

## Can non-polar hydrogen atoms accept hydrogen bonds?†‡

Lixu Yang, Thomas A. Hubbard and Scott L. Cockroft\*

Cite this: *Chem. Commun.*, 2014, 50, 5212Received 7th August 2013,  
Accepted 20th September 2013

DOI: 10.1039/c3cc46048g

www.rsc.org/chemcomm

We examine an unusual case where a neutral hydrogen atom acts as a hydrogen-bond acceptor. The association constant between trihexylsilane and perfluoro-*tert*-butanol was measured as  $\sim 0.8 \text{ M}^{-1}$  in cyclohexane. Computations and experimental NMR data are consistent with a weak, but favourable Si–H $\cdots$ HO interaction.

The ubiquitous hydrogen bond continues to intrigue chemists whilst its definition continues to evolve.<sup>1</sup> The classic H-bond acceptors nitrogen and oxygen (Fig. 1a) are now considered alongside less conventional acceptors<sup>2</sup> such as arenes,<sup>3</sup> halogens,<sup>4</sup> and even the noble gases.<sup>5</sup> While C–H $\cdots$ H–C contacts are not classified as H-bonds<sup>1</sup> and can be attributed as arising through solvophobic<sup>6,7</sup> and van der Waals dispersion forces,<sup>8,9</sup> examples of favourable MH $\cdots$ H contacts (where M is an element that is less electronegative than H) have been identified that have since become known as ‘dihydrogen bonds’ (Fig. 1b).<sup>10–12</sup> Most dihydrogen bonds identified to date involve metals or Lewis adducts that convey strong hydridic character to adjacent hydrogen atoms (Fig. 2a–c).<sup>10–13</sup> Thus, the question remains whether non-polar, neutral hydrogen atoms are able to act as H-bond acceptors (Fig. 1c).

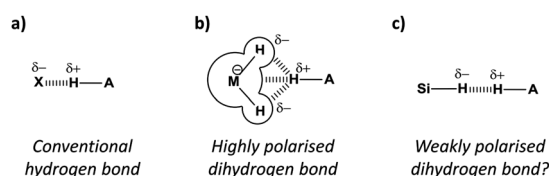


Fig. 1 (a) Conventional hydrogen bond compared with (b) a dihydrogen bond involving hydridic hydrogen atoms bonded to a strongly polarising element M. (c) A proposed dihydrogen bond where a weakly polarised hydrogen atom acts as a hydrogen-bond acceptor.

EaStCHEM School of Chemistry, University of Edinburgh, King's Buildings, West Mains Rd, Edinburgh, EH9 3JJ, UK. E-mail: scott.cockroft@ed.ac.uk; Tel: +44(0)131 650 4758

† Celebrating 300 years of Chemistry at Edinburgh.

‡ Electronic supplementary information (ESI) available: Experimental and computational methods and tabulated data. See DOI: 10.1039/c3cc46048g

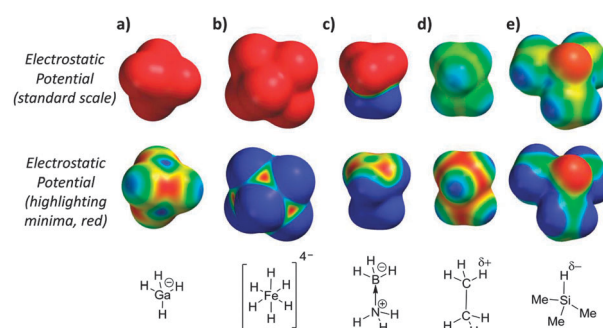


Fig. 2 Electrostatic surface potentials (ESPs) of representative dihydrogen bond acceptors. Electrostatic potentials on the top row are scaled using a standard scale from  $-50 \text{ kJ mol}^{-1}$  (red) to  $+50 \text{ kJ mol}^{-1}$  (blue), while those on the bottom row are scaled to highlight the ESP minimum (ESP<sub>min</sub>, red) to ESP<sub>min</sub> +  $50 \text{ kJ mol}^{-1}$  (blue). Minimised geometries and ESPs were calculated using B3LYP/LACVP.

Here we present a combined solution-phase and theoretical investigation of Si–H as a fully neutral H-bond acceptor (Fig. 1c, 2e, 3–5), whilst examining the utility of electrostatic potentials for rationalising the physicochemical origins and geometric preferences of dihydrogen bonds (Fig. 1 and 2).

Where conventional H-bonds are mostly linear (Fig. 1a), most dihydrogen bonds possess bent or bifurcated geometries (Fig. 1b).<sup>12</sup> As a result, some authors have ascribed M–H  $\sigma$ -bonds as being dihydrogen bond acceptors rather than the hydridic H atoms themselves,<sup>10</sup> while in other cases there is clear evidence of interactions being formed with adjacent metal centres (Fig. 1b).<sup>12</sup>

The calculated electrostatic potential surfaces of some previously identified dihydrogen bond acceptors<sup>10–13</sup> provide an alternative view consistent with the structural characteristics of polar dihydrogen bonds and the involvement of both M $\cdots$ H and H $\cdots$ H interactions (Fig. 2). Notably, the electrostatic minima (highlighted in red on the second row of molecular surfaces) in Fig. 2a and b, are found over the metal centres, between the hydridic hydrogen atoms, and not over the hydridic positions themselves. Similarly, although BH<sub>3</sub>·NH<sub>3</sub> is overall neutral and often declared to be ‘isoelectronic’ with ethane,<sup>10</sup> the electrostatic potential surface of this Lewis adduct



(Fig. 2c) is quite unlike ethane (Fig. 2d);  $\text{BH}_3 \cdot \text{NH}_3$  is highly polar and electrostatic minima are localised over the boron atom, which bears a formal negative charge (Fig. 2c). The magnitudes and locations of the electrostatic minima in the highly polarised examples shown in Fig. 2a–c provide a simple rationalisation for the formation of favourable electrostatic interactions with  $\delta^+$  H-bond donors with geometries similar to that shown in Fig. 1b. However, the ability of much more weakly polarised hydrogen atoms to accept H-bonds in solution is yet to be examined.

Calculated electrostatic potentials reveal the trialkyl Si–H group as a potential H-bond acceptor (Fig. 2e). Since silicon has a lower Pauling electronegativity than hydrogen (1.90 vs. 2.20),<sup>14</sup> the hydrogen atom directly bonded to the silicon centre is slightly negatively charged. In line with this prediction, recent spectroscopic studies and computations have shown that trialkyl-silanes are able to accept H-bonds in the gas-phase, but no thermodynamic data, nor the relevance of such interactions in solution have been determined to date.<sup>15</sup> Trimethyl silane, as shown in Fig. 2e, is a gas at room temperature, and thus is not suited to the experimental investigation of its H-bond properties in solution. However, trihexylsilane (Fig. 3) has a very similar electrostatic surface potential (with a minimum of  $-47 \text{ kJ mol}^{-1}$ , Fig. S3, ESI $\ddagger$ ), but also has a boiling point of  $160 \text{ }^\circ\text{C}$  and good solubility in organic solvents. The anticipated weakness of the H-bond acceptor ability of the Si–H group in trihexylsilane presents a number of challenges to the measurement of such a weak interaction using a titration-based approach.

With regards to the selection of a suitable H-bond donor, perfluoro-*tert*-butanol 2 (Fig. 3) is one of the strongest known single H-bond donors (with an ESP maximum of  $+292 \text{ kJ mol}^{-1}$  Fig. S2, ESI $\ddagger$ ) and has been employed extensively in previous studies of H-bonds by Hunter, Abraham and co-workers.<sup>16–18</sup> Meanwhile, cyclohexane is one of the least competitive solvents known that maintains reasonable solubility of many organic molecules.<sup>17</sup> Furthermore, potential secondary H-bonding or dispersion interactions involving the alkyl chains of trialkylsilane would cancel out due to similarity of the competitive interactions with the alkane solvent.<sup>6–8</sup> Thus, perfluoro-*tert*-butanol and cyclohexane were selected as the H-bond donor

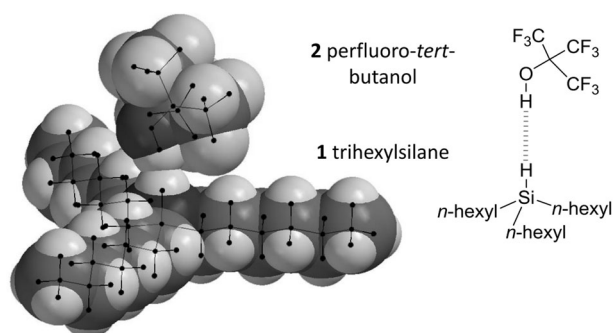


Fig. 3 Minimised geometry of the perfluoro-*tert*-butanol (**2**) and trihexylsilane (**1**) complex showing overlap of the van der Waals radii of hydrogen atoms involved in the Si–H...HO interaction. Geometry data for the complex shown (calculated using B3LYP/cc-pVTZ) and those for other theory/basis set combinations are provided in the ESI $\ddagger$ .

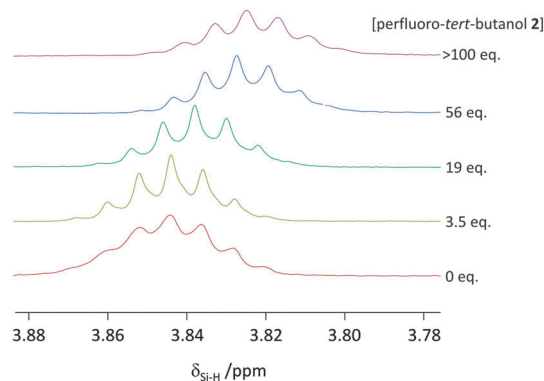


Fig. 4 Upfield  $^1\text{H-NMR}$  chemical shift seen for the Si–H in trihexylsilane (20 mM) in cyclohexane- $d_{12}$  upon increasing the concentration of perfluoro-*tert*-butanol at 298 K.

and solvent respectively for examining the H-bond acceptor ability of the Si–H group found in trihexylsilane (Fig. 3).

When a hydrogen atom acts as a classic H-bond donor, a downfield shift is observed in  $^1\text{H-NMR}$  spectra. On the contrary, when a hydrogen atom acts as a H-bond acceptor then a characteristic upfield  $^1\text{H-NMR}$  shift is observed.<sup>19</sup> An upfield shift was observed in  $\delta_{\text{Si-H}}$  of trihexylsilane upon addition of perfluoro-*tert*-butanol in cyclohexane- $d_{12}$  (Fig. 4). Although small, this upfield shift provides evidence that the Si–H is acting as a (very weak) H-bond acceptor rather than a H-bond donor. It is also worth mentioning, that in contrast to electron-rich metal hydrides (which often have  $^1\text{H-NMR}$  chemical shifts between  $\delta -5$  and  $-25 \text{ ppm}$ <sup>20</sup>), the  $\delta_{\text{Si-H}}$  of trihexylsilane has a chemical shift of approximately  $+3.8 \text{ ppm}$  (Fig. 4), in line with its weak nucleophilicity in the absence of an activating base.<sup>21</sup>

Due to the small change in the observed  $^1\text{H-NMR}$  chemical shift of the Si–H signal upon complexation by perfluoro-*tert*-butanol, and since  $\delta_{\text{F}}$  is more sensitive to electronic changes than  $\delta_{\text{H}}$ ,<sup>19</sup>  $^{19}\text{F-NMR}$  was employed for determining thermodynamic data. Perfluoro-*tert*-butanol shows no evidence of self-association even in very apolar solvents such as perfluorohexane.<sup>16</sup> Thus, the low, constant concentration of perfluoro-*tert*-butanol means that self-association of the perfluoro-*tert*-butanol does not complicate the titration data. The  $\delta_{\text{F}}$  of perfluoro-*tert*-butanol (at a constant concentration of  $1 \text{ mM}$ ) moved  $0.6 \text{ ppm}$  downfield as the concentration of trihexylsilane (**1**) was increased in cyclohexane implicating the formation of the desired complex (Fig. 3 and 5). In contrast, a control titration in

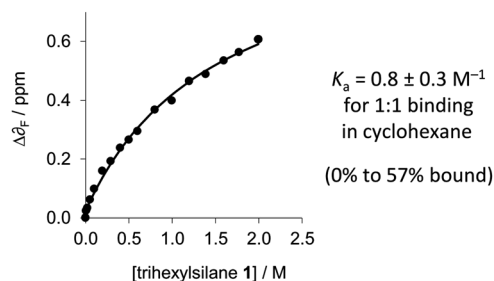


Fig. 5 1:1 binding data for the titration of trihexylsilane (**1**) against a constant concentration of perfluoro-*tert*-butanol (**2**) ( $1 \text{ mM}$  in cyclohexane) at 298 K. Additional data are presented in the ESI $\ddagger$ .



which trihexylsilane (**1**) was substituted with tetraethylsilane (which lacks an Si–H acceptor) gave a  $\delta_F$  of only 0.04 ppm (Fig. S2, ESI†) over the same concentration range. Data from three titrations with trihexylsilane (**1**) were found to fit a simple 1:1 binding model, from which a binding constant of  $0.8 \pm 0.3 \text{ M}^{-1}$  (95% confidence interval) was determined in cyclohexane at 298 K (Table S1, ESI†). The small binding constant combined with the downfield  $\delta_F$  shift and the upfield  $\delta_{\text{Si-H}}$  shift indicates that Si–H is indeed a very weak H-bond acceptor in solution.

Further support for the existence of a favourable dihydrogen bond between trihexylsilane and perfluoro-*tert*-butanol was provided by a series of HF, MP2, DFT (B3LYP), and DFT-D (M06 and  $\omega$ B97X-D) geometry optimisations (Fig. 3, left and Table S2, ESI†). Interaction energies for the trihexylsilane and perfluoro-*tert*-butanol complex were calculated as the difference between the energies of the complex and the isolated complex components, and were corrected for basis set superposition errors *via* the standard counterpoise method.<sup>22</sup> Given that the approximate +6 kJ mol<sup>-1</sup> entropic cost associated with biomolecular association in solution<sup>23</sup> is not taken into account *in silico*, the calculated gas-phase interaction energies of  $-6.5 \pm 2.8 \text{ kJ mol}^{-1}$  for the HF and DFT calculations (Table S2, ESI†) are remarkably consistent with our experimentally determined association constant of  $K_a = 0.8 \pm 0.3 \text{ M}^{-1}$  (*i.e.*  $\Delta G$  in cyclohexane  $\approx 0$ ). In contrast, but in line with other findings,<sup>6</sup> the MP2 and DFT-D methods significantly overestimate the complexation energy by several to tens of kJ mol<sup>-1</sup>. This is because gas-phase MP2 and DFT-D calculations approximate the quite significant attractive dispersion interactions between the complex components, but fail to consider the damping effects of competitive dispersion interactions with the solvent.<sup>6</sup>

Minimised SiH...HO distances varied between 1.69 Å and 1.99 Å (Table S2, ESI†) depending on the level of theory, falling within the range of classical H-bond lengths (1.6–2.0 Å).<sup>24</sup> Though such distances are well within the combined van der Waals radii of two hydrogen atoms (2.4 Å, Fig. 3),<sup>25</sup> the use of such a metric for the identification of H-bonding interactions has been cautioned against.<sup>1</sup> The Si–H...H angles observed in the calculations covered the range  $153 \pm 26^\circ$ , though it should be added that these angles are likely to be strongly influenced by steric interactions between the bulky alkyl chains and the CF<sub>3</sub> groups, rather than arising as a consequence of electronic effects. The O–H and Si–H bonds were also elongated in the bound state compared to the free state ( $+0.004 \pm 0.003 \text{ Å}$  and  $+0.009 \pm 0.003 \text{ Å}$  respectively), the former, but not the necessarily the latter, being concomitant with H-bond formation.<sup>1</sup>

In conclusion, the binding constant between a weakly polarised Si–H H-bond acceptor and a strong H-bond donor was determined for the first time in solution. Experimental NMR and computational

data are consistent with a weak Si–H...HO hydrogen bond.<sup>1</sup> Thus, the answer to the question in the title of this paper is, yes, non-polar hydrogen atoms can accept H-bonds, but only just. In addition, we show that electrostatic potential surfaces provide a simple tool for rationalising the geometric preferences of dihydrogen bonds involving hydrogen acceptors with pronounced hydridic character (Fig. 2).

We thank the EPSRC (EP/H02056X/1), Afton Chemical Ltd, MTEM and the School of Chemistry for financial support.

## Notes and references

- G. R. Desiraju, *Angew. Chem., Int. Ed.*, 2011, **50**, 52–59.
- I. Alkorta and J. Elguero, *Chem. Soc. Rev.*, 1998, **27**, 163–170.
- H. Adams, K. D. M. Harris, G. A. Hembury, C. A. Hunter, D. Livingstone and J. F. McCabe, *Chem. Commun.*, 1996, 2531–2532.
- A. Kovács and Z. Varga, *Coord. Chem. Rev.*, 2006, **250**, 710–727; J. D. Dunitz and R. Taylor, *Chem.–Eur. J.*, 1997, **3**, 89–98; J. A. K. Howard, V. J. Hoy, D. O'Hagan and G. T. Smith, *Tetrahedron*, 1996, **52**, 12613–12622.
- V. Aquilanti, E. Cornicchi, M. Moix Teixidor, N. Saendig, F. Pirani and D. Cappelletti, *Angew. Chem., Int. Ed.*, 2005, **44**, 2356–2360.
- L. Yang, C. Adam, G. S. Nichol and S. L. Cockroft, *Nat. Chem.*, 2013, DOI: 10.1038/nchem.1779.
- S. Otto, *Chem. Sci.*, 2013, **4**, 2953–2959.
- C. A. Hunter, *Chem. Sci.*, 2013, **4**, 834–848.
- P. R. Schreiner, L. V. Chernish, P. A. Gunchenko, E. Y. Tikhonchuk, H. Hausmann, M. Serafin, S. Schlecht, J. E. P. Dahl, R. M. K. Carlson and A. A. Fokin, *Nature*, 2011, **477**, 308–311; T. Steiner and G. R. Desiraju, *Chem. Commun.*, 1998, 891–892.
- R. Custelcean and J. E. Jackson, *Chem. Rev.*, 2001, **101**, 1963–1980.
- R. H. Crabtree, P. E. M. Siegbahn, O. Eisenstein, A. L. Rheingold and T. F. Koetzle, *Acc. Chem. Res.*, 1996, **29**, 348–354.
- N. V. Belkova, E. S. Shubina and L. M. Epstein, *Acc. Chem. Res.*, 2005, **38**, 624–631.
- N. V. Belkova, O. A. Filippov, A. M. Filin, L. N. Teplitskaya, Y. V. Shmyrova, V. V. Gavrilenko, L. M. Golubinskaya, V. I. Bregadze, L. M. Epstein and E. S. Shubina, *Eur. J. Inorg. Chem.*, 2004, 3453–3461.
- A. L. Allred, *J. Inorg. Nucl. Chem.*, 1961, **17**, 215–221.
- H. Ishikawa, A. Saito, M. Sugiyama and N. Mikami, *J. Chem. Phys.*, 2005, **123**, 224309; E. J. Juárez-Pérez, C. Viñas, F. Teixidor and R. Núñez, *J. Organomet. Chem.*, 2009, **694**, 1764–1770.
- J. L. Cook, C. A. Hunter, C. M. Low, A. Perez-Velasco and J. G. Vinter, *Angew. Chem., Int. Ed.*, 2007, **46**, 3706–3709; M. H. Abraham, P. L. Grellier, D. V. Prior, P. P. Duce, J. J. Morris and P. J. Taylor, *J. Chem. Soc., Perkin Trans. 2*, 1989, 699–711.
- R. Cabot, C. A. Hunter and L. M. Varley, *Org. Biomol. Chem.*, 2010, **8**, 1455–1462.
- R. Cabot and C. A. Hunter, *Org. Biomol. Chem.*, 2010, **8**, 1943–1950.
- E. S. Shubina, N. V. Belkova, A. N. Krylov, E. V. Vorontsov, L. M. Epstein, D. G. Gusev, M. Niedermann and H. Berke, *J. Am. Chem. Soc.*, 1996, **118**, 1105–1112; D. Donghi, T. Beringhelli, G. D'Alfonso and M. Mondini, *Chem.–Eur. J.*, 2006, **12**, 1016–1025.
- P. Hrobárik, V. Hrobáriková, F. Meier, M. Repiský, S. Komorovský and M. Kaupp, *J. Phys. Chem. A*, 2011, **115**, 5654–5659.
- A. D. Dilman and S. L. Ioffe, *Chem. Rev.*, 2003, **103**, 733–772.
- S. F. Boys and F. Bernardi, *Mol. Phys.*, 1970, **19**, 553–566.
- C. A. Hunter, *Angew. Chem., Int. Ed.*, 2004, **43**, 5310–5324.
- T. Steiner, *J. Phys. Chem. A*, 1998, **102**, 7041–7052.
- S. Grimme and P. R. Schreiner, *Angew. Chem., Int. Ed.*, 2011, **50**, 12639–12642.

