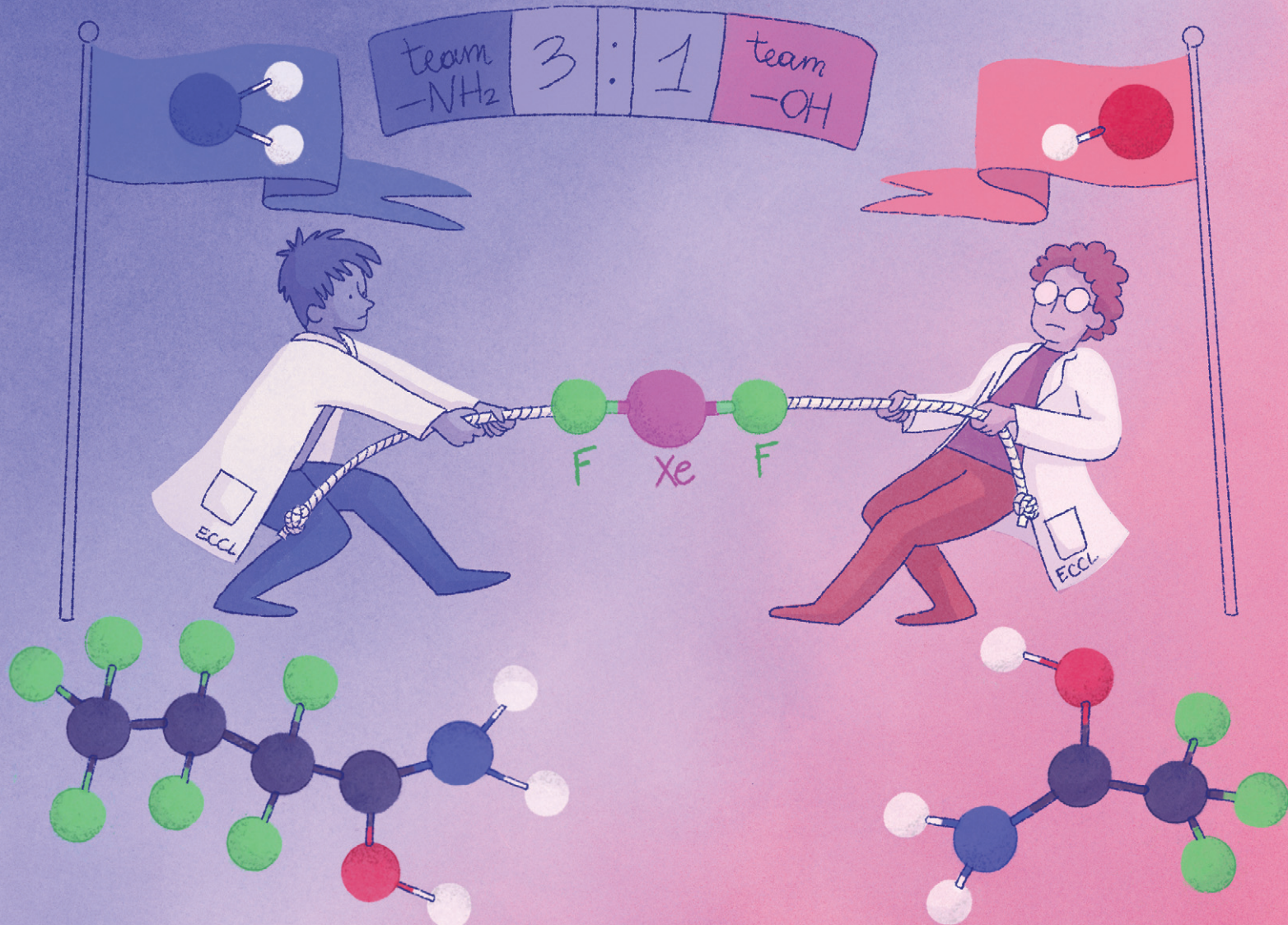


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**PAPER**

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## Hydrogen-bonded salt cocrystals of xenon difluoride and protonated perfluoroamides†

Erik Uran <sup>ab</sup> and Matic Lozinšek <sup>\*ab</sup>

The hydrogen-bonding ability of XeF<sub>2</sub> is an important factor influencing its chemical properties and reactivity, yet structurally characterised examples of hydrogen-bonded xenon fluorides remain rare. In this work, three salt cocrystals containing hydrogen-bonded xenon difluoride and hexafluoroarsenate salts of protonated perfluoroamides—CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>].XeF<sub>2</sub>, C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>].XeF<sub>2</sub>, and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>].XeF<sub>2</sub>—were synthesised and structurally characterised. Diverse hydrogen-bonding motifs were observed, and the first crystallographically characterised examples of N–H⋯FXeF hydrogen bonds are presented. In total, eleven new crystal structures are reported, including two perfluoroamides, three protonated and two hemiprotonated perfluoroamides, and one salt cocrystal containing an oxonium ion. The XeF<sub>2</sub>-containing cocrystals demonstrate that XeF<sub>2</sub> reliably functions as a hydrogen-bond acceptor and readily forms hydrogen-bonded cocrystals. These findings broaden the scope of noble-gas chemistry and highlight the potential of noble-gas fluorides for cocrystal formation.

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### Introduction

Xenon difluoride (XeF<sub>2</sub>) is the most common and extensively studied binary noble-gas fluoride and serves as a precursor to a wide range of xenon compounds.<sup>1,2</sup> It is a nonpolar molecular compound with linear geometry. XeF<sub>2</sub> is a good fluoride-ion donor<sup>3</sup> and thus forms a variety of Lewis acid–base adducts<sup>4</sup> and a plethora of coordination compounds with metal cations.<sup>1,2,5</sup>

The ability of XeF<sub>2</sub> to act as a hydrogen-bond acceptor strongly influences its physical and chemical properties. As a nonpolar molecule, it is highly soluble in the polar protic solvent anhydrous HF (aHF) (167 g/100 g at 30 °C).<sup>6</sup> This unusually high solubility arises from the formation of FXe–F⋯HF hydrogen bonds.<sup>7–9</sup> Furthermore, XeF<sub>2</sub> dissolved in aHF is a considerably more potent oxidiser than pure XeF<sub>2</sub>, and even trace amounts of HF can catalyse its reactions with organic substrates through the hydrogen-bonding induced polarisation [FXe<sup>δ+</sup>–F<sup>δ–</sup>⋯HF].<sup>10</sup> HF also facilitates fluorine exchange in XeF<sub>2</sub>, enabling the synthesis of <sup>18</sup>F-radiolabelled XeF<sub>2</sub>.<sup>11,12</sup> In certain cases, the influence of HF is so pronounced that it can unexpectedly alter reaction outcomes, even when inadvertently generated by reaction with the vessel material.<sup>13</sup>

Despite the ability of XeF<sub>2</sub> to act as a hydrogen-bond acceptor, systematic crystallographic investigations are absent, and reported solid-state examples remain scarce. To date, only a handful of crystallographically characterised examples of hydrogen-bonded XeF<sub>2</sub> have been described. These include O–H⋯FXeF hydrogen bonds observed in H<sub>3</sub>O[AsF<sub>6</sub>].2XeF<sub>2</sub> and in HNO<sub>3</sub>.XeF<sub>2</sub> cocrystals,<sup>14,15</sup> as well as an F–H⋯FXeF interaction observed in the coordination complex [Cd(HF)<sub>2</sub>(XeF<sub>2</sub>)(MF<sub>6</sub>)<sub>2</sub>] (M = Ta, Nb).<sup>5,16</sup>

It has also been shown spectroscopically that protonated trifluoroacetamide (CF<sub>3</sub>CONH<sub>2</sub>) forms a hydrogen-bonded salt cocrystal<sup>17</sup> with XeF<sub>2</sub>, CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>].XeF<sub>2</sub>.xHF.<sup>18</sup> This cocrystal is particularly noteworthy, as it may feature both =OH<sup>+</sup> and –NH<sub>2</sub> groups as hydrogen-bond donors,<sup>18</sup> potentially offering insight into the hydrogen-bonding preferences of XeF<sub>2</sub>.

To investigate the hydrogen-bonding propensity of XeF<sub>2</sub> in the solid state and its tendency to form cocrystals with NH and OH hydrogen-bond donors, the crystal structures of XeF<sub>2</sub> salt cocrystals with protonated CF<sub>3</sub>CONH<sub>2</sub>, C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub>, and C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub> were studied in this work. The perfluoroamides were selected because of their anticipated resistance to oxidative-fluorination by XeF<sub>2</sub>.

### Results and discussion

#### Crystal structures of CF<sub>3</sub>CF<sub>2</sub>CONH<sub>2</sub> and CF<sub>3</sub>CF<sub>2</sub>CF<sub>2</sub>CONH<sub>2</sub>

The crystal structures of pentafluoropropionamide (C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub>) and heptafluorobutyramide (C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>) were elucidated by low-temperature single-crystal X-ray diffraction (LT SCXRD)

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† Dedicated to Professor Resnati, celebrating a career in fluorine and noncovalent chemistry on the occasion of his 70th birthday.



**Table 1** Summary of crystal data and refinement results for crystal structures of amides and protonated amides

Compound	C <sub>2</sub> F <sub>5</sub> CONH <sub>2</sub>	C <sub>3</sub> F <sub>7</sub> CONH <sub>2</sub>	CF <sub>3</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ]	C <sub>2</sub> F <sub>5</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ]	C <sub>3</sub> F <sub>7</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ]
Space group	<i>C2/c</i>	<i>P</i> $\bar{1}$	<i>P2</i> <sub>1</sub> / <i>c</i>	<i>Pccn</i>	<i>P2</i> <sub>1</sub> / <i>c</i>
<i>a</i> (Å)	21.7871(5)	5.11713(18)	9.81910(18)	8.12957(13)	6.17592(15)
<i>b</i> (Å)	5.11704(12)	5.27137(14)	7.90095(13)	25.2768(4)	7.94187(19)
<i>c</i> (Å)	10.0754(3)	12.7768(3)	20.5015(4)	9.34322(16)	21.7914(5)
$\alpha$ (°)	90	95.467(2)	90	90	90
$\beta$ (°)	98.140(2)	91.890(3)	98.6498(18)	90	96.014(2)
$\gamma$ (°)	90	105.584(3)	90	90	90
<i>V</i> (Å <sup>3</sup> )	1111.94(5)	329.847(18)	1572.42(5)	1919.93(5)	1062.95(4)
<i>M</i>	163.06	213.07	302.97	352.98	402.99
<i>Z</i>	8	2	8	8	4
<i>T</i> (K)	100	100	100	100	100
<i>R</i> [ <i>F</i> <sup>2</sup> > 2σ( <i>F</i> <sup>2</sup> )]	0.048	0.028	0.033	0.027	0.027
w <i>R</i> ( <i>F</i> <sup>2</sup> )	0.138	0.076	0.087	0.058	0.068

(Tables 1 and S1), whereas the crystal structure of CF<sub>3</sub>CONH<sub>2</sub> has been previously reported at 295 K and 110 K.<sup>19,20</sup> For comparison of bond lengths (Table S2), only the structure obtained at 110 K was considered.<sup>20</sup>

C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub> (Fig. 1a and S1) crystallizes in the monoclinic space group *C2/c* with *Z* = 8. The C=O bond length (1.2323(19) Å) is comparable to the distances observed in the crystal structures of other primary amides, and the same applies to the C–N bond (1.317(2) Å).<sup>21</sup> Two N–H⋯O hydrogen bonds (2.912(2) Å, 171(2)°; 2.8396(17) Å, 149(2)°; Table S3) in the crystal structure form R<sub>2</sub><sup>2</sup>(8) and R<sub>6</sub><sup>4</sup>(16) hydrogen-bonding motifs,<sup>22</sup> which assemble into a corrugated layer parallel to the *bc* plane (Fig. S2 and S3).

C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub> (Fig. 1b and S4) crystallizes in the triclinic space group *P* $\bar{1}$  with *Z* = 2. The C=O bond distance (1.2293(15) Å) is essentially identical to that in C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub>, as is the C–N bond (1.3162(16) Å). These bond distances are shorter than the corresponding ones observed in non-fluorinated secondary amides, such as capsaicin.<sup>23</sup> Two N–H⋯O hydrogen bonds

(Table S4) are present in the crystal structure (2.9313(14) Å, 174.3(16)°; 2.8495(14) Å, 140.4(15)°), which fall within the typical range for amide molecules.<sup>21</sup> The R<sub>2</sub><sup>2</sup>(8) and R<sub>4</sub><sup>4</sup>(8) hydrogen-bond motifs link the molecules into a ladder along the *a*-crystallographic axis (Fig. S5).

#### Protonation of amides in superacidic media HF–AsF<sub>5</sub>

All amides are soluble in aHF and readily undergo protonation upon addition of AsF<sub>5</sub>. In all cases, protonation occurs at the oxygen atom, consistent with previous observations.<sup>18,24–26</sup> Low-temperature crystallisation from aHF afforded crystals of suitable quality for SCXRD.

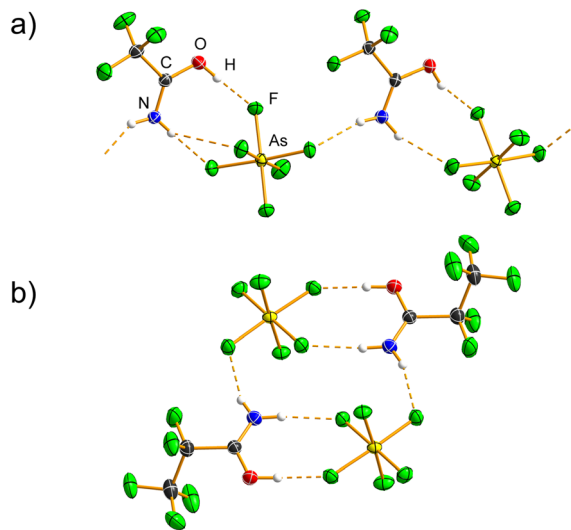
CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>−</sup> (Tables 1, S1 and S2; Fig. S6) crystallises in the monoclinic space group *P2*<sub>1</sub>/*c* with *Z* = 8 and *Z'* = 2. Upon protonation, the C=O bonds (1.2795(19), 1.282(2) Å) lengthen and the C–N bonds (1.279(2), 1.281(2) Å) shorten relative to those in CF<sub>3</sub>CONH<sub>2</sub> (1.2304(12) and 1.3164(13) Å, respectively).<sup>20</sup> These changes in the C=O and C–N bond lengths are consistent with previous crystallographic studies of protonated amides.<sup>25–27</sup> The O–H⋯F hydrogen bonds (2.5860(16) Å, 2.6530(18) Å; Table S5) bracket the value observed in CF<sub>3</sub>C(OH)NH<sub>2</sub>[SbF<sub>6</sub>]<sup>−</sup> (2.600(1) Å), whereas the N–H⋯F hydrogen bonds (2.8236(18)–3.0797(18) Å) are comparable to those in CF<sub>3</sub>C(OH)NH<sub>2</sub>[SbF<sub>6</sub>]<sup>−</sup> (2.884(2), 2.933(2) Å).<sup>26</sup> All hydrogen-bond angles (121(2)–179(3)°) fall within the typical range. The [AsF<sub>6</sub>]<sup>−</sup> anions deviate from ideal octahedral geometry, with the longest As–F bonds (1.7524(10), 1.7557(10) Å) participating in hydrogen bonding with =OH<sup>+</sup> group. In the crystal structure, cations and anions are linked through O–H⋯F and N–H⋯F hydrogen-bonded chains (Fig. 2a and S7).

C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>−</sup> (Tables 1, S1 and S2) crystallises in the orthorhombic space group *Pccn* with *Z* = 8 and features a disordered –C<sub>2</sub>F<sub>5</sub> moiety (Fig. S8). Perfluorinated alkyl chains frequently exhibit disorder in the crystalline state,<sup>28</sup> as F⋯F interactions are relatively weak,<sup>29</sup> and can therefore adopt various conformations. The C=O (1.2821(15) Å) and C–N (1.2772(16) Å) bonds are longer and shorter, respectively, than those in C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub>. The [AsF<sub>6</sub>]<sup>−</sup> anion deviates from ideal octahedral geometry, with the *mer*-As–F bonds involved



**Fig. 1** R<sub>2</sub><sup>2</sup>(8) hydrogen-bonding motifs in the crystal structures of (a) C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub> and (b) C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>. Hydrogen bonds are shown as dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.



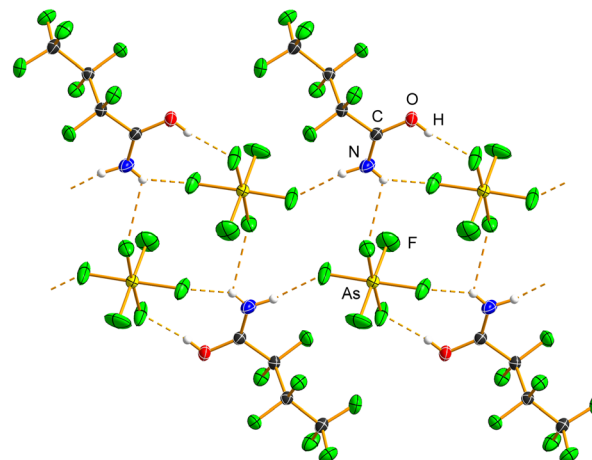


**Fig. 2** Crystal structure of (a) the hydrogen-bonded chain in  $\text{CF}_3\text{-C(OH)NH}_2[\text{AsF}_6]$  and (b) the discrete hydrogen-bonded cluster in  $\text{C}_2\text{F}_5\text{C(OH)NH}_2[\text{AsF}_6]$  (only one orientation of the disordered  $-\text{C}_2\text{F}_5$  unit is shown). Hydrogen bonds are shown as dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.

in hydrogen bonding being longer (1.7253(8)–1.7453(8) Å) than the remaining As–F bonds (1.6976(8)–1.7111(8) Å). The hydrogen bonds (Table S6) between the  $=\text{OH}^+$  and  $-\text{NH}_2$  groups and the  $[\text{AsF}_6]^-$  anions ( $\text{O(H)}\cdots\text{F}$ , 2.6006(12) Å,  $172(2)^\circ$ ;  $\text{N(H)}\cdots\text{F}$ , 2.8309(13) Å,  $174(2)^\circ$  and 2.8316(14) Å,  $161.3(19)^\circ$ ) lead to the formation of discrete units (Fig. 2b, S8 and S9), exhibiting  $\text{R}_2^2(8)$  and  $\text{R}_4^4(12)$  hydrogen-bonding motifs.

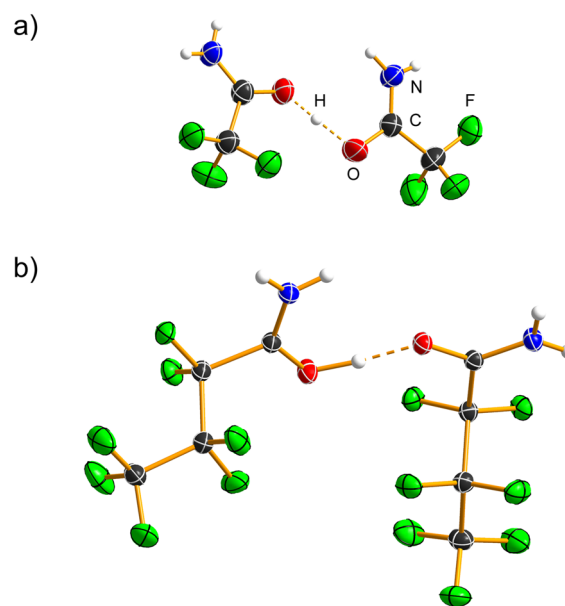
$\text{C}_3\text{F}_7\text{C(OH)NH}_2[\text{AsF}_6]$  (Tables 1, S1 and S2) crystallises in the monoclinic space group  $P2_1/c$  with  $Z = 4$ , with the  $[\text{AsF}_6]^-$  anion disordered over two positions (Fig. S10). The C=O bond (1.2797(14) Å) is elongated, and the C–N bond (1.2841(16) Å) is shortened compared to those in  $\text{C}_3\text{F}_7\text{CONH}_2$ . A similar C=O(H) bond distance (1.274(2) Å) was observed in the crystal structure of  $(\text{C}_6\text{F}_5)_2\text{COH}[\text{AsF}_6]$ .<sup>30</sup> Hydrogen bonds (Table S7) are formed between the  $=\text{OH}^+$  group (2.541(3), 2.557(3) Å;  $157(3)$ ,  $165(3)^\circ$ ) or the  $-\text{NH}_2$  group (2.737(4)–3.179(5) Å,  $118.0(19)$ – $168.1(19)^\circ$ ) and the  $[\text{AsF}_6]^-$  anions. The  $\text{O}\cdots\text{F}$  hydrogen bond is the shortest among the protonated amides in this study, and also shorter than those in  $\text{CH}_3\text{C(OH)NH}_2[\text{AsF}_6]$ <sup>25</sup> and  $\text{CF}_3\text{C(OH)NH}_2[\text{SbF}_6]$ .<sup>26</sup> The  $[\text{AsF}_6]^-$  anion deviates from ideal octahedral geometry (1.642(3)–1.795(2) Å). The  $\text{C}_3\text{F}_7\text{C(OH)NH}_2^+$  cations and  $[\text{AsF}_6]^-$  anions are linked into a hydrogen-bonded ribbon (Fig. 3 and S11), exhibiting conjoined  $\text{R}_4^4(12)$ ,  $\text{R}_2^2(8)$  and  $\text{R}_2^2(8)$  motifs.

Two hemiprotonated salts,  $(\text{CF}_3\text{CONH}_2)_2\text{H}[\text{AsF}_6]$  and  $(\text{C}_3\text{F}_7\text{CONH}_2)_2\text{H}[\text{AsF}_6]$  (Fig. 4 and S12–S17; Tables 2, S1 and S2), were also crystallographically characterised. The former was inadvertently found during the low-temperature crystal selection and mounting of the  $\text{CF}_3\text{C(OH)NH}_2[\text{AsF}_6]\cdot\text{XeF}_2$  sample, whereas the latter was identified as an impurity in



**Fig. 3** Hydrogen-bonded ribbon in  $\text{C}_3\text{F}_7\text{C(OH)NH}_2[\text{AsF}_6]$  (only one orientation of the disordered  $[\text{AsF}_6]^-$  anion is shown). Hydrogen bonds are shown as dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.

the sample of  $\text{C}_3\text{F}_7\text{C(OH)NH}_2[\text{AsF}_6]\cdot\text{XeF}_2$  salt cocrystals. Both compounds crystallise in the triclinic space group  $P\bar{1}$  with  $Z = 2$ . In both structures, the C=O bonds are elongated (1.283(6) Å in  $(\text{CF}_3\text{CONH}_2)_2\text{H}[\text{AsF}_6]$ ; 1.2652(9), 1.2459(9) Å in  $(\text{C}_3\text{F}_7\text{CONH}_2)_2\text{H}[\text{AsF}_6]$ ), whereas the C–N bonds are shortened (1.274(7) Å in  $(\text{CF}_3\text{CONH}_2)_2\text{H}[\text{AsF}_6]$ ; 1.2942(10), 1.3027(10) Å in  $(\text{C}_3\text{F}_7\text{CONH}_2)_2\text{H}[\text{AsF}_6]$ ) compared to the non-protonated amides.<sup>20</sup> The values for one of the amide molecules in  $(\text{CF}_3\text{CONH}_2)_2\text{H}[\text{AsF}_6]$  fall within the



**Fig. 4** Hydrogen-bonded dimers in the crystal structure of (a)  $(\text{CF}_3\text{CONH}_2)_2\text{H}[\text{AsF}_6]$  and (b)  $(\text{C}_3\text{F}_7\text{CONH}_2)_2\text{H}[\text{AsF}_6]$ . The short  $\text{O}\cdots\text{H}\cdots\text{O}=\text{C}$  hydrogen bonds are shown as dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.



**Table 2** Summary of crystal data and refinement results for hemiprotonated amides and H<sub>3</sub>O[AsF<sub>6</sub>] $\cdot$ 2CF<sub>3</sub>CONH<sub>2</sub> salt cocrystal

Compound	(CF <sub>3</sub> CONH <sub>2</sub> ) <sub>2</sub> H[AsF <sub>6</sub> ]	(C <sub>3</sub> F <sub>7</sub> CONH <sub>2</sub> ) <sub>2</sub> H[AsF <sub>6</sub> ]	H <sub>3</sub> O[AsF <sub>6</sub> ] $\cdot$ 2CF <sub>3</sub> CONH <sub>2</sub>
Space group	<i>P</i> $\bar{1}$	<i>P</i> $\bar{1}$	<i>Pnma</i>
<i>a</i> (Å)	5.2815(3)	5.32051(4)	11.5349(2)
<i>b</i> (Å)	10.1517(6)	10.45222(9)	14.0649(3)
<i>c</i> (Å)	12.4911(6)	16.10170(12)	7.78994(14)
$\alpha$ (°)	108.936(5)	90.5740(6)	90
$\beta$ (°)	93.107(5)	90.8760(6)	90
$\gamma$ (°)	102.904(5)	103.0402(7)	90
<i>V</i> (Å <sup>3</sup> )	611.63(6)	872.156(12)	1263.82(4)
<i>M</i>	416.02	616.06	434.04
<i>Z</i>	2	2	4
<i>T</i> (K)	100	100	100
$R[F^2 > 2\sigma(F^2)]$	0.047	0.022	0.037
$wR(F^2)$	0.125	0.055	0.085

range for neutral amide,<sup>20</sup> owing to the relatively high standard uncertainties of the bond lengths. The O(H) $\cdots$ O hydrogen bond length in (CF<sub>3</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>] (2.426(5) Å, 170(8)°) is essentially identical to that in (C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>] (2.4174(9) Å, 172(2)°) (Tables S8 and S9), and comparable to literature values for such hydrogen-bonded systems.<sup>21</sup> The nearly equidistant position of the hydrogen atom (O–H, H $\cdots$ O: 1.13(9), 1.31(9) Å in (CF<sub>3</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>]; 1.06(2), 1.36(2) Å in (C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>]), together with the relatively short O $\cdots$ O distances, indicates strong, positive charge-assisted hydrogen bonding, (+) CAHB.<sup>31</sup> These structures represent rare examples of proton sharing between two primary amide molecules,<sup>21,32</sup> a motif more commonly observed in secondary and tertiary amides.<sup>21</sup> The –NH<sub>2</sub> groups are hydrogen-bonded to [AsF<sub>6</sub>]<sup>–</sup> anions (N $\cdots$ F, 2.644(5)–3.005(5) Å in (CF<sub>3</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>]; 2.8168(9)–3.1046(9) Å in (C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>]) (Fig. S13–S17), resulting in the formation of ribbons that are further interconnected by the anions into layers parallel to the *ab* plane.

A crystal of H<sub>3</sub>O[AsF<sub>6</sub>] $\cdot$ 2CF<sub>3</sub>CONH<sub>2</sub> (Tables 2, S1 and S2; Fig. 5 and S18–S20) was fortuitously found during the low-temperature crystal selection and mounting of the CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>] $\cdot$ XeF<sub>2</sub> sample. It crystallises in the orthorhombic space group *Pnma* with *Z* = 4. The amide molecule is not protonated, resulting in C=O (1.236(4) Å) and C–N (1.304(4) Å) bond lengths that are close to those in CF<sub>3</sub>CONH<sub>2</sub>.<sup>20</sup> The amide molecule acts as both a hydrogen-bond donor and acceptor (Table S10, Fig. S19), forming N–H $\cdots$ F(As) and N–H $\cdots$ O(C) hydrogen bonds. An R<sub>2</sub><sup>2</sup>(8) motif is observed between two amide molecules, with the N $\cdots$ O hydrogen bond (2.959(4) Å, 171(4)°) comparable to that found in CF<sub>3</sub>CONH<sub>2</sub>.<sup>20</sup> The H<sub>3</sub>O<sup>+</sup> cation forms three hydrogen bonds: two symmetrically equivalent O–H $\cdots$ O(C) (2.525(3) Å, 167(4)°) and one O–H $\cdots$ F hydrogen bond (2.657(5) Å, 173(7)°) with the [AsF<sub>6</sub>]<sup>–</sup> anion. Together, these hydrogen bonds form a hydrogen-bonded cluster represented by R<sub>6</sub><sup>6</sup>(20), R<sub>4</sub><sup>4</sup>(14) and R<sub>2</sub><sup>2</sup>(8) graph-set motifs<sup>22</sup> (Fig. 5), which further extend into a layer parallel to the *bc* plane (Fig. S20).

### Hydrogen-bonded salt cocrystals of XeF<sub>2</sub>

The reaction of amides with equimolar amounts of [XeF]<sup>+</sup>[AsF<sub>6</sub>]<sup>–</sup> at temperatures down to –30 °C leads to the formation of RC(OH)NH<sub>2</sub>[AsF<sub>6</sub>] $\cdot$ XeF<sub>2</sub> salt cocrystals. This indicates that a proton from HF is transferred to the amide, generating a protonated amide, while the resulting fluoride anion reacts with [XeF]<sup>+</sup> to form XeF<sub>2</sub>. This behaviour was also reported in a previous study of the CF<sub>3</sub>CONH<sub>2</sub>–[XeF][AsF<sub>6</sub>] system.<sup>18</sup>

The salt cocrystals (Tables 3, S1 and S2) thus feature protonated amides cocrystallised with XeF<sub>2</sub> and exhibit a rare O–H $\cdots$ FXeF hydrogen bond, as well as the first crystallographically characterised examples of N–H $\cdots$ FXeF hydrogen bonds.

CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>] $\cdot$ XeF<sub>2</sub> (Fig. 6 and S21) crystallises in the monoclinic space group *P2*<sub>1</sub>/*n* with *Z* = 4. The XeF<sub>2</sub> molecule exhibits slight asymmetry in Xe–F bond distances (1.9669(10), 2.0237(9) Å) compared to pure XeF<sub>2</sub> (1.999(4) Å),<sup>33</sup> and it remains linear (178.10(5) Å). The asymmetry of XeF<sub>2</sub> is slightly smaller than that observed in XeF<sub>2</sub> $\cdot$ HNO<sub>3</sub> (1.9737(8), 2.0506(8) Å).<sup>15</sup>



**Fig. 5** R<sub>6</sub><sup>6</sup>(14) hydrogen-bonded cluster in the crystal structure of the H<sub>3</sub>O[AsF<sub>6</sub>] $\cdot$ 2CF<sub>3</sub>CONH<sub>2</sub> salt cocrystal. Hydrogen bonds are shown by dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.



**Table 3** Summary of crystal data and refinement results for salt cocrystals of protonated amides with XeF<sub>2</sub>

Compound	CF <sub>3</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ] <sup>-</sup> ·XeF <sub>2</sub>	C <sub>2</sub> F <sub>5</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ] <sup>-</sup> ·XeF <sub>2</sub>	C <sub>3</sub> F <sub>7</sub> C(OH)NH <sub>2</sub> [AsF <sub>6</sub> ] <sup>-</sup> ·XeF <sub>2</sub>
Space group	<i>P</i> 2 <sub>1</sub> / <i>n</i>	<i>Aea</i> 2	<i>Pnna</i>
<i>a</i> (Å)	7.41785(9)	8.67561(10)	8.62011(14)
<i>b</i> (Å)	9.84875(11)	31.0125(4)	35.5418(5)
<i>c</i> (Å)	14.90113(17)	8.65174(9)	8.71910(12)
$\alpha$ (°)	90	90	90
$\beta$ (°)	99.4517(11)	90	90
$\gamma$ (°)	90	90	90
<i>V</i> (Å <sup>3</sup> )	1073.85(2)	2327.77(4)	2671.31(7)
<i>M</i>	472.27	522.28	572.29
<i>Z</i>	4	8	8
<i>T</i> (K)	100	100	100
$R[F^2 > 2\sigma(F^2)]$	0.025	0.018	0.026
$wR(F^2)$	0.067	0.038	0.066



**Fig. 6** Hydrogen-bonded ribbon in the crystal structure of the salt cocrystal CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub>. Only one orientation of the disordered -CF<sub>3</sub> moiety is shown. Hydrogen bonds are shown as dashed orange lines. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.

CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> is the only salt cocrystal in this series that exhibits both O-H...F(Xe) and N-H...F(Xe) hydrogen bonds. One fluorine atom of XeF<sub>2</sub> is acting as a bifurcated acceptor (Fig. 6 and S21; Table S11). The O-H...F(Xe) hydrogen bond (2.5467(14) Å, 171(3)°) is shorter than that in H<sub>3</sub>O[AsF<sub>6</sub>]<sup>-</sup>·2XeF<sub>2</sub> (2.571(3) Å)<sup>14</sup> and HNO<sub>3</sub>·XeF<sub>2</sub> (2.690(1) Å).<sup>15</sup> It is also significantly shorter than the O-H...F(As) hydrogen bonds in CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup> and C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>, but comparable to that in C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>. The N-H...F(Xe) hydrogen bonds (2.7865(15) Å, 151(3)°; 3.0894(16), 124(2)°), which involve a single bifurcated donor, are longer than those observed in the other two salt cocrystals described in this study. The C=O (1.2773(15) Å) and C-N (1.2772(16) Å) bond lengths are essentially identical to those in the protonated salts,<sup>26</sup> indicating a negligible influence of hydrogen bonding on the overall geometry of the CF<sub>3</sub>C(OH)NH<sub>2</sub><sup>+</sup> cation. The -CF<sub>3</sub> moiety is disordered, as also observed in the crystal structure of CF<sub>3</sub>CONH<sub>2</sub>.<sup>20</sup> The [AsF<sub>6</sub>]<sup>-</sup> anion participates in hydrogen bonding with the -NH<sub>2</sub> group (2.8270(18) Å, 176(3)°; 3.0594(15) Å, 111(2)°), resulting in a slight deviation from ideal octahedral geometry (As-F, 1.7006(12)–1.7417(11) Å).

Hydrogen bonds between CF<sub>3</sub>C(OH)NH<sub>2</sub><sup>+</sup> and XeF<sub>2</sub> form a zigzag chain parallel to the *b*-crystallographic axis, with pendant [AsF<sub>6</sub>]<sup>-</sup> anions connected to the chain *via* N-H...F(As) hydrogen bonds, giving rise to a ribbon-like structure (Fig. 6 and S22).

Both C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> (Tables 3, S1 and S2; Fig. S23–S26) crystallise in orthorhombic space groups, *Aea*2 and *Pnna*, respectively, with *Z* = 8. The asymmetry of the Xe-F bond lengths in C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> (1.9734(14), 2.0061(15) Å) and in C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> (1.9674(15), 2.0135(16) Å) is comparable. The shorter Xe-F bonds are similar to that observed in the trifluoroacetamide analogue, whereas the longer Xe-F bonds are significantly shorter. In both cocrystals, the F-Xe-F angle is essentially linear (179.88(9)°; 179.57(7)°).

The N-H...F(Xe) hydrogen bonds (Tables S12 and S13) in C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> (2.688(3), 2.729(3) Å) and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> (2.692(3), 2.737(3) Å) are comparable and are significantly shorter than the corresponding hydrogen bonds in CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub>. The C=O (1.289(2), 1.285(3) Å) and C-N (1.279(3), 1.280(3) Å) bond lengths in C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> are almost identical to those observed in the corresponding protonated salts.

The protonated oxygen atom acts as a hydrogen-bond donor towards the [AsF<sub>6</sub>]<sup>-</sup> anions, forming bifurcated hydrogen bonds (Fig. 7, S23 and S25), which are longer than the O-H...F(As) hydrogen bonds observed in the parent protonated salts.

The packing in both C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub> consists of hydrogen-bonded ribbons composed of alternating protonated amide and XeF<sub>2</sub> molecules, similar to those observed in CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub>. These ribbons are further connected by O-H...F(As) hydrogen bonds, and in the case of C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]<sup>-</sup>·XeF<sub>2</sub>, also by N-H...F(As) hydrogen bonds (Fig. S24 and S26).

The relatively small difference in Xe-F bond lengths in the present XeF<sub>2</sub> cocrystals suggests that hydrogen bonding has only a minor influence on XeF<sub>2</sub> ionisation (XeF<sub>2</sub> → XeF<sup>+</sup> +





Fig. 7 Hydrogen bonds (dashed orange lines) in the crystal structures of the salt cococrystals (a)  $C_2F_5C(OH)NH_2[AsF_6] \cdot XeF_2$  and (b)  $C_3F_7C(OH)NH_2[AsF_6] \cdot XeF_2$ . The OH groups are bifurcated hydrogen-bond donors; however, two  $[AsF_6]^-$  anions have been omitted for clarity. Displacement ellipsoids are drawn at the 50% probability level, and hydrogen atoms are represented as spheres of arbitrary radius.

$F^-$ ).<sup>34</sup> In particular, the shorter Xe–F bonds (1.9669(10)–1.9734(14) Å) are considerably longer than those found in  $[XeF]^+$  tight ion pairs<sup>33,35,36</sup> and in  $[Xe_2F_3]^+$  salts.<sup>35,37</sup> They are comparable to the shortest Xe–F bond lengths in  $XeF_2$  adduct-salts with  $[BrOF_2]^+$  (1.956(5), 1.960(4) Å)<sup>38</sup> and  $[BrO_2]^+$  cations (1.970(4)–1.978(3) Å).<sup>39</sup> Nevertheless, the distortion of hydrogen-bonded  $XeF_2$  observed in the present salt cococrystals is significant when compared with Xe–F bond distances observed in the crystal structures containing cococrystallised  $XeF_2$ , e.g.,  $3XeF_2 \cdot 2MnF_4$  (1.9933(7) Å),<sup>36</sup> and in the molecular cococrystals  $XeF_2 \cdot XeF_4$  (1.9940(9) Å)<sup>37</sup> and  $XeF_2 \cdot XeOF_4$  (2.014(5) Å),<sup>40</sup> in which  $XeF_2$  is centrosymmetric.

### Vibrational spectroscopy

To corroborate the findings from LT SCXRD and to gain further insight into the ionisation of  $XeF_2$ , low-temperature Raman spectra were measured (Fig. 8 and S27–S40). Two bands at 457–475 and 528–535  $cm^{-1}$  are observed in all  $XeF_2$  salt cococrystals in this study, corresponding to the elongated and shortened Xe–F bond, respectively. These bands are



Fig. 8 Raman spectra of  $XeF_2$  salt cococrystals with protonated amides recorded at low temperatures ( $-90^\circ C$ ). The green dashed line is placed at the position of free  $XeF_2$  ( $497\text{ cm}^{-1}$ )<sup>41</sup> which was observed as an impurity in the reactions.

significantly shifted from that of pure  $XeF_2$  ( $497\text{ cm}^{-1}$ )<sup>41</sup> and from values observed when cococrystallised  $XeF_2$  does not participate in significant intermolecular interactions, such as in  $XeF_2 \cdot XeOF_4$  ( $494, 503\text{ cm}^{-1}$ ),<sup>40</sup>  $XeF_2 \cdot XeF_4$  ( $505\text{ cm}^{-1}$ ),<sup>42</sup>  $3XeF_2 \cdot 2MnF_4$  ( $508\text{ cm}^{-1}$ ),<sup>36</sup> and  $XeF_2 \cdot N_2O_4$  ( $509\text{ cm}^{-1}$ ).<sup>15</sup> The value of the higher-frequency band is comparable to the Raman shifts reported for the adduct salts  $[BrOF_2]^- [AsF_6]^- XeF_2$  ( $531, 543, 559\text{ cm}^{-1}$ ),<sup>38</sup>  $[BrO_2]^- [AsF_6]^- nXeF_2$  ( $n = 1, 2; 516\text{--}546\text{ cm}^{-1}$ ),<sup>39</sup> and for the hydrogen-bonded cococrystals  $H_3O[AsF_6] \cdot 2XeF_2$  ( $552\text{ cm}^{-1}$ )<sup>14</sup> and  $HNO_3 \cdot XeF_2$  ( $529\text{ cm}^{-1}$ ).<sup>15</sup> However, these shifts are significantly smaller than those observed in  $[XeF]^+$  tight-ion pair salts ( $>600\text{ cm}^{-1}$ ) and  $[Xe_2F_3]^+$  cations ( $580\text{--}600\text{ cm}^{-1}$ ).<sup>1,33,35,36,43,44</sup> The band around  $535\text{ cm}^{-1}$  is particularly noteworthy, as this value coincides with that observed for  $XeF_2$  dissolved in aHF, which has been attributed to the  $FXe-F \cdots HF$  hydrogen bonds.<sup>7–9</sup>

In addition to the bands attributed to  $XeF_2$ , those arising from  $[AsF_6]^-$  anions are observed around 375 and 680  $cm^{-1}$ .<sup>18,25,26,45</sup> Vibrations from the protonated amide molecules are also present, including an intense band around  $800\text{ cm}^{-1}$  corresponding to  $\nu(C-C)$ ,<sup>25,26</sup> and peaks near  $1100\text{ cm}^{-1}$  attributed to C–F vibrations.<sup>18,26,46</sup> In all protonated amides, the N–H stretching vibrations were observed in  $3150\text{--}3400\text{ cm}^{-1}$  range.<sup>18,26,46</sup>



## Experimental

**Caution!** Anhydrous HF, AsF<sub>5</sub>, XeF<sub>2</sub>, [XeF][AsF<sub>6</sub>] and the compounds prepared in this study are highly reactive and hazardous. The amides used may cause skin, eye, and respiratory irritation. Contact with the skin must be avoided, and all compounds should be handled exclusively in a well-ventilated fume hood.

Appropriate safety precautions must be observed at all times, and working with minimal quantities is strongly recommended.

## Materials and methods

Reactions were carried out in fluorinated ethylene propylene (FEP) vessels equipped with Kel-F or PTFE valves. All vessels were passivated with fluorine prior to use. Volatile substances were handled using a fluorine-resistant metal vacuum line, whereas solids were manipulated inside an N<sub>2</sub>-filled glovebox. Detailed synthetic procedures are provided in the SI. Characterisation was performed by low-temperature single-crystal X-ray diffraction and low-temperature Raman spectroscopy. Single-crystal selection and mounting were carried out using a low-temperature crystal-mounting apparatus, as described previously (SI).<sup>30,36,47</sup> Low-temperature Raman spectra were recorded directly on the aluminium trough used for mounting single crystals for X-ray diffraction measurements.

## Conclusions

In this work, the perfluoroamides trifluoroacetamide (CF<sub>3</sub>-CONH<sub>2</sub>), pentafluoropropionamide (C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub>), and heptafluorobutyramide (C<sub>3</sub>F<sub>7</sub>-CONH<sub>2</sub>), were protonated in superacidic medium HF-AsF<sub>5</sub>, and the crystal structures of the resulting salts, CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>], C<sub>2</sub>F<sub>5</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>], and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>] were elucidated. Protonation at the carbonyl oxygen atom is consistently observed. In addition, the crystal structures of the amides C<sub>2</sub>F<sub>5</sub>CONH<sub>2</sub> and C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>, the hemiprotonated salts (CF<sub>3</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>] and (C<sub>3</sub>F<sub>7</sub>CONH<sub>2</sub>)<sub>2</sub>H[AsF<sub>6</sub>], and the oxonium salt cocrystal H<sub>3</sub>O[AsF<sub>6</sub>]·2CF<sub>3</sub>CONH<sub>2</sub> were determined. Low-temperature reactions of the perfluoroamides with [XeF][AsF<sub>6</sub>] in aHF yielded rare XeF<sub>2</sub>-containing salt cocrystals: CF<sub>3</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]·XeF<sub>2</sub>, C<sub>2</sub>F<sub>5</sub>-C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]·XeF<sub>2</sub> and C<sub>3</sub>F<sub>7</sub>C(OH)NH<sub>2</sub>[AsF<sub>6</sub>]·XeF<sub>2</sub>. Their crystal structures reveal a rare example of O-H...FXeF and the first crystallographically characterised cases of N-H...FXeF hydrogen bonding. The XeF<sub>2</sub> molecule is slightly polarised, as indicated by the differences observed in Xe-F bond lengths compared with those in free XeF<sub>2</sub>; this finding is corroborated by low-temperature Raman spectroscopy. The reported crystal structures display diverse hydrogen-bonding motifs involving O-H...F(Xe), N-H...F(Xe), O-H...F(As) and N-H...F(As) interactions. The salt cocrystals prepared and structurally characterised in

this study demonstrate that XeF<sub>2</sub> readily forms hydrogen-bonded cocrystals and serves as a reliable hydrogen-bond acceptor. These results open new possibilities for the exploration of cocrystal formation with noble-gas fluorides and the expansion of noble-gas chemistry.

## Author contributions

Conceptualization, data curation, formal analysis, investigation, visualization, writing – original draft: EU; funding acquisition, methodology, project administration, resources, supervision: ML; validation, writing – review & editing: EU, ML. Both authors agreed on the final version of the article.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

Supplementary information: crystallographic details, Raman spectra, experimental details. See DOI: <https://doi.org/10.1039/d5ce00956a>.

Crystallographic data for all reported crystal structures has been deposited at the Cambridge Crystallographic Data Centre (CCDC) under deposition numbers 2493130–2493140.<sup>48a–k</sup>

Data for this article, including SCXRD datasets and Raman spectra are available at Zenodo open repository at <https://doi.org/10.5281/zenodo.17432981>.

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## Notes and references

- 1 M. Tramšek and B. Žemva, *Acta Chim. Slov.*, 2006, **53**, 105–116.
- 2 D. S. Brock, G. J. Schrobilgen and B. Žemva in *Comprehensive Inorganic Chemistry II, From Elements to Applications*, ed. J. Reedijk and K. Poeppelmeier, Elsevier, Netherlands, 2013, vol. 1, pp. 755–822.
- 3 N. Bartlett and F. O. Sladky, *J. Am. Chem. Soc.*, 1968, **90**, 5316–5317.
- 4 B. Žemva, *Croat. Chem. Acta*, 1988, **61**, 163–187.
- 5 G. Tavčar and M. Tramšek, *J. Fluorine Chem.*, 2015, **174**, 14–21.
- 6 J. G. Malm, H. Selig, J. Jortner and S. A. Rice, *Chem. Rev.*, 1965, **65**, 199–236.
- 7 S. S. Nabiev and I. I. Ostroukhova, *Spectrochim. Acta, Part A*, 1993, **49**, 1537–1546.
- 8 S. S. Nabiev, *Russ. Chem. Bull.*, 1998, **47**, 535–559.



- 9 S. M.-J. Yeh, *PhD Thesis*, University of California, 1984, (available at: <https://escholarship.org/uc/item/9b1195zf>).
- 10 M. A. Tius, *Tetrahedron*, 1995, **51**, 6605–6634.
- 11 G. Schrobilgen, G. Firna, R. Chirakal and E. S. Garnett, *J. Chem. Soc., Chem. Commun.*, 1981, 198–199.
- 12 N. Vasdev, B. E. Pointner, R. Chirakal and G. J. Schrobilgen, *J. Am. Chem. Soc.*, 2002, **124**, 12863–12868.
- 13 M. M. Nielsen and C. M. Pedersen, *Chem. Sci.*, 2022, **33**, 6181–6196.
- 14 M. Gerken, M. D. Moran, H. P. A. Mercier, B. E. Pointner, G. J. Schrobilgen, B. Hoge, K. O. Christe and J. A. Boatz, *J. Am. Chem. Soc.*, 2009, **131**, 13474–13489.
- 15 M. D. Moran, D. S. Brock, H. P. A. Mercier and G. J. Schrobilgen, *J. Am. Chem. Soc.*, 2010, **132**, 13823–13839.
- 16 M. Lozinšek, *PhD Thesis*, University of Ljubljana, 2013.
- 17 S. Aitipamula, R. Banerjee, A. K. Bansal, K. Biradha, M. L. Cheney, A. R. Choudhury, G. Desiraju, A. Dikundwar, R. Dubey, N. Duggirala, P. P. Ghogale, S. Ghosh, P. K. Goswami, N. R. Goud, R. K. R. Jetti, P. Karpinski, P. Kaushik, D. Kumar, V. Kumar, B. Moulton, A. Mukherjee, H. Mukherjee, A. S. Myerson, V. Puri, A. Ramanan, T. Rajamannar, C. M. Reddy, N. Rodriguez-Horned, R. D. Rogers, T. N. G. Row, P. Sanphui, N. Shan, A. Singh, C. C. Sun, J. A. Swift, R. Thaimattam, T. S. Thakur, R. K. Thaper, S. P. Thomas, S. Tothadi, V. R. Vangala, P. Vishweshwar, D. R. Weyna and M. J. Zaworotko, *Cryst. Growth Des.*, 2012, **12**, 2147–2152.
- 18 G. J. Schrobilgen and J. M. Whalen, *Inorg. Chem.*, 1994, **33**, 5207–5218.
- 19 B. Kalyanaraman, L. D. Kispert and J. L. Atwood, *Acta Crystallogr., Sect. B*, 1978, **34**, 1131–1136.
- 20 F. R. Fronczek and N. H. Fischer, CCDC 174252: Experimental Crystal Structure Determination, 2002, DOI: [10.5517/cc5vb1c](https://doi.org/10.5517/cc5vb1c).
- 21 C. R. Groom, I. J. Bruno, M. P. Lightfoot and S. C. Ward, *Acta Crystallogr., Sect. B: Struct. Sci., Cryst. Eng. Mater.*, 2016, **72**, 171–179.
- 22 M. C. Etter, J. C. MacDonald and J. Bernstein, *Acta Crystallogr., Sect. B: Struct. Sci.*, 1990, **46**, 256–262.
- 23 M. Lozinšek, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2025, **81**, 188–192.
- 24 R. J. Gillespie and T. Birchall, *Can. J. Chem.*, 1963, **41**, 148–155.
- 25 J. Axhausen, C. Ritter, K. Lux and A. Kornath, *Z. Anorg. Allg. Chem.*, 2013, **639**, 65–72.
- 26 T. Saal, R. Haiges and K. O. Christe, *Dalton Trans.*, 2023, **52**, 18143–18147.
- 27 M. C. Bayer, N. Greither, V. Bockmair, A. Nitzer and A. J. Kornath, *Eur. J. Inorg. Chem.*, 2022, **2022**, e202200501.
- 28 A. Dey, P. Metrangolo, T. Pilati, G. Resnati, G. Terraneo and I. Wlassics, *J. Fluorine Chem.*, 2009, **130**, 816–823.
- 29 Sakshi, Y. Gupta, C. M. Robertson, P. Munshi and A. R. Choudhury, *CrystEngComm*, 2025, **27**, 478–487.
- 30 E. Uran and M. Lozinšek, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2025, **81**, 577–583.
- 31 P. Gilli, L. Pretto, V. Bertolasi and G. Gilli, *Acc. Chem. Res.*, 2009, **42**, 33–44.
- 32 A. I. Gubin, M. Z. Buranbaev and N. N. Nurakhmetov, *Kristallografiya*, 1988, **33**, 506–508.
- 33 H. S. A. Elliot, J. F. Lehmann, H. P. A. Mercier, H. D. B. Jenkins and G. J. Schrobilgen, *Inorg. Chem.*, 2010, **49**, 8504–8523.
- 34 B. Žemva, A. Jesih, D. H. Templeton, A. Zalkin, A. K. Cheetham and N. Bartlett, *J. Am. Chem. Soc.*, 1987, **109**, 7420–7427.
- 35 K. Radan, E. Goresnik and B. Žemva, *Angew. Chem., Int. Ed.*, 2014, **53**, 13715–13719 (*Angew. Chem.*, 2014, **126**, 13935–13939).
- 36 K. Motaln, K. Gurung, P. Brázda, A. Kokalj, K. Radan, M. Dragomir, B. Žemva, L. Palatinus and M. Lozinšek, *ACS Cent. Sci.*, 2024, **10**, 1733–1741.
- 37 M. R. Bortolus, H. P. A. Mercier, B. Nguyen and G. J. Schrobilgen, *Angew. Chem., Int. Ed.*, 2021, **60**, 23678–23686 (*Angew. Chem.*, 2021, **133**, 23871–23879).
- 38 D. S. Brock, J. J. Casalis de Puty, H. P. A. Mercier, G. J. Schrobilgen and B. Silvi, *Inorg. Chem.*, 2010, **49**, 6673–6689.
- 39 M. R. Bortolus, G. J. LaChapelle, H. P. A. Mercier, J. Haner and G. J. Schrobilgen, *Inorg. Chem.*, 2023, **62**, 8761–8771.
- 40 M. J. Hughes, D. S. Brock, H. P. A. Mercier and G. J. Schrobilgen, *J. Fluorine Chem.*, 2011, **132**, 660–668.
- 41 P. A. Agron, G. M. Begun, H. A. Levy, A. A. Mason, C. G. Jones and D. F. Smith, *Science*, 1963, **139**, 842–844.
- 42 C. J. Adams, *J. Raman Spectrosc.*, 1974, **2**, 391–397.
- 43 F. O. Sladky, P. A. Bulliner and N. Bartlett, *J. Chem. Soc. A*, 1969, 2179–2188.
- 44 K. Motaln, K. Gurung, M. Dragomir, D. Kurzydowski, L. Palatinus and M. Lozinšek, *Inorg. Chem.*, 2025, **64**, 14968–14976.
- 45 K. Nakamoto, *Infrared and Raman Spectra of Inorganic and Coordination Compounds: Part A: Theory and Application in Inorganic Chemistry*, John Wiley & Sons, Hoboken, New Jersey, 2009.
- 46 E. K. Murthy and G. R. Rao, *J. Raman Spectrosc.*, 1988, **19**, 359–363.
- 47 M. Lozinšek, H. P. A. Mercier and G. J. Schrobilgen, *Angew. Chem., Int. Ed.*, 2021, **60**, 8149–8156 (*Angew. Chem.*, 2021, **133**, 8230–8237).
- 48 (a) CCDC 2493130: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9k8](https://doi.org/10.5517/ccdc.csd.cc2pp9k8); (b) CCDC 2493131: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9l9](https://doi.org/10.5517/ccdc.csd.cc2pp9l9); (c) CCDC 2493132: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9mb](https://doi.org/10.5517/ccdc.csd.cc2pp9mb); (d) CCDC 2493133: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9nc](https://doi.org/10.5517/ccdc.csd.cc2pp9nc); (e) CCDC 2493134: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9pd](https://doi.org/10.5517/ccdc.csd.cc2pp9pd); (f) CCDC 2493135: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9qf](https://doi.org/10.5517/ccdc.csd.cc2pp9qf); (g) CCDC 2493136: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9rg](https://doi.org/10.5517/ccdc.csd.cc2pp9rg); (h) CCDC 2493137: Experimental Crystal Structure Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9sh](https://doi.org/10.5517/ccdc.csd.cc2pp9sh); (i)



CCDC 2493138: Experimental Crystal Structure  
Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9tj](https://doi.org/10.5517/ccdc.csd.cc2pp9tj); (j)  
CCDC 2493139: Experimental Crystal Structure

Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9vk](https://doi.org/10.5517/ccdc.csd.cc2pp9vk); (k)  
CCDC 2493140: Experimental Crystal Structure  
Determination, 2025, DOI: [10.5517/ccdc.csd.cc2pp9wl](https://doi.org/10.5517/ccdc.csd.cc2pp9wl).

