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The atypical solubility of silver(I) pentafluorobenzoate and its use as a catch-and-release agent†

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Silver(I) pentafluorobenzoate (**1**) possesses an atypical solubility for a silver(I) carboxylate, demonstrating solubility in a range of etheric and aromatic solvents. This versatile solubility has been probed crystallographically by single-crystal X-ray diffraction (SCXRD) due to the myriad of solid-state structures that were isolated from a variety of solvents, providing insights into how **1** can be utilised as a reusable crystallisation aid. This potential as a reusable ‘catch-and-release’ agent to obtain the solid-state structures of liquid natural products was screened with the racemic dihydrocarvone, achiral eucalyptol, and the chiral *R*-carvone, and found to be viable. The results of these screenings demonstrated improved crystallographic outcomes than other prior attempts utilising different methodologies (e.g., encapsulation).

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Introduction

The chemistry of silver is rich, diverse, and stretches back centuries. The flexibility of silver to accommodate a myriad of coordination numbers (commonly ranging from 2–6),¹ its straightforward incorporation into molecules, its ability to cleanly form insoluble halide salts *via* metathesis, and commercial availability have all contributed to its continued utility in the present.

A renewed interest in silver(I) carboxylates has been driven by the recent revival of carbonyl hypoiodite chemistry, for which the analogous silver(I) carboxylates are their synthetic precursors.^{2–7} The intrinsic heteroleptic nature of stabilised carbonyl hypoiodites has facilitated the closest observation of the elusive iodine(I)-pyridine cation to date,³ a pursuit that might be furthered through the use of strongly electron-withdrawing substituents on the carboxylate being incorporated into the hypoiodite moiety. Known since 1967,⁸ silver(I) pentafluorobenzoate has already found utility being generated *in situ* for incorporation into coordination polymers and clusters,^{9,10} and therefore presented itself as a natural progression for future hypoiodite studies.

Silver(I) carboxylates typically have negligible solubility, necessitating strongly polar solvents like (CH₃)₂SO (DMSO) to effect dissolution for subsequent solution-based studies, though

this solubility is concomitant with decomposition of the complex on a timescale of hours to sometimes minutes. Attempts to generate more soluble silver(I) carboxylates have included the silver(I) derivative of pivalic acid, silver(I) pivalate (AgCO₂^tBu),⁵ though even this complex still required strongly polar solvents to dissolve it. The recalcitrant nature of silver(I) carboxylates is aptly explained *via* the solid-state structure of silver(I) pivalate, which revealed a polymeric bilayer structure of Ag⋯O interactions (Fig. 1).⁵ However, the effective insolubility of silver(I) carboxylates does contribute to their straightforward synthesis in H₂O, given that all starting materials, side-products, and potential impurities can be washed out of the products, nor did it hinder their use as intermediates toward the formation of carbonyl hypoiodites.^{3–6}

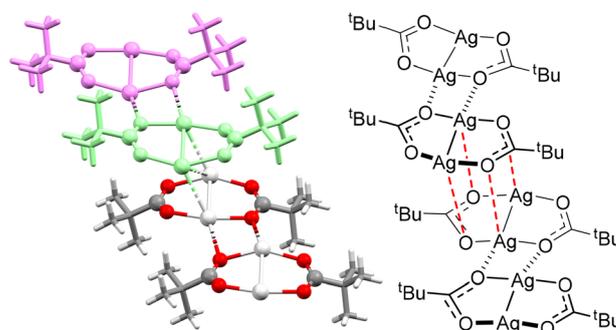
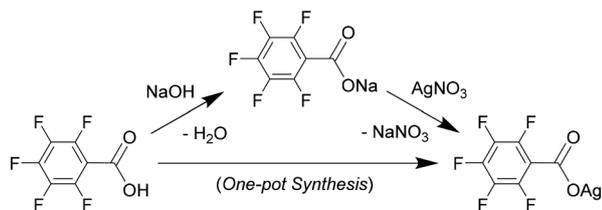


Fig. 1 The solid-state (left; dimeric subunits of the upper layer block coloured for clarity) and structural (right; dashed bonds highlighting the connectivity of dimeric subunits along the layer [black] and between the layers [red]) representations of the polymeric bilayer structure previously observed for (AgCO₂^tBu)_n.⁵

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Scheme 1 The one-pot synthesis of complex **1** from pentafluorobenzoic acid *via* its sodium salt by cation exchange using AgNO₃.

Results and discussion

Synthesis and solubility

Silver(I) pentafluorobenzoate (**1**) was synthesised in effectively quantitative yield from pentafluorobenzoic acid *via* formation of the sodium salt using NaOH, followed by cation exchange of the sodium with silver(I) using AgNO₃ (Scheme 1). However, given the atypically soluble nature of **1**, the work-up diverges from that of other silver(I) carboxylates by Et₂O extraction of the product from the H₂O reaction mixture, yielding the product in 99% yield upon evaporation under reduced pressure of the Et₂O extract. This method can be completed in its entirety on a gram scale in under an hour, and is therefore preferred over prior methods in the literature.^{11–13}

Initially intrigued by the unexpected solubility of **1** in both Et₂O and H₂O,[‡] a range of other ethers were explored. Complex **1** was found to be soluble in Et₂O, ¹Pr₂O, ⁴BuOMe, Ph₂O, Bn₂O, THF, and dioxane, which with the exception of Bn₂O, were all without decomposition. The more polar solvents MeOH, EtOH, ¹PrOH, MeOAc, EtOAc, acetone, MeCN, pyridine, (CH₃)₂NC(O)H (DMF), (CH₃)₂SO were also found to dissolve **1**. However, visually-apparent decomposition in solution of the silver(I) to silver(0) was observed (sometimes within seconds) for MeOH, EtOH, MeCN, (CH₃)₂NC(O)H, and (CH₃)₂SO. Given the well-established precedence of arene adducts of AgClO₄,^{14,15} the aromatic solvents benzene, toluene, *p*-xylene, chlorobenzene, and nitrobenzene were also tested, and all were found to dissolve **1**. It would perhaps be more expedient to list the common solvents in which **1** was found to have negligible solubility: alkanes, CH₂Cl₂, and CHCl₃. Though it might be tempting to think that the atypical solubility of **1** is a property that would be shared by other analogous strongly electron-withdrawing silver(I) carboxylates, the silver(I) complexes of CF₃CO₂, 4-BrC₆H₄CO₂, and 4-NO₂C₆H₄CO₂ only demonstrate partial solubility in strongly polar solvents such as (CH₃)₂SO.³ Whilst the good solubility of **1** is unusual for a silver(I) carboxylate, it is reminiscent of silver(I) trifluoromethanesulfonate (AgOTf), for which there are numerous prior examples of its derivatives serendipitously forming adducts *via* the silver(I) centre with organic solvents (*e.g.*, Et₂O, tetrahydrofuran) in the solid state.^{1,16–19}

[‡] **1** as the monohydrate is known (QOJDAV).⁵⁶

Solution studies

Despite the lack of hydrogen atoms, NMR spectroscopy was initially thought to be an ideal technique to study the effects of potential adduct formation, however, ¹³C NMR studies in multiple different solvents failed to observe the adduct or even the C₆F₅ resonances, suggesting fluxionality on the NMR timescale. Nevertheless, the ¹⁹F NMR resonances were clearly observed, though differences originating from the formation of potentially different adducts was difficult to differentiate from simple solvent effects, with differences of ~1–3 ppm for the various solvents attempted (C₆D₆, d₈-THF, CD₃CN, and (CD₃)₂SO). The solid-state IR of **1** was similarly limited, due to the absence of the strong C=O stretch observed at 1710 cm⁻¹ for pentafluorobenzoic acid, and the unchanged stretch at 1648 cm⁻¹ observed in both the precursor acid and in **1**. It should be noted that the desolvating nature of the adducts of **1**, though minimal for most of the solvents tested herein, hindered some solid-state studies (*e.g.*, IR, TGA) of these combinations.

Solid-state studies

Crystal engineering is an important aspect of silver(I) salts,^{20–23} especially with respect to the formation of silver(I) arene complexes.^{24,25} Investigating new types of crystalline silver(I) materials by incorporating so called super-weak anions such as [B₁₂F₁₂]²⁻, is widely pursued toward ultimately preparing significantly more reactive metal centres than with traditional anions (*e.g.*, BF₄, PF₆, ClO₄, *etc.*).^{24,26} Owing to the widely soluble nature of **1**, a myriad of solid-state structures of solvent adducts of **1** could be prepared by vapour diffusion and determined *via* single-crystal X-ray crystallography (SCXRD; Fig. 2 and Tables S1 and S2[†]). This offered a rare opportunity to study a large subset of solvated solid-state structures of the exact same parent complex, from which direct comparisons could be drawn with respect to the differing coordinating groups of the solvents explored. However, whilst a few of the solid-state structures herein can be considered as solvated structures of **1**, *i.e.*, with solvent molecules trapped in a cavity or coordinated by very weak interactions, the majority of the solid-state structures are adducts of **1**, the advantages of which will be discussed below.

The solid-state structures of **1** could be broadly differentiated into two groups: oxygen-adducts and π -adducts. The ethers Et₂O, ⁴BuOMe, THF, and dioxane were found to be straightforward oxygen-coordinated adducts to the silver(I) metal centre. ¹Pr₂O was also presumably behaving in an analogous fashion as it was able to dissolve **1**, though it could not be confirmed in the solid state. These oxygen-bound adducts featured similar Ag–O bond lengths of 2.476(6) Å (**1**·Et₂O), 2.482(1) Å (**1**·THF), 2.518(1) Å (**1**·⁴BuOMe), and 2.464(2)–2.7747(19) Å (**1**·Dioxane), as expected for their similar electronic and steric environments. The adducts **1**·Et₂O, **1**·THF, and **1**·⁴BuOMe crystallised in 1:1 [**1**]:[adduct] ratios, and retained the



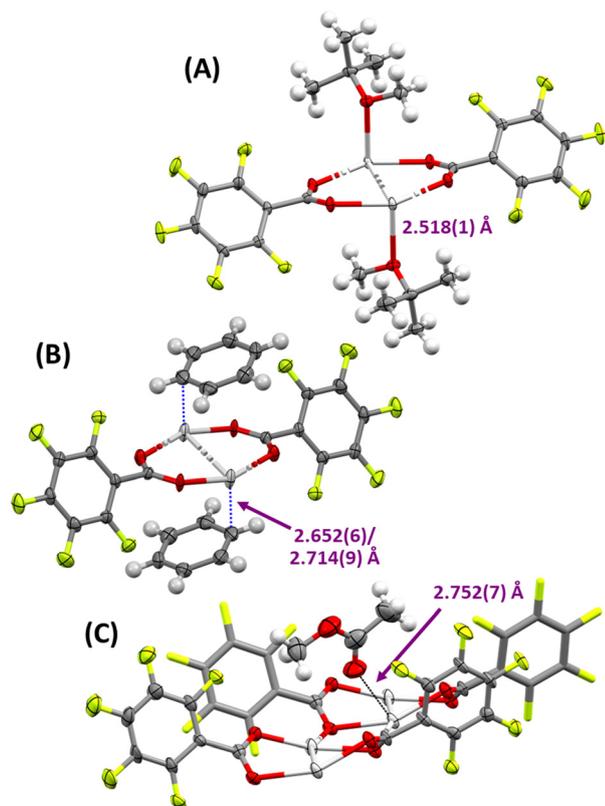


Fig. 2 The solid-state structures of A) **1**-^tBuOMe, B) **1**-Benzene, and C) **1**-MeOAc with the silver(I)-adduct interaction distances annotated, as representative examples of the functional group variability (ethers, arenes/alkenes, ketone/esters, respectively) possible for adduct formation with **1** (thermal ellipsoids at 50% probability).

dimeric subunits of **1** in their packing as 1D chains *via* Ag...O interactions to two neighbouring dimeric subunits. However, whilst the bidentate nature of dioxane yielded the same [1]:[adduct] ratio of 1:1 in **1**-Dioxane (not including an additional 0.125 molecules of solvated dioxane that were also present), it was observed as 2D sheets in their packing, with dioxane adducts bridging across two different dimeric subunits of **1**.

To test if **1** could be used productively to trap previously unknown liquid substrates, the tetrahydropyran derivative isochroman was attempted. This proved successful in an analogous fashion to the other etheric substrates, yielding **1**-Isochroman, which is the first time isochroman has been observed in the solid state (Fig. 3). Interestingly, the solid-state structure contained both an adduct and a solvate molecule of isochroman per molecule of **1**. However, though both crystallographically independent molecules were observed with minor disorder of the flexible heterocyclic ring, the benefits of adduct formation over simple solvation were demonstrated by the more reliable assignment of the oxygen atom in the adduct due to the presence of the Ag–O coordination bond.

The arene solvents all featured close η^1 Ag...C(H) interactions of 2.650(2) Å (**1**-*p*-Xylene), 2.652(6) Å (**1**-Benzene),

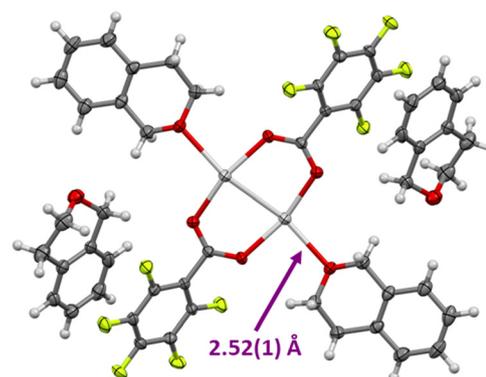


Fig. 3 The solid-state structure of **1**-Isochroman, showing the coordinated and solvated molecules of isochroman (minor disordered positions omitted for clarity; thermal ellipsoids at 50% probability).

2.710(8) Å (**1**-Toluene), and 2.97(2) Å (**1**-Chlorobenzene),§ which were both longer and comparable to the η^2 -silver(I) arene complexes with the OTf anion (*cf.* 2.387(2)/2.698(4) Å for [Ag(benzene)]OTf or 2.452(3)/2.474(3) Å for [Ag(toluene)]OTf,²⁷ and only slightly longer than the Ag...C(H) of 2.563(1)/2.563(1) Å observed for [Ag(benzene)]ClO₄.²⁸ The solid-state packing of **1**-Benzene and **1**-*p*-Xylene were found to be comparable to their etheric counterparts, with 1D chains comprised of the same dimeric subunits of **1** coordinating *via* Ag...O interactions and [1]:[adduct] ratios of 1:1. However, the structure of **1**-Toluene was found adopt a more involved network. The solid-state structure of **1**-Toluene comprised of only a single toluene adduct arranged axially along one side of the 1D chain of dimeric subunits of **1**, and with a lower [1]:[adduct] ratio of 2:1 compared to the 1:1 ratios of **1**-Benzene and **1**-*p*-Xylene that demonstrated alternating axial adducts on each side of the dimeric subunits of **1**. However, unlike benzene and *p*-xylene in **1**-Benzene and **1**-*p*-Xylene, respectively, the toluene molecules in **1**-Toluene bridged across two 1D chains *via* an η^2 Ag...aryl interaction with Ag...C(H) distances of 2.776(8)/2.811(8) Å (Fig. 4), which were only slightly longer than the η^1 Ag...C(H) (*cf.* 2.710(8) Å). However, the cause of this interesting bridging interaction was not readily apparent, given that the less sterically hindered benzene and more electron-rich arene system of *p*-xylene do not demonstrate the same behaviour, and instead mimic the solid-state packing of their etheric analogues (*cf.* **1**-Et₂O, **1**-THF, **1**-^tBuOMe).

The solid-state structure of **1** grown from acetone remained elusive due to rapid desolvation, and that of EtOAc was observed to have heavily disordered channels of overlapping partial-occupancy EtOAc molecules, negating the reliability of its solid-state structure. Nevertheless, the solid-

§ The solid-state structures of **1**-Chlorobenzene featured a symmetry-disordered and partial-occupancy chlorobenzene molecule, which by necessity had to be modelled isotropically. Therefore, the reliability of the value of its Ag...C(H) interaction is diminished in comparison to the other 1-[arene] structures.



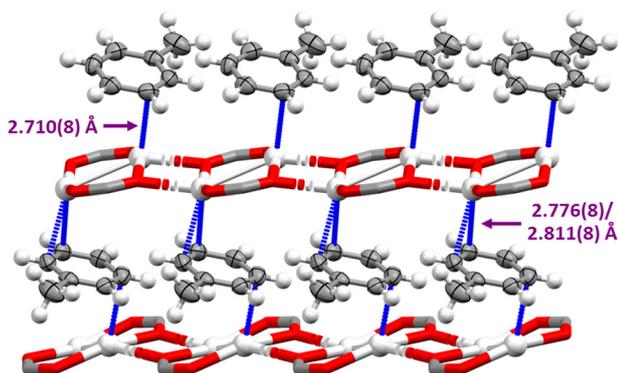


Fig. 4 The bridging toluene molecules between 1D chains of dimeric subunits of **1** via π -interactions (dashed blue lines) in the solid-state packing of eight molecules of **1**·Toluene (C_6F_5 substituents omitted for clarity; thermal ellipsoids at 50% probability).

state structure of **1**·MeOAc, which straddled two adjacent silver(I) dimers via $C=O \cdots Ag$ interactions of 2.752(7) and 3.253(7) Å, definitively confirmed that **1** could also be successfully implemented to immobilise ketones and esters. However, the MeOAc in **1**·MeOAc could be described as a solvate rather than an adduct given the long $Ag \cdots O$ interactions (*cf.* $Ag-O \approx 2.2-2.4$ Å) and minimal MeOAc present per molecule of **1** ($[1]:[solvate] = 4:1$).

The mixed etheric-aromatic solvents Ph_2O and Bn_2O were tested to probe the steric limitations and preferences of **1**, given their duality in potentially forming both oxygen- and π -adducts with **1**. Unfortunately, Bn_2O was found to react with **1** yielding **1**·AgBenzoate, which was not pursued further. Nevertheless, Ph_2O did successfully form an adduct with **1**, which was revealed to be an η^1 -adduct with a $Ag \cdots C(H)$ of 2.684(4) Å, comparable to the η^1 -adducts of **1** observed for benzene, toluene, and *p*-xylene.

Given the option to potentially form both oxygen- and π -adducts, it is surprising that **1**· Ph_2O would assume what is considered the weaker of the two interactions in the solid state. However, this can be attributed to the steric bulk of the phenyl rings preventing an oxygen-adduct from forming within the preferred packing of the 1D chain of the dimeric subunits of **1**. Another factor is the ability of both phenyl rings of Ph_2O to form π -adducts (Fig. 5), even if the second π -adduct is an η^2 $Ag \cdots$ arene interaction with comparably longer $Ag \cdots C(H)$ distances of 3.147(4)/3.255(5) Å compared to the primary η^1 -adduct of the other phenyl ring (2.684(4) Å), and similar to the two sets of π -adducts observed for each molecule of toluene in **1**·Toluene.

A general examination of the packing architectures of the solid-state adducts reveal that they are predominantly comprised of argentophilic dimeric subunits ($Ag \cdots Ag = 2.7370(11)-3.0674(11)$ Å) that pack as 1D chains via pairs of $Ag \cdots O$ interactions to two neighbouring dimeric subunits, with adducts situated axially on the silver(I) atoms of the chain alternating above and below the chain in an overall ratio of 1:1 for $[1]:[adduct]$ (Fig. 6). The exceptions to the aforementioned packing included **1**·Dioxane, which formed

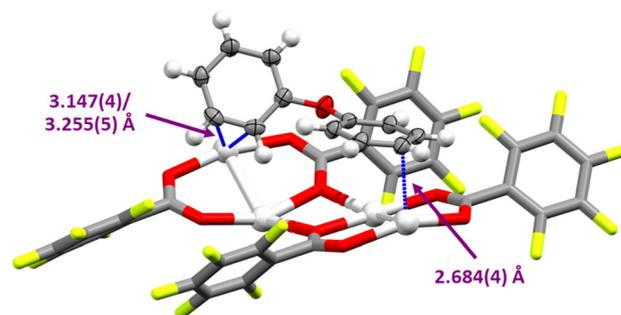


Fig. 5 The η^1 and η^2 π -adducts (dashed blue lines) formed by a single molecule of Ph_2O over two dimeric subunits of **1** in **1**· Ph_2O (second molecule of Ph_2O omitted for clarity; thermal ellipsoids at 50% probability).

adducts via both oxygen atoms to give 2D sheets connected via the same $Ag \cdots O$ interactions in an 8:9 ratio of $[1]:[dioxane]$, though that includes a single molecule of non-coordinating dioxane (*i.e.*, still a 1:1 ratio for adduct formation with **1**). Other exceptions to the 1:1 ratio of $[1]:[adduct]$ were **1**·Toluene (2:1), **1**·Chlorobenzene (1:0.33), **1**· Ph_2O (2:1, though this can be considered a 1:1 ratio per coordinating phenyl ring), **1**·MeOAc (4:1), and **1**·Eucalyptol (3:2; *vide infra*). Given the wide range of adducts observed for **1** (including arene adducts), and despite it possessing a historically strongly-coordinating carboxylate-based ‘anion’, it might be apt to regard **1** as more akin to the super-weak anions currently under research for crystal engineering applications.

Catch-and-release agent

The co-crystallisation of liquid-phase natural products, *e.g.*, essential oils, to obtain their solid-state structures is an established method to circumvent, as Fujita aptly stated, “the intrinsic limitation that the target molecules must be obtained as single crystals”.²⁹ Nevertheless, such methods have historically relied upon the formation of strong intermolecular interactions like hydrogen bonding,^{30,31} which necessitates the presence of hydrogen-bond capable donor

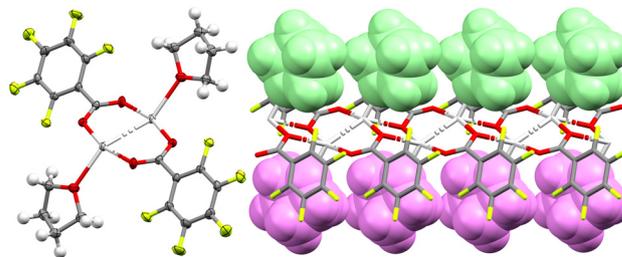


Fig. 6 A dimeric subunit of **1**·THF (left; thermal ellipsoids at 50% probability) and the alternating axial coordination of the THF adducts in a 1D chain of eight molecules of **1**·THF above and below the plane (block coloured green and purple, respectively) of the polymeric chain of dimeric subunits of **1**.



and acceptor functional groups. In the absence of such functional groups, and their strong intermolecular forces directing the solid-state packing of these co-crystallisations, methodologies have instead shifted to trapping the liquid substrates within cavities or inside porous networks.^{29,32–36} These methodologies are impressive in the scope of their substrates successfully co-crystallised, especially for substrates that lack any readily-exploited functional groups (e.g., alkanes).³³ However, the required porosity of such networks can result in large unit cells comprised of primarily light-atom molecules ($Z < \text{Si}$) can act as double-edged swords, given that such systems can necessitate the use of powerful synchrotron radiation sources.³⁷ This in turn results in other common limitations of macro-molecular crystallography, e.g., high R -factors ($R[F^2 > 2\sigma(F^2)] > 5\%$) relative to the crystallography of small discrete molecules. It is in this respect that **1** may offer advantages over other prior co-crystallisation methodologies.

The adduct formation of **1** with ethers, arenes, and ketones/esters demonstrates the same ability to direct the orderly packing of the adduct co-crystals with **1** as observed for hydrogen-bond directed co-crystals, but without the presence of a strong intermolecular interaction such as hydrogen bonding. As a result of this weak adduct formation, a surprising outcome of co-crystallisations of **1** was the ability to recover it after adduct formation (for solvents where decomposition had not been observed) through simple evaporation. Similar recovery and reuse had been previously reported for the closely-related AgOTf salt when it was utilised as a catalyst.³⁸ Isolation of the liquid substrates could also be envisioned through the same evaporation process, if desired, though was not attempted herein. Subsequent reuse of the recovered **1** with other solvents could then generate new adducts. A test sample was successfully reused five times to form various etheric or aromatic adducts, as confirmed from the crystallographic unit cells by SCXRD, and most of the X-ray datasets reported herein were themselves grown from recovered samples of **1**. It was envisioned that this ability would be an attractive feature for organic chemists to crystallise liquid-phase natural products, without the need for low-temperature crystallisation or advanced *in situ* crystallisation techniques, followed by sample recovery *via* evaporation. This ‘catch-and-release’³⁹ agent would minimise the amount of substrate consumed to attempt crystallisation, which would be ideal for natural products where the amount of sample is limited (e.g., if they are difficult to synthesise or isolate). In this way, **1** can be favourably likened to Fujita’s crystalline sponges,^{29,35,36} with **1** being easier to produce, and the resulting crystallographic modelling being generally more straightforward. Additional benefits would include the presence of an atom of greater atomic number than Si (*i.e.*, Ag), which would permit the absolute stereochemistry to be assigned for exclusively light-atom molecules even using Mo radiation X-ray diffractometers, which may prove challenging with the crystalline sponge methodology.^{40,41}

To test this potential of **1**, four additional liquid substrates were screened: the cyclic ethers 2-methyltetrahydrofuran

(2-MeTHF; racemic) and eucalyptol, as well as the ketone- and alkene-containing dihydrocarvone (racemic) and *R*-carvone. All substrates were able to successfully dissolve a sample of **1**, with crystals successfully obtained for all of them (**1**·2-MeTHF, **1**·Eucalyptol, **1**·Dihydrocarvone, and **1**·*R*-Carvone) *via* pentane vapour diffusion. The 2-MeTHF adduct in **1**·2-MeTHF was coordinated *via* the oxygen atom (Ag–O = 2.40(2)/2.42(1) Å) and did not display any narcissistic self-sorting of the *R* and *S* enantiomers, with both crystallising together in the solid-state structure. Similarly, dihydrocarvone also crystallised as the racemate in a 1 : 1 ratio with **1** (**1**·Dihydrocarvone), coordinating *via* the oxygen atom of its carbonyl group with a Ag–O bond length of 2.490(2) Å. No solid-state structure of dihydrocarvone previously existed in the literature, highlighting the potential of **1** to elucidate new structures. Whilst **1** had some solubility in neat dihydrocarvone, screening tests revealed that the use of a 1 : 1 dihydrocarvone : acetone mixture proved best to yield single crystals, with acetone being seen as an ideal co-solvent given its excellent solubility toward **1**, coupled with its aforementioned highly-labile nature as an adduct/solvate of **1**. This further demonstrates the potential applicability of **1** with substrates in which it has limited solubility through the use of co-solvents like acetone that form highly-labile adducts/solvates and are therefore easily out-competed by other substrates. Additionally, the use of a mixture of solvents can help reduce the amount of substrate used, should they be expensive or difficult to synthesise. The achiral **1**·Eucalyptol was also observed as an oxygen adduct (Ag–O = 2.367(3)/2.409(3) Å). The solid-state structure of eucalyptol was previously obtained *via in situ* crystallisation and co-crystallisation with 1,4-dihydroxybenzene.^{1,42,43} However, the crystals of **1**·Eucalyptol could be prepared overnight without the need for low-temperature crystallisation, and the stronger Ag–O adduct formation provided crystals that did not suffer from the rapid desolvation reported for the 1,4-dihydroxybenzene co-crystal,⁴³ which is beneficial for subsequent handling during SCXRD studies. Lastly, the enantiopure configuration of *R*-carvone in **1**·*R*-Carvone was successfully determined crystallographically *via* refinement of the Flack parameter in the non-centrosymmetric C_2 space group. The *R*-carvone was observed to coordinate *via* both its oxygen and terminal alkene,[¶] bridging across two 1D chains of **1** (Fig. 7), reminiscent of the previously discussed coordination of **1**·Dioxane.

The solid-state packing of these natural-product co-crystals of **1**, as observed for the other simpler adducts described previously, retained their dimeric subunits of **1** upon adduct formation, with Ag⋯Ag distances of 2.8598(4)/2.8835(4) Å for **1**·Eucalyptol, 2.8451(4) Å for **1**·Dihydrocarvone, and 2.795(1)/2.982(1) Å for **1**·*R*-Carvone. The structures were also analogously observed as 1D chains in their packing *via* Ag⋯O interactions between neighbouring dimeric subunits of **1**, with Ag⋯O distances of 2.461(3)–2.541(2) Å (**1**·Eucalyptol), 2.496(2) Å (**1**·Dihydrocarvone), and 2.384(3)–2.522(3) Å (**1**·*R*-

¶ There was disorder in the position of the silver(i) atoms in **1**·*R*-Carvone, which created ambiguity between the distances of the Ag⋯O and Ag⋯C(H₂ = C) interactions of the *R*-carvone with **1**.



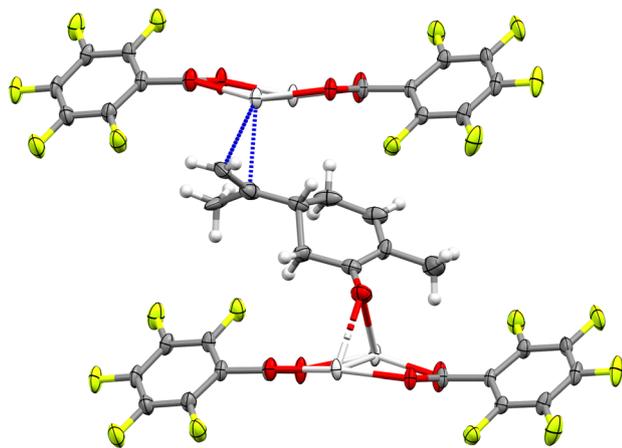


Fig. 7 The solid-state structure of **1-R-Carvone** illustrating the *R*-carvone molecule bridging across two 1D chains of **1** via both its terminal alkene (dashed blue lines) and oxygen atom (minor disordered position omitted for clarity; thermal ellipsoids at 50% probability).

Carvone). Finally, it should be noted that the natural-product crystals of eucalyptol or *R*-carvone with **1**, despite crystallographic disorder being present in both, gave excellent SCXRD structures with R_1/wR_2 values of 3.1/8.0% and 2.2/5.3%, respectively, which have lower R_1 values for both substrates in comparison to all prior literature structures.¹ The structure of dihydrocarvone (**1-Dihydrocarvone**) was without any apparent disorder present and was also of excellent quality ($R_1/wR_2 = 2.7/6.0\%$), though there were no literature examples for comparison. The **1-Eucalyptol** structure (3.1%) is comparable to the R_1 values of the several hydrogen-bonded co-crystals of eucalyptol (ranging from 4.6 to 8.3%)^{43,44} and the low-temperature isolated structure (4.7%),⁴² and a notable improvement over encapsulation within a porous organic cage (11.3%).⁴⁵ Similarly, **1-R-Carvone** (2.2%) is comparable to the R_1 values of hydrogen-bonded (4.2%)⁴⁶ or organic co-crystals (4.3 and 5.1%)³³ of *R*-carvone, as well as the low-temperature isolated structure (3.0%),⁴⁷ whilst again being an improvement over co-crystallisation within porous organic cages (ranging from 7.1 to 15.3%).^{45,48,49}

Conclusions

In conclusion, the easy to synthesise complex silver(i) pentafluorobenzoate (**1**) was found to be soluble in a myriad of organic solvents, which was atypical when compared to other silver(i) carboxylates, even those incorporating similarly strong electron-withdrawing substituents on the carboxylates. Solid-state adducts of **1** could be straightforwardly crystallised for liquid substrates coordinated *via* a range of functional groups including ethers, ketones, esters, alkenes, and arenes. The ease and low cost of synthesising **1** in bulk also makes it ideal for the inexpensive, large-scale screening of substrates (including determination of their absolute stereochemistry), such as liquid-phase natural products that

are of interest to researchers. The possibility of using substrate:acetone solvent mixtures for crystallisation also present an economic pathway should the natural products be expensive or difficult to synthesise. After SCXRD studies, **1** could often be recovered from these substrates by evaporation and subsequently reused; a favourable trait toward the pursuit of greener chemistry. The adducts of **1** with isochroman and dihydrocarvone also served to highlight the potential of **1** to yield previously unreported solid-state structures of liquid substrates. The solid-state structures of the natural-product adducts of **1** tested herein demonstrated excellent crystallographic metrics, which were found to have improved outcomes compared to those from other methodologies like encapsulation. Just as mechanochemical techniques minimise environmental impact by circumventing the use of toxic solvents like CH_2Cl_2 ,^{50–52} the recovery and reuse of crystallisation materials, as demonstrated with **1** herein, could offer an environmentally-friendly alternative when attempting to prepare suitable crystals for SCXRD.

Experimental

General considerations

All reagents and solvents were obtained from commercial suppliers and used without further purification. For structural NMR assignments, ^{19}F NMR spectra were recorded on a Bruker Avance 300 MHz spectrometer to an external KF standard, with chemical shifts reported on the δ scale in ppm to a CFCl_3 standard calculated from the following conversion: [chemical shift referenced to CFCl_3] = $(-125.3 \text{ ppm} + [\text{chemical shift referenced to KF}])$. Multiplicities are denoted as: s (singlet), d (doublet), t (triplet), q (quartet), m (multiplet), and br (broad). Infrared (IR) spectra were recorded on a Bruker Tensor 27 FT-IR spectrometer.

The single crystal X-ray diffraction (SCXRD) data were collected at 120 K using mirror-monochromated $\text{Cu-K}\alpha$ ($\lambda = 1.54184 \text{ \AA}$) radiation on a Rigaku XtaLAB Synergy-R diffractometer with a HyPix-Arc 100 detector. All structures were solved by intrinsic phasing (SHELXT)⁵³ and refined by full-matrix least squares on F^2 using Olex2,⁵⁴ utilising the SHELXL module.⁵⁵ Anisotropic displacement parameters were assigned to non-H atoms and isotropic displacement parameters for all H atoms were constrained to multiples of the equivalent displacement parameters of their parent atoms with $U_{\text{iso}}(\text{H}) = 1.2 U_{\text{eq}}(\text{CH})$ or $1.5 U_{\text{eq}}(\text{CH}_2, \text{CH}_3, \text{OH})$ of their respective parent atoms. The X-ray single crystal data and CCDC numbers (2310151–2310162, 2311134) of all new structures are included below, excluding the connectivity model for **1-Chlorobenzene**.

Synthesis and characterisation

AgCO₂C₆F₅ (1). The synthesis of **1** was performed as previously described for analogous silver(i) carboxylates,^{3,5} but with a different work-up due to its atypical solubility. A solution of NaOH (0.200 g, 5 mmol) in H_2O (5 mL) was added to a white suspension of pentafluorobenzoic acid (1.061 g, 5



mmol) in H₂O (10 mL) to give a colourless solution. A solution of AgNO₃ (0.850 g, 5 mmol) in H₂O (10 mL) was added to the reaction mixture to give some white precipitate. Stirred for 15 minutes, then the product was extracted with Et₂O (4 × 50 mL). All volatiles were removed under reduced pressure from the Et₂O extract to give the product as a white solid. Yield = 1.575 g (4.94 mmol, 99%).

1 (in (CD₃)₂SO). ¹⁹F NMR (282 MHz, (CD₃)₂SO) δ -202.23 (dd, *J* = 25.7, 8.1 Hz), -218.13 (t, *J* = 21.6 Hz), -221.44 (td, *J* = 25.6, 8.4 Hz). N.B. Even though the sample was collected within 5 minutes of dissolution in (CD₃)₂SO, two minor additional sets of ¹⁹F NMR resonances were observed due to decomposition of **1**.

1 (in CD₃CN). ¹⁹F NMR (282 MHz, CD₃CN) δ -203.54 (dd, *J* = 23.6, 8.4 Hz), -219.89 (t, *J* = 19.8 Hz), -223.37 (td, *J* = 23.5, 8.5 Hz).

1 (in d₈-THF). ¹⁹F NMR (282 MHz, d₈-THF) δ -201.39 (dd, *J* = 23.0, 8.2 Hz), -217.62 (t, *J* = 20.3 Hz), -223.33 (td, *J* = 23.1, 8.4 Hz).

1 (in C₆D₆). ¹⁹F NMR (282 MHz, C₆D₆) δ -200.53 (dd, *J* = 23.6, 7.8 Hz), -214.33 (t, *J* = 21.2 Hz), -221.02 (td, *J* = 22.7, 7.2 Hz).

SCXRD

Adduct synthesis. The synthesis of the crystalline adducts were all performed in the same general manner (with the exception of **1-Et₂O** and **1-Dihydrocarvone**; *vide infra*): ~5 mg of **1** would be dissolved into the desired adduct (1 mL), then set-up as a vapour diffusion crystallisation with pentane (4 mL) as the anti-solvent for 1–3 days at ambient temperature.

(AgCO₂C₆F₅-C₄H₁₀O)_n (1-Et₂O). Crystals suitable for single crystal X-ray diffraction were obtained by partial evaporation of a Et₂O solution of **1** at ambient temperature (N.B. crystals of **1-Et₂O** rapidly desolvate (<30 seconds), and therefore complete evaporation of the sample will not yield crystalline samples). Crystal data for **1-Et₂O**: CCDC-2310151, C₁₁H₁₀AgF₅O₃, *M* = 393.06, colourless plate, 0.01 × 0.06 × 0.15 mm, triclinic, space group *P* $\bar{1}$ (no. 2), *a* = 5.4941(2) Å, *b* = 9.8998(3) Å, *c* = 12.7267(5) Å, α = 82.824(3)°, β = 80.970(3)°, γ = 75.189(3)°, *V* = 658.31(4) Å³, *Z* = 2, *D*_{calc} = 1.983 g cm⁻³, *F*(000) = 384, μ = 12.94 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 2554 independent reflections, 2411 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.071, 2554 data, 216 parameters, 89 restraints, GooF = 1.07, 1.50 > Δρ > -0.82 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.045, *wR*(*F*²) = 0.131.

(AgCO₂C₆F₅-C₅H₁₂O)_n (1-^tBuOMe). Crystals suitable for single crystal X-ray diffraction were obtained from a ^tBuOMe solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-^tBuOMe**: CCDC-2310152, C₁₂H₁₂AgF₅O₃, *M* = 407.09, colourless plate, 0.02 × 0.06 × 0.07 mm, triclinic, space group *P* $\bar{1}$ (no. 2), *a* = 5.59808(5) Å, *b* = 10.23969(10) Å, *c* = 12.79989(10) Å, α = 97.6393(7)°, β = 96.3916(7)°, γ = 102.3278(7)°, *V* = 703.01(1) Å³, *Z* = 2, *D*_{calc} = 1.923 g cm⁻³, *F*(000) = 400, μ = 12.15 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 2849 independent reflections, 2791 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.027, 2849 data, 194 parameters, no restraints, GooF = 1.06, 0.39 > Δρ > -0.37 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.014, *wR*(*F*²) = 0.033.

(AgCO₂C₆F₅-C₄H₈O)_n (1-THF). Crystals suitable for single crystal X-ray diffraction were obtained from a THF solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-THF**: CCDC-2310153, C₁₁H₈AgF₅O₃, *M* = 391.04, colourless block, 0.05 × 0.07 × 0.12 mm, triclinic, space group *P* $\bar{1}$ (no. 2), *a* = 5.5278(1) Å, *b* = 8.6031(1) Å, *c* = 8.6031(1) Å, α = 103.225(1)°, β = 94.823(2)°, γ = 100.843(2)°, *V* = 593.50(2) Å³, *Z* = 2, *D*_{calc} = 2.188 g cm⁻³, *F*(000) = 380, μ = 14.35 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 2418 independent reflections, 2357 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.028, 2418 data, 181 parameters, no restraints, GooF = 1.12, 0.39 > Δρ > -0.46 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.016, *wR*(*F*²) = 0.042.

(AgCO₂C₆F₅-C₉H₁₀O)₂·2(C₉H₁₀O) (1-Isochroman). Crystals suitable for single crystal X-ray diffraction were obtained from an isochroman solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Isochroman**: CCDC-2334781, C₃₂H₂₀Ag₂F₁₀O₆·2(C₉H₁₀O), *M* = 1174.56, colourless plate, 0.03 × 0.03 × 0.11 mm, triclinic, space group *P* $\bar{1}$ (no. 2), *a* = 7.2534(3) Å, *b* = 11.0121(5) Å, *c* = 13.8225(6) Å, α = 85.841(4)°, β = 85.713(3)°, γ = 84.263(3)°, *V* = 1093.11(8) Å³, *Z* = 1, *D*_{calc} = 1.784 g cm⁻³, *F*(000) = 588, μ = 8.08 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 4439 independent reflections, 3829 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.052, 4439 data, 363 parameters, 38 restraints, GooF = 1.05, 1.53 > Δρ > -1.43 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.036, *wR*(*F*²) = 0.079.

(AgCO₂C₆F₅-C₄H₈O)_n·0.125(C₄H₈O)₂ (1-Dioxane). Crystals suitable for single crystal X-ray diffraction were obtained from a dioxane solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Dioxane**: CCDC-2310154, 2(C₂₂H₁₆Ag₂F₁₀O₈)·0.5(C₄H₈O)₂, *M* = 1672.23, colourless plate, 0.03 × 0.09 × 0.16 mm, monoclinic, space group *P*2₁/*n*, *a* = 14.4142(1) Å, *b* = 23.8477(1) Å, *c* = 15.1372(1) Å, β = 93.326(1)°, *V* = 5194.58(5) Å³, *Z* = 4, *D*_{calc} = 2.138 g cm⁻³, *F*(000) = 3264, μ = 13.25 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 10 607 independent reflections, 9585 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.025, 10 607 data, 815 parameters, 36 restraints, GooF = 1.08, 0.39 > Δρ > -0.64 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.020, *wR*(*F*²) = 0.054.

(AgCO₂C₆F₅-C₆H₆)_n (1-Benzene). Crystals suitable for single crystal X-ray diffraction were obtained from a benzene solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Benzene**: CCDC-2310155, C₂₆H₁₂Ag₂F₁₀O₄, *M* = 794.10, colourless needle, 0.01 × 0.01 × 0.13 mm, triclinic, space group *P* $\bar{1}$ (no. 2), *a* = 5.69188(13) Å, *b* = 12.5817(3) Å, *c* = 17.9282(6) Å, α = 105.107(3)°, β = 90.214(2)°, γ = 90.163(2)°, *V* = 1239.51(6) Å³, *Z* = 2, *D*_{calc} = 2.128 g cm⁻³, *F*(000) = 768, μ = 13.71 mm⁻¹, *T* = 120.0(1) K, θ_{\max} = 74.5°, 5044 independent reflections, 4271 with *I*_o > 2σ(*I*_o), *R*_{int} = 0.040, 5044 data, 379 parameters, no restraints, GooF = 1.15, 1.91 > Δρ > -1.39 e Å⁻³, *R*[*F*² > 2σ(*F*²)] = 0.044, *wR*(*F*²) = 0.109.

2(AgCO₂C₆F₅-C₇H₈)_n (1-Toluene). Crystals suitable for single crystal X-ray diffraction were obtained from a toluene solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Toluene**: CCDC-2310156, C₄₂H₁₆Ag₄F₂₀O₈, *M* = 1460.03, colourless needle, 0.01 × 0.03 × 0.12 mm, monoclinic, space group *P*2₁/*n*, *a* = 5.6097(1) Å, *b* = 55.4509(15) Å, *c* = 13.6451(4) Å, β = 97.194(2)°, *V* = 4211.07(18)



\AA^3 , $Z = 4$, $D_{\text{calc}} = 2.303 \text{ g cm}^{-3}$, $F(000) = 2800$, $\mu = 16.05 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 8525 independent reflections, 6833 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.063$, 8525 data, 669 parameters, no restraints, $\text{GooF} = 1.05$, $0.85 > \Delta\rho > -1.00 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.053$, $wR(F^2) = 0.133$.

(AgCO₂C₆F₅-C₈H₁₀O)_n (1-*p*-Xylene). Crystals suitable for single crystal X-ray diffraction were obtained from a *p*-xylene solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-*p*-Xylene**: CCDC-2310157, C₁₅H₁₀AgF₅O₂, $M = 425.10$, colourless needle, $0.02 \times 0.03 \times 0.31 \text{ mm}$, monoclinic, space group $P2_1/n$, $a = 5.6950(1) \text{ \AA}$, $b = 17.0008(3) \text{ \AA}$, $c = 14.7939(3) \text{ \AA}$, $\beta = 97.036(2)^\circ$, $V = 1421.55(5) \text{ \AA}^3$, $Z = 4$, $D_{\text{calc}} = 1.986 \text{ g cm}^{-3}$, $F(000) = 832$, $\mu = 12.00 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 2902 independent reflections, 2660 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.043$, 2902 data, 210 parameters, no restraints, $\text{GooF} = 1.05$, $0.46 > \Delta\rho > -0.69 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.023$, $wR(F^2) = 0.059$.

(4(AgCO₂C₆F₅)-C₃H₆O₂)_n (1-MeOAc). Crystals suitable for single crystal X-ray diffraction were obtained from a MeOAc solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-MeOAc**: CCDC-2310158, C₃₁H₆Ag₄F₂₀O₁₀, $M = 1349.84$, colourless needle, $0.01 \times 0.01 \times 0.10 \text{ mm}$, triclinic, space group $P\bar{1}$ (no. 2), $a = 5.5552(2) \text{ \AA}$, $b = 16.0610(7) \text{ \AA}$, $c = 20.9338(6) \text{ \AA}$, $\alpha = 74.645(3)^\circ$, $\beta = 89.103(3)^\circ$, $\gamma = 80.350(3)^\circ$, $V = 1774.79(12) \text{ \AA}^3$, $Z = 2$, $D_{\text{calc}} = 2.526 \text{ g cm}^{-3}$, $F(000) = 1280$, $\mu = 19.00 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 7191 independent reflections, 4740 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.090$, 7191 data, 588 parameters, no restraints, $\text{GooF} = 1.01$, $0.80 > \Delta\rho > -1.07 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.047$, $wR(F^2) = 0.092$.

(2(AgCO₂C₆F₅)-C₁₂H₁₀O)_n (1-Ph₂O). Crystals suitable for single crystal X-ray diffraction were obtained from a Ph₂O solution of **1** vapour diffused with pentane maintained at 303 K (Ph₂O melting point = 298–300 K). Crystal data for **1-Ph₂O**: CCDC-2310159, C₂₆H₁₀Ag₂F₁₀O₅, $M = 808.08$, colourless needle, $0.01 \times 0.03 \times 0.06 \text{ mm}$, monoclinic, space group $P2_1/n$, $a = 14.6080(4) \text{ \AA}$, $b = 9.2325(2) \text{ \AA}$, $c = 18.5937(4) \text{ \AA}$, $\beta = 103.760(2)^\circ$, $V = 2435.73(10) \text{ \AA}^3$, $Z = 4$, $D_{\text{calc}} = 2.204 \text{ g cm}^{-3}$, $F(000) = 1560$, $\mu = 14.00 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 4976 independent reflections, 4086 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.045$, 4976 data, 388 parameters, no restraints, $\text{GooF} = 1.02$, $1.60 < \Delta\rho < -1.07 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.033$, $wR(F^2) = 0.079$.

(AgCO₂C₆F₅-AgCO₂C₆H₅)_n (1-AgBenzoate). The use of Bn₂O as a crystallisation solvent for **1** resulted in the isolation of crystals of a silver(i) dimer consisting of a 1:1 mixture of benzoate:pentafluorobenzoate. Crystals suitable for single crystal X-ray diffraction were obtained from a Bn₂O solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-AgBenzoate**: CCDC-2310160, C₁₄H₅Ag₂F₅O₄, $M = 547.92$, colourless needle, $0.01 \times 0.01 \times 0.17 \text{ mm}$, monoclinic, space group $P2_1/n$, $a = 6.9141(3) \text{ \AA}$, $b = 5.5610(2) \text{ \AA}$, $c = 37.1068(11) \text{ \AA}$, $\beta = 93.837(3)^\circ$, $V = 1423.53(9) \text{ \AA}^3$, $Z = 4$, $D_{\text{calc}} = 2.557 \text{ g cm}^{-3}$, $F(000) = 1040$, $\mu = 22.86 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 2894 independent reflections, 2336 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.051$, 2894 data, 226 parameters, no restraints, $\text{GooF} = 1.04$, $1.26 > \Delta\rho > -1.14 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.047$, $wR(F^2) = 0.120$.

(AgCO₂C₆F₅-C₅H₁₀O)_n (1-2-MeTHF). Crystals suitable for single crystal X-ray diffraction were obtained from a 2-MeTHF (racemic) solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-2-MeTHF**: CCDC-2310161, C₁₂H₁₀AgF₅O₃, $M = 405.07$, colourless needle, $0.02 \times 0.02 \times 0.16 \text{ mm}$, triclinic, space group $P\bar{1}$ (no. 2), $a = 5.51822(13) \text{ \AA}$, $b = 9.8945(5) \text{ \AA}$, $c = 13.0489(3) \text{ \AA}$, $\alpha = 75.275(3)^\circ$, $\beta = 85.880(2)^\circ$, $\gamma = 79.845(3)^\circ$, $V = 678.00(4) \text{ \AA}^3$, $Z = 2$, $D_{\text{calc}} = 1.984 \text{ g cm}^{-3}$, $F(000) = 396$, $\mu = 12.59 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 2742 independent reflections, 2470 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.045$, 2742 data, 241 parameters, 132 restraints, $\text{GooF} = 1.09$, $0.98 > \Delta\rho > -0.70 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.037$, $wR(F^2) = 0.106$.

(AgCO₂C₆F₅-C₁₀H₁₆O)_n (1-Dihydrocarvone). Crystals suitable for single crystal X-ray diffraction were obtained from a dihydrocarvone:acetone (1:1) solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Dihydrocarvone**: CCDC-2352000, C₁₇H₁₆AgF₅O₃, $M = 471.17$, colourless plate, $0.01 \times 0.04 \times 0.22 \text{ mm}$, monoclinic, space group $P2_1/n$, $a = 13.9355(2) \text{ \AA}$, $b = 5.5125(1) \text{ \AA}$, $c = 22.5133(2) \text{ \AA}$, $\beta = 93.604(1)^\circ$, $V = 1726.04(4) \text{ \AA}^3$, $Z = 4$, $D_{\text{calc}} = 1.813 \text{ g cm}^{-3}$, $F(000) = 936$, $\mu = 10.00 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 3513 independent reflections, 3104 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.045$, 3513 data, 237 parameters, no restraints, $\text{GooF} = 1.05$, $0.55 > \Delta\rho > -0.53 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.027$, $wR(F^2) = 0.060$.

(3(AgCO₂C₆F₅)-2(C₁₀H₁₈O))_n (1-Eucalyptol). Crystals suitable for single crystal X-ray diffraction were obtained from a eucalyptol solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-Eucalyptol**: CCDC-2310162, C₄₁H₃₆Ag₃F₁₅O₈, $M = 1265.31$, colourless block, $0.06 \times 0.16 \times 0.18 \text{ mm}$, triclinic, space group $P\bar{1}$ (no. 2), $a = 12.4062(3) \text{ \AA}$, $b = 13.45738(19) \text{ \AA}$, $c = 13.9666(3) \text{ \AA}$, $\alpha = 85.3721(16)^\circ$, $\beta = 70.664(2)^\circ$, $\gamma = 86.9586(16)^\circ$, $V = 2192.24(8) \text{ \AA}^3$, $Z = 2$, $D_{\text{calc}} = 1.917 \text{ g cm}^{-3}$, $F(000) = 1244$, $\mu = 11.70 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 8953 independent reflections, 8507 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.029$, 8953 data, 925 parameters, 705 restraints, $\text{GooF} = 1.03$, $2.54 > \Delta\rho > -0.70 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.031$, $wR(F^2) = 0.080$.

(2(AgCO₂C₆F₅)-C₁₀H₁₄O)_n (1-*R*-Carvone). Crystals suitable for single crystal X-ray diffraction were obtained from an *R*-carvone solution of **1** vapour diffused with pentane at ambient temperature. Crystal data for **1-*R*-Carvone**: CCDC-2311134, C₂₄H₁₄Ag₂F₁₀O₅, $M = 788.09$, colourless plate, $0.03 \times 0.05 \times 0.22 \text{ mm}$, monoclinic, space group $C2$, $a = 17.79831(19) \text{ \AA}$, $b = 5.58069(5) \text{ \AA}$, $c = 25.9618(2) \text{ \AA}$, $\beta = 104.5758(10)^\circ$, $V = 2495.71(4) \text{ \AA}^3$, $Z = 4$, $D_{\text{calc}} = 2.097 \text{ g cm}^{-3}$, $F(000) = 1528$, $\mu = 13.63 \text{ mm}^{-1}$, $T = 120.0(1) \text{ K}$, $\theta_{\text{max}} = 74.5^\circ$, 5084 independent reflections, 4748 with $I_o > 2\sigma(I_o)$, $R_{\text{int}} = 0.051$, 5084 data, 511 parameters, 43 restraints, $\text{GooF} = 1.02$, $0.28 > \Delta\rho > -0.29 \text{ e \AA}^{-3}$, $R[F^2 > 2\sigma(F^2)] = 0.022$, $wR(F^2) = 0.053$.

Conflicts of interest

There are no conflicts to declare.



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