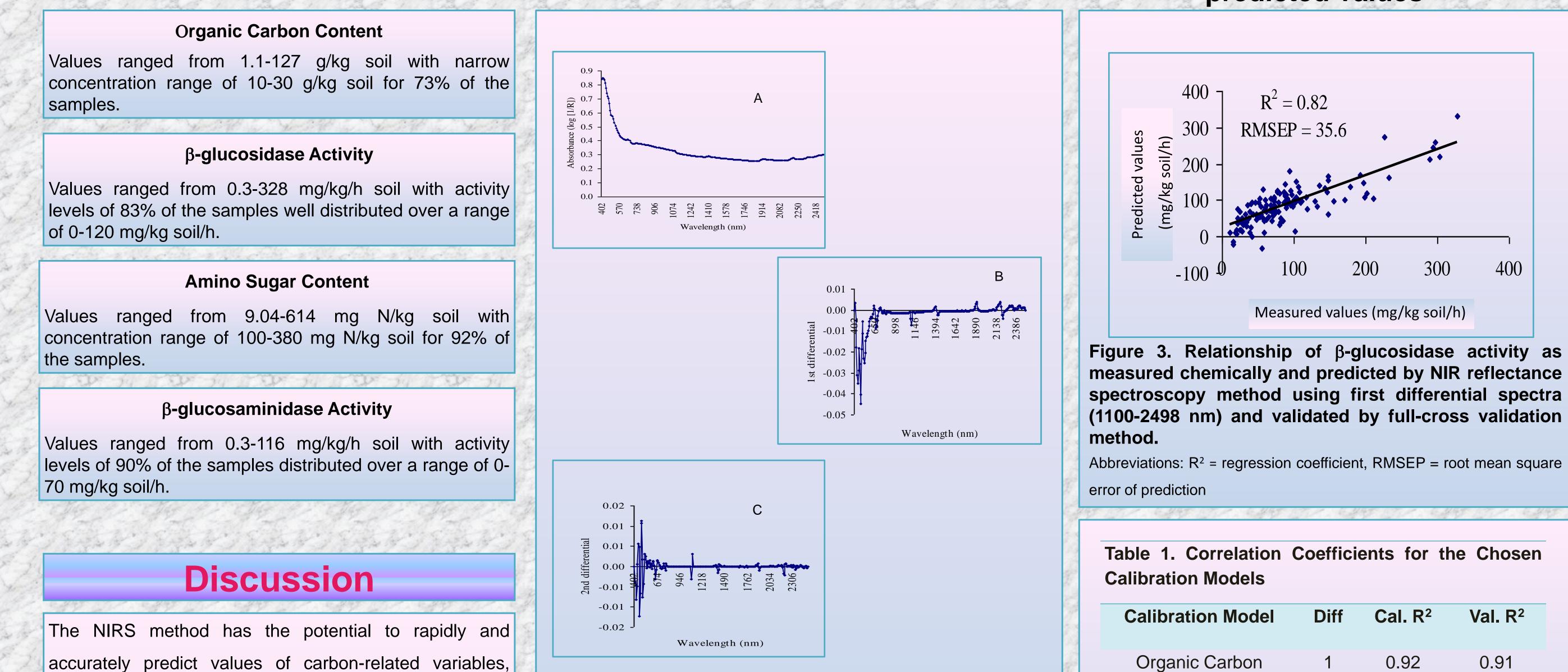
Figure 1. Schematics of soil spectrum analysis

Results

Soil Chemical Analysis

Spectral Data Processing

Correlation of measured vs. predicted values



including the carbon enzymes β -glucosidase and β glucosaminidase, in soils. The equipment needed is not expensive and the NIRS method can be used where very large numbers of environmental samples need to be rapidly analyzed. Indeed, the prediction equations can be constantly improved as more data points are entered into the correlations between laboratory-measured values and NIRS values.

Figure 2. Spectrum of a soil sample. (A) Raw spectral

data. (B) First differential of spectral data. (C) Second

differential of spectral data.

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References Storer, D.A. 1984. A simple high sample volume ashing procedure for determining soil organic matter Commun. Soil. Sci. Plant Anal. 15:759-772. 2. Khan, S.A., R.L. Mulvaney, and R.G. Hoeft. 2001. A simple test for detecting sites that are nonresponsive to nitrogen fertilization. Soil Sci. Soc. Am. J. 65:1751-1760. Eivazi, F. and M.A. Tabatabai. 1988. Glucosidases and galactosidases in soils. Soil Biol. Biochem. 20:601-606. Parham, J.A. and S.P. Deng. 2000. Detection, quantification and characterization of b-glucosaminidase activity in soil. Soil Biol Biochem. 32:1183-1190. CAMO Inc. 2003. The Unscrambler user manual. CAMO Inc., Oslo, Norway.

	β –glucosidase	1	0.90	0.82	
	Amino sugar	1	0.92	0.90	
	β-glucosaminidase	1	0.82	0.89	
Abbreviation: Diff = Differential transformation used, Cal= Calibration and Val= Validation					
Acknowledgments					
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