

The Hydrogen Bond. A Hundred Years and Counting

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Abstract

Since its original inception, a great deal has been learned about the nature, properties, and applications of the H-bond. This review summarizes some of the unexpected paths that inquiry into this phenomenon has taken researchers. The transfer of the bridging proton from one molecule to another can occur not only in the ground electronic state, but in various excited states as well. Study of the latter process has developed insights into the relationships between the nature of the state, the strength of the H-bond, and the height of the transfer barrier. The enormous broadening of the range of atoms that can act as both proton donor and acceptor has led to the concept of the $\text{CH}\cdots\text{O}$ HB, whose properties are of immense importance in biomolecular structure and function. The idea that the central bridging proton can be replaced by any of various electronegative atoms has fostered the rapidly growing exploration of related noncovalent bonds that include halogen, chalcogen, pnictogen, and tetrel bonds.

Keywords: halogen bond; chalcogen bond; pnictogen bond; tetrel bond; excited state proton transfer; $\text{CH}\cdots\text{O}$ H-bond

INTRODUCTION

Over the course of its century of study following its earliest conceptual formulation ¹⁻², the hydrogen bond (HB) has surrendered many of the mysteries of its source of stability and its myriad occurrences. Indeed, one might be hard pressed to think of chemical or biological systems which are completely free of the effects of HBs. Proteins, carbohydrates, and nucleic acids alike owe much of their structure to this phenomenon. The many catalytic functions of enzymes are heavily dependent upon HBs between amino acid residues and substrates. In fact, water would not even exist as a liquid at room and biological temperatures were it not for H-bonding. And of course, one would not be able to understand the many chemical and biological processes that take place in aqueous environment without a thorough treatment of the HBs that occur in each such system.

The HB owes a large segment of its stability to simple Coulombic forces. The normal polarity of a A-H bond, wherein A is an electronegative atom like O or N, places a partial positive charge on the proton. The latter can thus attract the partial negative charge of an approaching nucleophile D. Another important factor resides in the perturbation of the electronic structure of the two species as they approach one another. These alterations are typically referred to as induction, amongst other labels. At least conceptually, the perturbations can be categorized as internal and external. That is, charge shifts within a given molecule can result in polarization energy, while charge transfer refers to any electron density that crosses an imaginary border that separates the two entities. For example, it is common to speak of $n \rightarrow \sigma^*$ transfer by which some density is shifted from the nonbonding, i.e. lone pair, orbital of the nucleophile into the $\sigma^*(A-H)$ antibonding orbital of the proton donor. This accumulation of density in an antibonding orbital has been taken as the source of the weakening of the A-H covalent bond, and the resulting red shift of its stretching frequency, itself a hallmark of H-bonding. The latter density shifts, both internal and external, are sometimes referred to as “covalent” contributions, although this particular sobriquet can be rather vague. To all of the preceding terms, a last attractive contribution arises in connection with London dispersion, a factor which is common to all molecular interactions, HBs being no exception. Of course, if all components of the interaction were attractive, an $AH \cdots B$ HB would collapse into a single molecule. Such a collapse is prevented by Pauli exchange forces, similar in nature to what are colloquially referred to as steric repulsion.

While the earliest concept of a $AH \cdots D$ HB was predicated on only the very electronegative O, N, and F atoms as A or D participants, the list of atoms that can serve in this capacity has greatly expanded over the years. All of the halogens have been shown ³⁻⁷ capable, as have numerous chalcogen ⁸⁻¹⁷ and pnictogen atoms ¹⁸⁻²⁰. The list has been extended to metal atoms as well ²¹⁻²⁸, both

as proton donors and acceptors. In terms of nucleophilic proton acceptors, the original idea of lone pairs has also been extended, now to include π -systems of units such as alkenes or aromatics²⁹⁻³⁵. Even the σ -bonds of molecules such as H_2 can serve³⁶⁻⁴⁰ this function. Yet another new concept is connected⁴¹ with the ability of a through-space α -interaction between two lone pairs on different atoms to strengthen the H-bond with a CH donor. In fact, the definition of the HB has expanded so much over the past few decades that a IUPAC group has established a new set of guidelines⁴² that are quite general.

Since there are already available scores of works on HBs, including a number of extensive monographs⁴³⁻⁵⁰ that concentrate on the central issues of this phenomenon, this review focuses on some of the currently developing frontiers of the concept. As proton transfers within HBs have received a good deal of attention, and much has been written about them, the first topic considered here is the extension of this idea to proton transfers within excited states of HBs. Although of some importance in a number of areas, such as laser development, far less is known about this process in any of its excited states than its ground state analogue. A second frontier discussed here is the ability of the C atom to participate as a HB proton donor. Despite the low electronegativity of C, that generally precludes the normal $A-H^+$ polarization, there is growing evidence of $CH\cdots O$ HBs, and their importance in numerous phenomena. Lastly, discussion turns to a rapidly evolving field of close cousins of HBs, which in an apparent paradox, do not involve a H atom at all.

EXCITED STATE PROTON TRANSFER

Perhaps the first observation of a proton transfer in an excited state dates to 1956 and refers to the intramolecular HB within a methyl salicylate molecule⁵¹. The first documented case of excited state concerted double proton transfer occurred in the 7-azaindole dimer⁵². Dual fluorescence in 3-hydroxyflavone was explained on the basis of this process in 1979⁵³, and this process was placed in the context of a photoinduced proton transfer laser as the process required only 8 ps⁵⁴. In addition to lasers, the excited state proton transfer process has implications for data storage device and optical switching⁵⁵⁻⁵⁷, Raman filters and scintillation counters⁵⁸, triplet quenchers⁵⁹⁻⁶⁰, and polymer photostabilizers⁶¹⁻⁶².

The poster boy for examining intramolecular proton transfer, partly due to its simplicity, is the malonaldehyde molecule which contains an internal $OH\cdots O$ HB within a conjugated ring structure, as pictured in Fig 1. The bridging proton can transfer across to the other O atom which results in a symmetrically equivalent system. The transition state (TS) for this transfer is a symmetric

configuration with the proton equidistant between the two O atoms. An early study of this transfer⁶³ comprised both the ground state S_0 and the first excited $\pi \rightarrow \pi^*$ triplet state, T_1 . The excitation into this state required on the order of 95 kcal/mol, and had several effects. It reduced the acidity of the OH group, as well as the basicity of the other O, and weakened the internal HB. Another result of the excitation is the addition of antibonding character to the C=O bond, which causes it to elongate. The bottom line is a higher pT barrier in T_1 than in S_0 , 13.6 vs 3.6 kcal/mol. This concept was expanded⁶⁴ to include other excited states as well. The $^3\pi\pi^*$, $^3n\pi^*$, $^1\pi\pi^*$, and $^1n\pi^*$ states all displayed a rise in pT barrier vs the ground state, and indeed, a number showed barrierless pT. The height of the barrier was inversely correlated with the strength of the internal HB.

Research has progressed on malonaldehyde and its related systems over the ensuing years and continues unabated⁶⁵⁻⁶⁷. The excited state proton transfer process is apparently important even in terms of the ground state. A very recent examination of the proton transfer⁶⁸ implicated the two lowest excited singlet states in the properties of the ground state transfer via conical intersections. Measurements and later calculations place the height of the ground state proton transfer barrier at 5 kcal/mol, while the transfer in the $^1\pi\pi^*$ state is barrierless, and a high barrier occurs in the $^1n\pi^*$ state⁶⁸⁻⁷⁰, similar to the results obtained years earlier⁶⁴.

A number of modifications to the basic malonaldehyde conjugated ring structure were examined next in the context of glyoxalmonohydrazine. Replacement of the OCCCO ring by OCCNN raised the possibility⁷¹ of different energies for the enol and keto structures, i.e. proton on O or N, as indicated in Fig 2. The ground state pT potential contains two minima, **O** and **N**, with the latter lower by some 9 kcal/mol. The pT from N to O must pass over an energy barrier of only 2 kcal/mol. The situation is generally similar in the $^1\pi\pi^*$ and $^3\pi\pi^*$. The situation is reversed in the singlet and triplet $n\pi^*$ states, where **O** is preferred to **N** by 1 kcal/mol in $^1\pi\pi^*$ and by 14 in the triplet. Unlike the malonaldehyde analogue, most of the excited states of glyoxalmonohydrazine favor a nonplanar geometry. The specific distortion mode is different for each state, as is the force toward nonplanarity. Permitting full distortion has a profound influence upon the energetics of pT, switching the relative stability of **N** and **O** in both excited singlets, as compared to the transfer in the planar case. The situation was modified by replacing the H on the N by a $=CH_2$ group⁷². This perturbation leads to an interesting situation wherein the keto tautomer does not represent a minimum in the ground electronic state, but the reverse is true for either the first excited singlet or triplet, where it is only the keto minimum that is present. This system thus represents a pT that is forced by the excitation.

The entire system was remade⁷³ into a symmetric system, wherein both O atoms of malonaldehyde are replaced by NH groups, as displayed in Fig 3. As in the unsubstituted malonaldehyde, a correlation is noted in that the stronger the HB, the lower the barrier to proton transfer. Nonplanar distortions are different for each excited state, but because the distortions have similar energetic consequences for the equilibrium and transition state structures, the pT barrier of the $^1n\pi^*$ state is little affected by permitting such deformations. As for malonaldehyde molecule, the transfer barriers in either case obey the order $^1\pi\pi^* < S_0 < ^3\pi\pi^* < ^1n\pi^* < ^3n\pi^*$. The barriers are uniformly higher for the internitrogen transfers than for the OH \cdots O interaction in malonaldehyde, which is attributed to the longer HBs. For any given HB, the interoxygen transfer has a slightly higher barrier than does NH - N by 2-3 kcal/mol.

The effects of the 5-membered ring size of malonaldehyde change to both 4 and 6, coupled with addition of a negative charge, while holding the HB atoms to O, was examined⁷⁴ in a series of systems pictured in Fig 4. The pT barriers correlate strongly with various geometric and energetic markers of the strength of the HB. The H-B is weakened by $n\rightarrow\pi^*$ excitation, particularly for the neutral molecule, resulting in a higher barrier. In the case of the two anions, excitation to $^3\pi\pi^*$ strengthens the HB, while the result is more ambiguous for the $^1\pi\pi^*$ state. This trend is reversed in malonaldehyde where the singlet is strengthened by the excitation and the triplet weakened. Some of these patterns were traced directly to the nature of the pertinent orbitals and the density shifts arising from the excitation.

The malonaldehyde theme was added to a phenyl ring in such a way as to perturb the intrinsic symmetry of the pT process, which was augmented by inclusion of a methyl group in o-hydroxyacetophenone⁷⁵, as illustrated in Fig 5. The correlated pT potentials for the ground and first excited singlet states each contain a single minimum, but they differ in placement of the proton. Hence the $S_0\rightarrow S_1$ excitation yields a spontaneous pT, from the hydroxyl O to the carbonyl O. Attachment to a phenyl ring was also examined⁷⁶ in the closely related o-hydroxybenzaldehyde (oHBA), wherein the methyl group of hydroxyacetophenone is removed. In most respects, the addition of the aromatic system exerts little influence upon the properties of malonaldehyde. With the exception of the $^1\pi\pi^*$ state, electronic excitation weakens the HB and simultaneously raises the pT barrier in either system. Unlike the symmetric transfer potential in malonaldehyde, the enol and keto tautomers of oHBA are chemically distinct. $\pi\pi^*$ excitation reverses the preference for the enol tautomer in the ground state. This reversal is connected with the changing degree of aromaticity in the phenyl ring of oHBA. The asymmetric transfer potential in oHBA leads to forward and reverse

barriers of different magnitude. When this factor is accounted for by an averaging procedure, the transfer barriers in *o*HBA are similar to those of the corresponding states of malonaldehyde.

Another variation of the theme shrunk the HB segment down to a 4-membered OCCO ring, which is then attached to a 7-membered hydrocarbon ring as in tropolone, illustrated in Fig 6. The pT process in the S_1 state⁷⁷ is characterized by a low barrier, such that only one doublet of the OH stretching frequency lies below the peak of the pT barrier. A careful analysis of the tunneling splitting revealed that bending vibrations play only a minor role in the pT process so that a 2-dimensional stretching model, involving only $O\cdots O$ and O-H stretches ought to be adequate.

Attachment to a phenyl ring was also considered in the context of salicylaldimine, which contains the symmetric OCCCN ring, and in particular how the pT potential is affected by F substitution⁷⁸ as indicated in Fig 7. Many of the effects are inductive; the electronegative F makes the proximate N or O atom a stronger acid or weaker base and thereby modulates the preferred position of the proton. The agnitude of the perturbation diminishes as the site of substitution is further removed from the HB. This principle also controls the manner in which F affects the geometry and strength of the intramolecular HB in both the enol and keto tautomers. Whereas these notions apply fairly consistently to the ground state and excited $\pi\pi^*$ singlet and triplet, a number of anomalous patterns emerged in the $^1n\pi^*$ state. In general, the effects of fluorosubstitution are smaller in magnitude than the changes that occur in the pT properties as a result of electronic excitation.

While the results described above concerned intramolecular pT through an internal HB, it is also of interest to consider intermolecular processes. An example is the transfer of a proton from phenol to ammonia⁷⁹ which would morph the system from a neutral $\text{PhOH}\cdots\text{NH}_3$ pair to a $\text{PhO}^-\cdots^+\text{H}\text{NH}_3$ ion pair. While such a transfer is highly disfavored in the ground state, the situation changes dramatically in the first excited singlet state where the pT potential develops a second minimum, with the two configurations roughly equal in energy.

More details about these results, and the specific methods applied can be found in the original papers as well as a summarizing Feature Article⁸⁰. Work has certainly not ceased in this field, which continues apace⁸¹⁻⁸³ from both computational and experimental perspectives.

CH \cdots O H-BONDS

Another frontier in the definition and properties of H-bonding lies on the weak end of the spectrum. The C-H group is so pervasive in chemistry and biochemistry, that its ability to participate in a HB is of utmost importance. While a simple alkane does not provide a sufficiently polar CH

group to act in this fashion, it is well documented that a HB is formed if the C changes its hybridization⁸⁴⁻⁸⁵ from sp^3 to sp , as in $HC\equiv CH$ or $N\equiv CH$. Another means to amplify the CH polarity is the placement of electron-withdrawing substituents on the C, as would naturally occur in a protein where each $C^\alpha H$ is flanked by a pair of peptide groups. As work proceeded on $CH\cdots O$ HBs, it was soon apparent that some of them have an unusual quirk. Instead of shifting the A-H stretching frequency to the red as had been taken as a necessary condition of a $AH\cdots D$ HB, a certain subset of $CH\cdots O$ interactions shifted the C-H stretch to the blue⁸⁶⁻⁹². While there were some initial complaints that such a shift in the wrong direction ought to disqualify this interaction as being a true HB, it fulfilled all other typical criteria. This anomaly was soon determined to result from the fact that the direction of shift arises from a delicate balance between forces, some tending to shift to the red, and others to the blue⁹³⁻¹⁰¹. While the former tend to win out in most HBs, the subtle balance simply shifts in the opposite direction for some $CH\cdots O$ HBs¹⁰²⁻¹⁰⁶.

Within the realm of proteins, the most common CH group capable to participate in such a bond is the $C^\alpha H$ group which is surrounded by a pair of electron-withdrawing peptide groups. Calculations have demonstrated the strength of these bonds¹⁰⁷ as just less than that of a standard $NH\cdots O$ interaction. With respect to sidechains containing an aromatic group, e.g. Tyr or Im, the CH of the aromatic ring was also a viable proton donor¹⁰⁸.

It had been part of conventional wisdom that it is the $NH\cdots O$ HBs between strands that hold the β -sheet together. But study of the atomic positions of a β -sheet in Fig 8 shows that CH groups are also in position to donate a proton to the peptide O of the neighboring strand. Quantum calculations¹⁰⁹ showed that these putative interstrand $CH\cdots O$ HBs are competitive in strength with $NH\cdots O$, and serve as an integral component in the stability of the β -sheet, a finding that has since been confirmed by others¹¹⁰⁻¹¹⁵.

Despite a longstanding notion that the strength of a HB between two given groups depends only upon their relative geometry, i.e. HB length and angles, calculations showed this to only be part of the story. Even when a pair of peptide groups is locked into a given configuration¹¹⁶, the interaction energy is highly sensitive to the overall structure of the polypeptide chain on which they reside. In particular, extended conformations of a polypeptide are capable of only weak $NH\cdots O$ HBs, and the interstrand $NH\cdots O$ H-bonds in β -sheets are weaker than those found in other conformations, such as helices, ribbons, and β -bends, even if the specific HB geometries are similar. In a related vein, the $CH\cdots O$ HB is even stronger than $NH\cdots O$ within the context of a simple dipeptide¹⁰⁶ when in a C_5 geometry, a small model somewhat similar to the β -sheet. These trends are not restricted only to *in*

vacuo settings, but retain their integrity within the context of a dielectric continuum model of a protein interior ¹¹⁷.

The importance of the CH \cdots O HB is not limited to structural issues, but also plays a role in various enzymatic mechanisms. These ideas were tested within the context of the serine proteinase family of enzymes ¹¹⁸. Earlier workers had suggested what they called a “ring-flip” hypothesis involving a 180° rotation of a key His residue as a vital step in the catalysis. This mechanism relied on the presence of a CH \cdots O HB in order to stabilize one of the intermediates in the formation of the tetrahedral intermediate. The calculations were generally supportive of this idea but raised some important discrepancies that required resolution before its acceptance. This sort of HB has implications in other enzymatic mechanisms as well ¹¹⁹. There are also contributions of this weak HB as a determining factor ¹²⁰⁻¹²² in the conformation of certain organic systems. Needless to say, even normally weak HBs can be strengthened by the acquisition of charge on either the proton donor or acceptor group ¹²³⁻¹²⁵.

As experimentalists continue to examine systems for the presence of CH \cdots O HBs, they require certain trademark or fingerprint characteristics for which to search. In addition to geometric aspects which are already fairly well understood, it is common to apply spectroscopic methods to these biological systems. Quantum calculations have provided some such characteristics for which to search ¹²⁶⁻¹²⁸. It was noted earlier that CH stretching frequencies can shift in either direction; nonetheless a blue shift would be a valuable indicator as it would not occur in the absence of such a bond. A downfield shift of the bridging proton’s NMR signal would reinforce this supposition. With respect to the proton acceptor, a large upfield shift of the O chemical shift, by as much as 16 ppm, can serve as another indicator.

COUSINS OF THE H-BOND

Another HB frontier that has been approached is its definition as an interaction involving a bridging proton. Suppose this H were to be replaced by a different atom, but the various properties were to remain largely intact. Early work had suggested such a phenomenon, wherein H could be replaced by any of several halogen (X) atoms ¹²⁹⁻¹³¹. The ability of the X to serve in this capacity rests on the highly anisotropic charge distribution which surrounds it. While its electronegativity imparts to it an overall negative partial charge, there is a reduced density along the extension of the R-X bond, which has been termed a “polar flattening”, which in turn causes a region of positive

electrostatic potential in this region, commonly referred to as a σ -hole¹³²⁻¹³⁶. It is this localized positive region which can attract a nucleophile, in much the same way as does the H in a $AH\cdots D$ HB.

This idea of a halogen bond (XB) is not restricted only to halogen atoms, but is common also to chalcogen, pnictogen, and even tetrel atoms, in their eponymously named bonds. There are certain fine point differences amongst these bonds. For example, while a univalent X atom displays a single σ -hole lying along the extension of the R-X bond, a divalent Y chalcogen atom will typically contain two such σ -holes, each lying along an extension of the two R-Y bonds. These holes will not be able to lie directly opposite the bond, since the high electron density of the two Y lone pairs will tend to push the holes away from them. This distinction is illustrated in the comparison of FCl with HFS in **Figs 9a and 9b**, respectively. In each case, the blue oval represents the σ^* antibonding orbital, FCl in a and FS in b, along which the reduced electron density would tend toward a σ -hole. The Cl atom of FCl contains three lone pairs, represented by red ovals, whose density would push the positive potential, i.e. the σ -hole, away from themselves. Due to their symmetric disposition, the blue region of the potential lies directly along the F-Cl axis and the σ^* orbital direction. On the other hand, there are only two lone pairs on the S atom in **Fig 9b**, which together push the σ -hole down away from them. A nucleophile would thus tend toward this blue region σ -hole, and a nonlinear $FS\cdots Nuc$ alignment. A similar nonlinear arrangement between the F-P bond extension and the σ -hole would be expected for a pnictogen bond (ZB), in **Fig 9c**, due to a single lone pair. The absence of any lone pairs on the Si atom in **Fig 9d** would allow the blue positive potential σ -hole to align perfectly with the $\sigma^*(F-Si)$ antibonding orbital, resulting in a linear tetrel bond (TB). It must be noted that since the high electron density of lone pairs mitigate the positively charged σ -hole, the progressive decrease of lone pair number in the sequence halogen > chalcogen > pnictogen > tetrel would tend to enhance the σ -hole intensity in the same order.

After an initial study that demonstrated that a $P\cdots N$ interaction is energetically preferred to a $PH\cdots N$ HB¹³⁷, more detailed study¹³⁸ showed this to be a characteristic of pnictogen bonds in general. Part of the interaction arises from the donation of charge from the N lone pair into the $\sigma^*(PH)$ antibonding orbital. This transfer is identical to that in a $PH\cdots B$ HB, except that it is the P-end of this orbital which points toward the N, rather than the H-end. The strength of such a ZB is heightened when the H is replaced by an electron-withdrawing agent such as F¹³⁹. **Fig 10** shows how the interaction energy ΔE rises as the substituent's electron-withdrawing power increases from CH_3 and H up to F and NO_2 for the $XH_2P\cdots NH_3$ series¹⁴⁰. Along with this rise in binding is a

concomitant amount of charge transferred from the base to the Lewis acid, as measured either by NBO values of $E(2)$ or the total charge on the entire subunit Δq .

In fact, this substitution is even capable of making first-row N capable of accepting charge in a $N\cdots N$ pnictogen bond¹⁴¹ despite the reluctance of first-row atoms to engage in such bonds. With respect to the electronegativity and polarizability of the pnictogen atom, larger atoms yield stronger ZBs^{140, 142-143 144} in the order $P < As < Sb$. This trend has no parallel to HBs as it is always the proton that acts as bridge.

As one might anticipate, since halogen and pnictogen atoms can replace the proton in HBs, the same idea can be extended to chalcogen (S, Se, etc) atoms as well. Work by our group¹⁴⁵⁻¹⁴⁹ as well as numerous others¹⁵⁰⁻¹⁵⁶ elaborated on these ideas. **Indeed, there is currently a IUPAC group tasked**

with adopting a working definition of a chalcogen bond, with others to follow later for the other sorts.

The extension to tetrel atoms (the Si family) occurred soon thereafter, showing many of the same controlling factors that are present for X, Y, and Z atoms¹⁵⁷⁻¹⁶⁰. The normally tetravalent tetrel atoms introduced a new factor which had been less prominent in the other sorts of bonds. In order for a base to approach the central tetrel atom along a face of the tetrahedron, the three proximate substituents must “peel back” away from this base, changing the originally tetrahedral structure into something akin to a trigonal bipyramid. There is thus a good deal of deformation energy that must be surmounted¹⁶¹⁻¹⁶³ if this tetrel bond is to form. This deformation energy makes the tetrel bond formation less exothermic than it would otherwise be, and can even control the particular site at which the base can attack.

The idea of tetrel bonds brought up an interesting issue. It had typically been considered that a nucleophile lying along the R-C extension of a $R-CH_3$ group constituted a trifurcated HB, i.e. interaction with three H atoms. And there are certainly many such geometrical dispositions of this sort, in both chemical and biological systems¹⁶⁴. But how can one distinguish this idea of a trifurcated HB from the newer concept of a $R-C\cdots D$ tetrel bond? Indeed there are spectroscopic markers that are different for the two sorts of interactions¹⁶⁵⁻¹⁶⁶, and it is hoped that the future will witness attempts to distinguish these two types of interactions.

As work has progressed in this area, it has become recognized that the positive regions are not limited only to σ -holes lying along the extension of a particular covalent bond. There are π -holes as well, wherein the positive potential sits above the plane of a molecule, as in H_2SiO for example, in the vicinity of the electronic π -cloud. This broadening of the idea has been probed extensively and shown that while the π -hole interactions are usually weaker than their σ parallels, this trend is

sometimes reversed^{158, 167-169}, especially when the π -hole lies above a triel atom such as B or Al¹⁷⁰⁻¹⁷³. Of course, such π -hole interactions do not have a H-bonding parallel.

These relatives of the HB are hardly exotic academic novelties, but have a wide range of applications, such as serving as synthons in self-assembling networks¹⁷⁴, biological catalysis¹⁷⁵, oxidative addition¹⁷⁶, self-assembled monolayers¹⁷⁷, S_N2 reaction catalysis¹⁷⁸, design of functional mesomorphic materials¹⁷⁹, and even directed construction of supramolecular quadruple and double helices¹⁸⁰. One of the more interesting uses concerns selective binding of anions¹⁸¹⁻¹⁸⁸. It was realized that the replacement of the H atom of certain multidentate anion receptors with a halogen atom allowed them to engage in halogen bonds with an anion, which in turn strengthened the interaction, and enhanced the selectivity for certain anions over others.

Calculations were applied to this idea, and were able to suggest certain options that ought to enhance these abilities. Optimal choices of particular halogen atoms were proposed, along with identification of chemical groups to which they ought to be bonded, spacer groups between the halogen bonding groups, and overall charge¹⁸⁹⁻¹⁹¹. Subsequent work broadened this idea beyond simply halogen bonds, but considered their chalcogen, pnictogen, and tetrel counterparts¹⁹²⁻¹⁹⁵. It was concluded that tetrel bonds offered a particularly tempting choice for their interactions with a halide, furnishing both very strong interactions, and a marked preference for F⁻ over other halides.

PERSPECTIVE

It would seem then that even after a full century of study, which has provided a wealth of information and insights about the H-bond, we are nowhere near the end of learning its secrets. Its fundamental nature is a template for a much broader set of interactions. Far from the initial thoughts that the A and B atoms in the AH \cdots B interaction are limited to O, N, and F, the set of participating atoms has broadened so widely over the years, such that an atom that *cannot* serve in this capacity would be the exception, not the rule. In particular, the entry of the CH group into the club of proton-donating members, has opened wide new vistas concerning the structure and function of large molecules including proteins and nucleic acids, vistas that are only recently beginning to be explored. The possibility of proton transfer within a given HB has enormous implications as well, not only in the ground electronic state but also in various excited states which open new sets of practical applications. And the further broadening of the original concept of a proton-bridging HB to systems where this central proton is replaced by any of a large number of electronegative atoms has

introduced an entire new area that encompasses halogen, chalcogen, pnictogen, and tetrel bonds, again an area whose impact is only beginning to emerge.

Given all that has transpired in the last century, it would be foolish to presume that we have reached the final border of what the H-bond has to teach us. One can only hope that the next century of inquiry will be as fruitful as the first.

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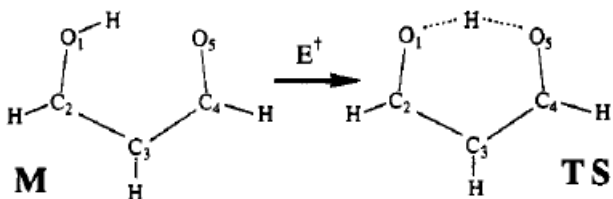


Fig 1. Structure of malonaldehyde (M) and the transition state (TS) for the transfer of the proton between O atoms.

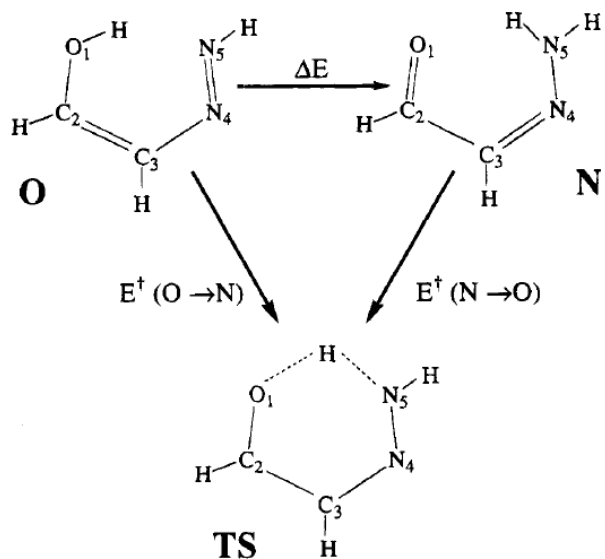


Fig 2. **O** and **N** tautomers related to asymmetric pT in glyoxalmonohydrazine

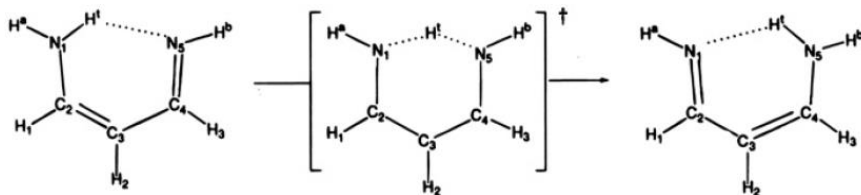


Fig 3. Symmetric proton transfer in 1,5-Diaza-1,3-pentadiene.

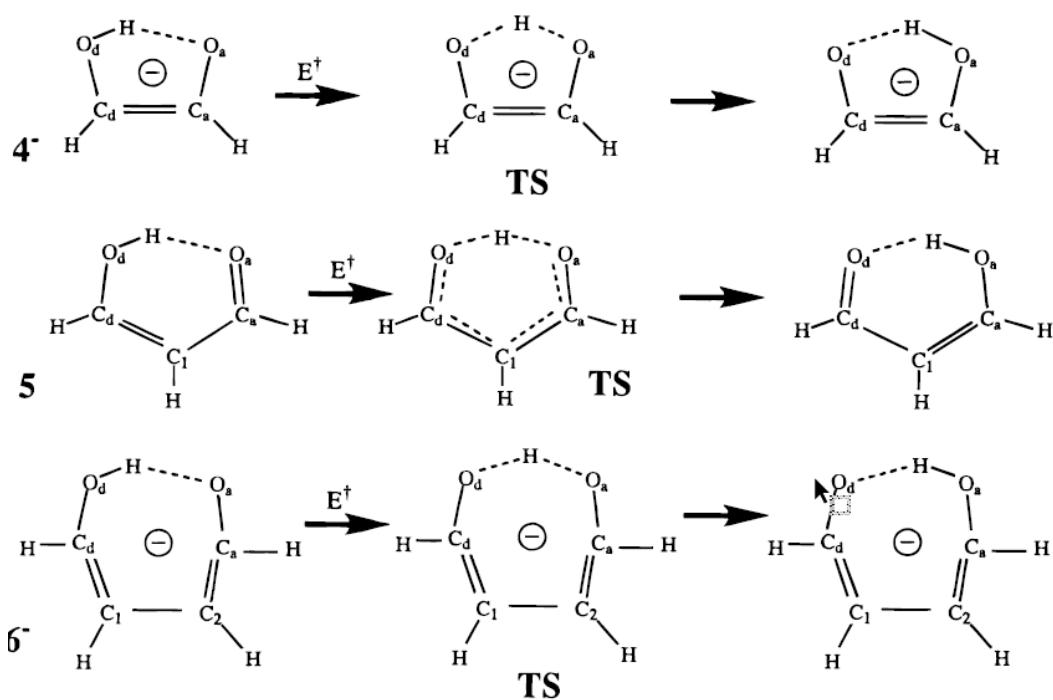


Fig 4. Four (4') and six (6') membered anionic ring analogues of malonaldehyde (5).

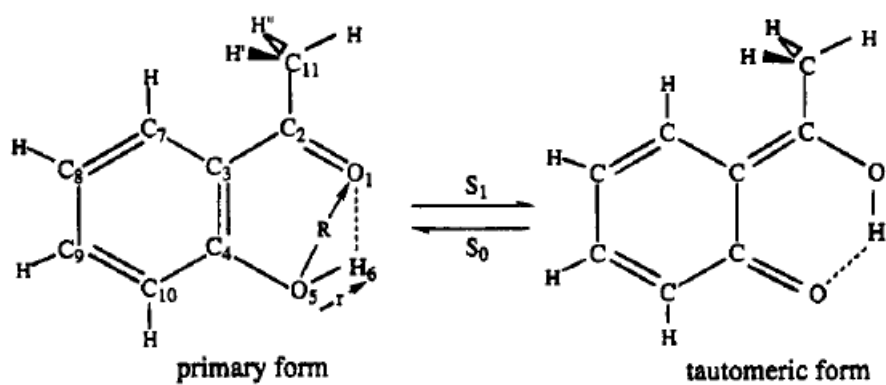


Fig 5. Primary and tautomeric forms of o-hydroxyacetophenone.

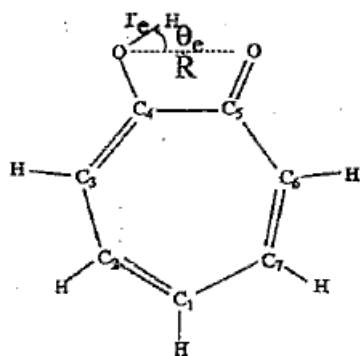


Fig 6. Four-membered OCCO ring attached to 7-membered ring in tropolone system.

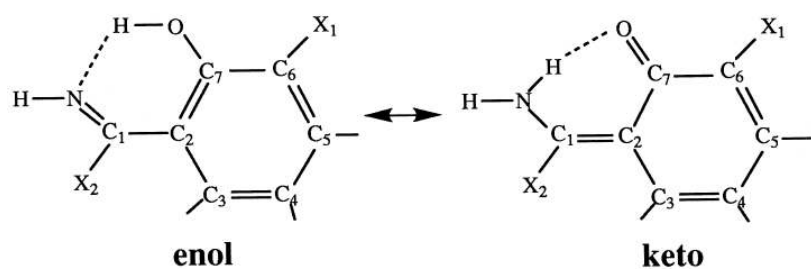


Fig 7. Placement of a F-substituent into salicylaldehyde in either X₁ or X₂ positions.

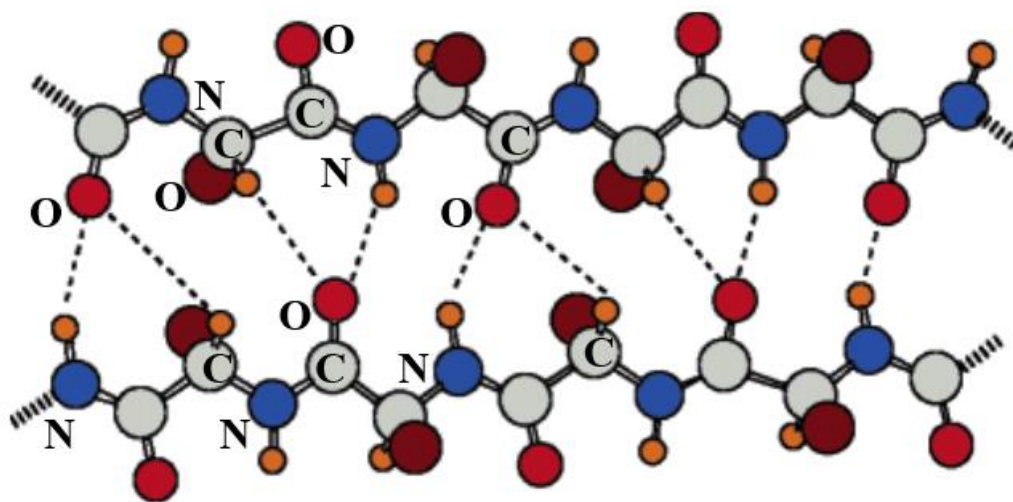


Fig 8. Pair of polypeptide strands in the antiparallel β -sheet arrangement, indicating putative HBs by broken lines. Brown atoms represent generic R groups of amino acids.

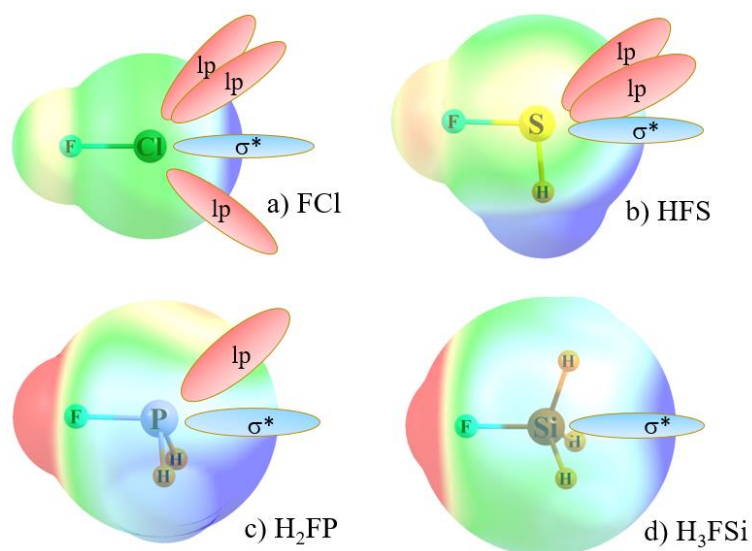


Fig 9. Molecular electrostatic potential surrounding indicated molecules, with blue indicating positive and negative represented by red regions. Light blue oval designates the $\sigma^*(\text{F-A})$ antibonding orbital that lies directly opposite the F-A covalent bond. Lone pairs on each central A atom are indicated by red ovals.

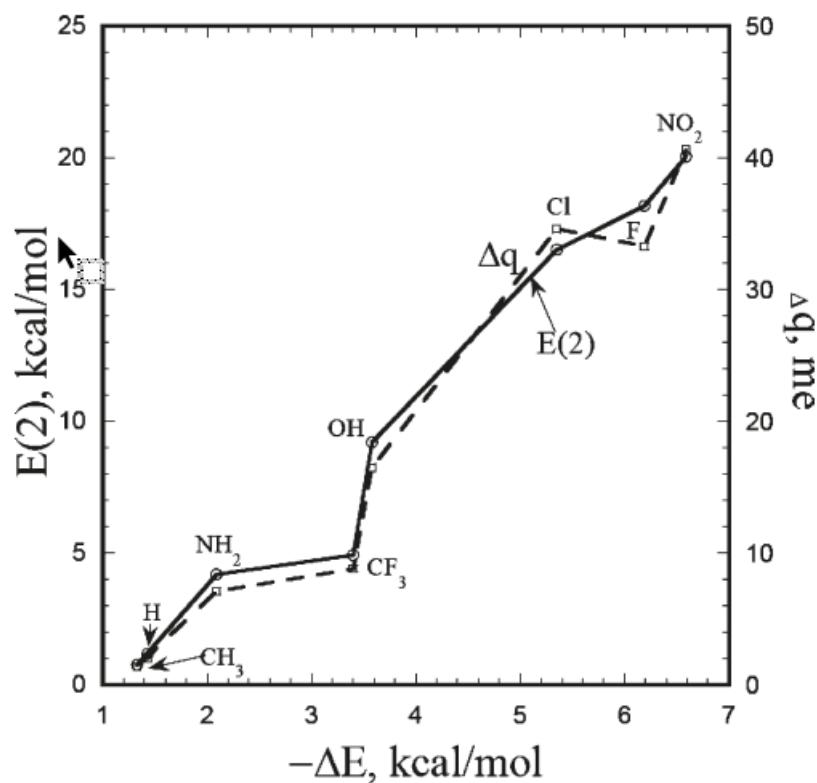


Fig 10. Variation of second-order energy $E(2)$ and amount of charge transferred from N lone pair of NH_3 to $\sigma^*(\text{XP})$ antibonding orbital of H_2PX as a function of interaction energy ΔE

Steve Scheiner is a professor of computational chemistry at Utah State University. After receiving his PhD from Harvard University in 1976, and proceeding to a Weizmann Postdoctoral Fellowship at Ohio State University, he started his independent academic career at Southern Illinois University, Carbondale. He moved to Utah State University in 2000. His research has centered around various aspects of H-bonds, from their fundamental underpinning to their role in enzymatic activity, from the proton transfers that take place within to their cousins, known as halogen, chalcogen, pnictogen, and tetrel bonds.

