Estimation of soil properties with mid-infrared soil spectroscopy across yam production landscapes in West Africa

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Abstract. Low soil fertility is challenging the sustainable production of yam and other staple crops in the yam belt of West Africa. Quantitative soil measures are needed to assess soil fertility decline and to improve crop nutrient supply in the region. We developed and tested a mid-infrared (mid-IR) soil spectral library to enable timely and cost-efficient assessments of soil properties. Our collection included 80 soil samples from four landscapes (10 km×10 km) and 20 fields/landscape across a gradient from humid forest to savannah, and 14 additional samples from one landscape that had been sampled within the Land Health Degradation Framework. We derived partial least square regression models to spectrally estimate soil properties. The models produced accurate cross-validated estimates of total carbon, total nitrogen, total sulfur, total iron, total aluminum, total potassium, total calcium, exchangeable calcium, effective cation exchange capacity, diethylenetriaminepentaacetic acid (DTPA) extractable iron and clay content ($R^2 > 0.75$). The estimates of total zinc, pH, exchangeable magnesium, bioavailable copper and manganese were less predictable ($R^2 > 0.50$). Our results confirm that mid-IR spectroscopy is a reliable and quick method to assess the regional-level variation of most soil properties, especially the ones closely associated with soil organic matter. Although the relatively small mid-IR library shows satisfactory performance, we expect that frequent but small model updates will be needed to adapt the library to the variation of soil quality within individual fields in the regions and their temporal fluctuations.

1 Introduction

Yam (*Dioscorea* spp.) is an important food and cash crop in West Africa. The yam belt of West Africa spans across the central zone of coastal countries in West Africa, located across the humid forest zone and northern Guinean savanna. It contributes to about 92% of total world yam production, e.g. a total yield of 73 million metric tons in 2017 (Food and Agriculture Organization of the United Nations, 2019). The cropping area in the West African yam belt has been expanded with accelerated population

growth. The deforestation and expansion of agricultural land has in many places caused soil degradation. Furthermore, there has been a trend of shortened fallow periods in the cropping areas of West Africa over the last decades, which has further exacerbated the decline in soil fertility across the yam belt. Traditionally, yam is grown without external input in these areas. Therefore, the production of yam and other crops grown in the region depends on soil organic matter (SOM) status (Padwick, 1983), which serves as a main pool of plant-available nutrients and provides cation exchange surfaces for soil nutrients (Syers et al., 1970; Soares and Alleoni, 2008). A particularly strong positive relationship between high organic matter stocks and yam productivity is reported after fallow and when no fertilizer is added (Diby et al., 2009; Kassi et al., 2017). Thus, maintaining or increasing SOM and available nutrient levels is of utmost importance for sustainable production of yam and other crops in West Africa (Carsky et al., 2010). Furthermore, linking soil properties and yam yields (Frossard et al., 2017) and accounting for soil macro- and micronutrient status (O'Sullivan and Jenner, 2006) is fundamental to improving crop yields and soil management strategies.

Soil fertility is an integrative measure of soil attributes and their interactions that support the long-term agricultural production potential. Soil fertility is commonly decomposed into the physical, chemical and biological major components (Abbott and Murphy, 2007). Here, it is important to interpret soil fertility in the form of soil conditions and functions at an adequate resolution over time and space, and in relation to the crop of interest. For yam, low tuber yields are often attributed to an unbalanced ratio of essential nutrients (i.e. N, P, K) available in the soil (Enyi, 1972) and a fast mineralization and hence depletion of organic matter (Carsky et al., 2010; Hgaza et al., 2011). Yet, the relationship between soil properties and tuber yield is not fully understood (Frossard et al., 2017). The reason is that the response of yam to mineral fertilization is highly variable because of confounding environmental and management variables, such as climate, soil type, inherent soil fertility, micronutrient deficiencies, tillage, seed tuber quality, planting date and density, staking and disease pressure across the yam belt (Kang and Wilson, 1981; O'Sullivan and Jenner, 2006; Cornet et al., 2016; Enesi et al., 2018). Further, there are no soil fertility recommendations specific for yam under West African conditions. For this reason, establishing yam field trials designed with different organic and mineral fertilization strategies within different yam growing regions is required to optimize yam nutrient supply targeting regional soil and environmental conditions (Frossard et al., 2017). Despite the importance of soil fertility, it is challenging to quantify soil measures at sufficient temporal and spatial resolution to relate them to yam productivity together with other management effects.

To quickly assess key soil properties, such as soil organic carbon (SOC) and cation exchange capacity (CEC), we need more cost- and time-efficient methods in addition to the traditional wet chemistry laboratory analyses that are often cost-intensive and time consuming. Proximal sensing is a method that can provide reliable, rapid, and inexpensive soil measurements (UNEP, 2012). Soil visible and near infrared (vis-NIR), and mid-infrared (mid-IR) diffuse reflectance spectroscopy has gained popularity over the past 30 years to assess soil properties in a complementary manner to conventional laboratory analytical methods (Nocita et al., 2015). For model development and calibration but as importantly also for validation purposes, soil IR spectroscopy requires laboratory reference analysis data. Previous studies have shown successful spectroscopic predictions of soil properties, such as organic C, texture, cation exchange capacity (CEC), and exchangeable K (Viscarra Rossel et al., 2006; Cécillon et al., 2009; Nocita et al., 2015; Sila et al., 2016). Many soil chemical and physical properties, such as soil mineralogy,

the concentration, forms and distribution of SOM, are closely associated with IR spectral diversity. However, for determining a range of extraction-based soil properties, the predictive capability seems variable. This can be caused by complex surface chemical processes that are not directly related to soil organic matter and/or insufficient densities available at local scale to represent such locally complex relationships (Viscarra Rossel et al., 2006; Abdi et al., 2012; Sanderman et al., 2020). Further, a library that includes a broad range of soil biophysical conditions found in the region in which it is used needs to be established. Depending on the geographical extent of the study — field (e.g., Cambou et al., 2016), region, country (e.g., Clairotte et al., 2016), continent (e.g., Sila et al., 2016)), world (e.g., Viscarra Rossel et al., 2016) — various statistical predictive modeling strategies are typically employed to account for geographically regional variability in soil properties and determine empirical relationships between spectra and soil attributes. Particular subsets of and features in spectra are characteristic for functional groups of soil components and thus, elucidating spectral features that are important for the prediction of a particular soil attribute helps to understand and validate the mechanisms based on which the empirically models predict the soil properties.

In this work, we aim to develop mid-IR spectroscopy as a diagnostic tool for key analytical soil variables within four climatically, ecologically, and agriculturally distinct landscapes in Burkina Faso and Ivory Coast. For yam and other cash crops, there is a lack of soil diagnostic tools to identify factors limiting yields and to derive site-specific fertilizer recommendations within and across landscapes. In these regions, yam has substantial economic importance for small-holder farmers. As land management and soil status is a key factor not only for yam but also other high-value crops in the region, quick and cost-effective soil status assessments should be transferable to other crops with similar nutrient demands. Thus, the main objectives of this study were to (1) develop and evaluate openly accessible and re-usable mid-IR spectroscopic models to estimate soil properties for selected landscapes representing major soil and climatic conditions in the West African yam belt, (2) to determine important spectral features for specific soil properties, and (3) to build a new soil spectral library in four landscapes of the West African yam belt for soil prediction and assessment. Finally, we make specific recommendations on whether and how specific mid-infrared diagnostic measures are applicable for different soil management and screening purposes. We also discuss the spectroscopic evaluation of the soil's capacity to retain and release nutrients for sustained and improved cropping in the region.

2 Materials and methods

2.1 Landscapes and soil sampling

Our study area covered the climatic and soil biophysical conditions representative of the West African yam belt. We selected four landscapes, two in Ivory Coast and two in Burkina Faso. Each landscape (approximately 10 km x 10 km) represents a diverse geographic ecoregion. The landscapes cover a gradient between humid forest and the northern Guinean savannah. Specifically, the landscape Liliyo in Ivory Coast is at $5.88°$N and in the humid forest zone. The predominant soil type is Ferralsol (FAO, 2014). The landscape Tieningboué in Ivory Coast is at $8.14°$N and belongs to the forest savannah transitional zone. The soils are dominated by Nitisols and Lixisols (FAO, 2014). The landscape Midebdo is at $9.97°$N and in the sub-humid savannah of Burkina Faso. Its dominant soil types include Lixisols, Gleysols, and Leptosols (FAO, 2014). The landscape Léo

is at 11.07°N and in the northern Guinean savannah of Burkina Faso and has Lixisols and Vertisols as the dominant soil type (FAO, 2014). The mean annual rainfall was approximately 1300 mm in Liliyo, and 900 mm in Tiéingboué, Midebdo, and Léo.

During July and August 2016, we sampled the soil from a total of 80 fields under yam cultivation across the four landscapes, i.e. 20 yam fields in each landscape. The fields were selected in advance by taking into account visual variation in soil color and texture across the landscape. The yam fields selected contained the maximum soil variability based on soil colour and cropping history, taking into account both local farmers' knowledge on soil fertility and agronomic extension expertise. Yam is typically planted on soil mounds, ranging from 5000 to 10000 mounds per hectare with a single yam plant per mound. Within each field, we sampled the soil at four adjacent mounds in square arrangement, which were spaced between 0.5 and 2 m. At each mound, 6 to 8 auger cores (25 mm in diameter) to the 0.3 m depth were taken at a radius between 0.15 and 0.3 m away from the center of a mound, depending on the size of the mounds. Then the soils from the four mounds were combined into one composite sample per field (around 500 to 1000 g of soil).

An additional set of 14 composite soil samples was collected by the International Center for Research in Agroforestry (ICRAF) at Liliyo from one sentinel site called "Petit-Bouaké" (UNEP, 2012). Sampling took place between 25 and 29 August, 2015 at positions that were previously selected for the Land Degradation Surveillance Framework (LDSF) in a spatially stratified manner (Vagen et al., 2010). The soil samples received from ICRAF were within the same landscape as the sampled soils in Liliyo within YAMSYS, but sampled from different positions. All soil samples were air-dried and stored in plastic bags until further analysis.

2.2 Soil reference analyses

The air-dried soil samples were crushed and sieved at 2 mm. About 60 to 70 g of the sieved soil was oven-dried at $60°C$ for 24 hours, of which 20 g were ball-milled. All chemical analyses except soil pH were conducted both on the soils sampled in yam fields ($n = 80$) and the LDSF soils obtained from ICRAF ($n = 14$).

The milled soils were analyzed for total C and macronutrient (N and S) concentrations using an elemental analyzer (vario PYRO cube, Elementar Analysensysteme GmbH, Germany). For each of the four landscapes, two soils were selected and analyzed based on three analytical replicates for quantifying within-sample variance of the elemental analysis. For the remaining samples, the analysis was not repeated. Sulfanilamide was used as a calibration standard for the dry combustion. For pH determination 10 g of air-dried soil per sample was placed in a 50 mL Falcon tube and 20 mL of de-ionized water was added. The samples were shaken in a horizontal shaker for 1.5 hours and measured for pH using a pH electrode (Benchtop pH/ISE meter model 720A, Orion Research Inc., USA).

Bioavailable micronutrient (Fe, Mn, Zn, and Cu) concentrations in soils were determined with the diethylenetriaminepentaacetic acid (DTPA) extraction method, as described in Lindsay and Norvell (1978). The extracting solution consisted of 0.0005 M DTPA, 0.01 M $CaCl_2$, and 0.1 M triethanolamine. Briefly, 10 g of the sieved (<2 mm) soils were extracted with 20 mL of DTPA solution. Micronutrient concentrations in the filtrates were measured by inductively coupled plasma optical emission spectroscopy (ICP-OES, a Shimandzu Plasma Atomic Emission Spectrometer ICPE-9820). Final DTPA extractable

concentrations of Fe, Mn, Zn, and Cu were calculated back to per kg dry soil. For each landscape, two soils were selected and analyzed in triplicates to assess analytical errors. For the remaining soils the analysis was not repeated.

For each sample, the concentrations of total element (Fe, Si, Al, K, Ca, P, Zn, Cu, and Mn) in the soil was assessed by energy dispersive X-ray fluorescence spectrometry (ED-XRF) measurements on 4 g of the milled soil with a SPECTRO XEPHOS instrument (SPECTRO Analytical Instruments GmbH, Germany). The soil was mixed with equal amount of wax using a ball mill and pressed into pellets. Exchangeable cations (Ca^{2+}, Mg^{2+}, K^+, Na^+, and Al^{3+}) were determined with the $BaCl_2$ method (Hendershot and Duquette, 1986). About 2 g of the air-dried soil (<2 mm) were extracted by shaking for 2 hours with 30 mL of 0.1 M $BaCl_2$ on a horizontal shaker (120 cycles min^{-1}). The suspension was filtered through no. 40 filter paper (Whatman, Brentford, UK). For each landscape, two soils were analyzed in analytical triplicates. The concentrations of exchangeable cations in the $BaCl_2$ extract were measured by inductively coupled plasma optical emission spectroscopy (ICP-OES, Shimandzu Plasma Atomic Emission Spectrometer ICPE-9820). Different $BaCl_2$ extract dilutions were used in order to obtain an optimal signal intensity for the quantification of specific elements across all samples. Concentration of H^+ per kg dry soil was calculated based on the pH measured in the $BaCl_2$ extractant. The $BaCl_2$ extraction does only slightly modify pH and is therefore an appropriate method to calculate effective CEC (CEC_{eff}) at native soil pH. Using the concentrations of the $BaCl_2$-extractable cations (i.e. Ca^{2+}, Mg^{2+}, K^+, Na^+, Al^{3+} and H^+), CEC_{eff} was calculated as sum of exchangeable cations in cmol of cation charge per kg dry soil. Exchangeable acidity was defined by the sum of exchangeable Al^{3+} and H^+. Base saturation in % was calculated as ratio of the sum of basic cations (Ca^{2+}, Mg^{2+}, K^+, and Na^+) in cmol(+) per kg soil to the CEC_{eff} multiplied by 100.

Particle size analysis was conducted by the International Institute of Tropical Agriculture (IITA) in Cameroon, as described in Bouyoucos (1951). Briefly, 50 g of dried 2 mm sieved soil was stirred with 50 mL 4 % sodium hexametaphosphate and 100 mL of deionized water in a mixer, to break down the aggregates into into individual particles. Readings with a hydrometer (ASTM 152 H, Thermco, New Jersey, USA) were taken after letting it stand in the suspension for 30 minutes. The silt content was calculated by subtracting the measured proportion of sand and clay from 100%.

Spectroscopic measurements

The milled soils ($n = 94$) were measured on a Bruker ALPHA DRIFT spectrometer (Bruker Optics GmbH, Ettingen, Germany), which was equipped with a ZnSe optics device, a KBr beamsplitter, and a DTGS (deuterated tri-glycine sulfate) detector. Mid-IR spectra were recorded between 4000 cm^{-1} and 500 cm^{-1} with a spectral resolution of 4 cm^{-1} and a sampling resolution of 2 cm^{-1}. Reflectance (R) spectra were transformed to apparent absorbance (A) using $A = \log_{10}(1/R)$ and corrected for atmospheric CO_2 using macros within the OPUS spectrometer software (Bruker Corporation, US). The spectra were referenced to a IR-grade fine ground potassium bromide (KBr) powder spectrum, which was measured prior to the first soil sample and measured every hour again. All spectra were recorded by averaging 128 scans (internal measurements) to improve the signal-to-noise ratio for each of the three independent replicate samples of each soil.

2.3 Spectroscopic modeling

2.3.1 Processing of soil spectra

Three replicates of spectra were averaged for each sample. The spectra were transformed by using a Savitzky-Golay smoothed first derivative using a third-order polynomial and a window size of 21 points (42 cm^{-1} at spectrum interval of 2 cm^{-1}) (Savitzky and Golay, 1964). Prior to spectral modeling, Savitzky-Golay preprocessed spectra were further mean centered and scaled (divided by standard deviation) at each wavenumber.

2.3.2 Model development and validation

The measured soil properties were modeled by applying partial least squares regression (PLSR) (Wold et al., 1983) with the preprocessed spectra as predictors. The models were fitted using the orthogonal scores PLSR algorithm. 5-times repeated 10-fold cross-validation was performed to provide unbiased and precise assessment of PLSR model performance (Molinaro et al., 2005; Kim, 2009). For each individual soil property, the number of factors for the most accurate PLSR model was tuned separately. For each soil property model, the sample set was repeatedly randomly split into $k = 10$ (approximately) equally-sized subsets without replacement for all repeats $r = 1, 2, .., 5$ and all candidate values in the tuning grid with the number of PLSR factors (ncomp) $= 1, 2, ..., 10$. Within each of the $r \times \text{ncomp} = 5 \times 10 = 50$ resampling data set splits, each of the 10 possible held-out and model fitting set combinations (folds) was subjected to candidate model building at the respective ncomp, using $k - 1 = 9$ out of 10 subsets and remaining held-out samples were predicted based on the fitted models. The root mean square error (RMSE, eq. (1)) of the held-out samples was calculated by aggregating all repeated *K*-fold cross-validation predictions ($\hat{y}_i$) and corresponding observed values (y_i) grouped by ncomp, which resulted in a cross-validated performance profile RMSE vs. ncomp.

$$\text{RMSE} = \sqrt{\frac{\sum_{i=1}^{n}(\hat{y}_i - y_i)^2}{n}} \tag{1}$$

Based on this performance profile, the minimal ncomp among the models whose performance was within a single standard error ("One standard error" rule, (Breiman et al., 1984)) of the lowest numerical value of RMSE was selected.

Model assessment was done with the best factors for each property using cross-validation hold outs. We reported the cross-validated measures RMSE, R^2 (coefficient of determination) obtained via linear least-squares regression, and ratio of performance to deviation (RPD), after averaging predictions across repeats. The RPD index is the ratio of the chemical reference data standard deviation (s_y) to the RMSE of prediction.

$$\text{RPD} = \frac{s_y}{\text{RMSE}} \tag{2}$$

Besides calculating the above listed performance measures, the uncertainty of spectral estimates was graphically reported for each soil sample, using prediction means and 95% confidence intervals derived from cross-validation repeats ($n = r = 5$; Eq. 3 and 4).

$$S_n^2 = \frac{1}{n-1} \sum_{i=1}^{n} \left(y_i - \overline{\hat{y}_i} \right)^2 \tag{3}$$

$$\overline{\hat{y}_i} \pm t(n-1, 1-\alpha/2) \frac{S_n}{\sqrt{n}}; \alpha = 0.05 \tag{4}$$

To cover the full training data space in the models for future sample predictions, the final PLSR models were rebuilt using the entire training set and the respective values of optimal final number of PLSR components determined by the procedure described above.

2.3.3 Model interpretation

The mid-IR spectra contain complex information about soil composition and properties. To establish a predictive relationship, statistical models need to find relevant spectral features for each soil property. Model interpretation requires a variable importance assessment to decide on the contribution of spectral variables to prediction and to explain spectral mechanisms. Therefore, we conducted model interpretation based on the variable importance in projection (VIP) method (Wold et al., 1993; Chong and Jun, 2005), using the model at respective best number of factors (ncomp). The VIP measure v_j was calculated for each wavenumber variable j as

$$v_j = \sqrt{p \sum_{a=1}^{A} \left[\mathrm{SS}_a \left(w_{aj} / \|w_{aj}\| \right)^2 \right] / \sum_{a=1}^{A} (\mathrm{SS}_a)} \tag{5}$$

where w_{aj} are the PLSR weights for the ath component for each of the wavenumber variables and SS_a is the sum of squares explained by the ath component:

$$\mathrm{SS}_a = q_a^2 t_a^{\mathsf{T}} t_a \tag{6}$$

where q_a are the scores of the predicted variable y and t_a are the scores of the predictors X. These VIP scores account for multicollinearity found in spectra and are considered as robust measure to identify relevant predictors. Important wavenumbers were classified with a VIP score above 1. A variable with VIP above 1 contributes more than the average to the model prediction. For model interpretation, we only computed VIP at the respective finally chosen number of PLS components a_{final} for each considered model. We focused on a selection of three well performing models with $R^2 \geq 0.8$ (RPD ≥ 2.3) to illustrate model interpretation. These were total C, total N and clay content.

2.4 Statistical software

The entire analysis was performed using the R statistical computing language and environment (version 3.6.0) (R Core Team, 2017). We used the pls (Mevik et al., 2019) package for PLSR, as described by Martens and Naes (1989). Cross-validation resampling, model tuning, and assessment was done using the caret package (Kuhn et al., 2019). Custom functions from the simplerspec package were used for spectroscopic modeling (Baumann, 2019). All data and code to reproduce the results of this study is available online via Zenodo (Baumann, 2020).

3 Results

Measured properties and mid-IR estimates of yam soils

The distribution of soil properties of the yam fields showed a wide variation across the landscapes (Figure 1). Total C concentrations across all fields ranged from 2.4 g C kg^{-1} soil to 24.7 g C kg^{-1} soil. Total C values at the landscape scale were the lowest (median) in Léo and the highest in Tiéningboué. Soils from yam fields in the two landscapes from Ivory Coast (13.0 ± 5.4 g C kg^{-1} soil; mean ± standard deviation) had relatively higher total C compared with the fields in the landscapes in Burkina Faso (6.1 ± 3.6 g C kg^{-1} soil). The median value and variation of CEC$_{eff}$ exhibited similar patterns across the landscapes to total C. Total N concentrations across all fields ranged from 0.18 g N kg^{-1} soil to 2.48 g N kg^{-1} soil. Total N within and across the four landscapes exhibited a similar pattern as total C. Generally, the landscapes in Burkina Faso were low in total N compared to those from Ivory Coast (0.44 ± 0.24 g N kg^{-1} soil vs. 1.09 ± 0.46 g N kg^{-1} soil). Median total N concentrations were almost identical for Liliyo and Tiéningboué with 1.1 g N kg^{-1} soil). Total S concentrations varied between 41 mg S kg^{-1} soil to 242 mg S kg^{-1} soil across all fields, and showed a similar pattern as total C and N. The yam fields in the landscapes of Bukina Faso had on average more than two times higher total S than the other landscapes. Total P concentrations were in a similar range for the landscapes Léo, Midebdo, and Liliyo. In Tiéningboué, total P values were almost two times higher than the other fields (817 mg S kg^{-1} soil vs. 453 mg P kg^{-1} soil), with more within-landscape variation.

The concentrations of total Fe, total Al, total Ca, total Zn, and total Cu in the soil tended to be higher for the landscapes in Ivory Coast than in Burkina Faso (Figure 1). To give an example, median concentrations of total Ca were 2.16 g Ca kg^{-1} soil in fields sampled from the Tiéningboué region, and similar in Liliyo (= 1.90 g Ca kg^{-1} soil), while they were markedly lower in Léo and Midebdo (= 0.90 vs. 1.26 g Ca kg^{-1} soil). In general, the ranges for total micronutrient contents were more variable in the landscapes of Ivory Coast (e.g., range = 14.0–57.0 mg Zn kg^{-1} soil in Liliyo; lowest range in Léo = 12.2–19.7 mg Zn kg^{-1} soil). Total K concentration was highly variable within and across the landscapes (overall range = 0.5–34.1 g K kg^{-1} soil), and lowest in Midebdo (range = 0.9–8.9 g K kg^{-1} soil), while the highest total K median was measured in yam fields of Léo (range = 4.1–25.0 g K kg^{-1} soil).

Median extractable Fe and its interquartile ranges were comparable across the landscapes (see Figure 1). However, there were some fields where extractable Fe reached values higher than 100 mg Fe kg^{-1} soil. Median extractable Zn values showed a similar pattern as total C, with the highest median values and interquartile range in Tiéningboué and had the lowest in Léo. In
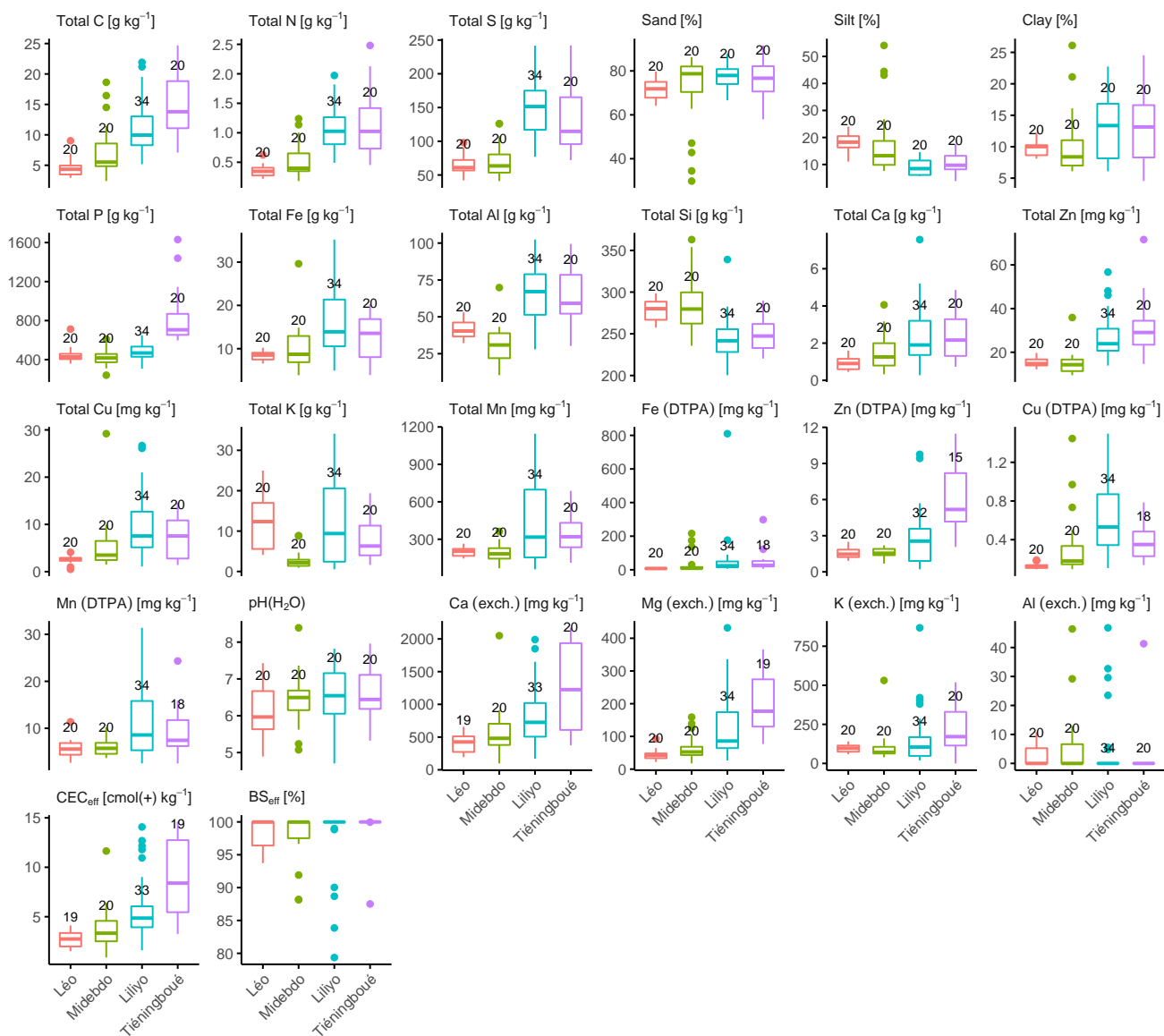
Figure 1. Reference measurements of soil chemical properties. Léo and Midebdo are two yam growing landscapes in Burkina Faso, and Lilyo and Tiéningboué are in Côte d'Ivoire. The chemically analyzed soils ($n = 94$) originated from 20 yam fields per landscape, and 14 additional soils from the Lilyo region were provided by the World Agroforestry Center (ICRAF). C = carbon, N = nitrogen, P = phosphorus, Fe = iron, Al = aluminum, Si = silicum, Ca = calcium, Zn = zinc, Cu = copper, K = potassium, Mn = manganese. Bioavailable micronutrients were measured by the diethylenetriaminepentaacetic acid (DTPA) extraction method. Ca(exch.), Mg(exch.), K(exch.), and Al(exch.) signify exchangeable elements determined with $BaCl_2$ extraction. CEC_{eff} = effective cation exchange capacity, BS_{eff} = effective base saturation. The number of soils analyzed for each individual property is indicated above the the 75% percentile.

comparison, the highest median values and interquartile range of extractable Cu and Mn were found in Liliyo. For extractable Zn, Cu, and Mn median values and interquartile range were higher in the two landscapes in Ivory Coast than the two landscapes in Burkina Faso.

Across all samples and landscapes, soil pH varied between 4.7 and 8.4. Median pH was comparable in Tiéningboué (= 6.4), Liliyo (= 6.5), and Midebdo (= 6.5). Median pH of yam fields in Léo (= 6.0) was lower than in the other landscapes. Exchangeable K, Ca, and Mg concentrations showed similar patterns across the four landscapes. In Burkina Faso, each of the exchangeable cations showed relatively low median concentrations across the fields and less landscape-level variation than in Ivory Coast. In general, the highest median and variation of exchangeable cations among the landscapes were measured in the yam field soils of Tiéningboué. Median exchangeable Al values were comparable among the landscapes, although there were some outliers with exchangeable Al > 20 mg kg^{-1} soil for Midebdo, Liliyo, and Tiéningboué. The CEC$_{eff}$ ranged from 0.9 cmol(+) kg^{-1} soil to 14.6 cmol(+) kg^{-1} soil across all fields and landscapes. Median CEC$_{eff}$ tended to decreases in the following order across landscapes: Léo > Midebdo > Liliyo > Tiéningboué. The interquartile range of CEC$_{eff}$ was also the greatest in Tiéningboué and the smallest in Léo.

Reference measurements for total N, S, exchangeable Ca, exchangeable Mg and CEC$_{eff.}$ were closely correlated with total C (Figure 2; $0.71 \leq r \leq 0.92$ (CEC$_{eff.}$)). Also, total Ca, Al, and clay content correlated closely with total C ($r > 0.70$). Clay contents were weakly related to silt ($r = 0.21$), while sand had a markedly negative relationship with silt ($r = -0.89$). Bioavailable Zn (DTPA) was co-varying with both CEC$_{eff.}$ ($r = 0.58$) and total Zn ($r = 0.59$). Bioavailable Cu (DTPA) had a strongly positive association to total Cu ($r = 0.90$). Exchangeable K (BaCl$_2$) had the strongest relationship with total C and CEC$_{eff.}$ ($r = 0.63$, and $r = 0.64$).

3.1 Soil mid-IR spectroscopic models

Among the measured soil properties, mid-IR PLSR models for total K ($R^2 = 0.96$) and total Al ($R^2 = 0.97$) were best performing (Table 1). Out of a total of 27 soil attributes, 11 were well quantified by the models ($R^2_{cv} \geqslant 0.75$; Figure 3). The confidence intervals derived from cross-validation prediction were very narrow, showing that all PLSR models were stable. Within this group of stable models, four soil attributes are directly related to the mineralogy (total Fe, Al, K and Ca), three are related to soil organic matter (total C, N and S), one to texture (clay fraction), one to plant nutrition (exchangeable Fe), and two related to mineralogy and plant nutrition (exchangeable Ca and CEC$_{eff}$). More specifically, total C was accurately predicted, with an R^2 of 0.92 and a RMSE of 1.6 g C kg^{-1} soil. The models were also able to predict total N well ($R^2 = 0.89$; RMSE = 0.16 g N kg^{-1} soil). Prediction accuracy of total S was slightly lower than for total C, but its goodness-of-fit and RMSE suggest that the model was reliable for prediction. However, exchangeable K ($R^2 = 0.28$) and BS$_{eff}$ ($R^2 = 0.24$) were poorly predicted (Table 1). Predictions for percent clay were reliable ($R^2 = 0.81$; RMSE = 2.1%), whereas predictions for percent sand ($R^2 = 0.45$; RMSE = 8.1%) and percent silt ($R^2 = 0.41$; RMSE = 6.5%) were not accurate. Finally, chosen models of all soil attributes had between 1 and 9 PLSR components.
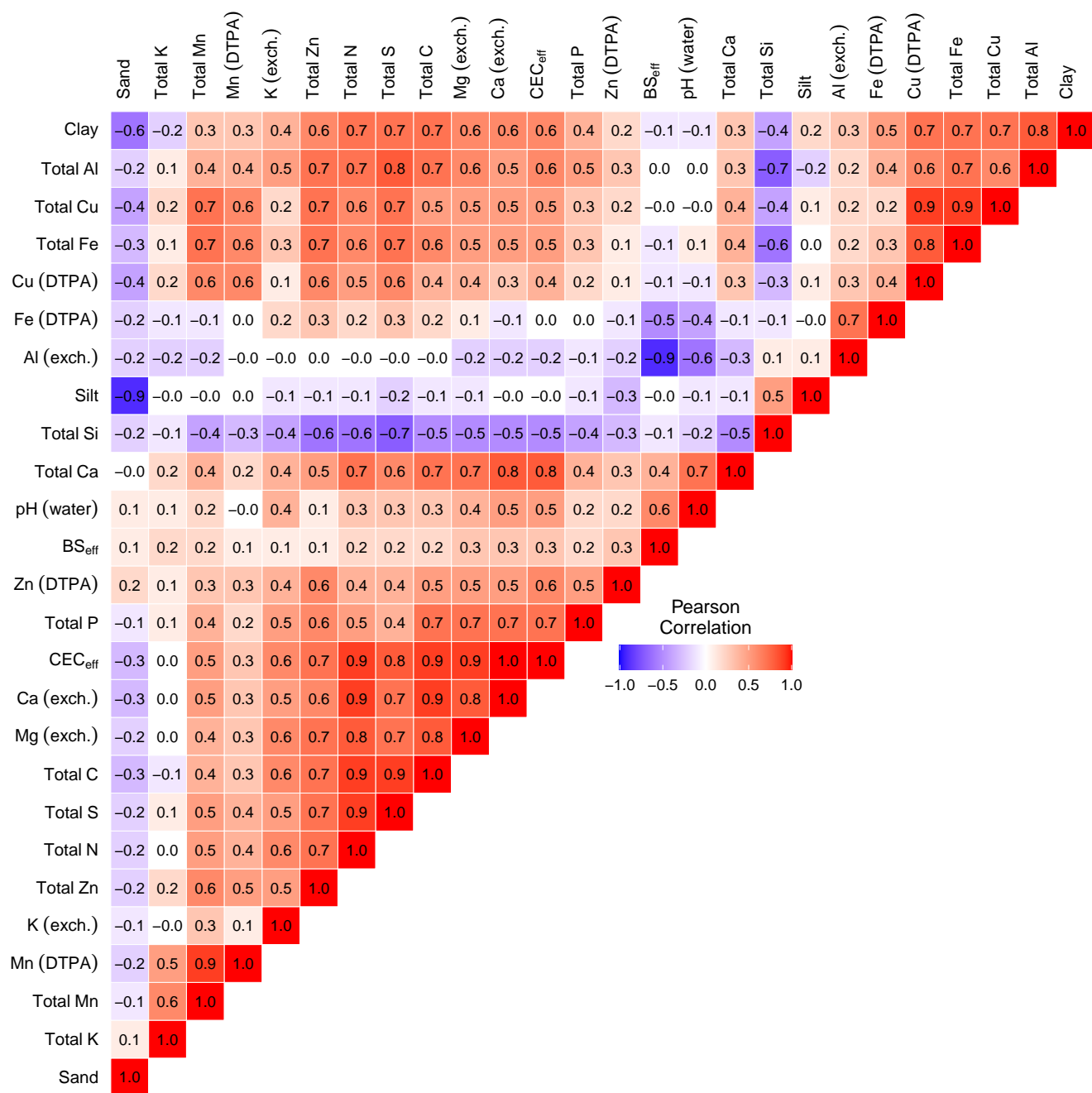
Figure 2. Correlation matrix of soil properties measured on each 20 soils sampled from individual yam fields per landscape, and 14 additional agricultural soils received from the World Agroforestry Center ($n = 94$; see Figure 1 for further details and abbreviated chemical properties). Pearson correlation coefficients (r) were rounded to 1 digit.

Table 1. Descriptive summary of measured (meas.) soil reference data (see Figure 1) and evaluation results of cross-validated PLSR models. All samples across the four landscapes were aggregated into a single model per respective soil property. Model evaluation was done on held-out predictions of 5 times repeated 10-fold cross-validation (cv) at the finally selected number of PLSR components (ncomp). CV = coefficient of variation, RMSE = root mean square error, RPD = ratio of performance to deviation. C = carbon, N = nitrogen, P = phosphorus, Fe = iron, Al = aluminum, Si = silicum, Ca = calcium, Zn = zinc, Cu = copper, K = potassium, Mn = manganese. Bioavailable micronutrients were measured by diethylenetriaminepentaacetic acid (DTPA) extraction. Ca(exch.), Mg(exch.), K(exch.), and Al(exch.) signify exchangeable elements determined with $BaCl_2$ extraction. CEC_{eff} = effective cation exchange capacity, BS_{eff} = effective base saturation.

Soil attribute	n	$Min_{meas.}$	$Max_{meas.}$	$Med_{meas.}$	$Mean_{meas.}$	$CV_{meas.}$	ncomp	$RMSE_{cv}$	R^2_{cv}	RPD_{cv}
Total C [g kg^{-1}]	94	2.4	24.7	8.5	9.9	58	6	1.6	0.92	3.6
Total N [g kg^{-1}]	94	0.18	2.48	0.72	0.81	61	6	0.16	0.89	3.0
Total S [mg kg^{-1}]	94	41	242	99	111	46	2	20	0.85	2.6
Sand [%]	80	29.8	91.6	75.6	74.2	14	2	8.1	0.42	1.3
Silt [%]	80	3.9	54.1	12.0	14.1	60	2	6.5	0.41	1.3
Clay [%]	80	4.5	26.1	10.1	11.6	42	2	2.1	0.81	2.3
Total P [mg kg^{-1}]	94	240	1631	467	530	40	3	131	0.61	1.6
Total Fe [g kg^{-1}]	94	4	35	10	12	54	5	3	0.81	2.3
Total Al [g kg^{-1}]	94	10	102	48	53	42	5	4	0.97	6.0
Total Si [g kg^{-1}]	94	200	363	262	262	12	3	20	0.59	1.6
Total Ca [g kg^{-1}]	94	0.3	7.6	1.4	1.9	70	5	0.6	0.78	2.2
Total Zn [mg kg^{-1}]	94	9.5	71.6	19.1	22.6	49	1	6.7	0.63	1.7
Total Cu [mg kg^{-1}]	94	0.5	29.2	4.7	6.8	87	7	3.2	0.71	1.9
Total K [g kg^{-1}]	94	0.5	34.1	5.8	9.5	91	7	1.7	0.96	5.1
Total Mn [mg kg^{-1}]	94	59.2	1146.0	221.5	308.0	74	5	116.4	0.74	2.0
log(Fe(DTPA)) [mg kg^{-1}]	92	1.0	6.7	2.7	2.9	38	9	0.5	0.77	2.0
Zn (DTPA) [mg kg^{-1}]	87	0.2	11.5	1.9	2.8	89	3	2.1	0.25	1.1
Cu (DTPA) [mg kg^{-1}]	92	0.1	1.5	0.2	0.4	89	6	0.2	0.74	2.0
Mn (DTPA) [mg kg^{-1}]	92	2.5	31.4	6.5	8.6	69	3	4.0	0.55	1.5
pH_{H_2O}	80	4.7	8.4	6.4	6.4	11	8	0.5	0.61	1.6
Ca (exch.) [mg kg^{-1}]	92	98	2170	604	774	70	5	237	0.81	2.3
Mg (exch.) [mg kg^{-1}]	93	18	432	76	113	84	3	58	0.62	1.6
K (exch.) [mg kg^{-1}]	94	0	868	104	145	95	1	120	0.28	1.2
Al (exch.) [mg kg^{-1}]	94	0	47	0	4	258	2	9	0.21	1.1
CEC_{eff} [cmol(+) kg^{-1}]	91	0.9	14.6	4.2	5.3	67	6	1.4	0.84	2.5
BS_{eff} [%]	91	79	100	100	98	4	2	3	0.24	1.1

3.1.1 Model interpretation

A large proportion of absorptions had VIP > 1 for each the total C, total N and clay models (Figure 4). Important wavenumbers
270 (VIP > 1) for total C were mostly between 3140 cm^{-1} and 1230 cm^{-1}. Besides clear absorption peaks, there were relatively
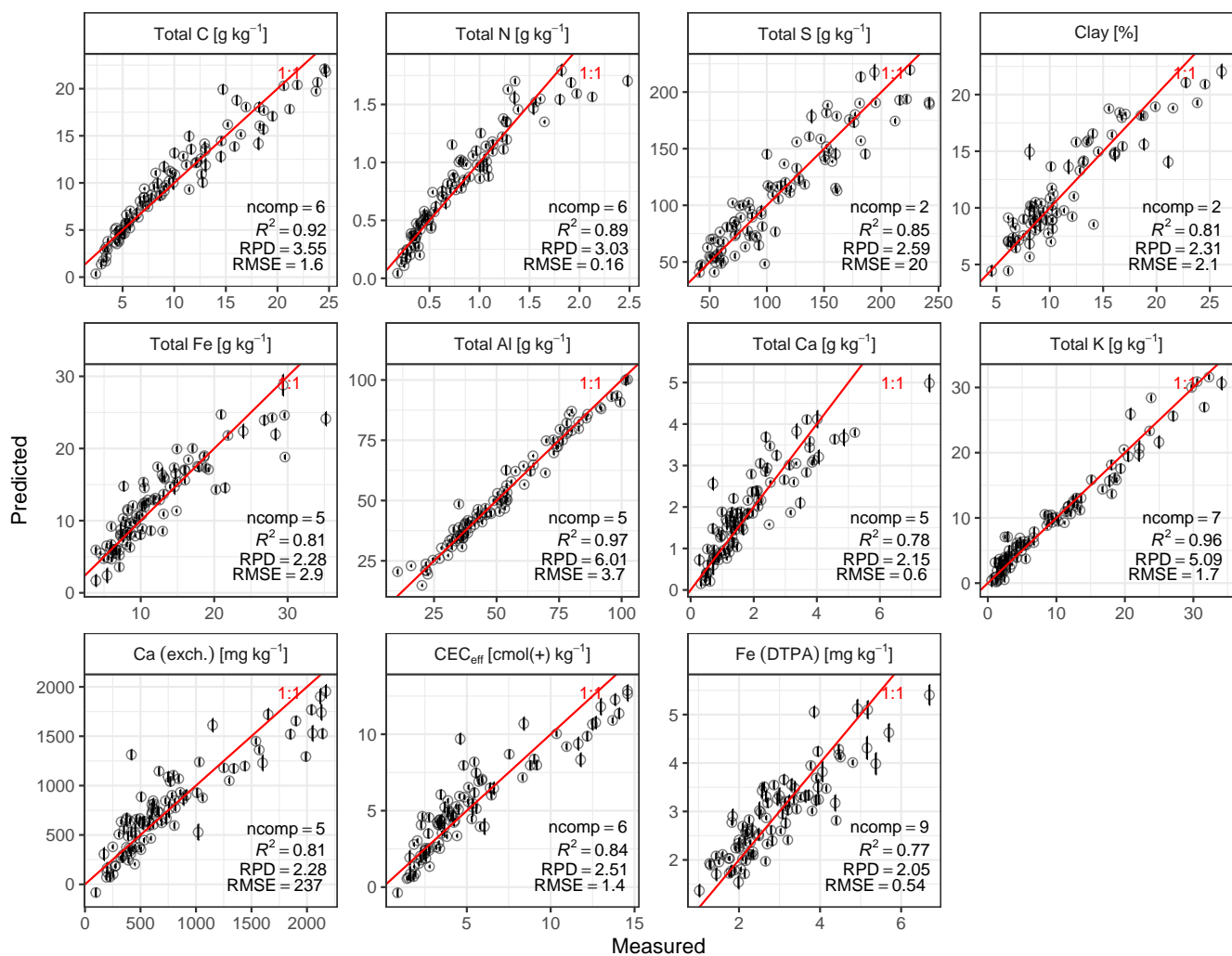
Figure 3. Cross-validated predictions of soil properties derived from best mid-infrared (mid-IR) partial least squares regression (PLSR) models vs. laboratory reference measurements (see Figure 1). Average estimates, their confidence intervals (error bars), and evaluation metrics were derived with $5 \times$ repeated 10-fold cross-validation. ncomp = number of PLSR components of most accurate final models, RSME = root mean square error, RPD = ratio of performance to deviation. Only soil properties modeled with $R^2 > 0.75$ are shown. CEC$_{\text{eff}}$ = effective cation exchange capacity. Exchangeable (exch.) elements were determined with $BaCl_2$. Bioavailable Fe was determined diethylenetriamine-pentaacetic acid (DTPA) extraction.

continuous spectral features that were important to the models. For example, the relatively continuous and smooth spectral region between the alkyl $C-H$ vibrations at 2855 cm^{-1} and 2362 cm^{-1} had comparable contribution to the model as peak regions associated with total C prediction. The VIP patterns across wavenumbers were almost identical for total C and N models, and its reference measurements were strongly correlated ($r = 0.94$; Figure 2). In contrast, the clay content model deviated from the total C model in particular regions, for example around the kaolinite $OH-$ feature at 3620 cm^{-1} or at kaolinite $Al-O-H$ vibrations at 934 cm^{-1} and 914 cm^{-1}.
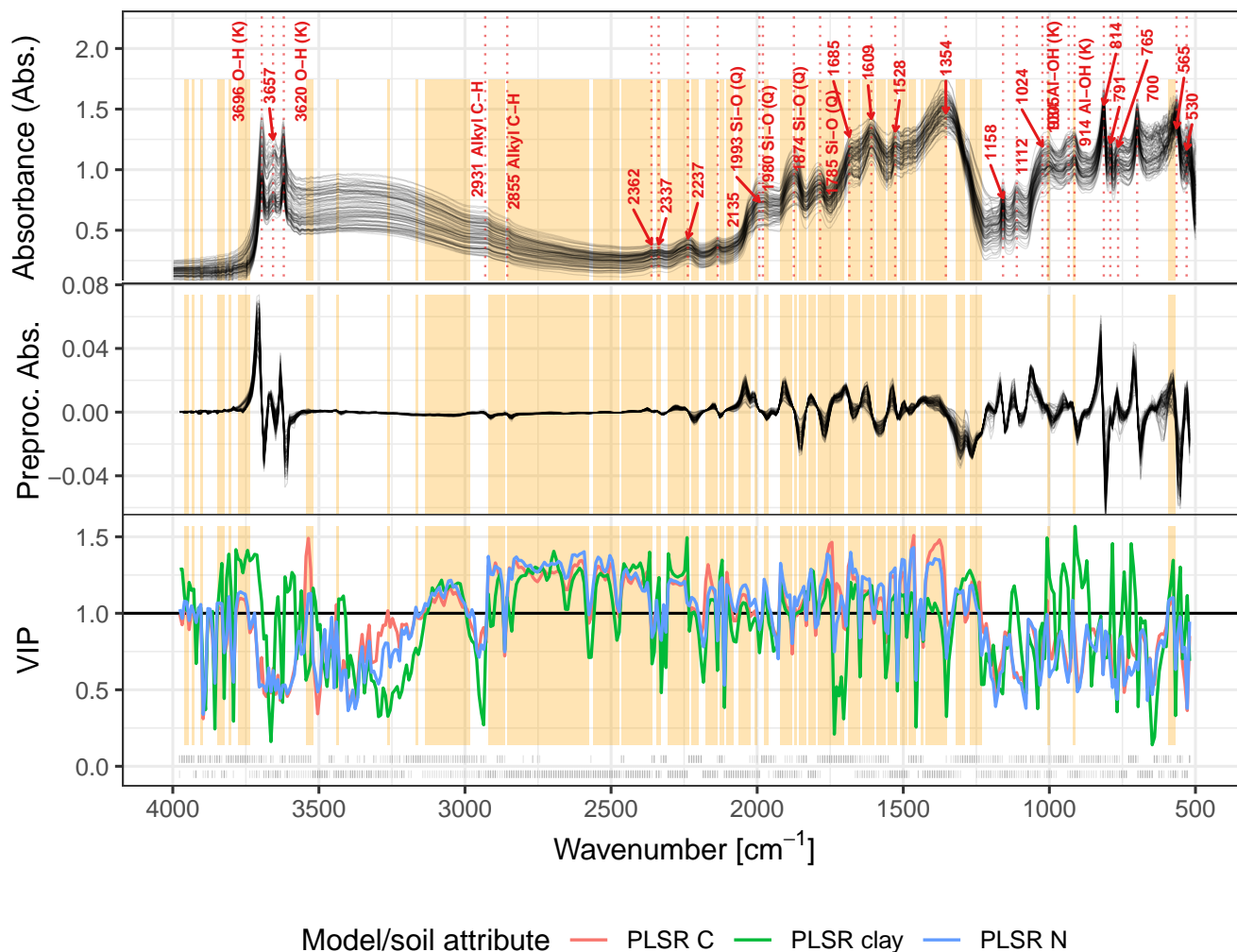
Figure 4. Variable importance analysis of partial least squares regression (PLSR) models for the concentrations of total soil C and total N, and clay content, including overlaid raw and preprocessed spectra. Top panel shows resampled mean sample absorbance spectra ($n = 94$). Prominent peaks were identified as local maxima with a span of 10 points 20 cm^{-1} for the selected wavenumbers. Fundamental mid-IR vibrations that are well described in the literature (e.g., Madejová et al., 2002; Rossel and Behrens, 2010; Stevens et al., 2013) were added as labels when identified peaks matched literature assignments. (Q) stands for quartz and (K) for kaolinite. The middle panel depicts preprocessed spectra (Savitzky-Golay first derivative with a window size of 21 points (42 cm^{-1}); 3rd order polynomial fit). The bottom panel shows variable importance in the projection (VIP) for three selected well performing PLSR models (total C, total N and clay; $R^2 > 0.81$). The black horizontal line at VIP $= 1$ indicates the threshold above where absorbance at the wavenumbers explain more than average to the prediction of a certain soil property. Dashed points closely below the $y = 0$ line of the VIP graph visualize positive (above $y = 0$) and negative (below $y = 0$) PLSR β coefficients.

Discussion

3.2 Accuracy and relevance of mid-IR spectroscopy for agronomic diagnostics

Timely and accurate estimates of multiple soil properties are required to better understand and predict soil constraints across
the yam belt in West Africa. The soil spectral library from our study, which includes four landscapes of the yam belt, can
be practical to diagnose and monitor (and eventually manage) soil fertility that is considered to be low and therefore being
a major constraint to yam production in West Africa. Specifically, our results show that properties closely related to organic
matter — total amount of C, (micro)-nutrients, and exchangeable cations — can be accurately estimated using mid-IR spectra
and in the selected yam growing landscapes (Figure 3). Soil organic matter plays a crucial role during vegetative growth
and tuber formation phases of yam, as it guarantees among many other functions the storage and availability of essential
nutrients and water needed for yam and tuber growth throughout the season, and as well prevents soil erosion due to it's
structural stabilization capacity. It promotes soil aggregation, which stabilizes soil organic matter and protects it from microbial
decomposition (Six et al., 2006).

Fertilizers are becoming more essential to replenish mineral nutrients for prolonged cropping. Nevertheless, soil organic mat-
ter is at high risk of depletion in the regions because of the increasing land use frequencies and shorter fallows to restore the soil
organic C pools. While it is pivotal to develop innovative crop and soil management solutions to this problem (O'Sullivan and
Jenner, 2006; Frossard et al., 2017; Kiba et al., 2020), it is also crucial to perform a separate but complementary activity to give
feedback on potential soil changes: developing and applying soil conventional and proximal sensing methods. When testing
sustainable soil and crop management options, for example to derive region-specific and farm-adapted nutrient management
strategies, putting both validated quantitative statements on the status of soil organic carbon and local farmers' soil knowledge
into the equation is crucial (Wawire et al., 2021). Inevitably, both determining the inherent soil status (i.e., soil texture and
organic carbon) and measuring the chemical and physical environment that regulates nutrient availability at trial sites (e.g.,
pH), is of agronomic and environmental importance (Foster, 1981). Maintaining and improving soil quality attributes will be
paramount to sustain soils' ecosystem functions and crop yields over time. Activities to maintain and improve soil properties
can for example be oriented towards fostering nutrient recycling.

Quick and reasonably accurate soil estimates derived from mid-IR spectra and empiric models as for example outlined
in this study can inform site-adapted timing, placing and form of nutrient supply based on local soil conditions. To give a
specific example, yam requires relatively large quantities of N and K (e.g., O'Sullivan, 2010); on light-textured soils, yam
can attainhigh tuber yields, but at a high risk of loosing large proportions of applied N and K to the environment (e.g., Diby
et al., 2011). Therefore, spectral estimates of texture can give an indication that applying larger amounts of N and K at once
would not improve yield potential under such situations. Hence, more frequent and local mineral applications of these nutrients
after crop emergence, eventually combined with organic mulch, could improve the fertilizer efficiency and mitigate negative
environmental impacts under these soil conditions. To estimate the availability of specific (micro)nutrients, however, more
efforts need to be made to measure them at fine temporal and spatial resolution.

310   The mid-IR model accurately estimated C (RMSE = 1.6 g kg^{-1} soil; Table 1; Figure 3). Mostly, only field-scale spectroscopic models achieve such accuracy (Nocita et al., 2015; Guerrero et al., 2016), whereas the predictive accuracy reported for larger-scale application of spectroscopic models is lower than for our model (Rossel and Webster, 2012; Stevens et al., 2013; Sila et al., 2016). Models covering a wide geographical range of soils often result in high prediction errors (Stenberg and Rossel, 2010). Despite different soil types and climate regimes across a wide geographic spacing between the calibration fields, we

315 achieved an accurate spectroscopic estimation of total C. The model was also able to reliably estimate a range of other important soil properties than total C. Specifically, other soil variables eligible for a mid-IR quantification include total N, total S, total Ca, total K, total Al, exchangeable Ca, Fe DTPA, CEC$_{\text{eff.}}$, and clay content ($R^2 > 0.75$). The close correlations of total C with N, S, exchangeable Ca, exchangeable Mg, CEC$_{\text{eff.}}$, total Ca, Al, and clay content (Figure 2) are consistent with Johnson et al. (2019), who reported very similar associations of clay content and exchangeable cations (Ca, Mg, K) as well as CEC$_{\text{eff.}}$ in soils

320 from rice fields ($0.54 \leq r \leq 0.65$) — nevertheless they spectrally modeled a considerable soil variability (20 countries in sub-Saharan Africa; 42 study sites) and a larger sample size ($n = 285$) using PLS regression. At the same time, the measured range and the error in spectral estimates of CEC were larger compared to ours (RMSE = 6.7 cmol(+) kg^{-1} vs. 1.4 cmol(+) kg^{-1}; range = 1.9–66.5 cmol(+) kg^{-1} vs. 0.9–14.6 cmol(+) kg). Even though, total K and Fe(DTPA) were poorly correlated with total C, their spectroscopic estimates were relatively accurate. This suggests that the mid-IR prediction of other soil properties is

325 largely based on their correlation with total C as well as other absorption features of many organic and mineral soil components having a specific IR adsorption.

    We also found reasonable prediction accuracy for Cu(DTPA) ($R^2 = 0.74$) and Mn(DTPA) ($R^2 = 0.55$), although soil nutrients that are extraction-based or dependent on surface chemistry usually have variable predictive performance (Janik et al., 1998). Since relationships between soil composition and soil matrix exchange processes are typically complex, some properties may

330 not be represented in the models in a straight-forward manner (Janik et al., 1998; Nocita et al., 2015).

    Although total elements are not necessarily a direct proxy for plant-available nutrients — with exception of total C from organic matter — they can be related to mineralogical status, which is influenced by weathering and nutrient supply. For example, total Fe from iron oxides can be important in controlling the availability of P (Parfitt et al., 1975) and total P can be correlated to available P in other cases. For yam — which is an understudied crop with a relatively large yield gap —

335 fertilizer response on N, P, and K is often absent on soils that have been under long fallow periods (O'Sullivan, 2010). Even more importantly, the number of thoroughly conduced yam fertilizer trials in a region and for distinct soil types are often not sufficient for site-specific calibration of soil tests with regard to fertilizer response and recommendations (O'Sullivan and Jenner, 2006).

3.3   Interpretation of spectral features

340 All mid-IR spectra that we measured for soils in the four landscapes exhibited a similar pattern of absorbance (Figure 4). The O-Si-O absorptions in quartz at 1080 cm^{-1}, 800–780 cm^{-1} and 700 cm^{-1} were a prominent feature in the spectra due to relatively high sand contents across the landscapes (range 30% to 92%, median 76%). Our spectra further had hydroxyl (OH) absorptions, which are typical for kaolin minerals, at 3695 cm^{-1} (surface OH), 3620 cm^{-1} (inner OH), 914 cm^{-1} (inner

OH), and 936 cm^{-1} (surface OH) (Madejová et al., 2002). The spectral pattern between the hydroxyl bands at 3695 cm^{-1} and 3620 cm^{-1} was relatively consistent and the intensity ratio of these flanking peaks was close to 1. This is typical for halloysite (0.8–0.9) while the ratio for kaolinite is often higher (1.2–1.5) and dickite lower (0.6–0.8) (Lyon and Tuddenham, 1960). The two weak intermediary stretching absorptions at around 3657 cm^{-1} and 3670 cm^{-1} indicate surface hydroxyls. Together with the absorption at 936 cm^{-1}, the spectra would suggest the presence of rather well-ordered prismatic halloysite (Hillier et al., 2016). This aligns well with the spectral patterns of soils that were assigned to the Halloysite archetype through similarity mapping (by comparison to the pure mineral spectra) by Sila et al. (2016). Our spectra confirm the presence of kaolin minerals, which reflects the advanced state of mineral weathering in these tropical soil types.

Our accurate predictions, which are comparable to field-scale calibrations, are most likely because of the relatively uniform mid-IR spectra we obtained from our samples and their linear relationships to some of the key properties. This suggests a relatively homogeneous soil chemical composition, particularly with regard to the mineralogy of the sampled soils. Still, the data set presented here is relatively small and no randomized spatial sampling strategy was used for selecting field locations. Therefore, we propose to implement a spectroscopy-driven approach to diagnose soils in more yam growing areas, as an effort to broaden the library to achieve better spatial coverage of soil variability.

4   Conclusions

We developed models with mid-IR spectra to estimate soil chemical and physical properties relevant to production of yam and other staple crops in four landscapes in the yam belt of West Africa. We tested the models for the important soil properties that are applied widely for agronomic performance evaluation. We showed that mid-IR spectroscopy models have the potential to cost-effectively and rapidly determine the distribution and variability of important soil properties across highly variable yam production landscapes in West Africa. Specifically, total C, total N, total S, total Fe, total Al, total K, total Ca, exchangeable Ca, CEC$_{eff}$, bioavailable Fe, and clay content can be quantified with RPD > 2 and $R^2 > 0.75$ when aiming to predict in the range of soil property values found in the environmental conditions covered by this study. We achieved spectral estimates with quite small uncertainties, that are typically reported for libraries at the geographical extent of a field or farm. The correlation analysis of measured values together with spectral inference helps improve our understanding of how soil properties are interrelated with soil functional composition. This study delivered parsimonious, unbiased and accurate mid-IR spectroscopy-based models to monitor and predict soil quality and to manage crop nutrition. Hence, we envision this pilot study as being a starting point to continuously update and adapt the mid-IR model library for more efficient site-specific and agronomically relevant soil estimates in the West African yam belt. This can bring better capacity to diagnose and and long-term monitor soils compared with traditional wet chemistry, and will hopefully ameliorate the soil conditions for sustainably meeting the demand of yam and other important staple crops in the regions.

Code and data availability.   All data and code to reproduce the results of this publication are publicly available under GNU General Public License v3.0, and can be accessed via the Zenodo archive and the corresponding github public repository (Baumann, 2020)

Author contributions.   Philipp Baumann carried out the research and analysis (soil sampling, sample preparation, soil chemical analysis, infrared spectroscopy, statistical modeling) under continuous support of the YAMSYS project team, and took the lead in writing the manuscript. All co-authors helped to improve the manuscript. Johan Six, Juhwan Lee and Emmanuel Frossard framed the idea of delivering validated models for soil properties relevant for yam growth in the four pilot regions in Burkina Faso and Ivory Coast. Valérie Hgaza and Delwende Kiba contributed to the selection representative yam fields that were sampled for our work.

Competing interests.   We declare that we have no conflict of interest.

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