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Halogen bonds of halonium ions

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Due to their electron deficiency, halonium ions act as particularly strong halogen bond donors. By accepting electrons in both lobes of their empty p-orbital, they are capable of simultaneously interacting with two Lewis bases. The interaction presumes the formation of three molecular orbitals and is accordingly typically entitled as a three-center halogen bond. In analogy to the [D–H–D]⁺ hydrogen bonds, which are at times entitled as short and strong bonds, the [D–X–D]⁺ halogen bonds of halonium ions show Bondi normalized interatomic distances of 0.6–0.7 and possess both charge transfer and electrostatic characteristics. The three-center halogen bond of halonium ions shows distinct differences in its properties from coordinative bonds of transition metals and is therefore applicable as a complementary synthon in supramolecular chemistry. The three-center halogen bond modulates the reactivity of halonium ions and is hence a useful tool for synthetic organic chemistry. Following the discussion of the nature and properties of halonium ions' halogen bonds, this tutorial review provides an overview of their current applications to stimulate future developments.

Introduction

Halogen bonding (XB) is the directional, non-covalent interaction of an electron-poor region of a halogen and a Lewis base.¹ It

resembles hydrogen bonding and has accordingly been proposed to be applicable as a complementary tool for the rational modulation of molecular recognition events in chemistry and biology.² Being a weak interaction, halogen bonding is typically difficult to detect, especially in disordered phases.³ The past two decades of halogen bonding research has consequently focused on the investigation of strong and thus

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Fig. 1 Schematic description of (a) conventional and (b) three-center halogen bonds. The former halogen, X, is neutral and is covalently bound, typically to a carbon, and has a σ -hole that may receive electron density from a Lewis base, D. The latter, X^+ , in contrast, has an empty p-orbital and consequently a positive charge and two partially positively charged regions, p-holes. It may receive electron density from two Lewis bases simultaneously and thereby form a three-center halogen bond in which the $X \cdots D$ interactions possess both covalent and secondary characteristics.

comparably easy to detect complexes, in which a large electrophilic area on the halogen, typically called a σ -hole, has been promoted by perfluorination of the nearby carbon atoms (Fig. 1a).³ An alternative to this classical approach that involves a positively charged halogen, X^+ , formally a halonium ion or halogen(i), as a halogen bond donor has also been introduced (Fig. 1b) recently, yet so far has scarcely been discussed.⁴ In a conventional halogen bond,² $D_1-X \cdots D$, a neutral halogen, X, typically covalently attached to a carbon (D_1), interacts with an electron donor, D, Lewis base. In contrast, the halonium ions, X^+ , tend to form three-center halogen bonds, in which they simultaneously interact with two Lewis bases, each contributing a lone pair of electrons to the interaction.⁴⁻⁷ This charge transfer to the “p-holes”, the empty p-orbital of X^+ , results in the formation of a linear $[D_1 \cdots X \cdots D]^+$ arrangement.⁴ Here, the two Lewis bases, D and D_1 , may or may not be identical. The halogen and the two electron donors are held together by four electrons and thus form a three-center, four-electron bond that is remarkably short ($R_{XB} = 0.65-0.69$, where $R_{XB} = d_{XB}/(X_{vdw} + B_{vdw})$)^{6,8} and strong (up to 180 kJ mol^{-1}).^{3,9}

Similar to most halogen bond complexes,² these were initially often characterized as charge-transfer complexes.¹⁰

The main aim of this tutorial review is to summarize and interpret the past decade's studies of halonium ions' halogen bonding, review its utility in supramolecular chemistry and discuss its emerging applicability, for instance in synthetic chemistry. Despite its short history, the investigations of halonium ions' halogen bonds have provided unique insights into the fundamentals of chemical bonding. This bond has been proven to be among the strongest halogen bonds, and has consequently rapidly found applications.

The nature of halonium ions' halogen bonds

Out of the four halogens, iodine(i), bromine(i) and chlorine(i) are capable of forming three-center, four-electron bonds, $[D \cdots X \cdots D]^+$, whereas fluorine(i) has been demonstrated to prefer conventional halogen bonds, $D^+-F \cdots D$, with a distinct short and strong covalent D-F bond and a second, weaker and longer, $D \cdots F$ halogen bond (Fig. 2).⁸ Fluorine centered halogen bonds are rare and only form when fluorine is directly attached to a highly electron withdrawing functionality, such as positively charged nitrogen, which enhances its σ -hole. The former complexes may be formed with variety of Lewis bases including nitrogenous, sulphurous, selenious and tellurous electron donors, which may be of sp^3 , sp^2 or sp character.⁵

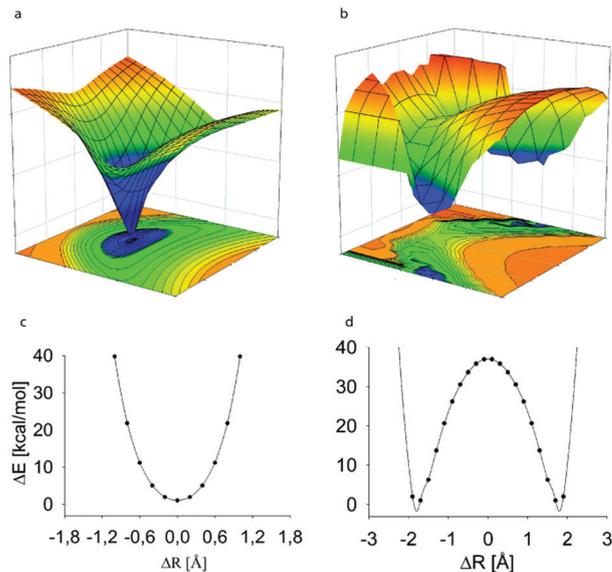


Fig. 2 The computed potential energy surface of the halogen-bonded complexes of halonium ions indicates the preference for (a) symmetric $[D \cdots X \cdots D]^+$ geometries for the complexes of iodonium, bromonium and chloronium ions, whereas (b) asymmetric $D^+-X \cdots D$ arrangement for the fluoronium ion complexes. The potential energy curves (c and d), that is the 2D-slices corresponding to the D–D distance with the global energy minimum of the complexes, possess a single energy minimum for the larger halogens, whereas that of the fluoronium complex shows two equal minima with a high energy barrier. Here, the potential energy is shown as a function of the donor–donor, r_{DD} , and the halogen–donor, D–X, distance, where ΔR is the difference between the D_1-X and D_2-X distances at a given r_{NN} . The data of [bis(pyridine)iodonium]⁺ and [bis(pyridine)fluoronium]⁺ complexes are shown as representative examples.⁸ This figure has been adapted from ref. 8 with permission from the Royal Society of Chemistry, copyright 2014.

A classical, two-center halogen bond is commonly seen to involve two atoms, namely the halogen, X, and the electron donor, D. Its halogen is in oxidation state 0 or formally -1 , and possesses two vastly different bonds: a conventional two-center covalent, D–X, and a classical two-center halogen bond, $D \cdots X$ (Fig. 1a). In contrast, the halogen of three-center, four-electron halogen bonds involves a halogen(i) in oxidation state $+1$, which forms two $D \cdots X$ bonds that simultaneously possess covalent and secondary bond characteristics (Fig. 1b). The $D \cdots X$ bond distances remain in between the sum of the van der Waals radii and the sum of the covalent radii of the participating atoms. They are similar in length, within $<10\%$ difference between the $D \cdots X$ distances. Breaking one of them typically destabilizes the second $D \cdots X$ bond as well, yielding a reactive halonium ion and two equivalents of Lewis bases.

Viewing from an orbital perspective, the three-center halogen bond is formed by the empty p-orbital of an X^+ simultaneously accepting electron density from two Lewis bases, with the valence shell of the central X^+ having an s^2p^4 electronic configuration.¹¹ Induced by the electrostatic field of the two Lewis bases, its p-orbitals are occupied in the spin paired $p_x p_y p_z$ arrangement. The two lobes of its empty p_z orbital are available for interaction with the electron pairs



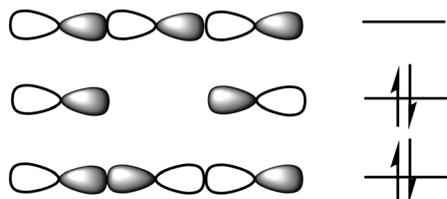


Fig. 3 The molecular orbital system of three-center, four-electron bonds, $[D \cdots X \cdots D]^+$. Two of the electrons are on the bonding orbital, whereas two are on the non-bonding level, and are thus predominantly located on the terminal electron donor atoms, which act as halogen bond acceptors. The central atom, here a halogen, is electron poor.

of two Lewis bases, D, in a linear $D \cdots X \cdots D$ fashion. These three atomic orbitals combine into three molecular orbitals that hold four electrons, and thus act as a three-center, four-electron system (Fig. 3). The three-center bond of halonium ions has been computationally thoroughly assessed,^{8,12–16} and concluded to be dominantly of covalent and electrostatic characteristics, with a smaller contribution from dispersion forces. The origin of the covalency is charge transfer, whereas the electrostatic characteristic is the result of Coulomb attraction and polarization. The partial covalent character of the three-center halogen bond increases as the size of the halogen decreases,^{8,13} and simultaneously, the strength of the bond decreases as the strength mainly originates from the electrostatic contribution (Fig. 4). This trend has been demonstrated both experimentally and computationally for $[N \cdots X \cdots N]^+$ systems, for which the chloronium-centered bond was observed to be the weakest despite its largest covalent character, whereas the least covalent idonium-centered system had the strongest halogen bond.⁸ The correlation of covalency and the three-center character of halogen bonding has been investigated at the CCSD(T) level of theory.¹⁶

Three-center halogen bond complexes have been proposed to be termed “coordination complexes”^{5,12,17} to emphasize the dative nature of their ionic bond.⁵ It has also been suggested that only the bond of I^+ should be termed coordinative, whereas that of Br^+ and Cl^+ should rather be seen as bonds with covalent character, and the analogous complexes of F^+ considered as ion–molecule complexes.¹² A combined NMR, X-ray diffraction and computational study has, however, indicated that I^+ complexes behave differently to typical transition metal complexes, such as those of $Ag(I)$,¹⁸ regarding solvation and counterion coordination. Therefore, the term “coordination complex” for

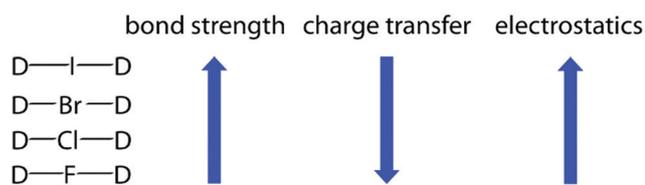


Fig. 4 The halogen dependence of the nature and characteristics of the three-center halogen bond: the larger the halogen, the stronger and the more electrostatic it is. With a smaller halogen, the bond is shorter and more covalent.

such halogen bond complexes is misleading and should not be used.¹⁹

$[D \cdots X \cdots D]^+$ compounds may also be interpreted as hypercoordinate, hypervalent, or 10-X-2 complexes of halogen(i).²⁰ Historically hypervalent bonding has been considered to involve the expansion of the octet valence electrons, whether through σ -hole interactions, electrostatic interactions or charge transfer, but this expansion is now viewed as being largely formal. The hypervalent bonding model has the advantage of allowing a smooth comparison of the structure and reactivity of electronically related complexes of different research fields.

Bond symmetry

The three-center halogen bond is isoelectronic to the $[D-H-D]^+$ hydrogen bond,⁴ as both have 4 electrons shared over 3 atomic centers. The three-center hydrogen bond has been postulated to be symmetric, to possess unusually short D–H bond length and to have exceptional bond strength. Accordingly, it is often entitled as a short, strong, or low barrier bond. The strength and symmetry of such hydrogen bonds has, however, been controversial. Using solution NMR, Perrin has shown that such hydrogen bonds are typically asymmetric in solution and do not possess unusual bond strength.²¹ The interpretation of Perrin’s NMR data has been questioned, for example, by Singleton.²² Whereas the nature of the $[D-H-D]^+$ hydrogen bonds has been debated for decades, studies of the analogous $[D-X-D]^+$ halogen bond have been initiated within the last 10 years.^{9,23}

A three-center halogen bond may prefer (a) a static and centrosymmetric, $[D \cdots X \cdots D]^+$ geometry with the central halogen forming two equally strong and long D \cdots X bonds (Fig. 5). The motion of the halogen in such a system is best described by a single-well energy potential. Alternatively, the halogen bond of halonium ions may be asymmetric, $[D-X \cdots D]^+$, with the halogen forming one shorter and stronger D–X bond, and a weaker and longer D \cdots X bond to the other donor. Compared to the centrosymmetric geometry, the covalent character of the D–X and D \cdots X bonds of the asymmetric complex is increased and decreased, respectively. The motion of the halogen in such a system may be described by (b) a symmetric double-well potential, (c) an asymmetric single-well potential, or (d) an asymmetric double-well potential. Analogous to the term prototropy, which has been coined to describe tautomeric exchange processes involving H^+ transfer, the motion of X^+ in such halogen-bonded systems is called halotropy.²⁴ A symmetric double-well potential (b) reflects a pair of asymmetric isomers, $[D-X \cdots D]^+$ and $[D \cdots X-D]^+$. When the Lewis basicity of the electron donors is comparable and the asymmetric geometries are separated by a shallow energy barrier, these may interconvert and exist as a dynamic mixture, $[D-X \cdots D]^+ \rightleftharpoons [D \cdots X-D]^+$. A system described by a symmetric double-well potential with a high energy barrier results in a static asymmetric geometry, $D^+-X \cdots D$, which corresponds to a conventional weak σ -hole halogen bond complex. If the electron donors have slightly different Lewis basicities, the complex may become static and asymmetric (c), with the halogen showing a higher preference to one of the electron donors.



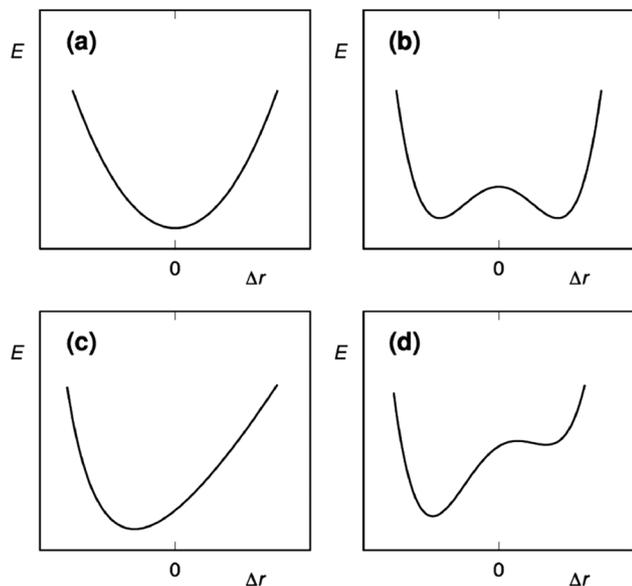


Fig. 5 The possible energy potentials of halogen motion in a three-center $[D \cdots X \cdots D]^+$ halogen bond: (a) a symmetric single-well energy potential describes the halogen motion as a static and symmetric $[D \cdots X \cdots D]^+$ system, in which the $D \cdots X$ bonds are equal. A (b) symmetric double-well potential may either reflect a pair of asymmetric isomers, $[D \cdots X - D]^+ \rightleftharpoons [D - X \cdots D]^+$, in dynamic equilibrium, when the energy barrier between the minima is shallow. Each form of this geometry has a shorter and stronger $D-X$ and a longer and weaker $D \cdots X$ bond. If the energy barrier between the two minima is high, the system becomes static and asymmetric. For systems possessing Lewis bases, D , with different electron densities, the halogen motion may either follow (c) an asymmetric single-well potential, or (d) an asymmetric double-well potential. The latter has a strong preference for a shorter and stronger bond towards one of the electron donors. The potential energy variation is shown here as a function of Δr , the displacement of X^+ from the symmetrical position.²⁴ This figure has been adapted from ref. 24 with permission from the American Chemical Society, copyright 2018.²⁵

Its motion between the electron donors is described by an asymmetric single-well. Alternatively, it forms asymmetric geometries that are of different energy (d), and the halogen motion is then described by an asymmetric double-well potential, where one of the minima is of lower energy and of higher stability. The symmetry of the halogen bond of halonium ions in the solid state, in solution and *in silico* has been reviewed in 2015, using systems possessing nitrogen Lewis bases as a representative example.⁴ There has been significant progress since then, with several experimental and computational studies addressing the factors possibly influencing halogen bond symmetry.^{8,15,19,24,26}

Halogen bond symmetry has been studied on [bis(pyridine)-halogen(i)]⁺ complexes and their geometrically restrained [(1,2-bis(pyridin-2-ylethynyl)benzene)halogen(i)]⁺ analogues (Fig. 6). The bromine(i) and iodine(i) centered model systems prefer static, symmetric geometries in solution, as demonstrated by the lower temperature dependence of their secondary isotope effects, as compared to the analogous dynamic $[N-H \cdots N]^+ \rightleftharpoons [N \cdots H-N]^+$ hydrogen bond complexes, using the isotopic perturbation of equilibrium method of Siehl.²⁷ These results were

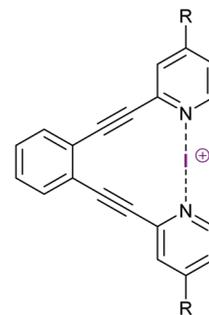


Fig. 6 [Bis(4-*R*-pyridine)iodine(i)]⁺ BF₄⁻, and geometrically restrained [1,2-bis((4-*R*-pyridine-2-ylethynyl)benzene)iodine(i)]⁺ BF₄⁻ complexes were used as model systems for assessment of the influence of electron density alterations on the $[N-I-N]^+$ halogen bond. The bis(pyridine)-type complex (left) permits free rotation and adjustment of $N-I$ distances for the most favorable interaction, whereas the 1,2-diethynylbenzene backbone of the complex (right) inhibits rotation around the $N-I-N$ axis and imposes some strain in the $N-I$ bonds to reach a geometrically optimal $[N-I-N]^+$ interaction. A systematic alteration of the electron density of the $[N-I-N]^+$ halogen bond was achieved by variation of the *R*-substituents, resulting in complexes with an increasing electron density in the order $NO_2 < CF_3 < H < F < Me < OMe < NMe_2$. Symmetric substitution led to centrosymmetric complexes with varying halogen bond strengths,²⁶ whereas asymmetric substitution led to static asymmetric complexes.²⁴

corroborated by ¹H, ¹³C and ¹⁵N NMR spectroscopy, and by computations at the DFT level.^{9,13} [Bis(pyridine)iodine(i)]⁺ and the analogous bromine(i) complexes are symmetric also in the solid state,^{4,7,19} as shown by X-ray diffraction, although these data have originally mistakenly been interpreted as displaying asymmetric structures⁹ due to the minor, <5%, difference in $N-X$ bond lengths. This difference is explained by crystal packing forces and is insignificant, and thus the X-ray data show symmetric complexes in the solid state. The analogous [bis(pyridine)chlorine(i)]⁺ complex was shown to also prefer a symmetric geometry in dichloromethane solution by DFT/MP2 studies (Fig. 7), supported by ¹H and ¹³C NMR studies.⁸ This complex is only stable below -70 °C, and undergoes rapid

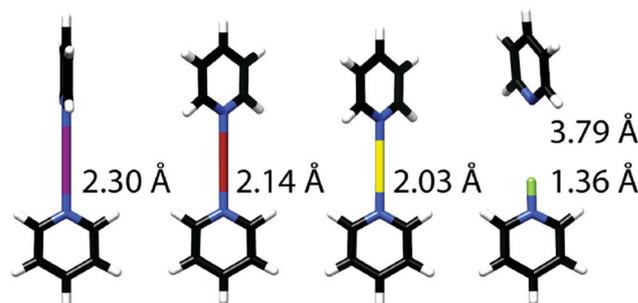


Fig. 7 The optimized (B3LYP/LANL08d) geometries of the I, Br, Cl and F-centred [bis(pyridine)halogen(i)]⁺ complexes, shown from left to right.⁸ Formation of the three-centred bond results in Bondi normalized interatomic distances 0.65 (N...I), 0.63 (N-Br) and 0.62 (N-Cl), with thus > 30% shortening of the sum of van der Waals radii of the involved atoms indicating the exceptional strengths of these halogen bonds. This figure has been adapted from ref. 8 with permission from the Royal Society of Chemistry, copyright 2014.



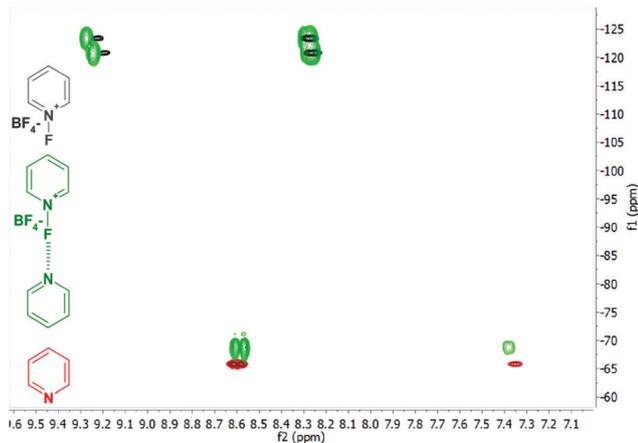


Fig. 8 The overlaid ^1H , ^{15}N HMBC of *N*-fluoropyridinium tetrafluoroborate (black), of pyridine (red) and of their halogen bond complex (green). Observation of -3 ppm ^{15}N coordination shift of the pyridine nitrogen, along with 0 ppm ^{15}N but 1.1 ppm ^{19}F coordination shift of the *N*-fluoropyridinium ion, indicates formation of a weak F-centred conventional halogen bond.⁸ The different chemical shifts of the two nitrogens of the complex are compatible with a static asymmetric $\text{N}^+-\text{F}\cdots\text{N}$ bond, which was also corroborated by DFT calculations. This figure has been adapted from ref. 8 with permission from the Royal Society of Chemistry, copyright 2014.

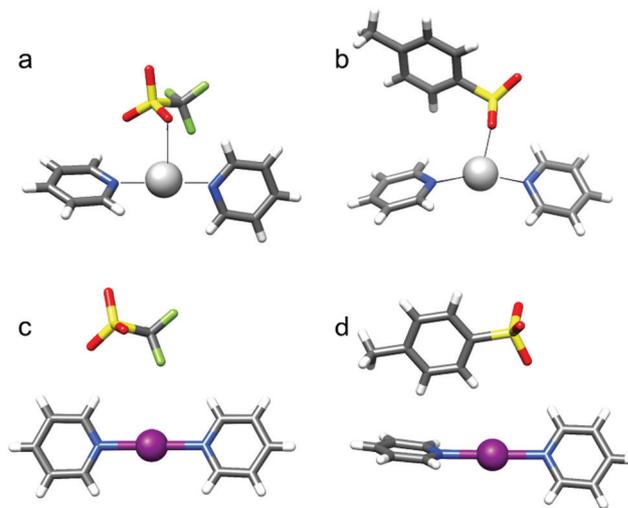


Fig. 9 The solid state structures of the [bis(pyridine)silver]⁺ (a and b) and [bis(pyridine)iodine(I)]⁺ (c and d) complexes possessing triflate (a and c) and tosylate (b and d), obtained by X-ray crystallography, reveal direct coordination of the counterion to silver(I) but not to iodine(I). The latter forms linear, bis-coordinated complexes that remain symmetric along the N–I–N three-center bond, even when the counterion is asymmetrically positioned. This figure has been adapted from ref. 26 with permission from the Royal Society of Chemistry, copyright 2016.

relaxation (T_1 0.08 s of the proton closest to the $[\text{N}\cdots\text{Cl}\cdots\text{N}]^+$ bond), preventing isotopic perturbation of equilibrium and ^{15}N NMR studies. Complex formation is corroborated by the quick relaxation of the nearby pyridine protons of the chlorine(I) complex (T_1 0.08 s) as compared to that of the corresponding Ag^+ (T_1 1.49 s) complex of pyridine (T_1 1.61 s). This quick relaxation of nearby protons to the chlorine may be explained by scalar relaxation of the second kind (SC2). The corresponding fluorine-centered bis(pyridine) complex was demonstrated to prefer a static asymmetric $\text{N}-\text{F}\cdots\text{N}$ geometry, with a strongly covalent $\text{N}-\text{F}$ bond and a weak, yet detectable, fluorine-centered $\text{N}\cdots\text{F}$ halogen bond, as demonstrated by diffusion NMR and by 2–3 ppm ^{15}N and ^{19}F NMR coordination shifts, *i.e.* chemical shift changes upon complexation (Fig. 8).⁸

The preference for a symmetric three-center bond was demonstrated to be independent of solvent polarity.¹¹ Hence, the iodine(I) and bromine(I)-centered complexes showed comparable ^{15}N NMR chemical shifts (2–6% alteration) in dichloromethane and acetonitrile solution,¹¹ whereas the corresponding [bis(pyridine)silver]⁺ complex, for example, showed 48% ^{15}N NMR shift alteration upon this solvent change.¹⁸ DFT (B3LYP/LANL08d) calculations indicated neither solvent dependence in the extent of charge transfer (28% positive charge on Br^+ and 41% on I^+ , respectively) or in bond lengths (2.14 and 2.30 Å), corroborating the experimental evidence.

Besides solvent polarity, the interaction with an asymmetrically positioned counterion (Fig. 9) has been proposed as a plausible inductor of halogen bond asymmetry.¹⁹ A solution NMR investigation, including isotopic perturbation of equilibrium, diffusion and ^{15}N NMR studies of iodine-centered complexes possessing spherical and weakly coordinating BF_4^- , ClO_4^- , PF_6^- , SbF_6^- ,

moderately coordinating OTf^- and OTs^- , and small and strongly coordinating NO_3^- and CF_3COO^- counterions, suggested that the preferred halogen bond symmetry is retained independent of the size, charge distribution or coordination strength of the counterion.¹⁹ These complexes showed a centrosymmetric structure in the solid state as well, with their geometry having been confirmed by DFT calculations.

It has further been studied whether the alteration of the electron density of the three-center halogen bond, other than upon changing the halogen,⁸ but by instead altering the electron density of the Lewis bases, may influence its geometry and strength.^{24,26} In case the Lewis bases involved have a low electron density, the central halogen(I) may be unable to efficiently accept electrons from both donors simultaneously and choose to compensate for its electron depletion by forming a strong covalent bond to one of the donors, and a weak halogen bond to the second, rather than forming a three-center halogen bond involving both. This would result in an asymmetric geometry. Alternatively, upon an increased electron density of the Lewis bases involved, the halogen(I) may prefer to form a strong bond to one electron donor, as it may better compensate for its electron depletion by connecting to one Lewis base covalently, than to two through a secondary bond, resulting in an asymmetric geometry. Changing the electron density by symmetric alteration of the pyridine *para*-hydrogen to NO_2 , CF_3 , F , Me , OMe , NMe_2 functionalities (Fig. 6), the centrosymmetric three-center halogen bond geometry is retained both in solution and in the solid state, as demonstrated by NMR, X-ray and DFT computations.²⁶ Whereas the bond symmetry, the N–I bond lengths and the ^{15}N NMR coordination shifts remained virtually unaltered, the $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ bond strength varied by



Table 1 N–I and N–N distances, N–I–N angles, ^{15}N NMR coordination shifts and DFT predicted stabilization energies for $[(1,2\text{-bis(pyridine-2-ylethynyl)benzene}iodine(I))^+]$ complexes shown in Fig. 6. Further details are given in ref. 26

4-R	$r(\text{NI})$ (Å)	$r(\text{NN})$ (Å)	$\angle(\text{NIN})$ ($^\circ$)	δ $^{15}\text{N}_{\text{coord}}$	ΔE_{stab} (kJ mol $^{-1}$)
H	2.3034	4.5934	175.7	−101.0	0
Me	2.3011	4.5875	175.4	−101.0	11.6
OMe	2.2982	4.5832	175.7	−100.0	39.8
NMe $_2$	2.2930	4.5715	175.4	−100.7	69.2
F	2.3027	4.5921	175.7	−101.4	16.9
CF $_3$	2.3041	4.5939	175.5	−106.0	−11.4
NO $_2$	2.3043	4.5952	175.7	N.A.	−19.1

~ 100 kJ mol $^{-1}$ upon altering the substituents, for both the $[\text{bis(pyridine}iodine(I))^+]$ and the $[\text{1,2-bis((pyridine-2-ylethynyl)benzene}iodine(I))^+]$ model systems (Table 1). This suggests that any general conclusions on the correlation of bond strength with bond lengths, and bond strength with NMR coordination shifts should be made with utmost caution.

Upon asymmetric modulation of the substituents of $[(1,2\text{-bis(pyridin-2-ylethynyl)benzene}iodine(I))^+]$, by introduction of a CH $_3$ and a CF $_3$ group in the *para*-position to the pyridine nitrogens (Fig. 6 and 10a), a static asymmetric $[\text{N-I} \cdots \text{N}]^+$ bond geometry (Fig. 5c) was obtained, with the halogen being closer to the nitrogen of higher electron density ($\Delta d_{\text{N-I}} = 4\%$).²⁴ This was demonstrated by the experimentally observed 20 ppm difference of the ^{15}N NMR coordination shifts of the two nitrogens, and by the low temperature dependence of the secondary deuterium isotope effects of the pyridine carbons, as compared to those of the reference model system involved in a dynamic equilibrium. This $[\text{N-I} \cdots \text{N}]^+$ complex is slightly destabilized (9.7 kJ mol $^{-1}$) as compared to its symmetric and unsubstituted analogue. The asymmetric three-center bond remains strong and the complex is stable in solution at room temperature for days. The above conclusions on the substituent effect were confirmed by an independent theoretical investigation.^{14,15}

In an analogous study,²⁹ the halogen position within a $[\text{D-X-D}]^+$ bond was assessed using pyridine complexes of *N*-halosuccinimides and *N*-halosaccharins (Fig. 10b–e). Due to the comparable Lewis basicity of imides and electron rich

pyridines, the position of X $^+$ (Fig. 6) is modulated by a slight adjustment of the electron density of the participating nitrogens. Co-crystallization of *N*-iodosuccinimide with 4-picoline and 4-dimethylaminopyridine (Fig. 10b and c) yielded complexes, in which the I $^+$ is closer to the nitrogen of succinimide whereas it is further away from the pyridine nitrogen ($\Delta d_{\text{N-I}} = 12\text{--}13\%$). As a consequence of the lower electron density of the saccharin anion as compared to the succinimide anion, the analogous *N*-iodosaccharin–4-dimethylaminopyridine complex showed a shorter N–I bond to the pyridine nitrogen and a longer bond to the imine nitrogen. Decreasing the electron density of the pyridine by NMe $_2$ to CH $_3$ substitution (Fig. 9d and e) inverted the geometry, with I $^+$ moving closer to the saccharin nitrogen ($\Delta d_{\text{N-I}} = 4\%$). It should be noted that the difference of N–I bond lengths of the latter two complexes, $\Delta d_{\text{N-I}} = 3\text{--}4\%$ (Fig. 10d and e), is comparable to that of the asymmetric $[(4\text{-methyl-2-}((2\text{-}((4\text{-}(\text{trifluoromethyl})\text{-pyridin-2-yl})\text{-ethynyl})\text{phenyl})\text{ethynyl})\text{pyridine}iodine(I))^+]$ complex (4%)²⁴ (Fig. 10a). Both N–I bonds of these complexes are undoubtedly part of a three-center complex, and thus possess both covalent and secondary characteristics.^{8,23} Formally, the shorter bonds can be seen to possess a “more covalent” whereas the longer a “more secondary” bond character. The central halogen of such complexes carries a partial positive charge; however, much of the formal charge of the halogen(I) is distributed into the conjugated electronic system of the coordinating Lewis bases.^{8,23} Such complexes may be assigned “more ionic” or “more neutral” character,²⁹ although the alteration of the charge distribution is minor. In addition to the Lewis basicity of the halogen bond acceptors, crystal packing forces were suggested to influence the position of the halogen for the *N*-iodosaccharin–pyridine complex.^{28,29} Interestingly, whereas the I $^+$ of the *N*-iodosaccharin–4-dimethylaminopyridine is slightly ($\Delta d_{\text{N-I}} = 4\%$) closer to the imide nitrogen, the Br $^+$ of the analogous *N*-bromosaccharin complex shows the opposite preference ($d_{\text{Nimide-Br}} = 2.231$ Å, $d_{\text{Npyridine-Br}} = 2.301$ Å, $\Delta d_{\text{N-I}} = 4\%$).²⁸ This was explained to be a consequence of the crystalline molecular environment that was computationally predicted to have a 26% larger influence on the Br $^+$ -centered complex. In agreement with previous literature (Fig. 4),⁸ the $[\text{N-Br} \cdots \text{N}]^+$ complex was shown (DFT) to have a larger covalent character than the corresponding $[\text{N-I} \cdots \text{N}]^+$ complex.²⁸

Halotropy, that is the low barrier $[\text{D-X} \cdots \text{D}]^+ \rightleftharpoons [\text{D} \cdots \text{X-D}]^+$ interconversion of asymmetric halogen bond geometries, has a fundamental importance not just for the understanding of the bonding phenomenon itself, but also for that of reactions involving halogen transfer. Electrophilic halogenations (Fig. 11), for instance, involve an analogous X $^+$ transfer by the initial weakening of a C–X, N–X or X–X covalent bond of the halogen transfer agent while forming a conventional halogen bond to an alkene. This is followed by formation of a three-center complex consisting of a nucleophile, a halogen(I) center and the leaving group. The three-center intermediate collapses to the product by forming a covalent halogen–nucleophile bond and eliminating the leaving group. The $[\text{D-X} \cdots \text{D}]^+$, $[\text{D} \cdots \text{X} \cdots \text{D}]^+$, and $[\text{D} \cdots \text{X-D}]^+$ species can be seen to represent the stages of any halogen transfer process from one halogen bond acceptor to another.

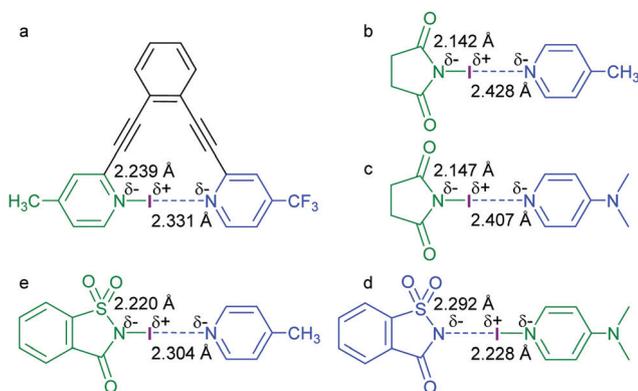


Fig. 10 The N–I distances (Å) of a variety of asymmetric N–I–N complexes, determined by DFT calculations and X-ray diffraction studies.^{24,28}



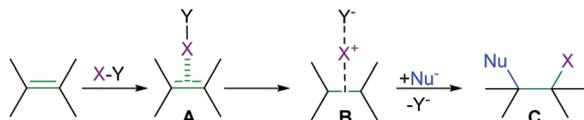


Fig. 11 The mechanism of electrophilic halogenation. The reaction is initiated by formation of a weak halogen bond with the alkene acting as a halogen bond acceptor (A). This is followed by formation of an intermediate (B) in which the central halogen possesses two halogen bonds, to Y- and to the π electrons of the double bond, and thus forms an analogous complex to a classical three-centre bond. The reaction reaches completion (C) by disconnection of the leaving group Y, which can be any group that binds a halogen X.

These have varying D–X bond lengths and possess a varying degree of covalency. Understanding whether a halogen transfer process between two halogen bond acceptors follows a single-well or double-well potential with a shallow barrier, whether it takes place at all, and whether the dynamic mixture of asymmetric geometries, the symmetric transition geometry or any of the asymmetric forms has the highest stability, is of fundamental interest.^{30–33}

To model the halogen transition between two halogen bond acceptors, and thus establish a dynamic mixture $[D \cdots X \cdots D]^+ \rightleftharpoons [D \cdots X \cdots D]^+$, the energy barrier between the two complexes must be shallow. This may be obtained with molecular systems that have a longer-than-optimal donor–donor distance, as predicted by DFT computations.²⁴ For $[N \cdots I \cdots N]^+$ -type systems, the global energy minimum is centrosymmetric and is at 4.51 Å nitrogen–nitrogen distance (Fig. 12). Upon increasing this distance above 4.88 Å, the asymmetric $[N \cdots I \cdots N]^+$ and $[N \cdots I \cdots N]^+$ arrangements become energetically more favorable than the alternative symmetric $[N \cdots I \cdots N]^+$ one. With an increasing nitrogen–nitrogen distance, the energy barrier of the $[N \cdots I \cdots N]^+ \rightleftharpoons [N \cdots I \cdots N]^+$ interconversion increases just as the energy of the most stable asymmetric geometries does, as compared to that of the geometry at electron donor–electron donor distance to a value above the global energy minimum ($r_{NN} = 4.51$ Å). A promising approach to obtain a model system in which an iodine can move between two halogen acceptors is thus to adjust the bifurcation point (4.88 Å), yet below the distance that would provide too high an energy barrier for interconversion of the asymmetric geometries. This has been confirmed by further computations of Scheiner, who used $H_3N \cdots X^+ \cdots NH_3$ and $H_3N \cdots X^+ \cdots OH_2$ complexes to estimate transfer barriers for various donor–donor distances in $[D \cdots X \cdots D]^+$ systems.³⁵ 2,2'-(9,10-Dimethoxyphenanthrene-3,6-diyl)dipyridine²⁴ was proposed as a bidentate ligand for a dynamic system; however, as a result of its suboptimally large N–N distance, 5.24 Å, this system was shown to be unstable as a monomeric complex and showed preference for the formation of a dimer instead, which encompasses two static and symmetric $[N \cdots I \cdots N]^+$ bonds.

Analogous elongation of the covalent bond of the interhalogens ICl (7%) and IBr (4%) was reported upon their halogen bonding to the sulfur of *N*-methylbenzothiazole-2-thione and 2(3)*H*-benzothiazole-thione (Fig. 13).³⁴ These complexes are slightly asymmetric ($\Delta d_{S-I/I-X} = 3\text{--}7\%$), with their S \cdots I halogen

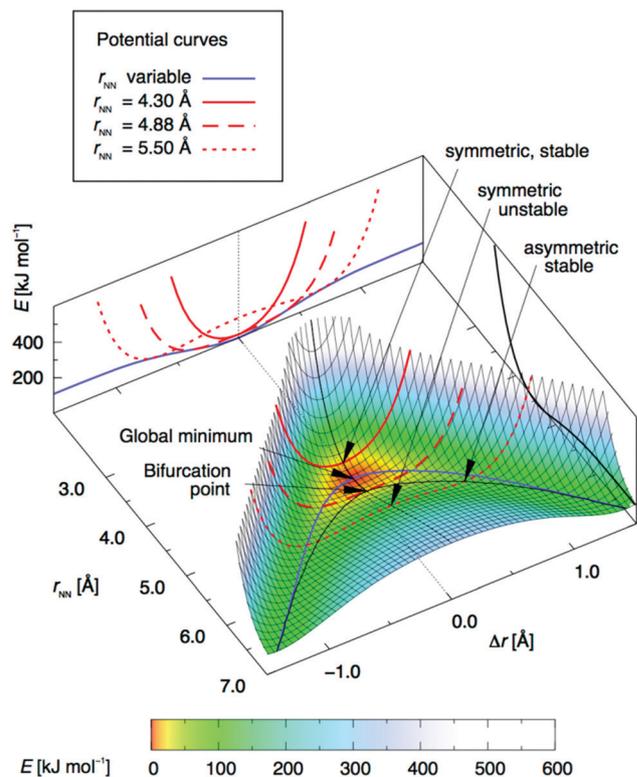


Fig. 12 The variation of electronic energy as a function of the position of the iodine in an $[N \cdots I \cdots N]^+$ halogen bond, upon varying the N–N distance, r_{NN} (Å), and the position of the iodine as described by Δr , the elongation of the iodine from the geometrical midpoint of the nitrogen–nitrogen distance. Thus, at $\Delta r = 0$ the iodine is centered between the nitrogens, whereas at $\Delta r = 0.5$ Å it is 1 Å closer to one of them. The blue line shows the potential curve, corresponding to Fig. 1a, for variable r_{NN} , whilst the red lines show the potential curves for r_{NN} kept fixed at 4.30 Å (solid), 4.88 Å (long-dashed) and 5.50 Å (short-dashed), respectively, shown both in the 3D plot and projected on the rear border plane. The black line on the right border plane shows the dissociation curve of the $[N \cdots I \cdots N]^+$ bond model system. DFT calculations were performed using the M06 exchange and correlation functional and a mixed-level (double zeta/triple-zeta/augmented triple-zeta) basis set. This figure has been adapted from ref. 24 with permission from the American Chemical Society, copyright 2018.²⁵

bonds being remarkably strong (-50 to -82 kJ mol^{-1}) as indicated by their S \cdots I distances being shortened to 66–72% of the sum of the van der Waals radii of the participating atoms. The interacting iodine of these complexes carries 0.323 (ICl) and 0.187 (IBr) charge, which is somewhat lower than the 0.41 charge that was reported for the analogous $[\text{bis}(\text{pyridine})\text{iodine}(\text{I})]^+$ complexes.⁸ Natural bond critical point analyses of these complexes showed $|V_{\text{BCP}}/G_{\text{BCP}}|$ values from 1.31 to 1.60 (here, V is the potential energy density and G the kinetic energy density) for the S \cdots I contacts, indicating them to have a partially covalent character ($|V_{\text{BCP}}/G_{\text{BCP}}|$ ratio typically has values < 1 for non-covalent interactions and > 2 for covalent bonds).

A three-center $[S \cdots I \cdots S]^+$ halogen bond was reported based on the crystallographic and computational investigation of $[\text{bis}(\text{imidazolidinethione})\text{iodine}(\text{I})]^+$ with an R_{XB} ratio of 0.7, $d_{S-I} = 2.6297$ Å, $|V_{\text{BCP}}/G_{\text{BCP}}|$ ratio of 1.58 and interaction energy of -60 kJ mol^{-1} .³⁶ The partial covalent character and



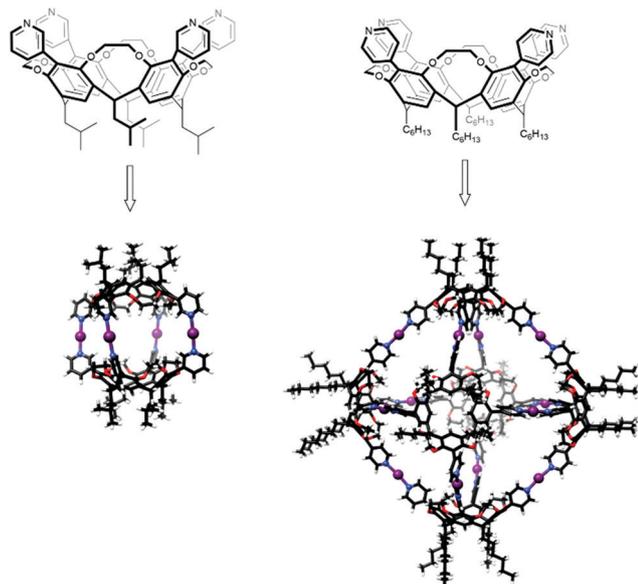


Fig. 16 Three-center halogen-bonded dimeric and hexameric capsules.^{46,48}

capsules were shown to exist as a mixture of at least two different dimeric capsules due to the coordinative nature of Ag^+ , whereas only one symmetric $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded dimeric capsule was observed. The capsules were characterized in solution by DOSY NMR, which confirmed the formation of the larger assemblies. Compared to the free cavitands ($7.5 \times 10^{-10} \text{ m}^2 \text{ s}^{-1}$ and $6.9 \times 10^{-10} \text{ m}^2 \text{ s}^{-1}$) significantly lower diffusion coefficients ($4.8 \times 10^{-10} \text{ m}^2 \text{ s}^{-1}$ and $4.6 \times 10^{-10} \text{ m}^2 \text{ s}^{-1}$) were observed for the complexes. The capsules were also characterized by ESI-MS that further confirmed the formation of the four symmetrical $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen bond stabilized supramolecular capsules.

In the formation of capsular assemblies through three-center $[\text{D}\cdots\text{X}\cdots\text{D}]^+$ halogen bonds, the geometry and the spatial arrangement of the Lewis bases play a critical role. Due to the linearity of the three-center $[\text{D}\cdots\text{X}\cdots\text{D}]^+$ systems, a small structural difference in the Lewis base can enable a construction of assemblies with completely different geometries. Ethylene-bridged tetrakis(pyridyl)cavitands were found to provide a suitable core structure to adjust the position of the electron donor, D, in the way that the construction of capsular assemblies with different geometries and sizes could be achieved.⁴⁶ By changing the position of the pyridyl N-atoms of the ethylene-bridged cavitands from *meta* to *para* (Fig. 16), $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded octahedral hexameric capsules could be constructed. DOSY measurements confirmed the formation of a single larger species with a diffusion coefficient of $2.2 \times 10^{-10} \text{ m}^2 \text{ s}^{-1}$. ESI-MS measurements supported the selective formation of the halogen-bonded capsules with twelve $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ bonds.⁴⁶ The structures of the $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded dimeric and hexameric capsules were further confirmed and studied by a combination of ESI-MS, DT-IMS and theoretical calculations.¹⁷ Rearrangement of the hexameric into pentameric capsules upon changing the solvent from chloroform to

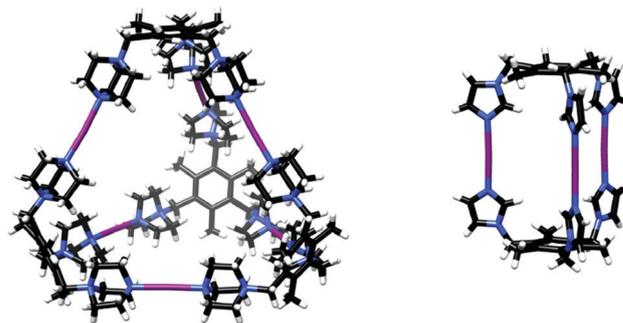


Fig. 17 The solid state structures of M_6L_4 (left) and M_3L_2 (right) $[\text{N}-\text{I}-\text{N}]$ cages.⁴⁹

dichloromethane was observed.¹⁷ This stimuli-responsive rearrangement was rationalized by a combination of micro-solvation, space-filling, and entropic contributions.

Tripodal N-donor ligands were used to form supramolecular M_3L_2 and M_6L_4 $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded cages.⁴⁷ A dimeric M_3L_2 halogen-bonded cage was synthesized from the flexible 1,3,5-tris(imidazole-1-ylmethyl)-2,4,6-trimethylbenzene ligand. Even though this ligand is capable of adapting different geometries due to the flexibility of the imidazole arms, selective formation of the M_3L_2 $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded cage was observed in solution by NMR spectroscopy as well as in the solid state by single crystal X-ray diffraction (Fig. 17).

A tetrameric M_6L_4 cage was achieved using a rigidified cationic tris(1-methyl-1-azonia-4-azabicyclo[2.2.2]octane)-mesitylene ligand. Interestingly, even though the formation of the analogous Ag^+ -coordination cage was not confirmed in either solution or in the gas phase, the subsequent $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded cage was successfully generated from the precursor solution of the Ag^+ -complex. The structure of the tetrameric $[\text{N}\cdots\text{I}\cdots\text{N}]^+$ halogen-bonded cage was confirmed by single crystal X-ray diffraction (Fig. 17).

Halonium-ion-based single and double turn helical complexes were synthesized using oligo-aryl/pyridylene-ethynylene backbones (Fig. 18),⁵⁰ and characterized by X-ray crystallography and solution NMR spectroscopy, corroborated by DFT calculations. The helical structures were stabilized by three-center halogen bonds, and by π - π interactions. These were shown to preserve their helicity in solution by NOESY measurements, as indicated by the proximity of distinct pyridyl groups. The helices exhibited homochiral self-sorting upon crystallization, and were shown to transfer iodonium ions to alkenes,

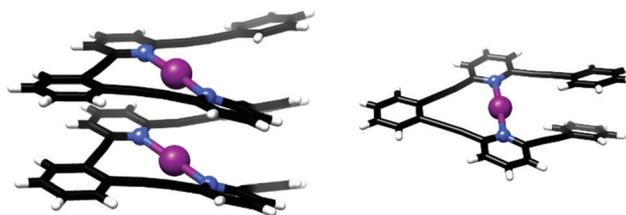


Fig. 18 The solid state structures of the three-center halogen-bonded helices.⁵⁰



demonstrated by iodocyclization of 4-penten-1-ol as a model reaction. As the larger helix has a remarkably short distance between its two iodonium ions (0.1 Å less than the sum of the van der Waals radii), it may provide inspiration for the development of highly compact halonium transfer agents with an unusually high halonium ion density.

Applications in organic synthesis

[Bis(pyridine)iodine(i)]⁺ tetrafluoroborate has been introduced by Barluenga as a mild iodine transfer reagent,⁵¹ and has been widely applied for alkenes' 1,2-iodofunctionalization⁵² and iodocyclizations^{53–55} as well as for alcohol oxidations,⁵⁶ among other applications. Due to the inertness of the tetrafluoroborate counteranion, this reagent allows halofunctionalizations with a variety of nucleophiles (Fig. 19). In the absence of added nucleophiles, 1,2-iodofluorination takes place.⁵⁷ A number of halocyclisation reactions have been reported, in which [bis(pyridine)iodine(i)]⁺ acts as a mild I⁺ transfer agent (Fig. 20). Halogen bond-stabilized halonium transfer reactions have recently been reviewed by Sekar *et al.*⁵⁸

Kinetics studies of halonium transfer from [bis(pyridine)-halogen]⁺ complexes to alkenes⁵⁹ (Fig. 21) suggest that the reactions proceed by reversible dissociation of the three-center complex into free pyridine and an *N*-halopyridinium ion, which is subsequently captured by an alkene, leading to formation of a new halogen-bonded complex, shown as intermediate A in Fig. 11 (where Y is pyridine). The subsequent steps

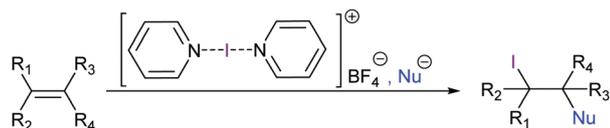


Fig. 19 Barluenga's reagent, [bis(pyridine)iodine(i)] BF₄, is a mild iodine transfer agent that allows the introduction of an iodine and an optional nucleophile into an alkene.

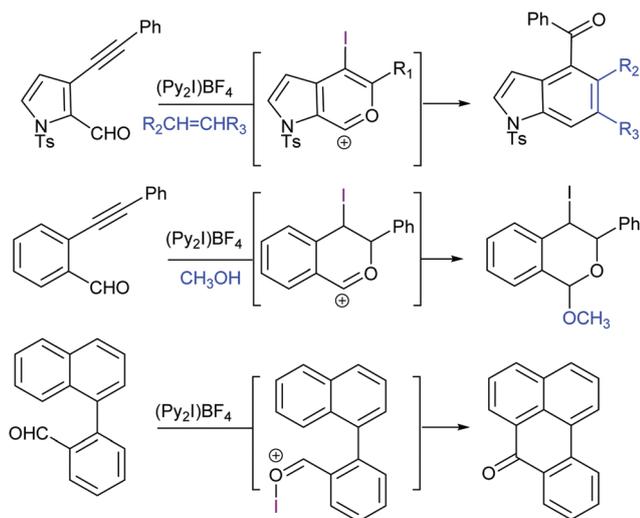


Fig. 20 Halocyclization reactions using Barluenga's reagent are useful in the generation of complex molecules, such as indoles.⁵⁸

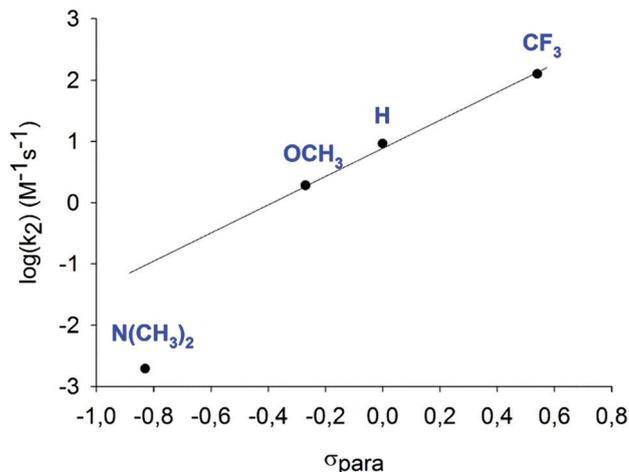


Fig. 21 The second-order rate constants (M⁻¹ s⁻¹) of iodocyclization reactions of [bis(4-*R*-pyridine)iodine(i)]BF₄ (R = H, OMe, NMe₂ or CF₃) reagents with 4-penten-1-ol, normalized to 1 mM, obtained in dry dichloroethane, shown as a function of the Hammett σ_{para} constants for the 4-substituents. A linear correlation is seen for all, but the most electron rich complex suggests that the latter provides I⁺ transfer through a different mechanism.²⁶

follow the general scheme depicted in Fig. 11 to provide various halofunctionalized products. Comparative kinetic investigation of substituted [bis(pyridine)iodine(I)]⁺ complexes revealed that the halogen transfer from electron rich complexes, such as that of 4-dimethylamino-pyridine, is slow and most likely proceeds through a different mechanism.

In addition to the complexes shown in Fig. 6, 2,4,6-collidine has been introduced as a Lewis base to form mild iodine(i) and bromine(i) transfer agents for halocyclisations (Fig. 22).⁶⁰ [Bis(pyridine)iodine(i)]⁺ and [bis(collidine)iodine(i)]⁺ were reported to be superior for some transformations as compared to *N*-iodo-succinimide.⁶⁰ Analogous dioxiodane reagents [R-COO-I-OOC-R]⁺ possessing a three-center halogen bond with oxygen as a halogen bond acceptor, [O-I-O]⁻, have also been reported to be applicable in halocyclization reactions.⁶¹ Furthermore, the

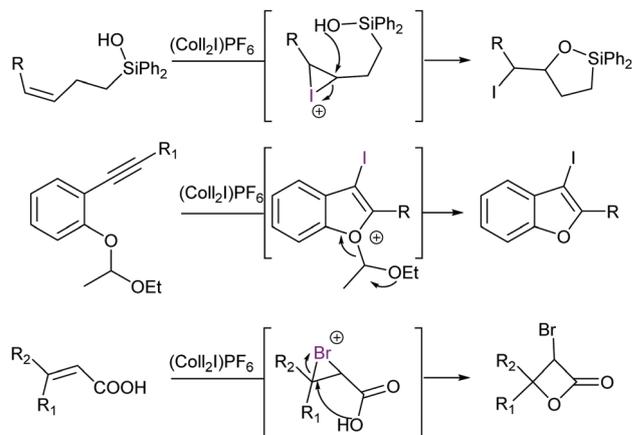


Fig. 22 Examples of halogen(i) transfer reactions of synthetic utility, using [bis(2,4,6-collidine)iodine(i)] PF₆ as a reagent (Coll = Collidine).⁵⁸



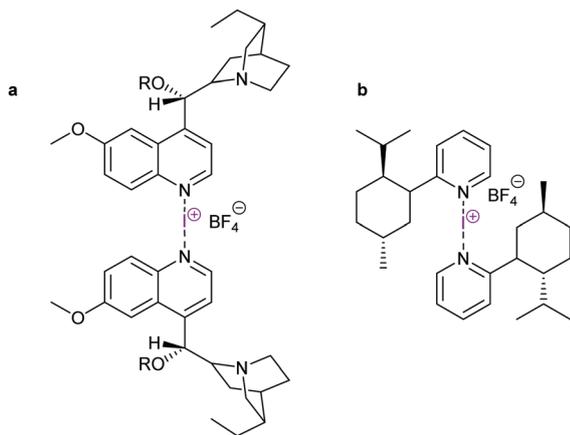


Fig. 23 Chiral [bis(pyridine)iodine(II)] reagents have so far given low enantioselectivities in halocyclizations.^{63,64}



Fig. 24 The proposed mechanism of an enantioselective halocyclisation involving an $[S-X-N]^+$ halonium transfer agent.⁶⁵

$[N-F-N]^+$ -type halogen bond of pyridine was reported as a plausible intermediate in fluorination initiated by Selectfluor.⁶²

Asymmetric halocyclisations using 3-center halogen bond complexes have also been attempted, yet have so far mostly given low enantioselectivities (<15%, Fig. 23),^{63,64} most likely due to the large distance of the reaction centre, that is I^+ (A and B in Fig. 11), from the chiral moieties which in turn also had pronounced flexibility. Using chiral phosphoric acid as an anion phase-transfer catalyst, in combination with an alkyl-DABCO-type $[N-I-N]^+$ or $[N-Br-N]^+$ complex, up to 99% ee has been achieved. Here, however, the chiral phosphate anion was

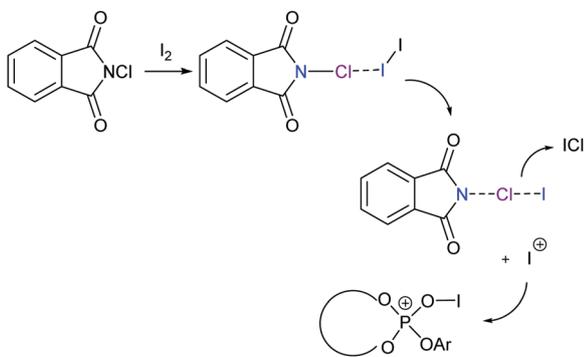


Fig. 25 The proposed mechanism of an enantioselective halocyclisation involving an I^+ generated from *N*-chlorosuccinimide and iodine through formation of a three-center halogen bond.^{67,68}

proposed to be the active catalyst, likely involving halogen bonding with an oxygen donor.⁶⁶ Asymmetric halocyclisation (93% ee) has also been achieved with an $[S-X-N]^+$ -type complex as the active halonium transfer (Fig. 24).⁶⁵

Chiral electrophilic olefin halocyclizations have also been achieved by using *N*-haloamides, for example, in the presence of a chiral ligand in catalytic amounts.^{67,68} In these examples the halonium transfer catalyst was generated upon formation of a three-center halogen bond complex (Fig. 25), with a halonium ion subsequently being transferred to a chiral phosphate catalyst.

Conclusions and future perspectives

Although the fundamentals of the three-center halogen bonding of halonium ions have been described already 70 years ago,⁴³ its experimental investigations have lately intensified and its applications have just begun to take off. The nature of their three-center bonding has over time been interpreted with a variety of theoretical models,²⁰ contributing to our current understanding of chemical bonding. The investigation of the symmetry of halonium ion's halogen bonds, that is the factors determining the position of a halonium ion in between two Lewis bases, has started to shed light on the factors governing halonium transfer processes. Subtle differences between the behavior of I^+ , Br^+ , Cl^+ and F^+ in three-center bonds have been reported, providing useful knowledge about their structure, stability and hence reactivity enabling novel future applications and opening up for studies aiming for a deeper understanding.

Over the past decade the factors possibly influencing the geometry and the stability of this bond have been systematically studied with spectroscopic techniques, providing the ground for future developments. Halonium ions' halogen bonds are remarkably strong and along with their directionality have been demonstrated to provide a robust tool for the design of complex supramolecular structures, such as capsules, cages and helices. Halogen bonds stabilize halonium ions, whose influence can be utilized by rational alteration of the electron density of the three-center system, to thereby modulate halonium ions' reactivity. Synthetic applications have initially been explored by Barluenga and have begun to flourish.⁵⁸ Accordingly, novel synthetic applications utilizing the gained knowledge for the development of selective halonium transfer agents and protocols are to be expected. Establishment of a conceptually new, general strategy for asymmetric halofunctionalization, with a detailed understanding of its mechanism will likely be one of the useful contributions of this research field to a major breakthrough in synthetic organic chemistry. Future applications of halonium ions' halogen bonds in supramolecular systems are expected to provide halonium transfer agents with an unusually high halonium ion loading, and to vastly complex supramolecular architectures with applications in a wide range of scientific fields.

Similar to the recent developments within the wide field of secondary bonding,^{69,70} a quick transfer of knowledge from three-center halogen bonds of halonium ions to related systems



is to be expected. Accordingly, three-center tetrel bonds have recently been discussed.^{44,71} While analogies to “short, strong” hydrogen bonds have been valuable for the discussion of the three-center halogen bonds of halonium ions, the more useful will be understanding their differences, and the differences from other types of hypervalent secondary bonds. This will provide a deeper understanding of the fundamentals of bonding, and allow identification of the most beneficial applications of each as well as utilizing their complementarity.

Overall, applications of the three-center bond of halonium ions' are at a very initial stage. Significant knowledge of their nature and properties that has lately been collected is expected to provide the solid ground for a quickly growing number of well-thought future applications.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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