Tunable chiral triazole-based halogen bond donors: assessment of donor strength in solution with nitrogen-containing acceptors†

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Strong halogen bond (XB) donors are needed for the activation of neutral substrates. We demonstrate that XB donor properties of iodo-triazoles can be significantly enhanced by quaternization in combination with varying the counterion and aromatic substituent, exemplified by association constants with quinuclidine as high as $1.1 \times 10^4$ M$^{-1}$.

Halogen bond (XB) based applications utilize the attractive interaction between a Lewis acidic halogen atom and a Lewis base. From the turn of the century numerous publications have focused on the use of XB in the solid state and more recently, in solution as well. Among these applications the potential of XBs in catalysis and anion recognition should be highlighted. As shown by Huber et al. these fields can be closely associated, exemplified by halide abstraction reactions.

Compared to anion recognition, the recognition of neutral species in solution has received less attention. Studies with neutral acceptors have a primary focus on the fundamental nature of halogen bonding, such as the influence of the solvent and the structure of the XB donor/acceptor on the strength of XB. Amines have usually been used as neutral acceptors in these studies for their high affinity towards XB donors. This property can potentially be utilized in the detection of biologically relevant amines by XBs. From the synthetic point of view, the activation of neutral species through XBs is also a topic of high interest. In general, compared to anions, neutral acceptors form weaker complexes with organic XB donors. Therefore, XB donors with stronger halogen bonding ability should be used for the activation of neutral compounds. From a catalyst design perspective, information on the extent different structural fragments affect XB donor ability is of great value.

Recently, we became interested in applying XBs in asymmetric catalysis. Chiral 5-halo-1,2,3-triazoles are among the best candidates of catalysts to achieve this goal. The triazoles are readily available through a Cu-catalysed click reaction and access to a broad range of alkynes with the possibility to quaternize the triazole core makes it feasible to enhance the donor ability of the triazole. In addition, the availability of many chiral azides offers wide opportunities for the design of new chiral XB donor systems. We have shown the potential of these compounds to interact with various possible substrates and in enantiodiscrimination. Due to relatively low affinity constants with thio urea acceptors, it was difficult to fully assess how structural modifications affect XB donors’ binding ability. Therefore, a stronger XB acceptor has to be selected for this kind of analysis. Anionic species are known to give large affinity constants with halo-triazolium salts, however, in these complexes charge attraction plays a key role in XB formation. We therefore chose quinuclidine, a neutral monodentate XB acceptor with a readily accessible lone pair, for screening of the effect of XB donor analogues (Fig. 1) on XB strength. Herein we describe the formation of complexes between triazole-based XB donors and quinuclidine in solution with emphasis on the influence of aromatic substituents and counterions on XB donor strength and investigate the XB donors’ ability to discriminate between enantiomers of chiral imines and amines.

A collection of monodentate XB donors shown in Fig. 1 were synthesized (see ESI† for details). To determine the effect of structural changes, the triazolium salts were modified by introducing a perfluorophenyl or a p-nitrophenyl substituent instead of a phenyl substituent, changing the counterions and varying the halogen atoms. The XB donor ability of the synthesized compounds was determined through their respective association constants with quinuclidine in CDCl$_3$, based on $^1$H NMR titration experiments. To evaluate the XB strength more accurately, the titration experiments were carried out in duplicate. The results are summarized in Table 1.
The given affinity constant between quinuclidine and 3-BARF is among the highest affinities reported so far for a neutral acceptor.\(^6\)

Usually XBs are stronger in apolar solvents than in more polar solvents\(^6\) and as a comparison, the association constant is only a magnitude smaller than that for the complex between quinuclidine and I\(_2\) measured in heptane.\(^7\)

The strength of the XB is known to decrease based on the polarization of the halogen atom and the increase of electronegativity in the order of I > Br > Cl > F.\(^3\) In our \(^1\)H NMR titration study, the iodo-triazolium analogue (1-OTf) displayed moderate affinity towards quinuclidine whereas the corresponding bromine analogue (4-OTf) did not show any affinity towards quinuclidine altogether (Table 1, entry 7, also see ESI† for details). The absence of complex formation with the bromine derivative agrees with a similar outcome in our previous investigation.\(^13\) In an attempt to obtain a complex containing a bromine atom as the donor, a bromo-triazolium salt 5-BARF with the strongly electronegative pentafluorophenyl substituent was synthesized. Nevertheless, the changes undertaken made the donor too labile and upon the titration experiment dehalogenation prevented the determination of the affinity constant (Table 1, entry 8, also see ESI† for details).

Quaterization of the triazole core has been critical to obtain compounds with sufficient XB donor ability.\(^13\) To ascertain the impact of charge in the triazole core, neutral perfluorinated triazole 7 was also titrated with quinuclidine. The obtained affinity constant is indeed very low. However, this result is of importance in its own right as there are only a few examples

![Fig. 1](image1)

**Table 1** Association constant \(K_a\) values\(^6\) of the XB donor–quinuclidine pairs

<table>
<thead>
<tr>
<th>Entry</th>
<th>XB donor</th>
<th>(K_a) M(^{-1})</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1-OTf</td>
<td>57 ± 5</td>
</tr>
<tr>
<td>2</td>
<td>1-BARF</td>
<td>1.23 ± 0.01 \times 10^3</td>
</tr>
<tr>
<td>3</td>
<td>2-OTf</td>
<td>257 ± 12</td>
</tr>
<tr>
<td>4</td>
<td>2-BF(_4)</td>
<td>284 ± 12</td>
</tr>
<tr>
<td>5</td>
<td>3-OTf</td>
<td>703 ± 6</td>
</tr>
<tr>
<td>6</td>
<td>3-BARF</td>
<td>(1.1 ± 0.3) \times 10^4</td>
</tr>
<tr>
<td>7</td>
<td>4-OTf</td>
<td>&lt;1</td>
</tr>
<tr>
<td>8</td>
<td>5-BARF</td>
<td>n.d(^b)</td>
</tr>
<tr>
<td>9</td>
<td>6-OTf</td>
<td>&lt;1</td>
</tr>
<tr>
<td>10</td>
<td>7</td>
<td>2.0 ± 0.3</td>
</tr>
</tbody>
</table>

\(^a\) Association constant \(K_a\) measured in CDCl\(_3\) at 298 K and determined by fitting the \(^1\)H NMR titration data to 1 : 1 binding isotherm of BindFit.\(^13\) The given \(K_a\) and standard error are the calculated mean values of two parallel experiments. Full details given in the ESI.\(^b\) \(K_a\) could not be determined due to the instability of XB donor during the experiment.

To evaluate the influence of substituents of the aromatic ring that connects directly to the triazolium core on XB formation ability of the triazolium salts, a perfluorinated and a nitro-substituted derivative (3-OTf and 2-OTf, respectively) were compared with the unsubstituted phenyl derivative 1-OTf (Table 1, entries 1, 3 and 5). The affinity towards quinuclidine decreases in the order of 3-OTf > 2-OTf > 1-OTf which corresponds to the decrease of the size of the σ-hole on the iodine atom.\(^13\) The perfluorinated XB donor had more than twice as high affinity towards quinuclidine as the NO\(_2\)-containing XB donor and over an order of magnitude higher affinity when compared to 1-OTf. The strong XB donating ability of perfluorinated XB donors is explained by its highly electronegative fluorine substituents that significantly increase the polarization of the C–X bond, therefore increasing the σ-hole.\(^7,14\) The electron-withdrawing nitro group in compound 2-OTf is similarly essential to enhance its XB donor ability, albeit less strongly compared to the more electron deficient perfluorophenyl group in 3-OTf. To determine that the changes in chemical shifts were indeed induced by halogen bonding, the nonhalogenated analogue 6-OTf was synthesized which expectedly did not interact favourably with quinuclidine (Table 1, entry 9).

The effects of anionic counterions were characterised based on triflate (1-OTf, 2-OTf, 3-OTf), tetrafluoroborate (2-BF\(_4\)) and tetrakis[3,5-bis(trifluoromethyl)phenyl]borate (1-BARF, 3-BARF) containing triazolium salts. Due to poor solubility, the comparison to 1-BF\(_4\) could not be made. In general, the change of the counterion affected XB strength substantially in accordance with their coordination ability.\(^6\) The less coordinating tetrafluoroborate containing triazolium salt 2-BF\(_4\) showed higher affinity towards quinuclidine compared to the triflate containing salt 2-OTf (Table 1, entries 3 and 4). The introduction of the BARF counterion increased XB strength by more than one order of magnitude (Table 1, comparing entries 1 and 5 to entries 2 and 6). To the best of our knowledge, the obtained affinity constant between quinuclidine and 3-BARF is among the highest affinities reported so far for a neutral acceptor.\(^6\)

Quaternization of the triazole core has been critical to obtain compounds with sufficient XB donor ability.\(^13\) To ascertain the impact of charge in the triazole core, neutral perfluorinated triazole 7 was also titrated with quinuclidine. The obtained affinity constant is indeed very low. However, this result is of importance in its own right as there are only a few examples.
describing complex formation in solution between a neutral
triazole and a neutral acceptor. The difference between
the neutral XB donor (7) and its charged derivative (3-OTf)
is more than two orders of magnitude (Table 1, entries 5 and 10).
However, if the counterion acts as an acceptor and competes
with quinuclidine for XB formation, triazole 7 should be
compared to 3-BARF, which has the less coordinating BARF
counterion and therefore provides a better representation for
a “naked” cationic backbone. In this case, the difference in
binding ability of four orders of magnitude was observed (Table
1, entries 6 and 10).

Furthermore, 1H NMR titration measurements were
performed using both enantiomers of chiral imine 8 and amine 9
(Fig. 2) to determine whether the XB donors are able to selectively
interact with chiral substances. For these experiments, 3-
BARF was chosen as the donor due to the highest binding
affinity towards quinuclidine. The XB donor showed no preference
towards either enantiomer of the selected acceptors
since no differences between the two enantiomers Ks values
were observed in either case (Table 2, entries 1 and 2; entries 3
and 4). Nevertheless, the affinity constant between amine 9 and
3-BARF is considerably higher compared to the only reported
example, where the XB strength between an organic XB donor
and secondary amine was measured. The affinity constant
between perfluorohexyl iodide and piperidine was <1 in all three
solvents used in that study. The difference compared to the
binding strength of quinuclidine can partly be explained by the
fact that cyclic amines are better acceptors than acyclic amines.

Calculations were performed on the CAM/B3LYP theoretical
level of theory using DEF2TZVP basis set to model the interaction
between both enantiomers of amine 9 and 3-OTf. The calculated
complexes in the vacuum and in CHCl3 had similar energy
values (see ESI for details). The substituents on the triazole
core are most likely not sufficiently bulky to differentiate
between the two enantiomers through steric repulsion or by
other noncovalent interactions (Fig. 3). This could also explain
our previously obtained results of enantiodiscrimination experiments,
where Takemoto’s catalyst was used as an acceptor and that suggest that both hydrogen and halogen
bonding interactions influenced the binding of enantiomers. Therefore, a more beneficial approach would be to use mul-
dentate or bifunctional XB donors that form more rigid
complexes. For example, Beer et al. has shown that chiral
didentate XB donors that contain at least two halo-triazole cores
are suitable for differentiating between enantiomers.

In conclusion, we have once again shown the pivotal role of
charge on XB donor strength. In addition, by changing the
aromatic substituent and the counterion, the XB donor prop-
erties of triazole-based donors can be enhanced even further.
This is exemplified by the fact that the donors form complexes
with quinuclidine with association constants covering almost
four orders of magnitude. To the best of our knowledge, the
reported association constants are comparable to the largest
described between an amine and an organic XB donor in
solution. Enantiodiscrimination of acceptors 8 and 9 by the
most powerful donor 3-BARF was not observed. However,
information obtained during this study can aid to move towards
more selective donors.

**Conflicts of interest**

There are no conflicts to declare.

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**Notes and references**

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