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SORPTION OF AROMA COMPOUNDS IN PET AND PVC DURING THE STORAGE OF A STRAWBERRY SYRUP

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2 **Sorption of aroma compounds in PET and PVC during the storage of a**
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4 **strawberry syrup**
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Abstract

The sorption of 14 aroma compounds into PET and PVC was monitored during storage of a strawberry syrup for 1 year. Concentrations in the syrup and in the polymer were determined during storage and compared with previously published results obtained with glass bottles. Apparent partition coefficients between the polymer and the syrup (noted K_{app}) were estimated from experimental kinetics without reaching equilibrium K_{app} values were optimally identified from the kinetic data obtained between 30 and 90 days. They exhibited a similar behaviour for both polymers with values were between $2 \cdot 10^{-5}$ and $2 \cdot 10^{-3}$, $4 \cdot 10^{-5}$ and $3 \cdot 10^{-2}$ respectively for PET and PVC. The variation of K_{app} values in PET was mainly correlated to the polarity of tested compounds as assessed by their logP values. By contrast, the variations in K_{app} values for PVC were mainly related to their chain lengths. Due to slightly higher partition coefficients and diffusion coefficients in PVC compared to PET, the amount of absorbed aroma was 4 times higher in PVC than in PET, however the amount of absorbed aroma compounds was less than 0.1% of the initial amount present into the syrup, except for octyl butanoate.. The variation in concentration in the syrup was interpreted as a combination of a degradation process and a transport process into the packaging material. Both effects were particularly noticeable for both PET and unstable aroma compounds.

Introduction

The absorption of aroma compounds by plastic materials may induce both a weakening of flavour and changes to the organic profile of packaged food products. This has mainly been reported for aqueous food, such as fruit juices, or model solutions, where the partition coefficient of aromas between food and packaging strongly favours sorption into the plastic material (Kwapong & Hotchkiss 1987, Sizer et al. 1988). However, there are many discrepancies in reported partition coefficient values (Kutty et al. 1994, Hernandez-Munoz et al. 2001, Tehrani & Desobry 2004). The possible reasons include: i) sorption kinetic studies performed without reaching the thermodynamic equilibrium (Gavara et al. 1996, Lebossé et al. 1997, Feigenbaum et al. 1998), ii) aroma used beyond their concentrations limits, which yield to deviations from Henry sorption equilibrium. Besides, partitioning results obtained with model solutions are difficult to extrapolate to the behaviour of aroma compounds in real foods, which are generally multi-constituent and possibly multi-phase. Recent papers have shown that in complex solutions simulating food matrices, the partition coefficients between aroma compounds and polymers depend also on the interactions with macromolecules (lipids, polyoligosides or proteins). In particular, lipids compete with polymers concerning the transfer of aroma compounds into polyolefins (Van Willige et al. 2000a,b). Moreover, during long periods of storage of up to a year, changes in aroma composition were detected in the food matrix without any contact with the polymer (Ducruet et al. 2001, Berlinet et al. 2005). In this case it is important to understand how the reactivity of the aroma compound in the matrix can influence the absorption of the aroma compounds by the packaging material.

Although PET and PVC are known to be good barriers to oxygen, in comparison with polyolefins, PET (one of the most common packaging polymers) and PVC (which is not widely used to package flavoured drinks), have seldom been tested (Ducruet et al. 2001, Van Willige et

1
2 al. 2002a, Berline et al. 2005) for their barrier properties to aroma. The objective of the present
3
4 study was to provide additional results on the sorption of aroma compounds in glassy polymers
5
6 such as PET and PVC during long storage periods and to compare the kinetic sorption process
7
8 with the degradation process, which also can reduce the amount of aroma. To make the results
9
10 valuable for industry, the sorption and degradation kinetics were assessed on real food products
11
12 and real packaging materials. Since the thermodynamic equilibrium cannot be obtained when the
13
14 rate constants of both processes have similar orders of magnitude, the apparent partition
15
16 coefficients were derived from an approach based on a new approximated solution proposed by
17
18 Vitrac and Hayert (2006) to identify transport properties (including the partition coefficient) from
19
20 censored kinetics (i.e. without reaching the thermodynamic equilibrium).
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29 **Materials and methods**

30 *Packaging materials*

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32 All the bottles were supplied from Teisseire S.A France. Strawberry syrups were packaged in rigid
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34 PVC bottles (PVC1, PVC2: average thickness: 1 mm) and in amorphous PET bottles (PET1 and
35
36 PET2; average thickness: 0.75 mm). PVC1 was made using GFG 52 D resin from Dorly; it is
37
38 suitable for contact with oils and is stabilized with an organotin stabilizer. PVC2 was
39
40 manufactured by Teisseire. PET1 was manufactured from Ramapet, and PET2 from Shinpet. The
41
42 surface area / volume in $\text{dm}^2 \text{ l}^{-1}$ ratio was approximately = 6. Packaging in glass bottles (GL) was
43
44 used as a reference to test the stability of the syrup without the influence of polymer contact.
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50

51 *Strawberry syrup*

52
53 Strawberry syrup was obtained by mixing a sucrose syrup ($^{\circ}\text{Brix}$: 64.5 +/- 1) and fruit juices
54
55 (strawberry, elder and lemon juices). Nature-identical flavouring substances were also included.
56
57
58
59 The pH of the strawberry syrup was 2.65 +/- 0.2 and its shelf life was 18 months.
60

1
2 After flash-pasteurisation at 105°C for 30 sec, syrup samples were packaged at ambient
3
4 temperature under aseptic conditions. All bottles (PET, PVC and glass) were sealed after filling
5
6 with aluminium foil in order to avoid a cap effect. The packaged syrups were stored at 20°C
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8

9
10 *Solvent extraction of the syrup*

11
12 200 ml of syrup, 200 ml of Milli Q water and 500 µl of tridecane solution (used as an internal
13
14 standard, 500 µl l⁻¹) were mixed in a 1 litre flask. Volatile compounds in the strawberry syrup
15
16 were extracted three times using distilled dichloromethane, with 80, 30 and 30 ml for 30 min, 15
17
18 min and 15 min respectively, at 0° C while stirring. Organic phases were separated from the
19
20 aqueous phase and pooled. The dichloromethane extracts were dried over anhydrous Na₂SO₄ and
21
22 concentrated using a Kuderna- Danish column. Analysis were done in triplicate.
23
24
25

26
27 *Solvent extraction of polymers*

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29
30 At the end of each period of contact, the strawberry syrup and its packaging were separated. The
31
32 plastic bottle (PET or PVC) was carefully and rapidly rinsed twice with ultra pure water, then with
33
34 ethanol, and finally wiped before cutting. Because of the heterogeneous thickness of bottles,
35
36 samples were taken by cutting pieces from three sections of a bottle (top, bottom and middle) and
37
38 then pooled. Approximately 7g of each pooled sample was then plunged immediately into a 100
39
40 ml flask containing distilled diethyl ether (95 ml). 2 ml of a solution of ethyl pentanoate in diethyl
41
42 ether (100 µl l⁻¹) were added as an internal standard. The flasks were tightly plugged with Teflon
43
44 caps and extraction was carried out for 2 hours at 15 °C with magnetic stirring. Solvent extracts
45
46 were dried with sodium sulfate and filtered over glass wool. Diethyl ether extracts were
47
48 concentrated using a Kuderna- Danish apparatus.
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54 Two plastic bottles were analyzed for each contact period. The extraction process was repeated
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56 twice for each bottle so as to enable analysis in duplicate of the plastic material after each period
57
58 of contact.
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Mass spectra

Identification of the constituents was achieved using gas chromatography-mass spectrometry (GC-MS). The GC-MS system consists of a Fisons GC-8000 chromatograph and a MD 800 mass spectrometer (Fisons Instruments, Les Ulis, France). Separations were performed on a Supelcowax fused silica capillary column (0.32 mm x 30 m; 0.5 μ m; Supelco, Bellefonte, PA). The linear flow velocity of helium was 32 cm sec⁻¹. The column was maintained at 30°C for 10 min and then programmed at 240°C for SPME analysis and 260°C for solvent extracts at 5°C min⁻¹. The split-splitless injector was at 230°C. Solvent extracts were injected into the splitless/split system (the split valve was closed after 30 sec). Electron ionization mass spectra were recorded under the following conditions:

capillary direct interface, 250°C; ion source, 200°C; ionization voltage, 70eV; mass range, 29-300 m/z; electron multiplier voltage, 450 V; scan rate 3 scans sec⁻¹.

Mass spectral matches were made by comparison with NIST and INRAMASS mass spectra libraries. Kovat's Indices of authentic compounds compiled in the INRAMASS library were used to confirm identification.

Estimation of apparent partition coefficients

The partition coefficient is defined from the ratio of concentrations of aroma at thermodynamic equilibrium, noted $C_P|_{eq}$ and $C_F|_{eq}$ respectively in the packaging material (P) and in Food (F):

$$K = \frac{C_P|_{eq}}{C_F|_{eq}} \quad (1)$$

It is emphasized that equation (1) is valid whatever the considered equilibrium. Since the equilibrium was not reachable due to insufficient contact time (even after one year) or due to aroma degradation during storage, the equilibrium was extrapolated from the kinetic phase diagram corresponding to the experimental sorption kinetic, which was recently proposed by

Vitrac and Hayert (2006). Only the main features of KPD relevant for the current application are reported here. A kinetic phase diagram, noted KPD, consists in plotting the sorption mass flux or equivalently the variation of the residual concentration in P with time, $\frac{dC_P|_t}{dt}$ versus the residual concentration in P, $C_P|_t$. During a sorption experiment, the expected concentration in P at equilibrium corresponds in the KPD space $\left(C_P|_t, \frac{dC_P|_t}{dt}\right)$ to the value $(C_P|_{eq}, 0)$. The interesting feature of the KPD space for the interpretation of sorption kinetics governed by diffusion in P and diffusion-convection in F is that after an initial rapid non-linear decay of $\frac{dC_P|_t}{dt}$ with $C_P|_t$, the variation of $\frac{dC_P|_t}{dt}$ is almost linear with $C_P|_t$. As a result, it is possible in the KPD space to extrapolate linearly the theoretical equilibrium state. Besides, a possible competition between sorption in P and degradation in F is easily detected by a change in curvature of the KPD. Thus a change in the decay of the KPD slope $\frac{d}{dC_P|_t}\left(\frac{dC_P|_t}{dt}\right)$ is interpreted as a degradation process, which tends to dominate the sorption process.

In this work, $C_P|_{eq}$ was lineally extrapolated from the last experimental point KPD, for which

$\frac{d}{dC_P|_t}\left(\frac{dC_P|_t}{dt}\right)$ was monotonous (i.e. increasing or constant). The same process was applied for

the concentration in the liquid. A key step in the KPD approach is calculation of the derivatives of the concentration with time when the experimental data are unevenly sampled and noisy. In this work, we use an efficient non deterministic filtering technique based on local polynomial approximants as detailed and discussed in Vitrac and Hayert (2006). By noting with a superscript “extrap” the extrapolated quantities, the apparent partition coefficient, K_{app} , corresponding to experimental data were calculated from:

$$K_{app} = \frac{C_P|_{eq}^{extrap}}{C_F|_{eq}^{extrap}} \quad (2)$$

It is emphasized that K_{app} may differ from the real K value, however it is the best estimator according to the data available and our knowledge of sorption mechanisms. Confidence intervals on K_{app} were derived by Monte Carlo sampling consisting in adding a white noise similar to the experimental error (relative error of 15 %) to kinetic data and in repeating the whole analysis. Each confidence interval was based on the 2.5th and 97.5th percentiles of at least 200 Monte Carlo trials.

Statistical analysis

Sorption experiments were analyzed using a one-way ANOVA program. When the differences were significant ($p < 0.05$), Duncan's test was used to check the differences between pairs of groups and was carried out using XLSTAT-Pro 7.0 software (Addinsoft, Paris, France).

Results and discussion

Sorption of the aroma compounds into PET and PVC

In order to monitor the sorption of aroma compounds, the strawberry syrup and the plastic material were extracted and analysed at different time points during the 330-day storage period. In the case of PVC, the presence in solvent extracts of additives and small oligomers along with absorbed volatile compounds produced more complex GC profiles than those seen with PET. Some of these additives were tentatively identified in the extracts of the syrup. (Table 1).

[Insert Table 1 here]

These compounds originate from the degradation of additives for example plasticizer, lubricants or modifiers (Gachter and Muller, 1990). Some can migrate into the syrup such as benzaldehyde,

1 acetophenone and 2-ethylhexanol. By solvent extraction of the syrup or the packaging, these
2
3 additives were extracted and co-eluted with the aroma compounds by GC. Thus, of the 38 aroma
4
5 compounds previously identified in the syrup at the initial time of storage (Ducruet et al. 2001),
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7 only 14 compounds can be quantified in both kinds of packaging and in the syrup over the 330-
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9 day storage period (Table 2).
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[Insert Table 2 here]

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20 The final concentration of the aroma compounds into the syrups stored in both PET and PVC
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22 bottles were compared to their initial concentrations (Figure 1). During shelf life, aroma
23
24 compounds decreased by hydrolysis irrespective of the polymer. The decrease was more for short
25
26 chain esters, and in particular for acetic acid esters and ethyl hexanoate (Figure 1A) compared to
27
28 more stable compounds (Figure 1B). This reactivity is not related to the contact with polymer as it
29
30 was previously found to occur during the storage of the syrup in a glass bottle (Ducruet et al,
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32 2001).
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[Insert Figure 1 here]

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42 The greater reactivity was observed for short chain esters (methyl or ethyl esters) and acetic acid
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44 esters and can be explained by the low steric hindrance around their carbonyl group in contrast to
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46 linear longer chain esters (butyl, hexyl or octyl esters) and specially to branched esters (ethyl 2-
47
48 methylbutanoate, ethyl 3-methylbutanoate and 3-methylbutyl 3-methylbutanoate). These results
49
50 demonstrated the instability of the aroma formulation for long periods of storage in acidic
51
52 condition such as it was also found in the case of citrus flavoured drinks (Sizer et al. 1988,
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54 Berlinet et al. 2005)
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59
60 The sorption of the aroma compounds from strawberry syrup into PET and PVC was monitored

1
2 with time. The sorption of the 14 aroma compounds over 330 days of storage was compared to
3 their initial concentration into the syrup (Table 2). The sorption into both kinds of packaging
4 materials was found to be weak. The sorption was generally lower than 0.1 % in all cases except
5
6 for octyl butanoate which reached 0.15 % in both PET samples, and 0.57 % or 0.26 % respectively
7
8 in PVC1 and in PVC2 samples. After one year of storage into both samples of PET and PVC, if
9
10 we consider the case of the octyl butanoate and hexyl acetate, as an example, their concentrations
11
12 decreased by 30-43 % and 67-81 % respectively (Figure 1) although the sorption of these
13
14 compounds into PET and PVC only represented 0.15-0.57 % and 0.013-0.023 % of the initial
15
16 amount, respectively (table 1). These glassy materials showed their resistance to sorption by
17
18 aroma and thus the main parameter which had an impact on the aroma formulation was the
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20 hydrolytic process into the syrup.
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28 Few papers reported quantitative measurement of the sorption of aroma compounds into PET for
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30 real foods over a long time storage especially for thick materials such as bottles. For soft drinks
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32 flavoured with orange oil and stored in PET during 12 weeks, Nielsen (1994) showed that only 2.1
33
34 % and 1.4 % of the initial concentrations of myrcene and of limonene were respectively sorbed by
35
36 the polymer. Berlinet et al. (2005) compared the initial amounts of these two aroma compounds in
37
38 orange juice and the quantities adsorbed in PET after 5 months of storage. They found an
39
40 adsorption of between 0.2 and 0.3% of the initial levels present. These data were in line with the
41
42 findings of Van Willige *et al.* (2003) who stated that only three flavour compounds (limonene, β -
43
44 myrcene and decanal) were absorbed by PET. The percentage absorption of PET from juice
45
46 containing limonene, β -myrcene, and decanal reached only 0.1%, 0.1% and 2.8% respectively,
47
48 after one month of storage. When orange juice was stored in PET, the sorption of these two
49
50 compounds was lower. The constituents of the juice (pulp, pectin), missing in the soft drink, may
51
52 play a competitive role against the sorption into PET. The difference in the behaviour of the aroma
53
54 compounds whether they are present in a simple solution or a more complex food was noticed by
55
56 Leufven & Hermansson (1994). The sorption of aroma compounds into PET was lower when
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1 these compounds were into a tomato juice in comparison with a model solution. These authors
2 suggested that the aroma compounds probably remain in the tomato juice because they are
3 retained on the natural juice constituents instead of being sorbed in the polymer.
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9
10 In the case of strawberry syrup, sucrose may reduce the diffusion and convection of flavour
11 compounds in the polymer during the early stages of storage, as it increases the viscosity of the
12 medium (Van Willige et al. 2000a). Moreover, sucrose molecules may create local hydrophobic
13 environments in the medium which favours inclusion or interaction with hydrophobic molecules
14 such as aroma compounds. This may lower the sorption of aroma compounds because of a
15 competition effect between the two hydrophobic phases (polymer and food matrix) (Roberts et al.
16 1996, Nawar 1971).
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27 *Typical kinetics of sorption into PET and PVC for the more stable compounds*
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32 [Insert Figure 2 here]
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37 In both PVC and particularly PVC1, the sorption of most aroma compounds increased with the
38 square root of time, following a Fickian mode, as we were able to show for more stable
39 compounds such as 3-methylbutyl-3-methylbutanoate. Their sorption kinetics and their
40 interpretation as kinetic phase diagrams are shown in figure 2. Open symbols plot experimental
41 values as measured (duplicate experiments) whereas filled symbols plot non-deterministically
42 filtered data as described by Vitrac and Hayert (2006) (figures 2 A-D). The continuous lines depict
43 a cubic spline model, whose values and first derivatives fit the filtered ones. Versus the square
44 root of time, the kinetics show a subsequent delay related to a possible external mass transfer
45 resistance, a linear segment followed by a part with a decreasing slope. The presence of a linear
46 segment confirmed that the sorption was controlled by the diffusion. The following sharp change
47 in slope with a possible negative slope is related to a dynamic modification of sorption conditions.
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2 This modification was created by the degradation of the aroma compound in the liquid phase,
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4 which modified in return the attainable equilibrium state. The expected equilibrium concentration
5
6 in the packaging material was consequently decreasing.
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10 These trends were more discernible in the kinetic phase diagram (KPD), which plots the first
11
12 derivative of the concentration in the packaging material with time versus its primitive (i.e. the
13
14 concentration itself) (figures 2E and 2F). Filled and open symbols plot respectively the filtered
15
16 values and the 95% confidence interval as assessed by Monte Carlo sampling. A non-diffusive
17
18 behaviour such as the one related to a reaction in the liquid phase is associated to a change in
19
20 convexity in the diagram. Indeed, a convex shape of the KPD or a sharp decrease of the absorbed
21
22 mass flux when the absorbed amount increase cannot be related anymore to diffusion (see Vitrac
23
24 and Hayert, 2006 for further details). This rule could not however be applied to the earlier stage of
25
26 desorption kinetics since the mass flux was not estimated with enough accuracy. The KPD were
27
28 used to predict the equilibrium, which would be reached if the sorption conditions as assessed
29
30 during the linear part of the concentration versus the square root of times were prolonged, since in
31
32 the parabolic regime of diffusion, the KPD is linear. The concentrations at equilibrium were
33
34 linearly extrapolated from the convex part of the KPD to the horizontal line $dC_p|_t/dt = 0$. The
35
36 linear extrapolation inferred from the confidence envelope of the KPD appears as dashed lines.
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47 *Affinity of aroma compounds for PVC and PET*

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49 Apparent partition coefficients K_{app} derived from the extrapolation of concentrations expected at
50
51 equilibrium in absence of reaction are plotted versus logP (octanol-water partition coefficient)
52
53 (figures 3A and 3B) and versus the molecular weight (figures 3C and 3D) for PET (figures 3A and
54
55 3C) and for PVS (figures 3B and 3D).
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57
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59
60

[Insert Figures 3 A, B, C, D here]

1
2 It is highlighted that for the set of tested aroma compounds, log P and molecular weight are not
3
4 independent quantities. Log (K_{app}) appeared almost linearly correlated to log P values. Whatever
5
6 the polymer, sorption was selective and increased with the chain length of the linear butanoic acid
7
8 esters (butyl butanoate, hexyl butanoate and octyl butanoate), a behaviour quite similar to that
9
10 reported by Shimoda et al. (1987) with respect to polyethylene. However the correlation was
11
12 poorer with the molecular mass than with the log P, due to branching into the molecules.
13
14 Branching or unsaturation in the backbone of the molecule (for an equal carbon number) caused
15
16 sorption to decrease, as can be seen for 3-methylbutyl-3-methylbutanoate by comparison with
17
18 hexyl butanoate in PET (Figure 3C).
19
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22

23
24 Quite similar partition coefficients were obtained for both PET (PET1 and PET2) and both PVC
25
26 (PVC1 and PV2). Both PET samples sorbed the esters to a lesser extent, PVC1 displayed the
27
28 highest level of sorption, three to ten-fold more than both samples of PET mainly for octyl
29
30 butanoate and 3-methylbutyl- 3-methylbutanoate Few studies have dealt with the sorption of
31
32 aroma compounds into PVC. Koszinowski & Piringner (1987), and DeLassus (1994) showed that
33
34 diffusion coefficients in PVC were several orders of magnitude lower than in polyethylene and
35
36 polypropylene. In contrast, their solubilities (S) were about one order of magnitude lower in
37
38 polyolefins. Barrier behaviour evaluated by the permeability coefficient P ($P=D \times S$), resulting from
39
40 the kinetic effect (coefficient of diffusion, D), and the thermodynamic effect (S), only gave a small
41
42 advantage for PVC over polyolefins.
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48 With PET, the coefficients of diffusion and solubility of the volatile compounds are lower
49
50 (Pennarun *et al.* 2004) than those measured for polyolefins, so a low interaction can be observed
51
52 with this polymer. Figure 3C and 3D shows that both PET samples were more or less equivalent
53
54 and were less sorbent than PVC 2, except for cyclic esters such as ethyl cinnamate and ethyl
55
56 salicylate The sorption of these compounds in PET reached the level of sorption observed in
57
58 PVC2. These compounds exhibited a differential affinity for PET, probably because of the
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60

1
2 similarity of their cyclic structure. This affinity could affect the typical flavour of the syrup related
3
4 to the “wild strawberry note” which is associated with ethyl cinnamate, ethyl salicylate and ethyl
5
6 benzoate.
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8
9
10 In conclusion, the variation of K_{app} values in PET was mainly correlated to the polarity of tested
11
12 compounds as assessed by their log P values. By contrast, the variations in K_{app} values for PVC
13
14 were mainly related to their chain lengths
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16 17 18 19 20 *Kinetics of sorption into PET and PVC for the unstable compounds* 21

22 The sorption kinetics of some compounds, (ethyl butanoate, hexyl acetate), showed the same
23
24 typical trends in both types of PET (Figure 4).
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26
27
28 [Insert Figure 4 here]
29

30 Sorption increased to a peak at 90 days and then declined until 150 days, reaching a state of
31
32 equilibrium until 330 days.
33

34
35 Several hypotheses can be proposed to explain this behaviour.
36

37
38 1. Durning & Russel (1985) proposed a model of diffusion with induced crystallization to
39
40 describe the sorption kinetics of organic liquids which might trigger PET crystallization. Berlinet
41
42 et al., (2005) showed that the absorption of aroma compounds (even at low levels) may be
43
44 responsible for increasing the crystallinity of PET, over 6 months of storage. This “solvent
45
46 induced crystallization” which was described in PET in contact with ethyl acetate by Hao et al.
47
48 (1999), could lead to a structure modification of PET during 3 months of storage and then induce
49
50 the release of the aroma compounds from the packaging. However this behaviour would have been
51
52 observed for all the compounds specially those which are more sorbed such as octyl butanoate,
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54 but it is not the case and this behaviour was only noticed for the more reactive compounds.
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59
60 2. The sorption behaviour into PET of the more reactive aroma compounds could be explained in

1
2 the following manner: during the first stage between 0 to 90 days, these compounds were slowly
3
4 sorbed by PET, and hydrolysis into alcohols and acids started in the syrup. During the second
5
6 period (90<t<150 days), degradation was the leading parameter, which resulted in desorption of
7
8 the volatiles from PET, reaching an equilibrium between both phases (syrup and polymer). No
9
10 alcohol or acid from the corresponding esters was identified in the PET extract. After 150 days,
11
12 the equilibrium of aroma compounds between packaging and syrup was reached, taking account of
13
14 degradation into the syrup. Thus, after 90 days, PET could play a positive role as a reserve for
15
16 some of these unstable compounds.
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21 As previously described (Ducruet et al 2001), esterified aroma compounds were hydrolysed in
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23 glass bottles and this was not influenced by polymer contact. The hydrolysis rate of volatile
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25 compounds displayed a first order kinetic. Stronger reactivity was observed for methyl or ethyl
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27 esters of acetic acid when compared with longer linear chain esters and particularly branched
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29 esters. These compounds were rapidly hydrolysed in the syrup packaged in glass bottles during the
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31 first 150 days. Same reactivity was observed in both kinds of polymers. As oxygen transfer
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33 through amorphous PET and rigid PVC were of the same order of magnitude, respectively $4.4 \cdot 10^{-11}$
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35 $\text{cm}^2 \text{sec}^{-1} \text{Pa}^{-1}$ and $3.4 \cdot 10^{-11} \text{cm}^2 \text{sec}^{-1} \text{Pa}^{-1}$ (Branrup et al. 1989), the oxidation of these
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37 compounds cannot explain the observed difference between PET and PVC.
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43 In the case of orange juice in contact with PET, recent results had already shown that, degradation
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45 due to the acid catalysis of aroma compounds (α -pinene, limonene, linalool) is the principal
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47 parameter controlling exchanges with the glassy material (Berlinet et al. 2005) and therefore the
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49 aromatic evolution of orange juice was not controlled by the packaging material but by reactions
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51 within the matrix itself. Polyolefins (polyethylene and polypropylene (in contact with a model
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53 medium containing highly reactive compounds such as limonene or pinene under acidic conditions
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55 may play a positive role in stabilising aroma compounds (Lebossé et al. 1997, Feigenbaum et al
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57 1998, Reynier et al. 2004). Because the degradation process in the model system is of the same
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2 order of magnitude as diffusion into the polymers, the two phenomena are in competition. In the
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4 case of PET, transfer is very slow, and many studies, which considered contact times of
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6 insufficient duration, were not able to demonstrate this effect. In the case of contact between real
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8 food products during their shelf life (up to one year) and unstable aroma compounds such as short
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10 chain esters, the degradation of aroma compounds is more rapid than diffusion into PET. Release
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12 is less rapid than with polyolefins, and some stabilization effects of PET may be observed after
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14 only 150 days of storage, in line with the type trend of sorption kinetics.
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19 In the context of food packaging interactions, the thermodynamic equilibrium of each aroma
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21 compound is expressed as the partition coefficient. Because the experimental determination of this
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23 partition coefficient may be prohibitive (Tehrany & Desobry 2004), modelling has been proposed
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25 as an alternative (Dekker et al. 2003, Tehrany & Desobry 2005). Models are available for
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27 polyolefins but extensions to glassy polymers such as PET or PVC are still pending. The main
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29 limitations consist in the availability of reference data, which are not subjected to bias due to a
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31 competition between slow sorption kinetics (longer than 150 days) and a fast degradation process.
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33 The production of apparent partition coefficient with minimum bias as assessed in this study using
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35 a kinetic phase diagram could be an alternative to the lack of published data. From the
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37 technological point of view, the loss of aroma for a particular food in contact with a glassy
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39 material must be envisaged as a consequence of both the sorption and degradation process. In the
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41 present case, the degradation of the aroma formulation of the syrup is mainly due to hydrolysis of
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43 esters.
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50 **Conclusion**

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52 This study shows that the interactions between aroma compounds and packaging may result in a
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54 dynamic and time-dependent change in food quality during shelf life. Two combined mechanisms
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56 contribute to a loss of aroma during long-term storage: the degradation process in the food product
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58 itself and the sorption process in the packaging material. Both effects were particularly noticeable
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2 for both PET samples and for unstable aroma compounds. It is emphasized that both processes are
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4 antagonist. Although the sorption rate is low in glassy polymers, the sorption process tends to
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6 prevent the aroma from the degradation process. By contrast, the degradation modifies the
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8 expected thermodynamic equilibrium of sorption of aroma and tends to limit the sorption of
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10 aroma. Both phenomena were simultaneously studied in this work and a general methodology was
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12 proposed to extract apparent partition coefficients, when the thermodynamic equilibrium cannot
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14 be reached due to a coupling with reactions. This methodology was used to derive the partition
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16 coefficients of 13 aroma compounds found in real fruit juices and two glassy polymers : PET and
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18 PVC. Besides, it was highlighted that the stability of the aroma profile in connection with a
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20 packaging material cannot be addressed with short contact times experiments. Additional studies
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22 and data appear desirable to help food industry to predict the behaviour of their aroma formulation
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24 during long term storage taking into account both the effects of the food matrix and the packaging.
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Table 1 Extraneous compounds tentatively identified in the syrup or in the packaging material extracts during storage

Packaging	Compounds initially present in the polymer	Migration into the syrup	Origin of compounds *
PVC	2-ethylhexanol	Yes after 250 days	thermal degradation of plasticizer
	2-ethylhexanol acetate	No	
	benzaldehyde	Yes	ABS or MBS impact modifier derived from an organotin stabilizer
	styrene	No	
	isooctyl thioglycolate		
acetophenone	Yes		
PET	2-ethylhexanol	Yes before 225 days	thermal degradation of plasticizer
	benzaldehyde	Yes	possible degradation products of the polyethylene waxes used as lubricants in PET
	naphthalene, linear aldehydes	No.	
		No	
acetophenone	Yes		

*Gachter & Muller, 1990

Table 2 : Amount of aroma compounds sorbed into PET and PVC after 90 days** and 330 days of storage at 20°C. Means of two determinations

Volatile Compound	Code	Co* mg l ⁻¹	PET1 µg / 6dm ²	% sorption	PET2 µg /6dm ²	% sorption	PVC1 330 days µg /6dm ²	% sorption	PVC2 330 days µg /6dm ²	% sorption
Butyl butanoate	BB	24.6	3.7	0.015 c	3.8	0.015 c	7.8	0.032 a	4.9	0.020 b
Hexyl butanoate	BH	9.6	2.3	0.024 c	2.6	0.027 c	11	0.115 a	5.1	0.053 b
Octyl butanoate	BO	8.4	13.1	0.156 c	12.8	0.150 c	47.8	0.569 a	22.1	0.263 b
Ethyl hexanoate	HE	167.5	24.5**	0.015 c	27.8**	0.017 b	54.3	0.032 a	29.1	0.017 b
3-Methylbutyl 3-methylbutanoate	3MB3MB	134.5	12	0.009 c	16.8	0.012 c	146.9	0.109 a	62.7	0.047 b
Ethyl 2-methylbutanoate	2MBE	257.5	8.8	0.003 c	8.1	0.003 c	23.2	0.009 a	16.4	0.006 b
Ethyl 3-methylbutanoate	3MBE	166.8	7.3	0.004 a	7.3	0.004 a	12.4	0.007 a	8.6	0.005 a
Hexyl acetate	AcH	152.5	23**	0.015 b	25.8**	0.017 b	34.6	0.023 a	19.4	0.013 c
3(Z)-Hexenyl acetate	Ac3H	142.2	11.4**	0.008 c	13**	0.009 b	21.3	0.015 a	12.7	0.009 b
Octyl acetate	AcO	10.4	9.1**	0.087 a	8.5**	0.082 a	9.2	0.088 a	4.3	0.041 a
Methyl cinnamate	CIME2	530.5	149.5	0.028 b	164.9	0.031 b	283.5	0.053 a	149.7	0.028 b
Ethyl salicylate	2OHBzE	120.4	45.5	0.038 b	40.8	0.034 b	85.7	0.071 a	42.9	0.036 b
Ethyl benzoate	BzE	47.5	7.2	0.015 bc	6.3	0.013 c	43	0.091 a	31.9	0.067 ab
γ-decalactone	GDL	339.2	21	0.006 c	13.6	0.005 c	132.3	0.039 a	43.2	0.013 b

* : concentration of the aroma compounds into glass bottle at t0 day

Different letters in the % sorption indicate significant differences at $p < 0.05$ (Duncan) for each aroma compound between packaging materials.

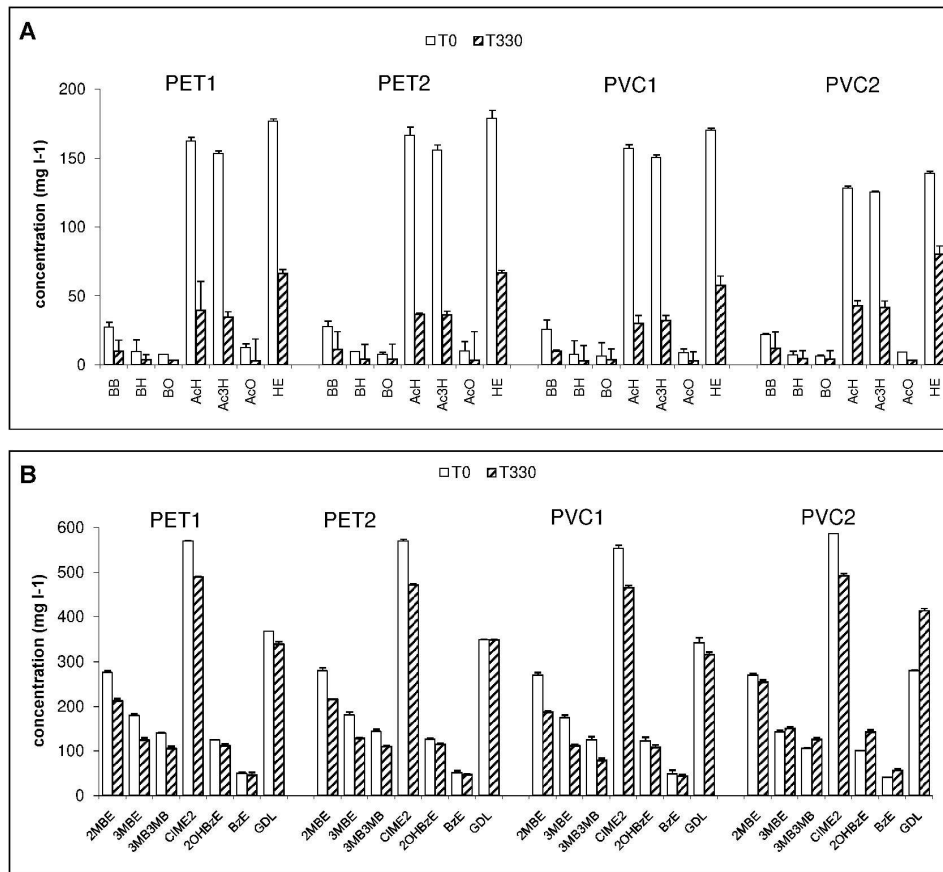


Figure 1 Concentration of the aroma compounds into the strawberry syrup at t 0 day and t330 days of storage into the 4 packaging materials (n=3)
149x141mm (600 x 600 DPI)

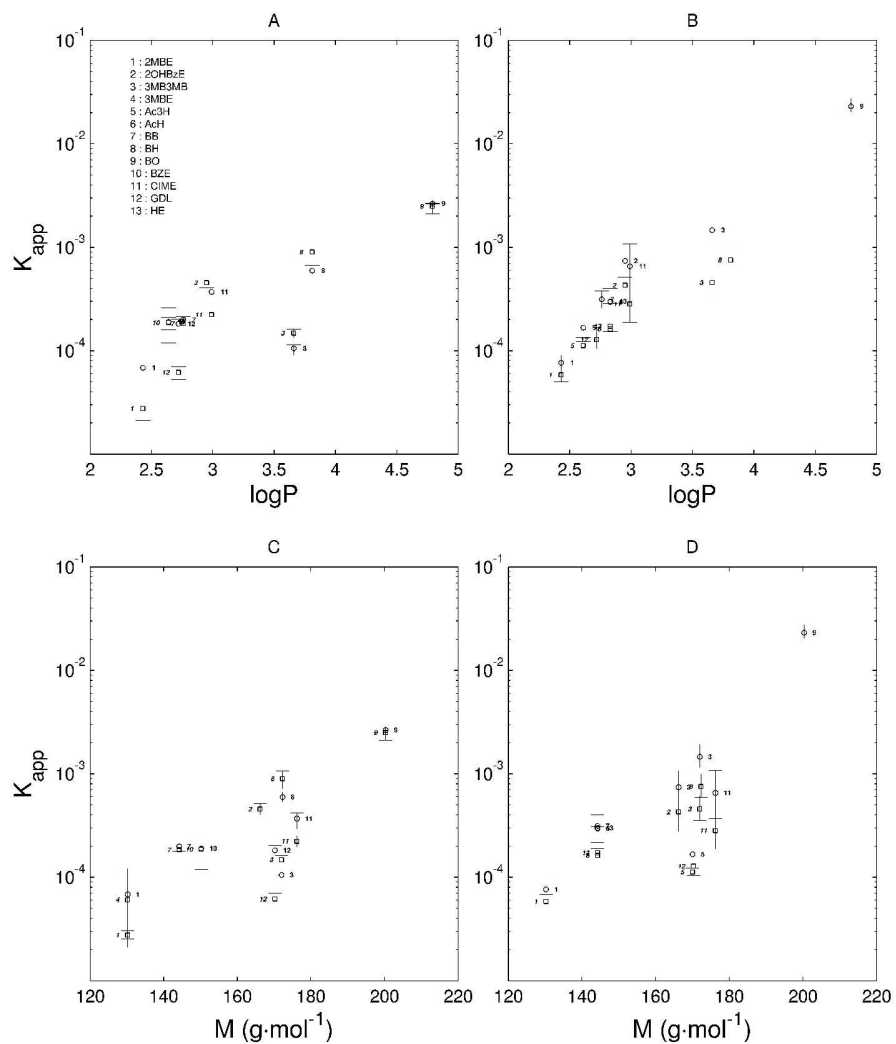


Figure 2 A-D) Sorption kinetics of 3-methyl butyl 3-methyl butanoate in PET and PVC on a linear scale (A,B) and versus the square root of time (C,D). E-F) Kinetic phase diagram corresponding to figures A and B respectively. 1695x1905mm (72 x 72 DPI)

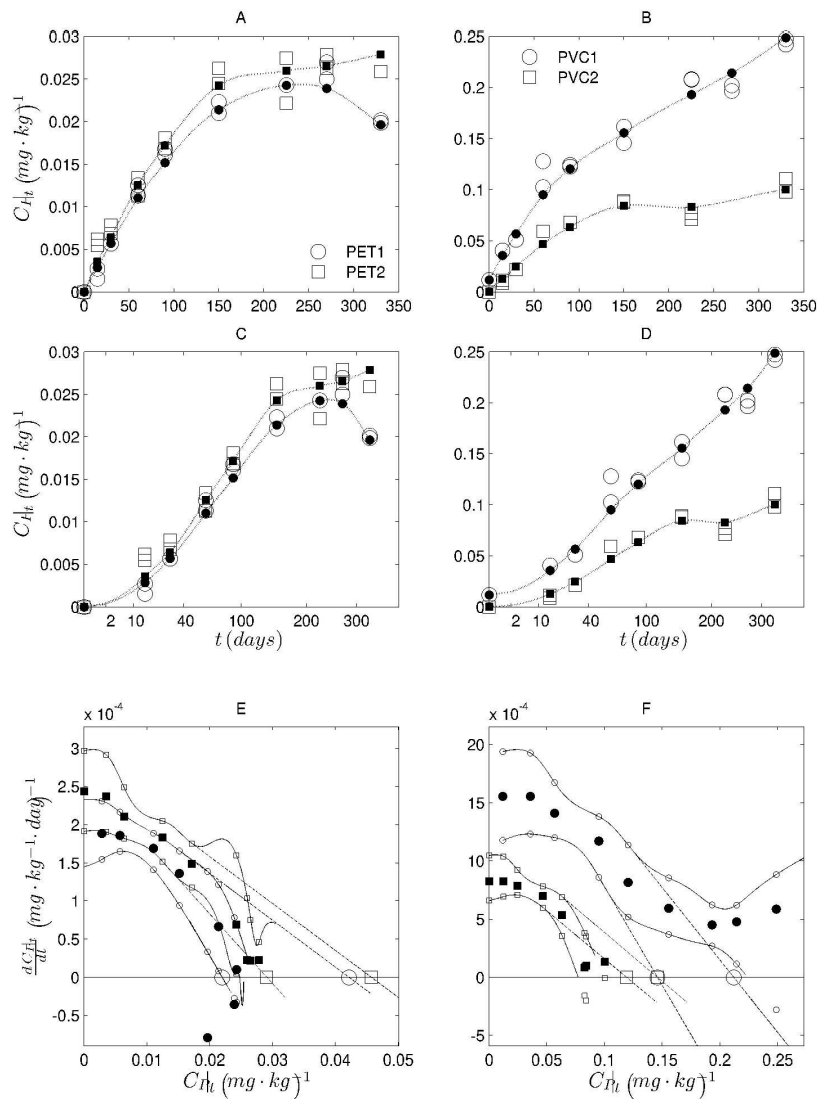


Figure 3 Apparent partition coefficients K_{app} , in PET (A,C) and PVC (B,D) versus $\log P$ (octanol-water partition coefficient) (A,B) and molecular weight (C,D) of aroma compounds. Normal and italic fonts are used for PET1 and PVC1, and for PET2 and PVC2 respectively. The codes of aroma compounds are defined in Table 2.

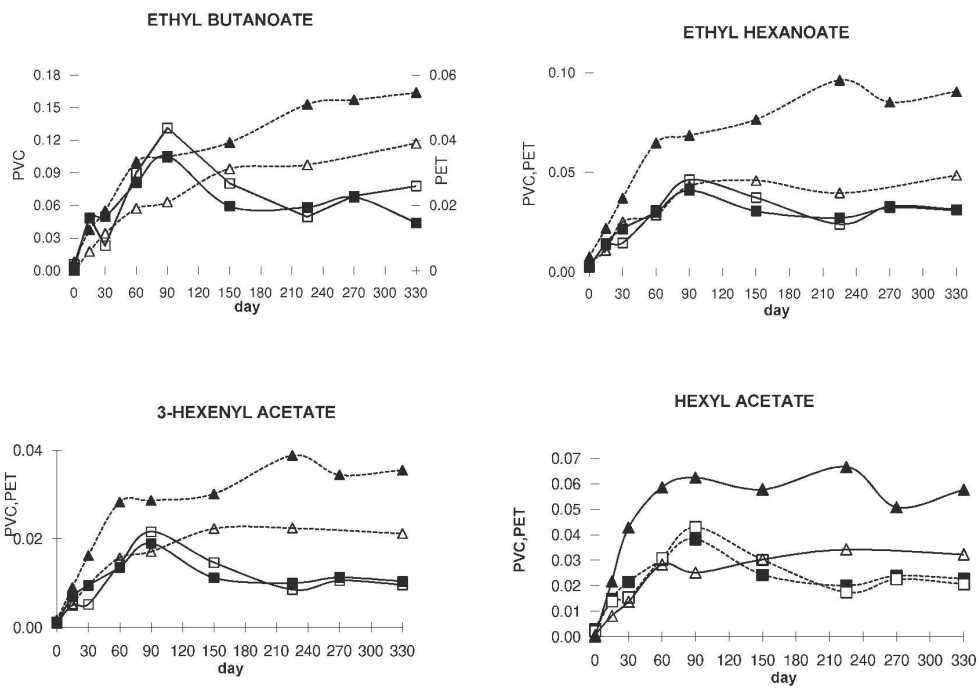


Figure 4 Sorption of aroma compounds ($\mu\text{g}/\text{cm}^2$) into PET1(■), PET2 (□) PVC1(▲) PVC2 (Δ) versus time
145x102mm (600 x 600 DPI)