

Quo Vadis high-resolution continuum source atomic/molecular absorption spectrometry?

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Abstract

After more than a decade since its commercial introduction, high-resolution continuum source atomic/molecular absorption spectrometry maybe facing a mid-life crisis. Certainly, it is no longer a novel technique full of unknown potential, so it would be already time to establish the fields for which it is most suitable. This is, however, not so simple for a number of reasons. In the first place, more than a technique what we are discussing herein is a type of instrumentation with the potential to use two different techniques (atomic or molecular absorption), making it somewhat unique. And both techniques have not been explored equally, as more research on the mechanisms of formation of diatomic molecules is clearly needed. In the second place, new possibilities have recently appeared in the literature that need to be weighed as well. And there is the still unfulfilled, but nowadays more technically feasible than ever, promise to significantly increase the multi-elemental capabilities.

This review critically examines the main research areas currently explored (namely, i) direct analysis of solids and complex liquid materials, and ii) determination of non-metals at trace levels *via* monitoring of molecular species)

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3 as well as the new venues (specifically, i) isotopic analysis *via* monitoring of
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5 molecular species, and ii) selective detection, quantification and sizing of
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7 nanoparticles) while also considering new instrumental developments, in an
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9 attempt to properly place high-resolution continuum source atomic/molecular
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11 absorption spectrometry in the field of trace element and isotopic analysis.
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1. Introduction

There is little doubt at this point that the arrival of high-resolution continuum source atomic absorption spectrometry (HR CS AAS) instrumentation has revitalized this field,¹ to the point it cannot be considered as only AAS anymore, because the use of molecular absorption spectrometry (MAS) in conjunction with this instrumentation provides some relevant advantages as well, particularly for the determination of non-metals.²⁻⁴

It is worth mentioning that different authors explored the use of continuum source to carry out AAS measurements over the years.⁵⁻⁹ Finally, the work of Becker-Ross and co-workers¹⁰⁻¹² lead to a device that was ultimately made commercially available in 2003, although it was equipped with a flame as atomizer only. An evolution of this device, incorporating also a graphite furnace, was released approximately five years later. The key components of such instruments are: (1) a high-pressure xenon short-arc lamp that can provide a high intensity in the visible and UV region, which is a significant difference with some previous models; (2) an optical system based on a double monochromator (using a prism first and an echelle grating afterwards); and (3) a linear CCD array that is used for detection.

The number of advantages brought by this type of instrument are quite significant, such that it is hard to argue that the already famous title coined by Dr. Welz “High-resolution continuum source AAS: the better way to perform atomic absorption Spectrometry” is quite accurate.¹³ These aspects have been covered in detail in several reviews,¹⁴⁻¹⁷ in addition to one book.¹ Let’s just stress that the critical difference with traditional line source AAS is the addition of one extra dimension (wavelength), that permits to actually see spectral

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3 overlaps, instead of trying to guess potential problems investigating 2D
4 (absorbance *versus* time) signals in various conditions. Moreover, it is feasible
5 to mathematically correct for some overlaps, and the high-spectral resolution
6 provided also permits to monitor molecular transitions, which are useful to
7 quantify elements for which the atomic lines are hardly accessible (e.g., non-
8 metals). Finally, the simultaneous monitoring of various lines is feasible,
9 although with significant restrictions, limiting the potential of the technique in its
10 current commercial configuration for multi-element analysis.^{18,19} This latter topic
11 will be discussed in more detail in section 4.3.

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22 At this point, HR CS AAS/MAS can no longer be considered as a novel
23 technique, but at the same time, it is still premature to consider it as fully
24 developed. In the authors' opinion, and bearing in mind that no general reviews
25 on this topic have been published since 2014,²⁰ it seems timely to review the
26 work carried out recently, discussing major research subjects, but particularly
27 trying to identify emerging trends and those subjects that deserve further
28 studies.

2. Analysis of the literature

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Figure 1 shows the evolution of publications dealing with high-resolution continuum source atomic or molecular absorption spectrometry as a function of different parameters. In general, it can be seen that the trend goes upwards. The decrease observed for 2018 simply obeys to the fact that only 8 months are accounted for such year at the time of writing this review.

If we focus on the type of atomizer, graphite furnace (GF) is the preferred one (selected in approximately 70% of publications), followed by flame (24%), as shown in **Figure 1a**. Use of other atomizers (quartz, tungsten) is only residual.

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3 This could be anticipated, since graphite furnace and flame are the only
4 commercially available atomizers, and graphite furnace offers superior potential
5 in terms of sensitivity and capability to analyze samples directly when compared
6 with flame. The latter is an important factor, as roughly 30% of the articles are
7 devoted to direct analysis of solid or of complex liquid samples. The number of
8 articles using graphite furnace still keep on growing, while the number of those
9 using flame seems to have reached a plateau over the last 4 years.

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12 Regarding the measurement technique (see **Figure 1b**), AAS articles are still
13 predominant. Use of MAS, with its possibilities to determine non-metals is,
14 however, investigated in approximately 27% of the publications up to now. This
15 type of research is still relatively new and the number of research groups
16 exploring it is still limited.

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19 Finally, an indication of the maturity of the technique is that, during the last
20 decade, a significant number of papers have appeared in more applied
21 Journals, to the point that the articles published in these Journals already
22 represent 15-20% of the total number of HR CS AAS/MAS publications. While
23 the term “applied” is always subjective, we are referring to Journals devoted to
24 food, clinical and environmental analysis. In particular, the number of articles
25 appearing in Journals devoted to food analysis is noteworthy. This number of
26 applied publications will presumably continue increasing.

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29 Overall, the field still shows potential to grow and has demonstrated possibilities
30 to be applied to various fields.

30 31 32 **3. Major research areas**

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35 This section will discuss the current two major areas of research, according to
36 the number of articles published, which are the development of methods that

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3 enable direct analysis of the samples (section 3.1.), and the development of
4 methods for the determination of non-metals (section 3.2.). Those papers of
5 higher novelty, in terms of methodology, will be highlighted. **Tables 1** and **2** will
6 present the publications in both fields, respectively, starting in 2015. It also
7 needs to be mentioned that those articles dealing with direct analysis while also
8 targeting the determination of non-metals will be included only in section 3.2.
9 and in **Table 2**, in order to avoid duplications while achieving a better balance
10 for both sections. For previous publications, the reader is referred to reviews on
11 both topics published in 2014.^{4,21}

22 3.1. Direct analysis of samples

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24 This remains the most popular research area. As mentioned in the previous
25 section, approximately 30% of all the papers devoted to HR CS AAS/MAS
26 focused on this topic. The percentage would be higher (roughly 40%) if only
27 graphite furnace articles are accounted for, since, obviously, flame is not as
28 suitable for direct analysis (although it also shows some potential: one work by
29 Leite *et al.*²² demonstrated that it is possible to carry out the direct analysis of
30 ethanol fuel, determining 9 analytes).

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32 It has to be stated that by direct analysis we are not referring to solid sampling
33 only, but also to those other papers in which generally complex liquid samples
34 can be analyzed directly. In fact, in such cases, the procedure used is
35 sometimes very similar to that of solid sampling (e.g., deposition of the sample
36 onto the external platform). However, in practice, almost 90% of the articles are
37 actually devoted to the analysis of solid materials.

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39 It certainly makes sense that solid sampling HR CS GFAAS/MAS is a major
40 area of research because this technique is particularly suitable for such an
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3 approach. This was, in fact, the topic of one previous review published in JAAS
4 in 2014.²¹ The temperature control, the potential of chemical modification, and
5 the possibilities of the technique to detect and deal with spectral overlaps make
6 it possible in many occasions to analyze difficult materials and even to simply
7 use aqueous standards to construct the calibration curve (an approach that will
8 be referred to as external calibration with aqueous standards from now on). This
9 characteristic is quite unique in comparison with many of other solid techniques
10 that require solid standards for calibration.²³ One may in fact wonder if this
11 technique could become the method of choice for trace elemental analysis of
12 solid samples, letting aside imaging applications, if higher multi-element
13 capabilities would be available at some point, but such topic will be discussed in
14 section 4.3.

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An examination of the recent literature on direct analysis using HR CS GFAAS/MAS (see **Table 1**) reveals a period of maturity. From a methodological point of view, practically all that was discussed in the review mentioned before²¹ holds true. The technique has shown potential to analyze very different types of materials, from some that could be considered as relatively easy for graphite furnace decomposition, such as polymers, biological materials or pharmaceuticals, to others that are quite refractory and, also, very difficult to bring into solution quantitatively, such as automobile catalysts,²⁴ fertilizers,²⁵ high-purity silicon²⁶ or silicon carbide,²⁷ in addition to many types of environmental samples.

A significant amount of the articles presented in **Table 1** (twelve works) have reported the simultaneous determination of two analytes, or else their sequential determination, but from the same sample aliquot. Thus, this is an

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3 aspect that continues being investigated, even though the main characteristics
4 that the analytes need to fulfill and the key aspects for optimizing the analytical
5 procedure for such type of determination have been already being well-
6 established before.²⁸
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11 The most interesting and elegant of these approaches, in the authors' opinion,
12 was published by Boschetti *et al.*,²⁹ as it takes advantage of various possibilities
13 to achieve the determination of four different elements in soil samples. In that
14 paper, the determination of Al, Cr, Cd, and Fe was aimed at. These elements
15 do not show atomic lines sufficiently close to be simultaneously monitored by
16 HR CS AAS instrumentation (see section 4.3. for further discussion on this
17 topic). However, as already discussed by Resano *et al.*,¹⁸ it is often not so
18 important to determine the analytes in truly simultaneous fashion, as it may be
19 to determine them from the very same sample aliquot. Since Cd is much more
20 volatile than the other analytes, it is possible to atomize first this element at
21 1700 °C and monitor it using the main Cd line (228.802 nm), and then change
22 the wavelength, increase the temperature and vaporize/atomize the rest. There
23 is an alternative line for Cr (425.433 nm) that is suitable for the expected
24 analyte contents, and very close to it there is an Fe line (425.076 nm), also
25 appropriate for the expected sample concentrations. That is not so surprising
26 giving the abundance of Fe lines over the whole UV-vis spectrum. There is no
27 Al line close, but there are rotational lines corresponding to AlH that are found
28 in the vicinity. It has to be remembered that molecular lines can be used not
29 only to determine non-metals, but in some occasions, it may be beneficial to
30 use them for metals as well, as shown in the past using AlF to determine Al.³⁰
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3 Summing up, by using both a sequential (Cd atomization first) and later a
4 simultaneous (Cr and Fe atomization) AAS approach, combined with MAS (AIH
5 simultaneous monitoring), the authors managed to determine four elements in
6 every single aliquot of soil samples directly, using external calibration with
7 aqueous standards, which is remarkable. It is certainly not common to be able
8 to deploy simultaneously two different techniques with the same instrument, but
9 that is an existing possibility that should be explored in more HR CS AAS/MAS
10 applications.
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20 The use of an internal standard (IS) is related to this topic of simultaneous
21 monitoring. A work by De Babos *et al.*³¹ demonstrated how the use of cobalt as
22 an internal standard helps in circumventing matrix effects for the simultaneous
23 determination of Ni and Mo in plants. In fact, the use of this IS was only
24 necessary for Ni, as the sensitivity obtained for aqueous solutions and for solid
25 samples was found to be significantly different. Co was selected as IS since it
26 shows an atomic line (313.221 nm) close to those used for Ni (313.410 nm) and
27 Mo (313.259 nm) and, also, because its thermochemical properties are very
28 similar to those of Ni. The suitability of Co as IS for Ni determination by HR CS
29 GFAAS has already been demonstrated in the literature.¹⁸ This is probably the
30 first time, however, that such an approach is used to perform direct solid
31 sampling *via* HR CS GFAAS. The IS was added in solution, and its use enabled
32 accurate results to be obtained for all the plants investigated using external
33 calibration with aqueous standards.
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50 Another topic that could be mentioned is the possibility to use solid sampling
51 HR CS GFAAS after some sample preparation. Obviously, in that way, some of
52 the intrinsic advantages of direct analysis are lost, but some benefits can also
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3 be obtained. For instance, it is possible to preconcentrate the analyte in a solid
4 phase and analyze such solid phase without any dilution. One example of this
5 approach was published by Dobrowolski *et al.*,³² who reported the use of
6 modified carbon nanotubes for the solid-phase extraction of Au in a digest of
7 geological samples, with the subsequent SS HR CS GFAAS analysis, resulting
8 in an enrichment factor of 100.

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16 A similar idea but adding speciation potential to the preconcentration efficiency
17 was described by López-García *et al.*³³ They proposed a micro-solid phase
18 extraction procedure, based on magnetic nanoparticles covered with Ag
19 nanoparticles and functionalized. This magnetic character enables an easy
20 separation of the solid phase containing the analytes, which can then be directly
21 analyzed *via* SS HR CS GFAAS, or else slurried and injected into the furnace,
22 achieving enrichment factors of 325 and 205, respectively. Moreover, by
23 controlling the pH during the extraction, quantitative information on both total Sb
24 (pH=2) and of Sb (III) only (at pH=9) can be achieved, such that the amount of
25 Sb (V) can be calculated by difference. A similar speciation approach, but not
26 using solid sampling this time, was reported by the same research group to
27 determine both total Cr and Cr (III) only (and thus, Cr (VI) by differences), which
28 were retained in a graphene oxide suspension.³⁴

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44 Use of nanomaterials/nanoparticles is, therefore, also enabling some
45 improvements in HR CS AAS procedures. Another recent example was
46 reported by Aramendía *et al.*,³⁵ who proposed the use of AuNPs aiming at the
47 direct determination of Hg in blood and urine *via* HR CS GFAAS. The rationale
48 is that one of the challenges for Hg determination is the extreme volatility of Hg
49 (0), for which use of an oxidizing agent (e.g., KMnO_4) as a chemical modifier is
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3 common. However, it is well-known that Hg (0) amalgamates with Au. Thus,
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5 instead of preventing Hg (0) formation, it may be beneficial to favor it, and have
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7 already Hg in metallic state, thermally stabilized by means of this interaction
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9 with AuNPs to avoid any losses during the drying step, but ready for fast
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11 atomization at a low temperature (500 °C – 700 °C), before the bulk of the
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13 matrix is released. This approach basically eliminates the pyrolysis step,
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15 resulting in a very fast temperature program and yet an improved sensitivity,
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17 because the samples are not diluted.
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20 HR CS GFAAS can also be deployed for quantifying and even for sizing NPs,
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22 but such topic will be covered in detail in section 4.2.
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25 The last topic worth mentioning in relation with direct analysis is the use of a
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27 disposable platform made of corn starch and sorbitol, as proposed by Colares
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29 *et al.*³⁶ This platform permits the transport of the solid sample to the graphite
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31 tube, but then it decomposes during the pyrolysis step, so the final effect is
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33 similar to using wall atomization. This is beneficial for refractory elements, as it
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35 minimizes tailing, enabling atomization in shorter times, thus enhancing the
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37 lifetime of the tubes. The reduction of memory effects also improves precision.
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39 Mo and V were chosen to prove these concepts and were successfully
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41 determined in different reference materials.
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43 44 3.2. HR CS MAS for determination of non-metals

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46 An examination of the recent literature (since 2015) shows that the elements
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48 more often determined *via* HR CS MAS are F, S, and Cl, in that particular order.
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50 The articles devoted to these elements represent approximately 80% of the
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52 total number in this period. These results can be explained considering the
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3 importance of the elements, the possibilities to determine them *via* HR CS MAS,
4 and the difficulties observed for competing techniques.

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7 For instance, F and S were already the most investigated elements in previous
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9 years using HR CS MAS.⁴ This is not so surprising as F forms very stable
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11 bonds (typically the most stable ones among halogens), favoring the formation
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13 and persistence of diatomic molecules at the high temperatures used in the
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15 atomizers. Moreover, the sensitivity of these molecules is rather high and the
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17 determination of F by competing techniques is complicated (e.g., F does not get
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19 positively ionized in an inductively coupled plasma -ICP-, making the use of
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21 ICP-mass spectrometry (MS) poorly suitable to determine this element at trace
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23 levels³⁷), while HR CS GFMAS can provide LODs of a few picograms only
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25 when the most sensitive molecule (GaF) is used.³⁸⁻⁴¹ Therefore, HR CS MAS is
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27 an important tool to quantify F traces.⁴² However, the literature shows that the
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29 popularity of such molecule has decreased very much in favor of CaF, which
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31 has been selected in more than 70% of the articles published since 2015. The
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33 main reasons for this are discussed by Morés *et al.* when they first reported on
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35 the use of CaF.⁴³ Targeting CaF does not require a complex mixture of
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37 modifiers, as in fact Ca acts as both modifier and molecule forming reagent.
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39 Moreover, CaF main absorption head is found in the visible (606.440 nm) region
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41 of the spectrum (GaF is measured at 211.248 nm), where spectral interferences
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43 are rare. Thus, unless required by the analyte contents (use of CaF results in
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45 roughly two orders of magnitude lower sensitivity), most works preferred the
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47 simplicity of monitoring CaF.

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49 In the case of S, the most favored molecule continues to be CS, except in a few
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51 specific cases. Possibilities of other molecules will be discussed in section
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3 4.3.⁴⁴ It is interesting to notice that the majority of articles discussed in this
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5 section propose the use of the graphite furnace, but in the case of S
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7 determination, several articles demonstrate the benefits of using flame or quartz
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9 vaporization. This is mostly because S needs not only to be determined at the
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11 trace levels, and a cheap and fast approach for S determination at higher levels
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13 is useful, particularly for food control.

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15 Investigations about P have significantly decreased in comparison with previous
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17 years.⁴ This is however not so rare because many of the previous P papers
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19 corresponded to basic research in which the problem of the potential
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21 appearance of both atomic and molecular (PO) lines was analyzed in detail
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23 depending on the type of modifiers, conditions used and/or BG correction
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25 method. In fact, the appearance of HR CS GFAAS instrumentation served to
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27 explain many of the issues associated with P monitoring by AAS.^{45,46} Since the
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29 situation is now clear and the conditions to determine this analyte, either as P or
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31 as PO and even using direct solid sampling, are well-established,⁴⁷ a lower
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33 production of articles can be seen as normal.

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37 Cl, on the other hand, has been the subject of more research in the period
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39 covered. This fact is very welcome. Actually, one review on the determination of
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41 non-metals published in 2014 noted the surprisingly low number of papers
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43 devoted to Cl determination via HR CS MAS, despite the importance of this
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45 element.⁴ Three new molecules have been proposed in this period to
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47 accompany the ones previously explored (AlCl,³⁸ InCl,⁴⁸ and SrCl⁴⁹), namely
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49 CaCl,⁵⁰ GaCl⁵¹ and MgCl.⁵²

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52 Having different molecules to choose from for every particular application is
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54 important. Nevertheless, the increasing popularity of Ca, already noted for F
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3 and also the main choice for Br determination, is noteworthy. CaX (where X
4 represents a halogen) molecules typically offer various (but not too many)
5 transitions of different sensitivity in the visible region, making it easy to set the
6 baseline, to adjust the working range and minimizing spectral interferences.
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8 Moreover, Ca is an element present at high contents in many types of samples
9 (e.g., biological), so it could be a strong interference if any other molecule is
10 targeted instead.⁵⁰ Use of Ca also represents some disadvantages in practice,
11 as this element is well-known to deteriorate the lifetime of the graphite parts.⁴³
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13 Determination of Br and, particularly, of I are more challenging because their
14 bonds with metals are much less strong than those of F and Cl. Thus,
15 determining them in samples in which the presence of F and/or Cl is significant
16 can be problematic.⁵³ A potential solution for such a situation based on the use
17 of isotope dilution will be discussed later.^{54,55} Recently, the use of TIBr has been
18 proposed for Br determination,⁵⁶ as well as the use of Cal and Srl for I
19 determination.⁵⁷ In any case, for these two elements, more research on the use
20 of different molecules is needed.
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37 Another topic that requires further research is the way in which the different
38 modifiers and molecule forming agents interact with the analyte. It has to be
39 considered that former studies on chemical modification were mainly carried out
40 to investigate the best ways to achieve the atomization of metals. Therefore,
41 there is not so much information on situations in which the analyte is a non-
42 metal and the goal is to ultimately form a molecule in the gas phase. It is not so
43 clear, for instance, when using a classical combination of modifiers (e.g., Pd
44 and Mg), plus Ca as molecule forming agent, which element will interact with
45 the target non-metal and how, and if the targeted molecule will be formed first in
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3 solid phase, or only later on in the gas phase.⁵⁸ Supplementary research in this
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5 area may enable the development of better general strategies to determine
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7 non-metals using HR CS MAS.

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9 Finally, a significant number of articles (fifteen, representing almost 30% of the
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11 works gathered in **Table 2**) have demonstrated the potential of the technique for
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13 direct analysis of solid and complex samples (and that is not counting
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15 approaches based on slurries), also when targeting non-metals. Direct analysis
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17 may be particularly beneficial in this field because sample pretreatment is
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19 particularly problematic owing to the volatility of many species in which the non-
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21 metals are usually found, and the problems associated with contamination. In
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23 fact, the signal obtained for non-metals might be significantly affected by the
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25 way in which these are found in the sample,⁵⁹ and the way in which such
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27 samples are pretreated. This is a challenge when aiming at bulk analysis, but it
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29 can be transformed into an advantage to perform some speciation studies. Two
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31 recent articles have reported on the determination of nitrite, nitrate, and p-
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33 nitrophenol⁶⁰ and of free and total sulfur (IV)⁶¹ *via* the selective formation of
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35 volatile species after proper sample treatment and the monitoring of NO and
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37 SO₂, respectively, using HR CS MAS with a quartz vaporizer in both cases.
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41 **4. New possibilities**

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43 Aside from these major areas, there are some new and intriguing possibilities
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45 that have appeared more recently and are covered in a few papers only. These
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47 are, however, worth commenting because they may have a significant effect in
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49 the future use of the technique.
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51 **4.1. Isotopic analysis *via* molecular monitoring**

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3 It has been well-documented for years that it is, in principle, possible to use
4 atomic absorption to attain isotopic information because the atomic lines of
5 different isotopes do not appear exactly at the same wavelength. However, in
6 practice, being able to appreciate these transitions separately is very
7 challenging. The differences among isotopic lines are very small (a few
8 picometers only) and, furthermore, even if such a high-resolution is attainable, it
9 has to be considered that broadening of the peaks due to different effects (i.e.,
10 collisional broadening, Doppler and Stark effects) occurs, further complicating
11 the separate monitoring of isotopic lines.
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22 It is therefore not surprising that only one paper to date has made use of HR CS
23 AAS for isotopic analysis. In that work, Wiltsche *et al.* focused on boron
24 isotopes.⁶² This selection is not casual. For such a light element, the mass
25 difference between ^{10}B and ^{11}B atoms is quite significant, and therefore, the
26 isotopic shift should be much larger. In fact, Wiltsche *et al.* discussed that it was
27 preferable to use the less sensitive atomic line observed in the vicinity of 208.9
28 nm because the isotopic shift of such transition was higher than that observed
29 for B main line, found at approximately 249.8 nm. This loss of sensitivity in order
30 to achieve better spectral resolution is a compromise that will be necessary also
31 when using MAS, as will be discussed below, and somewhat limited the
32 applicability of the method. The goal of the work was the determination of boron
33 isotopes in steel, which could be done for samples containing at least 1% of
34 boron, or else, a preconcentration was required.
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50 To provide an idea of the small differences that we are discussing when it
51 comes to atomic isotopic shifts, the line for pure ^{10}B has been reported to be
52 found at 208.95898 nm and, for pure ^{11}B , at 208.95650 nm. In practice, it was
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3 not possible to observe two fully resolved peaks with the HR CS AAS
4 instrument, as could be anticipated for the reasons discussed above
5
6 (broadening effects). What it could be appreciated was that the peak maximum
7
8 of the boron absorption line shifted as a function of the boron isotopic ratio,
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10 moving towards higher wavelengths with higher ^{10}B contents.
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13 It has to be mentioned that the authors used a flame as the atomizer. Use of a
14
15 graphite furnace could improve sensitivity, but it has to be remembered that
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17 boron is a defying analyte for GFAAS⁶³ and, also, that the authors
18
19 simultaneously monitored either Fe or Ni lines, together with those of boron,
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21 because the use of an internal spectral standard was recommended to correct
22
23 for instabilities of the monochromator (such as wavelength drifts and changes in
24
25 the optical resolution). This truly simultaneous monitoring is easier to perform
26
27 with a flame because with a graphite furnace there would be a delay between
28
29 the atomization of Fe or Ni and the atomization of a refractory element, such as
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31 boron.
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34
35 In the end, the method proposed was deemed satisfactory as a cheap and
36
37 simple approach for the identification of steel scrap enriched in ^{10}B , accepting
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39 an uncertainty level of 5% for the ^{10}B content.
40

41 However, it has been proposed more recently that, if instead of using HR CS
42
43 AAS, HR CS MAS is chosen, isotopic analysis potential could be enhanced.⁵⁴
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45 This seems like a logical evolution for two reasons: i) MAS is an important area
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47 of research in this particular field of high-resolution continuum source, as
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49 discussed before, because diatomic molecules can be formed and their spectra
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51 resolved; ii) it has been already shown for laser induced breakdown
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53 spectrometry (LIBS) by Russo *et al.* that the monitoring of molecular spectra
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provides higher isotopic shifts.^{64,65} Therefore, the same idea could also be applied to HR CS MAS.

And such idea was demonstrated for Cl using graphite furnace as the vaporizer.

Al was chosen as the counterion in solution, in order to form the AlCl molecule in the gas phase. The selection of Al was based on several aspects: i) AlCl is a stable molecule (bond dissociation energy, 511 kJ mol⁻¹), even at high temperatures; ii) it shows a good sensitivity, as reported before;³⁸ iii) Al is monoisotopic, which is always preferable, as the shifts will thus only depend on Cl isotopes; iv) the shift observed for Al³⁵Cl and Al³⁷Cl lines is significant.

This shift can be predicted with accuracy using the first three terms of the following equation, which was proposed by Herzberg:⁶⁶

$$\Delta v = (1 - \rho) \left[\omega'_e \left(v' + \frac{1}{2} \right) - \omega''_e \left(v'' + \frac{1}{2} \right) \right] - (1 - \rho^2) \left[\omega'_e x'_e \left(v' + \frac{1}{2} \right)^2 - \omega''_e x''_e \left(v'' + \frac{1}{2} \right)^2 \right] + (1 - \rho^3) \left[\omega'_e y'_e \left(v' + \frac{1}{2} \right)^3 - \omega''_e y''_e \left(v'' + \frac{1}{2} \right)^3 \right]$$

In that equation, Δv is the isotopic shift expressed in cm⁻¹, v stands for the vibrational quantum number, ω_e represents the harmonic frequency, $\omega_e x_e$ is the first anharmonic constant and $\omega_e y_e$ is the second anharmonic constant. As for the term ρ it equals $(\mu/\mu^i)^{1/2}$, where μ is the reduced mass of the molecule and i corresponds to the heavier isotope. The number of apostrophes denotes the electronic levels involved in the electronic transition: two for the lower one, and one for the upper one.

As shown in **Table 3**, the shift can be predicted with great accuracy, as demonstrated by the excellent agreement found between theoretical and experimental values.

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3 Unfortunately, the magnitude of the shift was found to be inversely proportional
4 to the sensitivity of the transition, as it is related to the vibrational level. Higher
5 vibrational levels produce higher shifts, but also lower sensitivities because their
6 population is also lower.
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11 This can be seen both in **Table 3** and, more visually, in **Figure 2**, which shows
12 the signals obtained for different transitions. The 261.418 nm transition, the
13 most sensitive one, is not useful for isotopic analysis, as the signals for the two
14 isotopes are not resolved (see **Figure 2A**). The next transition, 261.695 nm, still
15 does not show two different peaks, but the exact wavelength of the maximum
16 depends on the isotopic composition (see **Figure 2B**), shifting to higher values
17 with higher ^{35}Cl contents. The next transition already shows two separate peaks
18 (see **Figure 2C**), but they are not perfectly resolved. The resolution further
19 improves (in **Figure 2D** the ratio of the peaks is already correct, close to 1) for
20 the next transitions, although for the last one the sensitivity is too low for the Cl
21 contents used in the experiment.
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35 **Figure 3A** shows that the transition selected and the way in which signals are
36 processed may have an effect on the results obtained. Peak height seems to be
37 less affected by overlaps than peak area (therefore, peak height was selected
38 as measuring mode), and the number of detector pixels used also needs
39 optimization.
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46 Under optimized conditions, it is possible to carry out the isotopic analysis of Cl
47 in 20 mg L^{-1} solutions with precision values around 2% RSD (see **Figure 3B**),
48 without the need for any mass bias correction. This precision is not sufficient to
49 monitor Cl isotopic natural variations, but it may be enough if the goal is to carry
50 out tracer experiments, or to be able to calibrate by means of isotope dilution
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3 (ID). In fact, it was demonstrated in this work *via* analyses of natural water
4 CRMs and well as commercially available mineral water samples that the use of
5 ID serves to correct for interferences caused by chemical reactions (e.g.,
6 presence of high levels of Ca, thus leading to some Cl forming CaCl and not
7 AlCl), as these interferences should affect both isotopes in practically the same
8 way. This is one important conclusion of the work, as in GFMS the occurrence
9 of interferences of chemical origin is more common than in GFAAS.
10

11
12 A subsequent work from the same authors showed the potential of HR CS
13 GFMS for isotopic analysis of Br, using the CaBr molecule, after evaluating
14 some alternative ones (such as AlBr or BaBr). In that case, the spectrum
15 obtained was richer in lines, and there were several lines that responded
16 selectively to the presence of Ca⁷⁹Br and Ca⁸¹Br. Interestingly, there were other
17 lines in which both Ca⁷⁹Br and Ca⁸¹Br transitions overlapped, so they could be
18 used for total Br content with higher signals.
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20
21 The precision obtained for Br solutions of 10 mg L⁻¹ was of 2.6%. Again, this is
22 not sufficient for monitoring natural variations in Br isotopic composition.
23 However, it was demonstrated that it is feasible to directly determine Br in solid
24 samples despite the presence of interferences of chemical origin (e.g., very high
25 Cl levels, which decrease the formation of CaBr due to the formation of CaCl
26 instead⁵³) using ID for calibration. **Figure 4** shows the spectrum of a solid
27 biological (tomato leaves) reference material vaporized together with a liquid
28 spike. The peak profiles obtained are well-defined and unimodal, indicating that
29 proper isotopic equilibration between the Br present in the solid and in the spike
30 solution has occurred during the development of the temperature program, thus
31 enabling ID to work properly.
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3 The last work reporting on this approach has been published by Abad *et al.*⁶⁷ In
4 this work, B is the analyte, but instead of monitoring B atomic lines, BH was
5 formed and measured *via* HR CS GFMS. The expected difficulties for B
6 vaporization discussed before were circumvented by a mixture of liquid and gas
7 phase modifiers: an CaCl₂ solution with the aim to minimize the level of oxygen
8 in the furnace and decrease the formation of boron carbide; a mannitol solution
9 to prevent the formation of volatile boron species; X-100 triton as surfactant; 2%
10 CHF₃ in Ar to minimize memory effects; and 2% H₂ in Ar as primary gas.

11
12 Two different portions of the spectrum were measured, in the vicinity of 433 nm
13 and in the vicinity of 437 nm, both of them showing several lines, of which some
14 were completely resolved for ¹⁰BH and ¹¹BH in the latter spectral region. The
15 method finally developed was not very sensitive, requiring at least 1 g L⁻¹ of B.
16 However, by using partial least regression and building a library of BH
17 molecular spectra by monitoring solutions with different ¹¹B fractions, it was
18 possible to report precisions in the range between 0.013 and 0.05%, which may
19 suffice for monitoring B isotopic natural variations.

20 21 22 4.2. Nanoparticle characterization

23
24 HR CS AAS offers some possibilities for research in the nano field. In the
25 beginning, some early works were focused on the analysis of nanomaterials
26 such as carbon nanotubes,²⁸ or in the determination of nanoparticles (NPs) in
27 tissues.⁶⁸ These works, while relevant as applications, did not represent much
28 of a change in the working procedure in comparison with analysis of other
29 samples or other analytes. In other words, carrying out the direct analysis of
30 impurities in carbon nanotubes is not so different from performing the direct
31 analysis of impurities in other hard to dissolve carbonaceous materials, such as
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3 graphite. And those early works aiming at the determination of NPs were, in
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5 fact, determining only the total elemental content, which was the result of
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7 exposition to NPs.

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9 This situation has recently changed. It has been reported that it is feasible to
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11 obtain information on whether a particular element is found in the form of
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13 nanoparticles or in ionic form, and it may even be possible to calculate its size,
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15 when present as nanoparticles, all of these using GFAAS.

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17 The first article to report on this aspect was published by Gagné *et al.*⁶⁹ It was
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19 demonstrated for Ag that atomization of nanoparticles requires more energy,
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21 and thus higher atomization temperatures, than the atomization of ionic
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23 species. And the higher the particle size, the higher the temperature required.
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25 Thus, they proposed the approach of either determining the atomization
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27 temperature or monitoring the changes in absorbance at different atomization
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29 temperatures, to detect the presence of AgNPs in biological samples (after
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31 sample preparation) and to have a rough estimation of their size.

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33 This pioneering work did not make use of HR CS GFAAS but of line source
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35 GFAAS. However, there is an important reason for using HR GFAAS in that
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37 type of study, and that is the expanded possibility to directly analyze solid
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39 samples and complex materials in general.²¹ That is an intriguing aspect since
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41 many of the techniques that can be used for quantifying both the total content
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43 and the size distribution of nanoparticles, such as the increasingly popular
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45 single particle (SP)-ICP-MS, while offering several advantages, do not allow
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47 direct analysis of the samples. This could be an important issue because, as in
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49 every speciation/fractionation study, ensuring the preservation of the exact way
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3 in which the analyte is originally found in the sample during sample preparation
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5 is not trivial.

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7 The next papers on this topic explored this possibility. In a first work,
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9 Feichtmeier and Leopold,⁷⁰ showed the potential of SS HR CS GFAAS for the
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11 direct detection of Ag nanoparticles in parsley samples. They evaluated two
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13 different parameters, the so called atomization delay (the time period between
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15 the beginning of the atomization step and the appearance of the peak
16
17 maximum) and the atomization rate (the slope of a polynomial fitted curve
18
19 calculated at the first inflection point of the absorbance vs. time peak profile). In
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21 the end, they proposed an approach based on both aspects. It is possible to
22
23 assess if Ag is present in ionic form or as NPs simply by calculating this
24
25 atomization delay (normalized to the sample mass): a difference of 6.27 ± 0.96
26
27 s mg^{-1} between parsley samples containing ionic silver and AgNPs was found
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29 under optimum conditions. Furthermore, if AgNPs were found, the
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31 determination of the atomization rate permitted to differentiate between NPs of
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33 20, 60 and 80 nm.
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38 A second work expanded the applicability of this method to different types of
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40 biological samples (namely parsley, apple, pepper, cheese, onion, pasta, maize
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42 meal, and wheat flour).⁷¹ This work focused on the use of the atomization delay
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44 to establish the presence of either ionic silver or of AgNPs of 20 nm. It was
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46 found that the matrix plays an important role in the values found but,
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48 nevertheless, significant differences in this parameter between samples
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50 exposed to ionic silver or to AgNPs were observed in all cases investigated.
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54 It is important to stress that this type of research abandons some of the
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56 classical GFAAS paradigms. In fact, the whole concept of stabilized
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3 temperature platform furnace (STPF) tries to minimize any potential difference
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5 in the signals obtained for the analyte, even when it could be present in various
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7 forms in the sample. In this particular case, the goal is exactly the opposite, and
8
9 thus the conditions selected must be chosen with the aim of exacerbating the
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11 differences between the vaporization/atomization mechanisms of every species.
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13
14 The work by Resano *et al.*⁷² showed that HR CS GFAAS could be applied to
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16 other NPs beside Ag, which was the focus of the previous works. Also, the use
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18 of different parameters to quantify the presence and the size of NPs was
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20 discussed, and it was concluded that the time of appearance of the peak
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22 maximum was the preferred one in this case, as it did not depend on the
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24 concentration and permitted to appreciate clear differences according to the
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26 particle size, as shown in **Figure 5**. It can also be appreciated in such figure
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28 that, under the conditions used (no modifier added and a very slow atomization
29
30 ramp ($150\text{ }^{\circ}\text{C s}^{-1}$)) the signal profiles observed for ionic Au and for AuNPs were
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32 dissimilar: in the case of AuNPs, the signal is not only delayed according to its
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34 size, but its shape is practically Gaussian; on the other hand, the signals
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36 obtained for ionic Au shows much more tailing, to the point that a secondary,
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38 smaller peak is found. This fact not only hints at different atomization
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40 mechanisms, but permits to easily screen the presence of either NPs or ions
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42 from an unknown solution.
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48 Further work has basically confirmed the potential of the technique for sizing
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50 Au⁷³ and Au and Ag NPs,⁷⁴ respectively, despite some differences in the
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52 programs used by the different authors, and on the relation found between size
53
54 and time of appearance (completely linear or not). Finally, another work
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56 demonstrated⁷⁵ the potential of solid sampling HR CS GFAAS for determining
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3 both the total Fe content (*via* measurement of the peak area) and the particle
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5 size (*via* evaluation of the atomization rate) in solid magnetic NPs.
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8 Overall, this seems to be a relevant and growing field, where GFAAS can
9
10 provide at least complementary information to other techniques. So far it cannot
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12 provide the same type of information than SP-ICP-MS, where not only the mean
13
14 particle size but the whole number-size distribution is obtained. But, as
15
16 discussed before, GFAAS capability to analyze directly complex samples is very
17
18 interesting, and so it is its ability to detect very small NPs (below 10 nm), which
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20 are not easily detected *via* SP-ICP-MS.
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24 Still, more work is needed in order i) to develop a more unified methodology and
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26 even terminology; ii) to further improve the selectivity of the technique, as right
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28 now it is difficult to resolve mixtures of NPs or of NPs with ionic species,
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30 requiring the use of deconvolution approaches^{72,73} or of previous separations (in
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32 this regard, cloud point extraction has been reported to enable the selective
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34 quantification of AgNPs and of total Ag in waters with a LOD of 2 ng L⁻¹ using
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36 HR CS GFAAS⁷⁶); and, iii) to demonstrate the applicability of this approach to
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38 other types of NPs.
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41 42 4.3. New instrumental developments 43

44 Perhaps the most critical aspect to move forward the application of HR CS
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46 AAS/MAS would be enhancing its multi-element capabilities.
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48 In fact, one of the main historical motivations for the development of CS AAS
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50 was the possibility to implement higher multi-element capabilities, and when HR
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52 CS AAS instrumentation was made commercially available there were great
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54 expectations in this regard. It is necessary to state, however, that the potential
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3 of such commercial instrumentation to carry out the truly simultaneous
4 monitoring of lines corresponding to different elements is rather limited.
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7 The reason is simple and it is related with the optics and the detector
8 capabilities: besides the optical system, in order to cover the whole UV-vis
9 range with the resolution required (pm level), use of megapixel array detectors
10 with a sufficiently fast response for transient signals would be required. Instead,
11 the current instrument possesses a detector with only 588 pixels, but most of
12 those are used for corrections, and only 200 are used to monitor the analytical
13 signals. That is why the current instrumentation can measure simultaneously
14 only a very small portion of the spectrum (0.2–1.0 nm, depending on the
15 wavelength) with the required resolution.
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18 The limitations of this instrumentation and the possibilities for multi-line
19 monitoring are discussed in detail in previous works,^{18,19} so they will not be
20 repeated herein. Also, a paper discussing the evolution of CS AAS, with an
21 emphasis on detector requirements, was published by Becker-Ross *et al.*⁷⁷
22
23

24 It suffices to say that, at this point, with the commercially available
25 instrumentation it is in principle possible to determine simultaneously several
26 elements (a paper in which Co, Fe, Ni, and Pb were determined in solid
27 samples has been reported²⁸), but this is more the exception than the rule, and
28 most papers determine only one element and, in a few cases, two. It is also
29 important to stress that this limitation is more relevant for GFAAS than for flame
30 AAS, because in the latter case a fast sequential method may be sufficient in
31 most practical situations, and the current instrumentation already enables such
32 approach.²²
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3 The important point regarding this aspect is that several recent works
4 demonstrate that it is technically possible to monitor a much wider portion of the
5 spectrum simultaneously while still keeping the resolution required, which is
6 always key (monitoring the whole UV-vis with low resolution is not challenging
7 at this point, but such approach is hardly suitable for analysis of complex
8 samples⁷⁸). In order to do that, Geisler *et al.* proposed the use of laser-driven
9 xenon lamp as continuum source, a CCD detector, and a modular simultaneous
10 echelle spectrograph, denominated MOSES.⁷⁹

11
12 A scheme of such instrumentation is shown in **Figure 6**. This instrument can be
13 operated in two modes (panorama or zoom), the former providing a wider
14 spectral interval, and the latter more resolution to focus on particular lines.
15 Practically the whole UV spectra (from 193 nm to 390 nm) can be covered with
16 this instrument in four sub-areas (tiles): from 193.24 to 211.21 nm; from 209.83
17 to 236.00 nm; from 234.37 to 280.69 nm; and from 278.47 to 389.89 nm. These
18 are the spectral windows that can be simultaneously monitored. **Figure 7** shows
19 an example of all the elemental and molecular lines that can be seen using the
20 last tile. This obviously represents a significant step forward in terms of multi-
21 element possibilities. Again, as discussed in section 3.1., it is worth stressing
22 that this instrumentation offers the intriguing possibility of making simultaneous
23 use of two techniques, AAS and MAS.

24
25 In a second article of the same research group, this instrumentation was used
26 to evaluate complete spectra of different diatomic molecules that can be used
27 for S determination *via* HR CS GFMA⁴⁴. These molecules are GeS, SiS, SnS,
28 and PbS, and the authors report the benefits of choosing GeS over the most
29 popular today, CS, as it provides a (very slightly) lower characteristic mass (9.4
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3 ng for GeS vs. 12 ng for CS using their most sensitive transitions) and its
4 generation can be adjusted by controlling the Ge amount. Certainly, use of this
5 type of instrument is beneficial to help in selecting the best transitions when
6 very rich in lines molecular spectra are investigated.
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11 Also, the possibility to monitor P₂ when aiming at P determination was
12 investigated by Huang *et al.*⁸⁰ using the MOSES. The main advantage of
13 selecting this molecule, in comparison with the measurement of P atomic
14 absorption, is that it shows thousands of sharp transitions in the range between
15 196 and 245 nm that could be measured simultaneously and summed, in order
16 to provide a better LOD,⁸¹ in addition to the much more gentle temperature
17 program required.
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22 Finally, the most recent work of this research group features the use of the
23 MOSES to investigate CaF spectra,⁸² but also the use of minitubes of only 2
24 mm diameters to increase the sensitivity (an approach evaluated in a previous
25 work⁸³) and of an autosampler capable of delivering volumes down to 30 nL. In
26 this way, determination of F in sweat samples (12 to 49 mg L⁻¹) from cancer
27 patients with volumes below 1 μL was achieved. The possibilities of this
28 approach for situations in which sample volume is scarce or precious is
29 therefore remarkable.
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32 33 34 35 36 37 38 39 40 41 42 43 44 **5. Conclusions**

45
46 HR CS AAS/MAS is at a crossroads. No longer a young technique, but still too
47 young to be considered as fully matured. While many of the research articles
48 published transmit a sense of continuity and a trend towards more applied work,
49 there are also some promising novel areas, such as characterization of NPs or
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2 isotopic analysis, and a need for more fundamental studies, particularly in the
3 case of HR CS MAS.
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7 However, the final fate of the technique may well be related to the development
8 of instrumentation commercially available with more multi-element potential,
9 which seems possible from a technical point of view. It is however uncertain
10 when (and if) this will happen. The fact that to this day only a manufacturer
11 produces HR CS AAS/MAS instrumentation can be regarded as a drawback
12 because, in principle, more competition should favor the development of new
13 instrumental devices and also affect the price of acquisition of such instruments.
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Table 1. Applications reporting on direct analysis using HR CS AAS since 2015. All of them used graphite furnace except for ref. 22.

Analyte	λ (nm)	Chemical modifier	Type of sample	LOD / m_0	Remarks	Ref.
Ag	328.07	No modifier	Biological matrices	-- / --	Detection of Ag NPs based on atomization delay with respect to ionic silver. Study of the influence of the matrix with model matrices	71
Al Si	309.271 251.611	No modifier for Al; Pd(II) + Mg(II) nitrates for Si	Silicon carbide nanocrystals	0.5 $\mu\text{g L}^{-1}$ / 6.4 pg 3 $\mu\text{g L}^{-1}$ / 389 pg	External calibration with aqueous standards	27
Al (AlH) Cd Cr Fe	425.315 228.802 425.433 425.076	H ₂ SO ₄ 10%(v/v)	Soil samples	0.42 $\mu\text{g mg}^{-1}$ / 0.18 μg 7.3 pg mg^{-1} / 3.9 pg 0.13 ng mg^{-1} / 27 pg 0.07 $\mu\text{g mg}^{-1}$ / 0.09 μg	All elements determined in the same sample aliquot. Cd was determined first at $T_{\text{atom}}=1700$ °C. After changing the wavelength, Cr, Fe and Al (as AlH) were simultaneously determined at 2600°C	29
As	193.696	Pd(II) + Mg(II) nitrates + Triton X-100	Fish and seafood	0.05 $\mu\text{g kg}^{-1}$ / 20 pg	Maximum sample mass of 0.25 mg for avoiding matrix effects. Screening method for determination of inorganic As only. External calibration with aqueous standards	84
As	193.696	Zr (permanent modifier)	Soils	73 pg / 22 pg	External calibration with aqueous standards	85
As Cd Co Cr Cu Ni	193.696 228.802 240.725 359.349 327.396 341.477	Ir (permanent modifier)	Marine sediments	0.02 ng/- 0.01 ng/- 0.021 ng/- 0.015 ng/- 0.016 ng/- 0.011 ng/-	Calibration with solid reference materials. Fast temperature program with short pyrolysis step	86
As	193.696	Ir (permanent modifier)	Marine biota	0.03 ng / 40 pg	Fast temperature programs with short	87

Cd	228.802	modifier)	samples	0.014 ng / 10 pg	pyrolysis and drying steps. Calibration with solid reference materials. Least squares algorithm for correcting PO interference in As determination	
Co	240.725			0.017 ng / 77 pg		
Cr	357.868			0.07 ng / 66 pg		
Cu	327.396			0.004 ng / 77 pg		
Ni	232.003			0.03 ng / 12 pg		
Au	242.795	No modifier	Geological samples	$2.24 \cdot 10^{-6} \mu\text{g g}^{-1}$ / 6 pg	Determination of Au after preconcentration of digested samples in carbon nanotubes. External calibration with aqueous standards	32
Cd	228.8	Pd(II) + Mg(II) nitrates	Facial make-up	$0.0020 \mu\text{g kg}^{-1}$ / 0.56 pg	External calibration with aqueous standards	88
Cd	228.802	Pd(II) nitrate	Lipsticks	5.0 pg / 2.8 pg	External calibration with aqueous standards	89
Cd	228.801	No modifier	Ethanol fuels	$0.006 \text{ mg kg}^{-1}/C_0=0.01 \text{ mg L}^{-1}$	Use of flame atomizer. Multi-element fast sequential determination. External calibration with aqueous standards	22
Co	240.725			$0.02 \text{ mg kg}^{-1}/C_0=0.06 \text{ mg L}^{-1}$		
Cu	324.754			$0.001 \text{ mg kg}^{-1}/C_0=0.02 \text{ mg L}^{-1}$		
Fe	248.327			$0.02 \text{ mg kg}^{-1}/C_0=0.05 \text{ mg L}^{-1}$		
Mn	279.481			$0.003 \text{ mg kg}^{-1}/C_0=0.01 \text{ mg L}^{-1}$		
Na	589.592			$0.1 \text{ mg kg}^{-1}/C_0=0.03 \text{ mg L}^{-1}$		
Ni	232.003			$0.04 \text{ mg kg}^{-1}/C_0=0.08 \text{ mg L}^{-1}$		
Pb	217.000	$0.06 \text{ mg kg}^{-1}/C_0=0.10 \text{ mg L}^{-1}$				
Zn	213.857	$0.024 \text{ mg kg}^{-1}/C_0=0.01 \text{ mg L}^{-1}$				
Cd	228.802	No modifier	Tannin samples	$0.5 \mu\text{g kg}^{-1}$ / 0.3 pg	Sequential determination of Cd and Cr in the same sample aliquot. External calibration with aqueous standards	90
Cr	357.869			$17 \mu\text{g kg}^{-1}$ / 2.2 pg		
Cd	228.802	No modifier	Yerba mate	0.25 ng / 0.37 pg	Sequential determination for the same sample aliquot. External calibration with aqueous standards	91
Cr	357.869			0.72 ng / 2.4 pg		
Cd	228.802	Pd(II) + Mg(II) nitrates for Cd	Vegetables of the Solanaceae	0.78 ng g^{-1} / 0.42 pg	External calibration with aqueous standards	92
Cr	357.869			2.5 ng g^{-1} / 2.7 pg		

Cu	327.396		family	0.15 $\mu\text{g g}^{-1}$ / 19 pg		
Cd	228.802	Pd(II) + Mg(II)	Spices	0.2 ng g^{-1} / 0.6 pg	Air-assisted pyrolysis step needed for enabling external calibration with aqueous standards	93
Ni	232.003	nitrate for Cd; No		18 ng g^{-1} / 14.0 pg		
V	318.540	modifier for Ni, V		7 ng g^{-1} / 18.6 pg		
Co	240.725	No modifier	Wet animal feeds	0.051 $\mu\text{g g}^{-1}$ / 2.9 pg	External calibration with aqueous standards	94
Cr	357.869			0.009 $\mu\text{g g}^{-1}$ / 2.2 pg		
Co	252.136	No modifier	High purity silicon	0.39 mg kg^{-1} / 0.02 pg	External calibration with aqueous standards. Simultaneous determination of Fe and Ni	26
Fe	305.909			1.14 mg kg^{-1} / 0.09 pg		
Ni	305.760			5.71 mg kg^{-1} / 0.54 pg		
Co	304.400	Mg(II) nitrate for Ni. NH_4F for Co and V	Soil	0.045 ng / 0.093 ng	Co and V determined simultaneously. Different lines for Ni and Mo used for adjusting sensitivity. External calibration with aqueous standards feasible for all elements except Co. For Co, calibration with solid reference material was carried out	95
Mo	313.259			5.4 pg / 0.009 ng		
Ni	341.347			1.8 ng / 2.45 ng		
	341.477			0.021 ng / 0.045 ng		
V	304.355			0.43 ng / 0.43 ng		
	304.494	0.59 ng / 0.64 ng				
Cr	357.869	Mixture of H_2O_2 + ethanol + HNO_3 plus Mg(II) nitrate	Infant formulas	3.45 ng g^{-1} / 1.2 pg	Mixture of modifiers needed for reduction of carbonaceous residues. External calibration with aqueous standards feasible	96
Cr	357.868	No modifier	Pharmaceutical drugs	2.9 $\mu\text{g kg}^{-1}$ / 2.0 pg	External calibration with aqueous standards	97
Cr	360.532	Ru (permanent modifier) + Pd(II) + Mg(II) nitrates + Triton X-100 for Pb	Industrial waste of oil shale and various reference materials	102 $\mu\text{g g}^{-1}$ / 25 pg	External calibration with aqueous standards	98
Cu	327.396			39 $\mu\text{g g}^{-1}$ / 19 pg		
Pb	283.306			28 $\mu\text{g g}^{-1}$ / 24 pg		

Cr Pb	359.349 283.306	No modifier for Cr; Pd(II) + Mg(II) nitrates in Triton X-100 for Pb	Sunscreen samples	1.0 $\mu\text{g kg}^{-1}$ / 2.5 pg 3.0 $\mu\text{g kg}^{-1}$ / 10 pg	External calibration with aqueous standards	99
Cr Tl	427.5 276.8	Mg(II) nitrate in Triton X-100 (Cr); No modifier (Tl)	Fertilizer samples	60 ng g^{-1} / 88 pg 3 ng g^{-1} / 11 pg	HR-CS GF AAS used to confirm the absence of spectral interferences. Comparison with LS-GFAAS with Zeeman effect	100
Cu	249.215	No modifier	Micro-volumes of distilled alcoholic beverages	0.03 mg L^{-1} / --	Calibration with matrix-matched liquid standards. Internal standardization with Cr, Fe and/or Rh feasible	101
Cu Cu Hg	327.396 249.215 253.652	KMnO ₄ for Hg; No modifier for Cu	Phosphate fertilizers	19 $\mu\text{g g}^{-1}$ / 25 ng 0.012 $\mu\text{g g}^{-1}$ / 1.12 ng 0.015 $\mu\text{g g}^{-1}$ / 39 ng	External calibration with aqueous standards	25
Cu Fe	217.894 217.812	No modifier	Flours	0.11 mg kg^{-1} / 14 pg 0.03 mg kg^{-1} / 77 pg	Simultaneous determination of the two analytes. External calibration with aqueous standards	102
Cu Mn	324.754 403.076	No modifier	Infant formula	0.003 $\mu\text{g g}^{-1}$ / 8.0 pg 0.01 $\mu\text{g g}^{-1}$ / 8.0 pg	External calibration with aqueous standards	103
Fe	352.614	Pd(II) nitrate	Solid magnetic nanoparticles	--/--	External calibration with aqueous standards. Method used for characterization of magnetic NPs	104
Fe Ni	232.036 232.001	H ₂ gas admixed during pyrolysis	Fluoropolymers	221 ng g^{-1} / 370 pg 9.6 ng g^{-1} / 7.7 pg	Simultaneous determination of Fe and Ni. External calibration with aqueous standards	105
Fe Ni	232.036 232.003	No modifier	Vegetables of Solanaceae family	2 $\mu\text{g g}^{-1}$ / 340 pg 0.02 $\mu\text{g g}^{-1}$ / 95 pg	Simultaneous determination of Fe and Ni. Least squares algorithm for correcting interference of SiO. External calibration with aqueous standards	106

Fe Ni V	294.132 294.391 294.236	Ir (permanent modifier)	Fuel fly ash	3.19 ng /-- 1.20 ng /-- 24.13 ng /--	Simultaneous determination of Fe, Ni and V. External calibration with aqueous standards	107
Hg	253.652	Ir (permanent modifier)	Marine sediments and biota	0.025 ng sediments/- 0.096 ng biota/-	Calibration with solid reference materials. Fast temperature program with short pyrolysis step	108
Hg	253.652	Au NPs	Airborne particles	0.19 $\mu\text{g L}^{-1}$ / 0.004 $\mu\text{g L}^{-1}$	No pyrolysis step. Comparison of the effects of Ag and Au NPs as modifiers	109
Hg	253.652	Au nanoparticles	Blood and urine	2.3 $\mu\text{g L}^{-1}$ / 16 pg	External calibration with aqueous standards. Fast temperature program without pyrolysis	35
Li Na	610.353 285.301	No modifier	Yttrium oxyorthosilicate microsamples	20 $\mu\text{g g}^{-1}$ / 80 $\mu\text{g g}^{-1}$ /	Standard addition method for calibration	110
Mg	215.435	No modifier	Lithium niobate crystals	0.7 mg g^{-1} /--	Calibration with powdered lithium niobate crystal standards	111
Mn Ni Rb	403.076 231.096 420.018	No modifier	Meat samples	0.005 $\mu\text{g g}^{-1}$ / -- 0.002 $\mu\text{g g}^{-1}$ / -- 0.1 $\mu\text{g g}^{-1}$ / --	External calibration with aqueous standards for Mn. Calibration with solid reference materials for Ni and Rb. Least squares algorithm for correcting PO and SiO molecular interferences on determination of Ni	112
Mn Ni Rb Sr	403.076 231.096 420.018 407.771	No modifier	Powdered stimulant plants	0.05 $\mu\text{g g}^{-1}$ / 0.012 ng 0.02 $\mu\text{g g}^{-1}$ / 0.015 ng 0.1 $\mu\text{g g}^{-1}$ / 0.28 ng 0.01 $\mu\text{g g}^{-1}$ / 1.5 pg	External calibration with aqueous standards for Mn, Rb and Sr. Calibration with solid standards for Ni	113
Mo Ni	313.259 313.410	No modifier	Plant materials	0.018 ng / 0.12 ng 0.025 ng / 0.017 ng	Simultaneous determination of Mo and Ni. Co used as internal standard for avoiding	31

					matrix effects in Ni determination. External calibration with aqueous standards	
Mo V	313.259 318.398	No modifier	Reference materials of various natures	25 pg / 7 pg 130 pg / 18 pg	Atomization in disposable starch-based platforms made in-house for reducing tailing of the analyte signals. External calibration with aqueous standards	36
Pb	205.328 283.060	Pd(II) + Mg(II) nitrates	Glass	201.6 pg / 1.6ng 11.2 pg / 0.1ng	External calibration with aqueous standards compared with calibration with NIST 612 glass sample. Both approaches provide similar results	114
Pb	283.306	Pd(II) + Mg(II) nitrates	Eye shadow and blush	6 pg g ⁻¹ /-	External calibration with aqueous standards. Sample mass of 0.3 mg	115
Pb	217.001 205.328	Pd(II) + Mg(II) nitrates	Plastic food packaging	4.9 µg kg ⁻¹ / 5.0 pg 0.5 mg kg ⁻¹ / 0.4 ng	External calibration with aqueous standards. Secondary line at 205.328 nm used for adjusting sensitivity	116
Pb	217.005	Pd(II) + Mg(II) nitrates	Plastic toys	0.037 mg kg ⁻¹ / 9 pg	Ar gas flow during the atomization stage for adjusting sensitivity to the sample contents. External calibration with aqueous standards	117
Pb	217.001	Pt chloride	Dried Blood Spots	1 µg L ⁻¹ / 26.6 pg	Screening method developed for Pb in DBS from newborns and pregnant women. External calibration with aqueous standards	118
Pb	217.001 283.306	No modifier	Biomass and products of the pyrolysis process	5 pg / 4 pg 9 pg / 7 pg	Line at 217.001 nm could not be used for some samples due to spectral interferences. External calibration with aqueous standards	119
Pb	283.306	Pd(II) + Mg(II) nitrates	Incense	Rod: 5.6 pg / 8.2 pg Coating: 12 pg / 29 pg	Analysis of rod and coatings of incense sticks. External calibration with aqueous standards for rods, but with solid matrix-	120

					matched standards for coatings	
Pb	261.418 283.306	Pd(II) + Mg(II) nitrates + Triton X- 100	Road dust and soil	0.65 mg kg ⁻¹ / 1.20 ng 0.09 mg kg ⁻¹ / 0.38 ng (using side pixels +4, -4)	Investigation of molecules giving rise to spectral overlaps. Use of side pixels. External calibration with aqueous standards	121
Pd Pt Rh Rh	360.955 244.006 361.247 244.034	No modifier for pharmaceuticals; NH ₄ F·HF for catalysts	Automobile catalysts and active pharmaceutical ingredients	Pd Catalysts 6.5 µg g ⁻¹ / 0.44 ng Pharm. 0.08 µg g ⁻¹ / 0.18 ng Pt Catalysts 8.3 µg g ⁻¹ / 0.81 ng Pharm. 0.15 µg g ⁻¹ / 0.32 ng Rh (244.034 nm) Catalysts 9.3 µg g ⁻¹ / 0.84 ng Pharm. 0.10 µg g ⁻¹ / 0.30 ng Rh (361.247 nm) Catalysts -- / 1.58 ng Pharm. 0.26 µg g ⁻¹ / 0.68 ng	External calibration with aqueous standards for pharmaceuticals. Calibration with a solid reference material for the catalysts. Simultaneous determination of Pt+Rh and Pd+Rh feasible in both samples. An Ar flow of 0.1 L min ⁻¹ during atomization used for analysis of catalysts	24
Rh Ru	343.489 343.674	NH ₄ F·HF	Spiked river water, road runoff and municipal sewage	1.0 µg L ⁻¹ / 12.9 pg 1.9 µg L ⁻¹ / 71.7 pg	Simultaneous determination of Rh and Ru. External calibration with aqueous standards	122
Sb	217.581	Pd(II) + Mg(II) nitrates	Cosmetics	0.27 mg kg ⁻¹ /0.028 ng	Study of different precursors for generating reference spectra for correcting structural molecular background. Zeolite and mica were the most successful ones	123
Sb	217.582	Pd(II) nitrate	Waters subjected to micro-solid	0.02 µg L ⁻¹ / --	Total Sb determination and determination of Sb(III) achieved by two extractions carried out at different acidity. Sb(V) calculated by	33

			phase extraction with magnetic core-modified silver NPs		difference. Comparison with slurry approach. External calibration with aqueous standards	
Si	221.174	Rh (permanent modifier) plus Pd(II) + Mg(II) nitrates in Triton X-100	Plant materials	2.5 ng / 2.0 ng	Secondary line, Ar flow during atomization stage and absorbance in the center peak only used for adjusting sensitivity. External calibration with aqueous standards	124
Si	251.611 221.174	Rh (permanent modifier) plus Pd(II) + Mg(II) nitrates	Biomass and products of pyrolysis process	0.09 mg kg ⁻¹ / 0.3 ng 3 mg kg ⁻¹ / 2 ng	External calibration with aqueous standards	125
Tl	276.786	Pd(II) nitrate	Spruce needles	1.2 µg kg ⁻¹ / --	Comparison of four methods of analysis	126

Table 2. Applications reporting on the determination of non-metals using HR CS MAS since 2015.

Vaporizer	Analyte	Species monitored	λ (nm)	LOD / m_0	Chemical modifier	Type of sample	Remark	Ref.
Graphite furnace	Br	Ca ⁷⁹ Br Ca ⁸¹ Br CaBr	600.492 600.467 625.315	4.0 ng / --	Pd nanoparticles	CRMs: PVC and tomato leaves	Isotopic analysis of Br. Direct solid sampling with ID calibration	55
Graphite furnace	Br	CaBr	625.315	10 $\mu\text{g g}^{-1}$ (LOQ)	Zr (permanent modifier) + Pd	6 polymeric CRMs	Direct solid sampling. Comparison with LA-ICP-MS	127
Graphite furnace	Br	TlBr	342.982	0.3 ng / 4.4 ng	Ag	Water	Direct analysis. The modifier prevents losses of Br through precipitation as AgBr	56
Graphite furnace	Cl	Al ³⁵ Cl Al ³⁷ Cl	262.238 262.222	0.25 mg L ⁻¹ / --	Pd	Water CRMs and mineral waters	Isotopic analysis of Cl. ID for calibration	54
Graphite furnace	Cl	SrCl	635.862	0.85 ng / 0.24 ng	Zr (permanent modifier)	Coal (5 CRMs)	Direct solid sampling. External calibration with aqueous standards	128
Graphite furnace	Cl	AlCl InCl SrCl	261.418 267.281 635.862	2.1 ng / 0.28 ng 3.5 ng / 1.7 ng 0.7 ng / 1.0 ng	Zr (permanent modifier)	NIST 1634c fuel oil, Crude oils	Direct analysis. External calibration with aqueous standards. Sr added also to increase the sensitivity for AlCl and InCl molecules	129
Graphite furnace	Cl	InCl	267.217	0.10 ng / 0.21 ng	Pd + Mg	Water samples, a diclofenac pill and CRM QC 1060 anions-WP	Non-spectral interferences in the presence of other halogens	130
Graphite	Cl	SrCl	635.862	1.76 $\mu\text{g mL}^{-1}$ / --	Zr (permanent	Milk and one	Dilution of the sample to adjust	131

furnace					modifier)	CRM (waste water)	the sensitivity. External calibration with aqueous standards	
Graphite furnace	Cl	CaCl	620.862 377.501	0.75 ng / 0.072 ng 14.2 ng / 25.9 ng	Pd	CRMs of different nature	Direct solid sampling. External calibration with aqueous standards	50
Graphite furnace	Cl	CaCl	621.145	1.4 ng / --	No modifier	Cement	Direct solid sampling. External calibration with aqueous standards. Ca compounds present in cement samples acted as CaCl-forming reagent	132
Graphite furnace	Cl	SrCl	635.862	1.8 ng / 0.32 ng	Zr (permanent modifier)	Fish oil samples	Dilution with 1-propanol. External calibration with aqueous standards	133
Graphite furnace	Cl	MgCl	377.010	1.7 ng / 7.1 ng	W (permanent modifier) + Pd	Water samples from offshore oil platform and two CRMs	Wall vaporization. Direct analysis of water samples. External calibration with aqueous standards	52
Graphite furnace	Cl	MgCl	377.010	3.0 $\mu\text{g g}^{-1}$ / --	W (permanent modifier) + Pd	Crude oil samples and NIST 1634c (fuel oil) and NIST 1848 (lubricating oil)	Analysis after emulsification. External calibration with aqueous standards	134
Graphite	Cl	GaCl	249.060	2.3 ng / 6.5 ng	Zr (permanent	Milk, mineral	GaCl vaporizes at a low	51

furnace					modifier) Pd(II) + Mg(II) in solution	water and a certified wastewater sample	temperature (1100 °C)	
Graphite furnace	F	CaF	606.440	0.3 ng / 0.1 ng	Pd (permanent modifier)	Coal CRMs	Direct solid sampling. Comparison of different modifiers. External calibration with aqueous standards	135
Graphite furnace	F	CaF	606.440 + 606.231	0.18 ng / 0.058 ng	No modifier	Turkish wines	Summation of the absorbances at two λ in order to increase the sensitivity and reduce LOD	136
Flame C ₂ H ₂ /N ₂ O	F	AlF BF	227.66 195.59	-- / -- -- / --			Research on how the chemical form of F species affects the sensitivity of AlF	59
Graphite furnace	F	CaF	606.045	2 mg kg ⁻¹ / 0.68 ng	No modifier	Eye shadow samples and two CRMs	Direct solid sampling. External calibration with aqueous standards	137
Graphite furnace	F	CaF	606.440	0.20 ng / 0.17 ng	No modifier	Baby food and one CRM (bush branches)	Direct solid sampling. External calibration with aqueous standards	138
Graphite furnace	F	CaF					Thermodynamic simulation of the mechanism of formation of CaF	139
Graphite furnace	F	CaF	606.440	0.22 ng / 0.16 ng	No modifier	Flour samples and one CRM (bush branch NCS DC 73349)	Slurries sampling. External calibration with aqueous standards feasible	140

Graphite furnace	F	CaF	606.432	-- / --	Pd (permanent modifier)	Airborne particulate matter (PM ₁₀) and CRMs (NIST 1648 and 1648a)	Direct solid sampling. Determination of Cl, Br and I by ETV-ICP-MS	141
Graphite furnace	F	SrF	651.187	0.36 ng / --		PM2.5 airborne particulates and wastewater CRM	Determination of F in particulate loaded quartz filters	142
Graphite furnace	F	CaF	606.440	0.72 ng mg ⁻¹ / 0.13 ng	No modifier	Soil samples and one CRM (lake sediment)	Direct solid sampling. External calibration with aqueous standards	143
Graphite furnace	F	CaF	606.440	0.13 ng mL ⁻¹ / --	No modifier	Herbs, water samples and various CRMs	Use of slurries combined with preconcentration on nano-TiO ₂ prior to injection to HR CS MAS. External calibration with aqueous standards	144
Graphite furnace	F	GaF	211.248	8.1 µg L ⁻¹ / --	Pd/Mg + Zr + CH ₃ COONa	Surface waters and two river water CRMs	Comparison with ion chromatography. HR-CS-GFMAS offers higher throughput, a lower LOD and higher selectivity	145
Graphite furnace	F	SrF	651.187	30 µg L ⁻¹ / --		Mineral waters and a water CRM	Chemical interferences were caused only by high chlorine concentrations	146
Graphite	F	CaF	606.440	-- / --	No modifier	Tea leaves	Study of the mechanism of	58

furnace							formation of CaF	
Graphite furnace	F	CaF	583.069 606.440	0.1 ng / -- 0.01 ng / 0.1 ng	Zr (permanent modifier)	Sweat samples of cancer patients	Use of mini graphite tubes with a low sample consumption (down to 30 nL)	82
Graphite furnace	I	CaI SrI	638.904 677.692	0.041 mg g ⁻¹ / 23 ng 0.021 mg g ⁻¹ / 18 ng	Pd/Mg	Pharmaceutical (one liquid drug and sodium levothyroxine pills)	Even a small Cl concentration reduces the absorbance of both molecules significantly. External calibration with aqueous standards and standard addition (pills)	57
Graphite furnace	NO ₂ ⁻ + NO ₃ ⁻	NO	215.360	0.16 mg dm ⁻³ / --	Ca	Water CRM	Total NO ₂ ⁻ + NO ₃ ⁻ content monitoring NO	147
Graphite furnace	NO ₂ ⁻ + NO ₃ ⁻	NO	215.371	0.3 mg L ⁻¹ / --	Ca	Herbal infusions	Standard addition method for calibration	148
Quartz cell	NO ₂ ⁻ + NO ₃ ⁻ + PNP	NO	215.360	0.16 mg dm ⁻³ / --	Ca	Food and water samples	Speciation method for nitrite, nitrate and p-nitrophenol via photochemical vapor generation. Nitrate and p-nitrophenol need to be reduced to nitrite first to be determined	60
Graphite furnace	P	P ₂	204.205	7 ng / 10 ng	Zr (permanent modifier) plus sodium tetraborate	CRMs (Spinach leaves and pine needles) and serum reference materials	Use of P ₂ enables gentle temperature conditions and multi-line evaluation to improve LOD	80
Flame	P	PO	327.04	1.0 mg L ⁻¹ / --	Al	Ferrovanadium	Ca and Mg interfere, but Al react	149

C ₂ H ₂ /air						sample and CRM	with them forming stable compounds and eliminating the interference	
Graphite furnace	P	PO	213.561 213.526 213.617 213.637	2.35 mg g ⁻¹ / --	Mg	Soybean lecithin samples	Samples diluted in MIBK. The signal was obtained by summing four PO lines surrounding 213 nm to increase the sensitivity	150
Flame C ₂ H ₂ /air	S	CS	258.056	1.5 mg g ⁻¹ / --		Food samples and CRMs	Microwave-assisted digestion of samples	151
Graphite furnace	S	CS	258.056	7.5 ng /8.7 ng	Ru (permanent modifier) plus Pd and citric acid	Vegetables and CRMs	Direct solid sampling. External calibration with aqueous standards prepared with thioacetamide	152
Graphite furnace	S	CS	258.056	1.4 mg kg ⁻¹ /17 ± 3 ng	Ir (permanent modifier) plus Pd/Mg	Diesel fuel samples and CRMs	Dilution of samples with 1-propanol. External calibration with aqueous standards prepared with L-cysteine	153
Flame C ₂ H ₂ /air	S	CS	257.961	52.4 mg kg ⁻¹ / --		Preserved fruits	A method to detect SO ₂ in this kind of sample	154
Flame C ₂ H ₂ /air	S	CS	258.056	25 mg L ⁻¹ / --		Bovine serum albumin and L-cysteine	During MW digestion S is converted to SO ₄ ²⁻ previous to the determination. Standard addition method for calibration	155
Flame C ₂ H ₂ /air	S	CS	258.056	11.6 mg L ⁻¹ / --		Vinegars	Due to non-spectral interference, all sulfur species were oxidized to SO ₄ ²⁻ . Standard addition method for calibration	156

Graphite furnace	S	CS	258.056	0.0001 % (w/v) / --		Particulate loaded filter samples and one CRM	PM2.5 airborne particulates collected in quartz filters	157
Graphite furnace	S	SnS	271.624	5.8 ng / 13.3 ng	Pd	Crude oil samples and CRMs	Samples were prepared as micro-emulsion due to their high viscosity. External calibration with aqueous standards	158
Graphite furnace	S	CS	258.056	0.018 % / --	No modifier	Human hair samples and two hair CRMs	Correlation of sulfur levels of autistic children's hair with total protein and albumin	159
Graphite furnace	S	SiS GeS SnS PbS	282.910 295.209 271.578 335.085	-- / 15.7 ng -- / 9.4 ng -- / 20 ng -- / 220 ng	Zr (permanent modifier) plus sodium tetraborate		GeS provides the best sensitivity while PbS offers the lowest sensitivity and a low bond strength	44
Graphite furnace	S	CS	258.033	0.30 mg g ⁻¹ / --	W (Permanent modifier) plus KOH	Petroleum green coke samples and a CRM (NIST 2718)	Slurry sampling. External calibration with aqueous standards prepared with thiourea	160
Quartz cell	Free and total S (IV)	SO ₂	215.356 215.416 215.486	0.36 mg L ⁻¹ / --		Coconut water	Measurement of SO ₂ generated after the addition of HCl. For total S, NaOH was added before to release the sulfite bound to organics. Summation of three SO ₂ transitions to improve sensitivity	61

Graphite furnace	S	CS	258.056	6 ng / --	Zr (permanent modifier) plus Ca and Pd	Milk, orange juice, swamp water, urine and two biological CRMs	Direct analysis of liquid samples and digestion of the solid CRMs. External calibration with aqueous standards	161
Graphite furnace	S (as proxy for cysteine)	CS	258.034 258.056	0.4 mg kg ⁻¹ / --	Pd/Mg	Pharmaceutical formulations and a CRM (NIST 1573a)	Summation of two CS transitions	162
Graphite furnace	S	CS	257.958	23 ng / --	Pd/Mg	Apricot and grape samples and a CRM (NIST 1568a)	258.056 nm line not used due to potential overlap with Fe line	163

Table 3. Theoretical and experimental isotopic shifts calculated and observed *via* HR CS GFMS, respectively, for the analytical band heads of the AlCl molecule ($X^1\Sigma^+ \rightarrow A^1\Pi$ electronic transition) for different vibrational transitions (v', v''). Reproduced with permission of the RSC (<http://pubs.rsc.org/en/content/articlehtml/2015/ja/c5ja00055f>).⁵⁴

λ/nm	$\lambda_{\text{exp}}/\text{nm}$	v', v''	$\Delta\lambda_{\text{calc}}/\text{pm}$	$\Delta\lambda_{\text{exp}}/\text{pm}$	Relative sensitivity / %
261.44	261.418	0,0	1.37	---	100
261.70	261.695	1,1	4.86	4.8	62
261.82	261.819	2,2	9.61	9.6	30
262.24	262.238	3,3	16.0	15.6	26
262.70	262.697	4,4	24.2	24.3	16
263.22	263.216	5,5	34.8	35.3	7.4
263.81	263.807	6,6	48.0	48.1	3.2
264.49	264.490	7,7	64.3	64.5	1.1

Figure captions

Figure 1. Number of articles reporting on the use of HR CS depending on: A) the type of atomizer; and B) the technique. Source: Scopus, September 2018.

The search was carried out using the general term “high resolution continuum source” and the results were curated manually, in order to select only those related with the topic of the review.

Figure 2. Spectra of AlCl when vaporizing 400 ng of Cl from an approximately 1:1 $^{35}\text{Cl}/^{37}\text{Cl}$ molar ratio solution, in the presence of 10 μg Al and 20 μg Pd, and monitoring the following wavelengths *via* HR CS GFMS: A) 261.418 nm; B) 261.695 nm; C) 261.819 nm; D) 262.238 nm; E) 262.697 nm; F) 263.216 nm; G) 263.807 nm; and H) 264.490 nm. In Figure 1B, the spectrum obtained for CRM NIST 975a (^{35}Cl 75.774 %) is shown in black, and the one obtained for CRM AE642 (^{37}Cl 98.122 %) is highlighted in red. Reproduced with permission of the RSC (<http://pubs.rsc.org/en/content/articlehtml/2015/ja/c5ja00055f>).⁵⁴

Figure 3. A) Quantification of the Cl ratio for a solution (200 ng Cl) of CRM NIST 975a ($^{35}\text{Cl}/^{37}\text{Cl}=3.1279 \pm 0.0047$) for different AlCl transitions. Values in blue were obtained using peak area, and values in green using peak height. In both cases, a lighter color intensity indicates a higher number of detector pixels (1, 3, 5 or 7). Error bars represent the standard deviation ($n = 5$). B) Peak height values of 20 replicates of the same solution obtained for 262.222 (Al ^{37}Cl) and 262.238 (Al ^{35}Cl) nm transitions (left y-axis), and Cl ratios calculated from the same set of data (right y-axis). The red dashed line indicates the certified value. The blue line and the blue interval surrounding it represent the average value of the 20 replicates and its standard deviation, respectively. Reproduced with

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5 (<http://pubs.rsc.org/en/content/articlehtml/2015/ja/c5ja00055f>).⁵⁴
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7 **Figure 4.** CaBr 3-D spectrum observed in the region of 600.4 nm when using
8 HR CS GFMS for the direct vaporization of a solid sample (SRM NIST 1573a
9 tomato leaves) containing approximately 1.0 μg Br, mixed with a spike enriched
10 in ^{81}Br . The main band heads are highlighted. Reproduced with permission of
11 the RSC <http://pubs.rsc.org/en/content/articlehtml/2016/JA/C6JA00114A>.⁵⁵
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17 **Figure 5.** Time-resolved absorbance for an Au aqueous standard and for
18 aqueous suspensions of AuNPs of 20 nm (diameter 19.6 nm; standard
19 deviation: 2.1 nm, as estimated *via* TEM by the manufacturer) and 76 nm
20 (diameter 75.7; standard deviation: 10.1 nm), respectively. In all cases, a
21 content of 50 $\mu\text{g L}^{-1}$, an atomization temperature of 2000 $^{\circ}\text{C}$ and a heating ramp
22 of 150 $^{\circ}\text{C s}^{-1}$ were used. Reproduced with permission of the RSC
23 <http://pubs.rsc.org/en/Content/ArticleLanding/2016/JA/C6JA00280C>.⁷²
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32 **Figure 6.** Schematic diagram of the experimental set-up including MOSES
33 spectrograph: (1) laser-driven continuum source, (2, 4) transfer optics
34 consisting of two elliptical mirrors, (3) atomizer, (5) entrance slit, (6) folding
35 mirror, (7) off-axis parabolic mirror, (8) prism stage with two Littrow prisms, (9a,
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from Elsevier
(<https://www.sciencedirect.com/science/article/pii/S0584854715000543>).⁷⁹

Figure 7. Temporally and spectrally resolved simultaneous multi-element
absorption signals obtained by the MOSES spectrograph in tile No. 4 setup, as
obtained for 20 μL of a standard solution containing: 50 $\mu\text{g L}^{-1}$ of Cr and Cu; 100
 $\mu\text{g L}^{-1}$ of Co; 200 $\mu\text{g L}^{-1}$ of As, Mo, Se, and V; 2000 $\mu\text{g L}^{-1}$ of Ir. Reproduced with

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5 (<https://www.sciencedirect.com/science/article/pii/S0584854715000543>).⁷⁹
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15 **Figure 1**

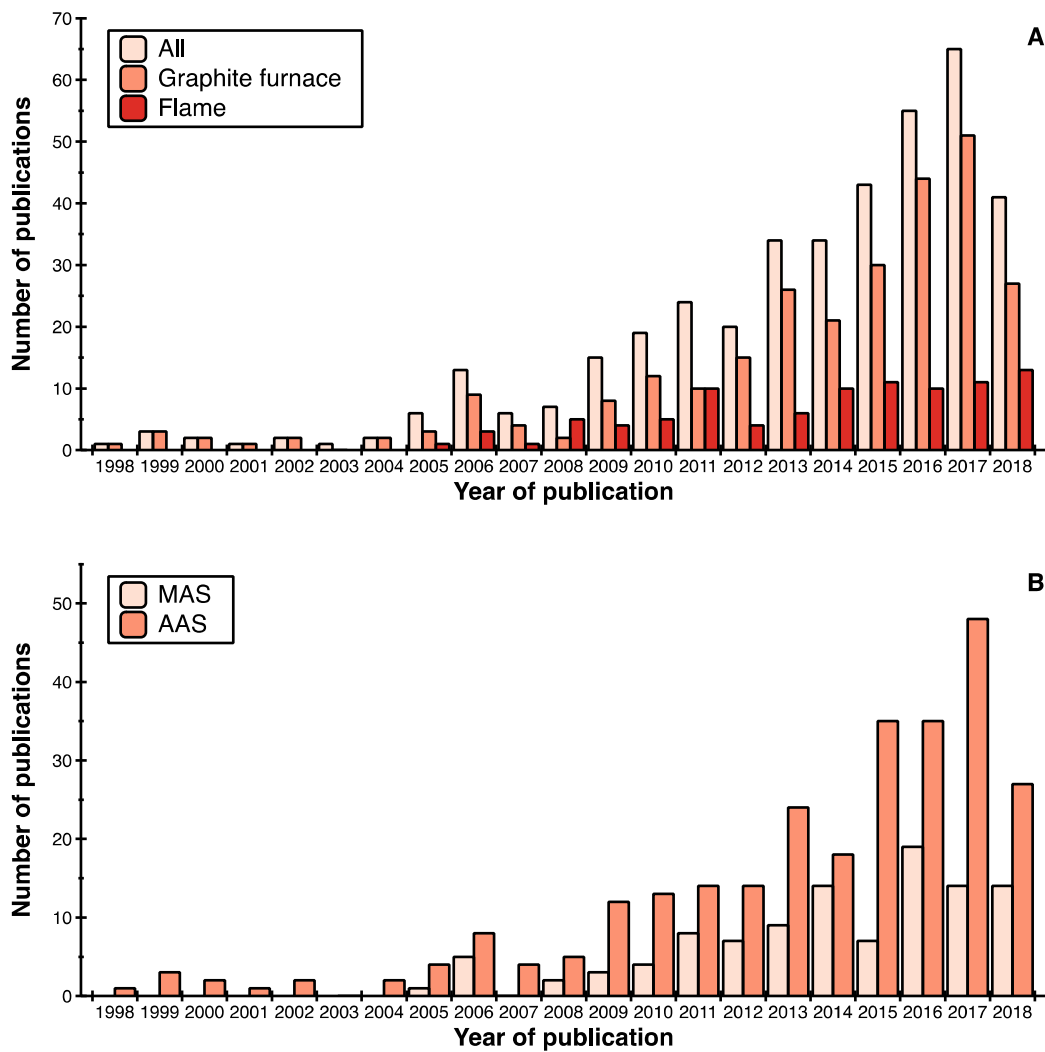


Figure 2

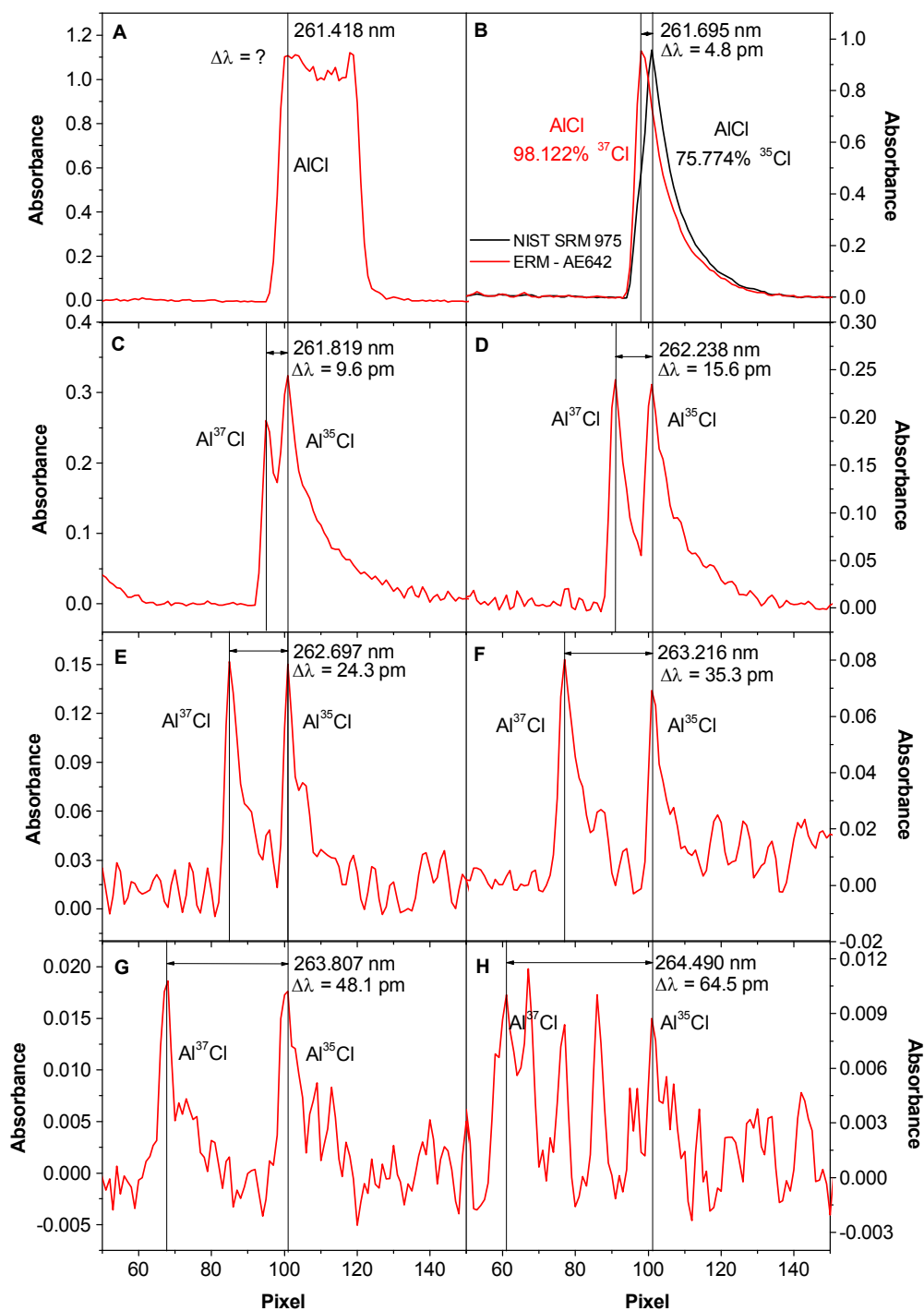


Figure 3

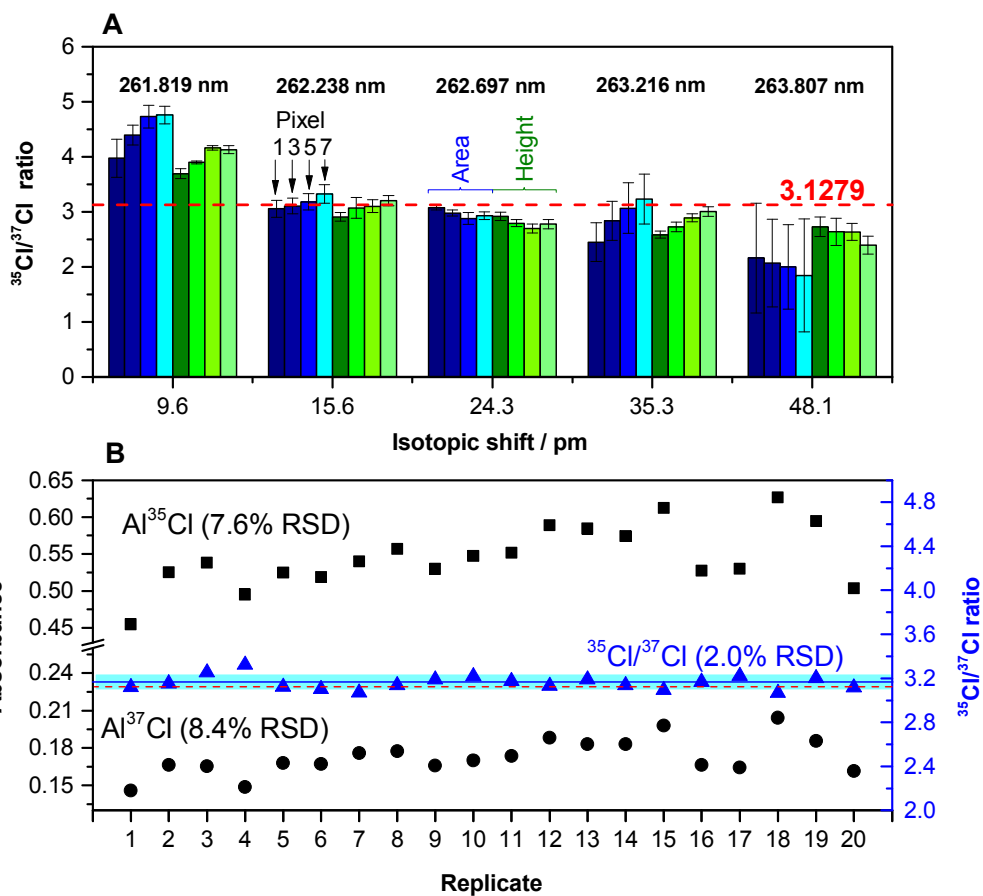


Figure 4

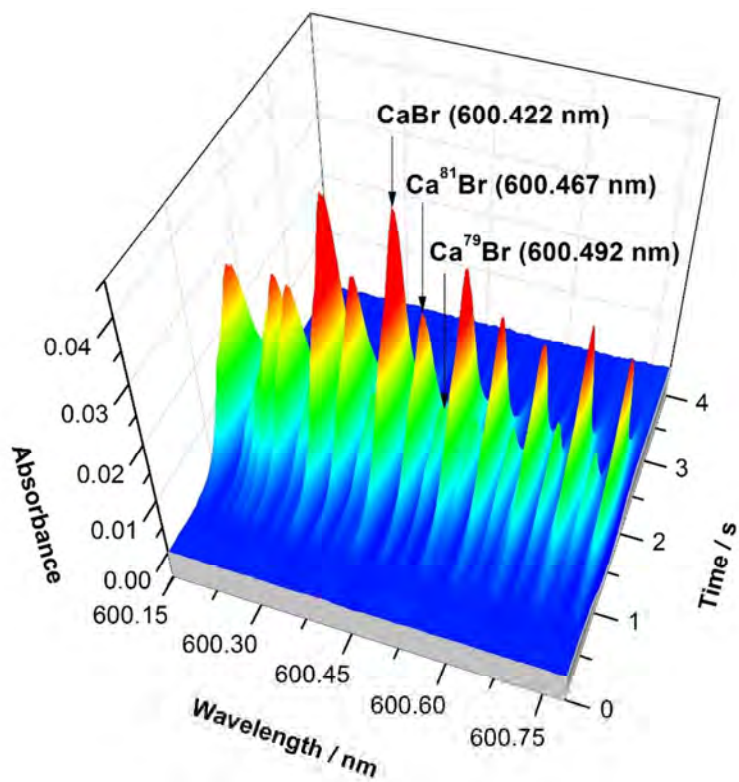


Figure 5

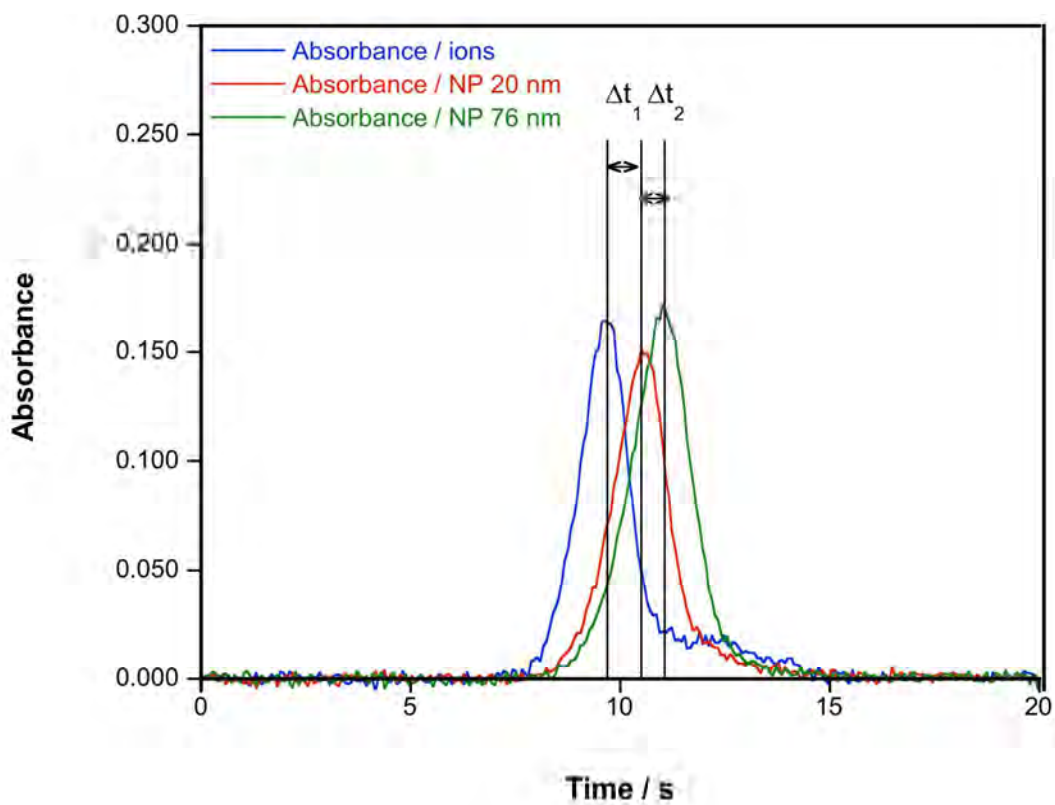


Figure 6

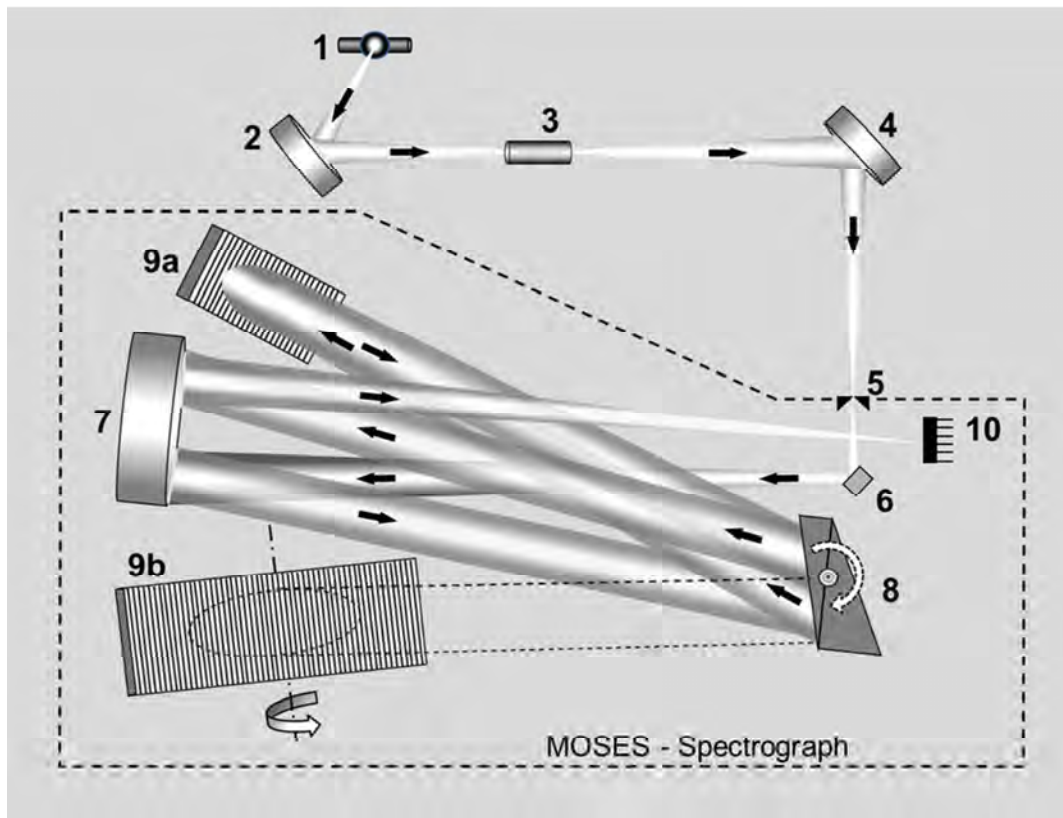


Figure 7

