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Monitoring the non-specific interactions of catechin through diffusion measurements based on pulsed-field gradients

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The self-association of aqueous catechin as a function of concentration was monitored through variations in ¹H chemical shifts, proton T_1 and T_2 data and translational diffusion coefficients obtained with the pulsed-field gradient spin-echo method. The latter approach is very efficient and it is not restricted to aromatic compounds. Equilibrium constants were estimated for various models of self-association and the apparent enthalpy of dissociation was measured with isothermal titration calorimetry. Comparison of the latter parameter with thermodynamic data reported for various types of non-specific interactions suggests that such phenomena could be studied using this approach.

INTRODUCTION

Non-specific interactions play a major role in the aggregation phenomena that govern the physical state of molecular mixtures. This is particularly true of plant extracts where control of colloid formation and/or precipitation is a major industrial concern. Although much effort has been devoted to devising NMR experiments for studying specific interactions such as ligand binding, non-specific interactions have received less attention ^{1,2}. The aggregation phenomena of surfactants ^{3,4}, nucleotides ^{5,6}, caffeine ^{7,8}, polyphenol ⁸⁻¹³ and aromatic drug molecules ¹⁴⁻¹⁶ and the non-specific stacking interactions of caffeine with methyl gallate ⁸, polyphenols with peptides ^{10,13,17} and aroma substances ¹² are notable exceptions. The majority of these NMR studies of self- or hetero-association involve aromatic compounds and have relied on chemical shift titration versus concentration to quantify the equilibrium constants ^{7,12,14-16}. However, in several cases no significant chemical shift variations were observed in spite of the detection of line broadening and/or intermolecular NOEs that indicated hetero-association ^{13,16,17}.

For many types of molecules, where non-specific interactions are not expected to produce significant chemical shift changes, monitoring variations in molecular motion (either rotational ¹⁸ or translational diffusion ³⁻⁶) would seem to be an attractive alternative. Measurement of self-diffusion coefficients is rapid ^{19,20} and prediction of the variations in translational motion on the basis of hydrodynamic theory ²¹ is straightforward as long as the molecules are well represented by the Stokes-Einstein-Debye model ('stick regime') ²². In this work, the self-association of catechin was studied to explore the feasibility of such an approach.

Catechin (*2,3-trans*) and epicatechin (*2,3-cis*) (**Fig. 1**) are the basic monomeric units of condensed tannins which are ubiquitous secondary metabolites of higher plants. Tannins are thought to participate in plant defense by rendering the tissues of leaves, bark and fruit unpalatable for herbivores and omnivores. These polyphenols display a strong propensity to form complexes with other biomolecules such as proteins, polysaccharides and alkaloids. This phenomenon is responsible for the astringency (precipitation of salivary proteins) and poor nutritional quality (irreversible complexation of gut enzymes and dietary proteins, inhibition of iron absorption) of polyphenol-rich foods ¹⁰. The dimerization of catechin *in silico* has afforded both parallel and anti-parallel orientations of the target molecules ²³ corroborating the nonspecific nature of catechin self-association.

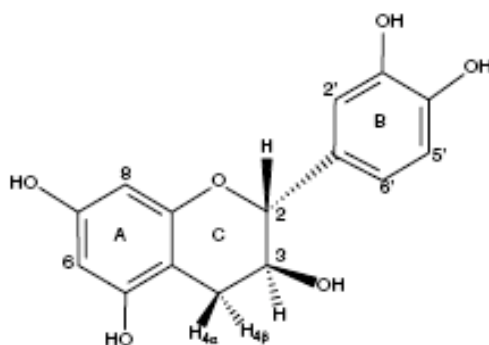
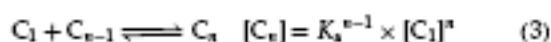
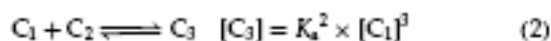
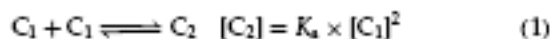


Figure 1. Structure of catechin with atom numbering

RESULTS AND DISCUSSION

Chemical shift titrations of catechin as a function of concentration have been investigated previously^{11,12} and self-association constants ($4.3 < K_a < 7.4$) were established from the latter data for the isodesmic model.¹² This model of solute self-association is based on the assumption that the solute molecules associate to form stacks (dimers, trimers, etc.) with an equilibrium constant, K_a , that is the same for each step. This leads to simple expressions for the fractional populations of the various species as a function of the monomer concentration $[C_1]$ and K_a :



The limiting value of the Taylor series for the sum of the fractional populations yields the following expression for the total concentration $[C_0]$ as long as the product $K_a \times [C_1]$ is < 1 :

$$[C_0] = [C_1]/(1 - K_a \times [C_1])^2 \quad (4)$$

A single chemical shift is observed for each spin, indicating that the various species are undergoing fast exchange. As the distances between vertically stacked molecules are in the 3–4 Å range and the ring current magnetic field giving the main contribution to nuclear shielding decreases rapidly with the distance ($\sim 1/r^3$), it is assumed that the upfield chemical shifts on complexation are due only to the influence of the neighboring molecule (within the limits of this model of molecular association all such contributions are assumed to be equal). The chemical shift of the C_n species and the chemical shift dependence on concentration for the isodesmic model are given by Eqs (5) and (6), respectively¹⁴:

$$\delta_{C_n} = (n - 1)\delta_{\max} + \delta_{C_1}/n \quad (5)$$

$$\delta_{\text{obs}} - \delta_{C_1} = (\Delta\delta_{\max}) \left(\frac{2K_a[C_0] + 1 - \sqrt{(4K_a[C_0] + 1)}}{2K_a[C_0]} \right) \quad (6)$$

A very similar expression is obtained for the dependence of δ_{obs} on concentration in the case of dimerization¹⁴:

$$\delta_{\text{obs}} - \delta_{C_1} = (\Delta\delta_{\max}) \left(\frac{4K_d[C_0] + 1 - \sqrt{(8K_d[C_0] + 1)}}{4K_d[C_0]} \right) \quad (7)$$

If it is assumed that each successive stack of the catechin monomer affords the same decrease in D_{obs} , ΔD_t , the dependence of D_{obs} on total concentration will be analogous to that of the chemical shifts for both the isodesmic (Eq 6) and dimerization (Eq 7) models.

¹H NMR chemical shifts of catechin as a function of $[C_0]$ are given in **Table 1**. The H-3, H-4 α and H-4 β signals display the most significant variations in δ_{obs} with increasing concentration. $\Delta\delta_{\max}$ and K_a values were calculated by leastsquares minimization of the experimental data ($\delta_{\text{obs}} - \delta_{C_1}$) with respect to the theoretical values in Eqs (6) and (7) using the simplex method²⁴ and the modeled data are given in **Table 2**.

Spin	2.3	4.6	8.0	11.5	17.2	23.0	27.6	34.4	41.3	45.9
H-2	—c	—c	—c	—c	—c	4.77	4.77	4.73	4.73	4.74
H-3	4.28	4.28	4.28	4.27	4.27	4.23	4.23	4.23	4.23	4.20
H-4a	2.90	2.90	2.90	2.90	2.90	2.89	2.88	2.89	2.88	2.87
H-4b	2.58	2.58	2.58	2.58	2.58	2.55	2.55	2.54	2.54	2.52
H-20	6.89	6.89	6.89	6.89	6.89	6.88	6.89	6.88	6.88	6.86

Table 1. ¹H NMR chemical shifts (ν_{obs} , ppm) for various concentrations of catechin^a Catechin concentration (mM)

^a Catechin (Sigma) was added to 1.75 mM solutions of acetic acid (SDS, purex 99.8%) in D₂O to afford solutions (0.75 ml) containing the above total concentrations, $[C_0]$. Both the HOD and acetic acid methyl signals were evaluated as internal references for the chemical shifts ($\delta_{\text{HOD}} = 4.800$ ppm was the internal standard for the data given above).

^b See Fig. 1 for the numbering.

^c Overlapping with the HOD signal.

Experimental probe	Self-association model	K_a (M^{-1})	Simulated values of
			$\Delta\delta_{\max}$ (ppm) or ΔD_t ($cm^2 s^{-1}$)
δ_{obs} H-3	I	4.7 ± 2	-0.42
	D	2.0 ± 0.8	-0.50
δ_{obs} H-4 α	I	2.0 ± 0.8	-0.30
	D	1.0 ± 0.9	-0.18
δ_{obs} H-4 β	I	3.2 ± 0.9	-0.33
	D	2.3 ± 1.3	-0.31
D_t^{obs} H-2'	I	8 ± 4.5	-6.7×10^{-6}
	D	8.5 ± 4	-4.4×10^{-6}

Table 2. Self-association constants of catechin, K_a , for isodesmic (I) and dimerization (D) models

The average value of K_a ($3.3 M^{-1}$ using all of the data for H-3, H-4, and H-4 β) is slightly lower than the average value ($5.6 M^{-1}$) reported by Dufour and Bayonove¹² for catechin dissolved in D_2O - H_2O -ethanol- d_6 (8 : 1 : 1) but the spread in K_a parameters ($\pm 50\%$) for the various spins is very similar. Comparison of the chemical shifts of the acetic acid methyl moiety with respect to those of the HOD resonance for the 10 samples listed in Table 1 yields an average value of 2.084 ± 0.008 ppm in spite of a digital resolution of ± 0.001 ppm. Unfortunately, in the present case, it is not feasible to acquire data for a much larger range of concentrations owing to the poor signal-to-noise ratio at lower concentrations and precipitation at higher concentrations. Hence it is difficult to improve the convergence of data from chemical shift titrations and the adjustable parameter $\Delta\delta_{\max}$ cannot be used to validate the simulations as it is not easily calculated.

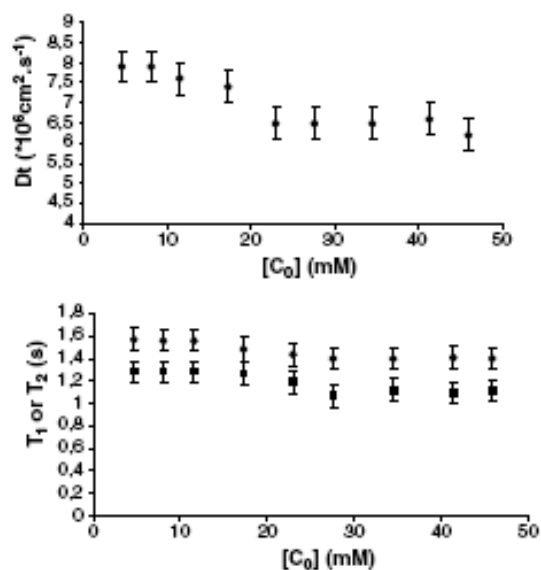


Figure 2. Plots of T_1 (\blacklozenge), T_2 (\blacksquare) and D_t for various total concentrations of catechin measured at 298 K and 400 MHz. Experimental error bars are indicated.

Plots of T_1 , T_2 and D_t as a function of concentration are shown in **Fig. 2**. The error bars for T_1 and T_2 are as large as the variations in the relaxation parameters, suggesting that rotational dynamics are not sufficiently modulated by association phenomena when the target molecules are of similar size. In contrast, more readily detectable variations in D_t indicate that translational diffusion is a better probe of the non-specific interactions of medium-sized molecules. In favorable cases, the experimental precision of D_t measurements is about 1%^{19,20} and standard deviations of $0.05 \times 10^{-6} cm^2 s^{-1}$ have been obtained from the attenuation of anomeric signals of carbohydrates ($1 < DP < 10$)²⁵ or from the B-ring signal (isolated singlets) of galloylated tannins. Owing to deuterium exchange with the solvent, the intensity of the catechin H-6 and H-8 singlets

evolved with time. Therefore, the attenuation of the H-2' signal was used to establish the D^{obs}_t values, but its intensity could not be evaluated with precision owing to the nearby H-5' and H-6' multiplets (the high-field H-3, H-4, and H-4' signals were even less suitable owing to phase distortion induced by J -modulation during the echo sequence), with the result that standard deviations for D^{obs}_t were almost an order of magnitude larger ($\pm 0.4 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$) than those referred to above. A K_a value of $\sim 8 \text{ M}^{-1}$ was obtained by leastsquares minimization for both isodesmic and dimerization models of self-association.

The D^{obs}_t coefficient of $7.9 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$ for the most dilute solution is much higher than the value of $3.6 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$ predicted on the basis of hydrodynamic theory²¹ (molecular models of catechin could be assimilated to a cylinder 14 Å in length with a radius of 3.8 Å), indicating that a microviscosity correction factor, f , of 0.45 would be necessary. The simulation protocol afforded ΔD_t values of -6.7×10^{-6} and $-4.4 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$ for the isodesmic and dimerization models of self-association, respectively. As the catechin monomer does not obey the Stokes–Einstein–Debye model, it is not possible to predict this adjustable parameter accurately. However, upper and lower limits for the microviscosity factor are known ($0.45 < f < 1.0$) and the D_t value for the dimer ($1.2 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$) that can be extrapolated from the ΔD_t parameter of the isodesmic model is clearly too low (with $f = 1.0$ this would correspond to a sphere of radius 16.6 Å, where $D_t = kT/6\zeta r_0$), whereas that of the dimerization model ($3.5 \times 10^{-6} \text{ cm}^2 \text{ s}^{-1}$) is reasonable.

A plot of the apparent enthalpy of dilution as a function of catechin concentration is shown in **Fig. 3**. Dissociation is endothermic and can be detected even at the lowest concentration (open squares). The apparent molar enthalpy of dilution (6.5 kJ mol^{-1}) includes both dilution and dissociation contributions and has not been treated quantitatively in this study.²⁶ Plots of theoretical monomer and dimer fractional populations (%) for both the isodesmic and dimerization models as a function of total catechin concentration are displayed in **Fig. 4**. The K_a value estimated from the translational diffusion profile (8 M^{-1}) was used to establish the fractional $[C_1]$ and $[C_2]$ populations. For both models of self-association, the predicted dimer population of the 2.3 mM sample is compatible with the reduced (3/410% of that of a 23 mM solution) apparent enthalpy of dilution that is observed experimentally. It appears that dimerization is followed by precipitation without significant accumulation of aggregates higher than dimers.

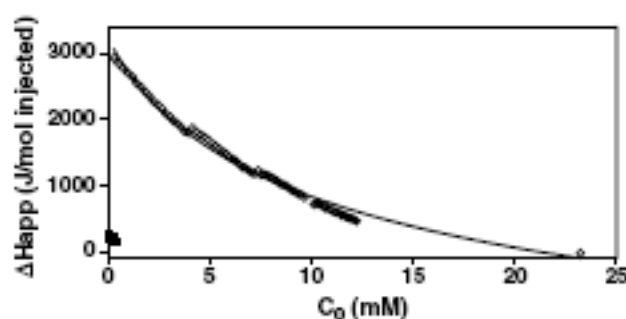


Figure 3. Plots of the apparent enthalpy change (J mol^{-1}) on diluting aqueous solutions of catechin [2.3 mM (\square) and 23.3 mM (\circ)] versus total catechin concentration $[C_0]$ in the cell.

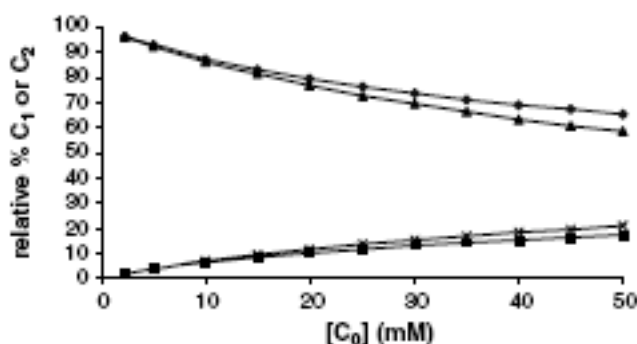


Figure 4. Plots of the relative percentage of monomer ($[C_1]$) and dimer ($[C_2]$) concentrations as a function of the total concentration ($[C_0]$) of catechin for both dimerization (\blacklozenge, \times) and isodesmic ($\blacksquare, \blacktriangle$) models characterized by an equilibrium constant, K_a , of 8.

In conclusion, it has been demonstrated that self-association of catechin can be monitored with translational diffusion measurements based on the pulsed-field gradient spin-echo method. The thermodynamic data for various non-specific interactions involving sugars (carbohydrate–surfactant and carbohydrate–carbohydrate interactions) has been reviewed recently²⁷ and comparison with results from the present work suggests that these systems should be amenable to study using translational diffusion measurements. In favorable cases (isolated NMR signals), the association constants should be more accurately defined than those obtained from chemical shift titrations and this approach is not limited to aromatic molecules. An added advantage of the D_t approach, in the case of moderately sized molecules (for example, tetrasaccharides²⁵), is that D_t can be related to the dimensions of the molecular assembly. As pointed out by a referee, other data treatment such as biexponential fitting²⁰ associated with the BPPLIED pulse sequence²⁸ (here the composite pulses provide self-compensation of induced eddy currents) might be useful for extracting the diffusion coefficients of the monomers and the dimers.

EXPERIMENTAL

Sample preparation

Various amounts of catechin (Sigma) were added to 1.75 mM solutions of acetic acid (SDS, purex 99.8%) in D₂O to afford solutions (0.75 ml) containing the total concentrations, [C₀] indicated in **Table 1**. The NMR tubes were sealed under argon after vacuum removal of dissolved oxygen.

Isothermal titration calorimetry

Calorimetric data were obtained using a VP-ITC microcalorimeter (MicroCal, USA). Experiments were performed at 298 K. Solutions were degassed under vacuum just prior to the experiment. Aliquots of 15 μl of catechin solution were automatically injected from a microsyringe at 5 min intervals into the sample cell (1.4192 ml) containing acetic acid solution. Efficient mixing of the solution was achieved by a paddle on the end of the syringe rotated at constant speed (310 rpm). The power versus time plot shows a peak as heat is absorbed upon equilibrium displacement by dilution, and then returns to baseline as thermal equilibrium is reached. The area under the peaks gives the amount of heat absorbed. The data were analysed using Microcal Origin software^{29,30}.

NMR

¹H NMR pulsed-gradient spin-echo experiments were conducted on a Bruker Avance 400 spectrometer at 298 K. The stimulated spin-echo sequence³¹ was used to measure the translational self-diffusion coefficients as described previously.²⁵ The gradient duration (δ) was varied from 1 to 20 ms while keeping its strength fixed at 9.15 G cm⁻¹. The intergradient delay (Δ) and gradient recovery delay were 5 ms and 50 μs, respectively. The translational self-diffusion coefficients were obtained by fitting the intensities of the H-2' signal in spectra acquired with various lengths of the gradient pulses (8–15 data points) to the Stejskal–Tanner equation with inhouse software :

$$M = 1/2M_0 \exp\left[-(\tau_2 - \tau_1)/T_1 - 2\tau_1/T_2 - (\tau_1 G \gamma)^2 D_1 [\tau_2 - (\tau_1/3)]\right] \quad (8)$$

All experiments were run at least four times. Spin–lattice relaxation times were measured with the inversion–recovery sequence. The recycle time was greater than 6xT₁ and data were collected for 10–20 τ values which varied from 5 ms to 2xT₁. Spin–spin relaxation times were obtained with the Carr–Purcell–Meiboom–Gill sequence and data were collected for roughly 20 spectra with various numbers of echos (total echo duration, 2n τ , varied from 2 ms to 1 s). The integrals of the peaks were fitted to a three- (T₁) or two- (T₂) parameter exponential function using spectrometer system software and inhouse software, respectively, and all relaxation experiments were also run at least twice.

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