



Evaluation of the Near Infrared Spectroscopy (NIRS) to predict chemical composition in *Ulva ohnoi*

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Abstract

Green algae of the genus *Ulva* have been identified as suitable organisms for biomass production and good candidates for the development of seaweed blue-biotech industries. The fluctuation of chemical composition during the growth of the algae, which depends largely on environmental factors, makes the development of rapid phenotyping protocols necessary. In this work the efficacy of Near Infrared Spectroscopy (NIRS) to predict dry matter, mineral fraction, nitrogen, and carbon contents directly from wet untreated samples and from dried samples was studied. Partial least square (PLS) models from spectra recorded on 80 samples were used to predict dry matter, and 44 samples to predict carbon, nitrogen and mineral fraction on a wet and dry weight basis. NIR models developed from spectra acquired on wet samples had good accuracy ($R^2 > 0.9$) for the prediction of N (on a ww and dw basis) and C (on a ww basis). Models with lower R^2 scores have been obtained for dry matter ($R^2 = 0.610$) and MF ($R^2 = 0.506-0.693$). The models developed to predict carbon and nitrogen directly on wet and untreated samples present NIRS as a valuable tool to determine these parameters in a rapid and low-cost way, allowing making decisions about the optimal harvesting time.

Keywords Near Infrared Spectroscopy (NIRS) · *Ulva ohnoi* · Chlorophyceae · Nitrogen · Carbon · Mineral Fraction

Introduction

An increasing interest in non-animal protein sources during the last decade has arisen and seaweeds have emerged as potential candidates (Pereira 2011). Seaweeds, also, have high potential as a source of bioactive compounds for human and animal consumption or pharmaceutical applications (Barrington et al. 2009; Holdt and Kraan 2011; Peña-Rodríguez et al. 2011; Rasyid 2017; Gomez-Zavaglia et al. 2019).

Biochemical composition in general, and protein content in particular, in wild-harvested seaweeds depends on species, harvesting location, and season (Pedersen et al. 2010; Pereira et al. 2012; Vasconcelos et al. 2022). Fluctuations in the composition of harvested seaweeds are a bottleneck for the seaweed industry development, which needs a stable supply with a stable composition.

The green algae of the genus *Ulva* have been identified as suitable organisms for biomass production and as good candidates for the development of seaweed blue-biotech industries. Moreover, regarding *Ulva ohnoi*, its high growth rates in temperate waters, and nitrogen and phosphorus biofiltration capacity present them as a suitable candidate for land-based cultivation (Lawton et al. 2013; Mata et al. 2016). When cultured in tanks, cultivation conditions (nutrients and light) affect growth rates and biochemical composition (Neori et al. 1991; Harrison and Hurd 2001; Msuya and Neori 2008; Angell et al. 2014; Oca et al. 2019; Toth et al. 2020); hence, cultivation conditions could be adapted from a food perspective (biochemical composition) or from a growth perspective (biomass yield) (Toth et al. 2020). Characterization of the biochemical composition in a fast way would be a useful tool to determine the best harvesting time to ensure specific seaweed properties.

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Biochemical composition determination in seaweed requires time, laborious methods with expensive equipment and trained personnel, and in general includes hazardous compounds. Near infrared spectroscopy (NIRS) is commonly used for routine analysis in food and feed industries (Manley and Williams 2021; Beć et al. 2022), and presents advantages such as their expeditiousness and versatility, which allows the determination of multiple parameters at once in different types of samples (powders, pellets, liquids, etc....) with little or no preparation. NIRS has also been shown as a valuable tool for measuring protein and other compounds in brown (Horn et al. 1999; Hay et al. 2010; Yang et al. 2021; Campbell et al. 2022; Cao et al. 2022) and red seaweeds (Tadmor et al. 2022).

In the present study near infrared spectroscopy (NIRS) was used to predict dry matter, mineral fraction, nitrogen, and carbon content in wet and dry samples of the green seaweed *U. ohnoi*.

Material and methods

Ulva ohnoi

Ulva ohnoi was collected at the Ebro Delta from the bioremediation ponds of an aquaculture facility in Sant Carles de la Ràpita, Spain (latitude, 40.62 N; longitude, 0.66 E). The species was genetically identified by DNA extraction and PCR amplification of the chloroplast *rbcl* gene following the protocol described in Hayden et al. (2003) with the primers used by Manhart (1994). It was maintained at the Aquaculture Laboratories of the Universitat Politècnica de Catalunya in Castelldefels (Spain) in indoor tanks fed by water coming from a *Solea senegalensis* recirculation aquaculture system (RAS) equipped with a biological and mechanical filter, water temperature control and aeration. *Ulva ohnoi* were cultivated in three circular tanks (28 cm water depth, 64 cm diameter) with bottom aeration to tumble the seaweeds. Seaweed tanks were illuminated by LED light sources (with photon flux densities on the water surface ranging from 160 to 1000 $\mu\text{mol photons m}^{-2} \text{s}^{-1}$). The seaweed stocking densities (0.8 to 3.0 kg m^{-2}) and the photon flux densities on the water surface were combined in order to obtain fronds with different ranges of dry matter, nitrogen and carbon content.

Seaweed sampling and analysis

Over the course of the study 80 samples of *U. ohnoi* were used (between February 2022 and May 2022). During the study the temperature range in the tanks was 12.5–19.5°C, pH 7.6–9.5, total ammonia nitrogen 0.01–0.20 mg L^{-1} , N-NO₃ 25–38 mg L^{-1} , P-PO₄ 0.7–4 mg L^{-1} and alkalinity 120–250 $\text{mg CaCO}_3 \text{L}^{-1}$. The NIR spectra for wet (ws) and

dry samples (ds) was acquired. In all the samples the dry matter (DM) was determined, and 44 out of the 80 samples were analyzed for mineral fraction (MF), carbon (C), and nitrogen (N) content.

Seaweed sampling and preparation

Seaweed fronds were randomly taken from the tanks and always the same protocol was followed to obtain wet and dry samples (ws and ds): excess water was removed in a mechanical spinner and fronds were rinsed (first with fresh water and after with distilled water) to remove epiphytes and salt; then, excess of water was removed again with the mechanical spinner (ws). NIR spectra on the wet samples (ws) were acquired and immediately the samples were weighed, obtaining the wet weight (ww) for dry matter determination. Samples were oven-dried at 60°C for 48 h, weighed (dry weight: dw), grounded in a mill (finely minced), and the NIR spectra on the dry samples (ds) were acquired. Samples were stored in closed containers and maintained in a dark and dry area until the chemical analysis.

Reference method analysis

Mineral fraction (MF) was quantified gravimetrically after incineration of the dry samples (around 1.5 g) for 14 h at 550°C in a muffle furnace (Lahaye and Jegou 1993). Carbon (C) and nitrogen (N) were analyzed using a CHN elemental analyser EA-CE 1108 (Thermo Fisher Scientific) with sulphanimide as a standard.

MF, C, and N were reported to the wet and dry weight of the seaweed (ww and dw).

Spectra acquisition

Since water has a high contribution in the NIR zone, sample spectra were acquired on wet (ws) and dry (ds) samples: one after the mechanical spinning (ws) and the other after being oven-dried and ground (ds) (see Sect. "Seaweed sampling and preparation"). In both cases, NIR spectra were acquired by diffuse reflectance, using a FT-NIR Antaris II (Thermo Fisher), equipped with a spinner module. Spectra were obtained as the average of 3 technical replicates acquired in the range between 10,000 and 4000 cm^{-1} , with a resolution of 8 cm^{-1} and 32 scans.

Data analysis

Statistical analysis

Chemical variables were analyzed to ensure that significant variation was captured among the samples. All analyses were performed using R software (v. 3.6.1; R Core Team,

Austria). Descriptive statistics (maximum, minimum, average) were calculated for each variable in order to describe the range of variation. The adjustment to a normal distribution was studied by means of the Shapiro test (“ggpubr” package). The correlation between all the variables was calculated with the Pearson correlation coefficient. Significant correlations ($p < 0.05$) were plotted in a correlation matrix using the “corrplot” package. Graphics were produced using the “ggplot2” package.

Multivariate model development

The whole spectral pre-treatment and data modeling process was performed by PLS Toolbox v8.9.1 software (Eigenvector research, Inc.). Different pre-treatments and their combinations were tested: Standard Normal Variate (SNV) and 1st (SG-1D) and 2nd (SG-2D) derivatives of Savitzky-Golay. Before the model's development, samples were divided into two sets: calibration set, for the model construction, and external validation or prediction set, to assess the model's predictive ability. These sets were automatically selected by using the Onion algorithm, selecting 66% of samples for calibration and the remaining 34% for external validation. After the automatic selection, for each parameter the sets were manually checked and modified if necessary, in order that all the samples with extreme reference values fall in the calibration set. To develop the predictive models for the studied parameters, Partial Least Squares (PLS) regression algorithm was used. It allows relating the spectra of all samples to their reference data, reducing considerably the number of variables. The new variables, known as latent variables (LVs), are spectral vectors directly related to the target parameter, which will be used for the prediction of new samples. The choice of the optimal number of LVs is a very important issue. If the LV number selected is too low, some important spectral information of the samples may not be sufficiently represented, while if it is too high, spectral noise information will be added to the model increasing the risk of overfitting. In the present work, for each model, the number of LVs was established according to their root mean square error (RMSE) decrease and accumulated variance and R^2 increase in each LV and regarding the previous one (in calibration and cross-validation). Cross-validation is an internal validation applied on the calibration samples. It has been applied automatically by the software, using the Venetian Blinds algorithm and splitting the samples in ten groups. Outliers of the model were discarded according to their Hotelling T^2 , Q residual values, and differences between measured and predicted values during the construction of PLS models. For each parameter and sample presentation (ws or ds), the best model was selected according to the higher determination coefficient (R^2) and lower root mean square error (RMSE), both in the calibration (C) and

prediction (P) set of samples. The predictive ability of the final models can be assessed by means of their reference-predicted scatterplots and the related statistics obtained in prediction (mainly R^2 and RMSEP).

Results

Chemical parameters

80 samples of *Ulva ohnoi* were used to determine DM and 44 out of the 80 to determine N, C, and MF. Since a good range of data from reference analyses is critical for developing robust calibration models, the distribution of individual values for each variable was analyzed. In Fig. 1 the range of the values (max and min) and the mean obtained for each parameter are shown. Individual values for DM, N (ww, dw), and C (ww) showed a good distribution along the range of variation, despite an isolated sample in the low part of the range in the case of N (dw) (Fig. 1). Individual values for C (dw) and MF (ww, dw) were concentrated around the mean (normal distribution). The Shapiro test for normality showed that C (dw) and MF (ww, dw) were adjusted to a normal distribution ($p > 0.01$), while the remaining variables were not normally distributed (DM showed a skewed distribution to high values, while N (ww, dw) and C (ww) showed non-symmetric and bimodal distributions).

Bivariate correlation analysis showed a significant positive correlation between DM and C and N contents (Fig. 2). Correlations among DM and N and C estimated on a ww basis were higher ($r = 0.97$ and $r = 0.98$, respectively) than on a dw basis ($r = 0.75$ and $r = 0.55$, respectively). MF on a ww basis showed a positive correlation with DM ($r = 0.86$), N ($r = 0.83$) and C ($r = 0.82$) (Fig. 2), i.e. samples with higher DM also accumulate more MF. However, on a dw basis MF content is negatively correlated with N (dw, $r = -0.42$) and C (dw, $r = -0.51$).

NIR spectra versus chemical composition

Figure 3 shows the raw and pretreated (SNV pretreatment) spectra measured from wet and dried samples. Spectra of wet samples differed from the dried samples ones, showing broad bands around 5000 cm^{-1} and 7000 cm^{-1} (Fig. 3a). SNV correct scattering effects (Fig. 3b), and SG-1D treatment (not shown) led to spectra with more defined peaks, although the absorbance of water remained very high in wet samples.

PLS models to estimate chemical variables were developed using different spectral pretreatments (SNV, SG-1D, SNV + SG-1D) on the entire spectral range (Table 1). The best pretreatment for each variable was selected based on the goodness of the resulting models. The performance

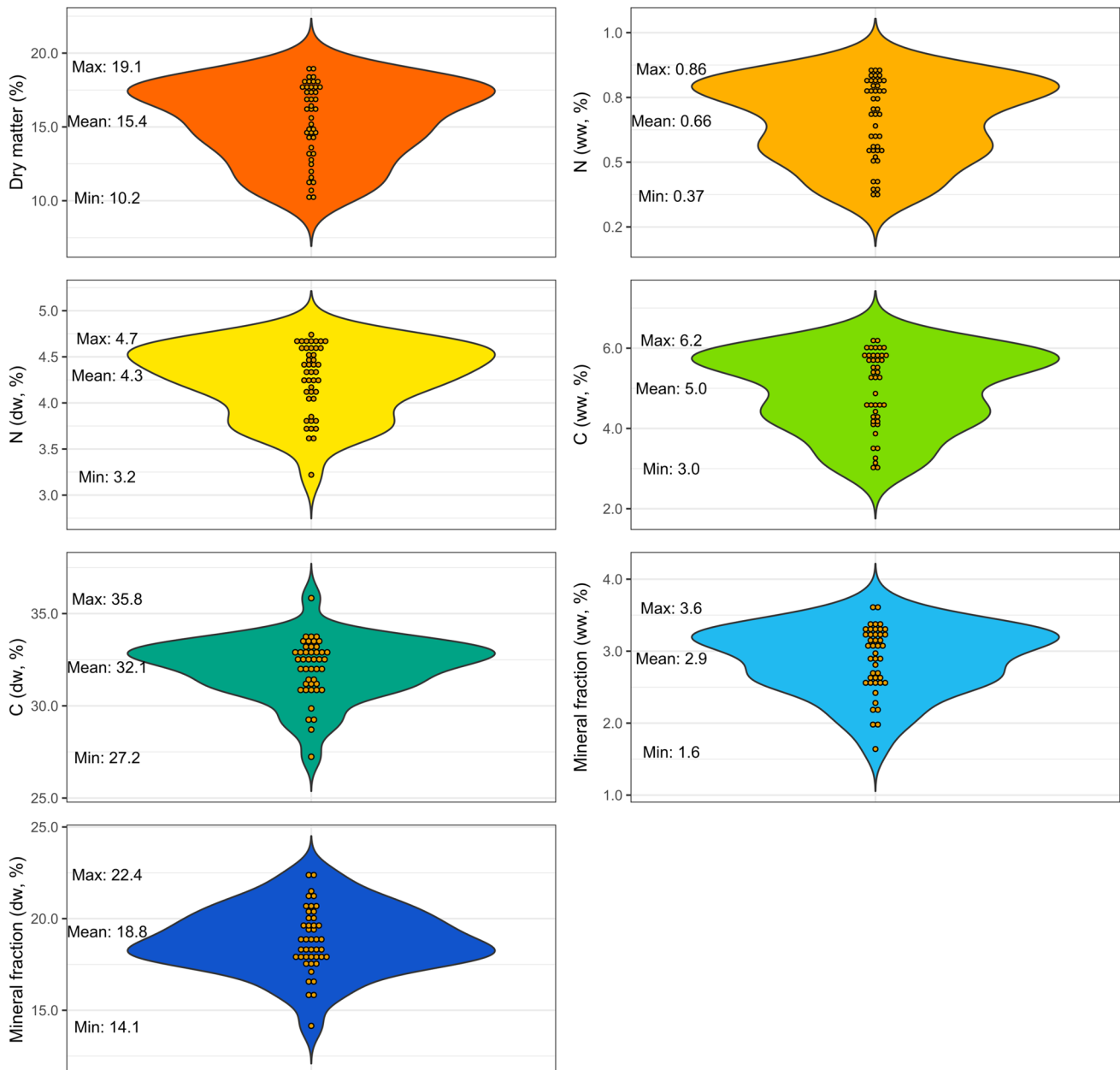


Fig. 1 Chemical composition of *Ulva ohnoi* samples used in this study: Violin plots showing the distribution of the samples along the range of variation for each parameter on a dry and wet weight basis

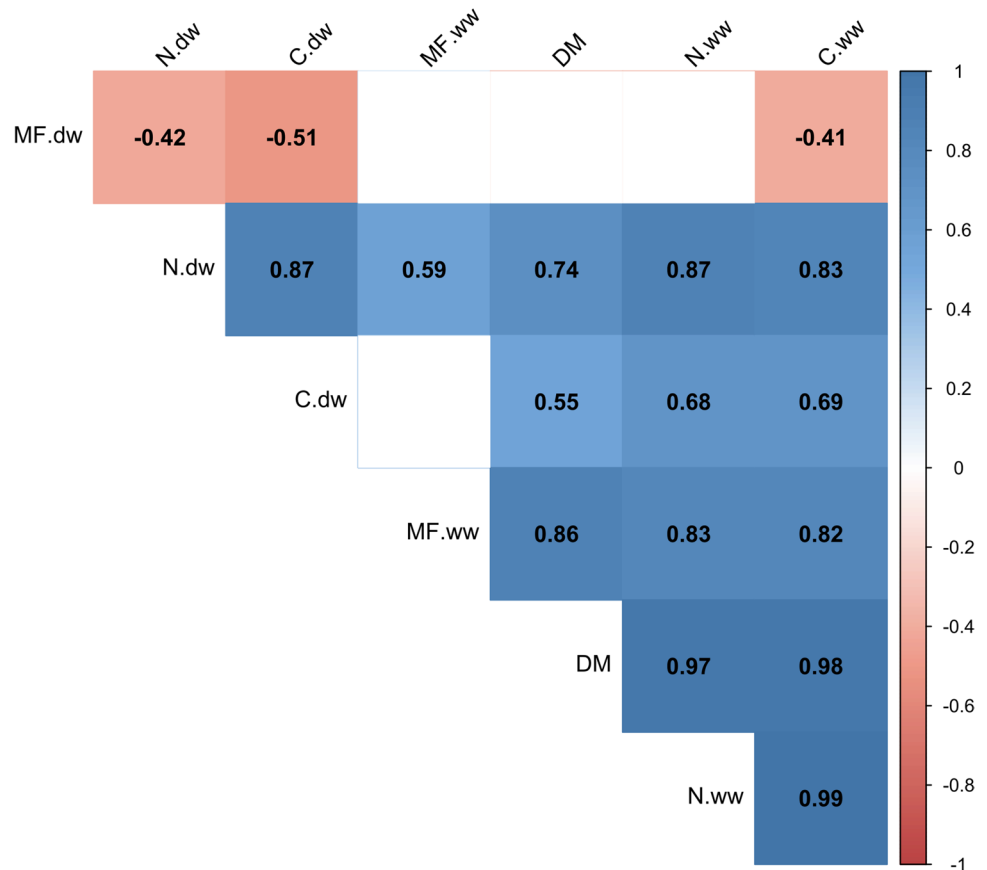
(dw and ww). Individual values of the samples are indicated with dots. Mean, Maximum, and Minimum values are indicated

of the models varied for each compound and for the two spectral acquisition ways (ws and ds: wet and dried samples). R^2 for the calibration models ranged 0.365–0.952. Models developed from NIR spectra acquired on wet samples (ws) present high R^2 for N (ww $R^2=0.915$, dw $R^2=0.904$) and C (ww $R^2=0.941$), moderate for C (dw $R^2=0.763$) and DM ($R^2=0.773$), and poor for MF (ww $R^2=0.642$, dw $R^2=0.365$) (Table 1). Regarding the models using NIR spectra acquired on dried samples (ds) it is shown that these were very similar for N (ww $R^2=0.952$,

dw $R^2=0.895$) and C (ww $R^2=0.912$, dw $R^2=0.865$), but differed widely for MF, where models developed from dry samples had better quality (ww $R^2=0.852$, dw $R^2=0.636$).

In general, prediction results (R^2) were slightly lower but similar to the calibration ones (Table 1). RMSEP values were, for most of the variables, equivalent to the corresponding RMSEC values, indicating good robustness of the NIR models.

Fig. 2 Bivariate correlations between parameters (only significant correlations are shown, $p < 0.05$). For each correlation, the Pearson coefficient is presented. The color scale indicates the correlation degree between parameters (blue, positive correlation; red, negative correlation)



Discussion

Chemical parameters

The parameters range of variation for the samples analyzed in our study is representative of the phenotypic diversity that can be found in cultured *Ulva* spp (Msuya and Neori 2008; Al-Hafedh et al. 2012; Angell et al. 2014; Mata et al. 2016). Chemical variables DM, C, and N showed non-normal distributions, with a uniform distribution of the reference values along the working ranges, which is favourable for the development of NIR models, according to Williams (2001). With regard to the correlations among variables, we identified significant and positive correlations between all the chemical variables expressed on a ww basis, whereas on a dw basis we identified a negative correlation between the inorganic (MF) and organic (C, N) fraction of the DM. The positive correlation between DM and C and N on a dry and wet weight basis is due to the fact that C is the main constituent of DM (C represents 27.2–35.8% dw) and N is an important element in the synthesis of biomass, despite being in a lower concentration (N represents 3.2–4.7% dw) (Fig. 1). The negative correlation on dry weight basis between MF and N ($r = -0.42$) and C ($r = -0.51$) indicates that the inorganic fraction of

the DM (MF) diminishes in samples with a high organic fraction (N, C).

NIR spectra versus chemical composition

The broad bands observed around 5000 and 7000 cm^{-1} in the raw spectra of wet samples (Fig. 3) are due to the combination bands and 1st overtone of O–H bonds vibration in water molecules. These bands were expected since the water has a strong contribution in NIR spectra and was the major constituent of wet samples, ranging its content between 80.9–89.8% (Fig. 1).

NIR models developed from spectra acquired on wet samples had good accuracy for the prediction of N (on a ww and dw basis) and C (on a ww basis), and thus can be used to characterize *Ulva ohnoi* samples directly after harvesting. Models with lower R^2 scores have been obtained for DM and MF. The low R^2 scores for the MF may be due to the lack of discriminating spectral bands specific for this parameter. Overall, our results show that NIR can be used to predict N (on a ww and dw basis) and C (on a ww basis) contents with good accuracy ($R^2 > 0.9$) from recently harvested seaweed samples (without requiring any treatment of the sample) (Fig. 4a–c), and to estimate roughly DM and C (on a dw basis) contents ($R^2 > 0.75$) (Fig. 4d–e).

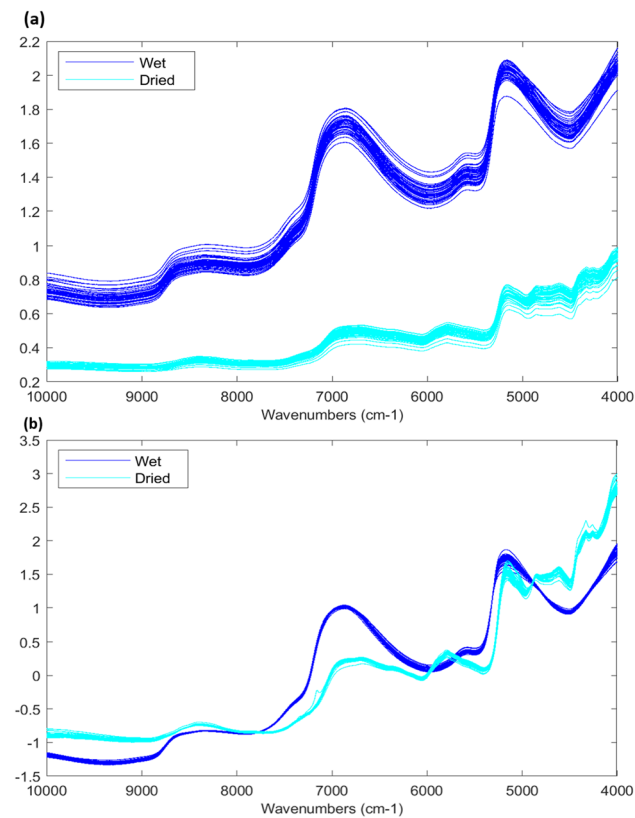


Fig. 3 NIR reflectance spectra of seaweed samples: (a) raw spectra from wet and dried samples; (b) SNV preprocessed spectra from wet and dried samples. Y axis represents arbitrary units

To the author's knowledge, this is the first full report on the efficacy of NIR to characterize the chemical composition of *U. ohnoi*. Previous works on brown algae yielded similar results to those reported in this study (Hay et al 2010; Campbell et al 2022; Tadmor et al. 2022). For instance, Hay et al. (2010) reported similar R^2 values for predicting C and N on *Sargassum flavicans*, and also observed a better fitness of the NIR models to predict N than C. More recently Campbell et al. (2022), working with 4 different brown seaweed species, also reported accurate models to predict the chemical composition, obtaining higher R^2 values than those obtained in our work, specifically for the MF. The preparation of the samples and the higher range of variation for MF studied by Campbell et al. (2022) can be the reason for this difference. These two studies used freeze-dried and lyophilized samples to develop the models, thus avoiding the signal noise impaired by water in the NIR spectra.

Conclusions

As one of the main benefits of NIR technology is the rapid characterization of samples with minimal sample processing time, the present research aimed to evaluate the potential of NIR applied either on wet and dried samples of the macroalgae *Ulva ohnoi*.

The effectiveness of NIR to predict dry matter, Nitrogen, Carbon and Mineral fraction in *Ulva ohnoi* samples have been explored, either directly on wet (recently harvested) or on dried samples.

Table 1 Calibration and prediction statistics obtained for the models developed with wet samples (ws) and dried samples (ds). All variables, as well as RMSEC/P, are expressed in %

Spectral acquisition	Variable	Pretreatment	LVs	Outliers	N	Calibration		Prediction				
						R^2	RMSEC	SDref	R^2	RMSEP	SEP	RPD
Wet samples (ws)	Dry matter	SNV + SG-1D	5	0	52	0.77	1.04	2.07	0.61	1.28	1.29	1.60
	Nitrogen (ww)	SG-1D	5	1	26	0.92	0.05	0.13	0.93	0.04	0.04	2.95
	Nitrogen (dw)	SNV	4	3	23	0.90	0.12	0.29	0.81	0.16	0.16	1.78
	Carbon (ww)	SNV + SG-1D	5	0	24	0.94	0.24	0.91	0.90	0.31	0.31	2.97
	Carbon (dw)	SNV + SG-1D	3	2	22	0.69	0.87	1.15	0.60	0.88	0.90	1.28
	Mineral fraction (ww)	1D + SNV	3	2	22	0.64	0.26	0.38	0.69	0.23	0.24	1.54
Dried samples (ds)	Mineral fraction (dw)	SNV	3	3	24	0.37	1.42	1.37	0.51	0.95	0.97	1.41
	Nitrogen (ww)	SG-1D	4	0	24	0.96	0.04	0.13	0.90	0.04	0.04	3.09
	Nitrogen (dw)	SNV + SG-1D	4	0	26	0.90	0.13	0.35	0.72	0.02	0.21	1.68
	Carbon (ww)	SG-1D	4	1	24	0.91	0.27	0.97	0.93	0.26	0.25	3.84
	Carbon (dw)	SG-1D + SNV	4	2	24	0.87	0.59	1.10	0.70	0.67	0.62	1.78
	Mineral fraction (ww)	1D	4	2	22	0.85	0.19	0.30	0.69	0.23	0.22	1.36
	Mineral fraction (dw)	SNV	3	2	27	0.64	1.10	1.31	0.54	1.37	1.06	1.24

SNV: standard normal variate; SG-1D: Savitzky-Golay first-order derivative; LVs: number of latent variables; N: number of samples; R^2 coefficient of determination of calibration and prediction sets; RMSEC: root mean square error of calibration; SDref: Standard Deviation of reference values; RMSEP: root mean square error of prediction; SEP: Standard Error of Prediction; RPD: ratio SD/SEP

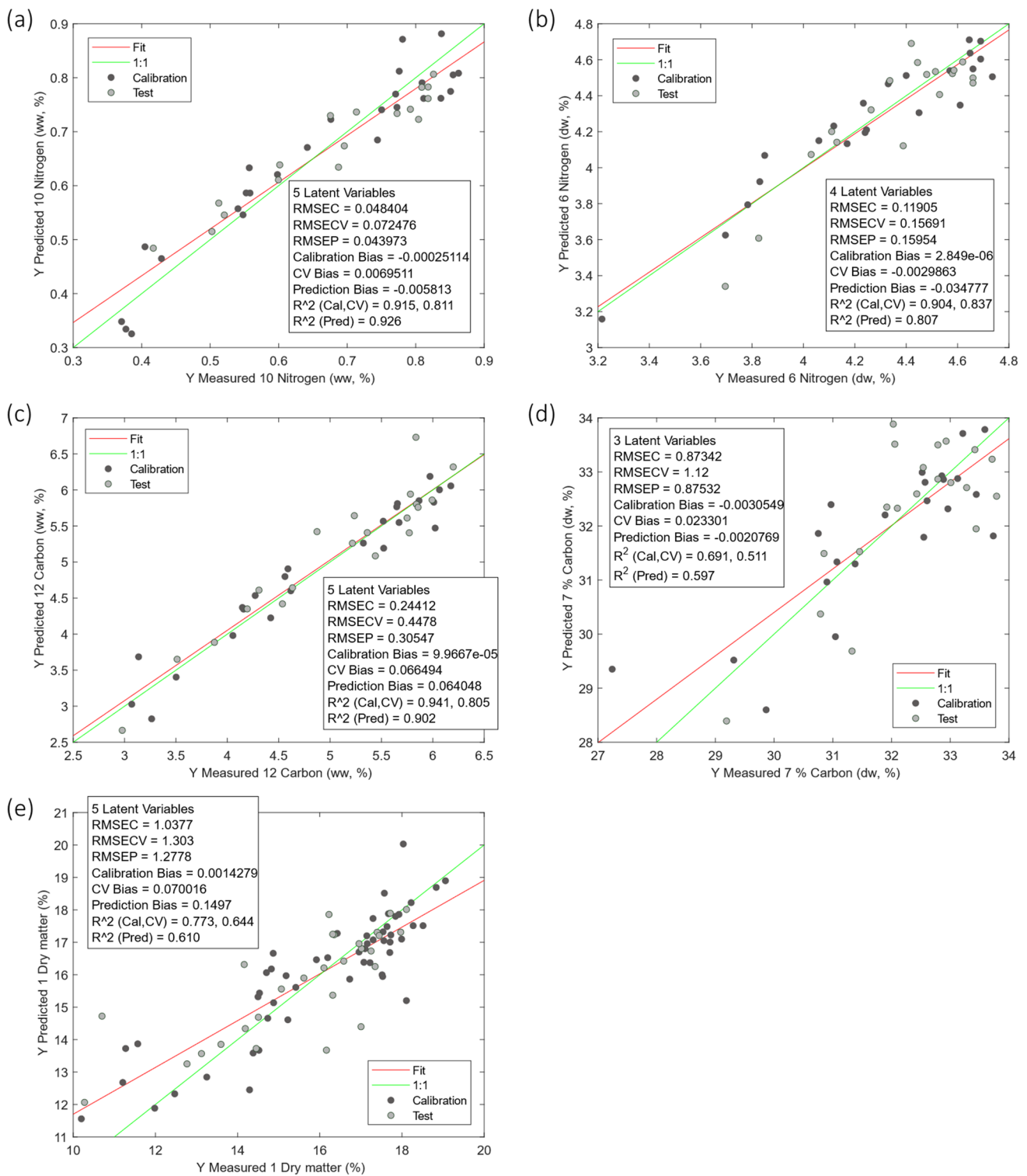


Fig. 4 Reference-predicted scatterplots for the chemical variables estimated with NIR in the wet samples. In each graph, the calibration (black dots) and validation (grey dots) sets are shown, as well as the

main statistics of the models. Panels: (a) Nitrogen (ww), (b) Nitrogen (dw), (c) Carbon (ww), (d) Carbon (dw), (e) Dry matter

Results show that NIR models to predict nitrogen and carbon on a wet weight basis directly on wet samples had similar robustness than on dried samples ($R^2 > 0.9$), while lower accuracies were obtained for MF and dry matter. Obtaining these parameters (carbon and nitrogen) in wet samples allows making rapid determinations, avoiding the processing of the samples needed to obtain the grounded dry matter.

Rapid and low-cost phenotyping of the chemical composition of *Ulva ohnoi* is a crucial step towards the quality monitoring of wild and/or cultivated algae and the decision of optimal harvesting time.

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Joan Casals: Conceptualization, Writing—original draft, Writing—review & editing.

Ingrid Masaló: Conceptualization, Writing—original draft, Writing—review & editing, Funding acquisition.

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Availability of data and material The authors declare that the data supporting the findings of this study are available within the article.

Code availability PLS Toolbox v8.9.1 (Eigenvector Research, Inc.) used through Matlab R2020b software (The MathWorks, Inc.).

Declarations

Conflicts of Interest The authors declare no conflict of interest.

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References

- Al-Hafedh YS, Alam A, Buschmann AH, Fitzsimmons KM (2012) Experiments on integrated aquaculture system (seaweeds and marine fish) on the Red Sea coast of Saudi Arabia: efficiency comparison of two local seaweed species for nutrient biofiltration and production. *Rev Aquacult* 4:21–31
- Angell AR, Mata L, de Nys R, Paul NA (2014) Variation in amino acid content and its relationship to nitrogen content and growth rate in *Ulva ohnoi* (Chlorophyta). *J Phycol* 50:216–226
- Barrington K, Chopin T, Robinson S (2009) Integrated multi-trophic aquaculture (IMTA) in marine temperate waters. In: Soto D (ed) *Integrated mariculture: a global review*. FAO Fisheries and Aquaculture Technical Paper. no. 529, pp. 7–46. Food and Agriculture Organization of the United Nations, Rome
- Beć KB, Grabska J, Huck CW (2022) Miniaturized NIR spectroscopy in food analysis and quality control: Promises, challenges, and perspectives. *Foods* 11:1465
- Campbell M, Ortuño J, Koidis A, Theodoridou K (2022) The use of near-infrared and mid-infrared spectroscopy to rapidly measure the nutrient composition and the in vitro rumen dry matter digestibility of brown seaweeds. *Anim Feed Sci Technol* 285:115239
- Cao X, Ding H, Yang L, Huang J, Zeng L, Tong H, Su L, Ji X, Wu M, Yang Y (2022) Near-infrared spectroscopy as a tool to assist *Sargassum fusiforme* quality grading: Harvest time discrimination and polyphenol prediction. *Postharvest Biol Technol* 192:112030
- Gomez-Zavaglia A, Prieto Lage MA, Jimenez-Lopez C, Mejuto JC, Simal-Gandara J (2019) The potential of seaweeds as a source of functional ingredients of prebiotic and antioxidant value. *Antioxidants (basel)* 8:406
- Harrison PJ, Hurd CL (2001) Nutrient physiology of seaweeds: application of concepts to aquaculture. *Cah Biol Mar* 42:71–82
- Hay KB, Millers KA, Poore AG, Lovelock E (2010) The use of near infrared reflectance spectrometry for characterization of brown algal tissue. *J Phycol* 46:937–946
- Hayden HS, Blomster J, Maggs CA, Silva PC, Stanhope MJ, Waaland J (2003) Linnaeus was right all along: *Ulva* and *Enteromorpha* are not distinct genera. *Eur J Phycol* 38:277–294
- Holdt SL, Kraan S (2011) Bioactive compounds in seaweed: functional food applications and legislation. *J Appl Phycol* 23:543–597
- Horn SJ, Moen E, Østgaard K (1999) Direct determination of alginate content in brown algae by near infra-red (NIR) spectroscopy. *J Appl Phycol* 11:9–13
- Lahaye M, Jegou D (1993) Chemical and physical-chemical characteristics of dietary fibres from *Ulva lactuca* (L) Thuret and *Enteromorpha compressa* (L.) Grev. *J Appl Phycol* 5:195–200
- Lawton RJ, Mata L, de Nys R, Paul NA (2013) Algal bioremediation of waste waters from land-based aquaculture using *Ulva*: Selecting target species and strains. *PLoS ONE* 8:e77344
- Manhart J (1994) Phylogenetic analysis of green plant rbcL sequences. *Mol Phylogenet Evol* 3:114–127
- Manley M, Williams PJ (2021) Applications: Food Science. In: Ozaki Y, Huck C, Tsuchikawa S, Engelsen SB (eds) *Near-infrared spectroscopy: Theory, spectral analysis, instrumentation, and applications*. Springer, Singapore, pp 347–359
- Mata L, Magnusson M, Paul NA, de Nys R (2016) The intensive land-based production of the green seaweeds *Derbesia tenuissima* and *Ulva ohnoi*: biomass and bioproducts. *J Appl Phycol* 28:365–375
- Msuya FE, Neori A (2008) Effect of water aeration and nutrient load level on biomass yield, N uptake and protein content of the seaweed *Ulva lactuca* cultured in seawater tanks. *J App Phycol* 20:1021–1031
- Neori A, Cohen I, Gordin H (1991) *Ulva lactuca* biofilter for marine fishpond effluents: II. Growth rate, yield and C: N ratio. *Bot Mar* 34:389–398
- Oca J, Cremades J, Jiménez P, Pintado J, Masaló I (2019) Culture of the seaweed *Ulva ohnoi* integrated in a *Solea senegalensis* recirculating system: influence of light and biomass stocking density on macroalgae productivity. *J Appl Phycol* 31:2461–2467
- Pedersen MF, Borum J, Fotel FL (2010) Phosphorus dynamics and limitation of fast- and slow-growing temperate seaweeds in Oslofjord, Norway. *Mar Ecol Prog Ser* 399:103–115

- Peña-Rodríguez A, Mawhinney TP, Ricque-Marie D, Cruz-Suárez LE (2011) Chemical composition of cultivated seaweed *Ulva clathrata* (Roth) C. Agardh Food Chem 129:491–498
- Pereira L (2011) A Review of the nutrient composition of selected edible seaweeds. In: Pomin VH (ed) Seaweed: Ecology, Nutrient Composition and Medicinal Uses. Nova Science Publishers, Coimbra, pp 15–47
- Pereira DC, Trigueiro TG, Colepicolo P, Marinho-Soriano E (2012) Seasonal changes in the pigment composition of natural population of *Gracilaria domingensis* (Gracilariales, Rhodophyta). Rev Bras Farmacogn 22:874–880
- Rasyid A (2017) Evaluation of nutritional composition of the dried seaweed *Ulva lactuca* from Pameungpeuk waters, Indonesia. Trop Life Sci Res 28:119–125
- Tadmor NS, Ghermandi A, Tchernov D, Shemesh E, Israel A, Brook A (2022) NIR spectroscopy and artificial neural network for seaweed protein content assessment in-situ. Comput Electron Agr 201:107304
- Toth GB, Harrysson H, Wahlström N, Olsson J, Oerbekke A, Steinhagen S, Kinnby A, White J, Albers E, Edlund U, Undeland I, Pavia H (2020) Effects of irradiance, temperature, nutrients, and pCO₂ on the growth and biochemical composition of cultivated *Ulva fenestrata*. J Appl Phycol 32:3243–3254
- Vasconcelos MMM, Marson GV, Turgeon SL, Tamigneaux E, Beaulieu L (2022) Environmental conditions influence on the physico-chemical properties of wild and cultivated *Palmaria palmata* in the Canadian Atlantic shore. J Appl Phycol 34:2565–2578
- Williams PC (2001) Implementation of near-Infrared technology In: P.C. Williams, K.H. Norris (eds) Near-Infrared Technology in the Agricultural and Food Industries. American Association of Cereal Chemists. Minnesota pp 145–171.
- Yang Y, Tong H, Yang L, Wu M (2021) Application of near-infrared spectroscopy and chemometrics for the rapid quality assessment of *Sargassum fusiforme*. Postharvest Biol Technol 173:111431

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