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Repurposing of the anti-HIV drug emtricitabine as a hydrogen-bonded cleft for bipyridines *via* cocrystallization†

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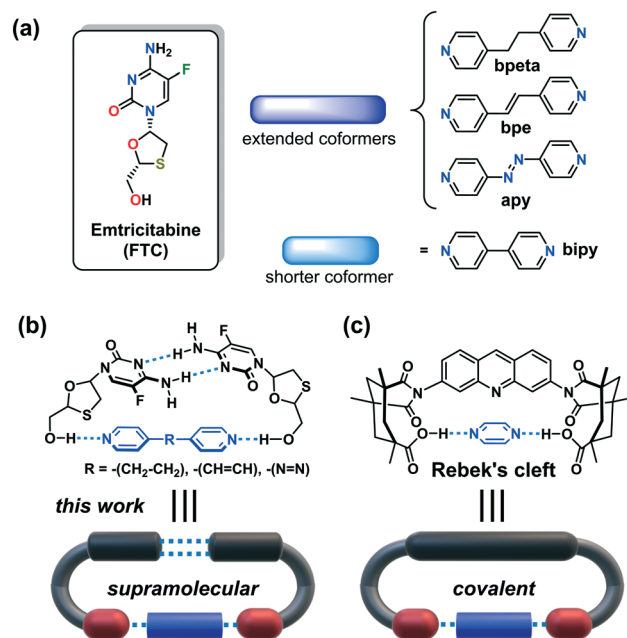
We report supramolecular repurposing of emtricitabine (FTC, trade name: Emtriva®), a blockbuster FDA-approved anti-HIV agent. FTC is revealed to act as a hydrogen-bonded cleft for bipyridine recognition. The supramolecular repurposing is realized by the generation of four cocrystals through liquid-assisted grinding. The clefts comprise discrete three-component assemblies sustained by a combination of hydrogen bonds and $\pi \cdots \pi$ interactions.

While the repurposing of pharmaceutical ingredients (APIs) is an important business model in drug development, the act of repurposing molecules for other applications (*e.g.* materials science) remains largely unexplored.¹ Generally, the tactic consists in finding new uses of existing molecules applied to molecular and/or supramolecular chemistry. Applications in the supramolecular domain are extremely rare and constitute sustainable research with promising applications. Diao, for example, has just applied supramolecular repurposing of DNA-binding agents to develop hydrogen-bonded organic semiconductors.²

Here, we describe supramolecular repurposing of the FDA-approved anti-HIV agent emtricitabine (FTC, trade name: Emtriva®). Specifically, we demonstrate the ability of FTC to function as an artificial supramolecular cleft-like receptor of bipyridines (Scheme 1a and b). The cleft is stabilized by complementary hydrogen bonding and through synthon competition. FTC is a major anti-HIV drug approved by the FDA included in the World Health Organization (WHO)'s list of essential medicines.³

Seminal work by Rebek described the ability of single molecules to function as molecular clefts that recognize

linear rod-shaped guests (Scheme 1c).⁴ Recognition is achieved owing to convergent hydrogen-bond donor sites. FTC self-assembles as a homodimer in the solid state, with axial hydroxyl groups engaged in N–H $\cdots$ N hydrogen bonds in a *transoid* conformation (Fig. S1, ESI†).⁵ Cocrystals of FTC are unknown, thus, as part of our program of study of multi-component solids,⁶ we asked whether FTC would form cocrystals with the rod-shaped bipyridines 1,2-bis(4-pyridyl)ethane (**bpeta**), 1,2-bis(4-pyridyl)ethylene (**bpe**), 4,4'-azopyridine (**apy**), and 4,4'-bipyridine (**bipy**). During our research, we discovered N–H $\cdots$ N hydrogen bonding of FTC to enable FTC to function as a self-assembled (*i.e.* supramolecular) cleft that recognizes **bpeta**, **bpe** and **apy**. The binary solids form an isostructural set of cocrystals. We are



Scheme 1 (a) FTC and 4-pyridyl-containing coformers, (b) supramolecular cleft recognition, and (c) cleft recognition.

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† Electronic supplementary information (ESI) available. CCDC 1981473–1981476 experimental conditions, powder X-ray diffractograms, additional SCXRD data. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d0ce00474j

unaware of a case wherein the behaviour of a unimolecular cleft has been expressed supramolecularly.

Initially, **FTC** (40 mg, 0.162 mmol) was combined with **bpeta** (14.9 mg, 0.081 mmol) through liquid-assisted grinding (LAG, methanol)⁷ for 15 min. Powder X-ray diffraction (PXRD) revealed a new solid phase by the appearance of a new set of peaks (*e.g.*, prominent peaks at $2\theta = 11.7$, 15.7 , 16.5 and 24.9°) when compared to the starting materials (Fig. S3, ESI†). Recrystallization of the solid in warm methanol (2 mL) afforded single crystals suitable for single-crystal X-ray diffraction (SCXRD) after slow evaporation over a period of 2 days. The composition of the solid was $(\text{FTC})_2 \cdot (\text{bpeta})$ by ^1H NMR spectroscopy (Fig. S7–S11, ESI†) and SCXRD (Table 1). PXRD confirmed the solid phases to agree with those of grounded solids. The LAG was necessary to form the solid phases.⁸

Structural analysis of $(\text{FTC})_2 \cdot (\text{bpeta})$ revealed the components to assemble in the monoclinic space group $P2_1$. **FTC** adopts a J-shape conformation and assembles into homodimers through $\text{N-H}\cdots\text{N}$ hydrogen bonds (Table S2, ESI†).^{5,9} The orientation of the NH_2 groups is also sustained by weak intramolecular $\text{N-H}\cdots\text{F}$ hydrogen bonds similarly to structurally related molecules.¹⁰ The formation of the homodimers results in the terminal hydroxyl groups adopting a *cisoid*, or convergent, orientation ($\text{O}\cdots\text{O}$ separation: $14.676(5)$ to $14.951(4)$ Å). The hydroxyl groups interact with **bpeta** through $\text{O-H}\cdots\text{N}_{\text{pyr}}$ hydrogen bonds to generate a discrete three-component assembly (Fig. 1a). The *cisoid* disposition of the hydroxyl groups contrasts a *transoid* orientation in pure **FTC**.⁵ The **FTC** molecules, thus, effectively chelate¹¹ **bpeta** as a supramolecular linker. Cocrystal formation is further stabilized by $\pi\cdots\pi$ contacts between the 5-fluorocytosine rings and bipyridines.¹² The assemblies form columns that run along the *a*-axis in an $(\text{ABA})_n$ pattern ($\text{A} = \text{FTC}$, $\text{B} = \text{bpeta}$) sustained by $\pi\cdots\pi$ contacts. $\text{C-H}\cdots\text{S}$ contacts support parallel alignment of the columns in the *ac*-plane (Fig. 1b).¹³ The columns define a herringbone arrangement

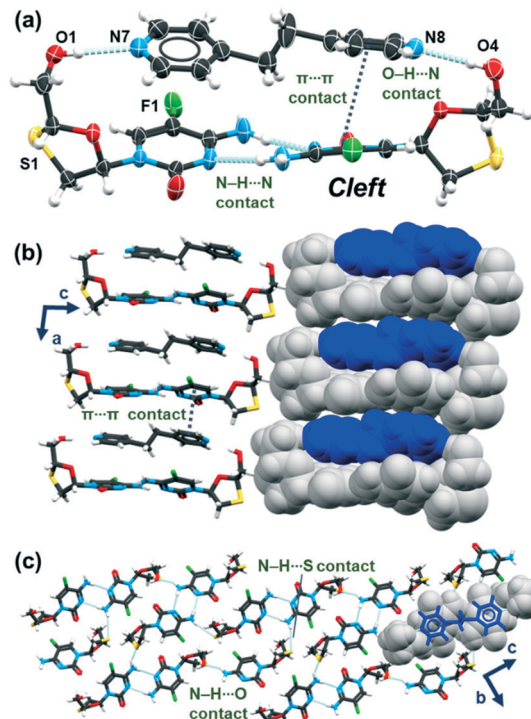


Fig. 1 X-ray structure $(\text{FTC})_2 \cdot (\text{bpeta})$: (a) three-component assembly, (b) 1D parallel columns, (c) ribbons.

in the *bc*-plane facilitated by $\text{N-H}\cdots\text{S}$ and $\text{N-H}\cdots\text{O}$ contacts (Fig. 1c).

Tolerance of the cleft-like behavior of **FTC** is evidenced by cocrystals $(\text{FTC})_2 \cdot (\text{bpe})$ and $(\text{FTC})_2 \cdot (\text{apy})$ (Fig. 2a and b). The solids, both of which are isostructural to $(\text{FTC})_2 \cdot (\text{bpeta})$, formed by LAG (methanol) as confirmed by PXRD analysis (Fig. S4–S6, ESI†). Single crystals suitable for SCXRD were obtained by recrystallization of the solids in warm methanol. The resulting clefts effectively “shrink-wrap” to accommodate the different bipyridines (Table 2).¹⁴ Thus, there is variability

Table 1 Crystallographic data $(\text{FTC})_2 \cdot (\text{bpeta})$, $(\text{FTC})_2 \cdot (\text{bpe})$, $(\text{FTC})_2 \cdot (\text{apy})$, and $(\text{FTC})_2 \cdot (\text{bipy})_2 \cdot \text{H}_2\text{O}$

Crystal data ^a	$(\text{FTC})_2 \cdot (\text{bpeta})$	$(\text{FTC})_2 \cdot (\text{bpe})$	$(\text{FTC})_2 \cdot (\text{apy})$	$(\text{FTC})_2 \cdot (\text{bipy})_2 \cdot \text{H}_2\text{O}$
Chemical formula	$(\text{C}_8\text{H}_{10}\text{FN}_3\text{O}_3\text{S})_2 \cdot (\text{C}_{12}\text{H}_{12}\text{N}_2)$	$(\text{C}_8\text{H}_{10}\text{FN}_3\text{O}_3\text{S})_2 \cdot (\text{C}_{12}\text{H}_{10}\text{N}_2)$	$(\text{C}_8\text{H}_{10}\text{FN}_3\text{O}_3\text{S})_2 \cdot (\text{C}_{12}\text{H}_8\text{N}_4)$	$(\text{C}_8\text{H}_{10}\text{FN}_3\text{O}_3\text{S})_2 \cdot (\text{C}_{10}\text{H}_8\text{N}_2)_2 \cdot \text{H}_2\text{O}$
MW (g mol ^{−1})	678.73	676.72	677.69	824.895
Space group	$P2_1$	$P2_1$	$P2_1$	$P2_1$
<i>a</i> (Å)	7.6333(8)	7.4883(7)	7.5024(8)	7.4668(10)
<i>b</i> (Å)	11.2359(11)	11.2957(11)	11.2194(11)	9.6700(14)
<i>c</i> (Å)	18.1058(18)	18.1735(18)	18.2134(18)	26.187(4)
<i>a</i> (deg)	90	90	90	90
<i>β</i> (deg)	99.640(5)	100.406(5)	100.538(5)	90.087(4)
<i>γ</i> (deg)	90	90	90	90
<i>V</i> (Å ³)	1531.0(3)	1511.9(3)	1507.2(3)	1890.8(5)
<i>Z</i>	2	2	2	2
<i>μ</i> (mm ^{−1})	0.243	0.246	0.249	0.215
<i>ρ</i> _{calcd} (g cm ^{−3})	1.472	1.486	1.493	1.449
<i>R</i> ₁ ^{b,c}	0.0292	0.0369	0.0487	0.0559
<i>wR</i> ₂ ^{d,e}	0.0700	0.0869	0.1157	0.1307
CCDC	1981473	1981474	1981475	1981476

^a $\lambda_{\text{MoK}\alpha} = 0.71073$ Å. ^b $F_o > 2\sigma(F_o)$. ^c $R_1 = \sum |F_o| - |F_c| / \sum |F_o|$. ^d All data. ^e $wR_2 = [\sum w(F_o^2 - F_c^2)^2 / \sum w(F_o^2)]^{1/2}$.

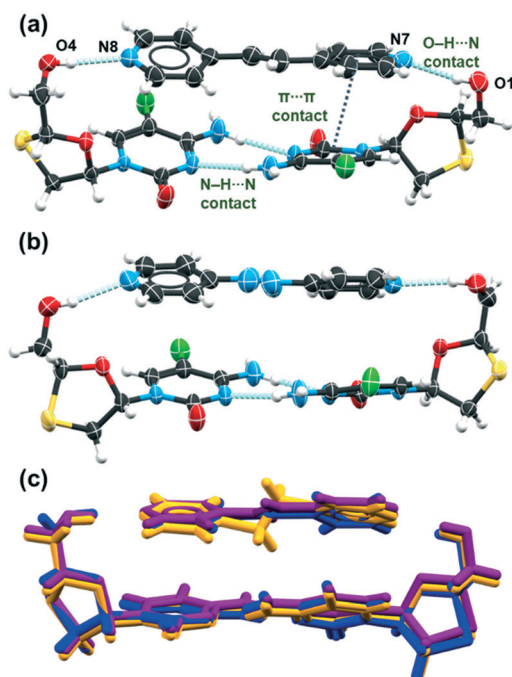


Fig. 2 X-ray structures: (a) $(\text{FTC})_2 \cdot (\text{bpe})$, (b) $(\text{FTC})_2 \cdot (\text{apy})$, and (c) overlay of $(\text{FTC})_2 \cdot (\text{bpe})$ (purple), $(\text{FTC})_2 \cdot (\text{apy})$ (blue), and $(\text{FTC})_2 \cdot (\text{bpeta})$ (orange).

Table 2 O–H...N_{pyr} hydrogen-bonds and selected $\pi \cdots \pi$ interactions of $(\text{FTC})_2 \cdot (\text{bpeta})$, $(\text{FTC})_2 \cdot (\text{bpe})$, $(\text{FTC})_2 \cdot (\text{apy})$ and $(\text{FTC})_2 \cdot (\text{bipy})_2 \cdot \text{H}_2\text{O}$

Cocrystal	$d_{\text{O} \cdots \text{H} \cdots \text{N}}$ (Å)	$d_{\text{O} \cdots \text{O}}$ (Å)	$d_{\text{N} \cdots \text{N}}$ (Å)
$(\text{FTC})_2 \cdot (\text{bpeta})$	2.838(4), 2.850(4)	14.889(4)	9.359(4)
$(\text{FTC})_2 \cdot (\text{bpe})$	2.863(4), 2.858(4)	14.951(4)	9.395(4)
$(\text{FTC})_2 \cdot (\text{apy})$	2.895(5), 2.887(5)	14.676(5)	9.024(5)

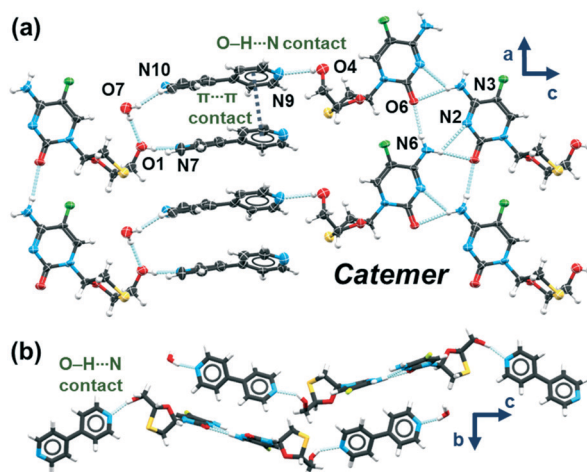
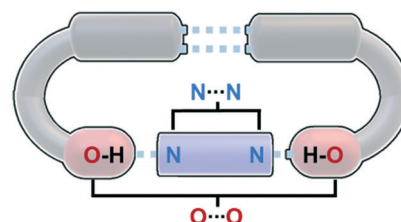


Fig. 3 X-ray structure $(\text{FTC})_2 \cdot (\text{bipy})_2 \cdot \text{H}_2\text{O}$: (a) hydrogen bonding and (b) adjacent 1D ribbons.

in spacer lengths ($\text{N} \cdots \text{N}$ 9.024(5) to 9.395(4) Å), which is then reflected in the hydroxyl group separations ($\text{O} \cdots \text{O}$ 14.676(5) to 14.951(4) Å) across the clefts. The three isostructural solids



Scheme 2 Selected metrics in $(\text{FTC})_2 \cdot (\text{bpeta})$, $(\text{FTC})_2 \cdot (\text{bpe})$ and $(\text{FTC})_2 \cdot (\text{apy})$.

exhibit a mutual crystal packing similarity index of 99.3% (Fig. 2c).¹⁵ UNI intermolecular potentials¹⁶ of isostructural cocrystals indicated a small variability in the total packing energies ($\Delta E_{\text{TOT}} = 11 \text{ kJ mol}^{-1}$), being $(\text{FTC})_2 \cdot (\text{apy})$ and $(\text{FTC})_2 \cdot (\text{bpeta})$ the lowest and highest energy packings (Table S3, ESI†).

An ‘error’ occurs in the self-assembly when **FTC** is introduced to the slightly smaller bipyridine **bipy** (Fig. 3). The components form the cocrystal hydrate $(\text{FTC})_2 \cdot (\text{bipy})_2 \cdot \text{H}_2\text{O}$ by LAG (methanol). Recrystallization of the solid in warm methanol afforded single crystals in the monoclinic space group $P2_1$ (Table S1, ESI†). **FTC** forms a hydrogen-bonded catemer (Fig. 3a) wherein the API, similar to pure **FTC**, adopts the *transoid* conformation (Fig. 3a).¹⁷ As a consequence of the assembly process, the components form 1D ribbons along the *c*-axis in an $(\text{ABCCB})_n$ pattern ($\text{A} = \text{H}_2\text{O}$, $\text{B} = \text{bipy}$, $\text{C} = \text{FTC}$). Adjacent ribbons interact by π -stacking of **bipy** along the *a*-axis (Fig. 3b). We attribute the formation of the ribbons to the shorter distance of the **bipy** guest ($\text{N} \cdots \text{N}$ 7.093(6), 7.117(8)). It is likely that the shorter length circumvents **bipy** to offer an element of preorganization to cleft formation, and thus allows a more efficiently packed hydrogen-bonded hydrated structure to form (Table S3, ESI†) (Scheme 2).

In this contribution, we have described supramolecular repurposing of the anti-HIV drug **FTC**. The API acts as a supramolecular cleft for bipyridine recognition in the solid state. Having identified the hydrogen-bonding recognition capabilities of **FTC**, we envisage possible supramolecular repurposing of other APIs,¹⁸ and similar molecules lined with multiple functional groups, to enable the formation of solids that exhibit additional properties (*e.g.*, photoactive, semiconductors, storage, separation).¹⁹

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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