Improving Calibration Strategy for LIBS Heavy Metals Analysis in Agriculture Applications

Vasily N. Lednev 1,*, Pavel A. Sdvizhenskii 1 ○, Mikhail Y. Grishin 1 ○, Evgeny A. Nikitin 2 ○, Sergey V. Gudkov 1 ○ and Sergey M. Pershin 1

1 Prokhorov General Physics Institute, Russian Academy of Sciences, 38 Vavilova Street, 119991 Moscow, Russia; s dvizhenskii.pa@misis.ru (P.A.S.); mikhail.grishin@kapella.gpi.ru (M.Y.G.); S_makary@rambler.ru (S.V.G.); pershin@kapella.gpi.ru (S.M.P.)
2 FSBSI “Federal Scientific Agroengineering Center VIM”, 1st Institutsky Proezd 5, 109428 Moscow, Russia; evgeniy.nicks@yandex.ru
* Correspondence: lednev@kapella.gpi.ru; Tel.: +74-(99)-5038777876

Abstract: A new calibration procedure, known as mapping conditional-calibration laser-induced breakdown spectroscopy (LIBS), has been suggested to improve analysis results for heterogeneous samples. The procedure is based on LIBS elemental mapping, followed by signal conditioning in every sampling spot to skip signal outliers, then by finalizing the calibration curve construction. The suggested mapping conditional calibration procedure was verified for zinc analysis in soybean grist samples. The laser parameters correspond to those of the hand-held LIBS instrument in order to estimate the influence of sample surface heterogeneity under on-site analysis conditions. The laser spot (60 µm) was equal to or smaller than the typical size of grist particles (40–500 µm) but laser crater dimensions were significantly greater and varied widely (150–450 µm). The LIBS mapping of different spectral signals (atomic and ionic lines for major and minor components) was achieved. Elemental maps were normalized to achieve signal maps that were conditionally spotted to skip signal outliers. It was demonstrated that the suggested mapping conditional-calibration LIBS provided 15 ppm RMSECV for zinc determination in heterogeneous samples, which is typical for agricultural products.

Keywords: laser-induced breakdown spectroscopy; express analysis; heavy metals; grist quality; cattle food diagnostics

1. Introduction

The control of concentrations of major and minor elements in agriculture has been of great interest during the last decade, due to requirements for the online management of nutrients and preventing contamination from toxic metal. The express detection of harmful elements in fresh plants, vegetables, or fruits can enable effective control in the food industry, resulting in safe and healthy food alimentation. Toxic elements and heavy metals in agricultural products can induce intoxication and even diseases, so express multi-elemental analysis techniques are required. Conventional laboratory techniques, such as inductively coupled plasma–atomic emission spectrometry (ICP-AES), inductively coupled plasma mass spectrometry (ICP-MS), graphite furnace atomic absorption spectrometry (GFAAS), flame atomic absorption spectrometry (FAAS), and X-ray fluorescence spectrometry (XRF) have been successfully adopted for food and agricultural products [1–7]. All the above-mentioned techniques require laboratory conditions, are time-consuming, and need sample preparation. However, high-quality agricultural production needs instruments that can provide onsite or inline control of heavy elements in plant or food materials, for example, in cattle.

Laser-induced breakdown spectroscopy (LIBS) is of growing interest in agriculture and food production in recent years, due to its capability of express multi-elemental analysis without any sample preparation [8–20]. LIBS was successfully utilized for the analysis.
of different food products like wheat [21,22], vegetables [23,24], milk powder [25–28], bread [29], oil [30] and wine [31]. For example, Beldjilali et al. [24,32] analyzed fresh potato flesh and skin without any sample preparation. They achieved good agreement between LIBS results and the values measured by conventional techniques (ICP-AES, ICP-MS and AAS) for both major and minor elements (Al, Mn, Ti, Li, and Si). Later, the same team suggested a two-step procedure for trace element analysis in food utilizing calibration-free LIBS [33].

Three different approaches are used in LIBS to quantify the elemental composition of organic materials [11,32,34]: (a) single or multiple variable calibrations with the standard samples; (b) calibration-free LIBS; (c) chemometrics. Calibration-free LIBS can perform analysis without any reference materials, but such an approach can provide reliable results only for major components, while errors of determination for additives are too large for meaningful analysis [35]. Chemometrics methods became very popular in LIBS within the last decade for both identification and quantitative analysis [36,37]. However, conventional calibration curve construction by reference materials is also widely used, due to its simplicity and good analytical performance [11].

Food and plant samples demonstrate complex composition and the heterogeneous distribution of elements, so LIBS calibration curve construction will be affected by large measurement errors. Because of the samples’ complex structure, every food item gives a different response to laser ablation due to differences in absorption coefficients, thermal diffusivity, oil/water concentration, etc. In order to improve LIBS measurements’ reproducibility and to achieve a flat surface on the sample, food or plant materials are pressed into tablets [11,19,38,39]. The granular sizes in plant material varied from a few microns up to a few millimeters, while an LIBS sampling spot is typically in the 10–200 µm range. In order to improve analysis reproducibility, different spots are ablated and the LIBS spectra are quantified and averaged across the sampling area. Typically, internal normalization on the matrix component is utilized to eliminate any unwanted shot-to-shot LIBS signal fluctuations. In such a case, LIBS spectra averaging is a poor choice since the variation of plasma properties caused by the heterogeneity of elements is diminished, so calibration curves will provide poorer analysis results compared to those samples with a homogeneous composition.

In the present paper, we suggest an alternative way of LIBS calibration to account for sample surface heterogeneity. The sampling procedure is based on LIBS elemental mapping and signals leveling in order to avoid large fluctuations of plasma properties during pulse-to-pulse measurements, so the calibration curve is constructed with reproducible spectra. Owing to the rapid progress in developing commercial hand-held LIBS analyzers [16,40], it would be most appropriate to test our mapping sampling strategy for the analysis of agriculture samples with such portable devices. Soybean grist samples were analyzed by LIBS, since such material is of high importance in the cattle industry and heavy metal concentrations must be controlled to assure high-quality milk products. Cattle for high-quality dairy products need a diet with high fat or protein contents, such as sunflower meal, soybeans, rapeseed, etc., which are the processed products of oilseed processing. In order to increase production efficiency, oil producers will utilize chemical extraction so the processed products (grist) may be contaminated by harmful substances. For this reason, the onsite/inline analysis of cattle food is of high importance.

2. Experimental Technique

2.1. Laser-Induced Breakdown Spectroscopy Setup

The experiments were carried out at a LIBS setup previously described in detail in [41] and will be described in this study hereafter (Figure 1). Laser parameters were set close to those for hand-held LIBS analyzers (1064 nm, 10 ns, 5 mJ, 5 Hz, TEM 00, 3 mm) in order to achieve the same conditions (power density, sampling spot size). The laser beam was focused by a quartz lens (F = 110 mm) placed orthogonally to the sample surface on the spot of 60 µm. Plasma emission was collected by a quartz lens (F = 60 mm) placed at two focal
lengths, then the plasma image was transferred at 1:1 magnification to the spectrograph input slit. The spectrograph (Shamrock 303i, Andor, Belfast, UK) was equipped with a gated camera (iStar, Andor, UK) to capture the plasma spectra. The sample holder was installed on a two-dimensional motorized stage (8MT173-30, Standa Ltd., Vilnius, Lithuania) that can be moved in a 25 mm range, with 0.2 \( \mu \text{m} \) position precision. Control and synchronization of the motorized stage, laser, and spectrometer were performed via custom-written software developed in the LabVIEW environment. Spectral data processing was also performed via custom-made LabVIEW programs. Here, we utilized a laser with parameters similar to those of hand-held LIBS systems.

Figure 1. Experiment setup scheme for grist tablet analysis by laser-induced breakdown spectroscopy (LIBS). ICCD—intensified charge-coupled device camera, PC—personal computer.

2.2. Samples

Commercially available soybean grist was purchased from Rusagro LLC (Tambov, Russia). The grist was milled in a milling machine (JSM 100, Herzeg, Nattheim, Germany) for 5 min and then the material was pressed into pellets under a load of 150 kN (Retsch PP25, Haan, Germany). A photograph of the grist pellets can be seen in Figure 2. After creating the pellets, they were analyzed with an X-ray fluorescence spectrometer (ARL 9900, Thermo Fisher Scientific, New York, NY, USA) [42], which revealed a low concentration of Zn (Table 1).

Figure 2. Soybean grist samples—color photo (a), milled and pressed soybean tablet photo (b), and microscopy image of the tablet surface (c).

Table 1. Soybean grist elemental composition (wt. %) as revealed by X-ray fluorescence spectrometry (ARL 9900, Thermo Fisher Scientific).

| Sample         | K   | Ca   | Mg   | Zn   | Na   | Pb     |
|----------------|-----|------|------|------|------|--------|
| sample1        | 1.67| 1.0500| 0.0849| 0.0645| 0.0058| 0.00487|
| sample2        | 1.70| 1.0600| 0.0812| 0.0395| 0.0057| 0.00260|
| sample3        | 1.69| 1.0600| 0.0820| 0.0520| 0.0051| 0.00387|
| sample4        | 1.68| 1.0300| 0.0822| 0.0270| 0.0047| 0.00103|
| sample5        | 1.72| 1.0400| 0.0792| 0.0195| 0.0049| 0.0061 |
| sample6 (raw material) | 1.71| 1.0500| 0.0826| 0.0170| 0.0059| <0.0002|
In order to calibrate the LIBS setup and estimate its analytical capabilities, a series of samples were prepared. The raw grist powder was dried in air heated to 99 °C for 1 h. The required amount of zinc and lead solutions at known concentrations had been added to the grist powder and then dried in air at 99 °C for 1 h. Different ratios of raw and “contaminated” grist powders were mixed to get 5 different samples. In order to obtain the uniform distribution of Zn and Pb additives, the prepared mixture was blended in an agate mortar for 30 min; it was then dried at 99 °C for 1 h to remove any non-bonded water. The obtained powders were pressed into tablets (40 mm diameter, 5 mm thickness) in air for 5 min under a load of 150 kN (Retsch PP25). The prepared tablets were analyzed with an X-ray fluorescence spectrometer (ARL 9900, Thermo Fisher Scientific). The tablet surface was examined using optical microscopy (Figure 2) to estimate sample non-homogeneity. The samples were fairly heterogeneous and different color particles had dimensions from 20 to 300 µm; this was equal to or smaller than the 60 µm diameter of the laser spot.

3. Results

The express elemental analysis of agriculture samples is necessary for modern farming, and hand-held LIBS systems can provide a good solution [16]. Such instruments are based on low-pulse-energy nanosecond lasers (1–10 mJ/pulse) that produce a small sampling spot of 30–100 µm in diameter. Typically, several spots are ablated to improve the sample's representative analysis, but the LIBS spectrum is averaged during this measurement. In this study, we reproduced the same conditions for laser ablation and studied in detail the analytical capabilities of LIBS for heavy metals analysis.

The sample was mapped by the LIBS system in the following way: the sample was moved, stopped, and then ablated using three laser pulses. Afterward, the translation stage was further moved to reveal a fresh sample surface for the next LIBS measurement. A series of 900 spots (30 × 30-spot grid) was ablated. The laser crater dimensions (diameter, depth) and shapes varied widely for different spots, as can be clearly observed by optical microscopy (Figure 3). Large variations in crater dimensions can be explained by the variation of particle absorption coefficients, as well as variations in particle size and bonding at nearby laser spots and nearby areas. For particles with a greater absorption coefficient (dark particles), the ablation efficiency will be greater; thus, plasma temperature and density will be greater, resulting in a higher intensity of lines in the spectrum. The expanding plasma plume will induce a shockwave, followed by a sound wave that will blow away powder particles, so the laser crater diameter will be much greater compared to that for a metal target [43]. The crater diameters varied from 150 to 450 µm, while the laser spot was only 60 µm in diameter.

Figure 3. Photograph of the soybean grist tablet surface after LIBS mapping (a), and a more detailed view of the craters (b, c), single laser crater photo (d), and laser beam spot (e).
of major elements or molecular bands; for this reason, we chose the spectral region at 200–220 nm, where resonant ionic and atomic lines are present: Zn II at 202.55, Zn II at 206.20, and Zn I at 213.86 nm. Additionally, calcium Ca II 210.32, Ca II 211.28, and phosphorus P I 213.62, P I 214.91 lines were observed in the spectrum. The most intensive zinc line Zn II 213.85 nm cannot be used as a signal (spectral line integral corrected for background) due to spectral interference with the P I 213.62 nm line. The evolution of laser-induced plasma spectra has been studied to reveal the optimal conditions for spectra acquisition. Plasma emissivity decreased rapidly within the first five microseconds (Figure 4), so the early moments of plasma emission were optimal for LIBS spectra acquisition (gate 1 us, delay 0.2 us) due to a better signal-to-noise ratio for Zn lines.

Figure 4. Laser plasma spectra at different moments in plasma evolution (a), the zinc (Zn II) 206.55 nm line and signal, and the signal-to-noise ratio at different moments of LIBS plasma expansion (b).

In order to provide representative LIBS analysis for a nonhomogeneous target, the sampling area has to be significantly greater compared to the typical particle size. To optimize the time required for a single LIBS measurement and to estimate the minimum sampling area for representative analysis, we have mapped the soybean grist tablets and have constructed maps for major elements (calcium, phosphorus) and zinc (Figure 5). The zinc distribution was fairly uniform, proving proper sample preparation (see Section 2.2); however, some spots with a higher signal can be traced in the map. The same distribution was observed for calcium and phosphorus. The continuous plasma emission (background) was also quantified and mapped since this signal can be used for analytical line normalization [44]. The corresponding maps of Zn II 206.55 nm, normalized on calcium, phosphorus and background signals, are presented in Figure 6.

Figure 5. Sample surface area photo after LIBS mapping (a) and corresponding atomic/ionic lines intensity maps for Zn II 206.55 (b), Ca II 211.28 (c), P I 214.91 (d) nm lines, and the plasma emission background (e).
In order to estimate the minimal sampling area needed for representative analysis, we have totaled the points in Figure 6a–c to achieve squared areas of different dimensions. Normalized signals had poor reproducibility but, after averaging over 300–400 pulses, the signal stabilized (Figure 6d). A better-normalized signal convergence was determined when normalizing on Ca II 211.28 nm, so it was convenient to use the Zn II 206.55 nm/Ca II 211.28 nm ratio for the construction of the calibration curve. To study this in detail, we have drawn normal box graphs for the Zn II 206.55 nm line intensity, as well as the signal normalized by phosphor, calcium lines and background level intensity. In order to compare the distribution function of normalized signals, we have also normalized such data by median normalization. According to Figure 7, the narrowest distribution was achieved when the normalization on the Ca II 211.28 nm line was chosen. Additionally, all normalized signal values beyond 2*sigma were considered as outliers, so data were conditioned before the calibration curve construction step.

Calibration curves for zinc with a different number of averaging spots have been constructed and are presented in Figure 8a. The calibration curves were fitted by linear functions. For viewability, only a few of the calibration curves are shown. LIBS analytical capabilities were compared in terms of precision, calibration curve linearity and analytical accuracy. The accuracy of the analysis was estimated by the leave-one-out cross-validation procedure, with the root mean square error of cross-validation (RMSECV) as the main index.
estimating the calibration model performance: 

$$\text{RMSECV} = \left( \frac{1}{n} \sum_{i=1}^{n} (c_i - c_{\text{ref}})^2 \right)^{1/2},$$

where $c_i$—predicted concentration, $c_{\text{ref}}$—reference concentration, and $n$—number of calibration samples.

For small-sized sampling areas (up to 225 points), the calibration curve cannot be established for meaningful LIBS analysis (Figure 8a): RMSECV values were in the 50–200 ppm concentration range, which was comparable to the concentration in the reference samples. Starting from 200–300 LIBS sampling spots, a good calibration curve can be achieved; however, additional data points in the calibration will further improve the analysis results. The coefficient of determination ($R^2$ factor) was systematically poorer for low-spots averaging. For 250-spot sampling (15 × 15 spots square), the $R^2$ reached 0.936, so meaningful calibration can be established, but 900 spot-averaging provided a value of $R^2 = 0.996$. The RMSECV value (Figure 8b) showed the same tendency, and after averaging for 300 spots, the accuracy of analysis was in the 15–20 ppm range. It should be noted that hand-held LIBS analyzers should be used with caution for heavy metals analysis in the heterogeneous samples (like grist) since such devices can only automatically map small areas (max spot of 8 × 8) and thus could not provide a representative sampling.

4. Conclusions

Here, we have presented a new approach for the analysis of heterogeneous samples by LIBS—mapping conditional calibration. The procedure is based on the LIBS elemental mapping of large areas, followed by signal conditioning to construct calibration curves. The suggested mapping conditional calibration procedure will be of great interest for the onsite analysis of heterogeneous samples, like the soybean grist pellets utilized in agriculture for cattle nutrition. Hand-held LIBS instruments are widely employed in farming due to their simplicity and the possibility of express analysis, so mapping conditional calibration LIBS was verified using the laser-sampling parameters that are typical for such compact devices (a laser spot diameter of 60 µm was equal to or smaller than the typical size of grist particles, 40–500 µm). Elemental maps for major components were obtained and normalized, while the signal was conditionally chosen to skip the signal outliers. It was demonstrated for zinc analysis in soybean grist that the suggested mapping conditional calibration LIBS provided an accuracy of determination of 15 ppm. Furthermore, employing LIBS with the mapping conditional calibration procedure will improve the results of an analysis in the case of homogeneous samples, since pulse-to-pulse plasma fluctuations can be traced.
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