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# Determination of the kinetics underlying the $pK_a$ shift for the 2-aminoanthracenium cation binding with cucurbit[7]uril†

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The binding dynamics of the 2-aminoanthracenium cation ( $AH^+$ ) and 2-aminoanthracene (A) with cucurbit[7]uril (CB[7]) was studied using stopped-flow experiments. The kinetics was followed by measuring the fluorescence changes over time for  $AH^+$  and A, which emit at different wavelengths. The studies at various pH values showed different mechanisms for the formation of the  $AH^+@CB[7]$  complex, with this complex formed either by the binding of  $AH^+$  or by the initial binding of A followed by protonation. In the latter case, it was possible to determine the protonation ( $(1.5 \pm 0.4) \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$ ) and deprotonation ( $89 \pm 7 \text{ s}^{-1}$ ) rate constants for complexed  $A/AH^+$ , which showed that the  $pK_a$  shift of +3.1 for  $A/AH^+$  in the complex is mainly due to a lower deprotonation rate constant.

## Introduction

Cucurbit[ $n$ ]urils (CB[ $n$ ]s) are macrocyclic host molecules formed from glycoluril units, which have a wide application as supramolecular hosts.<sup>1–7</sup> One of the key features of supramolecular systems is their dynamics,<sup>8</sup> which can be directly related to the intended function of a supramolecular system. CB[ $n$ ]s have been developed for a wide variety of applications, such as photocatalysis<sup>9</sup> or catalysis,<sup>10–13</sup> drug stabilization and delivery,<sup>14–17</sup> self-sorting and stimuli responsive systems,<sup>3,18–21</sup> tandem enzyme assays,<sup>22,23</sup> and control of supramolecular polymerization.<sup>24,25</sup>

The dynamics of guest complex formation with CB[7] occurs over a wide time range, from milliseconds to hours,<sup>10,26–35</sup> with rate constants for the association process as high as one order of magnitude below the diffusion controlled limit. The binding dynamics is affected by the size of the guest and the presence or

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absence of charges on the guest.<sup>10,26,30,34</sup> Exclusion complexes can be formed between the positive charge of the guest and the carbonyl moieties at the portals of CB[n] before inclusion of the guest into the interior of the CB[n] cavity.<sup>27,30,31</sup> However, in other cases, the formation of an exclusion complex is not detected in kinetic studies.<sup>32,34</sup> There are still too few reported studies on guest binding dynamics to develop a mechanistic understanding of how the guest binding dynamics can be controlled. Such an understanding is required for the rational design of systems containing CB[n]s where the dynamics is directly related to the intended function.

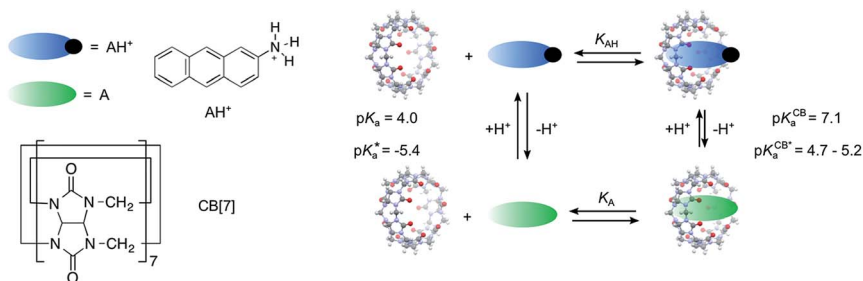
The objective of the current work is to explore one aspect of the guest–CB[n] dynamics, namely, the protonation and deprotonation reactions of a guest@CB[n] complex. This aspect is important because CB[n]s were shown to stabilize positively charged guests. In the case of protonated guests, such stabilization led to an increase of the  $pK_a$  of the CB[n]-complexed guest compared with the  $pK_a$  for the guest in water.<sup>4,15,36–38</sup> Therefore, CB[n]–guest complexation alters the protonation and deprotonation rates of bound guests compared with this reactivity in water. Understanding the origins for this change in kinetics is relevant to the design of supramolecular systems that alter chemical reactivity, including the ability to affect acid–base catalysis. In this context, CB[n]s were shown to catalyze the hydrolysis of included guests.<sup>39,40</sup>

The increase in the hydronium cation concentration at pH values where the guest is protonated led to slower kinetics when binding of the hydronium cation with CB[n] was competitive with guest binding.<sup>27,30,32</sup> The protonation state of the guest affects its binding dynamics with CB[n]s. For example, the association and dissociation rate constants of the cyclohexylmethylammonium cation with CB[6] are lower than the same rate constants for cyclohexylmethylamine because, in the case of the cation, an exclusion complex is formed before the guest is included within the cavity.<sup>30,31</sup> For pH values between the  $pK_a$  values of the guest in water and in the complex, the association process corresponds to the binding of the neutral amine followed by fast protonation of the amine@CB[6] complex, whereas dissociation corresponds to the exit of the ammonium cation from CB[6].<sup>30</sup> The kinetics for the cyclohexylmethylamine/CB[6] system occur on hour to day time scales, depending on the pH. Therefore, the acid–base equilibria for the guest in water or within the complex were fast compared to the complexation dynamics with CB[6]. On the other hand, no evidence for the formation of an exclusion complex was observed for the binding of the 2-naphthyl-1-ethylammonium cation with CB[7].<sup>32</sup>

In this work, we chose a guest, the 2-aminoanthracenium cation ( $AH^+$ ), with a lower  $pK_a$  (4.0)<sup>41</sup> than that of cyclohexylmethylamine (10.5)<sup>30</sup> to study the effect of pH on the guest binding dynamics with CB[7] (Scheme 1). The pH values of the solutions were varied between 2.0, where the guest is completely protonated ( $AH^+$ ), and pH 5.5, where the guest is mostly deprotonated (A). The singlet excited state of  $AH^+$ , which emits with a maximum at 422 nm (“blue” emission), has a much lower  $pK_a^*$  (–5.4 in 1 : 1 water : ethanol),<sup>42,43</sup> leading to the formation of singlet excited A, which emits with a maximum at 503 nm (“green” emission).<sup>38</sup> These photophysical properties provide a means of following the concentrations of  $AH^+$  and A separately.

$AH^+$  was shown to form a 1 : 1 complex with CB[7] resulting in a shift in the  $pK_a$  for ground state  $AH^+$  from 4.0 to 7.1, while the estimated excited state  $pK_a^*$  shifted





**Scheme 1** Structures of cucurbit[7]uril (CB[7]) and the 2-aminoanthracenium cation (AH<sup>+</sup>), equilibria between protonated AH<sup>+</sup>, neutral A, and CB[7], and pK<sub>a</sub> values for the ground (pK<sub>a</sub>, pK<sub>a</sub><sup>CB</sup>) and singlet excited state (pK<sub>a</sub><sup>\*</sup>, pK<sub>a</sub><sup>CB\*</sup>) of AH<sup>+</sup> in the absence<sup>41–43</sup> and presence of CB[7].<sup>38</sup>

from  $-5.4$  to between  $4.7$  and  $5.2$ .<sup>38</sup> In acidic solution, AH<sup>+</sup> was stabilized by complexation to CB[7], and its blue emission was observed.<sup>38</sup> The changes in the pK<sub>a</sub> values for free and CB[7]-complexed AH<sup>+</sup>, as well as the ability to independently follow the concentrations of AH<sup>+</sup> and A, were used to characterize the guest binding dynamics at pH values where only AH<sup>+</sup> or predominantly A were present. At low pH, the kinetics was slower than at high pH because AH<sup>+</sup> forms an encounter complex with CB[7], whereas at high pH, neutral A is in fast equilibrium with CB[7], and A@CB[7] is then protonated. Kinetic studies led to the determination of a protonation rate constant of  $(1.5 \pm 0.4) \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$  for the A@CB[7] complex and a deprotonation rate constant of  $89 \pm 7 \text{ s}^{-1}$  for AH<sup>+</sup>@CB[7], showing that the large pK<sub>a</sub> shift is mainly a reflection of the slow down of the deprotonation step.

## Experimental

### Materials

2-Aminoanthracene (Aldrich, 96%) was recrystallized from ethanol once. Sodium chloride (Sigma-Aldrich, BioUltra,  $\geq 99.5\%$ ), hydrochloric acid (Anachemia, ACS reagent grade), sodium hydroxide (Anachemia, ACS reagent grade), glacial acetic acid (ACP, ACS reagent grade), and methanol (Fisher, spectral grade,  $>99.9\%$ ) were used as received. Cucurbit[7]uril (CB[7]) was synthesized based on previous literature<sup>44–46</sup> and was purified according to the procedure described in the ESI.† Deionized water (Barnstead NANOpure deionizing systems  $\geq 17.8 \text{ M}\Omega \text{ cm}$ ) was used in the preparation of all aqueous solutions.

### Sample preparation

A 1 mM stock solution of 2-aminoanthracene was prepared in methanol. For the experiments at pH 2.0, 3.8, and 4.3, aqueous solutions were prepared by dissolving the required amounts of NaCl and 2.0 N HCl in water to achieve a final NaCl concentration of 20 mM and the required pH. The buffer solutions at pH 5.0 and 5.5 were prepared by adding the required quantity of 2.0 N NaOH to water to achieve a final sodium ion concentration of 20 mM and then titrating the solution with glacial acetic acid until the required pH was achieved. Aqueous AH<sup>+</sup> or A



solutions were prepared by diluting the methanol stock solution into the aqueous solutions of the required pH. CB[7] stock solutions (850  $\mu\text{M}$ ) were prepared by dissolving an appropriate amount of the solid in the aqueous solution of the required pH. The CB[7] stock solutions were titrated as described previously.<sup>47</sup> For the binding isotherm experiments, small aliquots of the CB[7] stock solution were injected directly into 3 mL of the  $\text{AH}^+$ /A solution with a gastight syringe. For stopped-flow experiments, a series of CB[7] solutions were prepared by diluting the stock solution into aqueous solutions of the required pH.

## Equipment

Absorption spectra were recorded on a Cary 100 UV-Vis spectrophotometer. Steady-state fluorescence measurements were performed on a PTI QM-40 spectrofluorimeter. Samples were excited at 365 nm and the emission was collected between 380 and 650 nm. A bandwidth of 2 nm was used for the excitation and emission monochromators. A baseline spectrum for a solution containing all chemicals except the fluorophore was subtracted from all emission spectra to obtain corrected spectra. All measurements were performed by maintaining the sample temperature at 20  $^{\circ}\text{C}$ . For the binding isotherm experiments, the area under each spectrum was integrated from 380 to 456 nm for the “blue” region and from 456 to 650 nm for the “green” region, corresponding to the emission of  $\text{AH}^+$  and A, respectively. These integrated intensities were then normalized by assuming unity for the integrated intensity in the absence of CB[7].

Time-resolved fluorescence decays were recorded with an Edinburgh OB920 single photon counting system. The excitation source was a light emitting diode (EPLD-360,  $\lambda_{\text{ex}} = 365 \text{ nm}$ ). The emission from the sample was collected at 405 or 510 nm using a monochromator with a bandwidth of 16 nm. The number of counts in the maximum intensity channel was 2000. The instrument response function (IRF) was recorded using a Ludox solution by collecting the emission at the excitation wavelength. The FAST (Edinburgh Instruments) software was used to fit the fluorescence decay traces. The IRF was deconvoluted with the decay during the fitting process. The quality of the fit was judged by the randomness of the residuals and the  $\chi^2$  values (0.9–1.2).<sup>48</sup> The data were fit to either a mono-exponential decay ( $i = 1$ , eqn (1)) or to a sum of two exponentials ( $i = 2$ , eqn (1)), where each species has a lifetime ( $\tau_i$ ) and a corresponding pre-exponential factor ( $A_i$ ). A  $10 \times 10 \text{ mm}$  quartz cell was used for the absorption, steady-state, and time-resolved fluorescence measurements.

$$I(t) = I_0 \sum_i A_i e^{-t/\tau_i} \quad (1)$$

Binding dynamics studies were carried out with an Applied Photophysics SX20 stopped-flow system. The solutions were excited at 365 nm with a Hg-Xe vapor lamp. This wavelength was chosen because it corresponds to a peak of the Hg-Xe lamp and leads to a higher excitation efficiency of the sample, increasing the signal-to-noise ratio for the measurements. The excitation monochromator bandwidth was set to 2 nm. The monochromator wavelength was calibrated by comparing the wavelength reading for the maximum intensity reading for water with the wavelength for the maximum intensity provided by the manufacturer of



the lamp. The emission was detected using an interference filter (385–423 nm) with a maximum at 405 nm for the “blue” region and a 515 nm cut-off filter for the green region. The solutions were mixed in a 1 : 1 ratio in the mixing chamber. The temperature of the solutions was maintained at 20 °C throughout the experiment, and the samples were incubated at this temperature for 10 min before the start of an experiment. A minimum of 25 traces were averaged for each experiment performed on the stopped-flow. The intensity of the stopped-flow for a solution containing all chemicals except the fluorophore and CB[7] was taken as the baseline and subtracted from the stopped-flow traces to obtain the corrected traces.

The stopped-flow traces were analyzed by fitting the individual traces to a sum of exponential functions (eqn (2)) or by using a global analysis method where all traces are simultaneously fit to a defined model. The fit to a sum of exponentials is defined by an offset ( $a_0$ ) and the sum of exponentials terms, each of which has a corresponding observed rate constant ( $k_{\text{obs}i}$ ) and an amplitude of  $a_i$ .

$$\Delta I = a_0 + a_1 e^{-k_{\text{obs}1}t} + a_2 e^{-k_{\text{obs}2}t} \quad (2)$$

The analysis of the individual stopped-flow traces at pH 2.0 and 3.8 was done as follows: to obtain the rate constant for the slow relaxation time, the traces were fit to a mono-exponential function by starting the fit at incrementally longer times until the residuals became random and the observed rate constant was constant. The rate constant for the fast relaxation time was then obtained by fitting the traces to a sum of two exponentials and fixing the rate constant for the slow relaxation process. The stopped-flow traces at pH 5.0 and 5.5 were fit to a mono-exponential function to yield one relaxation time. In the global analysis method, all the kinetic traces for a particular experiment were fit simultaneously to a model defined in the Prokineticist II software from Applied Photophysics. The goodness of the fit was judged by the randomness of the residuals.

## Results

The  $\text{AH}^+/\text{A}$  absorption and emission spectra depend on the solution's pH. At pH 2.0, where only  $\text{AH}^+$  is present in the ground state, a small emission intensity is observed from  $\text{AH}^+$  around 400 nm in addition to the predominant emission centered at 510 nm from A formed adiabatically from excited  $\text{AH}^+$  (Fig. 1a). The emission from  $\text{AH}^+$  is absent at pH 6.0, which is two pH units higher than the  $\text{pK}_a$  of  $\text{AH}^+/\text{A}$ . At this pH, only A is present in solution. The absorption spectrum shows sharp peaks at pH 2.0 (Fig. 1b), whereas a broad absorption around 400 nm appears as the pH is raised (Fig. 1c and d). This broad absorbance is related to the presence of A.<sup>41</sup> In the presence of 25  $\mu\text{M}$  CB[7], where 91%  $\text{AH}^+$  (5  $\mu\text{M}$ ) is bound at pH 2.0, a red shift was observed in the absorption spectrum of  $\text{AH}^+$ . At pH 3.8, with an approximately 3 : 2 mixture of  $\text{AH}^+$  and A in water, the shoulder around 400 nm decreased in the presence of CB[7], and the sharp peaks shifted to the same wavelengths observed at pH 2.0 in the presence of this host. These absorption spectra show that the equilibrium shifts toward  $\text{AH}^+$  when the guest is bound to CB[7]. At pH 5.5, the amount of  $\text{AH}^+$  is low (3%), and the absorption spectrum showed smaller changes in the presence of CB[7]. These results are consistent with the stabilization of  $\text{AH}^+$  when bound to CB[7].



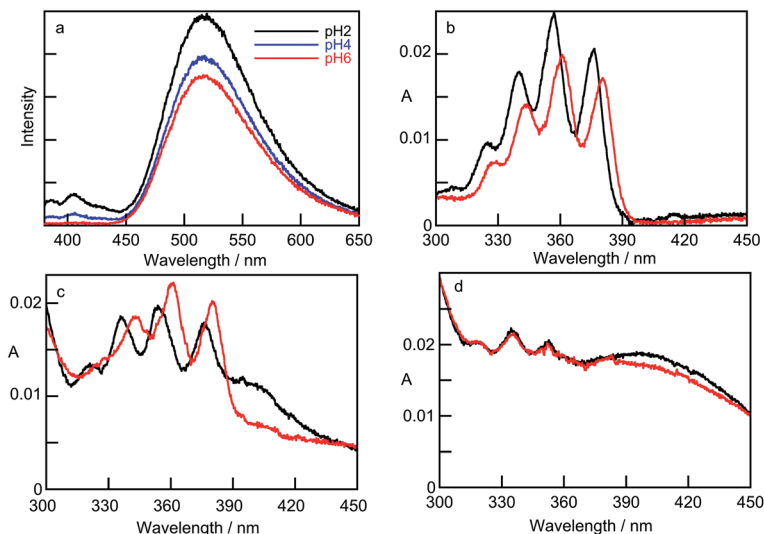


Fig. 1 (a) Emission spectra for AH<sup>+</sup>/A (1 μM) in water at different pH values: 2.0 (black), 4.0 (blue), and 6.0 (red). Absorption spectra of AH<sup>+</sup>/A (5 μM) at pH 2.0 (b), 3.8 (c), and 5.5 (d) in the absence (black) and presence of 25 μM CB[7] (red).

The binding of AH<sup>+</sup> with CB[7] was characterized at pH 2.0 where all of the guest molecules are protonated in water. Solubilization of CB[7] is enhanced in the presence of Na<sup>+</sup> because cations bind to the portals of CB[7].<sup>27,49–51</sup> The addition of Na<sup>+</sup> cations was also required to adjust the solution's pH. The system is described by the equilibria between Na<sup>+</sup> or AH<sup>+</sup> with CB[7] (Fig. 2a). The competitive binding of Na<sup>+</sup> to CB[7] was shown to slow the kinetics of AH<sup>+</sup> binding with CB[7] and to decrease the amplitude of the kinetics (Fig. 2b). This behavior is the same as that previously reported for the 2-naphthyl-1-ethylammonium cation binding to CB[7].<sup>32</sup> The smaller amplitude is due to the involvement of CB[7] in Na<sup>+</sup> bound complexes. The slow down of the relaxation kinetics is a consequence of a slower bimolecular association process because of the lower effective concentration of free CB[7] in the presence of Na<sup>+</sup> cations, while the dissociation is unaffected as it is a unimolecular reaction. Stopped-flow experiments were performed to determine the optimal Na<sup>+</sup> cation concentration for the kinetic studies. The kinetics were followed in the “blue” region where AH<sup>+</sup> emits because its excited state is not deprotonated owing to the higher pK<sub>a</sub><sup>\*</sup> of AH<sup>+</sup>@CB[7]. A concentration of 20 mM of Na<sup>+</sup> cations was chosen for all experiments because the kinetics was sufficiently slow to be detected in stopped-flow experiments with reasonable amplitude.

Any parameter that is dependent on the concentration of CB[7] is an overall or apparent parameter because a fraction of the CB[7] molecules was non-reactive owing to the formation of CB[7] complexes with Na<sup>+</sup> cations (Fig. 2a). The binding isotherms led to the determination of the overall binding constants ( $\beta$ ), while the bimolecular rate constants for the reactions involving CB[7] are apparent rate constants and denoted  $k'$ . The individual equilibrium constants ( $K$ ) or bimolecular rate constants ( $k$ ) can be obtained following the procedure previously



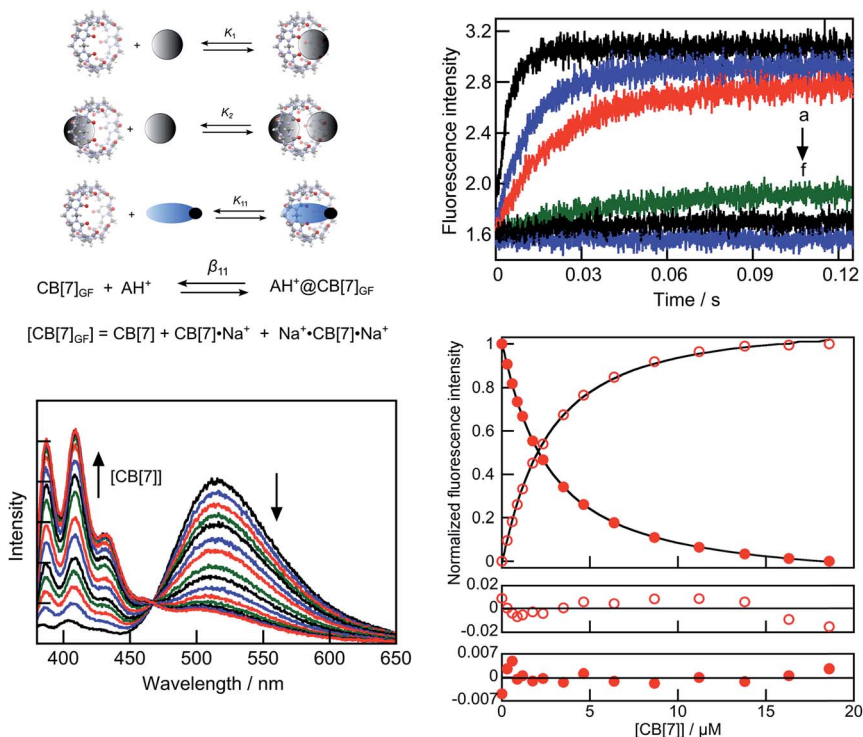


Fig. 2 Top left: Equilibria for CB[7] binding to Na<sup>+</sup> cations and AH<sup>+</sup>, and the definition of the overall equilibrium constant. Top right: Kinetics for the formation of the AH<sup>+</sup>@CB[7] complex ([AH<sup>+</sup>] = 2 μM, [CB[7]] = 7 μM) at pH 2.0 in the presence of increasing Na<sup>+</sup> cation concentrations: (a) 2, (b) 10, (c) 20, (d) 100, (e) 200 mM. Trace "f" corresponds to the baseline measurement in the absence of CB[7]. Bottom left: Fluorescence spectra for AH<sup>+</sup> at pH 2.0 in the presence of increasing CB[7] concentrations from 0 to 19 μM. Bottom right: Binding isotherms (top panel) for the intensity changes for the "blue" (integration from 380 to 456 nm, open circles) and "green" (integration from 456 to 650 nm, solid circles) emission. The black lines correspond to the numerical fits of the data. The residuals between the experimental data and calculated values are shown in the middle panel ("blue" emission) and lower panel ("green" emission).

described.<sup>32</sup> We chose to present the  $\beta$  and  $k'$  values in the results section as these are the values derived directly from the experiments.

The addition of increasing concentrations of CB[7] to AH<sup>+</sup> (1.0 μM) at pH 2.0 led to a decrease of the "green" emission of A around 510 nm and an increase of the AH<sup>+</sup> emission below 450 nm (Fig. 2c). The dependencies of the intensities in the "blue" and "green" regions with the CB[7] concentration (Fig. 2d) were numerically fit to an overall equilibrium constant ( $\beta_{11}$ , see ESI† for details) defined by the equations shown in Fig. 2a, where [CB[7]]<sub>GF</sub> corresponds to the guest-free CB[7] concentration that is not complexed to AH<sup>+</sup>. The residuals between the experimental data and the fits were random. The recovered average  $\beta_{11}$  values from two independent experiments were  $(4.92 \pm 0.09) \times 10^5 \text{ M}^{-1}$  when the emission intensity for A was followed and  $(4.8 \pm 0.2) \times 10^5 \text{ M}^{-1}$  when the intensity changes for AH<sup>+</sup> were measured, leading to an overall average  $\beta_{11}$  value





of  $(4.9 \pm 0.1) \times 10^5 \text{ M}^{-1}$ . The determination of the same  $\beta_{11}$  values for the two different emission bands and the presence of an isoemissive point support the assignment of the bands in the fluorescence spectra to  $\text{AH}^+$  and  $\text{AH}^+ @ \text{CB}[7]$ .

Binding isotherms were also measured at pH values of 3.8, 5.0, and 5.5 (Fig. S1 and Table S1 in the ESI†). The determined average  $\beta_{11}$  values from the binding isotherms measured for the “blue” and “green” emissions were  $(2.83 \pm 0.03) \times 10^5$ ,  $(3.49 \pm 0.09) \times 10^4$ , and  $(1.52 \pm 0.04) \times 10^4 \text{ M}^{-1}$  at pH values of 3.8, 5.0, and 5.5, respectively. At pH values of 3.8, 5.0, and 5.5, the percentage of  $\text{AH}^+$  is 61, 9, and 3%, while the percentage of A is 39, 91, and 97%, respectively. The decrease in the overall equilibrium constants as the pH was raised is a reflection of the lower equilibrium constant for the binding of A to CB[7] compared with that for the binding of  $\text{AH}^+$ . The equilibrium constant for the binding of A to CB[7] can be calculated from the thermodynamic cycle shown in Scheme 1 since three of the equilibrium constants are known. It is important to note that the thermodynamic cycle is valid for overall equilibrium constants where the value for  $\text{AH}^+$  ( $\beta_{11}^{\text{AH}}$ ) is the one determined at pH 2.0 and  $\beta_{11}^{\text{A}}$  is related to the other equilibrium constants (eqn (3), see ESI† for derivation), leading to a value for  $\beta_{11}^{\text{A}}$  of  $390 \pm 10 \text{ M}^{-1}$ , where the error is related to the measurement of  $\beta_{11}^{\text{AH}}$ . An attempt was made to measure directly the equilibrium constant between A and CB[7] at pH 12. The changes in the fluorescence intensity were small (Fig. S2 in the ESI†) and no saturation was achieved, which indicated the incomplete binding of A. The value of  $\beta_{11}^{\text{A}}$  and the quantum yield of A in  $\text{A} @ \text{CB}[7]$  are correlated, and for this reason, no unique value for  $\beta_{11}^{\text{A}}$  could be obtained. Adequate fits were observed for  $\beta_{11}^{\text{A}}$  values fixed between 100 and  $700 \text{ M}^{-1}$  (Fig. S3 in the ESI†), which are of the same order of magnitude as the value determined from the thermodynamic cycle.

$$\frac{10^{-pK_a^{\text{CB}}}}{10^{-pK_a}} = \frac{\beta_{11}^{\text{A}}}{\beta_{11}^{\text{AH}}} \quad (3)$$

Time-resolved fluorescence experiments are used to identify fluorophores with different lifetimes; in the case of supramolecular systems, the same fluorophore in different environments can have different lifetimes.<sup>8,52</sup> The decays for systems containing fluorophores with different lifetimes are fit to a sum of exponentials, where each term has an associated lifetime and pre-exponential factor  $A_i$  (eqn (1)). The  $A_i$  values are related to the abundance of each species; a positive value indicates the disappearance of the fluorophore, while a negative value indicates the formation of the fluorophore. The lifetime of excited A measured at 510 nm and pH 6.0 was  $24.8 \pm 0.1 \text{ ns}$ . This value is of the same order of magnitude as the lifetimes previously determined in water : ethanol (14 ns)<sup>43</sup> or cyclohexane (25–33 ns).<sup>53</sup> At pH 2.0, the kinetics at 510 nm for the emission of A showed a growth in kinetics with a lifetime of  $1.0 \pm 0.2 \text{ ns}$  followed by a decay with a lifetime of  $24.6 \pm 0.1 \text{ ns}$  (Fig. S4 and Table S2 in the ESI†). The growth corresponds to the adiabatic deprotonation of excited  $\text{AH}^+$  to form excited A, which decays with the same lifetime as excited A at the higher pH. This assignment at pH 2.0 is supported by the equal absolute values of the pre-exponential factors, which were  $-0.49 \pm 0.02$  and  $0.51 \pm 0.02$  for the short- and long-lived components, respectively. The equal pre-exponential factors indicated that all the excited states of A were formed from  $\text{AH}^+$ , as would be expected at a pH where the fluorophore is in the protonated form.





In the presence of 16  $\mu\text{M}$  CB[7] at pH 2.0, where all  $\text{AH}^+$  is bound, the fluorescence decay could only be measured at 410 nm because  $\text{AH}^+$  is stabilized in the complex and is the predominant species in the system. The decay was mono-exponential with a lifetime of  $7.6 \pm 0.1$  ns, which corresponds to the emission of excited state  $\text{AH}^+$  complexed with CB[7]. This lifetime is not limited by the deprotonation of  $\text{AH}^+@CB[7]$  because pH 2.0 is lower than both the  $\text{p}K_{\text{a}}^{\text{CB}}$  and  $\text{p}K_{\text{a}}^{\text{CB}^*}$  values. In the presence of 2.6  $\mu\text{M}$  CB[7], 50% of  $\text{AH}^+$  is bound to CB[7] and the remainder is free in water. For this reason, the fluorescence kinetics could be measured at 410 nm for the emission of  $\text{AH}^+$  and at 510 nm for the emission of excited A. The emission decay for  $\text{AH}^+$  at 410 nm led to the recovery of  $0.9 \pm 0.1$  and  $7.7 \pm 0.1$  ns lifetimes. The short lifetime corresponds to the deprotonation of  $\text{AH}^+$  in water, while the long lifetime corresponds to the emission of  $\text{AH}^+$  in the  $\text{AH}^+@CB[7]$  complex. At 510 nm, where excited A emits, a growth with a  $1.1 \pm 0.1$  ns lifetime was observed followed by a decay with a  $24.5 \pm 0.1$  ns lifetime. This kinetics is the same as that observed for  $\text{AH}^+$  in water, and the absence of a longer-lived growth with a lifetime close to 8 ns suggested that  $\text{AH}^+@CB[7]$  was not deprotonated during the excited state lifetime of  $\text{AH}^+$ . Therefore, the intensity changes at 510 nm are diagnostic for the changes in the  $\text{AH}^+$  concentration in water and do not have a contribution from the concentration changes for  $\text{AH}^+@CB[7]$ .

The longest lifetime observed for the  $\text{AH}^+/\text{A}$  system in the absence and presence of CB[7] is 25 ns, and the dynamics of the excited state occurs on a much faster time scale than the millisecond time scale for the formation of the  $\text{AH}^+@CB[7]$  complex (see below). Therefore, the changes in emission intensity can be seen as instantaneous when analyzing the kinetics of  $\text{AH}^+@CB[7]$  complex formation, and the excited state dynamics of  $\text{AH}^+$  is decoupled from the dynamics of CB[7] complex formation.

The kinetics for the formation of the  $\text{AH}^+@CB[7]$  complex was studied in stopped-flow experiments. Two solutions, one containing CB[7] and a second containing  $\text{AH}^+/\text{A}$ , were mixed in a 1 : 1 volume ratio. The concentrations stated are those for the final mixed solution. The kinetics was studied at pH 2.0, where only  $\text{AH}^+$  was present in water, and at pH 5.5, where A corresponds to 97% of the species present in water. A higher pH could not be used because the signals in the stopped-flow experiments became too small. The kinetics was also studied at intermediate pH values where a mixture of  $\text{AH}^+$  and A was present (pH 3.8: 61%  $\text{AH}^+$  and 39% A, pH 5.0: 9%  $\text{AH}^+$  and 91% A). At all pH values, mixing of the guest ( $\text{AH}^+/\text{A}$ ) with CB[7] led to a decrease of the emission intensity at 510 nm (Fig. 3a for pH 2.0 and Fig. S5 in the ESI†) and an increase of the emission intensity at 410 nm (Fig. S6 in the ESI†). It is important to note that at pH 2.0, the emission intensity at 510 nm corresponds to the concentration of  $\text{AH}^+$  in water, where excited A is formed from the deprotonation of excited  $\text{AH}^+$ , as  $\text{AH}^+$  in the CB[7] complex is not deprotonated. The intensity increase at 410 nm corresponds to the formation of  $\text{AH}^+@CB[7]$ . At pH 5.5, the emission intensity at 410 nm corresponds to the concentration of  $\text{AH}^+@CB[7]$  because, at this pH,  $\text{AH}^+$  in water deprotonates readily. The intensity at 510 nm corresponds to the sum of the intensities of A in water and  $\text{A}@CB[7]$ . At the intermediate pH values of 3.8 and 5.0, the intensity at 510 nm corresponds to the concentrations of A in water,  $\text{A}@CB[7]$ , and  $\text{AH}^+$  in water that forms excited A adiabatically. At all pH values, the same observed rate constants were recovered from the kinetics measured at 410 and



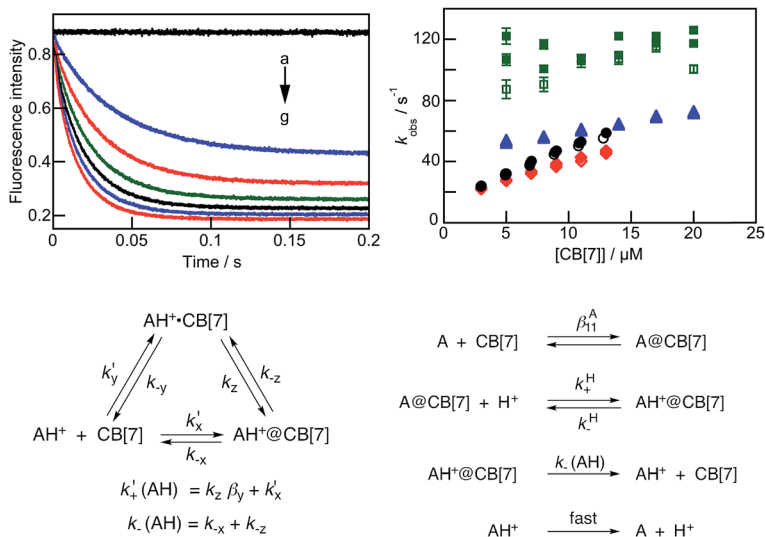


Fig. 3 Top left: Kinetic traces at pH 2.0 for the mixing of  $\text{AH}^+$  ( $1 \mu\text{M}$ ) with  $\text{CB}[7]$  ((a) 0, (b) 3, (c) 5, (d) 7, (e) 9, (f) 11, and (g)  $13 \mu\text{M}$ ) measured for the “green” emission. Top right: Dependence of the observed rate constant with the  $\text{CB}[7]$  concentration at different pH values (pH 2.0:  $\circ$ ,  $\bullet$ , black; pH 3.8:  $\diamond$ ,  $\blacklozenge$ , red; pH 5.0:  $\triangle$ ,  $\blacktriangle$ , blue; and pH 5.5:  $\square$ ,  $\blacksquare$ , green). The solid and open symbols are the values recovered for the kinetics measured for the “green” and “blue” emission, respectively. For pH values where the open symbols are not shown, they are the same as the closed symbols. The error bars are smaller than the symbols for all pH values, with the exception of pH 5.5. The observed rate constants for pH 2.0 and 3.8 correspond to the lowest values recovered from a fit of the kinetics to the sum of two exponentials. The kinetics for pH 5.0 and 5.5 were fit to a mono-exponential function. Bottom left: Mechanism used to analyze the kinetic data at pH 2.0. Bottom right: Mechanism used to analyze the kinetic data at pH 5.5.

510 nm, indicating that the kinetics are coupled, as would be expected for the relaxation kinetics of a system where the various species are in equilibrium.

Two control experiments were performed: (i) comparison of the amplitudes for the kinetic and binding isotherm measurements and (ii) kinetic measurements with buffered and unbuffered solutions.

(i) It is important to establish whether the kinetics are measured for a sufficiently long time for the system to reach equilibrium. The normalized amplitudes from the stopped-flow experiments at 0.2 s were the same as the amplitudes from the binding isotherm experiments at all pH values (Fig. S7 and S8 in the ESI†). This result shows that the system is equilibrated within 0.2 s. Kinetic processes faster than the 1 ms mixing time of the stopped-flow experiment appear as initial offsets in the kinetic traces. In the current system, such a fast process would involve  $\text{CB}[7]$ , and the amplitude of the offset would increase as the host concentration was raised. The kinetics at all pH values did not show a progressively increasing offset, indicating the absence of a relaxation process that occurred faster than 1 ms (Fig. S5 and S6 in the ESI†). Therefore, the whole kinetics of the system is captured in the stopped-flow experiments.

(ii) Experiments at pH 2.0 and 3.8 were performed in unbuffered solutions, where the pH was adjusted with the addition of HCl in the presence of 20 mM



NaCl, while measurements at pH 5.0 and 5.5 required the use of sodium acetate as a buffer ( $[\text{Na}^+] = 20 \text{ mM}$ ). A control experiment was performed at pH 4.3, where samples were prepared in the presence of HCl/NaCl or acetate buffer, and the kinetics for complex formation was measured. The dependence of the observed rate constants with CB[7] was similar, with a slightly lower slope observed for the experiments performed in acetate buffer (Fig. S9 in the ESI†). This difference is very similar to the variation observed between independent experiments, but it could also reflect the weak binding of acetic acid to CB[7]. However, this control experiment showed that the large differences in the dependencies of the observed rate constants with the CB[7] concentration (Fig. 3b) are not due to a counter-ion effect.

The kinetic behavior is different at the different pH values studied (Fig. 3b). A qualitative description will be provided first to guide the reader through the detailed analysis. The kinetics at pH 2.0 and 3.8 were fit to the sum of two exponentials, while the kinetics at pH 5.0 and 5.5 were mono-exponential. The relaxation kinetics for the lowest observed rate constant at pH 2.0 and 3.8 was slower than the relaxation kinetics at pH 5.5, and a steeper dependence of the observed rate constant with the CB[7] concentration was observed at the lower pH values. The errors for the measured rate constants are higher at pH 5.5 because the signal-to-noise ratio was lower, reflecting the smaller amount of CB[7] complex formed.

The slower kinetics observed at pH 2.0 is consistent with a mechanism where an exclusion complex is formed, denoted  $\text{AH}^+ \cdot \text{CB}[7]$ , in which the positively charged guest interacts with the carbonyl groups at the portal of CB[7] without the inclusion of the anthracene moiety into the CB[7] cavity in addition to a pathway where the anthracene moiety is included directly. Inclusion of the anthracene moiety to form  $\text{AH}^+ @ \text{CB}[7]$  occurs either from the exclusion complex (pathway “z”, Fig. 3c) or directly (pathway “x”).

At pH 5.5, deprotonated A is the predominant species in water (97%), but the hydronium ion concentration is sufficiently high to protonate the  $\text{A} @ \text{CB}[7]$  complex and form  $\text{AH}^+ @ \text{CB}[7]$ . The mechanism includes the fast equilibration between A and CB[7] followed by slow protonation and deprotonation steps (Fig. 3d). Deprotonation of  $\text{AH}^+ @ \text{CB}[7]$  and the exit of  $\text{AH}^+$  from  $\text{AH}^+ @ \text{CB}[7]$  are competitive, and the latter reaction needs to be accounted for in the fitting of the data.

The kinetics at pH 2.0 was fit to the sum of two exponentials, from which two observed rate constants,  $k_{\text{obs1}}$  and  $k_{\text{obs2}}$ , were obtained. Similar dependencies were observed for the  $k_{\text{obs2}}$  values with the CB[7] concentration when the kinetics were measured for the “blue” and “green” emission intensity changes. The values for  $k_{\text{obs1}}$  were scattered, and this pattern is a reflection of the small amplitude of this component at low CB[7] concentrations. To improve the precision of the fits, the kinetics was fit to a mono-exponential function by starting the fit at incrementally longer delays after the start of the reaction until a constant value for  $k_{\text{obs2}}$  was obtained and the residuals were random (Fig. S10 and Table S3, see ESI† for details). These  $k_{\text{obs2}}$  values were then fixed for the fit of the entire kinetic trace to recover the values for  $k_{\text{obs1}}$ .

In principle, the two relaxation times could correspond to two different processes. If  $k_{\text{obs2}}$  was related to the reaction of  $\text{AH}^+$  with CB[7] without the formation of an encounter complex, then the ratio of the slope of a linear



dependence between  $k_{\text{obs}2}$  and the CB[7] concentration ( $(3.4 \pm 0.2) \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ , average from two kinetic studies with the emission collected in the green region and one in the blue region) and the intercept ( $14.6 \pm 0.8 \text{ s}^{-1}$ ) should correspond to  $\beta_{11}^{\text{AH}}$ . However, this ratio ( $(2.3 \pm 0.2) \times 10^5 \text{ M}^{-1}$ ) is much lower than the  $\beta_{11}^{\text{AH}}$  value of  $(4.9 \pm 0.1) \times 10^5 \text{ M}^{-1}$  determined from the binding isotherm studies. This analysis showed that the kinetics for the two relaxation processes are coupled.

Based on the precedence for the formation of exclusion complexes followed by cavity inclusion with CB[n]s as the host,<sup>27,30</sup> the kinetics was analyzed using the mechanism shown in Fig. 3c. The formation of  $\text{AH}^+ \cdot \text{CB}[7]$  was assumed to be in fast equilibrium. With this assumption, the two relaxation processes are related to the three equilibria (eqn (4) and (5)), where  $k'_+(\text{AH})$  and  $k_-(\text{AH})$  are defined in Fig. 3c:<sup>54</sup>

$$k_{\text{obs}1} = k'_y[\text{CB}[7]] + k_{-y} \quad (4)$$

$$k_{\text{obs}2} = \frac{k'_+(\text{AH})[\text{CB}[7]]}{1 + \beta_y[\text{CB}[7]]} + k_-(\text{AH}) \quad (5)$$

Fits of the dependence of  $k_{\text{obs}2}$  with the CB[7] concentration to eqn (5) (see Fig. S11 and Table S4 in the ESI† for individual values) led to an average  $\beta_y$  value of  $(2.6 \pm 0.6) \times 10^4 \text{ M}^{-1}$ , a  $k'_+(\text{AH})$  value of  $(4.9 \pm 0.4) \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ , and a  $k_-(\text{AH})$  value of  $10 \pm 1 \text{ s}^{-1}$ . The ratio between  $k'_+(\text{AH})$  and  $k_-(\text{AH})$  is  $(4.9 \pm 0.6) \times 10^5 \text{ M}^{-1}$ , which is equal to the  $\beta_{11}^{\text{AH}}$  value ( $(4.9 \pm 0.1) \times 10^5 \text{ M}^{-1}$ ) determined from the binding isotherm. The equality of these values suggests that the mechanism proposed is consistent with the kinetic and the binding isotherm data.

The  $k_{\text{obs}1}$  values increased with the CB[7] concentration; however, the data had large errors and showed significant scatter, indicating that fitting of the data was not warranted (Fig. S12 in the ESI†). The value of  $k_{-y}$  is estimated to be between 100 and  $130 \text{ s}^{-1}$ , which is ten times higher than the rate constant for the exit of  $\text{AH}^+$  from the inclusion complex ( $k_-(\text{AH}) = 10 \text{ s}^{-1}$ ), supporting the assumption that the formation of the exclusion complex occurs as a fast equilibrium. The value for  $k_y$  estimated from the  $\beta_y$  and  $k_{-y}$  values is between  $2.7 \times 10^6$  and  $3.5 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ , which would lead to an increment for the  $k_{\text{obs}1}$  value of *ca.*  $30 \text{ s}^{-1}$  for a  $10 \text{ }\mu\text{M}$  increase in the CB[7] concentration. This increment is within the scatter observed for the experimental data.

The data at pH 5.5 were analyzed using a global analysis method (Scheme S1 in the ESI†), where the kinetics at all CB[7] concentrations for two independent experiments collected for the “green” emission were analyzed simultaneously. The data from the kinetics in the “blue” region were not used because of the poor signal-to-noise ratio.

The value for the equilibrium constant for  $\text{A}@\text{CB}[7]$  was fixed as  $390 \text{ M}^{-1}$ , and this equilibrium was assumed to be fast. Assuming that the association rate constant of A with CB[7] will be at least as high as the overall association rate constant for  $\text{AH}^+$  of  $4.9 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$ , then the dissociation rate constant for A from  $\text{A}@\text{CB}[7]$  will be at least  $1.3 \times 10^4 \text{ s}^{-1}$ , which is ten times faster than the time resolution of the stopped-flow experiment. This calculation is consistent with the assumption of a fast equilibrium for  $\text{A}@\text{CB}[7]$ .



The dissociation of  $\text{AH}^+\text{@CB}[7]$  needs to be included in the model because the value for the rate constant of this process ( $10 \text{ s}^{-1}$ ) corresponds to 10% of the observed rate constant. This value was also fixed in the model. The protonation and deprotonation reaction for  $\text{A/AH}^+$  in water is faster than the time resolution of the experiment, and the concentration of the hydronium ion is constant as the solution was buffered. For this reason, the deprotonation reaction for  $\text{AH}^+$  in water is not included in the model used for fitting the kinetics.

Global analysis of the kinetics at pH 5.5 (see Fig. S13 in the ESI† for the residuals) led to a recovery of the protonation rate constant ( $k^{\text{H}}$ ) of  $(1.5 \pm 0.4) \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$  for  $\text{A@CB}[7]$  and a deprotonation rate constant ( $k^{\text{H}}$ ) of  $89 \pm 7 \text{ s}^{-1}$  for  $\text{AH}^+\text{@CB}[7]$ . The  $\text{p}K_{\text{a}}^{\text{CB}}$  value calculated from these rate constants is  $7.3 \pm 0.2$ , which is in agreement with the reported value of  $7.1 \pm 0.2$  determined from a pH titration experiment.<sup>38</sup>

A quantitative analysis of the kinetics at pH 3.8 and 5 is not feasible because the model requires the inclusion of both the low and high pH mechanisms observed for the binding of  $\text{AH}^+$  and A; the model would include too many parameters to fit the data. However, the changes can be explained qualitatively based on the relative contributions from the formation of the exclusion complex  $\text{AH}^+\text{·CB}[7]$  and the complex with neutral A,  $\text{A@CB}[7]$ . The overall equilibrium constant for the formation of the former is 69 times higher than that for the latter. At pH 3.8, where 61% of the guest in water is in the protonated form and the remaining 39% is deprotonated, the kinetics is dominated by the binding of  $\text{AH}^+$  with  $\text{CB}[7]$  and the observed decay did not fit to a mono-exponential function. However, the amplitude of the fast component is smaller at pH 3.8 than at pH 2 (Table S5 in the ESI†). This difference increased from 11% for a  $\text{CB}[7]$  concentration of  $5 \text{ }\mu\text{M}$  to 32% for a  $\text{CB}[7]$  concentration of  $13 \text{ }\mu\text{M}$ . This increase is expected because the formation of  $\text{A@CB}[7]$  will be more prominent at a higher  $\text{CB}[7]$  concentration. The formation of  $\text{A@CB}[7]$  is followed by immediate protonation of the complex ( $\sim 2.4 \times 10^5 \text{ s}^{-1}$ ) because of the high hydronium ion concentration at this pH. This reaction path leads to a decrease in the contribution from  $\text{AH}^+\text{@CB}[7]$  formation through the exclusion complex. This effect is observed as a decrease in the slope for the dependence of  $k_{\text{obs}2}$  with the  $\text{CB}[7]$  concentration without a large change in the intercept.

The kinetics at pH 5 was adequately fit to a mono-exponential function. In water, 9% of the guest is in the form of  $\text{AH}^+$  while 91% corresponds to A. The kinetics is dominated by the binding of A, but a slow down was observed for the kinetics when compared with the kinetics at pH 5.5. This slow down was caused by the formation of the  $\text{AH}^+\text{·CB}[7]$  exclusion complex that removes A and free  $\text{CB}[7]$  from solution. The lower free concentrations of A and  $\text{CB}[7]$  decrease the amount of  $\text{AH}^+\text{@CB}[7]$  formed through the faster reaction pathway, which is the protonation of the  $\text{A@CB}[7]$  complex.

## Discussion

The guest binding dynamics of  $\text{AH}^+$  with  $\text{CB}[7]$  can be compared with the dynamics of previously studied guests. 2-Naphthyl-1-ethylammonium (Scheme 2) has a positive charge located in one extreme of the molecule in a similar fashion to  $\text{AH}^+$ , whereas the positive charge in berberine is centrally located. In both



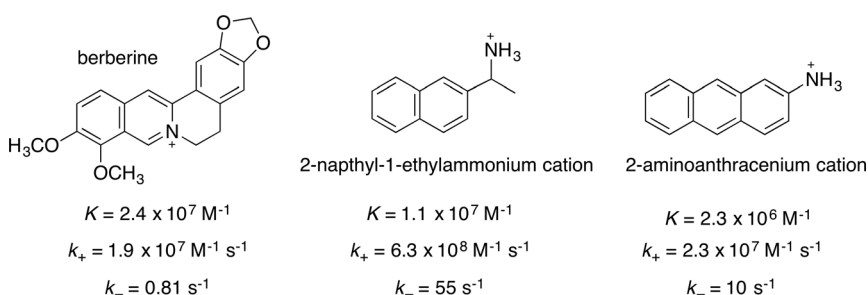
cases, the kinetics was associated to one relaxation process without the observing the formation of an exclusion complex.

The equilibrium constant  $K_{\text{AH}}$  was calculated to be  $(2.3 \pm 0.3) \times 10^6 \text{ M}^{-1}$  (eqn (6)) using the previously determined values of  $130 \pm 10 \text{ M}^{-1}$  and  $21 \pm 2 \text{ M}^{-1}$  for the binding of the first and second  $\text{Na}^+$  cation to CB[7],<sup>32</sup> and the average  $\beta_{11}$  value of  $(4.9 \pm 0.1) \times 10^5 \text{ M}^{-1}$ . The same multiplication factor was used to calculate the value of the association rate constant ( $k_+(\text{AH})$ ) as  $(2.3 \pm 0.2) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$  from the overall association rate constant  $k'_+(\text{AH})$ . It is important to note that the latter value includes the pathway for the formation of the exclusion complex.

$$K_{\text{AH}} = \beta_{11}(1 + K_{01}[\text{Na}] + K_{01}K_{02}[\text{Na}]^2) \quad (6)$$

A comparison of the binding of these three guests to CB[7] shows that the location of the positive charge and the size of the hydrophobic moiety of the guest influence the binding dynamics. The two modes of association of  $\text{AH}^+$  with CB[7], *i.e.* direct inclusion and formation of the exclusion complex, are probably related to the directionality of the approach of the guest with respect to the portal of CB[7]. Direct inclusion likely occurs when the anthracene moiety approaches the portal, whereas the exclusion complex is formed when the positive charge on the amino group interacts first with the carbonyl groups on the rim of the portal. In the case of the 2-naphthyl-1-ethylammonium cation, the guest is sufficiently small that if the exclusion complex is formed, the guest can rotate and enter the cavity. In this case, the rate constant of inclusion ( $k_z$  in Fig. 3c) is higher than the dissociation of the exclusion complex ( $k_{-y}$ ), the association rate constant is high, and the exclusion complex is not detected as a defined intermediate. The association rate constant for  $\text{AH}^+$  is a factor of  $\sim 25$  lower than that of the 2-naphthyl-1-ethylammonium cation, which accounts for the formation of an exclusion complex where the dissociation of the exclusion complex is competitive with inclusion. The dissociation rate constant is a factor of  $\sim 5$  lower for  $\text{AH}^+$  than for the 2-naphthyl-1-ethylammonium cation. This slow-down is probably related to the larger hydrophobic moiety of  $\text{AH}^+$ .

The aromatic moiety of berberine is included in the CB[7] cavity,<sup>34</sup> and the central location of the charge likely precludes the formation of an exclusion



**Scheme 2** Structure, equilibrium constants and association and dissociation rate constants for the binding of berberine,<sup>34</sup> the 2-naphthyl-1-ethylammonium cation<sup>32</sup> and  $\text{AH}^+$  with CB[7]. The values of  $K$  and  $k_+$  for  $\text{AH}^+ @ \text{CB}[7]$  were calculated from the overall values (see text).



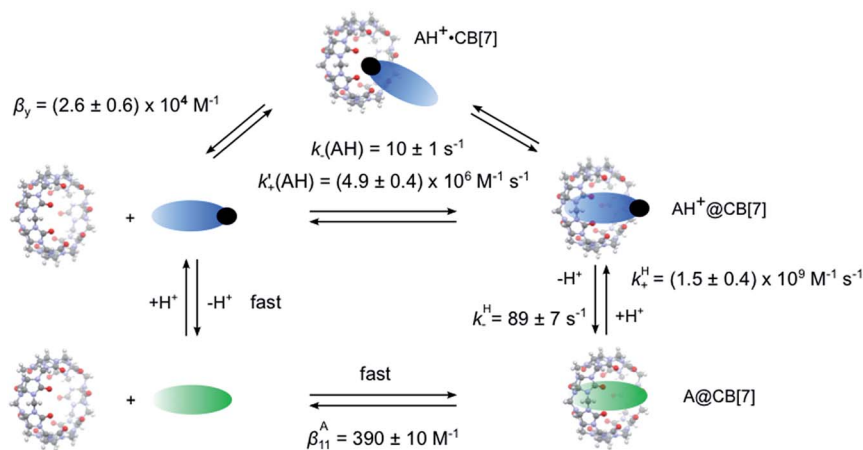
complex. The complexation of berberine to CB[7] was shown to occur through constrictive binding, where the CB[7] needs to be distorted for inclusion of the guest to occur. This stereochemical effect led to lower association and dissociation rate constants. The association rate constant for berberine and  $\text{AH}^+$  are the same, but different mechanisms led to these values being lower than the association rate constant for the 2-naphthyl-1-ethylammonium cation. Distortion of the host is also responsible for the lower dissociation rate constant observed for berberine compared to those of  $\text{AH}^+$  and the 2-naphthyl-1-ethylammonium cation. These results show that the binding dynamics cannot be predicted from the values of the equilibrium constants because the mechanism for binding differs for the three guests. Berberine has a similar equilibrium constant to the 2-naphthyl-1-ethylammonium cation, but the association and dissociation rate constants of berberine are decreased to a similar extent because of the constricted binding. On the other hand,  $\text{AH}^+$  has a lower equilibrium constant because of the slow down of the association process where the exclusion complex is formed, but the dissociation rate constant is affected to a lesser extent because no constricted binding is operating. It is important to note that in the case of cyclohexylmethylamine binding to the smaller CB[6], the formation of an exclusion complex and constrictive binding were postulated, leading to very slow kinetics,<sup>30,31</sup> showing that the binding dynamics with CB[n]s can occur over very different time scales.

The role of the protonation of the guest was previously studied for the cyclohexylmethylamine/CB[6] system, where the binding dynamics of the neutral and positively charged guest was uncoupled from the protonation/deprotonation dynamics.<sup>30,31</sup> In the case of  $\text{AH}^+$  and A binding to CB[7], the binding dynamics is coupled to the protonation/deprotonation reactions for the CB[7]-bound guest. In both systems, the dynamics is faster for the formation of the CB[n] complex with the neutral guest. However, in the case of  $\text{AH}^+$ /A binding to CB[7], it was possible to measure the kinetics of the system at different pH values. These studies led to the determination of the different mechanisms, where an exclusion complex is formed when  $\text{AH}^+$  binds to CB[7], whereas for A, complexation is fast and is followed by protonation of the complex (Scheme 3).

The protonation rate constant for ammonia in water is  $4.3 \times 10^{10} \text{ M}^{-1} \text{ s}^{-1}$ ,<sup>55</sup> and this value constitutes the highest possible rate constant for the protonation of A in water, which is unknown. The protonation rate constant for  $\text{A@CB[7]}$  of  $1.5 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$  is  $\sim 30$  times lower than this upper limit, suggesting that protonation of  $\text{A@CB[7]}$  is not significantly impeded, which is consistent with the location of the amino group at the portal of CB[7]. The lower value for the protonation rate constant for  $\text{A@CB[7]}$  is probably related to fact that the approach of the hydronium ion towards the sides of CB[7] or the portal that does not contain the amino group leads to unproductive encounter complexes. In this respect, the protonation rate constant for a guest where the acid/base group was exposed to the water phase while a portion of the guest was bound to CB[7] was the same as for the guest in water,<sup>56</sup> probably because the attack of the hydronium ion was not impeded. The deprotonation rate constant of  $\text{AH}^+\text{@CB[7]}$  is higher ( $89 \text{ s}^{-1}$ ) than the rate constant for the exit of  $\text{AH}^+$  from  $\text{AH}^+\text{@CB[7]}$  ( $10 \text{ s}^{-1}$ ). However, the deprotonation reaction is not observable at low pH values because the protonation process is too fast. For example, at pH 2.0, the protonation process for  $\text{A@CB[7]}$  has a pseudo-first order rate constant of  $1.5 \times 10^7 \text{ s}^{-1}$ .







**Scheme 3** Binding dynamics of  $\text{AH}^+$  and  $\text{A}$  with  $\text{CB}[7]$ , and rate constants for the protonation of  $\text{A@CB}[7]$  and deprotonation of  $\text{AH}^+\text{@CB}[7]$ . The kinetics for the binding of  $\text{AH}^+$  occurs through two pathways, and the association and dissociation rate constants are those for the combined pathways.

In conclusion, the studies on the binding dynamics of  $\text{AH}^+$  and  $\text{A}$  with  $\text{CB}[7]$  provided further evidence that different mechanisms occur for the binding of guests with  $\text{CB}[n]$ . The binding dynamics of a neutral guest was shown to be faster than that for the corresponding positively charged guest. However, the binding constant is higher for the charged guest than for the neutral guest, mainly because of the charge-dipole interaction between the guest and the carbonyl groups at the portal of  $\text{CB}[n]$ . The size of the guest and the position of the positive charge on the guest affect the type of mechanism for the binding dynamics and affect the magnitude of the association and dissociation rate constants. Positive charges located at one end of the guest are more likely to lead to the formation of exclusion complexes when compared to molecules with centrally located positive charges. In addition, the requirement for distortion of  $\text{CB}[n]$  to accommodate the guest can lead to a significant slow down of the dynamics, but has a much smaller effect on the equilibrium constant as the required distortions of the host will occur for both the association and dissociation processes.

The coupling of the binding dynamics of  $\text{A@CB}[7]$  and  $\text{AH}^+\text{@CB}[7]$  to the deprotonation and protonation reactions of these complexes made it possible to measure the protonation and deprotonation rate constants. The determination of these rate constants showed that the  $\text{pK}_{\text{a}}$  shift observed for the  $\text{CB}[7]$ -bound guest was due mainly to a decrease in the deprotonation rate constant because of the stabilization of the charged species. This result has direct implications for the use of  $\text{CB}[n]$ s in any application, such as in catalysis, where the lifetime of a charged species is important.

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