[Click here to view linked References](#)

**Postprint of: Environmental science and pollution research international
24(11): 10410-10420 (2017)**

**SLOW RELEASE FORMULATIONS OF THE HERBICIDE PICLORAM BY USING Fe Al-
PILLARED MONTMORILLONITE**

Jose L. Marco-Brown^{a,b†*}, Tomás Undabeytia^c, Rosa M. Torres Sánchez^d, María dos Santos Afonso^{a,b}

^aUniversidad de Buenos Aires, Facultad de Ciencias Exactas y Naturales, Departamento de Química Inorgánica, Analítica y Química Física, Ciudad Universitaria, Pabellón II, C1428EHA Buenos Aires, Argentina.

^bCONICET-Universidad de Buenos Aires, Instituto de Química Física de los Materiales, Medio Ambiente y Energía de la Facultad de Ciencias Exactas y Naturales (INQUIMAE), Buenos Aires, Argentina

^cInstitute of Natural Resources and Agrobiology (IRNAS-CSIC). Reina Mercedes 10. Apdo. 1052. 41080 Sevilla, España.

^dCETMIC (Centro de Tecnología en Recursos Minerales y Cerámica), CONICET-CCT La Plata-CIC, Camino Centenario y 506 CC (49) (B1897ZCA) M. B. Gonnet, Argentina.

*Corresponding author.

E-mail addresses: josemarcobrown@gmail.com, joseluis.marcobrown@unsam.edu.ar (J.L.M-B.), undabeytia@irnase.csic.es (T.U.), rosats@cetmic.unlp.edu.ar (R.M.T.S.), dosantos@qi.fcen.uba.ar (M.d.S.A.).

Present address:

[†]J.L.M-B: Instituto de Investigación e Ingeniería Ambiental (3iA), CONICET, Escuela de Ciencia y Tecnología, UNSAM, 25 de Mayo y Francia (1650) San Martín, Argentina.

Abstract

Slow release formulations of the herbicide picloram (PCM, 4-amino-3,5,6-trichloropyridine-2-carboxylic acid) were designed based on its adsorption on pillared clays (PILCs) for reducing the water-polluting risk derived from its use in conventional formulations. Fe-Al PILCs were synthesized by the reaction of Na⁺-montmorillonite (SWy-2) with base-hydrolyzed solutions of Fe and Al. The Fe/(Fe+Al) ratios used were 0.15 and 0.50. The PCM adsorption isotherms on Fe-Al PILCs were well-fitted to Langmuir and Freundlich models. The PCM adsorption capacity depended on the Fe content in the PILCs. Slow release formulations were prepared by enhanced adsorption of the herbicide from picloram-cyclodextrin (CD) complexes in solution. CDs were able to enhance up to 2.5-fold the solubility of PCM by the formation of inclusion complexes where the ring moiety of the herbicide was partially trapped within the CD cavity. Competitive adsorption of anions such as sulfate, phosphate and chloride as well as the FTIR analysis of PCM-PILC complexes provided evidence of formation of inner sphere complexes of picloram-cyclodextrin on Fe-Al-PILCs. Release of the herbicide in a sandy soil was lower from Fe-Al PILC formulations relative to a picloram commercial formulation.

Keywords: picloram, Fe-Al pillared montmorillonite, slow release formulations, adsorption

Acknowledgments

The authors acknowledge Universidad de Buenos Aires, Secretaria de Ciencia y Técnica Projects, Ministerio de Ciencia y Técnica, Agencia Nacional de Promoción Científica y Tecnológica, MINCyT-ANPCyT-FONCyT and Consejo Nacional de Investigaciones Científicas y Técnicas de la República Argentina (CONICET) for financial support. J.L.M-B acknowledges CONICET fellowship.

This research was also supported by the MEC Project CTM2013-42306-R and the Junta de Andalucía Project P12-RNM1897. Both projects received funding by the European Social Fund. The authors also acknowledge the Analytical Services of Seville University (CITIUS) for SEM-EDS measurements.

Introduction

A great variety of pesticides (herbicides, insecticides, fungicides) (Akelah 1996, Delcour et al. 2015, Kenawy et al. 1992, Qi & Donahoe 2008) have been produced and used in the last 50 years, aiming to protect and to increase the yield of agricultural production. Usually pesticides are sprayed as fine particles suspended in aqueous media, or in some cases in the presence of, organic compounds to form aqueous emulsions that offer a good contact surface with the crop. The excess of pesticide, non-absorbed by crops, is partially vaporized to the atmosphere and also leached below root zone by rain or irrigation. These losses in the amount of the applied pesticide would reach and contaminate groundwater and / or surface water bodies (Haynes et al. 2000, Loewy et al. 1999, Schipper et al. 2008). To minimize this effect, in the last decades, research has focused on the development of slow release formulations (SRFs) of herbicides. Slow release formulation of fertilizers had been studied too in order to reduce contamination by nitrogen and phosphorous (Entry & Sojka 2008). The benefits in using SRFs are: increased availability of pesticides in order to enhance the desired control of the crop, reduction of the frequency of application and leaching of the pesticide in the soil and therefore the risk of groundwater contamination (Maqueda et al. 2008, Undabeytia et al. 2010).

The low cost and easy access to natural inorganic materials such as clay minerals, has allowed their use as adsorbents and also as SRFs (Ali et al. 2012, Carrizosa et al. 2000, Galán-Jiménez et al. 2013, Gupta & Suhas 2009, Lagaly 2001, Sánchez-Jiménez et al. 2012). Montmorillonite (Mt) has relevant properties as its high surface area and laminar structure, that makes it an excellent matrix to attach different chemical species to generate SRFs systems: clay protein composites (Alromeed et al. 2015), phosphatidylcholine vesicles adsorbed on clay (Undabeytia et al. 2011, Undabeytia et al. 2012), alginate/exfoliated montmorillonite nanocomposite (Yan et al. 2015), etc. In the design of SRFs, the use of polysaccharides (Campos et al. 2014), chitosan (Kashyap et al. 2015), sepiolite

gels (Maqueda et al. 2008, 2009), and herbicide encapsulation by Poly(butyl acrylate-co-styrene) (Wang et al. 2015) have been also proposed.

Pillared clay minerals (PILCs) have been used as catalysts, adsorbents for the removal of pollutants and slow release formulations, due to its multiple loaded centers, large area, high thermal stability and interlayer space. Particularly, Fe-pillared clays were used for preparation of SRFs of the herbicide imazaquin (Undabeytia et al. 2013).

Picloram, 4-amino-3,5,6-trichloropyridine-2-carboxylic acid (PCM) is a synthetic organic chemical that belongs to the pyridine family. This organic compound has herbicide activity and is used to control annual broad leaf weeds at low rates and deeply rooted herbaceous weeds, in vines, woody plants and cereals crops such as wheat, barley and oats (FAO 2007, Palma et al. 2004). Kinetic and equilibrium processes of PCM adsorption on montmorillonite and Fe PILC have been previously reported (Marco-Brown et al. 2014, Marco-Brown et al. 2012), with a greater adsorption capacity on Fe PILC. With both clays, the adsorption of PCM on the mineral surfaces occurred between the pyridine and carboxylate functional moieties of groups in the PCM molecule and the metal centers at the mineral surface (Marco-Brown et al. 2015). PCM at the pH of most of the soils and water environments ($pK_a \approx 2.3$) present a very low adsorption on soil particles and extremely high soil mobility (Biggar et al. 1978, Celis et al. 2002, Cheung & Biggar 1974, Hall et al. 2015, Osteryoung & Whittaker 1980). Therefore, it would be of interest to obtain SRF of this herbicide based on modified clays such as PILCs to minimize its environmental pollution.

In general, the active ingredient loading on clay-based formulations is limited by the herbicide solubility, which prevents further adsorption (Undabeytia et al. 2013). Cyclodextrins (CDs) are cyclic oligosaccharides obtained from the enzymatic degradation of starch and formed by 6 to 12 units of α -D-glucopyranose linked through bonds α ($1 \rightarrow 4$). CDs are crystalline, highly soluble in water with a conformation of a truncated cone with a cavity therein (Szejtli 1998). This cavity is hydrophobic, because the methylene hydrogen atoms and glycosidic oxygen atoms are located inside the cavity whereas its outer surface is hydrophilic due to the hydroxyls 2, 3 and 6 of the glucanopyranose molecules. These features give to the CDs the ability to accommodate organic molecules as guest molecules in its interior, forming inclusion complexes, thus favoring the water solubility of the host. The CD inclusion complexes formation is a dynamic equilibrium process. CDs have been deeply investigated for their use as drug delivery systems, sequestering agents, solubility increase of compounds in aqueous solution, and soil remediation (Jozefaciuk et al. 2001, Morillo et al. 2014, Undabeytia et al. 2013, Wenz et al. 2006, Yáñez et al. 2012).

In this study, and due to the great sorption capacity of Fe PILC for picloram previously evidenced (Marco-Brown et al. 2012), the use of CDs for enhanced solubility of the herbicide picloram will be explored with the aim of increasing the active ingredient content in PILC-based SRFs.

The aims of this work are: (i) to evaluate PCM adsorption on Fe-Al PILCs for the preparation of SRFs; (ii) to study the solubility enhancement in water of PCM by the use of CDs; (iii) to evaluate PCM adsorption on Fe-Al PILCs in the presence of CDs and compare with PCM adsorption without CDs; (iv) to get insight into the mechanisms involved in the preparation of the SRFs; and finally (v) to test these formulations for slow release.

Materials and methods

Materials

Wyoming Na-Mt (SWy-2) was obtained from the Source Clays Repository of The Clay Minerals Society (Columbia, MO) (cation exchange capacity 0.8 mmol/g). $\text{Al}(\text{NO}_3)_3 \cdot 9\text{H}_2\text{O}$, $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$, NaCl, Na_2SO_4 , Na_3PO_4 , H_3PO_4 and NaOH were purchased from Sigma-Aldrich (Sigma Chemical Co., St Louis, MO). HPLC grade–acetonitrile was obtained from Teknokroma S.A. (Barcelona, Spain). Picloram (PCM 100% purity) was supplied by Sigma-Aldrich. The commercial formulation of picloram (Tordon®101) was supplied by PRINA AGROTÉCNICA, S.L. (Sevilla, Spain). Cyclodextrins (CDs) were purchased from Cyclolab (Budapest, Hungary). The CDs employed were: α -CD, β -CD, hydroxypropyl- β -CD (HP β), and randomly methylated- β -CD (RAME β). Figures 1 and 2 show the structural formulas of picloram and CDs, respectively.

Fig. 1 Structural formula of picloram (PCM)

Fig. 2 Structural formula of cyclodextrins (CD). α -CD, $n = 1$, $R = \text{H}$; β -CD, $n = 2$, $R = \text{H}$; HP- β , $n = 2$, $R = \text{H}$ or $\text{CH}_2\text{-CHOH-CH}_3$ and RAME β , $n = 2$, $R = \text{H}$ or CH_3

Soil samples were collected from the Ap Horizon (0-20 cm) of a Sandy Soil (SS) classified as Typic Xeropsamment. These soil samples were passed through a 2 mm sieve before use in PCM release studies. This soil has a pH 8.9 and 0.79% of organic matter content (more detailed physicochemical properties were reported in (Undabeytia et al. 2012)).

Pillared Clay Minerals Synthesis

Synthesis of pillared clay mineral was done as reported elsewhere (Undabeytia et al. 2013). Briefly, aluminum nitrate and iron chloride solutions in two Fe/(Fe+Al) molar ratios (0.15 and 0.5) were hydrolyzed by titration with 0.4 M NaOH under nitrogen atmosphere. The Al/OH molar ratio was 2.0. These pillaring solutions were aged for seven days. Then, 10 g of clay mineral were dispersed in 250 mL water, and the pillaring solutions were added slowly at a ratio of 10 mmol (Fe+Al)/g clay mineral with a peristaltic pump under nitrogen atmosphere. The final clay mineral content was 1% (w:w). The dispersions were shaken for 24 h, followed by centrifugation, removal of chloride anions by dialysis, dry freezing and further heating at 300°C under nitrogen atmosphere. The products were identified as P0.15 and P0.5 samples, where the numbers indicated the Fe/(Fe+Al) molar ratio used. The interlayer spaces for P0.15 and P0.5 were determined to be 0.9 and 4.3 nm, respectively, by X-ray diffraction (XRD) (Undabeytia et al. 2013). Nitrogen adsorption-desorption isotherms (see Fig. S1 in Supplementary Material) were recorded at 77 K using a Micromeritics ASAP 2010 instrument. P0.15 and P0.5 presented a type II nitrogen adsorption-desorption

isotherms (Fig. S1) with a type H3 hysteresis typically associated with materials formed by sheet agglomerations of lamellar particles yielding meso- and macropores.

Surface areas determined were 195 and 172 m²/g for P0.15 and P0.5 respectively (Undabeytia et al. 2013). Scanning electron microscopy (SEM) was performed using a field emission gun scanning electron microscope Zeiss (FEG-SEM Zeiss LEO 982 GEMINI) with combined Energy Dispersive X-ray Analyzer (EDS). The SEM images were taken by applying 3 kV voltage with 4000 times magnification for the clarification of surface. The surface morphology of the clay surface changed after pillaring to a more porous and fluffy appearance as the iron content increased revealing co-aggregation of iron polyoxocations on the clay mineral surface as noticed by EDS (See Fig. S2 in Supplementary Material).

Adsorption studies

PCM adsorption experiments were performed in duplicate by mixing 24 mg of the PILC with 15 mL of PCM solutions ranging up to 1.66 mM. Preliminary kinetic studies indicated that adsorption reached equilibrium after 24 h; thus, the dispersions were shaken for 24 h at 20°C. Subsequently, the dispersions were centrifuged and the concentration of herbicide in the supernatants was determined.

PCM adsorption on P0.5 was also done in the presence of RAME β and HP β . For these experiments, 24 mg of P0.5 were suspended in 15 mL of a solution of PCM and CD. A molar ratios of 4.97:100 for PCM:RAME β and 3.93:100 for PCM:HP β were used respectively.

Adsorption of PCM on P0.5 was also studied in the presence of NaCl, Na₂SO₄ and Na₃PO₄ electrolytes. Typically, 24 mg of PILC was mixed in a 1.66 mM PCM solution with a molar ratio of 1:1 PCM:electrolyte.

The PCM concentration was determined by a Shimadzu HPLC equipped with PDA detector set at a wavelength of 230 nm. An injection volume of 100 μ L was used. The mobile phase was a mixture of 50% acetonitrile and 50% water containing 1% acetic acid. A C18 reverse phase column (Kromasil 100, 150 mm length, 0.5 mm internal diameter) was used. The flow rate was 1 mL/min. PCM retention time was 1.5 min. The adsorption experiments were performed in duplicate. The standard error was always lower than 5%.

Experimental data for PCM adsorption were analyzed using Langmuir and Freundlich models.

Langmuir adsorption isotherms

The Langmuir model is used frequently to explain adsorption processes. This model assumes a reduction of the available interaction places as the adsorbate concentration increases. The Langmuir isotherm assumes monolayer adsorption and is represented by the following equation:

$$\Gamma = \frac{\Gamma_{\max} K_L C_{\text{eq}}}{(1 + K_L C_{\text{eq}})} \quad (1)$$

where, $\Gamma_{\max}$ is the maximum uptake and means the maximum concentration on the solid phase, C_{eq} is the adsorbate concentration at the equilibrium on the aqueous media and K_L is the Langmuir equilibrium constant which is related with the free energy of the reaction.

Freundlich adsorption isotherms

The Freundlich isotherm is an empirical equation that describes the adsorption on a heterogeneous surface with a non-uniform distribution of heat of adsorption that means that the surface on which the adsorbed molecules are interacting is energetically heterogeneous. This model does not predict any saturation of the adsorbent by the sorbate. Instead, infinite surface coverage is predicted, indicating multilayer adsorption on the surface. This isotherm is determined by the following equation:

$$\Gamma_e = K_F \times C_{eq}^{1/n} \quad (2)$$

where, K_F is the Freundlich constant indicating adsorption capacity and represents a measure of the surface area of the adsorbent while $1/n$ is an irrational fraction that varies between 0.1 and 1 and is a measure of the adsorption intensity.

Picloram solubility studies in aqueous phase in the presence of CD

Due to the low solubility of PCM in water, a solubility study in presence of CD was done. PCM (20 mg) was added in excess of its solubility at 25°C (430 mg/L) to aqueous solutions (10 mL) containing various concentrations of each CD. The concentration range was from 0 to 16 mM for β -CD and from 0 to 100 mM for α -CD, HP β and RAME β . CD solutions were previously prepared in order to obtain the desired concentrations and sonicated. The experiments were carried out in triplicate. Flasks containing the dispersions were sealed and shaken for 1 week. This reaction time was chosen after preliminary kinetic studies (data not shown). The dispersions were left to settle down for 24 h, and aliquots from the supernatants were then filtered using a syringe through a 0.22 μ m Millipore cellulose glass fiber membrane filter, and the concentration of PCM in the filtrate was determined by HPLC. For the calculations, as will be seen later, only the total CD concentration used as well as picloram concentration in the filtrates were needed.

Picloram formulations preparation

Two formulations were prepared by adsorption of PCM and PCM-HP β complexes on P0.5 at 1.6 g clay/L. One formulation was prepared by mixing 24 mg of P0.5 with 15 ml of 1.66 mM PCM solution. The other one was prepared in the same way, mixing 24 mg of P0.5 with 15 ml of a solution containing PCM and HP β with a concentration of 2.90 mM and 73.69 mM, respectively.

The dispersions were shaken for 24 h at 20°C. Then, the herbicide was analyzed in the removed supernatant, whereas the pellets were dry-frozen yielding the PILC formulations. They were

denoted as PCM-P0.5 and PCM-HP β -P0.5. The content of PCM in PCM-P0.5 and PCM-HP β -P0.5 formulations were 5.7 and 7.6% w:w, respectively.

Characterization of the adsorbent, adsorbates and adsorption products

Fourier-Transform Infrared (FTIR) Spectra of P0.5, PCM, HP β , PCM-HP β complex, PCM-HP β (PM) samples and PCM-P0.5 and PCM-HP β -P0.5 formulations were recorded on a FTIR Nicolet 8700 spectrometer equipped with a DTGS detector, using KBr discs, scanned in wave number range of 400-4000 cm⁻¹. The PCM-HP β complex was obtained by dry-freezing a solution of PCM 2.5mM and HP β 5.5 mM previously shaken at 20°C during 24 h. The physical mixture of PCM and HP β denoted as PCM-HP β (PM) was prepared by grinding in a mortar PCM and HP β at a 1:1 molar ratio.

Release of herbicide

The release of PCM from prepared formulations PCM-P0.5 and PCM-HP β -P0.5 and a commercial formulation (Tordon®101) were conducted by using Büchner funnels. In this procedure 98.9 g of sandy soil (SS) was added to a Büchner funnel (70.88 cm² of surface area) equipped with a paper filter on the bottom. The soil sample was homogenized to a 0.5 cm height. The herbicide formulations were uniformly spread over the soil at a rate of 2.0 kg of active ingredient (a.i.) per ha. The soil samples in each funnel were irrigated 15 times with 15 mL, each washing corresponding to 2.12 mm rain at 20 min intervals. The eluted volume after each watering was collected, and the concentration of herbicide in the eluent was determined by HPLC. All experiments were carried out by triplicate.

Results and discussion

PCM adsorption on PILCs

PCM adsorption isotherms on P0.5 and P0.15 were depicted in Figure 3. Adsorption coverage (Γ_{ads}) of PCM on P0.5 was about 4-fold higher than that found on P0.15. This adsorption increase can be ascribed to the surface site modifications originated by the iron pillaring process. In the development of pillared clay minerals with mixed oxides, the properties of the product depended on the iron content in the structure. In this sense, it was reported an increment in the trichloroethylene oxidation capacity by the use of a pillared clay (Mishra et al. 2008) and also an increase in the herbicide imazaquin adsorption capacity of Fe-Al pillared clay related to the structural Fe content (Undabeytia et al. 2013). The latter authors explained the high affinity of the herbicide to iron surface sites yielded an increase in imazaquin adsorption. Moreover, a Γ_{ads} increase for PCM adsorption on a Fe pillared montmorillonite relative to a raw montmorillonite was also found (Marco-Brown et al. 2012).

The adsorption isotherms are important criteria in the optimization of the use of adsorbent materials, because the correlation between experimental data and adsorption models may be related to different interactions (binding strength, heterogeneity/homogeneity of adsorbent sites, adsorbate

monolayer formation, etc.) between sorbate and adsorbent. Thus, analysis of experimentally obtained equilibrium data by either theoretical or empirical equations is useful for the practical design of adsorption systems aimed at slow release. Consequently, experimental data for PCM adsorption were analyzed using both Langmuir and Freundlich models.

The experimental data were well-fitted to the Langmuir and Freundlich models (Fig. 3, Table 1). In the Langmuir model, the affinity between sorbent and sorbate is represented by the constant K_L . In general for good sorbents high $\Gamma_{\max}$ and K_L are desirable.

The K_L for PCM sorption on P0.5 is within the range of values reported for strong interactions of organic molecules on clays: benzyltrimethylammonium on montmorillonite (5000 M^{-1}) (Nir et al. 2000); ethoxylated alcohols on sepiolite ($680\text{-}900 \text{ M}^{-1}$) (Galán-Jiménez et al. 2013). In the particular case of pillared clays, the existence of strong Brönsted and Lewis acid centers can promote the formation of inner sphere complexes with the loaded metal centers on the clay. No conclusion may be achieved by the examination of the K_L value of PCM. A higher K_L value than that of PCM was obtained for the sorption of the anionic herbicide sulfometuron on pillared clay (12000 M^{-1}), whose sorption mechanism was only dominated by strong electrostatic interactions (Nir et al. 2000).

Fig. 3 PCM adsorption isotherms on (●) P0.15, (○) P0.5, (▲) P0.5 in the presence of RAME β and (▼) P0.5 in the presence of HP β . Solid and dotted lines represent Langmuir and Freundlich fits respectively.

Table 1 Langmuir and Freundlich parameters for PCM adsorption onto P0.15 and P0.5 samples in presence and absence of CDs.

Solubility diagrams of picloram in the presence of cyclodextrins

The evaluation of PCM water solubility in the presence of β -CD, α -CD, HP β and RAME β , indicated an increase with CD concentration due to the formation of an inclusion complex picloram-cyclodextrin, PCM-CD (Fig. 4). In this PCM-CD complex, part of the ring of the herbicide molecule was trapped in the polar cavity of CD. The increase of PCM solubility with CD concentration as noticed with β -CD, HP β and RAME β , indicated the existence of an inclusion complex without a solubility limit (Higuchi & Connors 1965, Undabeytia et al. 2013). According to Higuchi and Connors (Higuchi & Connors 1965), these solubility diagrams correspond to the formation of an inclusion complex with an 1:1 stoichiometry. However, when α -CD was used, PCM solubility decreased after a 70 mM α -CD concentration, indicating a complex solubility limit above which the complex precipitated.

The inclusion complex formation is a dynamic equilibrium process (Higuchi & Connors 1965) and can be expressed by the following equation:

where S is the adsorbate (PCM), L is the ligand (CD), S_mL_n is the inclusion complex and $K_{m:n}$ is the apparent stability constant of the inclusion complex and can be calculated by the following equation:

$$K_{m:n} = \frac{[S_mL_n]}{[S]^m[L]^n} \quad (4)$$

where $[S]=S_0$, $[L]=L_t-n[S_mL_n]$, $[S_mL_n]=(S_t-S_0)/m$, S_0 is the adsorbate concentration in the absence of ligand L and in the presence of solid PCM, that is, the water solubility of S, S_t is the adsorbate concentration in solution and L_t is the total concentration of ligand added to the system. Combining these equations and applying them to known concentrations ($n=1$ and $m=1$) leads to the expression:

$$S_t = S_0 + \frac{K_{1:1}S_0L_t}{1 + K_{1:1}S_0} \quad (5)$$

By plotting S_t vs L_t , a straight line with intercept equal to S_0 is obtained, where the slope is defined by eq. 6:

$$\text{slope} = \frac{K_{1:1}S_0}{1 + K_{1:1}S_0} \quad (6)$$

Therefore, as stated earlier, the stability constant is given by the eq. 7:

$$K_{1:1} = \frac{\text{slope}}{S_0(1 - \text{slope})} \quad (7)$$

The graphical representation of PCM solubility (S_t) as a function of the added CD (L_t) gave a straight line in accordance to the mechanistic formation of a 1:1 PCM:CD complex (Fig. 4). In particular, the rising part of the solubility curve of PCM with α -CD was used for the calculation of K_c (Eq. 7). K_c values obtained for the PCM-CD complexes using β -CD, α -CD, HP β and RAME β cyclodextrins are shown in Table 2. The fitting was reasonably good, especially for RAME β and H β -CDs. The type of diagram found for α -CD can also indicate the formation of a more complex stoichiometry (Pérez-Martínez et al. 2000). In this case, a larger deviation in the goodness of the fit would have been noticed; however, a similar R^2 value to that of β -CD was obtained.

Fig. 4 Solubility diagrams of picloram in aqueous solutions in the presence of CD: (●) β -CD, (▲) RAME β , (▼) HP β and (■) α -CD.

Solubility efficiency (S_e) was also obtained from the solubility diagrams (Table 2), which is defined as the increment in the apparent solubility of PCM at a fixed CD concentration with respect to its solubility in the absence of CD. The inclusion complex of PCM with β -CD had the highest K_c value; however, due to the lower solubility of β -CD, the total concentration of solubilized PCM after the addition of β -CD was low compared to that reached with the addition of HP β or RAME β . The maximal increment in PCM solubility occurred with RAME β (up to 2.5-fold). The enhanced solubility of the herbicide was also remarkable with HP β , with solubility increments up to 2-fold. The percentage of encapsulation efficiency (E_{encap}) of PCM in CDs were obtained as $[S_m L_n] \times 100 / L_t$ at a fixed CD concentration (Table 2). In general, low encapsulation efficiency was obtained for the whole CDs evaluated.

Table 2 Apparent stability constants (K_c), regression coefficients (R^2), solubility increase (S_e , using 0–0.016 M for β -CD and 0–0.1 M for the other CD) and percentage of encapsulation efficiency (E_{encap}) for the inclusion compounds obtained between picloram and the different CDs. K_c is calculated based on the concentration of the species in equilibrium and not in terms of activities, which is indicated by the units in parentheses.

PCM adsorption on PILC in the presence of cyclodextrins

In the synthesis of delivery or slow release materials it is desirable to make that the content ratio of active ingredient / formulated mass has an appropriate value to the purpose thereof.

In the preparation of slow release formulations based on Al-Fe pillared clay minerals, the a.i. content of the formulation may be optimized by the use of ligands such as CDs that enhance the solubility of the herbicide in solution and on its turn permit to increase its sorption on the clay matrix.

This approach was tested with the pillared clay mineral P0.5 in the presence of RAME β and HP β CDs, which turned out to be the most effective for maximizing the concentration of PCM solution. The choice of P0.5 as adsorbent material was based on its higher sorption capacity (Fig. 3).

The PCM adsorption isotherms on P0.5 in the presence of RAME β and HP β obtained were shown in Figure 3, as well as the fit of the experimental data by adsorption models of Langmuir and Freundlich. The values of the fitting parameters to the experimental data of PCM adsorption isotherms on P0.5 in the presence of RAME β and HP β obtained from modeling by Langmuir and Freundlich are listed in table 1.

With the Langmuir model, the coincidence of K_L value obtained for PCM adsorption on P0.5 in the absence of cyclodextrins with that in the presence of RAME β and HP β (1460 M^{-1}) indicated that

cyclodextrins did not modify the mechanism of PCM interaction with the surface of the pillared clay mineral.

The maximum coating $\Gamma_{\max}$ increased slightly from 0.37 to 0.38 mol/kg, in presence of HP β . These results also pointed that the presence of HP β only increased the PCM concentration in solution but without participating in the adsorption process, achieving higher Γ_{ads} values than in the absence of CD.

The maximum coating $\Gamma_{\max}$ value was about 30% lower in the presence of RAME β than in the presence of HP β (Fig. 3 and Table 1). This result may indicate that the RAME β cyclodextrin was adsorbed onto the adsorbent surface, which resulted by either the competition with the PCM and occupation of the same sites or by steric hindrance preventing the PCM access to adsorption sites of the pillared clay. A RAME β strong interaction with bentonite, illite and kaolinite clay mineral surfaces were reported (Jozefaciuk et al. 2001), causing a decrease in surface area of the micro and mesopores.

From the Langmuir constant value K_L the standard free energy adsorption process was calculated $\Delta G^\circ = -18 \text{ kJmol}^{-1}$, which is an indicative of a spontaneous adsorption process.

These results indicated that the use of CDs enhanced the PCM solubility increasing its maximal adsorbed amount on P0.5 by the use of HP β . Therefore, the use of HP β is recommended in the preparation of PCM-Al-Fe PILCs formulations.

PCM adsorption in the presence of electrolytes

The use of HP β increased PCM sorption on P0.5 but it did not affect the mechanisms involved on its sorption on the clay because the analysis of the K_L values showed identical affinity. Then, the only function of this CD was to enhance the solubilization of the herbicide by the formation of an inclusion complex which is followed by its disruption and sorption of PCM. To elucidate the mechanisms involved on PCM sorption on P0.5, a study was performed in presence of competitive anions whose interaction modes with PILC were previously established. No influence of the HP β presence is expected, and for simplicity, this analysis was performed with PCM solutions in the absence of HP β .

PCM adsorption on P0.5 sample in the presence of chloride, sulfate and phosphate anions was also studied (Fig. 5). The presence of chloride ions did not change the amount of PCM adsorbed while the presence of sulfate and phosphate anions decreased drastically the adsorption of the herbicide.

Fig. 5 PCM adsorption on P0.5 sample in: the absence (a) or in the presence of the anions chloride (b), sulfate (c) and phosphate (d). The molar ratio used was 1:1.

Adsorption from strong anions such as Cl^- , NO_3^- , Br^- and ClO_4^- to functional groups on the surface of mineral acids occurs through outer sphere complexation processes driven mainly by electrostatic forces (Essington 2004).

The adsorption of phosphate ions on metal oxides has been reported to occur by inner sphere complexation processes, interacting with a metal center via formation of monodentate or bidentate complexes or with two adjacent metal centers by forming a bridge complex (He et al. 1997, Higuchi & Connors 1965)

On the contrary, the adsorption of sulfate anions on mineral surfaces such as γ -alumina and kaolinite was proposed to occur as outer sphere complexes (He et al. 1997). On soils, it was described as inner sphere complexation which was correlated with the soil iron oxides content (Kooner et al. 1995), which was in agreement with hematite oxides results (Eggleson et al. 1998). Adsorption of sulfate on mineral clays was assigned to the formation of inner sphere complex through one surface site or through a bridge complex (Sarbak 1994).

The results obtained for the PCM adsorption on P0.5 sample in the presence of different anions indicated competition on the adsorption sites when phosphate or sulfate were present. PCM adsorption on raw and pillared clay minerals were reported to occur by the formation of inner sphere complexes between PCM molecules and the mineral surface through the interaction of the carboxylic and pyridine nitrogen moieties of PCM and the metal centers of the pillars on the mineral surface (Marco-Brown et al. 2014, Marco-Brown et al. 2012, Marco-Brown et al. 2015). Therefore, these results are consistent with a PCM adsorption onto the surface of the PILC by forming an inner sphere complex with the metal centers of silicon, aluminum or iron.

A FTIR study was carried out in order to get an insight into the mechanisms that operate in the preparation of SRFs.

FTIR analysis

The FTIR spectra of PCM, HP β , P0.5, PCM-P0.5, PCM-HP β , PCM-HP β (PM) and PCM-HP β -P0.5 are shown in Figures 6 and 7. The band assignation of PCM, HP β and P0.5 are listed in Table 3 and Table 4.

Fig. 6 FT-IR spectra of: (—) PCM, (—) HP β , (—) P0.5, (—) HP β -P0.5, (—) PCM-P0.5, (—) PCM-HP β , (—) PCM-HP β (PM) and (—) PCM-HP β -P0.5 samples. Spectral region evaluated from 400 to 4000 cm^{-1} .

Fig. 7 FT-IR spectra of: (—) PCM, (—) HP β , (—) P0.5, (—) HP β -P0.5, (—) PCM-P0.5, (—) PCM-HP β , (—) PCM-HP β (PM) and (—) PCM-HP β -P0.5 samples. Spectral region evaluated from 400 to 1800 cm^{-1} .

426

427 **Table 3** Group assignments to the mean FT-IR peaks of PCM and HP β samples

428 **Table 4** Group assignments to the mean FT-IR peaks of P0.5 sample

429

430 **Interaction between PCM and P0.5 (PCM-P0.5 sample)**

431 The band of PCM at 1706 cm⁻¹ was not present which indicated that protonated carboxylic form of
432 PCM was absent in PCM-P0.5 sample. The PCM band at 1600 cm⁻¹ shifted to 1623 cm⁻¹ which
433 revealed interaction of PCM carboxylic group with the surface. The band of PILC at 1635 cm⁻¹
434 corresponding to interlayer water disappeared or diminished its intensity due to PCM entrance into
435 the clay galleries and the exit of surface water. The band of PCM at 1456 cm⁻¹ corresponding to
436 C=C pyridine-ring stretching vibrations shifted to 1427 cm⁻¹ in the adsorbed sample which
437 confirmed the PCM interaction with clay surface through pyridine nitrogen. The band of PCM at
438 1381 cm⁻¹ assigned to symmetric stretching vibration of the dissociated carboxyl group shifted to
439 1367 cm⁻¹ due to PCM interaction through its carboxylic group with the clay surface. All these
440 shifts could be attributed to the interaction of the PCM with the solid surface by formation of a
441 monodentate or bidentate coordination that involves the COO⁻ group and the nitrogen of the
442 pyridine ring as previously suggested for picolinic acid interactions with Al(III)-(hydr)oxide
443 minerals (Loring et al. 2000).

444 **PCM-HP β complex**

445 The bands of PCM at 1706 , 1600, 1238 and 906 cm⁻¹ corresponded to C=O stretching vibrations,
446 COO⁻ stretching and N-H bending vibrations, to C-N stretching vibrations and to N-H wagging
447 vibrations, respectively. These bands shifted to 1711, 1605, 1242 and 911 cm⁻¹ respectively in the
448 PCM-HP β complex due to PCM interactions with the CD. The bands of PCM at 1456, 1066, 991,
449 760 and 731 cm⁻¹ overlapped with CD bands. All the remaining bands of PCM were visible in the
450 PCM-HP β spectrum. As expected, an interaction between PCM and HP β occurred when both were
451 present in the aqueous solution, where PCM was located partly in the hydrophobic cavity of HP β .

452 **HP β -P0.5 sample**

453 HP β bands at 854, 1032, and 2929 cm⁻¹ appeared in HP β -P0.5 sample which indicated the absence
454 of HP β interaction with the clay surface. This behavior supported the previous PCM adsorptions
455 results where there was no difference between PCM adsorption in the presence or absence of HP β .

456 **Physical mixture of PCM and HP β (PCM-HP β (PM) sample)**

457 As expected the FTIR spectrum of the PCM and CD physical mixture was the sum of the separate
458 IR spectrums of PCM and CD. The absence of absorption peak shifts indicated no interaction
459 between both molecules when mixed physically.

460 **PCM adsorbed on P0.5 surface in presence of HP β (PCM-HP β -P0.5 sample)**

The shifts of the absorption bands observed are similar to those observed with the PCM-P0.5 sample. The bands of PCM at 1706 and 1600 cm^{-1} corresponding to C=O stretching vibrations and COO- stretching which shifted to 1698 (shoulder) and 1623 cm^{-1} respectively, indicating interaction of PCM carboxylic group with the surface. The band of PCM at 1456 cm^{-1} which corresponded to C=C pyridine-ring stretching vibrations shifted to 1427 cm^{-1} confirming an interaction of PCM with mineral surface through its pyridine nitrogen. The band of PCM at 1381 cm^{-1} was assigned to symmetric stretching vibration of the dissociated carboxyl group shifted to 1367 cm^{-1} due to PCM interaction through carboxylic group with clay surface groups. These results are in agreement with the formation of an inner-sphere complex through a bidentate interaction. The absence of any shift in the HP β bands at 1412, 1333, 1300, 1157, 1082, 947, 754 and 704 cm^{-1} in PCM-HP β -P0.5 sample, indicated no interaction between HP β and the clay surface, as stated previously. This behavior was also supported by the analysis of PCM adsorption in the presence/absence of HP β by the Langmuir model, and in line with findings of Jozefaciuk et al. (Jozefaciuk et al. 2001), who reported a very low affinity between HP β and clay surfaces. These results indicated PCM adsorption in the absence of PCM-HP β complex on the clay surface. On the one hand, the presence of HP β contributed to increase the PCM concentration in solution and on the other hand, the great affinity of PCM with the clay surface together with the low apparent stability constant of the PCM-HP β complex resulted in the rupture of the complex, thus increasing PCM loading on the clay surface.

Summarizing, the interaction between PCM and HP β molecules was established by FTIR analyses. The interaction absence between HP β and pillared mineral surface it was also confirmed from the adsorption study. FTIR results indicated that, even in the presence or absence of HP β , PCM interacted with clay surface sites through its carboxylic and pyridine nitrogen groups.

Release studies

PCM elution tests were made on a sandy soil by adding the PCM-P0.5 and PCM-HP β -P0.5 SRFs and the results were compared to those of a commercial solution applied at same rate. PCM concentration, expressed as % of PCM applied, in the eluates after 20 consecutive irrigations steps with water is shown in Figure 8. The commercial formulation showed a sharp peak indicative of a very rapid elution of PCM with almost negligible retention in this soil. On the contrary, the released amounts with the pillared-clay formulations showed flattened peaks consistent with a gradual release of the active ingredient.

After 13 irrigation steps, the commercial formulation eluted completely ($97.0 \pm 8.2\%$), whereas the total recoveries were: $70.1 \pm 12.1\%$ for PCM-HPB-0.5 and $76.9 \pm 9.8\%$ for PCM-P-0.5 samples, respectively, amounting to a total reduction of releases of 28 and 22%, respectively. The curves for the pillared clay based formulations were quite close to each other in agreement with the same mechanism of retention (electrostatic interactions with the pillars); the only difference between them was the higher a.i. content on PCM-HPB-0.5 (19% in excess).

Fig. 8 Herbicide release from the commercial and PCM-clay formulations.

Conclusions

Pillared clay mineral were synthesized varying the Fe/(Fe+Al) molar ratio in order to design slow release formulation of the herbicide picloram based on the high affinity of PCM observed with iron oxide surfaces. Therefore, an enhanced adsorption of PCM on iron modified clay and a slow release of it was achieved. The P0.5 sample evidenced a higher adsorption capacity of picloram than that found for P0.15 sample.

An increase in the loading of PCM on samples was achieved by using CDs as solubilizing agent due to the formation of inclusion complexes. The use of these complexes increased the amount of PCM adsorbed on the pillared clay. **The use of HP β in the preparation of PCM-Al-Fe PILCs formulations is recommended for the slow release of PCM.**

The interaction between PCM and HP β molecules was established by FTIR analyses. It was confirmed **a lack of** interaction between HP β and pillared mineral surface from the adsorption study. FTIR results suggested that, PCM interacted with clay surface sites through its carboxylic and pyridine nitrogen groups. Competitive adsorption of anions confirmed an inner-sphere complexation between PCM and clay surface. Release of the herbicide in a sandy soil from PCM-clay formulations was retarded compared to the commercial formulation Tordon®101.

References

- Akelah A (1996) Novel utilizations of conventional agrochemicals by controlled release formulations. *Mater. Sci. Eng. C* 4:83-98
- Ali I, Asim M, Khan TA (2012) Low cost adsorbents for the removal of organic pollutants from wastewater. *J. Environ. Manage.* 113:170-183
- Alromeed AA, Scrano L, Bufo SA, Undabeytia T (2015) Slow-release formulations of the herbicide MCPA by using clay-protein composites. *Pest Manag. Sci.* 71:1303-1310
- Biggar JW, Mingelgrin U, Cheung MW (1978) Equilibrium and kinetics of adsorption of picloram and parathion with soils. *J. Agr. Food Chem.* 26:1306-1312
- Campos EVR, de Oliveira JL, Fraceto LF, Singh B (2014) Polysaccharides as safer release systems for agrochemicals. *Agron. Sustain. Dev.* 35:47-66
- Carrizosa MJ, Calderón MJ, Hermosín MC, Cornejo J (2000) Organosmectites as sorbent and carrier of the herbicide bentazone. *Sci. Total Environ.* 247:285-293
- Celis R, Hermosin MC, Cornejo L, Carrizosa MJ, Cornejo J (2002) Clay-herbicide complexes to retard picloram leaching in soil. *Int. J. Environ. An. Ch.* 82:503-517
- Cheung MW, Biggar JW (1974) Solubility and Molecular-Structure of 4-Amino-3,5,6-Trichloropicolinic Acid in Relation to Ph and Temperature. *J. Agr. Food Chem.* 22:202-206
- Delcour I, Spanoghe P, Uyttendaele M (2015) Literature review: Impact of climate change on pesticide use. *Food Res. Int.* 68:7-15
- Eggleston CM, Hug S, Stumm W, Sulzberger B, dos Santos Afonso M (1998) Surface complexation of sulfate by hematite surfaces: FTIR and STM observations. *Geochim. Cosmochim. Ac.* 62:585-593
- Entry JA, Sojka RE (2008) Matrix based fertilizers reduce nitrogen and phosphorus leaching in three soils. *J. Environ. Manage.* 87:364-372

- Essington ME (2004): Soil and Water Chemistry, An Integrative Approach. CRC Press, Boca Raton, USA
- FAO FaAOotUN (2007): Specifications and evaluations for Agricultural Pesticides, Picloram. <http://www.fao.org>
- Galán-Jiménez MC, Mishael YG, Nir S, Morillo E, Undabeytia T (2013) Factors Affecting the Design of Slow Release Formulations of Herbicides Based on Clay-Surfactant Systems. A Methodological Approach. PLoS ONE 8:e59060
- Gupta VK, Suhas (2009) Application of low-cost adsorbents for dye removal - A review. J. Environ. Manage. 90:2313-2342
- Hall KE, Ray C, Ki SJ, Spokas KA, Koskinen WC (2015) Pesticide sorption and leaching potential on three Hawaiian soils. J. Environ. Manage. 159:227-234
- Haynes D, Müller J, Carter S (2000) Pesticide and herbicide residues in sediments and seagrasses from the Great Barrier Reef World Heritage Area and Queensland Coast. Mar. Pollut. Bull. 41:279-287
- He LM, Zelazny LW, Baligar VC, Ritchey KD, Martens DC (1997) Ionic strength effects on sulfate and phosphate adsorption on γ -alumina and kaolinite: Triple-layer model. Soil Sci. Soc. Am. J. 61:784-793
- Higuchi T, Connors K (1965): Phase solubility techniques. In: CN R (Editor), Advances in Analytical Chemistry Instrumentation. Interscience, New York, NY, pp. 117-212
- Jozefaciuk G, Muranyi A, Fenyvesi E (2001) Effect of cyclodextrins on surface and pore properties of soil clay minerals. Environ. Sci. Technol. 35:4947-4952
- Kashyap PL, Xiang X, Heiden P (2015) Chitosan nanoparticle based delivery systems for sustainable agriculture. Int. J. Biol. Macromol. 77:36-51
- Kenawy ER, Sherrington DC, Akelah A (1992) Controlled release of agrochemical molecules chemically bound to polymers. Eur. Polym. J. 28:841-862
- Kooner ZS, Jardine PM, Feldman S (1995) Competitive surface complexation reactions of sulfate and natural organic carbon on soil. J. Environ. Qual. 24:656-662
- Lagaly G (2001) Pesticide-clay interactions and formulations. Appl. Clay Sci. 18:205-209
- Loewy M, Kirs V, Carvajal G, Venturino A, Pechen De D'Angelo AM (1999) Groundwater contamination by azinphos methyl in the Northern Patagonic Region (Argentina). Sci. Total Environ. 225:211-218
- Loring JS, Karlsson M, Fawcett WR, Casey WH (2000) Attenuated total reflection-fourier-transform infrared and ^{27}Al -nuclear magnetic resonance investigation of speciation and complexation in aqueous Al(III) -picolinate solutions. Geochim. Cosmochim. Ac. 64:4115-4129
- Maqueda C, Villaverde J, Sopeña F, Undabeytia T, Morillo E (2008) Novel system for reducing leaching of the herbicide metribuzin using clay-gel-based formulations. J. Agr. Food Chem. 56:11941-11946
- Maqueda C, Villaverde J, Sopeña F, Undabeytia T, Morillo E (2009) Effects of soil characteristics on metribuzin dissipation using clay-gel-based formulations. J. Agr. Food Chem. 57:3273-3278
- Marco-Brown JL, Barbosa-Lema CM, Torres Sánchez RM, Mercader RC, dos Santos Afonso M (2012) Adsorption of picloram herbicide on iron oxide pillared montmorillonite. Appl. Clay Sci. 58:25-33
- Marco-Brown JL, Areco MM, Torres Sánchez RM, dos Santos Afonso M (2014) Adsorption of picloram herbicide on montmorillonite: Kinetic and equilibrium studies. Colloid. Surface A 449:121-128
- Marco-Brown JL, Trinelli MA, Gaigneaux EM, Torres Sánchez RM, Dos Santos Afonso M (2015) New insights on the structure of the picloram-montmorillonite surface complexes. J. Colloid Interf. Sci. 444:115-122

- Mishra T, Mohapatra P, Parida KM (2008) Synthesis, characterisation and catalytic evaluation of iron-manganese mixed oxide pillared clay for VOC decomposition reaction. *Appl. Catal., B* 79:279-285
- Morillo E, Sánchez-Trujillo MA, Villaverde J, Madrid F, Undabeytia T (2014) Effect of contact time and the use of hydroxypropyl- β -cyclodextrin in the removal of fluorene and fluoranthene from contaminated soils. *Sci. Total Environ.* 496:144-154
- Nir S, Undabeytia T, Yaron-Marcovich D, El-Nahhal Y, Polubesova T, Serban C, Rytwo G, Lagaly G, Rubin B (2000) Optimization of adsorption of hydrophobic herbicides on montmorillonite preadsorbed by monovalent organic cations: Interaction between phenyl rings. *Environ. Sci. Technol.* 34:1269-1274
- Osteryoung J, Whittaker JW (1980) Picloram - Solubility and Acid-Base Equilibria Determined by Normal Pulse Polarography. *J. Agr. Food Chem.* 28:95-97
- Palma G, Sanchez A, Olave Y, Encina F, Palma R, Barra R (2004) Pesticide levels in surface waters in an agricultural-forestry basin in Southern Chile. *Chemosphere* 57:763-770
- Pérez-Martínez JI, Ginés JM, Morillo E, Moyano JR (2000) ^1H -nuclear magnetic resonance and phase solubility studies of the stoichiometries in 2,4-D: α - and β -cyclodextrins inclusion complexes. *J. Inclusion Phenom.* 37:171-178
- Qi Y, Donahoe RJ (2008) The environmental fate of arsenic in surface soil contaminated by historical herbicide application. *Sci. Total Environ.* 405:246-254
- Sánchez-Jiménez N, Sevilla MT, Cuevas J, Rodríguez M, Procopio JR (2012) Interaction of organic contaminants with natural clay type geosorbents: Potential use as geologic barrier in urban landfill. *J. Environ. Manage.* 95, Supplement:S182-S187
- Sarbak Z (1994) Structural and surface properties of anion-activated clay. *Mater. Chem. Phys.* 39:91-97
- Schipper PNM, Vissers MJM, van der Linden AMA (2008): Pesticides in groundwater and drinking water wells: Overview of the situation in the Netherlands, *Water Science and Technology*, pp. 1277-1286
- Szejtli J (1998) Introduction and general overview of cyclodextrin chemistry. *Chem. Rev.* 98:1743-1753
- Undabeytia T, Sopena F, Sánchez-Verdejo T, Villaverde J, Nir S, Morillo E, Maqueda C (2010) Performance of slow-release formulations of alachlor. *Soil Sci. Soc. Am. J.* 74:898-905
- Undabeytia T, Recio E, Maqueda C, Morillo E, Gómez-Pantoja E, Sánchez-Verdejo T (2011) Reduced metribuzin pollution with phosphatidylcholine-clay formulations. *Pest Manag. Sci.* 67:271-278
- Undabeytia T, Recio E, Maqueda C, Sánchez-Verdejo T, Balek V (2012) Slow diuron release formulations based on clay-phosphatidylcholine complexes. *Appl. Clay Sci.* 55:53-61
- Undabeytia T, Galán-Jiménez MC, Gómez-Pantoja E, Vázquez J, Casal B, Bergaya F, Morillo E (2013) Fe-pillared clay mineral-based formulations of imazaquin for reduced leaching in soil. *Appl. Clay Sci.* 80-81:382-389
- Wang Y, Gao Z, Shen F, Li Y, Zhang S, Ren X, Hu S (2015) Physicochemical characteristics and slow release performances of chlorpyrifos encapsulated by poly(butyl acrylate- co -styrene) with the cross-linker ethylene glycol dimethacrylate. *J. Agr. Food Chem.* 63:5196-5204
- Wenz G, Han BH, Müller A (2006) Cyclodextrin rotaxanes and polyrotaxanes. *Chem. Rev.* 106:782-817
- Yan H, Chen X, Feng Y, Xiang F, Li J, Shi Z, Wang X, Lin Q (2015) Modification of montmorillonite by ball-milling method for immobilization and delivery of acetamiprid based on alginate/exfoliated montmorillonite nanocomposite. *Polym. Bull.* 73:1185-1206
- Yáñez C, Cañete-Rosales P, Castillo JP, Catalán N, Undabeytia T, Morillo E (2012) Cyclodextrin inclusion complex to improve physicochemical properties of herbicide bentazon: Exploring better formulations. *PLoS ONE* 7:e41072

1
2
3
4 641
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60
61
62
63
64
65

Fig. 1

Fig. 2

Fig 3

Fig. 3

Fig 4

Fig. 4

Fig. 5

Fig. 6

Fig. 7

Fig. 8

Table 1 Langmuir and Freundlich parameters for PCM adsorption onto P0.15 and P0.5 samples in presence and absence of CDs

Adsorbent	Langmuir			Freundlich		
	K_L (M^{-1})	Γ_{max} (mol/kg)	R^2	K_F	$1/n$	R^2
P0.15	410	0.19	0.9677	6.45	0.50	0.9570
P0.5	1460	0.37	0.9794	5.64	0.52	0.9728
P0.5+RAME β	1460	0.29	0.9862	9.97	0.40	0.9253
P0.5+HP β	1460	0.38	0.9714	15.49	0.38	0.9725

Table 2 Apparent stability constants (K_c), regression coefficients (R^2), solubility increase (S_e , using 0-0.016 M for β -CD and 0–0.1 M for the other CD) and efficiency of encapsulation (E_{encap}) for the inclusion compounds obtained between picloram and the different CDs. K_c is calculated based on the concentration of the species in equilibrium and not in terms of activities, which is indicated by the units in parentheses

Parameter	RAME β -CD	HP β -CD	α -CD	β -CD
K_c (L/mmol)	1.90×10^{-2}	1.05×10^{-2}	5.28×10^{-3}	1.97×10^{-2}
S_e	2.65	2.13	1.35	1.34
E_{encap} (%)	3.8	2.3	1.0	4.4
R^2	0.9880	0.9437	0.8086	0.7305

Table 3 Group assignments to the mean FT-IR peaks of PCM, HP β and P0.5 samples

Sample	Band (cm ⁻¹)	Assignment
PCM	3475	N-H stretching
	3373	N-H stretching
	1706	C=O stretching
	1600	COO ⁻ stretching and N-H bending
	1539	C=N pyridine-ring stretching
	1456	C=C pyridine-ring stretching
	1435	C=C pyridine-ring stretching
	1381	COO ⁻ stretching
	1306	C-Cl in-plane bending
	1284	C-Cl in-plane bending
	1238	C-N stretching
	1107	C-Cl in-plane bending
	1066	C-Cl in-plane bending
	991	C-Cl out-of-plane bending
	906	N-H wagging
	760	C-Cl stretching
	731	C-Cl stretching
HP β	3398	O-H stretching
	2929	CH ₂ stretching
	1645	H-O-H bending
	1458	C-H bending from CH ₂ and CH ₃
	1412	C-H bending from CH ₂ and CH ₃
	1369	C-H bending from CH ₃
	1334	C-C-H and C-O-H bending H-C-H bending
	1300	C-H in-plane bending
	1244	C-H in-plane bending
	1201	C-H in-plane bending
	1157	C-O-C stretching/O-H bending
	1082	coupled C-O/C-C stretching/O-H bending
	1032	coupled C-O/C-C stretching/O-H bending
	947	C-H out-of-plane bending
	854	C-H out-of-plane bending
	756	C-H wagging
	708	C-H wagging

Table 4 Group assignments to the mean FT-IR peaks of P0.5 sample

Sample	Band (cm ⁻¹)	Assignment
P0.5	3612	O-H stretching vibration
	3420	O-H stretching vibration
	1635	interlayer water
	1049	in-plane Si-O-Si stretching vibration
	920	Al-Al-OH deformation
	890	Si-O vibrations of some quartz impurities of the raw mineral
	797	Si-O vibrations of quartz
	777	Si-O vibrations of quartz