



Cite this: *Phys. Chem. Chem. Phys.*,
2019, 21, 2706

Microhydration of protonated 5-hydroxyindole revealed by infrared spectroscopy†

Johanna Klyne  and Otto Dopfer *

Controlled microsolvation of protonated aromatic biomolecules with water is fundamental to understand proton transfer reactions in aqueous environments. We measured infrared photodissociation (IRPD) spectra of mass-selected microhydrates of protonated 5-hydroxyindole ($5\text{HIH}^+ - W_n$, $W = \text{H}_2\text{O}$, $n = 1-3$) in the OH and NH stretch ranges ($2700-3800\text{ cm}^{-1}$), which are sensitive to the spectroscopic characteristics of interior solvation, water network formation, and proton transfer to solvent. Analysis of the IRPD spectra by dispersion-corrected density functional theory calculations (B3LYP-D3/aug-cc-pVTZ) reveals the coexistence of C3- and C4-protonated carbenium ions, $5\text{HIH}^+(\text{C3})$ and $5\text{HIH}^+(\text{C4})$, as well as the O-protonated oxonium ion, $5\text{HIH}^+(\text{O})$. Monohydrated $5\text{HIH}^+ - W$ clusters are formed by hydrogen-bonding (H-bonding) of the first water to the most acidic functional group, namely, the NH group in the case of $5\text{HIH}^+(\text{C3})$, the OH group for $5\text{HIH}^+(\text{C4})$, and the OH_2 group for $5\text{HIH}^+(\text{O})$. The latter benefits from its twofold degeneracy and the outstandingly high binding energy of $D_0 \sim 100\text{ kJ mol}^{-1}$. Larger $5\text{HIH}^+ - W_{2/3}$ clusters preferably grow (i) by H-bonding of the second water to the remaining vacant functional group and and/or (ii) by formation of W_2 water chains at the respective most acidic functional group. Our IRPD spectra of $5\text{HIH}^+ - W_n$ do not indicate any proton transfer to the solvent up to $n = 3$, in line with the proton affinities of 5HI and W_n . Comparison of $5\text{HIH}^+ - W_n$ to neutral $5\text{HI} - W$ and cationic $5\text{HI}^+ - W_n$ clusters elucidates the impact of different charge states on the topology of the initial solvation shell. Furthermore, to access the influence of the size of the arene ion and a second functional group, we draw a comparison to microhydration of protonated phenol.

Received 8th November 2018,
Accepted 7th January 2019

DOI: 10.1039/c8cp06950f

rsc.li/pccp

1. Introduction

The interaction of biomolecules, such as proteins, enzymes or hormones, with water (W) is crucial for their structure and function in living organisms. Controlled sequential microhydration of isolated biomolecules facilitates investigation of the interactions between solutes and solvents, which are blurred in the condensed phase. Infrared (IR) vibrational spectroscopy of size-selected hydrated clusters is particularly useful because it provides direct structural information. Combined with quantum chemistry, IR spectroscopy can elucidate the potential energy surface of hydration interactions.

Aromatic heterocyclic molecules are ubiquitous biomolecular building blocks.¹⁻³ For example, 5-hydroxyindole (5HI), the prototype chosen herein, consists of pyrrole fused to a phenol ring and occurs as a subunit in the neurotransmitter serotonin. The sequential microhydration of protonated aromatic molecules is particularly interesting because proton transfer from the

biomolecule to the solvent may happen in these systems. Both, protonation and hydration are found to enhance the acidity of the functional groups (OH/NH), and hence their ability to donate a proton.^{4,5} Moreover, the proton affinity (PA) of W (PA = 691 kJ mol^{-1})⁶ and small W_n clusters (PA = $808-879\text{ kJ mol}^{-1}$ for $n = 2-3$)^{7,8} is in the same range as those of aromatic hydrocarbons (*e.g.*, PA = 750 kJ mol^{-1} for benzene).⁶ Hence, formation of a hydronium ion (H_3O^+) or protonated water clusters $\text{H}^+(\text{H}_2\text{O})_n$ *via* proton transfer can eventually increase the interaction energy resulting in more stable hydrated clusters. For example, protonated benzene releases its excess proton already upon hydration by a single water molecule.^{9,10} Moreover, for the prototypical protonated arenes (A) naphthalene and benzaldehyde, proton transfer to water occurs at cluster sizes of $n = 2$ and 3, respectively, as revealed from IR photodissociation (IRPD) spectra of their microhydrated $[\text{A} - W_n]\text{H}^+$ clusters.^{11,12} In general, the topology of the potential energy surface of hydrated clusters strongly depends on their charge or protonation state.^{5,13-15}

The microhydration of the protonated phenol (PhH^+) subunit of 5HI has been studied by quantum chemistry and IRPD spectroscopy.^{14,16} While the theoretical study focused on the electronic structure of the carbenium ion,¹⁶ the IRPD study indicates the coexistence of *ortho/para* C-protonated carbenium

Institut für Optik und Atomare Physik, Technische Universität Berlin,
Hardenbergstr. 36, 10623 Berlin, Germany. E-mail: dopfer@physik.tu-berlin.de;
Fax: +49-30-31423018

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c8cp06950f



and O-protonated oxonium isomers, $\text{PhH}^+(o/p)$ and $\text{PhH}^+(\text{O})$, in a molecular beam.¹⁴ For both types of protomers, small W_n clusters are formed at the OH group of phenol. However, proton transfer is observed at different critical sizes of the hydration shell (n_c). While the W_n network accepts the excess proton already at $n_c = 3$ for $\text{PhH}^+(\text{O})-W_n$, proton transfer is only possible for $n \geq 4$ in $\text{PhH}^+(o/p)-W_n$. This change in n_c is explained by the geometry of the carbenium cluster. The protonated CH_2 group is simply too far away from the W_n network at the OH group. Proton transfer is possible only when the W_n cluster forms a ring bridging the OH and CH_2 groups at $n_c = 4$. The role of binding energies (D_0) of the PhH^+-W_n clusters or differences in the acidity of the OH groups have not been discussed. However, our preceding IRPD study of the sequential microsolvation of protonated 5-hydroxyindole (5HIH^+) by non-polar Ar and quadrupolar N_2 ligands (L) indicates a drastic difference in the ligand binding energies of the carbenium and oxonium protomers, $5\text{HIH}^+(\text{C})$ and $5\text{HIH}^+(\text{O})$.¹⁷ The 5HIH^+-L_n spectra ($n \leq 3$) reveal the coexistence of $5\text{HIH}^+(\text{C3})-L_n$, $5\text{HIH}^+(\text{C4})-L_n$, and $5\text{HIH}^+(\text{O})-L_n$ clusters with ion cores protonated at C3, C4, and O, respectively. This finding is surprising at first glance, because bare $5\text{HIH}^+(\text{O})$ is drastically less stable than the carbenium ions ($\Delta E_0 > 100 \text{ kJ mol}^{-1}$).^{5,17} The presence of clusters with an oxonium core can however be rationalized by the outstandingly large binding energies of $5\text{HIH}^+(\text{O})-L_n(\text{OH})$ clusters with ligands attached to the OH_2 group, the high barriers for isomerization from the oxonium to the carbenium ions (kinetic trapping), and the twofold degeneracy of these structures.^{14,17-20} Furthermore, the acidity of the NH and OH functional groups is found to strongly depend on the protonation site.¹⁷ Comparison with PhH^+ indicates an increase in the acidity of the OH group from 5HIH^+ to PhH^+ .¹⁷ Hence, we expect intracluster proton transfer in 5HIH^+-W_n clusters for a larger n_c value as compared to PhH^+-W_n .

To the best of our knowledge, only one theoretical study describes the microhydration of protonated 5HIH^+ ,⁵ and the protonation-induced change of the interaction potential compared to neutral hydrated clusters. This study reveals the preference of $\text{OH} \cdots \text{W}$ over $\text{NH} \cdots \text{W}$ H-bonds in the neutral ground state, in line with spectroscopic data.¹⁴ Both, $\text{OH} \cdots \text{W}$ and $\text{NH} \cdots \text{W}$ H-bonds are significantly strengthened upon protonation,⁵ whereas the increase in the acidity of the NH group is more pronounced than that of the OH group. As a result, the topology of the hydration shell is changed upon protonation and $\text{NH} \cdots \text{W}$ bonds are preferred. However, only C3-protonated $5\text{HIH}^+(\text{C3})-W$ clusters have been considered.⁵ Most likely, the topologies of $5\text{HIH}^+(\text{C4})-W$ and $5\text{HIH}^+(\text{O})-W$ clusters differ strongly.¹⁷ Thus, we address herein three main questions: (i) what is the protomer abundance within 5HIH^+-W_n clusters and how is it changed compared to 5HIH^+-L_n with $L = \text{Ar}$ and N_2 ; (ii) what is the structure of the initial solvation shell in 5HIH^+-W_n ; and (iii) do we observe proton transfer in the size range $n \leq 3$ and a protomer dependence of n_c ?

To this end, we analyze IRPD spectra of mass-selected hydrated clusters of 5HIH^+ with the aid of quantum chemical calculations. Vibrational spectroscopy in the NH and OH stretch range can determine the protonation site, discriminate interior ion solvation from water network formation, and signal

potential proton transfer from 5HIH^+ to the W_n solvent cluster. The comparison of our results on 5HIH^+-W to those obtained for 5HI^+-W and $5\text{HI}-W$ illustrates the influence of different charge states and protonation on the solvation of 5HI .

2. Experimental and computational techniques

The microsolvation of 5HIH^+ is studied by IRPD spectroscopy of mass-selected 5HIH^+-W_n ($n = 1-3$) clusters. Additional spectra of colder 5HIH^+-W are obtained by Ar and N_2 tagging. IRPD spectra are measured in the XH stretch range ($X = \text{N}$ and O , $2600-3800 \text{ cm}^{-1}$) with a quadrupole-octopole-quadrupole tandem mass spectrometer described elsewhere.¹⁸⁻²⁶ Briefly, protonated clusters are generated in an electron ionization source coupled to a pulsed molecular beam expansion. Solid 5HI (Sigma-Aldrich, 97%) is heated to $145 \text{ }^\circ\text{C}$ and the resulting vapor is seeded in Ar or N_2 carrier gas (8-9 bar) containing water. He/H_2 gas (90/10) is added to pure Ar (N_2) in a 1:1 ratio to enhance the protonation efficiency. The gas mixture is expanded into vacuum through a pulsed nozzle. Close to the nozzle orifice, electron and/or chemical ionization of X ($X = \text{Ar}$, N_2 , H_2 , or W_n) forms XH^+ ions, which subsequently protonate 5HI *via* proton transfer.^{18-20,24} Three-body collisions with W and/or carrier gas molecules lead to cluster formation. In the first quadrupole, the desired parent clusters are mass-selected. Pulsed IR radiation emitted from a tunable optical parametric oscillator (2-4 mJ pulse energy, 10 Hz repetition rate, 1 cm^{-1} bandwidth) is introduced into the adjacent octopole. Resonant vibrational excitation of the parent clusters leads to the loss of the least bonded ligand, *i.e.* single W molecules in the case of 5HIH^+-W_n or the loosely bound Ar/ N_2 tag for $5\text{HIH}^+-W-\text{Ar}/\text{N}_2$. The produced fragment ions are mass-selected with the second quadrupole and monitored by a Daly detector as a function of the IR laser frequency to derive the IRPD spectrum of the 5HIH^+-W_n or $5\text{HIH}^+-W-\text{Ar}/\text{N}_2$ parent clusters. The ion source is triggered at 20 Hz (twice the laser frequency) facilitating subtraction of the background signal (metastable decay). All IRPD spectra are normalized for laser intensity variations recorded with a pyroelectric detector. Collision-induced dissociation in the octopole confirms the composition of mass-selected parent clusters.^{22,23}

The protonation sites in 5HIH^+ have already been examined in a previous study.¹⁷ The 5HIH^+ protomers offer two competing H-bonding sites for W molecules, namely, their NH and OH functional groups. The 5HIH^+-W_n input structures for geometry optimization are constructed by hand, attaching W ligands successively to the NH and OH groups. Up to $n = 3$, this approach is still feasible to find all low-energy minima. For larger hydrates (and more flexible organic chromophores), it may be rather difficult to find geometries by hand and, as a consequence, systematic computational sampling techniques such as basin-hopping or molecular dynamics should be employed.²⁸⁻³⁰ Geometries, energies, and harmonic IR spectra of stable 5HIH^+-W_n ($n = 1-3$) and $5\text{HIH}^+-W-\text{Ar}/\text{N}_2$ (ESI^+) structures are calculated at the B3LYP-D3/aug-cc-pVTZ level using GAUSSIAN09.^{27,31-34} This hybrid density functional with



additive dispersion correction has proven to yield reliable results for related aromatic clusters.^{11,22,23,35–39} For example, the binding energies computed for the monohydrates of the benzene and naphthalene cations ($D_0 = 3209$ and 2773 cm^{-1}) compare favorably with the experimental values ($D_0 = 3290 \pm 120$ and 2800 ± 300 cm^{-1}).^{40,41} Similarly, the calculated binding energy of W_2 ($D_0 = 1108$ cm^{-1}) matches the measured value ($D_0 = 1105 \pm 10$ cm^{-1}).^{42,43} We optimize selected structures also at the PBE0-D3/aug-cc-pVTZ level to yield reference data on structural and spectroscopic properties. The PBE0 functional is frequently used to compute properties of molecular clusters and is less empirical than B3LYP.^{28,30,44–46} Furthermore, single-point energy calculations of selected optimized structures are performed at the CC2/aug-cc-pVDZ level. Comparison to neutral and cationic $s/a5\text{HIH}-W$ clusters previously studied at the same level (B3LYP-D3/aug-cc-pVTZ) yields the effects of protonation.^{22,23} Total binding energies (D_0) are derived by subtracting the zero-point corrected energies of the corresponding monomers from that of the cluster: $D_0 = E_0(5\text{HIH}^+-W_n) - E_0(5\text{HIH}^+) - n \cdot E_0(W)$. Binding energies of the Ar/N_2 tag in $5\text{HIH}^+-W-\text{Ar}/\text{N}_2$ are calculated in a similar way. For selected structures, we estimate the basis set superposition errors (BSSE) using the counterpoise method.^{47,48} Calculated harmonic vibrational frequencies are linearly scaled by a factor of 0.957 derived previously from fitting the free OH stretching frequency of cationic $s/a5\text{HIH}^+$.²² Yet, this scaling factor yields OH stretch frequencies of W ($\nu_{1/3} = 3635/3733$ cm^{-1}) systematically lower by ~ 20 cm^{-1} than the experimental values ($\nu_{1/3} = 3657/3756$ cm^{-1}).⁴⁹ For selected 5HIH^+-W clusters, anharmonic spectra are calculated at the B3LYP-D3/aug-cc-pVDZ level as implemented in GAUSSIAN09.⁵⁰

3. Results and discussion

3.1 IRPD spectra

An overview of the IRPD spectra of 5HIH^+-W_n with $n = 1-3$ measured in the XH stretch range is given in Fig. 1. The positions, widths, and suggested vibrational and isomer assignments of the transitions observed (A–L, X) are listed in Table 1, along with computed frequencies and IR oscillator strengths. The spectra cover the antisymmetric and symmetric OH stretching modes of water (ν_3 and ν_1 , bands A and B) and the OH and NH stretching modes of 5HIH^+ (ν_{OH} and ν_{NH}). Dotted lines indicate the correspondence of related bands in the different spectra. Bands G and H emerging in the spectra of 5HIH^+-W_n with $n \geq 2$ already indicate the formation of a H-bonded water network. In the following, the IRPD spectra are disentangled by comparison with IR spectra calculated for the possible isomers and taking into account the previous interpretation of the 5HIH^+-L_n spectra with $L = \text{Ar}$ and N_2 .¹⁷ IRPD spectra of tagged 5HIH^+-W-L clusters with $L = \text{Ar}$ and N_2 are shown in Fig. S1 in the ESI.†

3.2 5HIH^+-W

Internal OH rotation yields *syn* and *anti* rotamers of 5HIH^+ , denoted by $s5\text{HIH}^+$ and $a5\text{HIH}^+$, respectively. Our previous study of 5HIH^+-L_n demonstrates the coexistence of the most

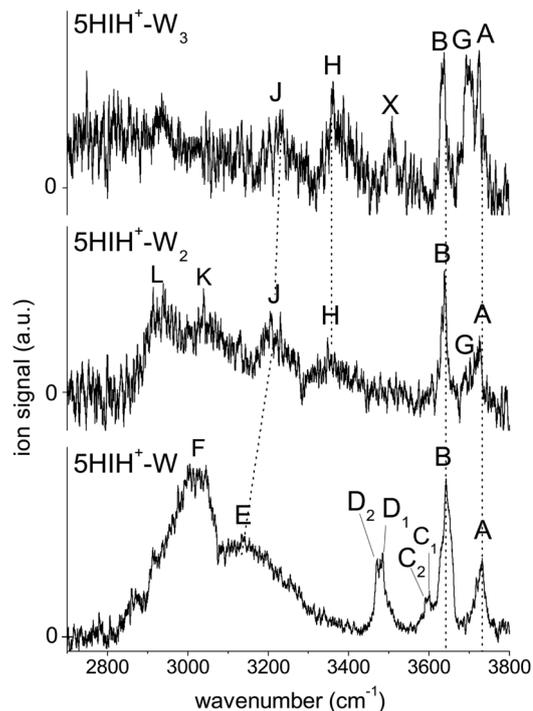


Fig. 1 Overview of the IRPD spectra of 5HIH^+-W_n ($n = 1-3$) measured in the XH stretch range ($2700-3800$ cm^{-1}). The positions, widths, and vibrational and isomer assignments of the transitions are listed in Table 1.

stable $s/a5\text{HIH}^+(\text{C}3)$ protomers ($\Delta E_0 = 0/1.8$ kJ mol^{-1}), as well as the $s/a5\text{HIH}^+(\text{C}4)$ and $s/a5\text{HIH}^+(\text{O})$ isomers ($\Delta E_0 = 5.1/11.1$ and $117.1/117.7$ kJ mol^{-1}).¹⁷ With the exception of $5\text{HIH}^+(\text{C}4)$, the *syn* and *anti* rotamers of the 5HIH^+ protomers have similar energies and IR spectra and thus cannot be distinguished at the current spectral resolution. A deep potential well ($\Delta E > 150$ kJ mol^{-1}) prevents the interconversion of the drastically less stable $5\text{HIH}^+(\text{O})$ isomers into $5\text{HIH}^+(\text{C}4)$.¹⁷ We compute proton affinities (PAs) of $905/902$ kJ mol^{-1} for $s/a5\text{HIH}^+(\text{C}3)$, $900/892$ kJ mol^{-1} for $s/a5\text{HIH}^+(\text{C}4)$, and $786/785$ kJ mol^{-1} for $s/a5\text{HIH}^+(\text{O})$.¹⁷ On the other hand, for W_n clusters we calculate PAs of 681 , 818 , and 898 kJ mol^{-1} for $n = 1-3$, respectively. Hence, the predicted PA of $s/a5\text{HIH}^+(\text{O})$ is smaller than that of the W_n cluster already at $n = 2$, and the proton may be transferred from this 5HIH^+ protomer to the W_n solvent cluster. The PAs of $s/a5\text{HIH}^+(\text{C}3/\text{C}4)$ are in the same range as that of the W_3 cluster. Yet, proton transfer may still occur if the solvation energy of the proton-transferred structure is higher. Due to the possible proton transfer to the W_n moiety, the notation $[5\text{HI}-W_n]\text{H}^+$ is more precise. However, for consistency, we keep the 5HIH^+-W_n notation for all clusters throughout this paper. Based on our previous results,¹⁷ we calculate herein hydrated clusters of $s/a5\text{HIH}^+(\text{C}3)$, $s/a5\text{HIH}^+(\text{C}4)$, and $s/a5\text{HIH}^+(\text{O})$ with W attached to the acidic functional $\text{OH}_{(2)}$ and NH groups, the π -electron cloud, or the protonated CH_2 group. In general, the binding motifs are very similar for the *syn* and *anti* rotamers of the 5HIH^+-W_n clusters, with $s5\text{HIH}^+-W_n$ being systematically more stable (Table S1, ESI†). Therefore, in the main text and



Table 1 Positions (in cm^{-1}) and widths (FWHM, in parentheses) of the bands observed in the IRPD spectra of 5HIIH^+-W_n ($n = 1-3$) (Fig. 1) compared to scaled harmonic frequencies (in cm^{-1}) and IR intensities (km mol^{-1} , in parentheses) of the assigned $s5\text{HIIH}^+-W_n$ isomers calculated at the B3LYP-D3/aug-cc-pVTZ level. Vibrational frequencies of the corresponding $a5\text{HIIH}^+-W_n$ clusters are listed in Table S1 (ESI)

Cluster	Exp.	Vibration	Calc.	Isomer	
5HIIH^+-W	A 3732 (30)	ν_3	3704 (119)	$s5\text{HIIH}^+(\text{C3})-\text{W}(\text{NH})$	
		ν_3	3708 (118)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{NH})$	
		ν_3	3697 (134)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{OH})$	
		ν_3	3685 (178)	$s5\text{HIIH}^+(\text{O})-\text{W}(\text{OH})$	
		B 3641 (25)	$\nu_{\text{OH}}^{\text{f}}$	3632 (164)	$s5\text{HIIH}^+(\text{C3})-\text{W}(\text{NH})$
			ν_1	3622 (54)	$s5\text{HIIH}^+(\text{C3})-\text{W}(\text{NH})$
	ν_1		3612 (52)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{OH})$	
	ν_1		3622 (40)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{NH})$	
	C ₁ 3600 (5)	$\nu_{\text{OH}}^{\text{f}}$	3600 (68)	$s5\text{HIIH}^+(\text{O})-\text{W}(\text{OH})$	
		$\nu_{\text{OH}}^{\text{f}}$	3588 (219)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{NH})$	
	C ₂ 3591 (5)	$\nu_{\text{OH}}^{\text{f}}$	3558 (199)	$s5\text{HIIH}^+(\text{O})-\text{W}(\text{OH})$	
	D ₁ 3485 (30)	$\nu_{\text{NH}}^{\text{f}}$	3494 (116)	$s5\text{HIIH}^+(\text{O})-\text{W}(\text{OH})$	
	D ₂ 3470 (30)	$\nu_{\text{NH}}^{\text{f}}$	3478 (139)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{OH})$	
	E 3130 (100)	$\nu_{\text{NH}}^{\text{b}}$	3191 (1184)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{NH})$	
	F 3015 (70)	$\nu_{\text{OH}}^{\text{b}}$	3035 (2283)	$s5\text{HIIH}^+(\text{C4})-\text{W}(\text{OH})$	
		$\nu_{\text{NH}}^{\text{b}}$	3011 (1399)	$s5\text{HIIH}^+(\text{C3})-\text{W}(\text{NH})$	
5HIIH^+-W_2	A 3725 (20)	ν_3	3711 (131)/3707 (113)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{OH})$	
		ν_3	3717 (114)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{W})$	
		ν_3	3711 (114)/3700 (128)	$s5\text{HIIH}^+(\text{C4})-W_2(\text{NH}-\text{OH})$	
	G 3692 (10)	$\nu_{\text{OH}}^{\text{f(W)}}$	3690 (105)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{W})$	
		B 3638 (15)	ν_1	3624 (49)/3619 (33)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{OH})$
	$\nu_{\text{OH}}^{\text{f}}$		3635 (156)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{W})$	
	H 3348 (> 50)	ν_1	3624 (37)/3615 (48)	$s5\text{HIIH}^+(\text{C4})-W_2(\text{NH}-\text{OH})$	
		$\nu_{\text{OH}}^{\text{b}}$	3283 (1565)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{OH})$	
		$\nu_{\text{OH}}^{\text{b(W)}}$	3308 (921)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{W})$	
	J 3205 (> 50)	$\nu_{\text{NH}}^{\text{b}}$	3218 (1094)	$s5\text{HIIH}^+(\text{C4})-W_2(\text{NH}-\text{OH})$	
		K 3040 (> 50)	$\nu_{\text{NH}}^{\text{b}}$	3041 (1331)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{OH})$
	$\nu_{\text{OH}}^{\text{b}}$		3083 (2234)	$s5\text{HIIH}^+(\text{C4})-W_2(\text{NH}-\text{OH})$	
	L 2940 (> 50)	$\nu_{\text{NH}}^{\text{b}}$	2787 (2249)	$s5\text{HIIH}^+(\text{C3})-W_2(\text{NH}-\text{W})$	
5HIIH^+-W_3		A 3725 (20)	ν_3	3719 (112)/3708 (114)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{OH}-\text{NH}-\text{W})$
	ν_3		3721 (111)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$	
	ν_3		3722 (208)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{W}-\text{NH}-\text{W})$	
	ν_3		3723 (106)/3702 (127)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{NH}-\text{W})$	
	ν_3		3722 (115)/3712 (112)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{W}-\text{NH})$	
	ν_3		3702 (147)/3695 (222)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W1})$	
	ν_3		3717 (128)/3704 (143)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W2})$	
	G 3692 (30)		$\nu_{\text{OH}}^{\text{f}}$	3692 (102)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{OH}-\text{NH}-\text{W})$
		$\nu_{\text{OH}}^{\text{f(W)}}$	3694 (70)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$	
		$\nu_{\text{OH}}^{\text{f(W)}}$	3693 (111)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$	
		$\nu_{\text{OH}}^{\text{f(W)}}$	3689 (92)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{NH}-\text{W})$	
		$\nu_{\text{OH}}^{\text{f(W)}}$	3679 (100)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{W}-\text{NH})$	
		$\nu_{\text{OH}}^{\text{f(W)}}$	3661 (140)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W1})$	
		$\nu_{\text{OH}}^{\text{f(W)}}$	3677 (97)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W2})$	
		B 3637 (20)	ν_1	3630 (27)/3620 (31)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{OH}-\text{NH}-\text{W})$
	$\nu_{\text{OH}}^{\text{f(W)}}$		3635 (151)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$	
ν_1	3631 (23)		$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$		
$\nu_{\text{OH}}^{\text{f}}$	3636 (147)		$s5\text{HIIH}^+(\text{C3})-W_3(\text{W}-\text{NH}-\text{W})$		
ν_1	3631 (29)		$s5\text{HIIH}^+(\text{C3})-W_3(\text{W}-\text{NH}-\text{W})$		
ν_1	3632 (24)/3616 (45)		$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{NH}-\text{W})$		
ν_1	3631 (29)/3624 (35)		$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{W}-\text{NH})$		
ν_1	3616 (54)/3586 (100)		$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W1})$		
X 3508 (30)	ν_1	3628 (36)/3617 (45)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W2})$		
	$\nu_{\text{NH}}^{\text{f}}$	3497 (108)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W1})$		
	$\nu_{\text{NH}}^{\text{f}}$	3500 (105)	$s5\text{HIIH}^+(\text{O})-W_3(\text{NH}-\text{OH}-\text{W2})$		
	$\nu_{\text{OH}}^{\text{b(W)}}$	3375 (723)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{NH}-\text{W})$		
H 3360 (> 50)	$\nu_{\text{OH}}^{\text{b(W)}}$	3306 (891)	$s5\text{HIIH}^+(\text{C4})-W_3(\text{OH}-\text{W}-\text{NH})$		
	$\nu_{\text{OH}}^{\text{b(W)}}$	3326 (867)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{OH}-\text{NH}-\text{W})$		
	$\nu_{\text{OH}}^{\text{b}}$	3301 (156)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{OH}-\text{NH}-\text{W})$		
	$\nu_{\text{OH}}^{\text{b(W)}}$	3378 (663)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{NH}-\text{W}-\text{W})$		
	$\nu_{\text{OH}}^{\text{b(as)}}$	3424 (380)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{W}-\text{NH}-\text{W})$		
	$\nu_{\text{OH}}^{\text{b(s)}}$	3381 (516)	$s5\text{HIIH}^+(\text{C3})-W_3(\text{W}-\text{NH}-\text{W})$		



Table 1 (continued)

Cluster	Exp.	Vibration	Calc.	Isomer
	J 3230 (> 50)	$\nu_{\text{OH}}^{\text{b}}$	3110 (1927)	$s5\text{HIH}^+(\text{C4})-\text{W}_3(\text{OH}-\text{NH}-\text{W})$
		$\nu_{\text{NH}}^{\text{b}}$	3078 (1906)	$s5\text{HIH}^+(\text{C4})-\text{W}_3(\text{OH}-\text{NH}-\text{W})$
		$\nu_{\text{NH}}^{\text{b}}$	3236 (1063)	$s5\text{HIH}^+(\text{C4})-\text{W}_3(\text{OH}-\text{W}-\text{NH})$
		$\nu_{\text{OH}}^{\text{b(W)}}$	3126 (1404)	$s5\text{HIH}^+(\text{C3})-\text{W}_3(\text{NH}-\text{W}-\text{W})$
		$\nu_{\text{OH}}^{\text{b}}$	3183 (1263)	$s5\text{HIH}^+(\text{O})-\text{W}_3(\text{NH}-\text{OH}-\text{W2})$
		$\nu_{\text{OH}}^{\text{b}}$	3183 (1263)	$s5\text{HIH}^+(\text{O})-\text{W}_3(\text{NH}-\text{OH}-\text{W2})$

figures, we concentrate on $s5\text{HIH}^+-\text{W}_n$. Corresponding data for $a5\text{HIH}^+-\text{W}_n$ are available in the ESI†.

Fig. 2 summarizes the calculated $s5\text{HIH}^+-\text{W}$ clusters relevant for the present study, along with their binding and relative energies (D_0 , E_0) and intermolecular H-bond lengths (R). Additional geometric and spectroscopic properties are listed in Table S1 (ESI†). Further stable binding motifs of $s5\text{HIH}^+-\text{W}$ are shown in Fig. S2, and corresponding structures of $a5\text{HIH}^+-\text{W}$ are given in Fig. S3 (Table S1, ESI†).

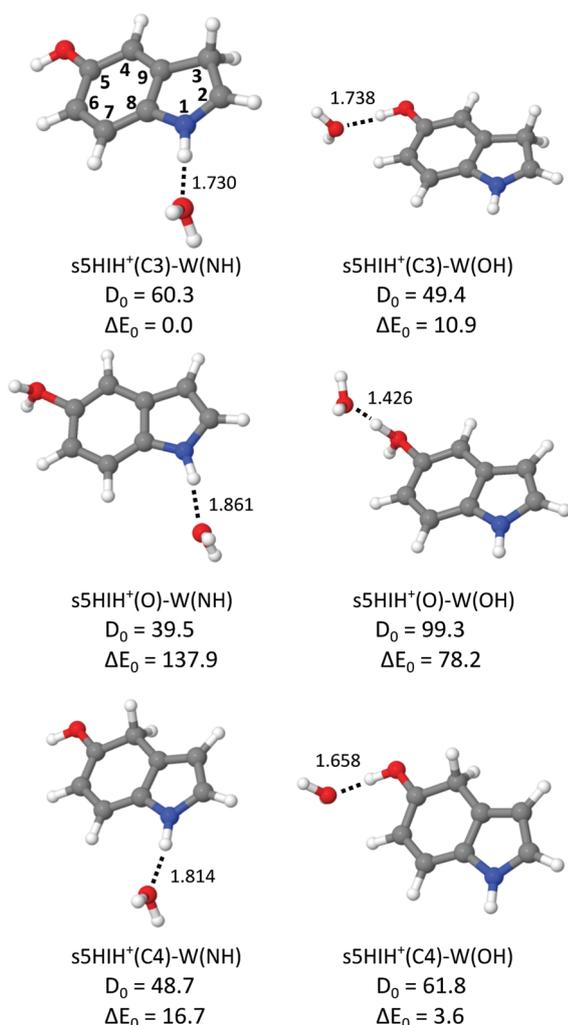


Fig. 2 Structures of selected $s5\text{HIH}^+-\text{W}$ clusters calculated at the B3LYP-D3/aug-cc-pVTZ level (Table S1, ESI†) along with binding energies and relative energies (D_0 and E_0 in kJ mol^{-1}) and intermolecular bond lengths (R in Å).

All 5HIH^+ protomers exhibit competing H-bonding sites, and most important are the acidic functional $\text{OH}_{(2)}$ and NH groups acting as proton donors in $\text{OH}\cdots\text{O}$ and $\text{NH}\cdots\text{O}$ H-bonds to W . As already observed for related heterocyclic arene cations,^{23,36,41,51–53} π -stacking of W is rather unfavorable, e.g., $D_0 = 33.3 \text{ kJ mol}^{-1}$ for $s5\text{HIH}^+(\text{C3})-\text{W}(\pi)$, and thus not considered further. Only one isomer with W attached to the protonated CH_2 group could be located, namely, $s5\text{HIH}^+(\text{C4})-\text{W}(\text{CH})$ with $D_0 = 28.6 \text{ kJ mol}^{-1}$. Because of the low binding energy of the $\text{CH}\cdots\text{O}$ H-bonds, this binding motif is also not considered further. Depending on the protonation site, the acidity of the OH and NH functional groups is changed,¹⁷ and the H-bond strengths vary in the same way. For example, the NH group is the strongest proton donor in $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ ($\Delta E_0 = 0$ and $D_0 = 60.3 \text{ kJ mol}^{-1}$, $R = 1.730 \text{ \AA}$ for $\text{NH}\cdots\text{O}$), while the OH group is preferred in $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ ($\Delta E_0 = 3.6$ and $D_0 = 61.8 \text{ kJ mol}^{-1}$, $R = 1.658 \text{ \AA}$ for $\text{OH}\cdots\text{O}$). The $\text{OH}\cdots\text{O}$ H-bond of the protonated OH_2 oxonium group is outstandingly strong, with $D_0 = 99.3 \text{ kJ mol}^{-1}$ and $R = 1.426 \text{ \AA}$ for $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$. This strong H-bond and the twofold degeneracy may thus again somewhat compensate for the large energy gap of $\Delta E_0 = 78.2 \text{ kJ mol}^{-1}$ between $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ and the $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ global minimum. The NH group of the oxonium protomer is a far less acidic than the OH_2 group, with $D_0 = 39.5 \text{ kJ mol}^{-1}$ and $R = 1.861 \text{ \AA}$ for the $\text{NH}\cdots\text{O}$ H-bond in $s5\text{HIH}^+(\text{O})-\text{W}(\text{NH})$. All considered $s5\text{HIH}^+-\text{W}$ isomers are readily distinguishable by their IR spectra in the investigated spectral range (Fig. 3 and Fig. S4, ESI†). Our calculated binding energies of $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ and $s5\text{HIH}^+(\text{C3})-\text{W}(\text{OH})$, $D_0 = 60.3$ and 49.4 kJ mol^{-1} , are substantially but systematically smaller than the corresponding H-bond energies reported earlier at the RI-MP2/aug-cc-pVDZ level, $D_e = 88.6$ and 76.8 kJ mol^{-1} .⁵ Similar to the MP2 results,⁵ the single-point energies computed at the CC2/aug-cc-pVDZ level yield the same energy hierarchy as predicted at the B3LYP-D3/aug-cc-pVTZ level, but the spread of relative energies is somewhat larger (Table S1, ESI†). In general, BSSE corrections of the computed binding energies evaluated at the B3LYP-D3/aug-cc-pVTZ level are small (on the order of 1% or less) because the aug-cc-pVTZ basis set is rather large (Table S1, ESI†). Reference calculations at the PBE0-D3/aug-cc-pVTZ level yield energies, structural and spectroscopic properties comparable to those obtained at the B3LYP-D3/aug-cc-pVTZ level (Table S1, ESI†). Indeed, the IR spectra computed at the B3LYP-D3/aug-cc-pVTZ level match somewhat better the experimental spectra (Fig. S5, ESI†).

In Fig. 3 the IRPD spectrum of $5\text{HIH}^+-\text{W}$ is compared to that of cationic $5\text{HI}^+-\text{W}$ and to linear IR absorption spectra calculated for relevant $s5\text{HIH}^+-\text{W}$ isomers (Table 1). The vibrational assignment



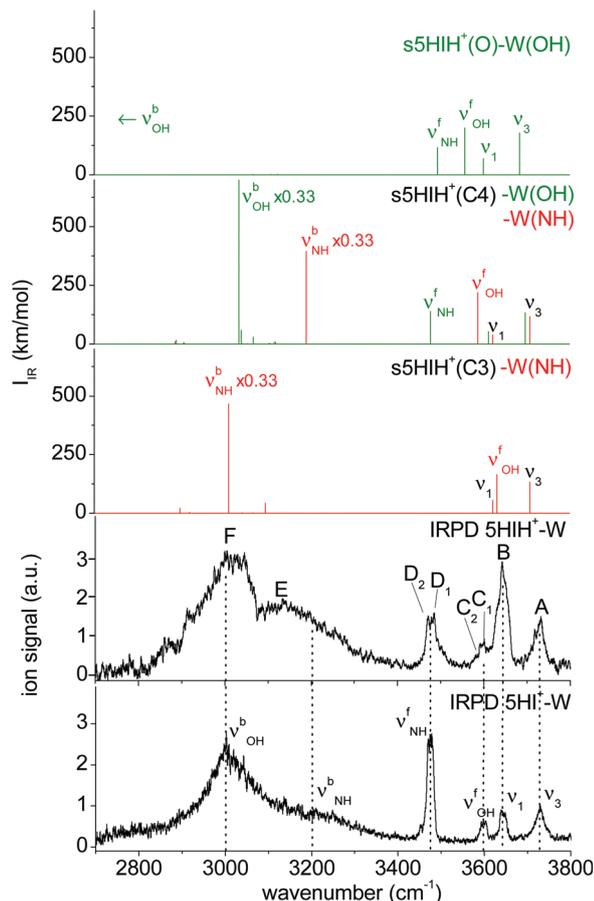


Fig. 3 Comparison of the IRPD spectrum of $5\text{HIH}^+-\text{W}$ to linear IR spectra calculated for the most stable isomers at the B3LYP-D3/aug-cc-pVTZ level. For comparison, the IRPD spectrum of $5\text{HI}^+-\text{W}$ is also shown.²³

is facilitated by comparing the spectra of $5\text{HIH}^+-\text{W}$ and $5\text{HI}^+-\text{W}$. Bands A and B at 3732 and 3641 cm^{-1} are readily assigned to the antisymmetric and symmetric free OH stretching modes of W (ν_3 and ν_1). Band A may also contain the red-shifted ν_3 mode of $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ and $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ predicted at 3685 and 3697 cm^{-1} , respectively. The relative intensity of band B is substantially enhanced for $5\text{HIH}^+-\text{W}$ as compared to $5\text{HI}^+-\text{W}$, while this is not the case for band A. Some contribution of the free phenolic OH stretching mode (ν_{OH}^f) of $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ predicted at 3632 cm^{-1} rationalizes the higher intensity of band B. Indeed, comparison of the IRPD spectra of $5\text{HIH}^+-\text{L}$ with $\text{L} = \text{Ar}, \text{N}_2$, and W further strengthens this assignment.¹⁷ The ν_{OH}^f band of $s5\text{HIH}^+(\text{C3})-\text{Ar}(\pi)$ is observed at 3635 cm^{-1} , that is only 6 cm^{-1} red-shifted from the maximum of peak B. By comparison to $5\text{HI}^+-\text{W}$, bands C_1/C_2 are clearly assigned to phenolic ν_{OH}^f modes. We attribute C_1 at 3600 cm^{-1} to ν_{OH}^f of $s5\text{HIH}^+(\text{C4})-\text{W}(\text{NH})$ predicted at 3588 cm^{-1} , and C_2 at 3591 cm^{-1} to ν_{OH}^f of $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ predicted at 3558 cm^{-1} . Another indication for the production of $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ is the doublet D_1/D_2 at $3485/3470\text{ cm}^{-1}$ ($\Delta\nu = 15\text{ cm}^{-1}$). By analogy to $5\text{HI}^+-\text{W}$, it is assigned to the free NH stretching mode (ν_{NH}^f). Here, D_1 is attributed to $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ and D_2 to $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ predicted at 3494 and 3478 cm^{-1} ($\Delta\nu = 16\text{ cm}^{-1}$), respectively.

The broad band E centered at 3130 cm^{-1} is interpreted as H-bonded NH stretch (ν_{NH}^b) of $s5\text{HIH}^+(\text{C4})-\text{W}(\text{NH})$ predicted at 3191 cm^{-1} . Band E is somewhat red-shifted compared to ν_{NH}^b in the IRPD spectrum of cationic $5\text{HI}^+-\text{W}$, indicating a destabilization of the NH bond induced by C4-protonation. The broad transition F at 3015 cm^{-1} contains both ν_{OH}^b of $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ and ν_{NH}^b of $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ predicted at 3035 and 3011 cm^{-1} . The IRPD spectrum shows no clear signature of the $s5\text{HIH}^+(\text{C3})-\text{W}(\text{OH})$ isomer (Fig. S4, ESI[†]). This finding is somewhat surprising because this isomer is predicted to be slightly more stable than $s5\text{HIH}^+(\text{C4})-\text{W}(\text{NH})$ ($\Delta E_0 = 10.9$ and $D_0 = 49.4\text{ kJ mol}^{-1}$ vs. $\Delta E_0 = 16.7$ and $D_0 = 48.7\text{ kJ mol}^{-1}$). We also rule out the presence of the $s5\text{HIH}^+(\text{O})-\text{W}(\text{NH})$ isomer, mostly on the basis of stability and spectroscopy (Fig. S4, ESI[†]).

In conclusion, we assign the measured IRPD spectrum of $5\text{HIH}^+-\text{W}$ to C3-protonated $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ with NH-bonded W, C4-protonated $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ and $s5\text{HIH}^+(\text{C4})-\text{W}(\text{NH})$ with OH- and NH-bonded W, and the OH-bonded $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ oxonium ion. This interpretation is strengthened by a comparison of the IRPD spectrum to exemplary anharmonic spectra computed for the monohydrates (Fig. S6 and Table S1, ESI[†]). Obviously, within the harmonic approximation, the red-shifts of the H-bond donor stretching vibrations (ν_{OH}^b and ν_{NH}^b) are underestimated by $60\text{--}110\text{ cm}^{-1}$, but both the anharmonic and scaled harmonic spectra yield the same assignment of the experimental transitions. At the current experimental resolution, we cannot discriminate *syn* and *anti* rotamers but assume that both are present in significant abundance for the assigned isomers. The estimation of population ratios for the three protomers (C3, C4, O) based on the comparison of measured and computed band intensities is not straightforward due to overlapping transitions. Furthermore, the binding energies of the $5\text{HIH}^+-\text{W}$ clusters are higher than the IR photon energies in the XH stretch range, such that absorption of a single photon will not lead to fragmentation from cold clusters. Hence, multiple-photon effects or ions with high internal energy must be considered. Moreover, due to the very different binding energies of the individual $5\text{HIH}^+-\text{W}$ clusters, the photodissociation cross sections may be different, too. Still, assuming similar fragmentation cross sections for the assigned isomers, a rough estimate of populations is possible. In contrast to the predictions for $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$, band C ($\text{C}_1 + \text{C}_2$, ν_{OH}^f) is significantly less intense than D ($\text{D}_1 + \text{D}_2$, ν_{NH}^f). This result suggests that we probe only a few oxonium ions ($\sim 10\%$), while the main contribution to band D arises from $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$. The comparison of bands C and D also indicates that significantly fewer $s5\text{HIH}^+(\text{C4})-\text{W}(\text{NH})$ than $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ isomers are probed. Band B is roughly twice as intense as band A, while ν_1 is predicted to be half as intense as ν_3 in any of the calculated IR spectra. Thus, we conclude that most signals of band B stem from ν_{OH}^f of $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$. Hence, we suggest that the major contribution (roughly 70%) to the IRPD spectrum arises from the most stable $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ and $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ isomers.

3.3 $5\text{HIH}^+-\text{W}-\text{Ar}/\text{N}_2$

Tagging with Ar or N_2 reduces the peak widths in the IRPD spectra of $5\text{HIH}^+-\text{W}$ because the binding energy of the least



bonded ligand limits the internal temperature of the cluster. Fig. S7 and S8 (ESI[†]) compare the IRPD spectra of 5HIH⁺-W-Ar and 5HIH⁺-W-N₂ to the IR spectra calculated for relevant tagged isomers. The IRPD spectra of 5HIH⁺-W-Ar/N₂ do not add any new information about the monohydrated clusters but confirm the assignment of s5HIH⁺(C3)-W(NH), s5HIH⁺(C4)-W(OH), s5HIH⁺(C4)-W(NH) and s5HIH⁺(O)-W(OH) being the predominant monohydrates in the molecular beam. The Ar tag is mainly attached to the W moiety or π -stacked. The N₂ ligand is H-bonded to either W or to the remaining functional group not occupied by W.

Selected structures of s5HIH⁺-W-Ar and s5HIH⁺-W-N₂ are depicted in Fig. S9 and S10 (ESI[†]), respectively, and Table S1 (ESI[†]) contains structural and spectroscopic information for all calculated 5HIH⁺-W-Ar/N₂ isomers. For a detailed discussion of the 5HIH⁺-W-L spectra, the interested reader is referred to the ESI.[†]

3.4 5HIH⁺-W₂

Fig. 4 shows the structures of selected s5HIH⁺-W₂ clusters along with binding energies and H-bond lengths. Additional

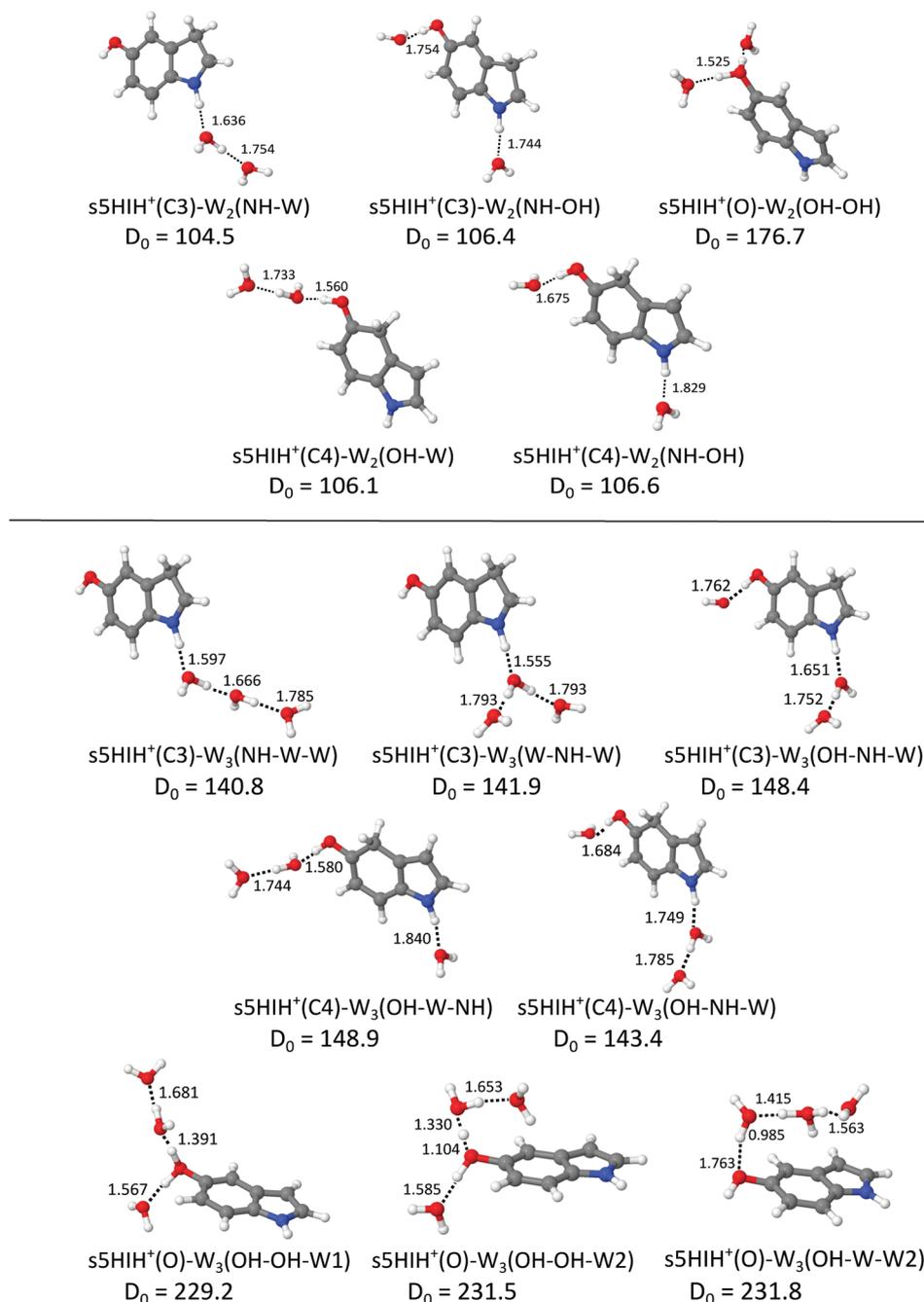


Fig. 4 Structures of selected s5HIH⁺-W_{2/3} clusters calculated at the B3LYP-D3/aug-cc-pVTZ level (Table S1, ESI[†]) along with binding energies (D_0 in kJ mol⁻¹) and intermolecular distances (R in Å).



stable $s5\text{HIH}^+-\text{W}_2$ isomers are shown in Fig. S11 (ESI[†]). Corresponding $a5\text{HIH}^+-\text{W}_2$ structures are depicted in Fig. S12 (ESI[†]). Selected structural and spectroscopic properties of all calculated $s/a5\text{HIH}^+-\text{W}_2$ isomers are listed in Table S1 (ESI[†]).

For $5\text{HIH}^+(\text{C3/C4})-\text{W}_2$, we consider three binding motifs. H-bonding of two individual W ligands to both functional groups yields the most stable $s5\text{HIH}^+(\text{C3/C4})-\text{W}_2(\text{NH}-\text{OH})$ with $D_0 = 106.4/106.6 \text{ kJ mol}^{-1}$. BSSE corrections are again on the order of 1% of the computed binding energies (Table S1, ESI[†]). Noncooperative three-body effects weaken the individual H-bonds compared to the monohydrates, as revealed by comparison of binding energies and H-bond lengths. For example, corresponding H-bonds elongate from $R = 1.730$ and 1.738 \AA in $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ and $s5\text{HIH}^+(\text{C3})-\text{W}(\text{OH})$ to $R = 1.744$ and 1.754 \AA in $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{OH})$. The same trend holds for $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{NH}-\text{OH})$ with $R = 1.814$ and 1.658 \AA compared to $R = 1.829$ and 1.675 \AA . This weakening of the H-bonds directly translates into reduced complexation-induced redshifts of the corresponding $\nu_{\text{NH}}^{\text{b}}$ and $\nu_{\text{OH}}^{\text{b}}$ modes (Table S1, ESI[†]). For example, $\nu_{\text{NH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ is predicted at 3011 cm^{-1} and that of $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{OH})$ at 3041 cm^{-1} . For the C3-protonated ion, formation of a H-bonded water network (*i.e.*, attachment of a H-bonded W_2) at the NH group is more favorable than at the OH group, with $D_0 = 104.5$ and 86.8 kJ mol^{-1} for $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$ and $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{OH}-\text{W})$, respectively. Due to cooperativity, the initial H-bond is shortened from $R = 1.730 \text{ \AA}$ in $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ to 1.636 \AA in $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$. In line with the altered acidity of the functional groups for the C4-protonated ion,¹⁷ $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH}-\text{W})$ is more stable than $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{NH}-\text{W})$, with $D_0 = 106.1$ and 87.6 kJ mol^{-1} , respectively. Again, network formation strengthens the first H-bond with $R = 1.658 \text{ \AA}$ in $s5\text{HIH}^+(\text{C4})-\text{W}(\text{OH})$ vs. 1.560 \AA in $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH}-\text{W})$. Formation of water networks yields characteristic spectroscopic features, most prominently, $\nu_{\text{OH}}^{\text{f(W)}}$ and $\nu_{\text{OH}}^{\text{b(W)}}$ of the solvated water in the W_2 unit, predicted at 3690 and 3307 cm^{-1} for $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$. The corresponding ν_3 and ν_1 of the terminal free water are blue-shifted from 3708 and 3622 cm^{-1} in $s5\text{HIH}^+(\text{C3})-\text{W}(\text{NH})$ to 3717 and 3628 cm^{-1} in $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$. The impact on the respective functional group by solvation with W_2 is largely enhanced. For example, $\nu_{\text{OH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH})$ is predicted at 3035 cm^{-1} and shifts down to 2710 cm^{-1} for $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH}-\text{W})$.

The largest binding energy of $D_0 = 176.7 \text{ kJ mol}^{-1}$ is calculated for $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{OH})$, *i.e.* the oxonium ion monohydrated at both OH groups. Yet, due to noncooperativity, this value is significantly lower than twice the binding energy of $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$, $D_0 = 99.3 \text{ kJ mol}^{-1}$. Formation of a water network at the OH_2 group yields $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}1)$ and $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}2)$ with $D_0 = 167.3$ and $169.7 \text{ kJ mol}^{-1}$ (Fig. S11, ESI[†]). The NH group of the oxonium ion is far less attractive, leading to $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{NH}-\text{OH})$ and $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{NH}-\text{W})$ with only $D_0 = 135.5$ and 73.4 kJ mol^{-1} . Again, BSSE corrections are on the order of only 1% of the computed binding energies (Table S1, ESI[†]). In $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}1)$, the water chain points away from the phenol ring. Proton transfer

to the W_2 chain is indicated, because the OH bond of $s5\text{HIH}^+(\text{O})$ is already longer ($r_{\text{OH}} = 1.262 \text{ \AA}$) than the $\text{OH} \cdots \text{W}$ bond ($R = 1.147 \text{ \AA}$). Thus, the ion may be better described by $s5\text{HI}-\text{H}_5\text{O}_2^+$. Yet, for consistency, we keep the introduced notation for the structures. On the other hand, in $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}2)$ the water chain is bent toward the aromatic ring facilitating an additional $\text{OH} \cdots \pi$ interaction with the aromatic π -electron cloud. In this structure, proton transfer is more pronounced, with $r_{\text{OH}} = 1.434$ and $R = 1.056 \text{ \AA}$.

The assignment of the IRPD spectrum of $5\text{HIH}^+-\text{W}_2$ (Table 1) is based on the comparison to the corresponding calculated spectra (Fig. 5 and Fig. S13, ESI[†]). Exemplary anharmonic spectra depicted in Fig. S14 (ESI[†]) are in line with the following assignments based on the harmonic spectra. Bands A and B at 3725 and 3638 cm^{-1} are readily assigned to ν_3 and ν_1 and are well reproduced by essentially all considered isomers. Band G at 3692 cm^{-1} is characteristic of an H-bonded W_2 network and appears at 3690 and 3676 cm^{-1} in the calculated spectra of $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$ and $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH}-\text{W})$, respectively. In line with the assignments above, $\nu_{\text{OH}}^{\text{f}}$ of $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$ predicted at 3635 cm^{-1} contributes to band B, which is again significantly more intense than band A. The $\nu_{\text{NH}}^{\text{f}}$ mode characteristic of $s5\text{HIH}^+(\text{C4})-\text{W}_2(\text{OH}-\text{W})$ at 3481 cm^{-1} is however not observed. Hence, we assign only $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{W})$. Consequently, band H centered at 3348 cm^{-1} is attributed to a superposition of $\nu_{\text{OH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C3})-\text{W}_2(\text{NH}-\text{OH})$ predicted at

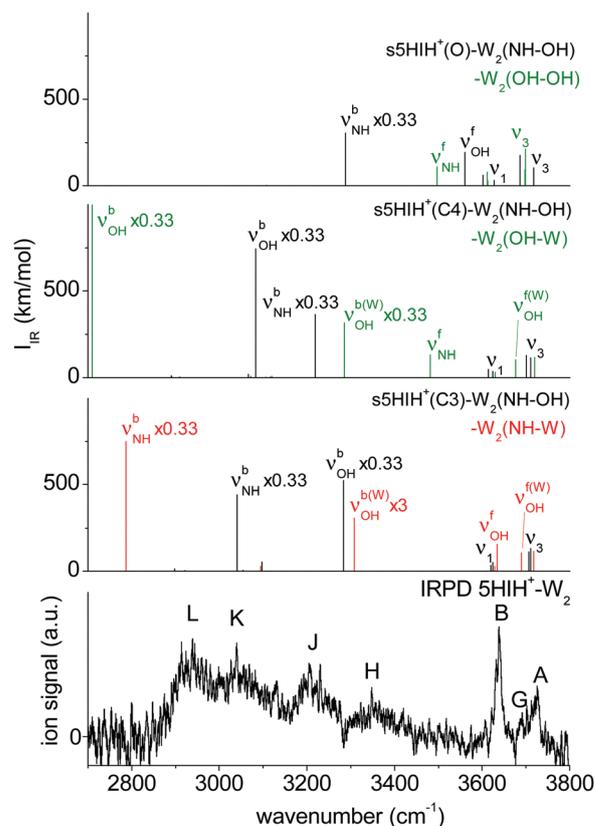


Fig. 5 Comparison of the IRPD spectrum of $5\text{HIH}^+-\text{W}_2$ to calculated IR spectra of the relevant isomers at the B3LYP-D3/aug-cc-pVTZ level.



3283 cm^{-1} and $\nu_{\text{OH}}^{\text{b}}$ of the W_2 unit in $s5\text{HIH}^+(\text{C}3)-\text{W}_2(\text{NH}-\text{W})$, which is predicted at 3308 cm^{-1} ($\nu_{\text{OH}}^{\text{b(W)}}$). Band J at 3205 cm^{-1} arises from $\nu_{\text{NH}}^{\text{b(W)}}$ of $s5\text{HIH}^+(\text{C}4)-\text{W}_2(\text{NH}-\text{OH})$ predicted at 3218 cm^{-1} . Its $\nu_{\text{OH}}^{\text{b}}$ mode at 3083 cm^{-1} and $\nu_{\text{NH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C}3)-\text{W}_2(\text{NH}-\text{OH})$ predicted at 3041 cm^{-1} account for band K observed at 3040 cm^{-1} . Finally, band L at 2940 cm^{-1} may be assigned to $\nu_{\text{NH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C}3)-\text{W}_2(\text{NH}-\text{W})$, which is predicted to be rather low (2787 cm^{-1}). However, below 2800 cm^{-1} , the laser power is very low such that this mode may also not be detected. We cannot safely assign any $5\text{HIH}^+(\text{O})-\text{W}_2$ oxonium isomer because there is no strong signal in the IRPD spectrum around 3500 cm^{-1} . In this range, the $\nu_{\text{NH}}^{\text{f}}$ and $\nu_{\text{OH}}^{\text{f}}$ modes of $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{OH})$ and $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{NH}-\text{OH})$ are predicted (3498 and 3560 cm^{-1} , Fig. 5). Furthermore, the $\nu_{\text{NH}}^{\text{f}}$ modes of the two proton-transferred structures $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}1)$ and $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}2)$ and their intense $\nu_{\text{OH}}^{\text{b}}$ modes of the H-bonded W_2 are expected around 3500 cm^{-1} (Fig. S13, ESI†). For the $5\text{HIH}^+-\text{W}$ cluster, the population of the OH-bound oxonium isomer $s5\text{HIH}^+(\text{O})-\text{W}(\text{OH})$ was most likely enhanced by its high binding energy of $D_0 = 99.3 \text{ kJ mol}^{-1}$ and the twofold degeneracy. However, degeneracy effects no longer promote the most stable $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{OH})$ cluster or the formation of W_2 chains in $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}1/\text{W}2)$. Thus, our spectra do not indicate proton transfer for any of the $5\text{HIH}^+-\text{W}_n$ clusters at $n = 2$.

3.5 $5\text{HIH}^+-\text{W}_3$

In $s/a5\text{HIH}^+-\text{W}_3$, interior ion solvation competes with water network formation. W_3 chains at either one of the functional groups (NH-W-W and OH-W-W) are comparably strong as bifurcated W_3 H-bonds (W-NH-W and W-OH-W). Selected structures of $s5\text{HIH}^+-\text{W}_3$ are shown in Fig. 4. Additional $s5\text{HIH}^+-\text{W}_3$ and selected $a5\text{HIH}^+-\text{W}_3$ isomers are depicted in Fig. S15 and S16 (ESI†). Corresponding structural and spectroscopic data are listed in Table S1 (ESI†). Again, we observe a strong modulation in the acidity of the functional groups depending on the protonation site. As the NH group is more acidic in $s5\text{HIH}^+(\text{C}3)$, formation of W_3 networks at the NH site ($D_0 \approx 141 \text{ kJ mol}^{-1}$) is favored over the OH site ($D_0 \approx 119 \text{ kJ mol}^{-1}$). The bifurcated W-NH-W H-bond ($D_0 = 141.9 \text{ kJ mol}^{-1}$) is slightly favored over the linear NH-W-W chain ($D_0 = 140.8 \text{ kJ mol}^{-1}$). However, interior ion solvation with one water molecule attached to the OH group and two to the NH group yields the most stable $s5\text{HIH}^+(\text{C}3)-\text{W}_3(\text{OH}-\text{NH}-\text{W})$ isomer with $D_0 = 148.4 \text{ kJ mol}^{-1}$. In contrast, W_3 networks at the OH site are preferred for $s5\text{HIH}^+(\text{C}4)$ with $D_0 \approx 142-145 \text{ kJ mol}^{-1}$ (Table S1, ESI†). Solvation of both functional groups yields $s5\text{HIH}^+(\text{C}4)-\text{W}_3(\text{OH}-\text{W}-\text{NH})$ and $s5\text{HIH}^+(\text{C}4)-\text{W}_3(\text{OH}-\text{NH}-\text{W})$ with $D_0 = 148.9$ and $143.4 \text{ kJ mol}^{-1}$ (Fig. 4). Noncooperative effects destabilize the individual W_2 chains. For example, comparing $s5\text{HIH}^+(\text{C}4)-\text{W}_2(\text{OH}-\text{W})$ and $s5\text{HIH}^+(\text{C}4)-\text{W}_3(\text{OH}-\text{W}-\text{NH})$, the H-bonds within the W_2 chain at the OH group elongate from $R = 1.560/1.733$ to $1.580/1.744 \text{ \AA}$. The same trend is observed for all four interior solvated $s5\text{HIH}^+-\text{W}_3$ isomers (Fig. 4 and Table S1, ESI†). In line with the weakening of the individual H-bonds, for $n = 3$ (interior solvation) the impact on the $s5\text{HIH}^+$ core is also weaker than for $n = 2$ (water chain).

This directly translates into blue-shifts of the affected proton donor XH stretches. For instance, $\nu_{\text{NH}}^{\text{b}}$ of $s5\text{HIH}^+(\text{C}3)-\text{W}_2(\text{NH}-\text{W})$ and $s5\text{HIH}^+(\text{C}3)-\text{W}_3(\text{OH}-\text{NH}-\text{W})$ are predicted at 2787 and 2834 cm^{-1} , respectively.

Solvation of both OH groups of the oxonium ion yields $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{OH}-\text{OH}-\text{W}1)$ and $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{OH}-\text{OH}-\text{W}2)$ isomers with $D_0 = 229.2$ and $231.5 \text{ kJ mol}^{-1}$, respectively. The latter is again stabilized by its additional interaction with the aromatic π -electron cloud. A W_2 chain is attached to one OH site, significantly elongating the affected OH bond ($r_{\text{OH}} = 1.068$ and 1.104 \AA). The $\text{OH} \cdots \text{W}$ bond is still longer ($R = 1.391$ and 1.330 \AA). Hence, in contrast to what has been predicted for the corresponding $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}1)$ and $s5\text{HIH}^+(\text{O})-\text{W}_2(\text{OH}-\text{W}2)$ clusters, the proton is not transferred due to the noncooperative character of this W_3 binding motif. However, proton-transferred structures cannot be neglected for $n = 3$. We find four proton-transferred structures, namely, $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{NH}-\text{OH}-\text{W}1/\text{W}2)$ with a W_2 chain at one OH group and a single W at the NH group as well as $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{OH}-\text{W}-\text{W}1/\text{W}2)$ with a W_3 chain at its OH group (Fig. 4 and Fig. S15, ESI†). Interestingly, $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{OH}-\text{W}-\text{W}2)$ is the most stable structure found with $D_0 = 231.8 \text{ kJ mol}^{-1}$. The W_3 chain is entirely detached from the OH group ($r_{\text{OH}} = 1.763$ and $R_{\text{OH}-\text{W}} = 0.985 \text{ \AA}$), and is arranged over the phenol ring of neutral 5HI (Fig. 4). This isomer is distinguished by its $\nu_{\text{OH}}^{\text{f}}$ and $\nu_{\text{NH}}^{\text{f}}$ predicted at 3633 and 3497 cm^{-1} , and three intense $\nu_{\text{OH}}^{\text{b(W)}}$ at 3300, 3243, and 2806 cm^{-1} . The other three structures are significantly less stable with $D_0 = 201.9$, 208.8, and 216.5 kJ mol^{-1} , respectively.

Fig. 6 compares the IRPD spectrum of $5\text{HIH}^+-\text{W}_3$ to the calculated IR spectra of selected $s5\text{HIH}^+-\text{W}_3$ isomers. Spectra calculated for some additional $s/a5\text{HIH}^+-\text{W}_3$ conformers are given in Fig. S17 (ESI†). The triplet A, G, and B at 3725, 3692, and 3637 cm^{-1} unambiguously reveals the predominance of chainlike W_2 and/or W_3 solvation structures. Band A is thus assigned to ν_3 , band B to ν_1 , and band G to $\nu_{\text{OH}}^{\text{f(W)}}$ of the single-donor water molecules. Candidates to explain this pattern are at least the isomers considered in Fig. 6, except for $s5\text{HIH}^+(\text{C}3)-\text{W}_3(\text{W}-\text{NH}-\text{W})$ with a bifurcated H-bond. However, a clear isomer assignment is challenging, because the IRPD spectrum is not well resolved below 3600 cm^{-1} , possibly due to overlapping transitions of several isomers. Still, we can draw some conclusions from the comparison with the IRPD spectra of $5\text{HIH}^+-\text{W}_{1/2}$ (Fig. 1). First, band X at 3508 cm^{-1} is considered to be a contamination band. It is not convincingly rationalized by any calculated mode and it occurs at the same position as band X in the spectrum of $5\text{HIH}^+-\text{W}-\text{N}_2$ (ESI†). Yet, one may argue that $\nu_{\text{NH}}^{\text{f}}$ of $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{NH}-\text{OH}-\text{W}1/\text{W}2)$ predicted at 3497/3500 cm^{-1} gives rise to band X. However, already for $5\text{HIH}^+-\text{W}_2$ it remains unclear whether oxonium clusters are probed. Any other modes of $s5\text{HIH}^+(\text{O})-\text{W}_3(\text{NH}-\text{OH}-\text{W}1/\text{W}2)$ are also not clearly observed. Second, bands H and J at 3360 and 3230 cm^{-1} can be related to corresponding transitions in the IRPD spectra of $5\text{HIH}^+-\text{W}_{1/2}$. In analogy to the spectrum of $5\text{HIH}^+-\text{W}_2$, band H may be assigned to $\nu_{\text{OH}}^{\text{b(W)}}$ of the solvated water and/or $\nu_{\text{OH}}^{\text{b}}$ of the $5\text{HIH}^+(\text{C}3)$ protomer. Corresponding transitions are predicted at 3225 and 3301 cm^{-1} for $s5\text{HIH}^+(\text{C}3)-\text{W}_3(\text{OH}-\text{NH}-\text{W})$. In the



Table 2 Observed frequencies of the proton donor stretching modes (ν_{XH} , X = N/O, in cm^{-1}) of 5HIH^+ and PhH^+ and corresponding complexation-induced frequency red shifts ($\Delta\nu_{\text{XH}}$) in $5\text{HIH}^+/\text{PhH}^+-\text{L}$ clusters (L = Ar, N_2 , W)

Isomer	ν_{XH}	$\Delta\nu_{\text{XH}}$		
		L = Ar	L = N_2	L = W
$\text{PhH}^+(\text{O})$	3552 (X = O) ^a	-18 ^a	-9 ^a	+33 ^d
	3477 (X = O) ^a	-148 ^a	-440 ^a	> -877 ^d
$\text{PhH}^+(\text{o/p})$	3554 (X = O) ^b	-61	-146	-654
$s/5\text{HIH}^+(\text{O})$	3555 (X = O) ^c	-20	-15	+36
	3480 (X = O) ^c	-115	-335	-1304 ^c
$s5\text{HIH}^+(\text{C}3)$	3503 (X = N)	Not observed	Not observed	Not observed
	3635 (X = O)	Not observed	Not observed	Not observed
$a5\text{HIH}^+(\text{C}4)$	3405 (X = N)	-40	-65	-390
	3598 (X = O)	-63	-118	-583
	3478 (X = N)	-24	-68	-348
$s5\text{HIH}^+(\text{C}4)$	3584 (X = O)	-49	-131	-569
	3478 (X = N)	-13	-68	-348

^a Values correspond to Ne-tagged $\text{PhH}^+(\text{O})-\text{Ne}(\text{OH})$,²⁰ which closely approximate those of bare $\text{PhH}^+(\text{O})$. Complexation-induced red-shifts are calculated relative to $\text{PhH}^+(\text{O})-\text{Ne}(\text{OH})$. ^b Value corresponds to π -bonded $\text{PhH}^+(\text{o/p})-\text{Ar}(\pi)$,²⁰ which closely approximates that of bare $\text{PhH}^+(\text{o/p})$. ^c Values calculated at the B3LYP-D3/aug-cc-pVTZ level. ^d Values taken from ref. 14.

protomers for 5HIH^+ , namely, $s/a5\text{HIH}^+(\text{C}3)$, $s/a5\text{HIH}^+(\text{C}4)$, and $s/a5\text{HIH}^+(\text{O})$.

In the neutral ground state (S_0), NH- and OH-bound hydrates of both *s*- and *a*5HI rotamers could be identified by isomer-selective IR spectroscopy.⁵⁶ While neutral *a*5HI-W clusters are more stable by $\Delta E_0 > 1.1 \text{ kJ mol}^{-1}$, protonation reverses this trend, such that protonated *s*5HIH⁺-W are more stable by $\Delta E_0 > 1.5 \text{ kJ mol}^{-1}$. In the S_0 state, OH...W bonds are stronger than NH...W bonds. Upon C3-protonation the neutral NH...W bond is strengthened from $D_e = 40.2$ to 88.6 kJ mol^{-1} .⁵ The strengthening of the OH...W bond is less pronounced with $D_e = 43.3$ vs. 76.8 kJ mol^{-1} .⁵ These results were rationalized by the large charge density on the pyrrole ring of $5\text{HIH}^+(\text{C}3)$, which affects the acidity of the OH and NH groups. Our recent analysis of the NBO charge distribution in $5\text{HIH}^+(\text{C}3)$, $5\text{HIH}^+(\text{C}4)$, and $5\text{HIH}^+(\text{O})$ quantifies this qualitative argument.¹⁷

The IRPD spectrum of $5\text{HI}^+-\text{W}$ is depicted in Fig. 3. Number and positions of the bands (A-F) are similar to those observed for $5\text{HIH}^+-\text{W}$, indicating comparable structures of the 5HI core and similar microhydration motifs (OH...W and NH...W H-bonds). However, band widths and intensities differ significantly. The spectrum of $5\text{HI}^+-\text{W}$ was explained by the coexistence of $5\text{HI}^+-\text{W}(\text{OH})$ and $5\text{HI}^+-\text{W}(\text{NH})$.²³ Also in the cationic ground state (D_0), OH...W bonds are stronger than NH...W bonds, yet the interaction strengths are significantly enhanced compared to the neutral clusters. Hence, OH...W bonds are strongly preferred over NH...W bonds by a factor of 10.²³ A very rough estimate of the population of the $5\text{HIH}^+-\text{W}$ protomers yields that $5\text{HIH}^+(\text{C}3)-\text{W}(\text{NH})$ and $5\text{HIH}^+(\text{C}4)-\text{W}(\text{OH})$ contribute $\approx 70\%$, $5\text{HIH}^+(\text{C}4)-\text{W}(\text{NH}) \approx 20\%$, and $5\text{HIH}^+(\text{O})-\text{W}(\text{OH}) \approx 10\%$. Thus, due to the strong variation of the acidity of the NH and OH groups upon protonation, there is no longer any preference for OH...W or NH...W bonds. In the D_0 state, clusters of *s*5HI⁺ are more stable than those of *a*5HI⁺, and the same is true for the protonated species.

IRPD spectroscopy of $5\text{HI}^+-\text{W}_n$ clusters reveals the competition between interior ion solvation and the formation of H-bonded water networks.²³ For $5\text{HI}^+-\text{W}_2$, interior ion solvation at both acidic groups (OH/NH) is strongly preferred ($\geq 90\%$). For $5\text{HI}^+-\text{W}_3$, two isomers coexist which bear one single W and one W_2 dimer H-bonded to either of the functional groups. The IRPD spectrum of $5\text{HI}^+-\text{W}_3$ suggests a strong preference for attachment of the W_2 dimer at the OH group, leading to an estimated population ratio of 10:1 for OH/W/NH:OH/NH/W.²³ Proton transfer from $5\text{HI}^+-\text{W}_n$ to the solvent was not observed for $n \leq 3$. These results are in line with our current findings on the $5\text{HIH}^+-\text{W}_n$ clusters.

3.8 Comparison to PhH^+-W_n

It is instructive to compare our results for $5\text{HIH}^+-\text{W}_n$ to those obtained by IRPD spectroscopy of the related PhH^+-W_n clusters because (i) similar protonation mechanisms have been evidenced for PhH^+ , resulting in carbenium $\text{PhH}^+(\text{p/o})$ and oxonium $\text{PhH}^+(\text{O})$ ions,^{14,18-20,24} and (ii) as a consequence, the microhydration is expected to be similar. In our previous study,¹⁷ we compared the protonation of 5HIH^+ and PhH^+ and their micro-solvation by Ar and N_2 , revealing that the acidity of the OH group increases as $s5\text{HIH}^+(\text{C}3) < a5\text{HIH}^+(\text{C}4) < s5\text{HIH}^+(\text{C}4) < \text{PhH}^+(\text{p/o}) < s5\text{HIH}^+(\text{O}) < a5\text{HIH}^+(\text{O}) < \text{PhH}^+(\text{O})$.¹⁷ The acidity of the functional group(s) correlates with the H-bond strength and is crucial for proton transfer.

The IRPD spectra of PhH^+-W_n with $n \leq 5$ were interpreted by $\text{PhH}^+(\text{p/o})-\text{W}_n$ and $\text{PhH}^+(\text{O})-\text{W}_n$ clusters, for which proton transfer occurs at different critical sizes of the hydration shell (n_c). In the case of $\text{PhH}^+(\text{O})-\text{W}_n$, the critical size is determined as $n_c = 3$, and for $\text{PhH}^+(\text{p/o})-\text{W}_n$ as $n_c = 4$.¹⁴ Most likely, the transferred proton is the excess proton, coming from the OH_2 group of $\text{PhH}^+(\text{O})$ and the CH_2 group of $\text{PhH}^+(\text{p/o})$. However, for all PhH^+-W_n exclusively the OH group is solvated. Hence, the W_n network has to bridge the OH group and the protonated CH_2 group in $\text{PhH}^+(\text{p/o})$ which is only possible for $\text{W}_{n \geq 4}$ chains. Our current IRPD spectra of $5\text{HIH}^+-\text{W}_n$ do not indicate proton transfer at $n \leq 3$ for any of the assigned protomers. This finding is interesting as we also observe hydrated oxonium ions, $5\text{HIH}^+(\text{O})-\text{W}_n$. To elucidate the microsolvation mechanism, we compare the acidity of the OH groups of $\text{PhH}^+(\text{o/p})$ and $\text{PhH}^+(\text{O})$ to those of the observed $5\text{HIH}^+(\text{C}3)$, $5\text{HIH}^+(\text{C}4)$, and $5\text{HIH}^+(\text{O})$ clusters with the aid of the measured complexation-induced red-shifts of the OH stretch ($\Delta\nu_{\text{OH}}$) (Table 2). Most obviously, the acidity of the OH group of $5\text{HIH}^+(\text{C}3)$ is the smallest, because its ν_{OH} has the highest measured frequency (3635 cm^{-1}),¹⁷ and we do not observe OH...W bonds for $5\text{HIH}^+(\text{C}3)-\text{W}$. Preferentially, $s/a5\text{HIH}^+(\text{C}4)$ are solvated at the OH group, yet the observed $\Delta\nu_{\text{OH}}$ shifts are of medium size ($\Delta\nu_{\text{OH}} = -569/-583 \text{ cm}^{-1}$). The OH...W H-bond in $\text{PhH}^+(\text{o/p})-\text{W}$ is comparably strong with $\Delta\nu_{\text{OH}} = -654 \text{ cm}^{-1}$. Unfortunately, $\Delta\nu_{\text{OH}}$ has not been measured for $\text{PhH}^+(\text{O})-\text{W}$,¹⁴ but is estimated to be larger than -877 cm^{-1} . Hence, we determine the largest value of $\Delta\nu_{\text{OH}} = -1304 \text{ cm}^{-1}$ for $a/s5\text{HIH}^+(\text{O})-\text{W}$. Considering the red-shifts induced by attachment of Ar and N_2 at the OH group of $\text{PhH}^+(\text{O})$ and $5\text{HIH}^+(\text{O})$, we again infer a somewhat



larger acidity of the OH group of $\text{PhH}^+(\text{O})$.¹⁷ Hence, proton transfer may occur at $n_c \geq 4$ for $5\text{HIH}^+(\text{O})\text{-W}_n$.

4. Conclusions

Herein, we investigate the initial microhydration of a prototypical protonated heteroaromatic biomolecule using IRPD spectroscopy of size-selected $5\text{HIH}^+\text{-W}_n$ ($\text{W} = \text{H}_2\text{O}$, $n = 1\text{--}3$) clusters in the XH stretching range and calculations at the B3LYP-D3/aug-cc-pVTZ level. Our results may be summarized as follows.

We observe clusters of the C3- and C4-protonated carbenium ions, $5\text{HIH}^+(\text{C3})$ and $5\text{HIH}^+(\text{C4})$, and the oxonium ion, $5\text{HIH}^+(\text{O})$. Detection of $5\text{HIH}^+(\text{O})\text{-W}_n$ clusters is surprising at first glance because they are significantly less stable ($\Delta E_0 > 75 \text{ kJ mol}^{-1}$). However, the H-bonds to the protonated OH_2 group are very strong ($D_0 > 99 \text{ kJ mol}^{-1}$). In line with our previous results obtained for $5\text{HIH}^+\text{-L}$ with $\text{L} = \text{Ar}$ and N_2 ,¹⁷ $5\text{HIH}^+(\text{O})\text{-W}_n$ clusters benefit from the strong $\text{OH}\cdots\text{W}$ bond and the twofold degeneracy of the corresponding minimum. At the current spectral resolution, *syn* and *anti* rotamers (*s*- and *a* 5HIH^+) are not distinguishable. As the energy differences between their clusters are rather small ($\Delta E_0 = 1\text{--}5 \text{ kJ mol}^{-1}$), we assume the production of both *s*- and *a* $5\text{HIH}^+\text{-W}_n$ clusters.

$5\text{HIH}^+\text{-W}_n$ grow by H-bonding of the first W ligand to the acidic NH and OH groups, and π -stacking of W is unlikely. The absolute and relative strengths of the $\text{NH}\cdots\text{W}$ and $\text{OH}\cdots\text{W}$ H-bonds strongly depend on the 5HIH^+ protomer. The strongest H-bond is found in $5\text{HIH}^+(\text{O})\text{-W}(\text{OH})$ with W attached to one of its OH groups with an outstandingly high binding energy of $D_0 = 99.3 \text{ kJ mol}^{-1}$. The acidity of the OH group (NH group) increases (decreases) in the order $5\text{HIH}^+(\text{C3}) < 5\text{HIH}^+(\text{C4}) < 5\text{HIH}^+(\text{O})$.¹⁷ Thus, we predominantly observe $5\text{HIH}^+(\text{C3})\text{-W}(\text{NH})$, both $5\text{HIH}^+(\text{C4})\text{-W}(\text{NH})$ and $5\text{HIH}^+(\text{C4})\text{-W}(\text{OH})$, and $5\text{HIH}^+(\text{O})\text{-W}(\text{OH})$ clusters. IRPD spectra of tagged $5\text{HIH}^+\text{-W-Ar/N}_2$ clusters confirm this assignment. Interior ion solvation and formation of water networks compete for $5\text{HIH}^+\text{-W}_2$. We assign carbenium clusters with both functional groups solvated, $s5\text{HIH}^+(\text{C3})\text{-W}_2(\text{NH-OH})$ and $s5\text{HIH}^+(\text{C4})\text{-W}_2(\text{NH-OH})$, and those with W_2 water chains at their respective most acidic functional group, $s5\text{HIH}^+(\text{C3})\text{-W}_2(\text{NH-W})$ and $s5\text{HIH}^+(\text{C4})\text{-W}_2(\text{OH-W})$. The IRPD spectrum of $5\text{HIH}^+\text{-W}_2$ does not clearly show features of any $5\text{HIH}^+(\text{O})\text{-W}_2$ clusters. The spectrum of $5\text{HIH}^+\text{-W}_3$ clearly indicates W_2 water chains at the NH and OH groups. It can readily be explained by the two most stable carbenium ions, $s5\text{HIH}^+(\text{C3})\text{-W}_3(\text{OH-NH-W})$ and $s5\text{HIH}^+(\text{C4})\text{-W}_3(\text{OH-W-NH})$. Isomer $s5\text{HIH}^+(\text{C3})\text{-W}_3(\text{W-NH-W})$ with a bifurcated H-bond may also contribute to the measured spectrum. Again, we exclude the oxonium $5\text{HIH}^+(\text{O})\text{-W}_3$ clusters. Future IR-UV or IR-IR double resonance spectroscopy may facilitate disentangling the isomer contribution to the IRPD spectrum of $5\text{HIH}^+\text{-W}_n$.

Compared to the growth of $5\text{HIH}^+\text{-L}_n$ clusters ($\text{L} = \text{Ar}$, N_2), H-bonding is strongly preferred for $\text{L} = \text{W}$, and W_n networks compete with interior ion solvation, which is not the case for $\text{L} = \text{Ar/N}_2$. The strength of individual H-bonds increases in the order $\text{Ar} < \text{N}_2 < \text{W}$ as shown by comparison of the respective

complexation-induced frequency red-shifts ($\Delta\nu_{\text{XH}}$) of the corresponding H-bonded proton donor stretching modes.

Protonation significantly strengthens the $\text{OH}\cdots\text{W}$ and $\text{NH}\cdots\text{W}$ H-bonds observed in neutral 5HI-W due to the excess charge.⁵ The distribution of the excess positive charge is very different in the individual protomers directly affecting the acidity of the OH and NH groups. In the neutral S_0 ground state, *a* 5HI-W clusters are more stable than *s* 5HI-W .⁵⁶ In contrast, ionization into the D_0 state and protonation reverse the relative stability of the rotamers.^{22,23} The mechanism of the growth of the initial solvation shell ($n < 4$) is very similar for cationic and protonated hydrates.

IRPD spectra of the PhH^+ subunit of 5HIH^+ revealed proton transferred at critical sizes $n_c = 3$ and 4 in the case of $\text{PhH}^+(\text{O})\text{-W}_n$ and $\text{PhH}^+(\text{o/p})\text{-W}_n$, respectively.¹⁴ In contrast, we do not observe any clear characteristics of proton-transferred structures in the $5\text{HIH}^+\text{-W}_n$ clusters up to $n = 3$. Yet, we observe an increase in the acidity of the OH group in the order $5\text{HIH}^+(\text{C3}) < 5\text{HIH}^+(\text{C4}) < \text{PhH}^+(\text{p/o}) < 5\text{HIH}^+(\text{O}) < \text{PhH}^+(\text{O})$. Hence, proton transfer may occur at $n_c \geq 4$ for $5\text{HIH}^+(\text{O})\text{-W}_n$.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This study was supported by Deutsche Forschungsgemeinschaft (DO 729/3). J. K. is grateful for a fellowship of the Studienstiftung des deutschen Volkes.

References

- 1 A. T. Balaban, D. C. Oniciu and A. R. Katritzky, *Chem. Rev.*, 2004, **104**, 2777–2812.
- 2 J. M. Berg, J. L. Tymoczko and L. Stryer, *Biochemistry*, Freeman, New York, 2002.
- 3 J. P. Schermann, *Spectroscopy and modeling of biomolecular building blocks*, Elsevier, 2007.
- 4 H.-S. Andrei, N. Solcà and O. Dopfer, *ChemPhysChem*, 2006, **7**, 107–110.
- 5 R. Omidyan, M. Omidyan and A. Mohammadzadeh, *RSC Adv.*, 2016, **6**, 33148–33158.
- 6 E. P. L. Hunter and S. G. Lias, *J. Phys. Chem.*, 1998, **27**, 413–656.
- 7 D. J. Goebbert and P. G. Wenthold, *Eur. J. Mass Spectrom.*, 2004, **10**, 837–845.
- 8 M. Miyazaki, A. Fujii, T. Ebata and N. Mikami, *Chem. Phys. Lett.*, 2004, **399**, 412–416.
- 9 E. S. Kryachko and M. T. Nguyen, *J. Phys. Chem. A*, 2001, **105**, 153–155.
- 10 T. C. Cheng, B. Bandyopadhyay, J. D. Mosley and M. A. Duncan, *J. Am. Chem. Soc.*, 2012, **134**, 13046–13055.
- 11 I. Alata, M. Broquier, C. Dedonder-Lardeux, C. Jouvét, M. Kim, W. Y. Sohn, S. Kim, H. Kang, M. Schütz and A. Patzer, *et al.*, *J. Chem. Phys.*, 2011, **134**, 74307.



- 12 O. Dopfer, A. Patzer, S. Chakraborty, I. Alata, R. Omidyan, M. Broquier, C. Dedonder and C. Jouvet, *J. Chem. Phys.*, 2014, **140**, 124314.
- 13 O. Dopfer and M. Fujii, *Chem. Rev.*, 2016, **116**, 5432–5463.
- 14 M. Katada and A. Fujii, *J. Phys. Chem. A*, 2018, **122**, 5822–5831.
- 15 K. Tanabe, M. Miyazaki, M. Schmies, A. Patzer, M. Schütz, H. Sekiya, M. Sakai, O. Dopfer and M. Fujii, *Angew. Chem., Int. Ed.*, 2012, **51**, 6604–6607.
- 16 M. Ataelahi and R. Omidyan, *J. Phys. Chem. A*, 2013, **117**, 12842–12850.
- 17 J. Klyne and O. Dopfer, *J. Phys. Chem. B*, 2018, **122**, 10700–10713.
- 18 N. Solcà and O. Dopfer, *Chem. Phys. Lett.*, 2001, **342**, 191–199.
- 19 N. Solcà and O. Dopfer, *J. Chem. Phys.*, 2004, **120**, 10470–10482.
- 20 N. Solcà and O. Dopfer, *J. Am. Chem. Soc.*, 2004, **126**, 1716–1725.
- 21 S. A. Nizkorodov, O. Dopfer, T. Ruchti, M. Meuwly, J. P. Maier and E. J. Bieske, *J. Phys. Chem.*, 1995, **99**, 17118–17129.
- 22 J. Klyne and O. Dopfer, *J. Mol. Spectrosc.*, 2017, **337**, 124–136.
- 23 J. Klyne, M. Miyazaki, M. Fujii and O. Dopfer, *Phys. Chem. Chem. Phys.*, 2018, **20**, 3092–3108.
- 24 N. Solcà and O. Dopfer, *J. Chem. Phys.*, 2004, **121**, 769–772.
- 25 O. Dopfer, *Int. Rev. Phys. Chem.*, 2003, **22**, 437–495.
- 26 O. Dopfer, *Z. Phys. Chem.*, 2005, **219**, 125–168.
- 27 M. J. Frisch, *et al.*, *GAUSSIAN09, Rev. D.01*, Gaussian, Inc., Wallingford, CT, 2009.
- 28 J. Klyne, A. Bouchet, S. Ishiuchi, M. Fujii, M. Schneider, C. Baldauf and O. Dopfer, *Phys. Chem. Chem. Phys.*, 2018, **20**, 28452–28464.
- 29 W. Fu and W. S. Hopkins, *J. Phys. Chem. A*, 2017, **122**, 167–171.
- 30 M. Rossi and C. Baldauf, *J. Phys.: Condens. Matter*, 2015, **27**, 493002.
- 31 A. D. Becke, *Phys. Rev. A: At., Mol., Opt. Phys.*, 1988, **38**, 3098.
- 32 A. D. Becke, *J. Chem. Phys.*, 1993, **98**, 5648–5652.
- 33 C. Lee, W. Yang and R. G. Parr, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1988, **37**, 785.
- 34 T. H. Dunning, *J. Chem. Phys.*, 1989, **90**, 1007–1023.
- 35 M. Miyazaki, A. Naito, T. Ikeda, J. Klyne, K. Sakota, H. Sekiya, O. Dopfer and M. Fujii, *Phys. Chem. Chem. Phys.*, 2018, **20**, 3079–3091.
- 36 M. Schütz, Y. Matsumoto, A. Bouchet, M. Öztürk and O. Dopfer, *Phys. Chem. Chem. Phys.*, 2017, **19**, 3970–3986.
- 37 Q. Zhang and L. Du, *Comput. Theor. Chem.*, 2016, **1078**, 123–128.
- 38 H. Zhao, J. Chang and L. Du, *Comput. Theor. Chem.*, 2016, **1084**, 126–132.
- 39 S. Grimme, J. Antony, S. Ehrlich and H. Krieg, *J. Chem. Phys.*, 2010, **132**, 154104.
- 40 M. Miyazaki, A. Fujii and N. Mikami, *J. Phys. Chem. A*, 2004, **108**, 8269–8272.
- 41 K. Chatterjee and O. Dopfer, *Phys. Chem. Chem. Phys.*, 2017, **19**, 32262–32271.
- 42 B. E. Rocher-Casterline, L. C. Ch'ng, A. K. Mollner and H. Reisler, *J. Chem. Phys.*, 2011, **134**, 211101.
- 43 K. Chatterjee and O. Dopfer, *Chem. Sci.*, 2018, **9**, 2301–2318.
- 44 C. Baldauf, K. Pagel, S. Warnke, G. von Helden, B. Koks, V. Blum and M. Scheffler, *Chem. – Eur. J.*, 2013, **19**, 11224–11234.
- 45 V. Scutelnic, M. A. S. Perez, M. Marianski, S. Warnke, A. Gregor, U. Rothlisberger, M. T. Bowers, C. Baldauf, G. von Helden and T. R. Rizzo, *et al.*, *J. Am. Chem. Soc.*, 2018, **140**, 7554–7560.
- 46 M. Ropo, M. Schneider, C. Baldauf and V. Blum, *Sci. Data*, 2016, **3**, 160009.
- 47 S. Simon, M. Duran and J. J. Dannenberg, *J. Chem. Phys.*, 1996, **105**, 11024–11031.
- 48 S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553–566.
- 49 G. Herzberg, *Molecular spectra and molecular structure*, Read Books Ltd, 2013.
- 50 V. Barone, *J. Chem. Phys.*, 2004, **122**, 14108.
- 51 J. Klyne, M. Schmies, M. Miyazaki, M. Fujii and O. Dopfer, *Phys. Chem. Chem. Phys.*, 2018, **20**, 3148–3164.
- 52 J. Klyne, M. Schmies, M. Fujii and O. Dopfer, *J. Phys. Chem. B*, 2015, **119**, 1388–1406.
- 53 N. Solcà and O. Dopfer, *Chem. Phys. Lett.*, 2001, **347**, 59–64.
- 54 O. Couronne and A. Ellinger, *Chem. Phys. Lett.*, 1999, **306**, 71–77.
- 55 E. A. Colbourn and A. E. Douglas, *J. Chem. Phys.*, 1976, **65**, 1741–1745.
- 56 T. Ikeda, K. Sakota and H. Sekiya, *J. Phys. Chem. A*, 2016, **120**, 1825–1832.

