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# **Quantification of levoglucosan and its isomers by High Performance Liquid Chromatography – Electrospray Ionization tandem Mass Spectrometry and its applications to atmospheric and soil samples**

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## Abstract

The determination of atmospheric concentrations of levoglucosan and its two isomers, unambiguous tracers of biomass burning emissions, became even more important with the development of wood as renewable energy for domestic heating. Many researches demonstrated the increase during recent years of atmospheric particulate matter load due to domestic biomass combustion in developed countries. Analysis of biomass burning tracers is traditionally performed with Gas Chromatography-Mass Spectrometry (GC-MS) technique after derivatization and requires an organic solvent extraction. A simpler and faster technique using Liquid Chromatography – Electrospray Ionisation – tandem Mass Spectrometry (LC-ESI-MS/MS) was optimized for the analysis of levoglucosan, mannosan and galactosan isomers after an aqueous extraction. This technique allows a good separation between the three compounds in a very reduced time (runtime ~5 min). LOD and LOQ of this method are  $30 \mu\text{g l}^{-1}$  and  $100 \mu\text{g l}^{-1}$  respectively, allowing the use of filters from low-volume sampler (as commonly used in routine campaigns). A comparison of simultaneous levoglucosan measurements by GC-MS and LC-ESI-MS/MS for about 50 samples coming from different types of sampling sites and seasons was realized and shows very good agreement between the two methods. Therefore LC-ESI-MS/MS method can be used as an alternative to GC-MS particularly for measurement campaigns in routine where analysis time is important and detection limit is reduced. This paper shows that this method is also applicable to other environmental sample types like soil.

## 1 Introduction

A growing number of scientific studies have recently focused on the apportionment of biomass burning emissions in ambient aerosol (Zheng et al., 2002; Puxbaum et al., 2007; Gaeggeler et al., 2008; Caseiro et al., 2009). This primary source emits high amounts of organic aerosol (OA) and can largely contribute to the OA mass of particulate matter (PM) in winter. For example in Europe, biomass burning contributions to OA

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are 20, 30, 35, 40 and 68 % in Vienna (Austria) (Caseiro et al., 2009), Oslo (Norway) (Yttri et al., 2009), Ghent (Belgium) (Zdrahal et al., 2002), Zürich (Switzerland) (Szidat et al., 2006) and Grenoble (France) (Favez et al., 2010) respectively. Contributions of this source to total PM mass in winter are about 20 % in Paris (France) (Favez et al., 2009) and 42 % in Grenoble (France) (Favez et al., 2010). Better source apportionment studies, especially addressing biomass burning contributions, will be mandatory in the near future in order to respect tougher European Union regulations of the aerosol mass (EU-Directive 2008/50/CE).

OA emitted by biomass burning is particularly rich in carcinogenic compounds, such as polycyclic aromatic hydrocarbons (Simoneit, 2002 and references therein). Among the myriad of molecular compounds emitted by biomass burning, the three isomeric anhydrous sugars levoglucosan (1,6-anhydro- $\beta$ -D-glucopyranose), mannosan, and galactosan, formed during pyrolysis of cellulose, are the predominant organic species (Simoneit et al., 1999). Levoglucosan is the most abundant anhydrous sugar among the monosaccharide anhydrides (Simoneit et al., 1999). In addition, levoglucosan is reasonably stable in the atmosphere (Fraser and Lakshmanan, 2000) and is specific to the burning of cellulose. So, it is considered since the 1980s as a key marker for the apportionment of biomass burning emissions (Hornig, 1985; Locker, 1988) and is largely used in CMB modelling (Fraser and Lakshmanan, 2000). Further, the ratios of levoglucosan-to-mannosan and levoglucosan-to-galactosan are somewhat specific of wood types, allowing the differentiation between hardwood and softwood combustion (Schmidl et al., 2008). For instance, levoglucosan-to-mannosan ratio is about 17 for American beech combustion and about 4 for White spruce combustion (Fine et al., 2004). So, the simultaneous analysis of the three monosaccharides is an important issue for biomass burning study, notably for the choice of wood burned profile in source apportionment models.

Very few studies deal with monosaccharide anhydrides in environmental compartments other than in the atmosphere: Simoneit et al. (2004) and Otto et al. (2006) studied soil samples, Schkolnik et al. (2005) looked at rainwater and Fabbri et al. (2008)

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focused on lignites. However, these studies seem to indicate that monosaccharide anhydrides could be used as proxies for the detection of the impact of biomass burning events in many types of matrices.

Analysis of molecular markers is traditionally performed using Gas Chromatography-Mass Spectrometry (GC-MS) technique after organic solvent extraction and derivatization steps (Bergauff et al., 2008). Widely used for the chemical characterization of atmospheric aerosol, this method is also used for the analysis of soil samples (Otto et al., 2006; Simoneit et al., 2004). Even though the reliability of this approach is demonstrated in several studies, it requires intensive and expensive sample preparation. In addition, the derivatization step usually based on a silylation reaction prevents the analysis of aqueous samples. Recently, other analytical methods without derivatization step were developed for monosaccharide anhydrides quantification using High Performance Liquid Chromatography (HPLC) coupled with various detectors, including pulsed amperometric detection (PAD) (Engling et al., 2006; Caseiro et al., 2007), aerosol charge detection (ACD) (Dixon and Baltzell, 2006), mass spectrometry (MS) (Dye and Yttri, 2005; Wan and Yu, 2006; Gambaro et al., 2008; Saarnio et al., 2010) and Electrospray Ionisation-tandem Mass Spectrometry (ESI-MS/MS) (Palma et al., 2004). Ion-exclusion chromatography coupled with a spectroscopic detection was also used to analyse directly rainwater (Schkolnik et al., 2005) and microchip capillary electrophoresis coupled with pulsed amperometric detection allows a quick analysis of levoglucosan (Garcia et al., 2004). However, only few papers have compared the analytical performance of these methods with the more widely used GC-MS technique (Engling et al., 2006; Schkolnik et al., 2005).

In this study, we present a new method based on coupling HPLC and Electrospray Ionisation-tandem Mass Spectrometry (ESI-MS/MS), which provides an appropriate separation of the monosaccharide anhydride isomers, a sensitive detection, and a fast and inexpensive analysis. Tandem Mass Spectrometry allows a better selectivity of compounds by selecting daughter ion characteristics of the studied compounds (levoglucosan and its isomers).

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This method allows the analysis in the aqueous phase and is therefore applicable to a wide variety of environmental samples including atmospheric aerosol, soil and water (rain, snow, ice) samples. Atmospheric samples from different sites and seasons were simultaneously analyzed with this direct method and with the derivatization-GC-MS method in order to compare their analytical performances. The application of the HPLC-ESI-MS/MS method (called LC-MS) to levoglucosan quantification in soil sample is also presented.

## 2 Material and methods

### 2.1 Reagents and materials

Authentic standards used in this study include levoglucosan (1,6-anhydro- $\beta$ -D-glucopyranose) 99.0% (CAS 498-07-7, Sigma-Aldrich, Steinheim, Germany), mannan (1,6-anhydro- $\beta$ -D-mannopyranose) (CAS 14168-65-1, Carbosynth, Compton, U.K.) and galactosan (1,6-anhydro- $\beta$ -D-galactopyranose) (CAS 644-76-8, TRC, Toronto, Canada). Standard solutions, sample extraction, and mobile phase solutions were prepared with ultrapure water (Purelab Ultra system, Elga, High Wycombe, UK). Stock solutions at  $10\text{ g l}^{-1}$  were prepared by dissolving 1.00 g of each compound in 100 ml of ultrapure water. These solutions were stored in amber glass bottles (SCHOTT® Duran®) at  $4^{\circ}\text{C}$ . Sodium hydroxide solutions for the mobile phase were prepared from a 50% (w/w) NaOH solution (J.T. Baker). Ultrapure water was degassed with He before NaOH addition in order to limit carbonate formation.

### 2.2 Sample collection and LC-MS extraction

Atmospheric particulate matter of less than  $10\text{ }\mu\text{m}$  and  $2.5\text{ }\mu\text{m}$  diameter ( $\text{PM}_{10}$ , and  $\text{PM}_{2.5}$ , respectively) were collected onto QM-A quartz fiber filters (Whatman) in a high-volume sampler (flow rate  $30\text{ m}^3\text{ h}^{-1}$ ) with collection times of 12 or 24 h. Samples were

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collected in two urban background sites in France: “Les Frênes” in Grenoble and “Cinq Avenues” in Marseille, during autumn to winter 2009 and summer 2008, respectively, during the FORMES program (Favez, 2010; El Haddad, 2011). After collection, samples were stored at  $-20^{\circ}\text{C}$ , with filters sealed in aluminum foil and polyethylene bags.

5 Soil samples were collected in the top soil horizon (between two and five cm depth) located under a charcoal burning two days after the end of the combustion, in the karstic Vercors massif (French Alps). After collection, they were air dried at room temperature and sieved at 2 mm.

Appropriate atmospheric sample fractions ( $3$  to  $12\text{ cm}^2$ ) and soil sample fractions 10 of  $5\text{ g}$  were extracted with  $15\text{ ml}$  and  $5\text{ ml}$ , respectively, of ultrapure water with a vortex agitation during  $20\text{ min}$ . Longer agitation and ultrasonic agitation were also tested. In order to evaluate extraction recoveries of the two extraction methods (ultrasonic or vortex agitation), blank Whatman QM-A filters were spiked in triplicate with a standard solution containing the three monosaccharides in aqueous solvent at low, medium and 15 high concentrations ( $100$ ,  $500$ , and  $1000\text{ }\mu\text{g l}^{-1}$ ). They were air dried at room temperature in order to evaporate the aqueous solvent. The results are discussed in Sect. 3.1.

Just before the analysis, extracts were further filtered using Acrodisc® filters (Pall, Gelmann) with a porosity of  $0.22\text{ }\mu\text{m}$  previously rinsed with  $40\text{ ml}$  of ultrapure water. Soil samples extracts were previously filtered using pleated filter cellulose paper.

### 20 2.3 LC-MS analysis

Sample was analyzed using High Performance Liquid Chromatography – Electrospray Ionisation – tandem mass spectrometry (HPLC-ESI-MS2) like presented by Piot et al. (2009). Liquid chromatography is performed with a Dionex pump (model DX500) mounted with Peek and vacuum degasser. Sample is injected by a  $449\text{ }\mu\text{l}$  injection 25 loop. The separation is carried out at room temperature (about  $20^{\circ}\text{C}$ ) using a Carbobac PA–1 anion-exchange analytical column ( $250\text{ mm} \times 4\text{ mm}$ , Dionex) coupled with a Carbobac PA–1 guard column ( $50\text{ mm} \times 4\text{ mm}$ , Dionex) like in Caseiro et al. (2007). Elution is achieved in isocratic mode at  $1.2\text{ ml min}^{-1}$  with  $0.5\text{ mM}$  sodium hydroxide solution.

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Columns are flushed and equilibrated between two samples with an elution gradient between  $0.5$  and  $3\text{ mM}$  sodium hydroxide at a  $1.2\text{ ml min}^{-1}$  flow rate (run time:  $9\text{ min}$ ). During this step, the mobile phase is not injected into the MS. Columns are washed overnight (after approximately 20 samples) with an elution gradient between  $0.5$  and 5  $200\text{ mM}$  sodium hydroxide at a  $0.5\text{ ml min}^{-1}$  flow rate (run time:  $15\text{ h}$ ).

A micrometric split valve is used to reduce the flow injected to the MS at  $0.8\text{ ml min}^{-1}$ . The analytical detector is an Electrospray Ionization Ion Trap MS (LCQ Fleet MS, Thermo Fisher Scientific). Detection is achieved in the negative ion mode like in Gambaro et al. (2008) with a  $m/z$  161 trap isolation. Parameters are optimized for the best 10 Collision Induced Dissociation (CID) efficiency with selective current in  $m/z$  101 and  $m/z$  113, characteristic of daughter ions of levoglucosan and its two isomers. Instrumental conditions are reported in Table 1. Chromatogram integration is realized on the selective current:  $m/z$   $101 \pm 0.5 + 113 \pm 0.5$ .

Calibration is performed twice, at the beginning of the analysis sequence and at the 15 end of the sequence, with standard solutions containing the three monosaccharides at  $100$ ,  $500$ , and  $1000\text{ }\mu\text{g l}^{-1}$ . Samples and standard solutions are injected twice for each analysis.

### 2.4 GC-MS analysis

Standards and atmospheric samples are simultaneous analyzed by GC-MS as 20 described in El Haddad et al. (2009). Authentic standard solutions were prepared in acetone and stored at  $4^{\circ}\text{C}$ . Briefly, sample fractions are extracted with a dichloromethane/acetone mix ( $1:1\text{ v/v}$ ) using an Accelerated Solvent Extractor (ASE 200, Dionex) and reduced to a volume of  $1\text{ ml}$ . A  $100\text{ }\mu\text{l}$  extract fraction is trimethylsilylated with  $100\text{ }\mu\text{l}$  of  $N,O$ -bis(trimethylsilyl)-trifluoroacetamide (BSTFA) containing  $1\%$  25 trimethylchlorosilane (TMCS) for two hours at  $50^{\circ}\text{C}$  before GC-MS analysis. Quantification is performed using selected ion current peak areas ( $204$  for levoglucosan and mannosan and  $217$  for galactosan) and calibration curves are established with authentic standards and a deuterated levoglucosan internal standard. Calibration was

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checked every 10 samples and is performed with 8 levels of concentration between 2 and 400 mg l<sup>-1</sup>.

### 3 Results and discussion

#### 3.1 Methods performance

5 The elution conditions used in the LC-MS method allow the detection of the three monosaccharides in less than 6 min with a very good separation (Fig. 1a). The levoglucosan retention time is about 2.3 min followed successively by mannosan, and galactosan. The chromatogram shows a high resolution ( $R_s$ : peak resolution) between the three peaks ( $R_s = 1.25$  for levoglucosan and mannosan and  $R_s = 1.65$  for mannosan and galactosan) in a very reduced time (runtime ~5 min). However, this method  
10 allows only the analysis of levoglucosan and its two isomers. All analytical performance and linear regression parameters for LC-MS calibration are presented in Table 2. Limit of detection (LOD) of the analytical method presented in this paper (3 times the standard deviation of the blank) is 30 µg l<sup>-1</sup> and the limit of quantification (LOQ) (10 times  
15 the standard deviation of the blank) is 100 µg l<sup>-1</sup> (Table 2). The analytical concentration range was 20 to 2000 µg l<sup>-1</sup>.

Calibration curves systematically show  $R^2$ -values above 0.996 for the three compounds. Analytical reproducibility, evaluated by the relative standard deviation (RSD) between five successive injections of the same standard solution at concentrations of  
20 500 and 1000 µg l<sup>-1</sup>, ranges between 5 to 10 %. In conditions of extraction allowing two injections and the analysis of a sample, mass LOD is 60 ng.

The analysis of levoglucosan by GC-MS is traditionally conducted after an organic solvent extraction using dichloromethane (Simoneit et al., 1999) or mixture of dichloromethane and methanol (Simoneit, 2002). In the case of HPLC analysis, some  
25 studies used a water extraction assisted by ultrasonic or short vortex agitation to extract saccharides because of their high solubility in water (Caseiro et al., 2007; Engling

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et al., 2006; Schkolnik et al., 2005). In this study, the efficiency of these two water extraction procedures was tested. Blank filters were spiked in triplicate (100 µl or 500 µl) with three standard aqueous solutions (more or less concentrated) containing the three monosaccharides at low, medium and high concentrations (representing 0.5, 2.5, and  
5 5 µg of each compound respectively), in order to cover the whole calibration range. After drying at room temperature, those filters were then extracted with 5 ml of aqueous solvent. Extraction was tested both by 20 min ultrasonic agitation and by 20 min short vortex agitation. With the latter, the average recoveries were 90 ± 9 %, 88 ± 28 %, and 99 ± 9 % for levoglucosan, mannosan, and galactosan respectively. Average ultrasonic  
10 agitation recovery was 13 ± 5 % lower than for the short vortex agitation method, for the three monosaccharides. Performance is not improved by longer short vortex agitation but extraction time seems to be important for ultrasonic agitation. Caseiro et al. (2007) showed the best reproducibility (100 ± 8 %) for an extraction time of 45 min. Therefore, all further work was performed with short vortex agitation and all results for levoglucosan were corrected using an average extraction efficiency of 92 %. In addition, with  
15 this method, the minimum solvent extraction volume is about 2 ml allowing the filtration step and two successive LC-MS analyses of the sample. In these conditions the maximum extracted filter fraction is 4.5 cm<sup>2</sup> representing 21 m<sup>3</sup> of collected air for a sampling at 30 m<sup>3</sup> h<sup>-1</sup> during 24 h onto a 150 mm quartz fiber filters.

GC-MS analysis was optimized to quantify about twenty compounds including monosaccharide anhydrides, acids, methoxyphenols and sterols (details in El Haddad et al., 2009). Levoglucosan retention time with GC-MS method (17.98 min) is much longer than with the LC-MS method (Fig. 1b). In our chromatographic conditions, elution of three monosaccharides is in the following order: galactosan, mannosan, and  
25 levoglucosan (Fig. 1b). The analytical concentration range is 100 µg l<sup>-1</sup> to 500 mg l<sup>-1</sup>. Calibration curves for the three monosaccharides show  $R^2$ -values above 0.963 (Table 2). RSD range between 3 % and 5 % for high and low concentrations analysis, respectively. LOD and LOQ of GC-MS analysis is 100 µg l<sup>-1</sup> and 333 µg l<sup>-1</sup>, respectively (Table 2). These concentrations correspond to a mass LOD of 100 ng for the

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extraction and analysis conditions described in the experimental part. The GC-MS analytical performances could be enhanced by increasing the preconcentration of the sample. However, this lowering of the final volume would hamper the successive analyses of derivatized and non derivatized samples to be performed for a full particulate organic matter characterization. In addition, a low volume sample may lead to a loss of reproducibility or performances.

In order to evaluate recoveries by the GC-MS analysis method, filters are spiked in duplicate with 500  $\mu\text{l}$  of levoglucosan standard solutions in acetone at low, medium, and high concentrations. Those concentrations represent 1, 10, and 250  $\mu\text{g}$  of each compound by filter, respectively, after solvent evaporation at room temperature. After ASE extraction the samples were concentrated to 1 ml solution before the derivatization step and the analysis. In these conditions, the recovery for levoglucosan extraction with dichloromethane/acetone mix solvent is  $73 \pm 8\%$ .

LOD and LOQ of both methods are in the same order of magnitude (Table 2) but LC-MS LOD is lower than the GC-MS one's. LC-MS shows better analytical performance for the quantification of the lower levoglucosan concentrations. However, for larger concentrations, this method has a lower reproducibility with a RSD value reaching 10%.

Minimum solvent extraction volume for the GC-MS method is about 60 ml of organic solvent mix that is thirty times more than LC-MS method (extraction with a minimum of 2 ml of aqueous solvent). In addition, LC-MS method uses aqueous solvent, thus minimizing the waste management of the analysis. Another advantage of the aqueous extraction is that the same water aliquot may be used for further compounds analysis like that of ions or of other water soluble organic compounds.

In addition, with these optimized extraction conditions ( $4.5\text{ cm}^2$  in 2 ml of solvent) and for a  $720\text{ m}^3$  sampling collected on QM-A quartz fiber filters (impacted surface =  $153.9\text{ cm}^2$ ), LC-MS shows an atmospheric concentration LOD of  $2\text{ ng m}^{-3}$ . For the same extraction surface area and sampling conditions, the GC-MS method has an atmospheric concentration LOD of  $5\text{ ng m}^{-3}$ . LC-MS allows to reach lower levels of atmospheric concentration for the same extracted surfaces of filter.

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### 3.2 Comparison of LC-MS and GC-MS analysis to atmospheric applications

Parallel analyses were conducted by LC-MS and GC-MS methods on the same fifty atmospheric samples (a different fraction of each sample was analyzed with each method). Samples were collected during different seasons between summer 2008 and winter 2009 in two urban background sites located in Marseille and Grenoble, the second and the sixteenth most populated city in France respectively. Sampling of  $360\text{ m}^3$  and  $720\text{ m}^3$  were collected with High-Volume samplers. Concentrations were corrected by extraction efficiencies. Levoglucosan concentrations covered a wide range from 4 to  $3200\text{ ng m}^{-3}$  and concentrations found are in the same range as previous measurements reported in Europe. For instance, Caseiro et al. (2009) measured concentrations ranging from 20 to  $400\text{ ng m}^{-3}$  of levoglucosan in Austrian Regions and Puxbaum et al. (2007) measured 0.3 to  $1651\text{ ng m}^{-3}$  in CARBOSOL sites.

Comparison between the two methods was only made for levoglucosan since concentrations of the other monosaccharide anhydrides (mannosan and galactosan) were lower than the detection limit for too many samples. Results show an excellent agreement between the LC-MS and GC-MS methods, with a slope of almost unity, within the uncertainty of the measurement, and  $R^2$ -values of 0.94 (Fig. 2). This comparison validates the LC-MS method versus the more traditional GC-MS method for the analysis of atmospheric levoglucosan. With a lower detection limit for atmospheric analysis and faster sample treatment, LC-MS method represents a very good alternative to the widely used GC-MS method. With this method, quantification of levoglucosan could be achieved in low-volume sampling conditions and for field campaigns with many samples.

Several such studies are in progress in our labs in different environment type (rural, urban, alpine sites...), including collections with low volume samplers ( $1\text{ m}^3\text{ h}^{-1}$ ) for week-long sampling, and for a year-long survey of eight urban background sites in the Rhône-Alpes Region (Piot, 2011; Piot et al., 2011) where measured levoglucosan concentrations range between  $4\text{ ng m}^{-3}$  (in summer) and  $1000\text{ ng m}^{-3}$  (in winter).

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### 3.3 Other environmental samples analysis

The use of aqueous solvent for levoglucosan extraction in the LC-MS method allows to consider the analysis of monosaccharide anhydrides in many environmental matrices. Levoglucosan and its two isomers were analyzed by the LC-MS method in soil samples collected under wood fire combustion (2 to 5 cm depth) two days after the end of a combustion performed to produce charcoal. Extraction with Soxhlet and dichloromethane, and analyses using GC-MS were performed but no monosaccharide anhydrides were observed in these analytical conditions. Water extraction (5 g of soil extract with 5.0 ml of water during 20 min of short vortex agitation) was undertaken and followed by LC-MS analysis. In these conditions, concentrations of 10.0, 1.5 and 0.6  $\mu\text{g g}^{-1}$ , were measured for levoglucosan, mannosan and galactosan, respectively, highlighting a noteworthy impact of fire combustion on soil. Otto et al. (2006) have analyzed charred pine forest surface soil samples in Canada by GC-MS after organic solvent extraction and have measured levoglucosan, mannosan and galactosan concentrations of 1.0, 0.6, and 0.3  $\mu\text{g g}^{-1}$ , respectively. Simoneit et al. (2004) measured levoglucosan concentrations of less than 0.1  $\mu\text{g g}^{-1}$  in soil or soil dust weakly impacted by biomass burning. Thus, data reported in the literature are much lower than concentrations measured in this study. This may be related to the type of soil samples, or maybe due to a better efficiency of water extraction than organic solvent extraction for soil samples. Additional tests would be necessary to compare aqueous and organic solvent extraction methods but test samples of soil with certified levoglucosan concentrations do not exist in order to quantify the extraction efficiency. However, the analysis of levoglucosan in soil, easily achievable with the LC-MS method with a low detection limit, is a promising way that can allow to evaluate the impact of forest fires in such environmental archives.

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## 4 Conclusions

Levoglucosan concentrations of atmospheric samples obtained with two independent methods (LC-MS and GC-MS) were compared and present extremely good correlation for a wide range of concentrations. This shows the validity of the LC-MS measurements. Whereas the GC-MS allows the detection of a large number of compounds and can handle large atmospheric concentration range, the LC-MS method allows only to measure water-soluble compounds like levoglucosan. Nevertheless, analytical performances are better for the LC-MS method (lower LOD, better recovery) than for the GC-MS method. Moreover one of the main advantages of the LC-MS method is its rapidity, allowing the processing of large sets of samples in order to obtain data for this biomass burning marker in large field campaigns. In fact, LC-MS allows the analysis of monosaccharide anhydrides in less than five min with a shorter time of sample preparation using a cheaper and very simple extraction technique with less impact on the environment. Finally, the extraction method used in LC-MS can be applied to many environmental types, as for example soil whose moisture does not allow organic solvent extraction.

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**Table 1.** Instrumental conditions.

Spray voltage (kV)	6.44
Spray current ( $\mu$ A)	4.46
Sheath gas flow rate	40.84
Auxillary gas flow rate	21.32
Sweep gas flow rate	12.03
Capillary temperature ( $^{\circ}$ C)	310.07

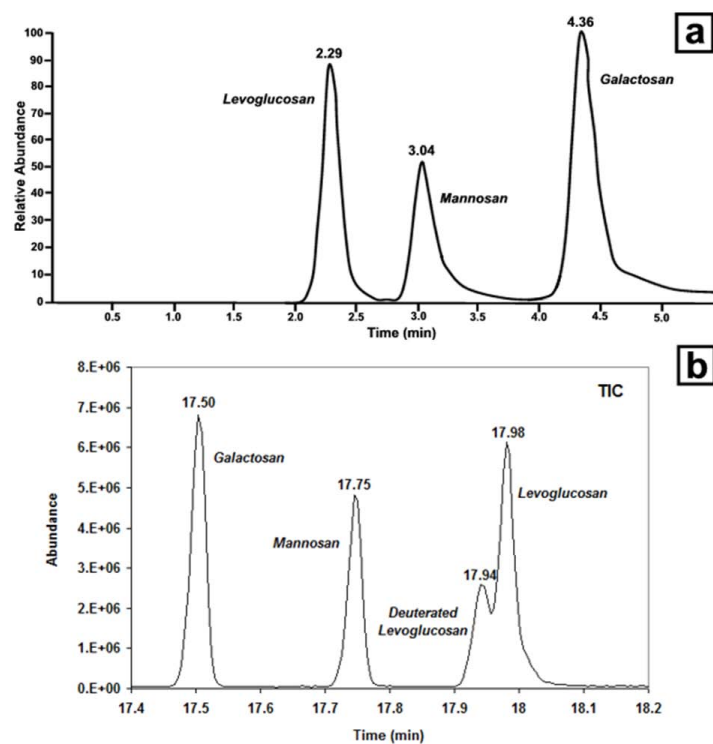
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**Table 2.** Analytical performances and linear regression parameters of levoglucosan (same performances for mannosan and galactosan).

	LC-MS	GC-MS
LOD <sup>a</sup> ( $\mu$ g l <sup>-1</sup> )	30	100
Masse LOD ( $\mu$ g)	0.06	0.1
LOQ <sup>b</sup> ( $\mu$ g l <sup>-1</sup> )	100	333
Analytical concentration range ( $\mu$ g l <sup>-1</sup> )	20–200	100–5.10 <sup>5</sup>
RSD for high concentration <sup>c</sup> (%)	10	3
RSD for low concentration <sup>c</sup> (%)	5	5
$R^2$	0.996	0.963

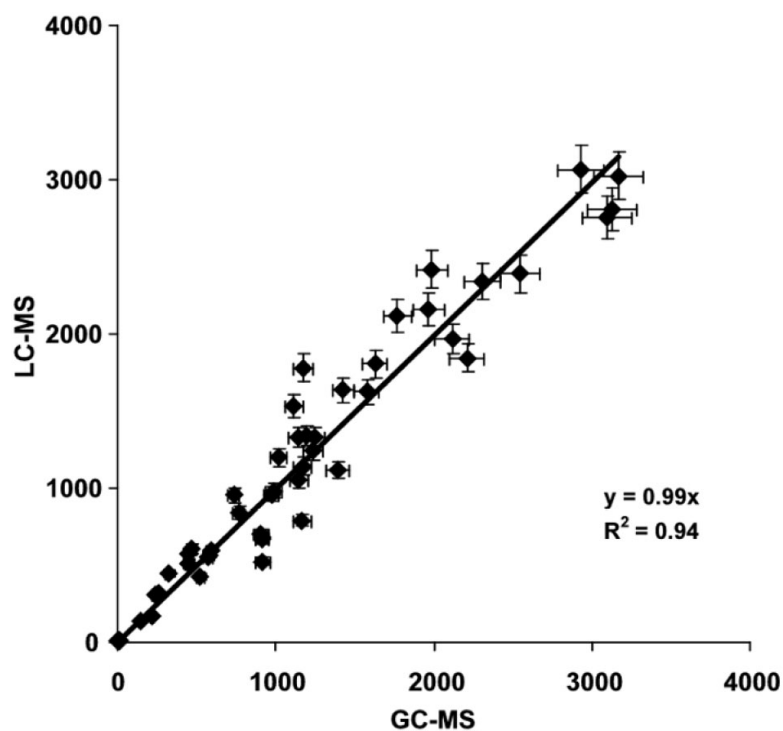
<sup>a</sup> 3  $\times$  standard deviation of the blank<sup>b</sup> 10  $\times$  standard deviation of the blank<sup>c</sup> successively 5 injections of standard solution.

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**Fig. 1.** LC-MS chromatogram (a) and GC-MS chromatogram on total ion current (TIC) (b).

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**Fig. 2.** Correlation between LC-MS results and GC-MS results for levoglucosan (50 samples analysed). Concentrations are in  $\text{ng m}^{-3}$  in air. Error bars represent the RSD.

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