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**Journal of The American Society for
Mass Spectrometry**

The official journal of The American
Society for Mass Spectrometry

ISSN 1044-0305

J. Am. Soc. Mass Spectrom.
DOI 10.1007/s13361-012-0488-6



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RESEARCH ARTICLE

Mass Spectrometry Parameters Optimization for the 46 Multiclass Pesticides Determination in Strawberries with Gas Chromatography Ion-Trap Tandem Mass Spectrometry

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Abstract

Multiclass analysis method was optimized in order to analyze pesticides traces by gas chromatography with ion-trap and tandem mass spectrometry (GC-MS/MS). The influence of some analytical parameters on pesticide signal response was explored. Five ion trap mass spectrometry (IT-MS) operating parameters, including isolation time (IT), excitation voltage (EV), excitation time (ET), maximum excitation energy or “*q*” value (*q*), and isolation mass window (IMW) were numerically tested in order to maximize the instrument analytical signal response. For this, multiple linear regression was used in data analysis to evaluate the influence of the five parameters on the analytical response in the ion trap mass spectrometer and to predict its response. The assessment of the five parameters based on the regression equations substantially increased the sensitivity of IT-MS/MS in the MS/MS mode. The results obtained show that for most of the pesticides, these parameters have a strong influence on both signal response and detection limit. Using the optimized method, a multiclass pesticide analysis was performed for 46 pesticides in a strawberry matrix. Levels higher than the limit established for strawberries by the European Union were found in some samples.

Key words: Optimization parameters, Multiclass pesticides, GC-MS/MS, Ion trap, Strawberries

Introduction

Pesticides are used to protect crops before and after harvest from infestation by pests and plant diseases. A

consequence of their use may be the presence of pesticide residues in treated products, fruits, vegetables, grains, and other commodities. Even after being washed, stored, processed, and prepared, some pesticide residues may remain in both fresh products and processed foods. Therefore, multiresidual methodologies that can determine a large number of pesticides simultaneously with satisfactory sensitivity and selectivity are highly required [1]. Stringent international food safety standards require monitoring some chemicals in a variety of food samples

Electronic supplementary material The online version of this article (doi:10.1007/s13361-012-0488-6) contains supplementary material, which is available to authorized users.

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at trace level with the lowest maximum residue limit (MRL) being set at 10 ng/g (UE) [2, 3]. Gas chromatography coupled to an electron capture detector (GC-ECD), a nitrogen phosphorus detector, and different other detectors are still widely used as analytical technique for the trace analysis of pesticides in various environmental and food matrices with high sensitivity [4–13]. Nowadays, the use of mass spectrometry (MS) detection for a proper and accurate analyte identification together with the electron ionization (EI) mode and the GC retention time information is essential [2, 14–16]. Ion trap (IT) GC-MS has the potential to identify analytes at trace levels and to avoid the influence of matrix components as well as allows for selective analysis by MS/MS. This could be achieved by collision-induced dissociation (CID) of a selected unique precursor ion that produces ions in sufficient abundance ratios specific to the detection of a particular molecule [2]. In order to attain high sensitivity to achieve low detection limits, the instrumental parameters of IT-MS affecting the performance of this system must be thoroughly optimized [14]. Although parameters for some compounds are available in MS/MS libraries, the optimization of GC-MS/MS parameters is indispensable in order to run with best efficiency [17–20]. There are a few papers describing the optimization and the best parameters for each analyte in a given apparatus [2]. The optimization of GC-MS/MS system requires assessing IT-MS parameters influence by the approach of changing one-factor-at-a-time (OFAT) [2]. This approach does not give information on interactions between factors, so it can miss the optimal settings when interactions do occur. Statistically designed experiments such as Plackett–Burman and central composite designs can help to optimize analytical parameters much more efficiently and in less experimental runs [19]. Optimization of MS/MS parameters is a hard work for each analyte; hence, knowledge of optimal values will save considerable time in analytical method development.

The aim of the present study was to establish an overall analytical method and optimization of a set of instrumental parameters in order to attain the highest possible sensitivity for pesticides determination. Particular attention was paid to the optimization of five IT-MS parameters, namely, the duration of the ion isolation waveform voltage [isolation time (IT)], the duration of the ion excitation [excitation time (ET)], the mass range window around the ion of interest [isolation mass window (IMW)], excitation voltage (EV) and the maximum excitation energy (q), which is defined as the amount of energy that holds a precursor ion in the ion trap during excitation.

In this work a visual basic for macro applications in Microsoft Excel was created to compute the signal response based on the best regression model for all possible combinations among the five parameters in a total of 2160 arrangements and save the maximum value. Bearing all this in mind, a GC-MS/MS based multiresidue analysis method

was developed covering 46 pesticides, including pyrethroids, organophosphorus, carbamates, strobilurines, pyrimidines, thiophthalimides, dinitrophenols derivative, azoles, anilides, dicarboximides, anilinopyrimidines, aryloxyphenoxy propionic acids, 2,6-dinitroanilines, triazines and a set of persistent ones such as organochlorines. The optimized method was then used in the screening program and the levels of these pesticides in strawberries are reported in this paper.

Materials and Methods

Chemicals and Preparation of Standard Solutions

Reference standards were purchased from Sigma Aldrich (Steinheim, Germany), Riedel-de Haen (Seelze, Germany), and Chem Service (West Chester, PA, USA). Stock standard solutions (approximately 2000 $\mu\text{g}\cdot\text{L}^{-1}$) were prepared by dissolving reference standards in n-hexane and methanol and were stored in a freezer at 4 °C. Working pesticide standard mixtures were prepared by dilution of stock solutions in n-hexane. n-Hexane and methanol were chromatographic grade and were supplied by Merck (Darmstadt, Germany).

Apparatus

GC-MS/MS instrument, TRACE GC Ultra (Thermo Fisher Scientific, Austin, TX, USA) gas chromatograph coupled with a Polaris Q ion trap mass spectrometer was used. The system included an AS-3000 autosampler. A ZB-XLB capillary column from Phenomenex (30 m \times 0.25 mm \times 0.25 μm) was used for chromatographic separation. The system was controlled by Xcalibur software, ver. 1.3. The interface line and ion source temperatures were maintained at 250 °C in all experiments. Injections (2 μL) were done in the splitless mode. MS/MS conditions such as isolation (wideband application (IMW), isolation time (IT)), fragmentation (excitation time (ET) and voltage (EV) and factor “ q ” were optimized for each analyte, beginning with the following base conditions: factor $q=0.45$, $EV=1$ V, $IT=12$ ms, $ET=15$ ms, $IMW=1$ and carrier flow 1.3 $\text{mL}\cdot\text{min}^{-1}$.

Gas Chromatograph Conditions

The column oven temperature was programmed as follows: initial temperature 40 °C (held for 1 min), increased by 30 °C/min to 220 °C (held for 5 min), increased by 10 °C/min to 250 °C and held at this temperature for 20 min and finally increased again by 5 °C/min to 285 °C and held at this temperature for 5 min. The mass spectrometer was operated in electron ionization (EI) mode at 70 eV with an external ionization source. The inlet temperature was 240 °C and helium (purity ≥ 99.999 %) was used as carrier gas at 1 mL/min and the injection volume was 2 μL . The ion source

temperature was 250 °C and the electron multiplier was operated at 2100 V (autotune to gain of 1×10^7).

Limit of Detection

The limit of detection (LOD), also defined as the lowest concentration that the analytical process can differentiate from background levels, was estimated for a signal-to-noise ratio (S/N) of three from the chromatograms analysis.

Multiple Linear Regression Analysis

For each pesticide, multiple linear regression tests were conducted in order to estimate the relationship between the statistically significant parameters of IT-MS/MS detection and the instrumental signal response by fitting a linear equation to observed data.

Macro Edition in Microsoft Excel

A visual basic for macro applications (VBA) was developed in Microsoft Excel in order to assess the maximum of $8 \text{ ET} \times 6 \text{ IT} \times 5 \text{ EV} \times 3 \text{ q} \times 3 \text{ IMW}$ combinations, available in the IT-MS/MS detector, based on the predicted value determined by the multiple regression equation.

Sampling

Strawberries were collected in the first week of May in 2009 and 2010 from crops in the center of Portugal. Different varieties of strawberries were collected including Siba, Camarosa, Festival, and Albion. Samples determined to be without any target analytes were used as blank strawberry samples. The samples were analyzed following the procedure described below.

Extraction and Cleanup Procedures The selected QuEChERS (EUMIV50CT-VP) and cleanup (CUMPS15C18CT) were obtained from UCT (Bristol, PA, USA). For the initial extraction step, an amount (10 g) of chopped strawberries was weighted into a 50 mL centrifuge tube and 50 μL of internal standard (IS) solution was added. The strawberry sample was left during 30 min at room temperature to let the *n*-hexane evaporate before the addition of 10 mL of acetonitrile.

The resulting solution was shaken for 1 min followed by the addition of 6 g anhydrous magnesium sulfate, 1.5 g sodium chloride, 1.5 g trisodium citrate dihydrate, and 0.75 g disodium hydrogenocitrate sesquihydrate. The centrifuge tube was capped and shaken vigorously for 1 min to prevent salt agglomeration before centrifugation at 3000 rpm for 5 min at room temperature. An aliquot of 1.5 mL was sampled from the upper layer into a 2 mL cleanup vial containing 150 mg primary secondary amine (PSA), 150 mg MgSO_4 , and 50 mg C18 and again

vortexed for 1 min and then centrifuged for 5 min at 4000 rpm at room temperature. From the upper layer of the prepared sample, an aliquot of 1.0 mL was transferred into a vial and put under a mild stream of nitrogen to dryness. Finally, 1 mL of *n*-hexane was added to dissolve the residue and then 2 μL of this solution was injected onto the gas chromatograph.

For fortification studies, 10 g of strawberry sample free of detectable pesticides were spiked and homogenized at 50, 200, and 400 $\mu\text{g} \cdot \text{kg}^{-1}$ levels for each pesticide. In the case of organochlorine pesticides 30, 90, and 180 $\mu\text{g} \cdot \text{kg}^{-1}$ levels were used. The mixture was left for 30 min before the beginning of the extraction process. Samples were then prepared according to the procedure aforementioned.

Results and Discussion

The optimization of MS/MS parameters in IT-MS was carried out in four steps: (1) isolation of precursor ion and subsequent product ion selection, (2) screening analysis, (3) multiple linear regression test to check the importance of each parameter on the signal response, fitting data to linear and second order models for the significant parameters and model discrimination, and finally (4) combinatorial optimization based on the best previously chosen model.

Isolation of Precursor Ion and Product Ions Selection

The most abundant ion from the spectra of the different pesticides was selected as precursor ion and it was then isolated in the ion trap and fragmented by collision induced dissociation (CID). The two most abundant product ions, Q_1 and Q_2 [1] were selected and monitored and the spectrum was obtained with the default operating parameters from the GC-MS/MS system ($\text{IT}=12 \text{ ms}$, $\text{ET}=15 \text{ ms}$, $\text{IMW}=1$, $\text{q}=0.45$, $\text{EV}=1.0 \text{ V}$) (Table 1).

Screening Analysis

Different values of EV, ET, IMW, q, and IT were set up by a one-factor-at-a-time (OFAT) method, meaning that when one parameter is changed the others are fixed at their default value. Regarding a multiclass pesticides signal maximization, the search of the best IT-MS parameters is necessary. Thus, the parameters were varied in order to determine the suitable values at which both absolute peak area of the product ions and signal to noise (S/N) ratio were maximized for the 46 pesticides under investigation. The concentration of the standard mixture solution used in this work was $150 \mu\text{g} \cdot \text{L}^{-1}$. The results obtained for the signal to noise ratio by OFAT experiments are shown in Figure 1. An example of OFAT study (HCB) concerning the dependence of q factor on area response is shown in Figure S1 (Supplementary Information). An important

Table 1. Precursor Ions and the Products Ions (Q_1 and Q_2) for the Selected Pesticides

Chemistry class	Pesticides	CAS	MW (g/mol)	Ion precursor	Q_1	Q_2	
Chloroacetanilide Organochlorine	Alachlor	15972-60-8	270	188	160	132	
	Aldrin	309-00-2	365	263	227	193	
	Dieldrin	60-57-1	381	243	211	176	
	Endrin	72-20-8	381	245	209	173	
	HCB	118-74-1	285	284	214	249	
	Lindane	58-89-9	291	183	109	181	
	o,p'-DDT	789-02-6	355	235	165	199	
	p,p'-DDD	72-54-8	320	235	165	199	
	p,p'-DDE	72-55-9	318	318	299	281	
	Methoxychlor	72-43-5	346	227	169	197	
	α,β,δ -HCH	608-73-1	291	183	179	177	
	α -Endosulfan	959-98-8	407	195	191	170	
	β -Endosulfan	33213-65-9	407	195	191	170	
	Triazines	Atrazine desethyl	6190-65-4	188	172	136	145
		Atrazine	1912-24-9	216	200	164	122
Simazine		122-34-9	202	200	172	164	
Strobin	Azoxystrobin	131860-33-8	403	344	327	325	
Pyrethroids	β -Cyfluthrin	68359-37-5	434	163	91	127	
	Bifenthrin	82657-04-3	423	181	165	164	
	Cypermethrin	52315-07-8	416	181	163	152	
	Deltamethrin	52918-63-5	505	172	141	88	
	λ -Cyhalothrin	68085-85-8	450	181	152	141	
	Permethrin	52645-53-1	391	183	165	153	
	Fenvalerate	51630-58-1	420	125	121	132	
	Fenpropathrin	39515-41-8	349	181	152	153	
Thiophthalimide	Captan	133-06-2	301	264	257	254	
	Folpet	133-07-3	297	260	151	189	
Organophosphorus	Chlorpyrifos	2921-88-2	351	197	114	148	
	Malathion	121-75-5	330	173	134	106	
	Diazinon	333-41-5	304	179	167	110	
Pyrimidine	Cyprodinil	121552-61-2	225	224	208	197	
	Bupirimate	121552-61-2	316	208	178	129	
	Mepanipyrim	110235-47-7	223	222	207	221	
	Pyrimethanil	53112-28-0	199	198	183	158	
Unclassified	Dazomet	533-74-4	162	162	89	44	
Anilide	Fenhexamid	126833-17-8	302	177	166	153	
Unclassified	Fludioxinil	131341-86-1	248	248	235	191	
N-methyl Carbamate	Methiocarb	2032-65-7	225	168	153	109	
2,6-Dinitroaniline	Pendimethalin	40487-42-1	281	252	191	162	
Aryloxyphenoxy propionic acid	Quizalofop-p-ethyl	100646-51-3	299	223	151	255	
	Fluazifop-p-butyl	79241-46-6	383	254	249	163	
Azole	Tetraconazole	112281-77-3	372	336	249	105	
	Myclobutanil	88671-89-0	289	152	150	151	
Phenylsulfamide	Tolylfluanid	731-27-1	347	138	137	136	
Dicarboximide	Vinclozolin	50471-44-8	286	212	172	145	
	Procymidone	32809-16-8	284	283	281	282	
	Iprodione	36734-19-7	330	314	271	153	

increase of area was observed when “ q ” factor increases from 0.225 to 0.45 (Figure 1q). This increase is more remarkable for HCB, vinclozolin and bifenthrin. In particular, bifenthrin achieved a LOD of $1.61 \mu\text{g.L}^{-1}$ for $q=0.255$ and a LOD of $0.49 \mu\text{g.L}^{-1}$ for $q=0.45$, whereas HCB attained higher sensitivity as “ q ” rose with a LOD decrease from $11.25 \mu\text{g.L}^{-1}$ to $0.68 \mu\text{g.L}^{-1}$.

Figure 1 (IMW) shows a variation in signal response of HCB, diazinon, pyrimethanil, and tolylfluanid as a result of IMW change.

In the case of IT, the variation in signal response was more pronounced for diazinon, bifenthrin, alachlor, and tolylfluanid [Figure 1(IT)]. The S/N values for pyrimethanil and cyprodinil showed a maximum for IT=12 ms.

As to EV dependence, the pesticides with higher variation in signal to noise value were diazinon, bifenthrin, alachlor, pendimethaline, and tolylfluanid [Figure 1(EV)].

As shown in Figure 1(ET), some pesticides (diazinon, bifenthrin, DDT, pyrimidine pesticides, pendimethaline, tolylfluanid, and myclobutanil) revealed a random variation with respect to ET influence.

Hence, for some pesticides, it seems that better LODs can be achieved by changing the IMW to detect neighboring ions or by varying other parameters of the ion trap (i.e., IT, ET, EV, or q).

This screening analysis proves that the variation of one parameter can mean a change in the response, so the importance of applying statistical studies may allow the prediction of the best parameter conditions.

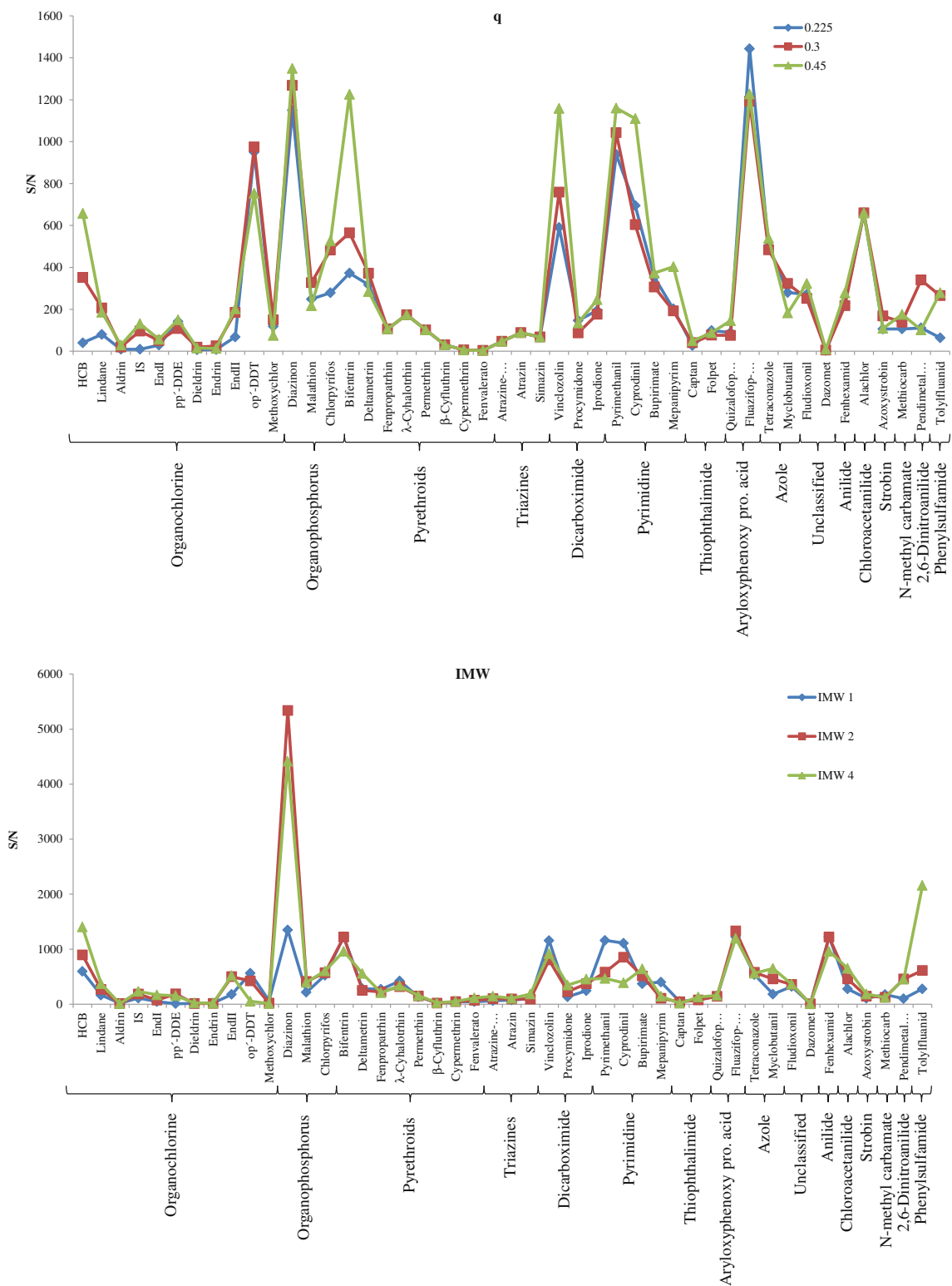


Figure 1. Plot of average signal to noise (S/N) versus, “q” value, IMW, IT, ET, and EV for the 46 multiclass pesticides

Multiple Linear Regression

Multiple linear regression tests were carried out in order to estimate the influence of each parameter up to five

on the IT-MS analytical response. In most cases, the true functional relationship is unknown. This study allowed finding the parameters that have significant

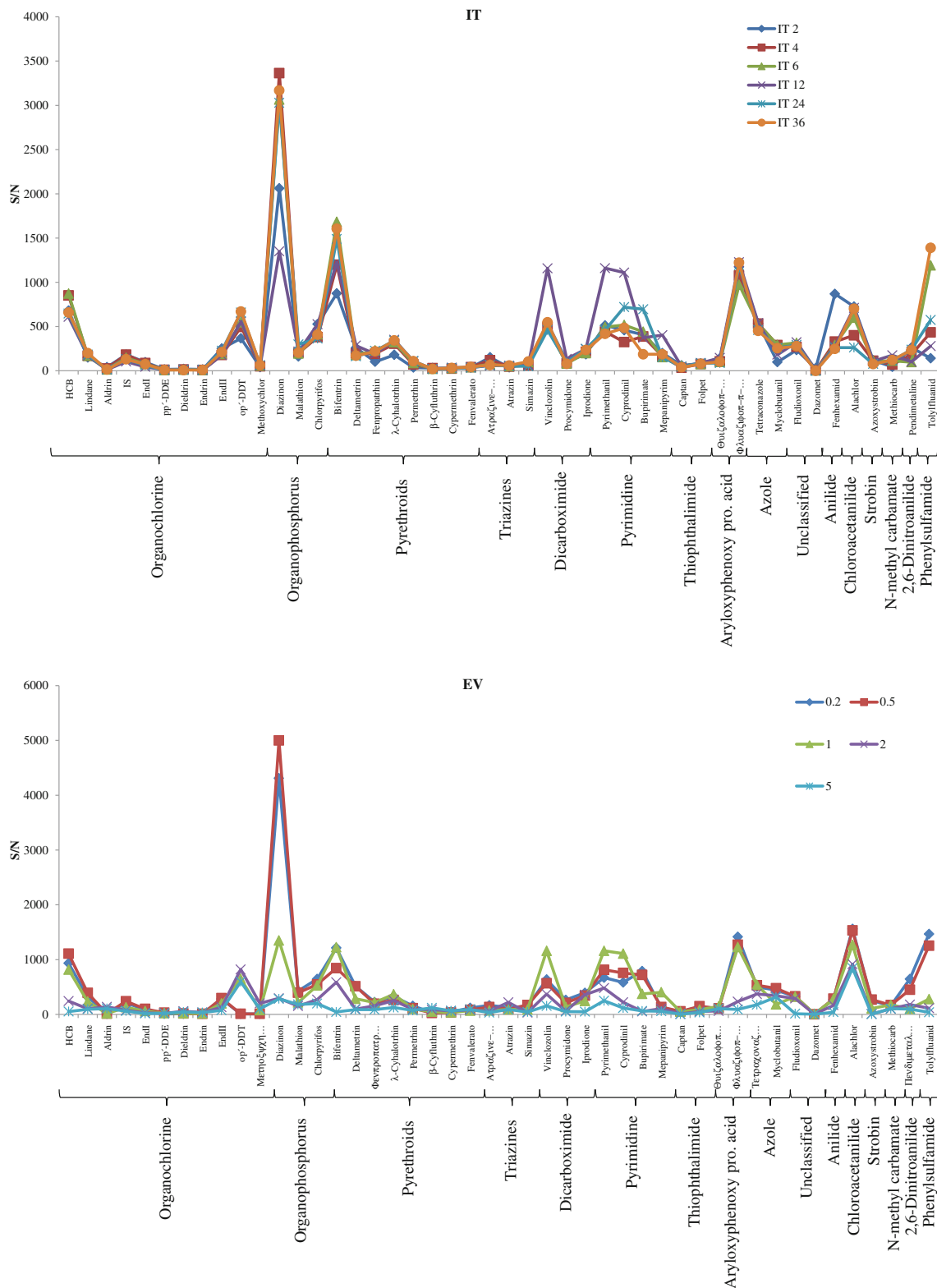


Figure 1. (continued)

influence on signal response. Hence the IMW, IT, ET, EV, and q values were screened in order to discard the irrelevant parameters keeping the remaining for subsequent optimization.

Model Discrimination To evaluate the “best” empirical regression equation, the analytical responses obtained were fitted to linear models that included only first order terms (Equation 1), and both first and second-order (curvature) terms (Equation 2).

Table 2. Coefficient of Determination for First and Second Order and Statistic T Used for Model Discrimination

	Pesticides	R ²		Regression test	
		First order	Second order	T (F _{critical})	
Chloroacetanilide	Alachlor	0.7353	0.9309	14.64 (2.41)	
	Organochlorines	Aldrin	0.5508	0.7658	5.66 (2.36)
		Dieldrin	0.4648	0.8566	16.84 (2.36)
		Endrin	0.5178	0.8789	18.39 (2.36)
	Hexachlorobenzene	0.7701	0.9424	18.45 (2.36)	
		Lindane	0.3551	0.4627	1.23 (2.36)*
		Methoxychlor	0.5623	0.9273	30.95 (2.36)
		o,p'-DDT	0.5349	0.9142	27.26 (2.36)
		p,p'-DDE	0.7250	0.8997	10.74 (2.36)
		α-Endosulfan	0.5335	0.8497	12.97 (2.36)
		β-Endosulfan	0.6853	0.8668	8.40 (2.36)
		IS	0.6132	0.8935	16.23 (2.36)
Triazines	Atrazin	0.5754	0.8770	12.67 (2.41)	
	Atrazine-desethyl	0.3382	0.7134	6.76 (2.41)	
		Simazine	0.6254	0.8886	12.20 (2.41)
Strobin	Azoxystrobin	0.5673	0.7993	5.97 (2.41)	
Pyrethroids	β-Cyfluthrin	0.3970	0.5919	2.47 (2.41)	
		Bifentrin	0.4433	0.6284	2.57 (2.41)
		Cypermethrin	0.6629	0.7695	2.39 (2.41)*
		Deltamethrin	0.4629	0.7100	4.40 (2.41)
		Fenpropathrin	0.7139	0.9092	11.11 (2.41)
		Fenvalerate	0.5954	0.9042	16.66 (2.41)
		λ-Cyhalothrin	0.7051	0.8915	8.88 (2.41)
		Permethrin	0.7417	0.8520	3.85 (2.41)
Pyrimidine	Bupirimate	0.5079	0.8150	9.01 (2.41)	
		Mepanipyrim	0.1960	0.2148	0.12 (2.41)*
		Cyprodinil	0.2138	0.6476	3.20 (2.41)
		Pyrimethanil	0.2931	0.4657	1.67 (2.41)*
Thiophthalimide	Captan	0.3501	0.4213	0.78 (2.41)*	
		Folpet	0.2792	0.5851	3.81 (2.41)
Organophosphorus	Chlorpyrifos	0.8862	0.8283	1.97 (2.41)	
		Diazinon	0.6283	0.8956	13.23 (2.41)
		Malathion	0.3761	0.7714	8.93 (2.41)
Unclassified	Dazomet	0.2796	0.6753	5.41 (2.41)	
		Fludioxinil	0.6123	0.8570	9.56 (2.41)
Anilide	Fenhexamid	0.5615	0.7244	3.05 (2.41)	
Aryloxyphenoxy propionic acid	Fluazifop-p-butyl	0.7402	0.8585	5.05 (2.41)	
	Quizalofop-p-ethyl	0.6995	0.7869	2.12 (2.41)*	
N-methyl carbamate	Methiocarb	0.3700	0.5948	2.87 (2.41)	
Azole	Myclobutanil	0.3235	0.6585	5.07 (2.41)	
		Tetraconazole	0.6504	0.7276	1.46 (2.41)*
2,6-Dinitroaniline	Pendimethalin	0.5366	0.7033	2.90 (2.41)	
Phenylsulfamide	Tolylfluanid	0.5951	0.8505	9.26 (2.41)	
Dicarboximide	Vincllozolin	0.5823	0.6084	0.34 (2.41)*	
		Procymidone	0.6050	0.8808	12.54 (2.41)
		Iprodione	0.4716	0.7527	5.87 (2.41)

*Second order model is not significantly better than first order model

more of the coefficients belonging to the general Equations 1 or 2. The statistics for testing the significance of any individual regression coefficient is:

$$t_0 = \frac{\hat{\beta}_j}{\hat{\sigma}^2 C_{jj}} \quad (5)$$

where C_{jj} is the diagonal element of the matrix variance-covariance $(X^T X)^{-1}$ for the corresponding $\hat{\beta}_j$. The denominator of the Equation 5 is often called the standard error of the regression coefficient $\hat{\beta}_j$, that is:

$$se(\hat{\beta}_j) = \sqrt{\hat{\sigma}^2 C_{jj}} \quad (6)$$

where $\hat{\sigma}$ refers to the standard error of regression calculated as:

$$\hat{\sigma} = \sqrt{\frac{SS_E}{n-k}} \quad (7)$$

The IMW is the mass range window around the ion of interest that is isolated for optimization of MS/MS parameters. The multiple regression test proved that the IMW parameter has significant effect in the area response of 1 chloroacetanilide, 4 organochlorines, 4 pyrethroids, 1 thiophthalimide, 1 unclassified, and 1 aryloxyphenoxy propionic acid pesticides.

The IMW parameter is not significant for triazines, strobil, pyrimidine, *n*-methyl carbamate, azole, 2,6-dinitroanilide, phenylsulfamide, and dicarboximide.

Another pertinent parameter for MS/MS optimization is the maximum excitation energy (*q*), which is defined as the amount of energy that holds a precursor ion in the ion trap during excitation and thus influences the usable mass range. Only three values (instrument characteristics) can be set for “*q*.” Figure S1 shows the chromatogram of HCB for all acceptable *q* values. The area response related to *q*=0.45 is 11.5 times higher than with *q*=0.225. Since higher “*q*” values allow more energy to be deposited in the precursor ion before its dissociation (CID), it can be concluded that these pesticides need more energy for fragmentation. Regarding multiple linear regression test, “*q*” has a significant effect on the signal response for most of the organochlorine pesticides, 4 pyrethroids, 1 organophosphorus and 1 unclassified pesticide.

The isolation time (IT) of an ion is defined as the duration of the ion isolation waveform voltage applied to isolate a selected precursor ion. An initial value of 12 ms was imposed, which was afterward changed for optimization purposes of the S/N ratio of individual compounds. The test on individual regression coefficients showed that more than half of the studied pesticides were affected by the variation of this parameter.

The EV parameter has significant effect in the area response of 1 chloroacetanilide, 10 organochlorines, 3 triazines, 6 pyrethroids, 1 pyrimidine, 1 thiophthalimide, 3 organophosphorus, 1 unclassified, 1*N*-methyl carbamate, 1 azole, 1 dicarboximide, 1 2,6-dinitroaniline, 1 aryloxyphenoxy propionic acid, and 1 phenylsulfamide pesticides.

As can be proven in the multiple linear regression test, the excitation time (ET) parameter has effect on the MS/MS determination of about 2/3 of the pesticides tested.

The results showed that an optimum ET parameter with a value of 60 ms holds only for some OCPs. In these cases, the ET value of 60 ms is always combined with an EV value of 2 v, and both these values are the higher ones of the ET and EV range chosen. The EV and ET were significant parameters for 70 % and 67 % of the 46 studied pesticides, respectively.

In summary, the statistical analysis allowed a quick evaluation of the variation in the signal response of certain pesticides as a result of ion trap parameters change. The obtained optimum conditions for the instrumental parameters were used to check the sensitivity of IT-MS. Due to the absence of background signals, high signal-to-noise (*S/N*) ratios were sought.

Combinatorial Optimization

The aim of fitting experimental data to a regression model was to predict the chromatographic signal response for other IT-MS parameter combinations that were not tested in the laboratory. When the second order linear model (Equation 2)

does not significantly fit data better than the first order one (Equation 1), the optimal matching corresponds to any of the extreme points of all allowed significant parameter values. Conversely, if the second order model fits better but the stationary points are not maxima or even if they are but do not match any possible combination of MS/MS parameters, it becomes necessary to screen all allowed combinations and to calculate the response in order to find the maximum S/N ratio. For this purpose, a combinatorial optimization was carried out. A visual basic for applications macro in Microsoft Excel environment was created to evaluate the regression model equation for a maximum of 8 ET×6 IT×5 EV×3 *q*×3 IMW combinations in a total of 2160 arrangements and save the best value. The maximum signal and optimal match of the MS/MS parameters are presented in Table 3.

Concerning the excitation voltage (EV) concerns, the value that maximizes the response when this parameter is significant was 0.2 V, except for four organochlorine pesticides (aldrin, dieldrin, methoxychlor, and *o,p'*-DDT took the value of 2.0 V) and several others where the EV had no significant effect on the signal response (e.g., lindane, azoxystrobin, β -cyfluthrin, tetraconazole, and iprodione). In general, the optimal ET value was 5 ms corresponding to the lowest allowed value except for some of the organochlorine pesticides for which a value of 60 ms was obtained. The “*q*” values that maximize the response were 0.30 and 0.45. It can also be extracted from Table 3 that there is a strong correlation between ET and EV parameters. When both parameters are significant there is a linear positive relationship between them.

Validation Process

Analyte fortification was carried out by applying a known concentration of the 46 pesticides to a known quantity of blank strawberry. Following fortification, spiked samples were allowed to equilibrate 30 min at room temperature prior to extraction. Linearity of the analytical method described was studied using calibration curves with six concentration levels ranging between 5 and 250 μgkg^{-1} . The calibration curves were linear given that the determination coefficients were always higher than 0.99 for all pesticides. For all 46 studied pesticides, the LODs were lower than 12 μgkg^{-1} . The results indicate that the LOD is adequate for food safety.

Recovery was studied in a spiked strawberry sample at three levels of concentration. The majority of the pesticides gave satisfactory recoveries (ranging from 70 % to 120 %). Despite diazinon (60 %) tolylfluanid (62 %), and *p,p'*-DDE (59 %) in the lower spiking level, and mepanipyrim (139 %), myclobutanil (154 %), tetraconazole (131 %) and fludioxonil (132 %) in the higher spiking level, all others had satisfactory recoveries (in a range from 70 % to 120 %) after QuEChERS and cleanup steps.

Table 3. The Best MS/MS Significant Parameter Combinations

	Pesticides	Best parameters by combinatorial optimization					
		ET	IT	EV	q	IMW	
Chloroacetanilide	Alachlor	5	24	0.2	–	4	
	Organochlorines	60	–	2.0	–	–	
Organochlorines	Aldrin	60	–	2.0	0.30	4	
	Dieldrin	60	–	2	0.30	1	
	Endrin	60	–	2	0.45	–	
	Hexachlorobenzene	–	2	0.2	–	–	
	Lindane	60	–	–	–	–	
	Methoxychlor	60	–	2.0	0.30	1	
	o,p'-DDT	60	36	2.0	0.30	–	
	p,p'-DDE	5	2	0.2	0.45	2	
	α-Endosulfan	5	2	0.2	0.30	–	
	β-Endosulfan	–	2	0.2	0.30	–	
	IS	5	2	0.2	0.45	–	
	Triazines	Atrazin	5	12	0.2	–	–
		Atrazine-desethyl	5	12	0.2	–	–
		Simazine	–	–	0.2	–	–
Strobin	Azoxystrobin	5	2	–	–	–	
Pyrethroids	β-Cyfluthrin	–	2	–	–	–	
	Bifentrin	5	–	0.2	0.30	–	
	Cypermethrin	–	–	–	0.45	–	
	Deltamethrin	5	–	0.2	–	1.0	
	Fenpropathrin	5	–	0.2	0.45	4	
	Fenvalerate	5	12	0.2	–	–	
	λ-Cyhalothrin	5	–	0.2	0.45	4	
	Permethrin	–	2	0.2	0.30	4	
	Pyrimidine	Bupirimate	5	24	0.2	–	–
		Mepanipyrim	–	–	–	–	–
		Cyprodinil	5	12	–	–	–
		Pyrimethanil	–	12	–	–	–
	Thiophthalimide	Captan	–	2	–	–	–
		Folpet	5	24	0.2	–	1.0
Organophosphorus	Chlorpyrifos	–	–	0.2	0.30	–	
	Diazinon	5	–	0.2	–	–	
	Malathion	5	12	0.2	–	–	
Unclassified	Dazomet	–	24	–	–	2	
	Fludioxinil	5	24	0.2	0.45	–	
Anilide	Fenhexamid	–	2	–	–	–	
Aryloxyphenoxy propionic acid	Fluazifop-p-butyl	5	12	0.2	–	4	
	Quizalofop-p-ethyl	5	–	–	–	–	
N-methyl carbamate	Methiocarb	–	–	0.2	–	–	
Azole	Myclobutanil	–	24	0.2	–	–	
	Tetraconazole	5	–	–	–	–	
	Pendimethalin	5	2	0.2	–	–	
2,6-Dinitroaniline	Tolyfluanid	5	–	0.2	–	–	
Phenylsulfamide	Vinclozolin	–	–	–	–	–	
Dicarboximide	Vinclozolin	–	–	–	–	–	
	Procymidone	5	12	0.2	–	–	
	Iprodione	5	24	–	–	–	

Default parameter

Screening of Unknown Samples

The optimized method using the 46 multiclass pesticide screening program was applied to Portuguese strawberry samples. In each unknown sample, the target (quantifier) and qualifier MRM transitions of each test compound (Table 1) were monitored. Five percent of all samples examined contained levels of organochlorine pesticides greater than the limit established by the European Union for strawberry samples. The highest value was 27.30 $\mu\text{g}\cdot\text{kg}^{-1}$ for lindane [5].

In only three samples, residues of fludioxonil, bifenthrin, mepanipyrim, tolyfluanid, cyprodinil, tetraconazole, iprodione, and malathion were detected, and the

concentrations were much lower than their harmonized EU MRL. The highest concentration was obtained for iprodione (1071 $\mu\text{g}\cdot\text{kg}^{-1}$), which is lower, however, than the MRL value. Looking for positive pesticide residues, except for organochlorine pesticides that are stable and very persistent in the environment, most of the others are commonly used in strawberries crops.

Conclusions

The selected optimization strategy based on multiple linear regression allows for an efficient method development. It is not only helpful to optimize all compounds or a group of them, as it can also help to establish the optimum values for

all parameters of the MS/MS system. In the case of OCPs (aldrin, dieldrin, endrin, lindane, and methoxychlor) that tend to be very persistent in the environment, it is proposed that because of their stable structure, higher ET (60 ms) and EV (2 v) are required to achieve the best signal response. The results showed that certain pesticides, which have low signal in MS/MS, also showed poorly to fit in the models. The applicability of the method was demonstrated by analysis of real samples (strawberries), resulting in good quality control data, thus making possible rapid and reliable determination of the targeted pesticides.

This paper shows that the statistics study is a useful tool to optimize this kind of analytical parameters. Particularly, multiple linear regression was an important tool to predict the “best” combination of IT-MS parameters to maximize the analytical response within the range of values experimentally tested in this work.

Acknowledgment

This research was supported by a Ph.D. grant (BD/47200/2008) from Fundação para a Ciência e a Tecnologia, co-financed by POPH/FSE (Programa Operacional Potencial Humano/Fundo Social Europeu), grant no. PEst-C/EQB/LA0006/201 and Ph.D. grant from programa Mundus.

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