**Hydrolysis optimization of mannan, curdlan and cell walls from *Endomyces fibuliger*
grown in mussel processing wastewaters**

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29 **ABSTRACT**

30 The aim of this report was to optimize the hydrolysis of the cell walls (CWs) from the yeast
31 *Endomyces fibuliger* grown in mussel processing wastewaters (MPW) to establish a more
32 accurate protocol for analyzing the composition of the monosaccharides in these CWs.
33 Therefore, a kinetic study of CW hydrolysis and polysaccharide standards (mannan and
34 curdlan) was performed to determine the effect of different temperatures and trifluoroacetic
35 acid (TFA) concentrations on this process. In all cases, the experimental data were fit
36 satisfactorily to Saeman's equation with an Arrhenius relation between rate constants and the
37 temperature effect. Optimal conditions for curdlan and mannan hydrolysis were achieved with
38 70% TFA at 100°C for 2.3 h and 50% TFA at 100°C for 2.6 h, respectively. The best
39 operating options for CW hydrolysis were 100°C/70% TFA for 4.58 h, 100°C/50% TFA for
40 4.08 h and 100°C/70% TFA for 3.27 h for the maximum production of glucose, mannose and
41 reducing sugars, respectively.

42

43 **Keywords:** yeast cell walls; mussel processing wastes; *Endomyces fibuliger*; trifluoroacetic
44 acid hydrolysis; mathematical modeling; curdlan and mannan

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46 **1. Introduction**

47 Cell walls (CWs) in yeast and fungi are dynamic, stratified structures that are in a constant
48 state of hydrolysis, and its biosynthesis is controlled by strict regulation. Changes in culture
49 conditions, different cellular stages and dimorphic transitions can influence the CW
50 composition [1]. The main components of CWs are 1) structural elements such as β -1,3 and
51 β -1,6 glucans (a polymer of D-glucose) and chitin (a polymer of N-acetylglucosamine), 2)
52 matrix components, such as α -glucans and glyco- or mannoproteins (polymers of mannose
53 linked to proteins), and 3) lipids and proteins [2-4].

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The study of active oligosaccharides and β -glucans has generated much interest over the last two decades because of the substantial empirical evidence that they have healthy effects, mainly as antitumor and immunomodulation agents, on different biological entities that range from invertebrates to mammals [5-7]. Although the mechanisms that underlie these properties are still not completely understood, it seems clear that β -glucans activate the leukocyte mediators of the immune response [8], acting as an antigenic stimulus of dendrite cells and/or developing the phagocytic activity of the leukocytes [9]. Their effects mainly depend on the molecular weight, degree of polymerization, proportion of 1-3, 1-4 and 1-6 bonds, presence of mannose and the formation of chitin or protein complexes [10]. For the latter case, there is still no clear consensus, but it is generally agreed that chitin and mannoprotein complexes develop the appropriate properties and also that a high molecular weight (low solubility) produces adverse effects, at least when intraperitoneal administration is used. Additionally, these factors are modified by the type of producer microorganism, the taxonomic group, the life-cycle phase and the medium growth conditions [1]. Therefore, it is important to have tools to identify and quantify the carbohydrates (i.e., polysaccharides of mannose, glucose and N-acetyl-D-glucosamine) in microbial CW.

The most common methods for studying the composition of these carbohydrates are based on chromatography techniques (e.g., gas chromatography (GC) or high-performance liquid chromatography (HPLC)), which allow the CW monosaccharides to be distinguished. In these techniques an initial hydrolysis step, based on a chemical reaction in acid media at high temperatures, is necessary [11,12]. This step is a destructive process and its effects depend on the structure, the CW composition and the conditions and type of chemical used [13,14]. Optimum hydrolysis conditions should maximize polysaccharide breakdown and minimize the destruction of the corresponding released monosaccharides. Therefore, we need to consider that monosaccharide stability with strong acids employed for hydrolysis varies depending on their chemical composition and the presence and proportion of amino groups.

Nevertheless, in the literature, there is a tendency to accept the optimal hydrolysis conditions used in one system for another. For example, using the conditions found in potato peel for the structural analysis of yeast CW carbohydrates in [13]. Using non-optimal procedures for these substrates leads to the partial destruction of some monosaccharides or the incomplete hydrolysis of the polysaccharides, which alters the final compositional results. It is therefore essential to optimize the hydrolysis conditions and study the joint effects of variables such as temperature and the oxidant compound for each substrate used to study its carbohydrate composition. To our knowledge, the optimal conditions for breaking down curdlan, mannan and yeast CW have not yet been reported.

The CWs of different yeasts have been used as substrates for β -glucans production [2,6]. The most common is *Saccharomyces cerevisiae*; however, amylolytic yeasts, which are able to grow in amylaceous and residual effluents, have not yet been studied as polysaccharides and β -glucans producers. The benefit of this proposal is especially interesting in our region because more than 25% of the world mussel production occurs on the Galician coast (NW, Spain). In the thermal process for canning mussels, approximately 1.6 million m³ of mussel processing wastewaters (MPW) are generated per year and dumped in the sea without previous depuration [16]. This residual effluent has been successfully used as a carbon substrate for several bioproductions including gibberellins [17], amylases [18], glucose oxidase [19], citric acid [20], pediocin [21], hyaluronic acid [22] and single cell proteins from *Endomyces fibuliger* [23]. This last amylolytic yeast is an excellent candidate for glucans production because its CW composition (i.e., 56% CW per biomass unit with 64% of total sugars per CW unit) obtained in our study is higher than that of other yeasts (i.e., 35% per biomass unit with 85% total sugars per CW unit) commonly used for this purpose and as reported in [24]. To our knowledge, *E. fibuliger* has not been previously studied for this purpose.

The main objective of this study was to investigate the effect of temperature and trifluoroacetic acid (TFA) concentration on the hydrolysis kinetics of commercial curdlan and mannan and CWs obtained from *Endomyces fibuliger* that was previously grown in MPW. These findings are essential for accurately quantifying CW monosaccharides. The experimental profiles were fit satisfactorily to the Saeman and Arrhenius equations for describing and defining the optimal conditions for the maximal breakdown of polysaccharides to monosaccharides and minimizing the destruction of these monosaccharides due to the thermal and chemical reactions.

2. Materials and Methods

2.1. Microbiological methods, media preparation and culture conditions

Endomyces fibuliger (CBS 2521) was used as the CW source in this study. MPW were kindly supplied by Marcelino S.A. (Galicia, Spain), and their chemical composition was as follows: 7 g/L glycogen, 0.10 g/L reducing sugars, 3.5 g/L proteins and 1.6 g/L total nitrogen. Sediments were not observed in these effluents, and the initial pH was 7.2.

To precipitate the proteins, the MPW was first treated with HCl (50% v/v) until a pH of 4.5 was reached, and the supernatant was subsequently concentrated with an ultrafiltration membrane [25,26]. Ultrafiltration was performed using 0.56 m² spiral polyethersulfone membranes (*Millipore Prepscale*) with a 100 kDa cutoff using an assembly with total recirculation at 30°C.

This concentrate, which contained 25 g/L of glycogen and 2 g/L of protein-Lowry, was supplemented with 400 mg/L of phosphorus (KH₂PO₄) and 1200 mg/L of nitrogen (NaNO₃:NH₄Cl in a 0.8:0.2 (w/w) ratio) to formulate the MPW based medium for yeast fermentation [23,27]. The kinetic analysis cultures were performed in triplicate in 300 mL Erlenmeyer flasks with 50 mL of medium at 30°C, 200 rpm and an initial pH of 5.0. For the

biomass and CW studies, 3 L flasks with 500 mL of culture medium were collected at the end of the growth phase (55 h). Inocula (2%, v/v), a cellular suspension of 48 h-old *E. fibuliger* fermentations in MPW based medium, were prepared to a final concentration of 2.5×10^6 cells/mL in the experimental units.

2.2. Fermentation sampling and analytical determinations

At pre-established times, each Erlenmeyer flask was removed from the shaker incubator and the post-incubation medium was centrifuged at 4500 g for 20 min. The sediment was washed using distilled water and centrifuged again to eliminate all medium components. The washed precipitate was used to determine the dry weight after it had been dried in an oven at 107°C. The following compounds were analyzed in the supernatant: total amylolytic activity [18], total sugars [28,29], proteins [30], total nitrogen [31] and reducing sugars [32].

The 3 L flasks were collected at 55 h and the fermented media were centrifuged at 10,000 g for 20 min. The sediment was separated by filtration using 0.45 µm glass microfiber filters (Whatman), washed with abundant distilled water, lyophilized and crushed with a pestle to determine its chemical composition. The obtained material was stored at -20°C after desiccation with KOH for CW treatment. The total sugars, reducing sugars, total nitrogen, ash (by calcination at 550°C until constant weight) and total lipids using Soxhlet extraction were determined in triplicate [33]. The CWs were obtained according to the methodology proposed by Kasahara [34]. However, the polysaccharides standards used as controls for the hydrolysis process, curdlan and mannan, were purchased from Sigma Aldrich (St. Louis, MO, USA).

2.3. Hydrolysis conditions and hydrolysates analysis

The substrates, CWs and polysaccharide standards were placed in 30 mL tubes with sealed Teflon caps and mixed with several diluted TFA concentrations that ranged from 10% to 70% (v/v) at different temperatures between 35 and 120°C with an initial solid/liquid ratio of 0.5

mg/mL. Samples were removed from the reaction media at the pre-established times. After hydrolysis and before the chemical analysis, the TFA was completely evaporated, and the hydrolysates were completely dried in a heating oven with a gas extractor for between 30 and 50 h at 30°C.

The amount of monosaccharides released by hydrolysis was determined using the following:

- 1) The 3,5-dinitrosalicylic reaction method, with spectrophotometric measurements taken at 540 nm for the hydrolysis of the standards and the CWs [32]
- 2) Gas chromatography-mass spectrometry (GC-MS) measuring alditol derivatives (only in the case of CW) to differentiate between the release of mannose and glucose [35].

This second method is based on forming alditol acetate compounds from monosaccharides present in the CWs and then quantifying GC with a flame ionization detector. Thus, 3 mL of pyridine anhydride (C_6H_5N) and 1 mL of acetic anhydride ($CH_3CO_2CH_3$) were added to 1 mg of the sample in solution (generating a reductive ambient), and the oxygen was removed by nitrogen flow. The reaction medium was agitated for 24 h at room temperature. Next, 3 mL of concentrated HCl was added drop by drop in an ice bath. The resulting phase was extracted with ethyl acetate three times until a volume of 5 mL was reached. The calibration curves of the monosaccharides (i.e., glucose and mannose) were prepared as pentacetates [36] using ribose obtained by the same procedure as the internal standard.

We used an HP 5850 GC with a selective mass detector HP 5971 (series J) in scan mode in the range 50-400 m/z, and a Supelco SP-2330 column (30 m × 0.25 mm). The temperatures were 200°C (injector), 280°C (detector), and a column programming range from 150°C to 250°C, which was maintained for 10 min, with a gradient of 7°C/min. Helium was used as the carrier gas and its flow was kept constant at 8 psi.

2.4. Mathematical models

The mathematical equation used to describe the experimental hydrolysis profiles was based on a pseudohomogeneous kinetic model in liquid phase with a first order reaction [37,38], according to the modifications reported in [39] and the variables studied in our report:

where k_h is the specific rate of monosaccharide production (h^{-1}) and k_d is the specific decomposition rate (h^{-1}). Integrating the corresponding differential equations led to the following explicit equation:

$$M = M_0 e^{-k_d t} + P_0 \frac{k_h}{k_d - k_h} (e^{-k_h t} - e^{-k_d t}) \quad (1)$$

where t is the time course of the reaction (min), M is the relative concentration of monosaccharide, glucose or mannose (%), M_0 is the initial concentration of the monosaccharide (%) and P_0 is the initial concentration of the polysaccharide (%). This equation was used to model the hydrolysis of the polysaccharide standards and the CWs from *E. fibuliger* in all the experimental conditions tested. The sugar values from the polysaccharide standards were calculated in relation to the maximum sugar concentration measured (in %). For *E. fibuliger* bioproduction, the sugar values were calculated in terms of the CW dry weight for each instance.

In all cases, M_0 was zero, and thus equation (1) is simplified to:

$$M = P_0 \frac{k_h}{k_d - k_h} (e^{-k_h t} - e^{-k_d t}) \quad (2)$$

216

217 The time necessary to obtain maximal monosaccharide production (t_m , in hours) was
 218 calculated by deriving equation (2) with respect to time and equaling to zero:

219

$$\left. \frac{dM}{dt} \right|_{t=t_m} = P_0 \frac{k_h}{k_d - k_h} (-k_h e^{-k_h t} + k_d e^{-k_d t}) = 0 \Rightarrow t_m = \frac{\ln\left(\frac{k_d}{k_h}\right)}{k_d - k_h} \quad (3)$$

221

222 If t_m is substituted in equation (2), we can obtain the maximum monosaccharide production
 223 (M_m , in %):

224

$$M_m = P_0 \frac{k_h}{k_d - k_h} (e^{-k_h t_m} - e^{-k_d t_m}) \quad (4)$$

226

227 The temperature dependence of the kinetic parameters from equation (1) is described by the
 228 Arrhenius equation:

229

$$k_i = k_{i0} \exp\left(-\frac{Ea_i}{RT}\right) \quad (5)$$

231

232 where k_i is the kinetic parameter (for $i = h$ or d), k_{i0} is the pre-exponential factor (h^{-1}), Ea_i is
 233 the activation energy (kJ/mol), R is the universal gas constant ($8.314 \times 10^{-3} \text{ kJ mol}^{-1} \text{ K}^{-1}$) and
 234 T is the absolute temperature (K).

235

236 By inserting equation (5) into equation (2), a bivariate model is used to calculate the
 237 activation energies, the pre-exponential factors and the relative concentration of
 238 monosaccharide (M) at any given time and temperature:

$$M = P_0 \frac{k_{h0} \exp\left(-\frac{Ea_h}{RT}\right)}{k_{d0} \exp\left(-\frac{Ea_d}{RT}\right) - k_{h0} \exp\left(-\frac{Ea_h}{RT}\right)} \left[\exp\left(-k_{h0} \exp\left(-\frac{Ea_h}{RT}\right)t\right) - \exp\left(-k_{d0} \exp\left(-\frac{Ea_d}{RT}\right)t\right) \right] \quad (6)$$

240

241 2.5. Numerical and statistical methods

242 The strategy for modeling the experimental data is summarized in the following steps:

243

244 1) Experimental data from each individual set of TFA and temperature were fit to equation
 245 (2). Because P_0 has to be equal for each substrate independently of the TFA and temperature
 246 conditions, a numerical estimate of this parameter was made only and jointly for all
 247 experimental series. The individual set of parameters (k_h and k_d) obtained for each TFA and
 248 temperature condition was used to calculate the monosaccharide production maximum and the
 249 time it occurs (equations 4 and 3, respectively).

250

251 2) The experimental data from each TFA concentration set at all tested temperatures were fit
 252 jointly to equation (6) to obtain the numerical values of Ea_h , k_{h0} , Ea_d and k_{d0} . The used P_0
 253 value was that obtained in previous adjustments.

254

255 The fitting procedures and parametric estimates from the experimental results were performed
 256 by minimizing the sum of quadratic differences between the observed and model-predicted
 257 values using the nonlinear least-squares (quasi-Newton) method provided by the ‘Solver’
 258 macro from Microsoft Excel spreadsheet. The confidence intervals of the best-fit values for
 259 the parametric estimates ($\alpha=0.05$), consistency of the mathematical models (Fisher’s F test;
 260 $p<0.05$) and covariance and correlation matrices were calculated using the ‘SolverAid’ macro,
 261 which is freely available from de Levie’s Excellaneous website:
 262 <http://www.bowdoin.edu/~rdelevie/excellaneous/>.

263

3. Results and Discussion

3.1. *E. fibuliger* culture and biomass composition

Figure 1 shows the experimental results of *E. fibuliger* fermentation in the MPW-based medium. Maximum growth was obtained at 55 h with 12 g/L of biomass (as dry weight). At that time, the glycogen from in residual culture media was completely consumed, and the pH increased from 5.3 to 6.8. At 30 h, the yeast amylases were deactivated due to a drop in the reducing sugar concentration. In addition, the yeast consumed 1.3 g/L of protein-Lowry. Therefore, 55 h of culture was selected to obtain the maximum biomass. This biomass was then used for compositional analyses and CW hydrolysis.

The chemical composition of the biomass after 55 h of cultivation was as follows (in %, dry basis (db)): 59.2 ± 2.5 total sugars, 7.63 ± 0.77 reducing sugars, 14.9 ± 0.7 total lipids, 1.6 ± 0.2 ash and 6.4 ± 0.4 proteins (as N $\times$ 6.25). The values for the CW analysis were the following (in %, db): 69.3 ± 1.2 total sugars, 25.9 ± 1.4 proteins, 0.7 ± 0.1 total lipids and 3.3 ± 0.2 ash. Reducing sugars were undetected in the CWs. A 54.7% CW yield was obtained from the *E. fibuliger* biomass. This value was higher than that reported in [24]. In this article, the yeasts *Kluyveromyces marxianus* and *Debaryomyces hansenii* had the highest proportion of CW in their biomass (32.5% and 32%, respectively). However, in all the microorganisms studied by these authors, the total sugars percentage was superior to the *E. fibuliger* results (more than 84% compared with 65% for *E. fibuliger*). These differences could be due to the different methods for obtaining CWs. Using autoclaving, a French pressure cell press, cell lysis by glass beads or a homogenizer led to different efficiency results and yields in the biomass breakdown of microorganisms [24].

3.2. Hydrolysis kinetics of curdlan and mannan

The hydrolysis results at different temperatures and several TFA concentrations using curdlan and mannan as substrates are shown in Figure 2. The profiles adjusted to the experimental

data according to model (2) are also shown. Table 1 lists the values of the kinetic parameters and the statistical analyses of the numerical fittings. In general, the proposed models were statistically robust (Fisher's F-test p -values < 0.05), the parametric estimations were significant (Student's t-test $\alpha = 0.05$), the residuals were randomly distributed and autocorrelations were not observed by the Durbin-Watson test (data not shown). The linear determination coefficients (R^2) between the predicted and observed values were always greater than 0.95.

Table 2 summarizes the parameters defined by equations (3) and (4), which are important for determining the optimal conditions for maximal sugar release from commercial polysaccharides or monosaccharide extraction from CW for chemical composition analysis. Using curdlan as the hydrolysis substrate, four different experimental conditions, 80°C/70% TFA, 80°C/50% TFA, 100°C/70% TFA and 100°C/50% TFA, were used to obtain the maximum concentrations of released glucose (98.42%, 94.76%, 94.10% and 94.23%, respectively), though with markedly different maximum hydrolysis times (10.7 h, 14.2 h, 2.3 h and 3.4 h, respectively). To shorten the processing time, we performed the hydrolysis at 100°C and 70% TFA, although almost 4% less glucose is produced under these conditions. Mannan hydrolysis demonstrated two optimal maxima in the 100°C/50% TFA and 80°C/50% TFA pairs with processing times of 2.6 h and 13.1 h, respectively, and the former option is the most useful. The decreased t_m was correlated by increasing the TFA concentration but without a clear tendency for significant modeling.

These results are in agreement with those previously reported by Freimund et al. [14]. These authors obtained the best conditions for glucan hydrolysis in the range of 92.5 to 100°C for 1.5 to 3 h and 72.5% TFA. For breaking down mannan, these intervals were 90 to 100°C for 1.75 to 4 h and with a higher TFA concentration (60%) than we propose. Although [14] presents a similar study of the combined effect of the dependent variables (i.e., temperature,

time and acid concentration), these data were obtained by individual observation and by combining the variables at their apparent maximum point without optimizing by mathematical modeling. This type of procedure is common in the literature, but it leads to a faulty understanding of the combined variable effect in terms of the response to maximization [40,41].

The numerical parameters for the bivariate equation (6) are shown in Table 3. In both hydrolysis and decomposition, the activation energies for curdlan were higher than that for mannan, which is in agreement with the shorter t_m obtained for this polysaccharide (Table 2). In addition, the increase in the TFA concentration led to a slight decrease in Ea_h and Ea_d . Indeed, with higher acidity in the reaction medium, less energy is needed to break the glycosidic bonds that link the monosaccharides in the polysaccharide skeleton.

3.3. *E. fibuliger* CW hydrolysis kinetics

Figure 3 shows the experimental data and modeling trends of the hydrolysis kinetics of the CWs produced by *E. fibuliger*. The statistical analyses of the relevant kinetic parameters are summarized in Table 4. The P_0 values were 59.4%, 41% and 24.5% for the ratios of RS, glucose and mannose per CW db, respectively. These percentages were similar to those reported earlier in section 3.1 (i.e., 69.3% total sugars). All of the parameters were again statistically significant (Student's t-test, $\alpha=0.05$), and equation (2) was consistent (Fisher's F-test, $\alpha=0.05$ and R^2).

Equations (3) and (4) permitted us to predict the hydrolysis time (t_m) necessary for obtaining the maximum percentages of glucose, mannose and reducing sugars released from the studied yeast CWs (Table 5). There were two options for recovering the maximum glucose: 100°C/70% TFA and 120°C/70% TFA, with reaction times of 4.58 h and 0.88 h, respectively. These conditions were different for mannose: 100°C/50% TFA for 4.08 h, 120°C/35% TFA

for 1 h and 120°C/50% TFA for 0.83 h. Therefore, the suggestion by Freimund et al. (2005) that for determining the monosaccharides present in the CWs, different treatment conditions should be applied seems reasonable. For compositional analyses of glucose and mannose in *E. fibuliger* CWs, we suggest the following operating conditions: 100°C/70% TFA for 4.58 h and 100°C/50% TFA for 4.08 h, respectively. The reducing sugars results show that maximum RS production is obtained at 100°C with 70% TFA for 3.27 h. However, if shorter processing times are necessary, the hydrolysis reaction could be performed at 120°C with 70% TFA for 1 h to generate 39% RS. To the best of our knowledge, this study is the first approach to mathematically model the thermal and acid hydrolysis of yeast CWs to optimize monosaccharide production experimental conditions. As in the previous study with standards, the time values decreased with an increase in the reaction media acidity. Moreover, the optimal temperature and TFA concentration for obtaining the maximum glucose, mannose and RS concentration from CW was similar: 100°C/70% TFA, 100°C/50% TFA and 100°C/70% TFA, respectively. These results confirm the validity of the proposed model and the developed methodology.

The regression coefficients that relate the temperature to the hydrolysis rates are summarized in Table 6. The correlations between the expected and experimental data were satisfactory ($R^2 > 0.95$) for all cases. The pre-exponential factor for both hydrolysis and decomposition did not show any differences in relation to the increase in the acid percentage used in the reaction. Nevertheless, the increase in acidic conditions generated a small, progressive decrease in the hydrolysis and decomposition activation energies. The Ea_d values for mannose were significantly lower than those obtained in the other cases. This shows that mannose is more sensitive to decomposition due to high temperatures and strong acid conditions.

3.4. Modeling the combined effect of temperature and TFA concentration on the hydrolytic process

The joint TFA concentration and temperature effects on the hydrolysis process were also evaluated. As the changes in Ea_h and Ea_d , in relation to the TFA concentrations, were not significant and the k_{h0} and k_{d0} values were quite similar (Tables 3 and 6), we modified the Arrhenius equation to include acid levels in the modeling process [42,43]:

$$k_s = C^{n_s} \exp(a_s) \exp\left(-\frac{Ea_{MS}}{RT}\right) \quad (7)$$

where a_s and n_s are the regression parameters (for $s = h$ or d), C is the TFA concentration (in % v/v), k_s is the kinetic parameter of the combined effect of temperature and acid (for $s = h$ or d), Ea_{MS} is the average of the activation energies (for $s = h$ or d , in kJ/mol), R is the universal gas constant (8.314×10^{-3} kJ mol⁻¹ K⁻¹) and T is the absolute temperature (K).

Thus, a global model is defined by inserting equation (7) into equation (2):

$$M=P_0 \frac{C^{n_h} e^{a_h} e^{\left(-\frac{Ea_{Mh}}{RT}\right)}}{C^{n_d} e^{a_d} e^{\left(-\frac{Ea_{Md}}{RT}\right)} - C^{n_h} e^{a_h} e^{\left(-\frac{Ea_{Mh}}{RT}\right)}} \left[\exp\left(-C^{n_h} e^{a_h} \exp\left(-\frac{Ea_{Mh}}{RT}\right)t\right) - \exp\left(-C^{n_d} e^{a_d} \exp\left(-\frac{Ea_{Md}}{RT}\right)t\right) \right] \quad (8)$$

All experimental data from each substrate were fit to this equation (8) to estimate the coefficients a_s and n_s . The activation energies (Ea_{Mh} and Ea_{Md}) were determined as the average of energies showed in Tables 3 and 6 and P_0 was the same previously calculated.

The numerical parameter values obtained from this model are shown in Table 7. Curdlan was the substrate hydrolyzed most easily (lowest Ea_{Mh} value), and its mannose monomers were the most difficult to decompose (highest Ea_{Md} value). The values of coefficients a_s and n_s are in agreement with those reported by other groups that have used a similar mechanistic approach, but worked with different substrate types [39,43-45]. Furthermore, the

determination coefficients from the fittings were always higher than 0.93. Finally, hydrolysis kinetics simulations were performed using equation (8) in a range from 50°C to 130°C with the TFA concentrations tested in this study (Figure 4). This representation allowed us to predict the kinetic profiles due to the joint effect of temperature and TFA on the hydrolysis of curdlan, mannan and the *E. fibuliger* CW.

4. Conclusions

The methods used for CW compositional analysis and the processes developed to produce active oligosaccharides and monosaccharides from complex polysaccharides are based on a chemical reaction in acid media at high temperatures. It is therefore necessary to optimize the effect of these variables (i.e., temperature and acid concentration) on the hydrolysis kinetics. However, the optimal conditions for breaking down curdlan, mannan and yeast CW to obtain the maximum production of glucose and mannose and to avoid the decomposition of these sugars have not yet been determined.

The experimental results showed that the mannose polymers, both standard and obtained from CWs, were more easily hydrolyzed (i.e., lowest Ea_{Mh} value) than those formed by glucose units. In addition, it was more difficult to destroy mannose than glucose (i.e., highest Ea_{Md} value). The most suitable conditions for maximal sugar release of the three studied substrates are between 50 and 70% TFA at 100°C, with a processing time interval of between 2.3 and 4.58 h. In all of the cases that were assessed, and with both a theoretical and empirical approach, the mathematical modeling of the hydrolysis reactions was statistically significant and consistent, and the equations accurately predicted the experimental profiles.

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Figure Captions

Figure 1: Time course of *E. fibuliger* culture on MPW-based medium. X: biomass (▲); Pr: proteins (○); Gl: glycogen (●); RS: reducing sugars (△); TAA: total amylolytic activity (◆) and pH (■). Error bars are the confidence intervals ($\alpha=0.05$, $n=2$).

Figure 2: The kinetics of glucose and mannose released (%) from the hydrolysis of polysaccharides mannan and curdlan at different temperatures (▲: 35°C; ●: 80°C; △: 100°C; ○: 120°C) and concentrations of TFA (%). The experimental data (points) were fit to equation (2) (solid lines). For clarity, confidence intervals (in all cases less than 5% of the experimental mean value; $\alpha=0.05$; $n=2$) were omitted.

Figure 3: The kinetics of glucose, mannose and reducing sugars released (%) from *E. fibuliger* CW hydrolysis at different temperatures (▲: 35°C; ●: 80°C; △: 100°C; ○: 120°C) and concentrations of TFA (%). The experimental data (points) were fit to equation (2) (solid lines). For clarity, confidence intervals (in all cases less than 5% of the experimental mean value; $\alpha=0.05$; $n=2$) were omitted.

Figure 4: Simulations of monosaccharide hydrolysis kinetics by equation (8) at different temperatures and TFA concentrations in the curdlan, mannan and CW substrates. 1: 130°C; 2: 120°C; 3: 110°C; 4: 100°C; 5: 90°C; 6: 80°C; 7: 70°C; 8: 60°C; 9: 50°C. For clarity, confidence intervals (in all cases less than 5% of the experimental mean value; $\alpha=0.05$; $n=2$) were omitted.

Table Captions

Table 1: Kinetic parameter values for equation (2) that describing the hydrolysis process of curdlan and mannan at different temperatures and TFA concentrations. Values $\pm$ Confidence Intervals for $\alpha=0.05$; p -value from Fisher's F -test ($\alpha=0.05$); R^2 : determination coefficients between experimental and predicted data; (-): zero value was obtained. NS: not significant.

Table 2: Numerical values of time for achieving the maximum monosaccharides concentration (t_m) and maximum monosaccharides concentration (M_m) at the different temperatures and TFA concentrations tested. These values were calculated using equations (3) and (4) with the parameters summarized in Table 1. (-): no value was obtained.

Table 3: Effect of temperature on the kinetic parameters (k_h and k_d) shown in Table 1 for curdlan and mannan hydrolysis. The bivariate equation (6) was used to fit these parameters. Values $\pm$ Confidence Intervals for $\alpha=0.05$; p -value from Fisher's F -test ($\alpha=0.05$); R^2 : determination coefficients between experimental and predicted data. NS: not significant.

Table 4: Kinetic parameters values for equation (2), which describes the hydrolysis process of *E. fibuliger* CW at different temperatures and TFA concentrations. The dependent variables were glucose, mannose and reducing sugars as measured by GC-MS and spectrophotometry, respectively. Values $\pm$ Confidence Intervals for $\alpha=0.05$; p -value from Fisher's F -test ($\alpha=0.05$); R^2 : determination coefficients between experimental and predicted data; (-): zero value was obtained. NS: not significant.

Table 5: Numerical values of the time for achieving the maximum monosaccharides concentration (t_m) and maximum monosaccharides concentration (M_m) at the different temperatures and TFA concentrations tested. These values were calculated using equations (3) and (4) with the parameters summarized in Table 4. (-): no value was obtained.

Table 6: Effect of temperature on the kinetic parameters (k_h and k_d) shown in Table 4 for the hydrolysis of *E. fibuliger* CW. The bivariate equation (6) has been employed to fit these parameters. Values $\pm$ Confidence Intervals for $\alpha=0.05$; p -value from Fisher's F -test ($\alpha=0.05$); R^2 : determination coefficients between experimental and predicted data. NS: not significant.

Table 7: Numerical estimates obtained using equation (8). Values $\pm$ Confidence Intervals for $\alpha=0.05$; p -value from Fisher's F -test ($\alpha=0.05$); R^2 : determination coefficients between experimental and predicted data.

TABLES

Table 1

PARAMETERS	TFA (%)				
	10	20	35	50	70
CURDLAN					
P_0 (%)	100 ± 2.79	100 ± 2.79	100 ± 2.79	100 ± 2.79	100 ± 2.79
k_h (h ⁻¹)-T=35°C	1 × 10 ⁻²⁰ (NS)	1 × 10 ⁻²⁰ (NS)	6 × 10 ⁻⁵ (NS)	7 × 10 ⁻⁶ (NS)	7 × 10 ⁻⁵ (NS)
k_h (h ⁻¹)-T=80°C	0.010 ± 0.005	0.058 ± 0.008	0.093 ± 0.012	0.320 ± 0.043	0.550 ± 0.095
k_h (h ⁻¹)-T=100°C	0.060 ± 0.009	0.330 ± 0.047	0.500 ± 0.083	1.291 ± 0.540	1.897 ± 0.196
k_h (h ⁻¹)-T=120°C	0.244 ± 0.036	0.671 ± 0.148	1.298 ± 0.660	2.879 ± 0.513	5.200 ± 1.916
k_d (h ⁻¹)-T=35°C	5.8 × 10 ⁻⁴ (NS)	5.8 × 10 ⁻⁴ (NS)	5.8 × 10 ⁻⁴ (NS)	5.8 × 10 ⁻⁴ (NS)	5.8 × 10 ⁻⁴ (NS)
k_d (h ⁻¹)-T=80°C	0.003 ± 0.002	0.004 ± 0.003	0.005 ± 0.003	0.004 ± 0.002	0.002 ± 0.000
k_d (h ⁻¹)-T=100°C	0.014 ± 0.005	0.005 ± 0.002	0.013 ± 0.002	0.018 ± 0.003	0.027 ± 0.003
k_d (h ⁻¹)-T=120°C	0.046 ± 0.006	0.053 ± 0.006	0.103 ± 0.012	0.133 ± 0.012	0.240 ± 0.070
R ²	0.967	0.993	0.995	0.986	0.992
p-values	<0.001	<0.001	<0.001	<0.001	<0.001
MANNAN					
P_0 (%)	100 ± 3.68	100 ± 3.68	100 ± 3.68	100 ± 3.68	100 ± 3.68
k_h (h ⁻¹)-T=35°C	2.6 × 10 ⁻⁵ (NS)	4.8 × 10 ⁻⁶ (NS)	4.8 × 10 ⁻⁶ (NS)	2.8 × 10 ⁻⁶ (NS)	4.8 × 10 ⁻⁶ (NS)
k_h (h ⁻¹)-T=80°C	0.025 ± 0.006	0.045 ± 0.008	0.172 ± 0.022	0.321 ± 0.047	0.197 ± 0.026
k_h (h ⁻¹)-T=100°C	0.129 ± 0.017	0.161 ± 0.022	0.689 ± 0.147	1.402 ± 0.692	2.282 ± 1.293
k_h (h ⁻¹)-T=120°C	0.580 ± 0.132	0.573 ± 0.130	1.819 ± 0.380	2.996 ± 2.160	5.438 ± 0.434
k_d (h ⁻¹)-T=35°C	6.8 × 10 ⁻⁶ (NS)	6.8 × 10 ⁻⁶ (NS)	6.8 × 10 ⁻⁶ (NS)	6.8 × 10 ⁻⁶ (NS)	6.8 × 10 ⁻⁶ (NS)
k_d (h ⁻¹)-T=80°C	0.001 ± 0.000	0.005 ± 0.001	0.006 ± 0.003	0.005 ± 0.003	0.017 ± 0.004
k_d (h ⁻¹)-T=100°C	0.026 ± 0.005	0.029 ± 0.005	0.031 ± 0.005	0.040 ± 0.006	0.087 ± 0.015
k_d (h ⁻¹)-T=120°C	0.160 ± 0.022	0.164 ± 0.022	0.162 ± 0.026	0.192 ± 0.052	0.500 ± 0.210
R ²	0.984	0.990	0.994	0.993	0.990
p-values	<0.001	<0.001	<0.001	<0.001	<0.001

Table 2

PARAMETERS	TFA (%)				
	10	20	35	50	70
CURDLAN					
t_m (h)–T=35°C	-	-	-	-	-
M_m (%)–T=35°C	-	-	-	-	-
t_m (h)–T=80°C	173.88	48.84	33.47	14.20	10.74
M_m (%)–T=80°C	59.67	81.19	85.02	94.76	98.42
t_m (h)–T=100°C	31.74	12.59	7.56	3.37	2.28
M_m (%)–T=100°C	64.06	93.57	90.72	94.23	94.10
t_m (h)–T=120°C	8.45	4.10	2.12	1.12	0.62
M_m (%)–T=120°C	67.88	80.36	80.39	86.17	86.17
MANNAN					
t_m (h)–T=35°C	-	-	-	-	-
M_m (%)–T=35°C	-	-	-	-	-
t_m (h)–T=80°C	144.03	54.92	20.09	13.10	13.69
M_m (%)–T=80°C	89.07	76.25	88.35	93.78	79.47
t_m (h)–T=100°C	15.54	12.83	4.70	2.60	1.49
M_m (%)–T=100°C	66.40	69.05	86.34	90.00	87.83
t_m (h)–T=120°C	3.05	3.07	1.48	0.98	0.48
M_m (%)–T=120°C	61.08	60.45	79.41	82.87	78.53

Table 3

PARAMETERS	TFA (%)				
	10	20	35	50	70
CURDLAN					
<i>Ea_h</i> (kJ/mol)	74.49 ± 13.79	71.25 ± 12.74	69.2 ± 13.13	66.55 ± 9.74	64.71 ± 8.85
<i>k_{h0}</i> (h ⁻¹)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)
<i>Ea_d</i> (kJ/mol)	119.3 ± 72.7	118.0 ± 40.9	117.0 ± 26.5	115.9 ± 24.1	114.0 ± 19.7
<i>k_{d0}</i> (h ⁻¹)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)
R ²	0.941	0.979	0.977	0.954	0.962
<i>p</i> -values	<0.001	<0.001	<0.001	<0.001	<0.001
MANNAN					
<i>Ea_h</i> (kJ/mol)	71.99 ± 19.55	71.98 ± 19.60	68.11 ± 17.04	66.40 ± 13.19	64.53 ± 13.11
<i>k_{h0}</i> (h ⁻¹)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)	2 × 10 ⁹ (NS)
<i>Ea_d</i> (kJ/mol)	115.3 ± 38.5	115.2 ± 38.3	115.3 ± 32.3	114.7 ± 29.2	111.6 ± 23.4
<i>k_{d0}</i> (h ⁻¹)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)
R ²	0.970	0.987	0.974	0.953	0.983
<i>p</i> -values	<0.001	<0.001	<0.001	<0.001	<0.001

Table 4

PARAMETERS	TFA (%)				
	10	20	35	50	70
GLUCOSE as dependent variable					
P_0 (%)	41.0 ± 5.0	41.0 ± 5.0	41.0 ± 5.0	41.0 ± 5.0	41.0 ± 5.0
k_h (h ⁻¹)-T=35°C	1.3 × 10 ⁻³ (NS)	8 × 10 ⁻⁴ (NS)	1 × 10 ⁻³ (NS)	8.6 × 10 ⁻⁴ (NS)	2.5 × 10 ⁻³ (NS)
k_h (h ⁻¹)-T=80°C	0.016 ± 0.010	0.020 ± 0.011	0.028 ± 0.012	0.049 ± 0.015	0.093 ± 0.022
k_h (h ⁻¹)-T=100°C	0.062 ± 0.018	0.151 ± 0.037	0.188 ± 0.047	0.204 ± 0.052	0.589 ± 0.219
k_h (h ⁻¹)-T=120°C	0.236 ± 0.068	0.429 ± 0.140	0.543 ± 0.219	0.612 ± 0.290	2.258 ± 0.139
k_d (h ⁻¹)-T=35°C	1.5 × 10 ⁻⁵ (NS)	1.4 × 10 ⁻⁶ (NS)	1.4 × 10 ⁻⁶ (NS)	7.5 × 10 ⁻⁶ (NS)	7.5 × 10 ⁻⁶ (NS)
k_d (h ⁻¹)-T=80°C	0.015 ± 0.000	0.025 ± 0.003	0.035 ± 0.024	0.043 ± 0.016	0.039 ± 0.011
k_d (h ⁻¹)-T=100°C	0.026 ± 0.007	0.078 ± 0.012	0.053 ± 0.013	0.053 ± 0.013	0.049 ± 0.013
k_d (h ⁻¹)-T=120°C	0.152 ± 0.029	0.204 ± 0.048	0.321 ± 0.075	0.437 ± 0.110	0.460 ± 0.448
R ²	0.963	0.961	0.978	0.991	0.990
p-values	<0.001	<0.001	<0.001	<0.001	<0.001
MANNOSE as dependent variable					
P_0 (%)	24.5 ± 1.5	24.5 ± 1.5	24.5 ± 1.5	24.5 ± 1.5	24.5 ± 1.5
k_h (h ⁻¹)-T=35°C	1.7 × 10 ⁻³ (NS)	4.3 × 10 ⁻³ (NS)	4.1 × 10 ⁻³ (NS)	3.4 × 10 ⁻³ (NS)	4.1 × 10 ⁻³ (NS)
k_h (h ⁻¹)-T=80°C	0.022 ± 0.010	0.030 ± 0.011	0.036 ± 0.011	0.084 ± 0.017	0.150 ± 0.029
k_h (h ⁻¹)-T=100°C	0.124 ± 0.024	0.335 ± 0.072	0.629 ± 0.180	0.790 ± 0.260	0.852 ± 0.310
k_h (h ⁻¹)-T=120°C	0.579 ± 0.179	0.981 ± 0.492	2.384 ± 1.300	2.835 ± 0.011	3.145 ± 0.027
k_d (h ⁻¹)-T=35°C	1.5 × 10 ⁻⁵ (NS)	3.2 × 10 ⁻³ (NS)	3.4 × 10 ⁻³ (NS)	1 × 10 ⁻¹¹ (NS)	3.9 × 10 ⁻⁷ (NS)
k_d (h ⁻¹)-T=80°C	0.015 ± 0.000	0.013 ± 0.002	0.016 ± 0.013	0.021 ± 0.007	0.040 ± 0.009
k_d (h ⁻¹)-T=100°C	0.026 ± 0.007	0.025 ± 0.006	0.032 ± 0.007	0.037 ± 0.008	0.055 ± 0.011
k_d (h ⁻¹)-T=120°C	0.152 ± 0.029	0.205 ± 0.038	0.298 ± 0.189	0.371 ± 0.192	0.461 ± 0.248
R ²	0.953	0.954	0.978	0.989	0.989
p-values	<0.001	<0.001	<0.001	<0.001	<0.001
REDUCING SUGARS as dependent variable					
P_0 (%)	56.6 ± 3.7	56.6 ± 3.7	56.6 ± 3.7	56.6 ± 3.7	56.6 ± 3.7
k_h (h ⁻¹)-T=35°C	4.3 × 10 ⁻⁴ (NS)	2.2 × 10 ⁻⁴ (NS)	7 × 10 ⁻⁴ (NS)	1.5 × 10 ⁻⁴ (NS)	3.6 × 10 ⁻³ (NS)
k_h (h ⁻¹)-T=80°C	0.030 ± 0.009	0.053 ± 0.012	0.072 ± 0.014	0.102 ± 0.016	0.173 ± 0.026
k_h (h ⁻¹)-T=100°C	0.154 ± 0.023	0.328 ± 0.053	0.477 ± 0.090	0.602 ± 0.130	1.150 ± 0.460
k_h (h ⁻¹)-T=120°C	0.411 ± 0.086	0.834 ± 0.279	1.126 ± 0.621	1.064 ± 0.590	2.054 ± 0.580
k_d (h ⁻¹)-T=35°C	3.2 × 10 ⁻³ (NS)	3.2 × 10 ⁻³ (NS)	3.2 × 10 ⁻³ (NS)	3.2 × 10 ⁻³ (NS)	3.2 × 10 ⁻³ (NS)
k_d (h ⁻¹)-T=80°C	0.013 ± 0.001	0.023 ± 0.009	0.029 ± 0.008	0.018 ± 0.005	0.021 ± 0.005
k_d (h ⁻¹)-T=100°C	0.021 ± 0.005	0.019 ± 0.004	0.022 ± 0.004	0.028 ± 0.005	0.029 ± 0.005
k_d (h ⁻¹)-T=120°C	0.081 ± 0.013	0.135 ± 0.020	0.214 ± 0.033	0.276 ± 0.044	0.380 ± 0.330
R ²	0.980	0.991	0.988	0.989	0.985
p-values	<0.001	<0.001	<0.001	<0.001	<0.001

Table 5

PARAMETERS	TFA (%)				
	10	20	35	50	70
GLUCOSE as variable dependent					
t_m (h)–T=35°C	-	-	-	-	-
M_m (%)–T=35°C	-	-	-	-	-
t_m (h)–T=80°C	56.83	44.17	31.94	21.84	16.14
M_m (%)–T=80°C	13.55	13.55	13.30	16.09	22.00
t_m (h)–T=100°C	17.15	11.13	9.38	8.95	4.58
M_m (%)–T=100°C	17.07	24.06	24.91	25.58	32.74
t_m (h)–T=120°C	5.45	3.30	2.37	1.93	0.88
M_m (%)–T=120°C	19.21	20.91	19.16	17.69	27.30
MANNOSE as variable dependent					
t_m (h)–T=35°C	-	-	-	-	-
M_m (%)–T=35°C	-	-	-	-	-
t_m (h)–T=80°C	54.74	48.98	40.13	21.88	12.04
M_m (%)–T=80°C	10.79	12.80	12.77	15.35	15.15
t_m (h)–T=100°C	15.94	8.41	5.00	4.08	3.44
M_m (%)–T=100°C	16.22	19.89	20.88	21.06	20.27
t_m (h)–T=120°C	3.13	2.02	1.00	0.83	0.72
M_m (%)–T=120°C	15.21	16.19	18.20	18.03	17.61
REDUCING SUGARS as variable dependent					
t_m (h)–T=35°C	-	-	-	-	-
M_m (%)–T=35°C	-	-	-	-	-
t_m (h)–T=80°C	49.74	27.74	21.07	20.80	13.85
M_m (%)–T=80°C	30.25	29.81	30.71	39.11	42.24
t_m (h)–T=100°C	15.02	9.23	6.78	5.36	3.27
M_m (%)–T=100°C	41.31	47.57	48.84	48.81	51.42
t_m (h)–T=120°C	4.93	2.61	1.82	1.71	1.01
M_m (%)–T=120°C	38.05	39.84	38.34	35.27	38.60

Table 6

PARAMETERS	TFA (%)				
	10	20	35	50	70
GLUCOSE as variable dependent					
Ea_h (kJ/mol)	84.93 ± 21.73	83.10 ± 19.64	81.84 ± 21.27	81.46 ± 23.06	77.27 ± 21.83
k_{ho} (h ⁻¹)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)
Ea_d (kJ/mol)	115.6 ± 50.9	114.5 ± 35.6	113.0 ± 33.0	112.0 ± 33.2	111.9 ± 23.3
k_{do} (h ⁻¹)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)	3 × 10 ¹⁴ (NS)
R ²	0.954	0.989	0.993	0.990	0.990
p-values	<0.001	<0.001	<0.001	<0.001	<0.001
MANNOSE as variable dependent					
Ea_h (kJ/mol)	81.74 ± 27.11	79.92 ± 30.55	77.10 ± 30.58	76.52 ± 28.30	76.19 ± 26.97
k_{ho} (h ⁻¹)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)	4 × 10 ¹⁰ (NS)
Ea_d (kJ/mol)	94.03 ± 45.87	93.10 ± 37.51	91.89 ± 28.31	91.18 ± 26.28	90.46 ± 25.12
k_{do} (h ⁻¹)	5 × 10 ¹¹ (NS)	5 × 10 ¹¹ (NS)	5 × 10 ¹¹ (NS)	5 × 10 ¹¹ (NS)	5 × 10 ¹¹ (NS)
R ²	0.992	0.986	0.984	0.982	0.981
p-values	<0.001	<0.001	<0.001	<0.001	<0.001
REDUCING SUGARS as variable dependent					
Ea_h (kJ/mol)	80.48 ± 17.00	78.15 ± 19.01	77.15 ± 21.31	77.21 ± 21.91	75.07 ± 20.24
k_{ho} (h ⁻¹)	2 × 10 ¹⁰ (NS)	2 × 10 ¹⁰ (NS)	2 × 10 ¹⁰ (NS)	2 × 10 ¹⁰ (NS)	2 × 10 ¹⁰ (NS)
Ea_d (kJ/mol)	119.9 ± 53.1	118.2 ± 34.8	116.8 ± 29.2	115.9 ± 28.3	114.9 ± 24.8
k_{do} (h ⁻¹)	7 × 10 ¹⁴ (NS)	7 × 10 ¹⁴ (NS)	7 × 10 ¹⁴ (NS)	7 × 10 ¹⁴ (NS)	7 × 10 ¹⁴ (NS)
R ²	0.989	0.977	0.979	0.993	0.989
p-values	<0.001	<0.001	<0.001	<0.001	<0.001

Table 7

	CURDLAN	MANNAN	CW (Glucose)	CW (Mannose)	CW (RS)
Ea_{Mh} (kJ/mol)	69.78 ± 0.05	70.03 ± 0.02	82.32 ± 0.16	79.92 ± 0.23	70.68 ± 0.11
a_h	16.14 ± 0.01	15.48 ± 0.01	15.10 ± 0.05	22.38 ± 0.07	18.74 ± 0.03
n_h	1.63 ± 0.00	1.79 ± 0.00	2.56 ± 0.00	0.77 ± 0.00	0.84 ± 0.00
Ea_{Md} (kJ/mol)	114.07 ± 0.08	117.27 ± 0.06	113.25 ± 0.20	92.63 ± 0.23	115.03 ± 0.23
a_d	29.69 ± 0.02	29.48 ± 0.02	31.33 ± 0.06	24.97 ± 0.07	30.77 ± 0.07
n_d	1.03 ± 0.00	1.16 ± 0.00	0.61 ± 0.01	0.60 ± 0.00	0.81 ± 0.01
R^2	0.983	0.992	0.946	0.980	0.931
p-values	<0.001	<0.001	<0.001	<0.001	<0.001

FIGURES

Figure 1

Figure 2

Figure 3

Figure 4