

Evaluation of degradation of organophosphate insecticide residues in stored corn (*Zea mays L.*) grains

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SUMMARY

The aims of this work were to monitor the dissipation dynamics of the organophosphate insecticides applied to corn grains stored in supervised experiments, establish the dissipation dynamics and determine the minimum period post application necessary to comply with the established commercial standards. The insecticides dichlorvos (DDVP), chlorpyrifos-methyl+deltamethrin (CPM) and pirimiphos-methyl (PMM) were applied to maize grains, which were then stored for 120 days. The samples were processed and extracted using the QuEChERS technique and the pesticide residues were identified and quantified by gas chromatography-mass spectrometry. Deltamethrin residues were not analyzed. After 90 days of storage, more than 80% of the initial residue had been degraded, and after 120 days the residues were below the detection limit (LoD). DDVP, CPM and PMM residues followed a first order degradation dynamics [$C_t = C_0 \times e^{-kt}$], with k values of 0.384, 0.025 and 0.020 and estimated half-lives of 1.8, 27.7 and 34.7 days, respectively. These results show that at recommended dosages, the insecticides CPM and PMM are safe for use on stored maize grains; DDVP results provide additional evidence further reasserting its prohibition, which dates back to 2018.

Key words: postharvest, grains, degradation kinetics, dichlorvos, chlorpyrifos-methyl, pirimiphos-methyl, residues

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RESUMEN

Los objetivos del trabajo fueron monitorear la degradación de insecticidas organofosforados aplicados a granos de maíz almacenados en ensayos experimentales, establecer las dinámicas de degradación y determinar el tiempo mínimo de espera post aplicación necesario para cumplir con los

estándares comerciales. Se aplicaron los insecticidas diclorvós (DDVP), clorpirifós-metil+deltametrina (CPM) y pirimifós-metil (PMM) a granos de maíz almacenados durante 120 días. Las muestras se procesaron y extrajeron mediante la técnica QuEChERS y los residuos de plaguicidas se identificaron y cuantificaron mediante cromatografía de gases-espectrometría de masas. Los residuos de deltametrina no fueron determinados. Luego de 90 días de almacenamiento, más del 80 % del contenido inicial de residuos se había degradado y a los 120 días se encontraban por debajo del límite de detección (LoD). Los residuos de DDVP CPM y PMM siguieron una dinámica de degradación de primer orden [$C_t = C_0 \times e^{-(k)t}$], con valores de k de 0,384, 0,025 y 0,020 y vidas medias estimadas de 1,8; 27,7 y 34,7 días respectivamente. Estos resultados permiten concluir que a la dosis recomendada, los insecticidas CPM y PMM se pueden usar de manera segura en granos de maíz almacenados, mientras que los resultados de DDVP aportan información adicional que ratifica su prohibición, que data del año 2018.

Palabras clave: postcosecha, granos, dinámica de degradación, diclorvós, clorpirifós-metil, pirimifós-metil, residuos

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INTRODUCTION

Argentina is one of the world leaders in maize (*Zea mays* L.) production, occupying the fourth place as producer with more than 56 Mt in 2019 and the third place as exporter after Brazil and USA with 36 Mt (FAO Stat, 2020). In the last 30 years, maize yields in Argentina have increased at a rate of 159 kg.ha⁻¹.yr⁻¹. In this country, maize cultivation is of great importance, due to the large cultivated area, annual production volume and multiple uses for human and animal consumption or industrialization. Most of the production is exported, providing important foreign currency revenues to the country.

During storage, maize grains can be attacked by insects, which are controlled using preventive measures, such as spraying with organophosphate

and pyrethroid insecticides (Abadia and Bartosik, 2013). However, to avoid the presence of residues in grains above the Maximum Residue Limits (MRLs) legally allowed both by the Argentinian regulations from Servicio Nacional de Sanidad y Calidad Agroalimentaria (SENASA, 2010) and by the international regulations of the *Codex alimentarius* (2020), insecticides should be applied appropriately. Thus, information about the dissipation kinetics of pesticides in crops is a key aspect in preventing risks and reducing the impact on health and trade. More restrictive MRLs in the exportation markets create obstacles to maize trade in big markets, such as the Europeans. Therefore, studies that include the analysis of residue dynamics in food are essential to support production and take decisions. The most used model for this is a pseudo-first-order kinetic model (Fantke and Juraske, 2013). For this type of model,

it is also important to calculate the maximum residual concentrations that can be allowed in food for safe human or animal consumption according to the MRLs. The minimum period of time between pesticide application and consumption is called "post-harvest interval" (POHI) and is defined as the time in days established between the last application in storage and consumption of the agricultural products. The requirements have determined that grains shall be completely free from insects but contain residues of no more than one tenth or less of the MRLs (Fleurat-Lessard et al., 1998). PMM and CPM are registered in Argentina for application in stored grains. Regarding DDVP, in 2018 SENASA through Resolution N° 149/2018 prohibited the importation, commercialization and use of the active principle and products that contain it for use in grains. This prohibition came as a response to the widespread use of DDVP in Argentina which established much higher MRLs than those required by the purchasing countries. Even though this active principle is banned in Argentina and its use restricted to only some agricultural products in foreign legislation, the results of this work can be considered an important contribution on the dynamics of residues degradation. The dissipation of insecticide residues in other cereal grains, like wheat, has been extensively studied (Pal and Shah, 2008; Uygun et al., 2008; Lucini and Molinari, 2011), but that for maize is poorly known (Alleoni and de Baptista, 2001; Hamacher et al., 2002; Sgarbiero et al., 2003; Silveira et al., 2009). Pesticide trading companies must carry out tests to determine the POHI necessary to reach values below the MRL, but their studies are not available to estimate the residue levels after a pre-market application. Since MRLs vary among different markets, they turn out to act as barriers to trade. As respects storage grains, for SENASA (2021) there is not a "post-harvest interval" (POHI) for PMM and CPM but in the case of grain production for exportation, the Limit of Maximum Residue in the country must be known and the application periods of these pesticides must be adapted to them. For CPM a period of 20-30 hours is recommended before the consumption. Regarding DDVP there is not a grace period since it is banned, but in the past it used to be 20 to 30 days. The information about the MRLs in Argentina, European Union (EU) and the *Codex Alimentarius* are available (Global MRL database, 2020; European Union Pesticides database, 2020; SENASA, 2021). The surface characteristics and the physical and chemical properties of the grain tegument of different cereals are very different and thus specific degradation models are required to describe the degradation of pesticide residues

in a specific cereal, stored in a specific country (Fleurat-Lessard et al., 1998).

Thus, the aims of this work were to monitor the degradation dynamics of the organophosphate insecticides applied to stored maize grains in Argentina, to establish the first order kinetics and to determine the POHI necessary to comply with different standards.

MATERIALS AND METHODS

Supervised experimental trials and analysis of pesticide residues

The experimental field assays were carried out during 120 days at Instituto Nacional de Tecnología Agropecuaria at the Estación Experimental Agropecuaria Manfredi (EEA INTA Manfredi), Córdoba province, Argentina. A completely randomized experimental design was used to simulate the typical grain storage conditions in Argentina. Three separate maize grain lots (13 kg) were treated with each pesticide and other three grain lots of the same size were used as controls (untreated). All lots were sampled at different time points to quantify the pesticides residues. The organophosphate insecticides formulated as emulsifiable concentrates were evaluated on the maize supervised experimental trials. The concentrations (and rates in μg of active ingredient per g of grain) were 100% (20 μg per g) for dichlorvos (DDVP), 14.5% + 0.65% (2.9 μg per g) for chlorpyrifos-methyl + deltamethrin respectively (CPM), and 50% (5 μg per g) for pirimiphos-methyl (PMM). The maize grains were treated with the rates indicated in the labels of the commercial products used.

For insecticide application, maize grains were spread in a layer, on a 100 μm polyethylene sheet for a homogeneous insecticide distribution; insecticides were applied using a hand 1.5l Giber sprayer (Giber SA, Argentina). The flow was previously tested with water. The grains were vigorously mixed, placed inside the duly labeled plastic containers, capped and stored. The lids were perforated to allow gas exchange.

The plastic containers were placed in shelves inside a room conditioned for grain storage. During the experimental assay, room temperature and relative humidity (RH) were maintained in relatively controlled conditions (within 20 to 25 °C and 40 to 60% RH).

The objective of the study was to put the samples in conditions similar to the real ones. Grain moisture content was 14%. Samples (500g) were collected

from each lot with a bag trier at 2, 30, 60, 90 and 120 days after application, placed in three-layered bags and taken to a freezer at -20°C until further analysis.

Grain samples were processed at the Laboratorio de Calidad de Granos at EEA INTA-Manfredi, Argentina. The milling was done with glass blenders. Insecticides were extracted from grains by using the QuEChERS technique (Anastassiades et al., 2003; Lehotay, 2007) adapted to dry matrices (Mastovska et al., 2010; Lucini and Molinari, 2011). Each sample (5 g) was extracted in a centrifuge tube by adding water (10 mL) and acetonitrile (15 mL) and 150 μL of internal standard solution of ethoprophos ($20\mu\text{g}\cdot\text{mL}^{-1}$), 6g of anhydrous magnesium sulfate and 1.5g of sodium chloride. The extract was then homogenized and centrifuged ($\text{RCF}=5000\text{ g}$, 5°C , 5 min). An aliquot (5 mL) of the organic phase supernatant was transferred to a clean-up tube containing 0.75 g of anhydrous magnesium sulfate, 0.2 g of bonded silica bulk, primary or secondary amine SUPELCO 40 μm (PSA) and 0.2 g of C_{18} , and then shaken and centrifuged ($\text{RCF}=5000\text{ g}$, 5°C , 5 min). An aliquot of the supernatant (2 mL) was transferred and evaporated to dryness under a stream of nitrogen with the water bath set at 35°C . The sample was recovered with toluene (500 μL) and shaken; 300 μL was collected and placed in a vial. Then, 50 μL of Triphenylphosphate (TPP) solution ($2\mu\text{g}\cdot\text{mL}^{-1}$) and 25 μL of toluene were added. Blank samples (without matrix with insecticide mixture) and fortified samples (with matrix and insecticide mixture) were prepared for quality control of the extraction technique.

Validations were also performed using calibration tests, matrix effect studies and recovery experiments to define the linear dynamic range and calculate recovery, expanded uncertainty, and limits of detection (LoD) and limits of quantification (LoQ). Recovery data were used to assess the accuracy and precision of the method (Document SANCO 10684/2009). Standard solutions of all the active pesticide compounds used in this study were prepared for validation assays by using stock solution concentrations of $1\text{ mg}\cdot\text{mL}^{-1}$ for DDVP, PMM and CPM, and $2\text{ mg}\cdot\text{mL}^{-1}$ for ethoprophos and TPP to derive working solutions. The fortification levels used for DDVP and PMM were 0.01, 0.05, 0.1, 1, 3 and $9\mu\text{g}\cdot\text{g}^{-1}$, whereas those for CPM were 0.01, 0.05, 0.1, 1 and $3\mu\text{g}\cdot\text{g}^{-1}$, using a matrix of organic maize flour (5 g) and three replicate tests per level.

Pesticide residues from maize samples of supervised experimental trials were identified and quantified by high-resolution gas chromatography coupled to mass spectrometry (GC-MS) at the

Laboratorio de Contaminantes Químicos of the Instituto de Tecnología de los Alimentos of INTA Castelar. Deltamethrin residue insecticide that was applied to the corn storage grains treatment, was neither able to be identified nor quantified.

A Perkin Elmer Clarus 600 gas chromatograph, with programmable temperature vaporizer (PTV) injection port was used in order to separate, identify and quantify each insecticide analyte. The PTV temperature program used was 35°C (4 min), then it was raised to 290°C ($200^{\circ}\text{C}\cdot\text{min}^{-1}$ for 4.5 min), and finally reduced to 35°C ($60^{\circ}\text{C}\cdot\text{min}^{-1}$). The injection volume was 25 μL : For separation of each analyte a Guard column fused silica of $5\text{ m} \times 0.25\text{ mm}$ (Supelco) with a Capillary column (Varian, Factor Four VF-5ms), $30\text{ m} \times 0.25\text{ mm}$ (id, $0.25\mu\text{m}$). Helium of 99.999 % purity was used as gas carrier. The initial pressure was 2 psi (3.9 min), reached 25 psi (at $25\text{ psi}\cdot\text{min}^{-1}$ for 7 min), and then decreased to 15 psi (at $25\text{ psi}\cdot\text{min}^{-1}$), which was maintained until the end of the assay (35 min). The initial split flow was $100\text{ mL}\cdot\text{min}^{-1}$; the flow was closed at min 3.9 ($0\text{ mL}\cdot\text{min}^{-1}$) and at min 10, the flow was set at $20\text{ mL}\cdot\text{min}^{-1}$. The oven temperature program used was 70°C (8 min), increased to 170°C ($25^{\circ}\text{C}\cdot\text{min}^{-1}$), 230°C ($5^{\circ}\text{C}\cdot\text{min}^{-1}$), reaching a maximum of 290°C (at $20^{\circ}\text{C}\cdot\text{min}^{-1}$) which was held for 10 minutes. The transfer line in the detector was held at 290°C and the collision cell at 200°C . Chromatographic data were obtained by electron impact (EI+), quadrupole analyzer and monitoring of specific ions (SIM, selected ion monitoring), one ion for quantitation and at least two for qualification for each analyte. High-purity analyte standards from Sigma Aldrich (USA) were used for validation assays and quality control of the extraction process: dichlorvos (99.9%), chlorpyrifos-methyl (99.9%) and pirimiphos-methyl (99.5%); and the internal standards ethoprophos (93.1%) and triphenylphosphate (TPP) (99%). Control samples (organic maize flour) were used for validation assays and as quality control of the extraction process. DDVP, CPM and PMM residues were recorded. The results of the analyses are expressed in micrograms of the DDVP, CPM and PMM insecticides per gram of grains ($\mu\text{g}\cdot\text{g}^{-1}$).

Statistical analyses

The daily dissipation rate (DDR) was calculated considering the decrease in the level of residues within two successive analytical determinations. A generalized linear mixed model with gamma distribution and logarithmic link function was adjusted to evaluate the effect of the days after application on the degradation of each insecticide.

The model included a random effect to induce experimental unit correlations between repeated measurements of the same lots of grains. The dynamics of the residues according to the days after application (dissipation kinetics) was modeled by the first-order kinetic model:

$$C_t = C_0 \times e^{-(K \cdot t)} \quad \text{Eq. (1)}$$

where C_t is the residue concentration at time t , C_0 is the initial concentration or nominal rate, and K is the dissipation constant. The parameters estimated (C_0 and K) were used to calculate using Eq. (2). Analyses were performed using the statistical software InfoStat (Di Rienzo et al., 2019).

$$HL \text{ or } t_{1/2}(\text{days}) = \ln 2 \cdot K^{-1} \quad \text{Eq. (2)}$$

RESULTS AND DISCUSSIONS

Validation of the QuEChERS+GC-MS methodology

The values of average recovery, relative standard deviation, expanded uncertainty, LOD and LOQ for DDVP, CPM and PMM by using QuEChERS+CG-MS are shown in Table 1. An effect of the matrix was observed for all analytes and the quantification was performed through calibration curves in the matrix, which were linear for the concentration range studied ($R^2 > 0.98$) (Document SANCO 10684/2009).

Mean recovery values for the three insecticides were greater than 70%, with adequate reproducibility. These results are consistent with previous validation of QuEChERS in cereal grains (Mastovska et al., 2010; Lucini and Molinari, 2011; Yu et al., 2014). The validation assays showed no residues in the control samples during the extraction process. Thus, we inferred that the process quality was adequate.

Dynamics of insecticide residues

All the treated samples showed residues of the insecticides studied, whereas the untreated control

samples were negative for residues. The curves describing the expected dynamics for the three insecticides are presented in Figure 1. The mean residue levels obtained in maize grains at 2, 30, 60, 90 and 120 days after application, the DDR, and the residue decline (%), estimated by the pseudo-first-order kinetic model, are presented in Table 2. The equations representing the dissipation dynamics are shown in Table 3.

DDVP, CPM and PMM residues showed lower levels as the number of days after application increased (Table 2), in response to a first-order kinetics, with high coefficient of determination (R^2) (Table 3). The number of days after application had a statistically significant effect ($p < 0.05$) on the levels of residues present in the grains for the three insecticides, with residues below the LoQ at 120 days (Table 2). The dissipation kinetics has three phases: the first or "stripping phase", which is due mainly to mechanical causes not involved in storage trails (wind and rain); the "degradation phase", which is due to physical and chemical causes; and the "persistence phase", which is determined by

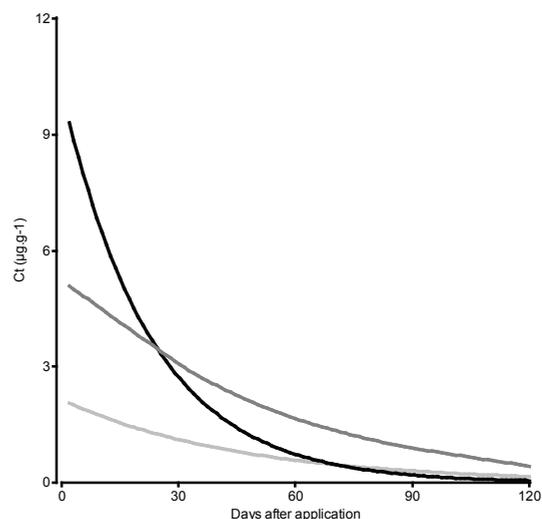


Figure 1. Curves describing the expected dynamics for dichlorvos (—), pirimiphos-methyl (—) and chlorpyrifos-methyl (—) applied in stored maize grains

Table 1. Parameters obtained during analytical method validation for dichlorvos (DDVP), chlorpyrifos-methyl (CPM) and pirimiphos-methyl (PMM) in stored maize grains

Insecticides	Spiked level ($\mu\text{g}\cdot\text{g}^{-1}$)	Average recovery (%)	RSD (%)	LoD ($\mu\text{g}\cdot\text{g}^{-1}$)	LoQ ($\mu\text{g}\cdot\text{g}^{-1}$)	Regression equation	R^2	U (%)
DDVP	0.01-9.00	78.0-102.8	6.6-13.1	0.003	0.01	$y = 0.37x - 0.04$	0.99	13.1-26.2
CPM	0.01-3.00	78.0-99.9	10.3-20.0	0.003	0.01	$y = 0.81x - 0.06$	0.98	20.5-31.8
PMM	0.01-9.00	70.0-95.7	2.3-25.0	0.003	0.01	$y = 0.95x + 0.27$	0.99	4.6-24.8

References: relative standard deviation (RSD), limit of detection (LoD), limit of quantification (LoQ), expanded uncertainty (U).

Table 2. Degradation parameters of dichlorvos (DDVP), chlorpyrifos-methyl (CPM) and pirimiphos-methyl (PMM) in stored maize grains obtained in experimental assays

Variable	Insecticides	Days after application				
		2	30	60	90	120
Residues ($\mu\text{g}\cdot\text{g}^{-1}$)	DDVP	9.27	2.97	0.38	0.02	<LoQ
	CPM	1.98	1.29	0.67	0.10	<LoQ
	PMM	5.10	3.54	1.52	0.72	<LoQ
DDR ($\mu\text{g}\cdot\text{g}^{-1}\cdot\text{day}^{-1}$)	DDVP	0.2250	0.0860	0.0120	0.0010	---
	CPM	0.0250	0.0210	0.0190	0.0030	---
	PMM	0.0560	0.0670	0.0027	0.0024	---
Residue decline (%)	DDVP	ND	68.0	95.9	99.8	100.0
	CPM	ND	34.8	66.2	94.9	100.0
	PMM	ND	30.6	70.2	85.9	100.0

References: daily dissipation rate (DDR), < LOQ = below limit of quantification. Nominal rate: $20 \mu\text{g}\cdot\text{g}^{-1}$ for DDVP, $2.9 \mu\text{g}\cdot\text{g}^{-1}$ for CPM and $5 \mu\text{g}\cdot\text{g}^{-1}$ for PMM. SD: no data.

Table 3. Equations for first-order kinetics of residue dynamics, for dichlorvos (DDVP), chlorpyrifos-methyl (CPM) and pirimiphos-methyl (PMM) in stored maize grains

Insecticide	Kinetics equation	R ²	HL or $t_{1/2}$ (days)
DDVP	$Ct = 20 \times e^{-0.384t}$	0.97	1.8
CPM	$Ct = 2.54 \times e^{-0.025t}$	0.88	27.7
PMM	$Ct = 5.27 \times e^{-0.020t}$	0.78	34.7

References: coefficient of determination (R²) and half-life (HL- $t_{1/2}$).

the retention residues in the material. In our study, the second and third phases were recognized for all insecticides.

With respect to experimental DDR, the number of days after application had a statistically significant effect ($p < 0.05$) on the levels of residues present in the grains for DDVP and PMM. The DDR for PMM were different until 60 days after application, whereas those for CPM were not statistically different. The apparent rates of degradation in the initial storage period were higher than those in the later stages. The values of DDR inferred from the fitted model were similar to the experimental ones in the case of PMM and CPM.

For DDVP, the DDR were high up to 30 days after application, which corresponds to the "degradation phase" of residues, whereas after that period, DDR decreased (Table 2). The K coefficient was 0.384 for DDVP, 0.025 for CPM and 0.020 for PMM (Table 3). For all the insecticides, 90 days after application, experimental residues showed a clear decline (Figure 2). The calculated HL was higher for PMM than for CPM and lowest for DDVP (Table 3).

The first-order degradation kinetics has been previously reported in other matrices for DDVP, CPM and PMM (Fleurat-Lessard et al., 1998; El-Behissy et al., 2001; Yu et al., 2014). In wheat and oat grains treated with PMM, for example, Lucini and Molinari (2011) did not detect residue degradation,

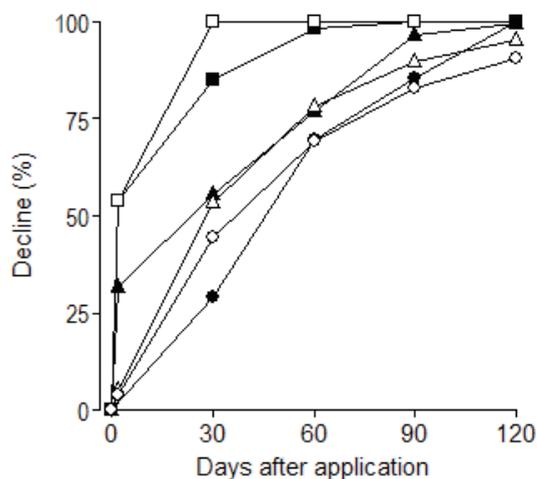


Figure 2. Experimental residue decline (%) of dichlorvos (■), chlorpyrifos-methyl (▲) and pirimiphos-methyl (●) and estimated decline of dichlorvos (□), chlorpyrifos-methyl (△) and pirimiphos-methyl (○) in maize grains at 2, 30, 60, 90 and 120 days after application

whereas in maize stored for 18 months. White et al. (1997) found similar results. In rice, Yu et al. (2014) determined K coefficient of 0.384 for DDVP, 0.025 for CPM and 0.020 for PPM.

In wheat, other authors found lower values for CPM, between 0.0038 and 0.0072 (Fleurat-Lessard et al., 1998; Pal and Shah, 2008). In the present study, the first-order kinetics for DDVP showed high average K coefficients, in agreement with results reported by other authors working with this insecticide (El-Behissy et al., 2001). The high value of K for DDVP may respond to the high volatility of this insecticide (El-Behissy, et al. 2001), which corresponds to a short period of protection. Then, the fitted model should be used to predict the

behavior of residues only in the initial stages of dissipation.

The residue decline (85 %) was consistent with other findings in cereal grains (Sgarbiero et al., 2003; Silveira et al., 2009; Strada et al., 2012), and in agreement with the physico-chemical characteristics of each active ingredient, as reported by Balinova et al. (2006). These authors determined that, after 90 days of storage of wheat, CPM residues decreased by 60 % and that PMM residues decreased by only 30 %. This decrease in PPM residues observed in maize 90 days after application was in consonance with the decrease observed by Hamacher et al. (2002) in wheat after 126 days of storage and by Yu et al. (2014) in rice after 60 days of storage. Fantke and Juraske (2013) determined that organophosphate pesticides have high variability in HL (0.7 to 55 days). In our study, the average half-life (HL or $t_{1/2}$ in days) of each insecticide in stored maize grains was higher for PMM than for CPM and the lowest for DDVP. In contrast, there has been slower dissipation of PMM residues reported in wheat (between 100 and 490 days) by Holland, Hamilton, Ohlim and Skidmore, 1994 and in maize (between 85 and 95 days) by Alleoni and de Baptista, 2001 and Sgarbiero et al., 2003.

Yu et al. (2014) reported HL values for PMM between 27 and 29 days in rice, where as other authors reported HL values lower than 14 days for DDVP in different matrices (Holland et al., 1994; El Behissy et al., 2001). In our study, DDVP was the insecticide with highest dissipation (1.8 days). Regarding CPM, the HL reported in stored wheat grains is of 60-70 days (Pal and Shah, 2008) or 93-169 days, depending on the grain moisture (Fleurat-Lessard et al., 1998; Holland et al., 1994). All these results confirm that the HL estimated for a pesticide depends on the matrix studied, the active ingredient, the application rate, and the environmental conditions during storage, and the grain moisture content. Likewise, the residue decline is slower when the insecticide is applied on stored products (Holland et al., 1994; Balinova et al., 2006) than on crops in the field, with residue persistence (Lucini and Molinari, 2011).

The MRLs for PMM according to the different legislation are: SENASA, 10 $\mu\text{g}\cdot\text{g}^{-1}$, *Codex Alimentarius*, 7 $\mu\text{g}\cdot\text{g}^{-1}$ for PMM on storage grains and UE legislation, 0.5 $\mu\text{g}\cdot\text{g}^{-1}$.

In the same way, the MRLs for CPM are 5 $\mu\text{g}\cdot\text{g}^{-1}$ according to SENASA and 0.01 $\mu\text{g}\cdot\text{g}^{-1}$ according to EU legislation. The MRLs for DDVP are: following the *Codex Alimentarius*, 7 $\mu\text{g}\cdot\text{g}^{-1}$ for DDVP (in rice and wheat) and following EU legislation, 0.01 $\mu\text{g}\cdot\text{g}^{-1}$

for DDVP EU pesticide database, 2020.

Some works conducted in stored grains have reported that storage is not efficient to reduce insecticide residues below the permitted MRLs (Uygun et al., 2008).

The results of our study suggest that the residues of PMM in maize grains under the storage conditions commonly used in Argentina do not exceed the MRLs established by SENASA and the *Codex Alimentarius*. In the present study, the PMM residues found two days after application were below the MRLs established in Argentina and suggested by the *Codex Alimentarius*; however, the values exceeded those established by the EU.

CPM residues were below the MRLs established in Argentina two days after application. With respect to CPM the POHI within the limits by SENASA. POHI to reach residue values experimentally established by the EU (0.01 $\mu\text{g}\cdot\text{g}^{-1}$) is 110 days for CPM and DDVP. Then, grains destined for this market should not be treated with these insecticides (Abadia and Bartosik, 2013).

CONCLUSIONS

A first-order kinetic model allowed explaining the decline of residue levels of DDVP, CPM and PMM applied for insect control on stored maize grains, with levels below the LoQ being found at 120 days. The half-lives of insecticide residues and DDR varied among these organophosphate pesticides. Regarding the POHI required for residues to reach the MRLs, the three insecticides exceeded the EU MRLs, whereas PMM and DDVP are in accordance with the regulations of Argentina and the *Codex Alimentarius*, because the residue levels were within the tolerated. Therefore, the results lead to the conclusion that, at the recommended dosage, the use of the insecticides PMM and CPM will be recommended in maize grains considering the LMR by SENASA. The knowledge of the dissipation curves in these three insecticides allows us to predict the POHI necessary to attain a certain level of residues in order to comply with current legislation.

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