

Tri- and tetrafluoropropionamides derived from chiral secondary amines – synthesis and the conformational studies†

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Cite this: *New J. Chem.*, 2014, 38, 3819

Received (in Montpellier, France)
3rd March 2014,
Accepted 2nd June 2014

DOI: 10.1039/c4nj00317a

www.rsc.org/njc

A convenient procedure for the preparation of tri- and tetrafluoropropionamides derived from *R*-(+)/*S*-(-)-*N*-methyl-1-phenylethylamine and cyclic pyrrolidine derivatives has been described. The X-ray analysis and the theoretical calculations have been used to study conformational analysis of obtained compounds. In contrast to single α -fluorine substituted amides, which preferred *anti* conformation around the F–C–C=O bond, for tetrafluorinated amides the additional trifluoromethyl group forces the conformation of the F–C–C=O bond as nearly *syn*.

Introduction

An amide bond is of key importance for all living organisms as the main chemical bond linking amino acids in peptides and proteins. Peptides play a crucial role in most of the biological processes, such as enzymatic catalysis, immune protection and others.^{1,2} Fluorinated amides are very interesting species in the design of peptide mimetics, as amino acid derivatives and as traditional fluorine modified drugs.^{3,4} It is well known that the introduction of fluorine or fluorine-containing groups into organic molecules can cause considerable changes in the physical, chemical and biological properties.^{5–7} Fluorination can change biological activity, as well as physical properties such as lipophilicity *etc.* Some examples of new drugs, being trifluoromethylated amido compounds, are efavirenz (anti-HIV)^{8,9} and odanacatib (inhibitor of cathepsin K)¹⁰ (Fig. 1). Moreover, α -hydroxy- α -trifluoromethylated amides arouse interest due to their structural analogy to some of the antiandrogens used in the treatment of prostate cancer in humans.¹¹

The introduction of a *gem*-difluoromethylene group has been proved to increase the biological activity of many pharmaceuticals containing β - and γ -lactam groups. For example, *gem*-difluoro- γ -lactam inhibits γ -lactamase, an enzyme responsible for bacterial resistance to γ -lactam antibiotics.¹² Moreover *gem*-difluoropropargyl amides are useful as suitable building blocks for the synthesis of fluorinated δ -lactams *via* cycloisomerization.¹³ Another example for application of the fluorine-containing compounds in synthesis

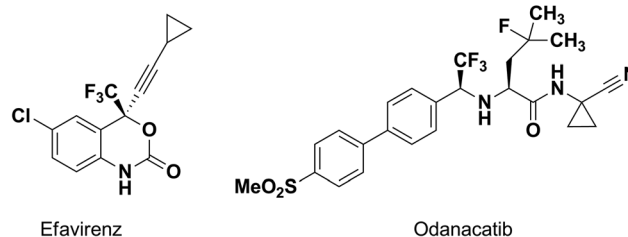


Fig. 1 Examples of biologically active trifluoromethylated amido compounds.

is trifluoroacetamide analogues that serve as efficient enantioselective nucleophilic trifluoromethylating reagents.^{14,15}

Recently, the tri- and tetrafluorinated amides have been studied in our group.¹⁶ As has been previously reported by Ishikawa,¹⁷ it has been shown that the hydrolysis of fluorinated enamines or the enamine–amine mixture has given the corresponding fluorinated amides. Additionally, the formation of the γ , δ -unsaturated amides, containing the fluorine atom or the CF₃ group at the α position to the carbonyl group, *via* Claisen rearrangement, has been reported.^{18–20}

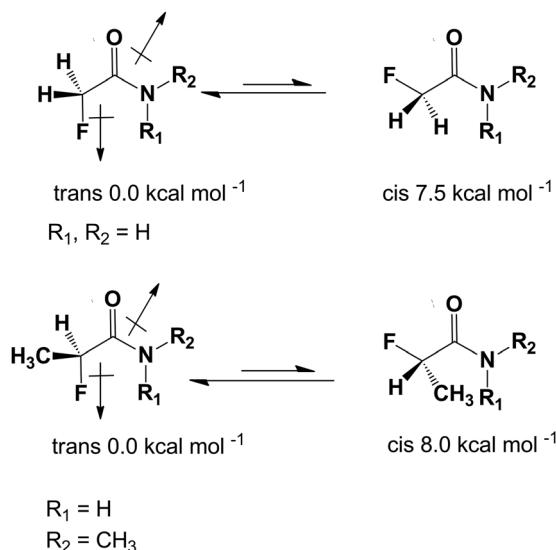
The presence of a fluorine atom in organic molecules can have a significant influence on the preferred conformation. O'Hagan *et al.* have shown recently that this effect is particularly noticeable in a group of fluorinated carbonyl compounds, especially in amides.^{21,22} When the fluorine atom is placed in the α position to the carbonyl group of an amide it adapts an *anti* conformation with respect to the C=O group and *syn* with respect to the C–N bond. This is a substantial effect of thermodynamic stability estimated to be about 7.5–8.0 kcal mol^{–1} (Scheme 1).⁷

Considering the presence of a fluorine atom in the β position with respect to the nitrogen atom of an amide, the fluorine

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† CCDC 989111. For crystallographic data in CIF or other electronic format see DOI: 10.1039/c4nj00317a





Scheme 1 The energy difference between the *trans* and *cis* conformers of α -fluoroamides.

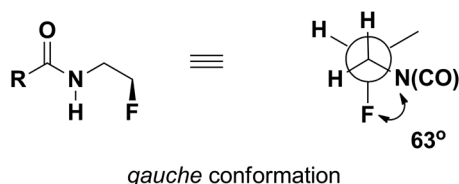


Fig. 2 Preferred orientation of a fluorine atom in the β position with respect to the nitrogen atom of an amide.

atom adopts a conformation in which the C–N and C–F bonds are oriented *gauche* to each other, with a dihedral angle of approximately 63° (Fig. 2).²³

In this paper we would like to focus on the synthesis and the conformational analysis of tri- and tetrafluoropropionic acid amides. Addition of hexafluoropropene and 1,1,3,3,3-pentafluoropropene to chiral secondary amines, distribution of reaction products, and their usefulness as the new reagents for deoxyfluorination will be published elsewhere.

Results and discussion

The synthesis of fluorinated amides has been based on the reactions of chiral secondary amines with commercially available 1,1,3,3,3-pentafluoropropene (PFP) or hexafluoropropene (HFP). As the model amines, two enantiomeric *R*-(+)/*S*-(–)-*N*-methyl-1-phenylethylamine (*R/S*-MPEA; series **a**) and cyclic pyrrolidine derivatives (series **b**) have been chosen. The first step of synthesis, in the case of reaction of PFP with MPEA, has given the tetrafluorinated enamine **1a** (Scheme 2) with a 73% yield, determined by ¹⁹F NMR.

Changing the solvents to THF or DCM, or increasing temperature (reflux) of this reaction had no influence on the yield. Analysis of ¹⁹F NMR spectra of **1a** confirmed the enamine formation,

where one signal of the CF₃ group appeared at δ : –52.7 ppm (dd) and the second signal from the vinylic fluorine atom at δ : –92.7 ppm (dq). The *Z*-configuration of enamine **1a** was established on the basis of vinylic fluorine and vicinal hydrogen atom coupling constant values. A characteristic coupling constant for *trans* arrangement is $J^3_{\text{F-H}} = 30.7$ Hz, whereas the second (q) coupling constant was equal to 15.2 Hz ($J^4_{\text{F-CF}_3}$). Usually for F and H in *trans* arrangement coupling constant J varies from 27–32 Hz, while for *cis* geometry of the double bond J ranges from 4–6 Hz.^{24–26} Exclusive formation of the *Z*-isomer of **1a** could be explained by preferential formation of the most stable zwitterion intermediate, having the CF₃ and NR₁,R₂ groups in antiperiplanar arrangement, followed by *anti* fluoride anion elimination (Scheme 3).^{17,27}

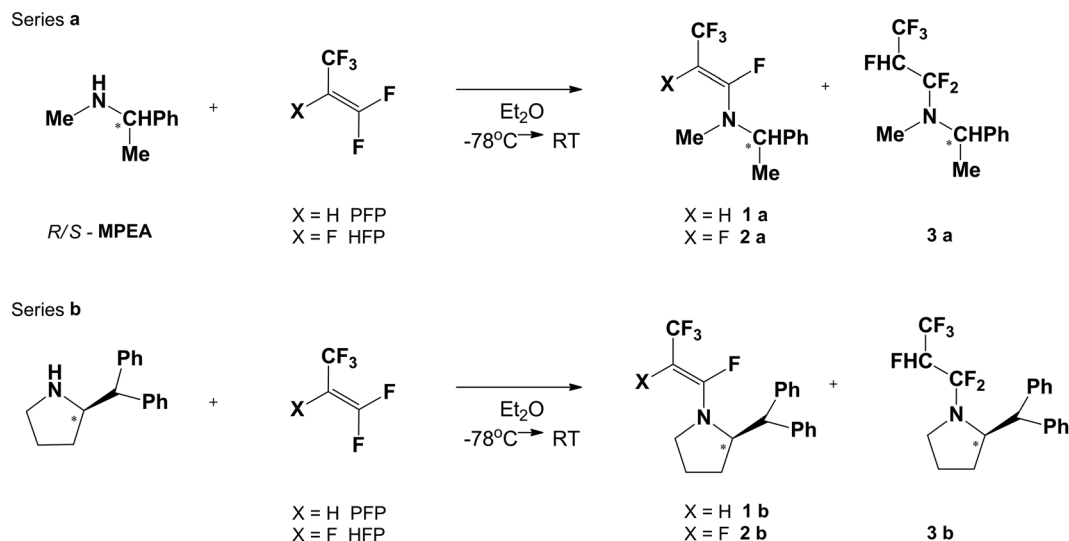
The high stereoselectivity of enamine formation is of striking difference from the previously reported results of the reaction of PFP with some pyrimidine or purine derivatives.²⁷ Thus, the reactions carried out in DMF (NaH, RT or 60–70 °C) yielded a high *Z/E* ratio of enamines in reaction mixtures. On the other hand, the reactions of PFP with Et₂NH or Me₂NH led to *Z*-enamines and only traces of corresponding saturated tertiary fluoroalkylamines.¹⁶ Analogous reaction of PFP with cyclic amine-pyrrolidine derivatives (series **b**) has given *Z*-enamines **1b** as the only product, with a 54% yield, which was determined by ¹⁹F NMR. The *Z*-geometry of a double bond in enamine **1b** was confirmed by chemical shift analysis of ¹⁹F NMR, *i.e.* δ : –51.7 ppm (CF₃ group), while a vinylic fluoride signal appeared at δ : –87.1 ppm with corresponding coupling constant $J^3_{\text{F-H}} = 29.2$ Hz (Table 1). These results are in good agreement with previously reported spectral characteristics of the PFP-pyrrolidine enamine product.¹⁶

Analogous reaction of appropriate chiral amine (**a** or **b**) with HFP has led to a mixture of enamine and fluorinated tertiary amine **2a–3a** with a 79% yield (the ratio of enamine/amine **2a/3a** was 1 : 2), while for **2b/3b** yield was 65% and the ratio was 1 : 32 (Scheme 3).

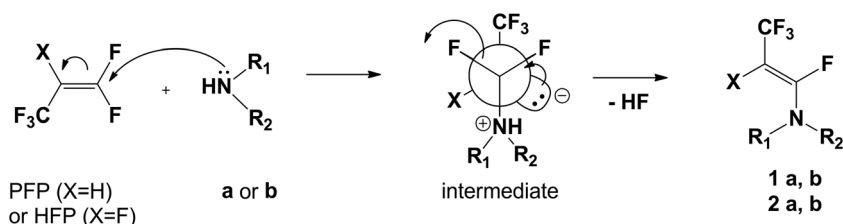
The analysis of ¹⁹F NMR spectra of **2a** has indicated the stereochemistry of the enamine double bond, similarly to the reaction with PFP, as *E*-geometry. It has been confirmed by higher values of coupling constants ($J^3_{\text{F-F}} = 118.7$ Hz, *trans*) between both vicinal fluorine atoms (δ : –114.5 and δ : –193.5), while signals of the CF₃ group have appeared at δ : –65.8 as dd ($J^4_{\text{CF}_3-\text{F}} = 13.6$ Hz, $J^3_{\text{CF}_3-\text{F}} = 23.1$ Hz). Analogous signals for **2b** have appeared at δ : –116.8 and –198.8 with $J^3_{\text{F-F}} = 116.4$ Hz (indicating *E*-enamine) and δ : –65.0 (CF₃). Noteworthy, the reaction of secondary amine with HFP has given mainly Michael type addition products – saturated amines **3a** and **3b**. The fluorine atom signals of pentafluorinated tertiary amine were located for **3a** at δ : –74.5 to –74.7 (CF₃) and diastereotopic (–CF₂–) fluorine atoms at δ : –81.5 to –87.5 and δ : –87.7 to –93.3 (multiplets), while for **3b** were situated at δ : –74.8 to –75.0 (CF₃), and at δ : –85.6 to –86.6 (1F) and at δ : –90.4 to –91.5 (1F). The high-field shifted fluorine atom signal of CHF₂CF₃ was situated at δ : –207.8 to –208.3 ppm (multiplet) for **3a**, and at δ : –205.5 to –206.0 for **3b** as a multiplet, respectively (Table 1).

The formations of major fluoroalkylamines and *E*-enamines, as well as their spectral properties are parallel with analogous





Scheme 2 The synthesis of enamines and mixtures of enamines–amines.



Scheme 3 The formation of the *Z*-isomer of **1a, b** (**2a, b**).

Table 1 ^{19}F NMR chemical shifts (ppm) and coupling constants of enamines **1a** (**1b**), **2a** (**2b**) and amines **3a** (**3b**)

Products	Yield ^a [%]	CF_3	F_α ^b	F_β ^b
Enamines	1a 73	−52.7 (dd, $J^4 = 15.2$ Hz, $J^3 = 7$ Hz)	−92.7 (dq, $J^3 = 30.7$ Hz, $J^4 = 15.2$ Hz)	n/a
	1b 54	−51.7 (dd, $J^4 = 14.6$ Hz, $J^3 = 7.1$ Hz)	−87.1 (dq, $J^3 = 29.2$ Hz, $J^4 = 14.6$ Hz)	n/a
	2a 24	−65.8 (dd, $J^4 = 13.6$ Hz, $J^3 = 23.1$ Hz)	−193.5 (dq, $J^3 = 118.7$ Hz, $J^4 = 13.5$ Hz)	−114.5 (dq, $J^3 = 118.8$ Hz, $J^4 = 22.6$ Hz)
	2b 2	−65.0 (dd, $J^4 = 14.4$ Hz, $J^3 = 22.8$ Hz)	−198.8 (dq, $J^3 = 118.1$ Hz, $J^4 = 13.4$ Hz)	−116.8 (dq, $J^3 = 116.4$ Hz, $J^4 = 22.8$ Hz)
Amines	3a 55	−74.5 to −74.7 (m)	−81.5 to −87.5 (m)	−207.8 to −208.3 (m)
	3b 63	−74.8 to −75.0 (m)	−87.7 to −93.3 (m) −85.6 to −86.6 (m) −90.4 to −91.5 (m)	−205.5 to −206.0 (m)

^a Yield determined by ^{19}F NMR using *m*-fluorotoluene as an internal standard ($\delta = -114.1$ (m, 1F)). ^b Positions α and β determined in relation to the nitrogen atom.

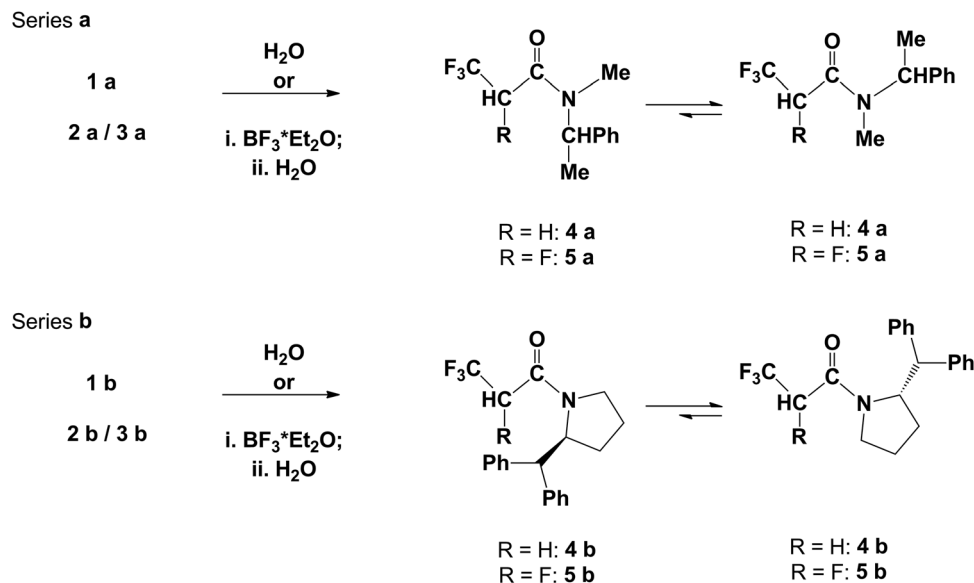
results reported by Ishikawa *et al.*¹⁷ Thus, reaction of diethylamine, *n*-Bu₂NH or piperidine with HFP yields an enamine–amine mixture with a different ratio depending on substituent bulkiness. In the case of piperidine as a starting material the only product is amine, while for Et₂NH as a substrate, the amine/enamine ratio is 1 : 3. On the other hand, the reaction of uracil, cytidine or guanine derivatives with HFP results in mixtures of *E/Z* isomeric enamines in an almost 1 : 1 ratio, while for adenine analogue the ratio is 2 : 1 – with a major product being *E*-geometry enamine.²⁷ Additionally, due to reaction conditions (secondary amine and sodium hydride)

fluorinated alkylamine (major product in our case) was not detected in the reaction mixture.

Subsequent H₂O addition to **1a**, **1b** enamine double bonds has given 3,3,3-trifluoropropionamide **4a**, **4b**, while reactions of water with **2a**, **2b** have given tetrafluorinated analogues **5a**, **5b**. Interestingly, contrary to prompt hydrolysis of enamines, in the case of hydrolysis of fluorinated amines **3a/3b** addition of Lewis acid such as BF₃·Et₂O has been necessary (Scheme 4).

Therefore, as a result of water addition to **1a** and **1b**, the trifluoropropionamide derivatives **4a** and **4b** were isolated from





Scheme 4 The hydrolysis reaction of 1a, 1b (2a, 3a/2b, 3b).

Table 2 ^1H NMR, ^{19}F NMR chemical shifts (ppm) of trifluoropropionamide derivatives and a reference amide IPMA

	Major					Minor				
	CF_3	CF_3-CH_2	N-R_1^a	N-R_2^a	CH_3^b	CF_3	CF_3-CH_2	N-R_1^a	N-R_2^a	CH_3^b
IPMA	n/a	n/a	2.83	4.52	1.03	n/a	n/a	2.70	3.92	1.15
4a	-62.9	3.18	2.60	5.99	1.42	-62.8	3.26	2.65	4.95	1.56
4b	-62.7	2.44	3.34	4.43	n/a	-63.3	2.49	3.56/3.87	3.17	n/a

^a IPMA: $\text{R}_1 = \text{CH}_3$, $\text{R}_2 = \text{CH}(\text{CH}_3)_2$; 4a: $\text{R}_1 = \text{CH}_3$, $\text{R}_2 = \text{CHCH}_3\text{Ph}$; 4b: $\text{R}_1 = \text{CH}_2\text{CH}_2$, $\text{R}_2 = \text{CHCH}(\text{Ph})_2$. ^b IPMA: $\text{CH}(\text{CH}_3)_2$; 4a: CHCH_3Ph .

reaction mixtures with 57% and 38% yield, respectively. Analysis of ^{19}F NMR spectra of 4a has indicated two triplets at δ : -62.9 and δ : -62.8 ($J_{\text{CF}_3-\text{H}} = 10.0$ Hz) in a 71:29 ratio. The phenomenon of the existence of two sets of signals in magnetic resonance spectra is well known in the case of amides. Thus, due to hindered rotation about a partially double amide C-N bond, both *N*-substituent groups are chemically non-equivalent and compounds can exist as two rotamers: cisoid and transoid, considering the orientation with respect to the carbonyl group. Alike, in the ^1H NMR spectrum of *N,N*-dimethylformamide (DMF) or *N,N*-dimethylacetamide (DMA), both *N*-methyl groups have separate signals.²⁸ The observed chemical shift values in the ^1H NMR spectrum of methyl groups are δ : 2.85 ppm and δ : 2.94 ppm for DMF, while for DMA they appear at δ : 2.93 ppm and δ : 3.03 ppm, respectively. In the case of *N,N*-dimethyl-3,3,3-trifluoropropionamide, in the ^1H NMR spectrum signals from H atoms of both *N*-neighboring methyl groups are observed at δ : 3.3 and δ : 3.4 ppm, while in ^{19}F NMR spectra the fluorine signal of the CF_3 group was located at δ : -63.2 ppm as a triplet.¹⁶ Similarly, the compound 4a can exist as two rotamers (Scheme 4): transoid (*trans* 4a) with a larger substituent at nitrogen located in the opposite direction to the carbonyl group and in the reverse direction as rotamer cisoid (*cis* 4a). The analysis of ^{19}F NMR spectra has shown two signals: the one of the major rotamer has been located at δ : -62.9, while for the minor

rotamer of 4a it has appeared at δ : -62.8 (CF_3 group). Considering the ^1H NMR spectra of the 4a major and minor rotamer, they indicated that the signals of *C*-methyl and *N*-methyl groups for the major rotamer have been located at δ : 1.42 and δ : 2.60, whereas for the minor rotamer have appeared at δ : 1.56 and δ : 2.65. Also methylene protons of the trifluoropropionic part for the major rotamer have been slightly shifted upfield to δ : 3.18, compared to δ : 3.26 for the minor one. Interestingly, while the chemical shift of aromatic protons for both rotamers was parallel, the difference in the chemical shift of benzylic protons for major and minor rotamers is $\Delta\delta = 1.03$ ppm (δ : 5.99 for the major and δ : 4.95 for the minor rotamer, respectively) (Table 2).

The relationship of the chemical shift values and *N*-neighboring group arrangement in disubstituted amides has already been reported. In the case of *N*-isopropyl-*N*-methyl acetamides (IPMA) the predominant *cis* geometry (*Z*) of the *i*-Pr and the carbonyl $\text{C}=\text{O}$ groups has been determined.²⁹ Thus, the *cis* rotamer signals for *N*-methyl and *N*-CH(CH₃)₂ protons occur at δ : 4.52 and δ : 2.83, whereas for the minor *trans* rotamer they occur at δ : 3.92 and δ : 2.70, respectively. The comparison of chemical shift values of benzylic and *N*-methyl protons for both rotamers of 4a, and similarity to IPMA relationships have helped to establish the main rotamer of 4a as having *cis* geometry, while the minor rotamer has been determined as having *trans* geometry.



Similarly, in ^{13}C NMR spectra of *cis* and *trans* isomers of **4a**, both sets of signals are typical and analogous for both isomers, with the exceptions of *N*-neighboring atoms like carbons of the $\text{C}-\text{CH}_3$ signal at δ : 15.4 for *cis* **4a** and at δ : 17.9 for *trans* **4a**, whereas the *N*-methyl group signal for *cis* **4a** has been located at δ : 30.1 and for *trans* **4a** at δ : 28.4. Simultaneously, similarly to ^1H NMR spectra, the largest chemical shift difference has been observed in the case of the benzyl carbon atom signals ($\text{C}-\text{Ph}$). Thus, the signal for major *cis* **4a** has been shifted downfield (δ : 50.8) compared to *trans* **4a** (δ : 55.7).

As a result of **1b** hydrolysis, the 3,3,3-trifluoropropionamide **4b**, as a mixture of two rotamers with a 53:47 ratio, has been isolated. The comparison of the signals in ^1H NMR spectra, due to the larger difference in chemical shifts of protons at chiral carbon, has helped matching the major rotamer with *N*-CHC proton shift δ : 4.43 as *cis* **4b**. The minor rotamer with the analogous proton signal situated at δ : 3.17 has been determined as *trans* **4b**. These data are also in agreement with spectral characteristics of pyrrolidine acetamide.^{30,31} While the chemical shifts of the remaining analogous protons in both **4b** rotamers have been similar, in ^{19}F NMR spectra the *cis* and *trans* **4b** rotamers of CF_3 group signals have appeared at δ : -62.7 and -63.3, respectively.

Next, the hydrolysis reactions of enamine-amine mixture **2a-3a** and **2b-3b** have been performed. The hydrolysis of the **2a-3a** mixture has led to formation of **5a** being a mixture of diastereomers (*R,R*)/(*R,S*) **5a** (in 49:51 ratio) with a 78% isolated yield. Separation of both diastereoisomers has given two fractions (after column chromatography) – crystals and the oil – each consisting of two rotameric forms (77:23 ratio).

Similarly to compounds **4a** and **4b** the comparison of the ^1H NMR spectra and the correlation of proton chemical shifts of the *N*-neighboring group in two rotamers with an amide geometry of IPMA and **5a** (with a ratio 77:23) have helped to determine the geometry of major rotamers of both diastereomers of **5a** as *cis* (*Z*) with the *N*-bulky group oriented in the same direction as the carbonyl group. Thus, the signals of benzylic protons for major rotamers *cis* **5a** have been down-field shifted and have been located at δ : 6.02 (*R,S*) and 6.05 (*R,R*) while those of *N*-methyl protons have been situated at δ : 2.74 (*R,S*) and 2.76 (*R,R*). The methylbenzyl $\text{C}-\text{CH}_3$ protons have been recorded at δ : 1.55 (*R,S*) and 1.54 (*R,R*). On the other side, the chemical shift of analogous benzylic (δ : 5.24 and 5.30 for both diastereoisomers) and methylic protons (δ : 2.71/2.72 and 1.67/1.66 for *R,S/R,R* diastereoisomers, respectively) in minor rotamers has

indicated the *trans* geometry of **5a** (Table 3). These data are consistent with ^1H NMR and ^{13}C NMR spectra reported already for *N,N*-dimethylfluoroacetamide (DMFA) and *N,N*-dimethyl- α -fluoropropionamide (DMFP), as well as for *N,N*-dimethyl-3,3,3-trifluoroacetamide.^{17,32}

Although, the largest difference between chemical shifts of both diastereomers of **5a** has been observed at stereogenic carbon atoms, these notices have not allowed us to clearly distinguish the configuration at the chiral centre in both diastereomers. While for major rotamers in both diastereoisomers **5a**, the chemical shifts of appropriate amine moiety protons had been similar ($\Delta\delta \leq 0.3$), the signals derived from tetrafluorinated amide parts have slightly varied *i.e.* α -fluorines (δ : -199.2 and -199.5) with α -protons (δ : 5.43 and 5.42) and for the trifluoromethyl group (δ : -75.6). In contrast, for minor rotamers, the signals of α -fluorine atoms have been observed at δ : -196.1 and -197.6. The chemical shifts of the rest of the appropriate protons in the whole molecule have been similar.

The comparison of chemical shift values for diastereomeric α -methoxy- α -(trifluoromethyl)phenyl acetic acid (MPTA) analogues could be applied, as has already been applied, in determining absolute configuration of secondary hydroxyl or amine derivatives.³³ Thus, for the (*R*)-MTPA amide derivatives of the (*R*)-1-phenylethylamine *e.g.* (*R,R*)-MTPAPEA, the α -positioned CF_3 group has been down-field shifted and has been recorded at δ : -70.98, compared to the analogous signal of the (*R,S*) diastereoisomer located at δ : -71.23 in ^{19}F NMR spectra. As a comparison, the signal of the α -fluorine atom, vicinal to the $\text{C}=\text{O}$ group in the crystal of *anti* **5a**, has been observed at δ : -196.1 and has resulted in determination of configuration of the **5a** diastereoisomer as (*R,R*). Analogously, a signal of α -fluorine in the ^{19}F NMR spectrum for *anti* **5a** has been placed at δ : -197.6, indicating the configuration (*R,S*) for an oily **5a**.

Analogous hydrolysis reaction of the **2b-3b** mixture has given two diastereoisomers **5b**, in a 53:47 ratio with 98% isolated yield, each isomer with a 77:23 rotamers ratio. The diastereoisomers have been inseparable using column chromatography. As the signals at δ : 4.08–4.17 ppm (m) and δ : 3.40–3.64 ppm (m) corresponding to *N*-CHC and *N*-CH₂–, respectively, have been matched by integration to the major rotamer having *cis* geometry of the *N*-bulky group and the $\text{C}=\text{O}$ bond in one diastereomer **5b**, the minor one has had the analogous signals located in the range δ : 4.00–4.08 ppm (m) and at δ : 3.40–3.64 ppm (m) (in the same range) indicating *trans* orientation analogous groups in **5b**. Unfortunately, for the major diastereoisomer of **5b**, the signals of

Table 3 ^1H NMR, ^{19}F NMR chemical shifts (ppm) of tetrafluoropropionamide derivatives and a reference amide IPMA

	Major					Minor				
	CF_3	CF/CH	$\text{N}-\text{R}_1^a$	$\text{N}-\text{R}_2^a$	CH_3^b	CF_3	CF/CH	$\text{N}-\text{R}_1^a$	$\text{N}-\text{R}_2^a$	CH_3^b
IPMA	n/a	n/a	2.83	4.52	1.03	n/a	n/a	2.70	3.92	1.15
5a (<i>R,R</i>)	-75.6	-199.2/5.43	2.76	6.05	1.54	-75.5	-196.1/5.52	2.72	5.30	1.66
5a (<i>R,S</i>)	-75.6	-199.5/5.42	2.74	6.02	1.55	-75.5	-197.6/5.53	2.71	5.24	1.67
5b (<i>S,S</i>)	-75.7	-200.7/5.11	3.08–3.28	3.74–3.82	n/a	-76.6	-202.7/5.05	3.08–3.28	3.74–3.82	n/a
5b (<i>S,R</i>)	-75.7	-200.6/4.92	3.40–3.64	4.08–4.17	n/a	-76.6	-203.7/4.93	3.40–3.64	4.00–4.08	n/a

^a IPMA: $\text{R}_1 = \text{CH}_3$, $\text{R}_2 = \text{CH}(\text{CH}_3)_2$; **5a**: $\text{R}_1 = \text{CH}_3$, $\text{R}_2 = \text{CHCH}_3\text{Ph}$; **5b**: $\text{R}_1 = \text{CH}_2\text{CH}_2$, $\text{R}_2 = \text{CHCH}(\text{Ph})_2$. ^b IPMA: $\text{CH}(\text{CH}_3)_2$; **5a**: CHCH_3Ph .



hydrogen atoms at chiral carbon, in the amine part, were situated in the same range (δ : 3.74–3.82 ppm) as well as the slightly shifted signal of the *N*-neighboring methylene group (δ : 3.08–3.28 ppm).

Additionally, for both major diastereomers *cis* **5b** the signals derived from tetrafluorinated amide parts have slightly varied *i.e.* α -fluorines (δ : –200.7 and 200.6), α -protons (δ : 5.11 and 4.92) and the trifluoromethyl group (δ : –75.7). In contrast, for minor *trans* rotamers of both diastereomers **5b**, the signals of α -fluorine atoms have been visible at δ : –202.7 and –203.7 ppm, while the chemical shifts of α -protons have been observed at δ : 5.05 and δ : 4.93 ppm. The signals of CF₃ groups have been situated at δ : –76.6 ppm. Therefore, aiming to absolute configuration assignment in both diastereomeric **5b** (as a pyrrolidine part has already had *S*-configuration), the correlation of fluorine chemical shift values of CHF(CF₃) in minor rotamers of **5b** has been performed. The signal of the α -fluorine atom, vicinal to the C=O group in *trans* **5b** has appeared at δ : –202.7 ppm and could determine the **5b** diastereomer as (*S,S*). Analogously, the signal of α -fluorine in the ¹⁹F NMR spectrum for *trans* **5b** has been located at δ : –203.7 ppm, indicating the configuration (*S,R*) **5b**. This assignment is in agreement with the ones performed for (*R,R*)/(*R,S*) **5a** and (*S,S*)/(*S,R*) **5a**, as well as (*R,R*)-MTPAPEA.³² However, further studies also involving the crystal structure determination would be necessary to the unambiguous assignment of absolute configuration of chiral atoms in the fluorinated amide part of both diastereomers **5b**.

Conformational analysis of tetrafluorinated amide – theoretical calculations

While the orientation of the amine part toward the carbonyl group is easily detectable based on NMR spectra, the influence of fluorine atoms on steric and electronic effects in molecules and preferable conformation of molecules is the well-known phenomenon. Thus, compound **5a** could exist in a few conformations taking into account only fluorine and CF₃ group orientation with respect to the carbonyl group (Fig. 3). The *cis/trans* notation deals with the configuration of partially double amide bond C(O)–N, while *syn/anti* notation is related to the F–C–C=O torsion angle (*syn* is 0°, *anti* is 180°).

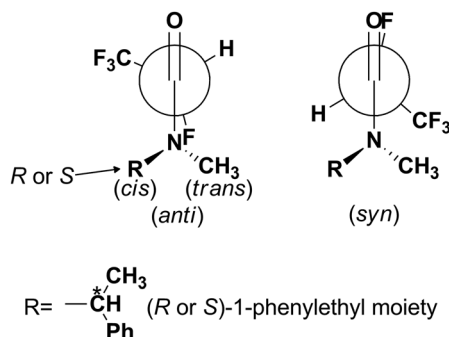


Fig. 3 Anti and syn conformers of **5a** *R,R cis* (or *R,S cis*).

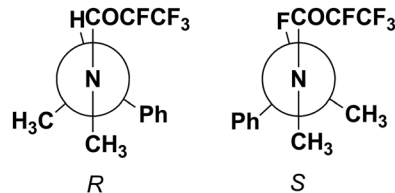


Fig. 4 Most stable conformational arrangement of the (*R* or *S*)-1-phenylethyl moiety, which was used during all optimizations at the DFT level to quantify the dependence of the conformation on the F–C–C=O torsion angle (Fig. 5).

The most stable conformational arrangement of the (*R* or *S*)-1-phenylethyl moiety has been found (Fig. 4) and used during all optimizations at the DFT level in order to quantify the dependence of the conformation on the F–C–C=O torsion angle.

The obtained potential energy surfaces (Fig. 5) for **5a**: *R,R trans*, *R,R cis*, *R,S trans* and *R,S cis* are very similar and all show three energy minima (Fig. 6): *minimum-1* (deepest minimum at F–C–C=O torsion angle about 150°), *minimum-2* (very shallow minimum at about 240°) and *minimum-3* (at about 330°), whose results are in agreement with the data obtained by Abraham *et al.* for *N,N*-dimethyl- α -fluoropropionamide (DMFP).³² In contrast to α -fluorinated *N*-monosubstituted amides, where the most stable conformation is that with the fluorine atom *anti* to the carbonyl group,²² in the case of α -fluorinated *N,N*-disubstituted amides conformation *anti* is unstable due to the steric congestion (Fig. 5). Two of the three potential energy minima obtained for the **5a** are considered to be stable conformers: *minimum-1* and *minimum-3* (with an F–C–C=O torsion angle of about 330° which corresponds to the X-ray structure angle of 331.3°) (Fig. 7). There, the X-ray structure is not the global minimum potential energy conformation in the isolated state. Further, the structure of *minimum-2* seems to be even a shoulder on the potential energy surface and it is considered to be the least stable minimum due to the relatively high potential energy and a very low energy barrier of rotation (relaxation) towards the global minimum structure (*minimum-1*). This, on the other hand, is in contrast with the calculation for DMFP, which has a CH₃ group instead of a CF₃ group.³² Then, the structure corresponding to the *minimum-2*, having the CH₃ group in the *syn* position to the carbonyl C=O, was the most stable one. In our case, the structure of *minimum-2* has the CF₃ group in the *syn* position to carbonyl which seems to be less preferred due to the repulsion between fluorine and oxygen atoms. This observation shows how additional fluorine atoms can influence the conformation equilibrium.

These data are in agreement with the crystallographic structure of solid **5a** (Fig. 7) and *R,R* absolute configuration determined from ¹⁹F NMR data. The X-ray structure shows that molecule is observed as the *syn* conformer, with the O(1)–C(1)–N(1)–C(4) dihedral angle of 355.6°.

Analysis of the X-ray structure of amide *R,R syn* **5a** shows that the C–F bond α to the amide carbonyl group is orientated nearly *anti* to the N–C amide bond, with the F(1)–C(2)–C(1)–N(1) dihedral angle of 151.9° and nearly *syn* to the C=O bond, with the F(1)–C(2)–C(1)–O(1) dihedral angle of 331.3° (Table 4).



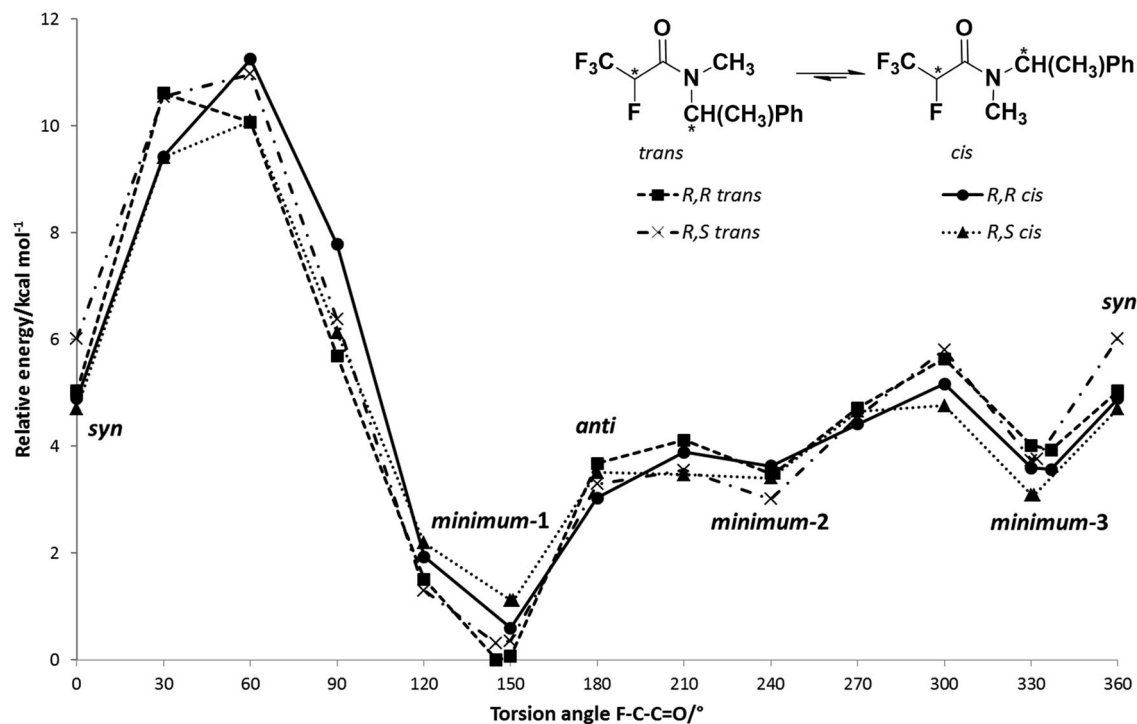


Fig. 5 Potential energy surfaces for **5a**: *R,R* trans, *R,R* cis, *R,S* trans and *R,S* cis at the wB97XD/cc-pVDZ level.

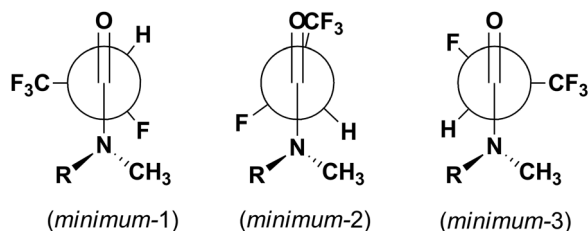


Fig. 6 The geometries of three energy minima on the potential energy surface for **5a** *R,R* cis (or *R,S* cis).

Table 4 Torsion angles (°) of amide (*R,R*)-syn **5a**

Torsion angles (°)	
C(3)–C(2)–C(1)–O(1)	88.2
H(2)A–C(2)–C(1)–O(1)	211.2
F(1)–C(2)–C(1)–O(1)	331.3
F(1)–C(2)–C(1)–N(1)	151.9
C(3)–C(2)–C(1)–N(1)	268.8
H(2)A–C(2)–C(1)–N(1)	31.8
O(1)–C(1)–N(1)–C(12)	176.3
O(1)–C(1)–N(1)–C(4)	355.6

Table 5 The short intermolecular contacts in the crystal of **5a** (Å and °)

D–H...A	D–H	H...A	D...A	<D–H...A	Symmetry code
C2–H2a...O1 ^a	0.960	2.296	3.143(5)	144.17	–x, 0.5 + y, 1.5 – z
C5–H5a...F1 ^a	0.960	2.624	3.470(5)	147.42	–x, y – 0.5, 1.5 – z
C12–H12c...O1 ^a	0.960	2.703	3.656(5)	171.71	1 – x, y, z

^a The H-acceptors transformed according to the symmetry code of the last column.

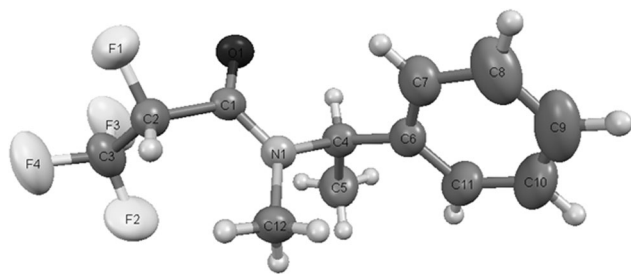


Fig. 7 A perspective drawing and atom labelling of molecule *R,R* syn **5a**. The thermal ellipsoids have been drawn at the 40% probability level. The hydrogen atoms are shown as small spheres of arbitrary diameter.

It is deviation from the preferred *anti* conformation of the F–C–C=O for a single fluorine substituent at the α -position with respect to a carbonyl group in α -fluoroamides.^{21,22} Following these observations it has been predicted that the preferred conformation around an amide bond could be caused by the

replacement of the CH₃ group by the trifluoromethyl group in the obtained compounds. Not only the fluorine atom but also the trifluoromethyl group dictates the conformation of these amides.

The shortest intermolecular contacts listed in Table 5 show that the strongest interactions can be associated with the proton at C2 in the most electronegative region of fluorine substituents. These and other shortest interactions bind the molecules along crystallographic plane (001) [Fig. 8].



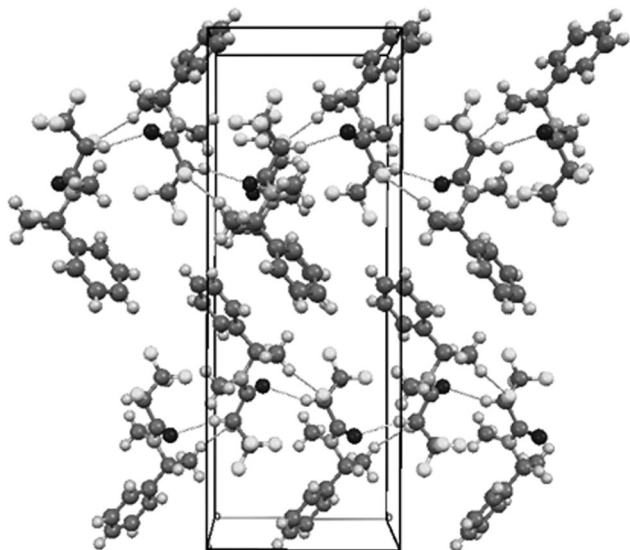


Fig. 8 Autostereographic projection of the crystal structure of *R,R* syn **5a** viewed down [100], with the shortest contacts indicated as the dashed lines.³⁴

IR spectra

The IR (C=O) wavenumbers in the series **4a–5a** have indicated the influence of the C–F bond polarization on the increasing positive charge density on carbon atoms, making the C=O bond shorter, due to the electrostatic stability of the C–F bond which is consistent with the results from the quantum mechanical calculations. As a reference point, the amide **4a** with a IR (C=O) ν^{-1} 1659 cm^{-1} has been selected, while for tetrafluorinated (*R,R*) **5a** the wavenumber has equaled to 1655 cm^{-1} . Analogously, for the (*R,S*) diastereomer **5a** the corresponding IR (C=O) ν^{-1} has equaled to 1667 cm^{-1} (Fig. 9).

Probably, in the case of the studied diastereoisomers, the absolute configuration of the carbon atom placed adjacent to the amide bond has an impact on the general polarization, creating the large difference in C=O stretching frequency between the diastereoisomers (*R,R*) and (*R,S*) of **5a**. These data are parallel with the results obtained for (*S*)-2-fluoro-*N*-(2-fluoroethyl)-propionamide (wavenumber value 1671 cm^{-1}),²¹ indicating an “alpha-fluoroamide effect” and adopting a preferred *gauche* conformation of the C–N(C=O) bonds in the α -fluoroamide moiety seen in the crystal structure.

Conclusion

Our experimental results have demonstrated that the hydrolysis reactions of enamines have given 3,3,3-trifluoropropionamides,

while an addition of water to enamine–amine mixtures has given tetrafluorinated analogues. The combined spectroscopic, X-ray analyses and theoretical calculations have given us evidence of the differences in the conformational geometry of the α -fluorine substituted amides and newly synthesized tri- and tetrafluorinated amide derivatives. According to the latest version of the Cambridge Structural Database, no other structures containing the C(F₃)C(HF)C(=O)N– have been determined. In contrast to single α -fluorine substituted amides which preferred *anti* conformation of F–C–C=O, for the tetrafluorinated amides the additional trifluoromethyl group has enforced that the C–F bond α to the amide carbonyl group is orientated nearly *anti* to the N–C amide bond, and nearly *syn* to the C=O bond. These data could be considered in a design of important CF₃-building blocks employed in the synthesis of useful compounds such as peptide analogs.

Experimental section

General procedures

¹H (Me₄Si) NMR spectra were determined with solutions in CDCl₃ at 300 MHz, ¹³C (Me₄Si) at 75 MHz and ¹⁹F NMR (CCl₃F) at 282 MHz. The yields of reaction products were conveniently evaluated by ¹⁹F NMR in CDCl₃ using *m*-fluorotoluene as an internal standard. Mass spectra were obtained by electron impact (MS-EI) techniques with an ionizing energy of 70 eV, unless otherwise noted. Optical rotations were measured at 20 °C in CHCl₃ using a 243 B Perkin-Elmer polarimeter. [α]_D values were determined at 589 nm. IR spectra were performed using a spectrometer FT-IR IFS 66/s from Bruker. Reagent grade chemicals were used and solvents were dried by refluxing with sodium metal-benzophenone (THF), with CaH₂ (CH₂Cl₂), with NaH (Et₂O) and distilled under an argon atmosphere. All moisture sensitive reactions were carried out under an argon atmosphere using oven-dried glassware. Reaction temperatures below 0 °C were obtained using a cooling bath (dry ice/isopropanol). Thin-layer chromatography (TLC) was performed on Merck Kieselgel 60-F₂₅₄ with EtOAc–hexane as developing systems. Visualization of the reaction products was achieved using UV light (254 nm) and a standard procedure (solution of phosphomolybdic acid or KMnO₄). Merck Kieselgel 60 (230–400 mesh) was used for column chromatography. 1,1,3,3,3-Pentafluoropropene (PFP) and hexafluoropropene (HFP) were provided to us by SynQuest Laboratories Ltd. *R*-(+)- and (*S*)-(–)-*N*, α -dimethylbenzylamine (**a**) and BF₃·Et₂O were supplied by Sigma Aldrich. 2(*S*)-2(Diphenylmethyl)pyrrolidine (**b**) was achieved according to the known literature method.^{35–37} All products

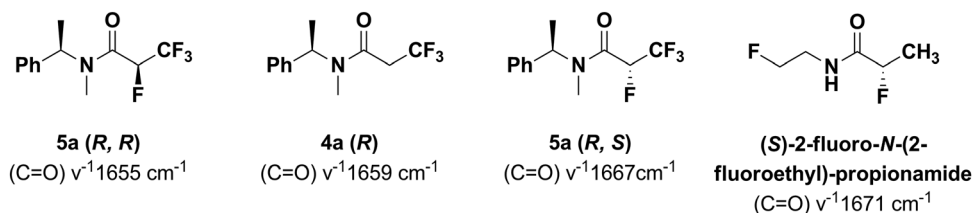


Fig. 9 IR (C=O) wavenumbers of series **4a–5a** in comparison with (*S*)-2-fluoro-*N*-(2-fluoroethyl)-propionamide.²¹



of the analogous reactions provided with (*R*) or (*S*) *N*, α -dimethylbenzylamine gave the same result with the identical spectral data (with an exception of $[\alpha]$ -typical for enantiomers).

Theoretical calculations. The quantum mechanical calculations of potential energy in a vacuum with the DFT (wB97XD) method have been performed using the GAUSSIAN03 program,³⁸ in order to systematically search for possible conformations. The long-range corrected hybrid functional wB97XD³⁹ with the correlation-consistent polarized basis set cc-pVDZ⁴⁰ has been used as the necessary compromise between the desired accuracy and the computational time needed.

Crystal structure analysis. The structure of compound *R,R* syn **5a** was determined by single-crystal diffraction. A KuMa KM-4 CCD four-circle diffractometer with MoK α radiation (0.71073 Å) was used. The structure was solved by direct methods with program SHELXS and refined by full-matrix least squares on F^2 by SHELXL.⁴¹ Hydrogen atoms were located from the molecular geometry at idealized positions and assigned isotropic thermal parameters depending on the equivalent displacement parameters of their carriers.

X-ray crystal structure analysis of **5a** (CCDC 989111):† formula C₁₂H₁₃F₄NO, $M = 263.23$, colourless crystal 0.30 × 0.10 × 0.10 mm, $a = 6.07446(13)$ Å, $b = 8.98778(19)$ Å, $c = 24.2410(6)$ Å, $\beta = 90^\circ$, $V = 1323.46(5)$ Å³, $\rho_{\text{calc}} = 1.321$ g cm⁻³, $\mu = 1.075$ mm⁻¹, $Z = 4$, orthorhombic, space group $P2_12_12_1$, $\lambda = 1.54184$ Å, $T = 293(2)$ K, θ scans, reflections collected/unique 21 223/2734 [$R(\text{int}) = 0.0301$], 164 refined parameters, $R = 0.0441$, $wR_2 = 0.1384$, goodness-of-fit on $F^2 = 1.106$, max. residual electron density 0.257 and -0.164 e Å⁻³.

General procedure for the synthesis of enamines

Chiral secondary amine (1 equiv.) and dried solvent (3 mL) were placed in a glass pressure-vessel. The solution was cooled to -78 °C and the pressure was lowered. Next, the excess of 1,1,3,3,3-pentafluoropropene (PFP) was transferred to the glass tube and the vessel was sealed. The cooling bath was removed and the mixture was brought to the room temperature with stirring overnight. After (approximately) 16–24 hours the crude mixture was used for the next step.

Note: The choice of solvent has depended on a solubility of amines. The elevated temperature of reaction (reflux) has not affected the yield.

(*R,Z*)-1,3,3,3-Tetrafluoro-*N*-methyl-*N*-(1-phenylethyl)prop-1-en-1-amine 1a. Reaction of PFP (180 mg, 1.36 mmol) and *R*-(+)-*N*, α -dimethylbenzylamine (0.1 mL, 92 mg, 0.68 mmol) in dried THF, according to the general procedure, gave a mixture of adduct **1a** (yield 73%) and traces of amide **4a**.

1a ¹⁹F NMR δ : -52.75 (3F, dd, $J_{\text{CF}_3-\text{F}} = 15.2$ Hz, $J_{\text{CF}_3-\text{H}} = 7.0$ Hz, CF₃), -92.73 (1F, dq, $J_{\text{F-H}} = 30.7$ Hz, $J_{\text{F-CF}_3} = 15.2$ Hz, F).

(*S,Z*)-2-Benzhydryl-1-(1,3,3,3-tetrafluoroprop-1-enyl)pyrrolidine 1b. Reaction of PFP (112 mg, 0.85 mmol) and 2(*S*)-2(diphenylmethyl)-pyrrolidine (100 mg, 0.42 mmol) in dried CH₂Cl₂, according to the general procedure led to a mixture of adduct **1b** (yield 54%) and traces of amide **4b**.

1b ¹⁹F NMR δ : -51.68 (3F, dd, $J_{\text{CF}_3-\text{F}} = 14.6$ Hz, $J_{\text{CF}_3-\text{H}} = 7.1$ Hz, CF₃), -87.15 (1F, dq, $J_{\text{F-H}} = 29.2$ Hz, $J_{\text{F-CF}_3} = 14.6$ Hz, F).

General procedure for the synthesis of enamine-amine mixtures

Chiral secondary amine (1 equiv.) and dried solvent (3 mL) were placed in a glass pressure-vessel. The solution was cooled to -78 °C and the pressure was lowered. Next, the excess of hexafluoropropene (HFP) was transferred to the glass tube and the vessel was sealed. The cooling bath was removed and the mixture was brought to the room temperature with stirring overnight. After (approximately) 16–24 hours the crude mixture was used for the next step.

Note: The choice of solvent has depended on a solubility of amines. The elevated temperature of reaction (reflux) has not affected the yield.

(*R,E*)-1,2,3,3,3-Pentafluoro-*N*-methyl-*N*-(1-phenylethyl)prop-1-en-1-amine 2a, 1,2,3,3,3-Pentafluoro-*N*-methyl-*N*-((*R*)-1-phenylethyl)propan-1-amine 3a. Reaction of HFP (180 mg, 1.2 mmol) and *R*-(+)-*N*, α -dimethylbenzylamine (0.1 mL, 92 mg, 0.68 mmol), according to the general procedure, gave a mixture of the enamine **2a** (yield 24%) and amine **3a** (yield 55%).

2a ¹⁹F NMR δ : -65.79 (3F, dd, $J_{\text{CF}_3-\text{F}} = 23.1$ Hz, $J_{\text{CF}_3-\text{F}} = 13.6$ Hz, CF₃), -114.48 (1F, dq, $J_{\text{F-F}} = 118.8$ Hz, $J_{\text{F-CF}_3} = 22.6$ Hz, F), -193.51 (1F, dq, $J_{\text{F-F}} = 118.7$ Hz, $J_{\text{F-CF}_3} = 13.5$ Hz, F).

3a ¹⁹F NMR δ : -74.50 to -74.69 (3F, m, CF₃), -81.48 to -87.53 (1F, m, F), -87.72 to -93.33 (1F, m, F), -207.82 to -208.26 (1F, m, F).

(2*S,E*)-(2-Benzhydryl)-1-(perfluoroprop-1-enyl)pyrrolidine 2b, (2*S*)-2-Benzhydryl-1-(1,2,3,3,3-pentafluoropropyl)pyrrolidine 3b. Reaction of HFP (126 mg, 0.84 mmol) and 2(*S*)-2(diphenylmethyl)-pyrrolidine (100 mg, 0.42 mmol) according to the general procedure gave a mixture of enamine **2b** (yield 2%) and amine **3b** (yield 63%).

2b ¹⁹F NMR δ : -65.01 (3F, dd, $J_{\text{CF}_3-\text{F}} = 22.8$ Hz, $J_{\text{CF}_3-\text{F}} = 14.4$ Hz, CF₃), -116.85 (1F, dq, $J_{\text{F-F}} = 116.4$ Hz, $J_{\text{F-CF}_3} = 22.8$ Hz, F), -198.85 (1F, dq, $J_{\text{F-F}} = 118.1$ Hz, $J_{\text{F-CF}_3} = 13.4$ Hz, F).

3b ¹⁹F NMR δ : -74.83 to -74.96 (3F, m, CF₃), -85.57 to -86.65 (1F, m, F), -90.42 to -91.51 (1F, m, F), -205.53 to -206.01 (1F, m, F).

General procedure for the synthesis of amide

Method A. The crude reaction mixture (enamines **1a**, **b**) obtained in the first step was readily hydrolyzed by addition of water (2 equiv.). Next, the reaction mixture was dried (Na₂SO₄), filtered and evaporated under reduced pressure. The product was purified on silica gel (AcOEt-hexane) to give an appropriated amide.

Method B. The crude reaction enamine-amine mixtures **2a-3a** and **2b-3b** obtained in the first step were readily hydrolyzed by addition of boron trifluoride diethyl etherate (2 equiv.) and water (2 equiv.). Next, the reaction mixture was dried (Na₂SO₄), filtered and evaporated under reduced pressure. The product was purified on silica gel (AcOEt-hexane) to give an appropriated amide.

3,3,3-Trifluoro-*N*-methyl-*N*-((*R*)-1-phenylethyl)propanamide 4a. An enamine **2a** was readily hydrolyzed by addition of water (2 equiv.). The product was purified on a silica gel (10% AcOEt-hexane) to give an amide **4a** (two rotamers with 71/29 ratio), as an oil (96 mg, 57%).



$[\alpha]_{\text{D}}^{25} + 172^\circ$ (c 0.4, CHCl_3).

IR (KBr/film) ν 3090, 3064, 3032, 2981, 2945, 2882, 1659, 1605, 1586, 1450, 1429, 1406, 1268, 1137, 1114, 1092, 922, 854, 784, 701, 644.

Rotamer *cis* **4a**: ^1H NMR δ : 1.42 (3H, d, $J = 7.1$ Hz, CH_3), 2.60 (3H, s, NCH_3), 3.18 (2H, q, $J_{\text{H}-\text{CF}_3} = 10.1$ Hz, CH_2), 5.99 (1H, q, $J = 7.1$ Hz, $\text{CH}(\text{CH}_3)$), 7.12–7.33 (m, 5H, Ph); ^{13}C NMR δ : 15.43 (CH_3), 30.08 (NCH_3), 38.64 (q, $J = 28.9$ Hz, CH_2), 50.80 (CH), 124.36 (q, $J = 276.6$ Hz, CF_3), 127.31 (Ph), 127.63 (Ph), 128.65 (Ph), 139.76 (Ph), 163.41 (q, $J = 3.1$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –62.90 (3F, t, $J_{\text{CF}_3-\text{H}} = 10.0$ Hz, CF_3).

Rotamer *trans* **4a**: ^1H NMR δ : 1.56 (3H, d, $J = 6.9$ Hz, CH_3), 2.65 (3H, s, NCH_3), 3.26 (2H, q, $J_{\text{H}-\text{CF}_3} = 10.0$ Hz, CH_2), 4.95 (1H, q, $J = 6.8$ Hz, $\text{CH}(\text{CH}_3)$), 7.12–7.33 (5H, m, Ph); ^{13}C NMR δ : 17.89 (CH_3), 28.41 (NCH_3), 38.24 (q, $J = 28.8$ Hz, CH_2), 55.68 (CH), 124.30 (q, $J = 276.7$ Hz, CF_3), 126.33 (Ph), 127.96 (Ph), 129.01 (Ph), 139.38 (Ph), 163.33 (q, $J = 2.8$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –62.78 (3F, t, $J_{\text{CF}_3-\text{H}} = 10.0$ Hz, CF_3).

MS m/z (rel. int.) 245 $[\text{M}]^+$ (50), 230 $[\text{M}-15]^+$ (20), 134 (5), 120 (100), 111 (25), 77 (50), 69 (5).

1-((S)-2-Benzhydrylpyrrolidin-1-yl)-3,3,3-trifluoropropan-1-one 4b. An enamine **1b** was readily hydrolyzed by addition of water (2 equiv.). The product was purified on a silica gel (20% AcOEt–hexane) to give an amide **4b** (two rotamers with 53/47 ratio) as an oil (55 mg, 38%).

$[\alpha]_{\text{D}}^{25} + 2.3^\circ$ (c 2.8, CHCl_3).

IR (KBr/film) ν 3086, 3061, 3027, 2980, 2889, 1652, 1601, 1582, 1494, 1449, 1440, 1350, 1267, 1252, 1218, 1189, 1113, 1032, 919, 851, 754, 701.

Rotamer *cis* **4b**: ^1H NMR δ : 1.65–1.80 (1H, m, CHH), 1.87–2.10 (3H, m, CHH), 2.44 (2H, q, $J = 10.2$ Hz, CH_2CF_3), 3.34 (2H, dt, $J = 9.9, 8.1$ Hz, CHHN), 4.43 (1H, dt, $J = 10.5, 5.9$ Hz, NCH), 4.63 (1H, d, $J = 5.5$ Hz, $\text{CH}(\text{Ph})_2$), 6.97–7.47 (m, 10H, Ph); ^{13}C NMR δ : 23.57 (CH_2), 27.20 (CH_2CH), 39.80 (q, $J = 28.8$ Hz, CH_2CF_3), 45.04 (CH_2N), 51.66 ($\text{CH}(\text{Ph})_2$), 60.25 (CH), 124.16 (q, $J = 276.9$ Hz, CF_3), 126.27 (Ph), 127.42 (Ph), 128.06 (Ph), 128.72 (Ph), 128.89 (Ph), 140.18 (Ph), 141.50 (Ph), 161.84 (q, $J = 3.3$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –62.74 (3F, t, $J_{\text{CF}_3-\text{H}} = 10.2$ Hz).

Rotamer *trans* **4b**: ^1H NMR δ : 1.65–1.80 (1H, m, CHH), 1.87–2.10 (3H, m, CHH), 2.49 (2H, q, $J = 10.3$ Hz, CH_2CF_3), 3.17 (1H, dt, $J = 9.9, 4.2$ Hz, NCH), 3.56 (1H, dt, $J = 12.6, 6.6$ Hz, CHHN), 3.87 (1H, dt, $J = 12.4, 8.1$ Hz, CHHN), 4.63 (1H, d, $J = 5.5$ Hz, $\text{CH}(\text{Ph})_2$), 6.97–7.47 (m, 10H, Ph); ^{13}C NMR δ : 21.08 (CH_2), 30.56 (CH_2CH), 37.63 (q, $J = 28.6$ Hz, CH_2CF_3), 47.58 (CH_2N), 53.94 ($\text{CH}(\text{Ph})_2$), 62.99 (CH), 123.98 (q, $J = 276.8$ Hz, CF_3), 126.77 (Ph), 127.17 (Ph), 128.31 (Ph), 128.84 (Ph), 129.61 (Ph), 141.08 (Ph), 141.76 (Ph), 162.49 (q, $J = 3.3$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –63.30 (3F, t, $J_{\text{CF}_3-\text{H}} = 10.3$ Hz).

MS m/z (rel. int.) 347 $[\text{M}]^+$ (3), 264 (3), 180 (65), 165 (27), 70 (100).

(S/R)-2,3,3,3-Tetrafluoro-N-methyl-N-((R)-1-phenylethyl)propanamide 5a. The enamine–amine mixture **2a–3a** was readily hydrolyzed by addition of water (2 equiv.) and boron trifluoride diethyl etherate (2 equiv.) giving compound **5a** (two diastereoisomers with 51/49 ratio). The product was purified on a silica gel (10% AcOEt–hexane) to give an amide (*R,S*) **5a** as an oil

(72 mg, 40%) and an amide (*R,R*) **5a** as a crystalline solid (68 mg, 38%), each diastereoisomer as the two rotamers with a 77/23 ratio.

(R,S) 5a. $[\alpha]_{\text{D}}^{25} + 118^\circ$ (c 0.2, CHCl_3).

IR (KBr/film) ν 3090, 3064, 3032, 2982, 2946, 2883, 1667, 1605, 1585, 1496, 1450, 1420, 1348, 1287, 1266, 1189, 1153, 1128, 1086, 1029, 947, 925, 870, 844, 784, 754, 699, 659.

Rotamer *cis* (*R,S*) **5a**: ^1H NMR δ : 1.55 (3H, d, $J = 7.1$ Hz, CH_3), 2.74 (3H, d, $^5J_{\text{CH}_3-\text{F}} = 2.3$ Hz, NCH_3), 5.42 (1H, dq, $J_{\text{H}-\text{F}} = 46.4$ Hz, $J_{\text{H}-\text{CF}_3} = 6.2$ Hz, CHF), 6.02 (1H, q, $J = 7.0$ Hz, $\text{CH}(\text{CH}_3)$), 7.20–7.44 (m, 5H, Ph), ^{13}C NMR δ : 15.13 (CH_3), 29.25 (NCH_3), 52.13 (CH), 84.54 (dq, $J = 198.7, 34.1$ Hz, CHF), 121.30 (dq, $J = 282.4, J = 26.2$ Hz, CF_3), 127.49 (Ph), 127.99 (Ph), 128.84 (Ph), 138.95 (Ph), 161.08 (d, $J = 19.4$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –75.60 (3F, dd, $J_{\text{CF}_3-\text{F}} = 12.9$ Hz, $J_{\text{CF}_3-\text{H}} = 6.2$ Hz, CF_3), –199.52 (1F, ddq, $J_{\text{F}-\text{H}} = 46.4$ Hz, $J_{\text{F}-\text{CF}_3} = 12.8$ Hz, $^5J_{\text{F}-\text{CH}_3} = 1.9$ Hz, F).

Rotamer *trans* (*R,S*) **5a**: ^1H NMR δ : 1.67 (3H, d, $J = 6.8$ Hz, CH_3), 2.71 (3H, d, $^5J_{\text{CH}_3-\text{F}} = 0.9$ Hz, NCH_3), 5.24 (1H, q, $J = 6.8$ Hz, $\text{CH}(\text{CH}_3)$), 5.53 (1H, dq, $J_{\text{H}-\text{F}} = 46.4$ Hz, $J_{\text{H}-\text{CF}_3} = 6.2$ Hz, CHF), 7.20–7.44 (m, 5H, Ph), ^{13}C NMR δ : 17.68 (CH_3), 29.84 (NCH_3), 54.64 (CH), 85.06 (dq, $J = 199.3, 34.0$ Hz, CHF), 121.18 (dq, $J = 282.5$ Hz, $J = 26.9$ Hz, CF_3), 126.74 (Ph), 128.18 (Ph), 129.05 (Ph), 138.65 (Ph), 160.96 (d, $J = 18.8$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –75.46 (3F, dd, $J_{\text{CF}_3-\text{F}} = 13.0$ Hz, $J_{\text{CF}_3-\text{H}} = 6.2$ Hz, CF_3), –197.58 (1F, ddq, $J_{\text{F}-\text{H}} = 46.2$ Hz, $J_{\text{F}-\text{CF}_3} = 13.0$ Hz, $^5J_{\text{F}-\text{CH}_3} = 0.7$ Hz, F).

(R,R) 5a. $[\alpha]_{\text{D}}^{25} + 182^\circ$ (c 0.3, CHCl_3).

IR (KBr/film) ν 3088, 3064, 3033, 3000, 2957, 1655, 1606, 1496, 1457, 1425, 1352, 1267, 1189, 1158, 1130, 1089, 1049, 1028, 926, 870, 839, 784, 749, 699, 671, 658.

Rotamer *cis* (*R,R*) **5a**: ^1H NMR δ : 1.54 (3H, d, $J = 7.1$ Hz, CH_3), 2.76 (3H, d, $^5J_{\text{CH}_3-\text{F}} = 2.6$ Hz, NCH_3), 5.43 (1H, dq, $J_{\text{H}-\text{F}} = 46.4$ Hz, $J_{\text{H}-\text{CF}_3} = 6.1$ Hz, CHF), 6.05 (1H, q, $J = 7.1$ Hz, $\text{CH}(\text{CH}_3)$), 7.12–7.56 (m, 5H, Ph), ^{13}C NMR δ : 15.18 (CH_3), 29.06 (NCH_3), 52.02 (CH), 84.63 (dq, $J = 198.9, 34.0$ Hz, CHF), 121.30 (dq, $J = 282.3, J = 26.3$ Hz, CF_3), 127.40 (Ph), 127.92 (Ph), 128.79 (Ph), 138.95 (Ph), 161.09 (d, $J = 19.1$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –75.58 (3F, dd, $J_{\text{CF}_3-\text{F}} = 12.8$ Hz, $J_{\text{CF}_3-\text{H}} = 6.2$ Hz, CF_3), –199.15 (1F, ddq, $J_{\text{F}-\text{H}} = 46.7$ Hz, $J_{\text{F}-\text{CF}_3} = 12.8$ Hz, $^5J_{\text{F}-\text{CH}_3} = 2.5$ Hz, F).

Rotamer *trans* (*R,R*) **5a**: ^1H NMR δ : 1.66 (3H, d, $J = 6.8$ Hz, CH_3), 2.72 (3H, d, $^5J_{\text{CH}_3-\text{F}} = 1.0$ Hz, NCH_3), 5.30 (1H, q, $J = 6.8$ Hz, $\text{CH}(\text{CH}_3)$), 5.52 (1H, dq, $J_{\text{H}-\text{F}} = 46.5$ Hz, $J_{\text{H}-\text{CF}_3} = 6.1$ Hz, CHF), 7.12–7.56 (m, 5H, Ph), ^{13}C NMR δ : 17.54 (CH_3), 29.12 (NCH_3), 54.48 (CH), 85.89 (dq, $J = 201.4, 34.0$ Hz, CHF), 121.18 (dq, $J = 283.9$ Hz, $J = 26.8$ Hz, CF_3), 126.74 (Ph), 128.07 (Ph), 128.95 (Ph), 138.83 (Ph), 160.98 (d, $J = 18.0$ Hz, $\text{C}=\text{O}$); ^{19}F NMR δ : –75.48 (3F, dd, $J_{\text{CF}_3-\text{F}} = 12.7$ Hz, $J_{\text{CF}_3-\text{H}} = 6.4$ Hz, CF_3), –196.08 (1F, ddq, $J_{\text{F}-\text{H}} = 46.4$ Hz, $J_{\text{F}-\text{CF}_3} = 12.6$ Hz, $^5J_{\text{F}-\text{CH}_3} = 1.0$ Hz, F).

MS m/z (rel. int.) 263 $[\text{M}]^+$ (35), 248 $[\text{M}-15]^+$ (15), 186 (5), 162 (15), 142 (50), 129 (5), 105 (100), 101 (18), 77 (40), 69 (5).

1-((S)-2-Benzhydrylpyrrolidin-1-yl)-(R/S)-2,3,3,3-tetrafluoropropan-1-one 5b. The enamine–amine mixture **2b–3b** were readily hydrolyzed by addition of water (2 equiv.) and boron trifluoride diethyl etherate (2 equiv.). The crude product was purified on a silica gel (20% AcOEt–hexane) giving mixture of two diastereoisomers with 53/47 ratio of **5b** as an oil (152 mg, 98%), each diastereoisomer as a mixture of two rotamers with a 77/23 ratio.



IR (KBr/film) ν 3086, 3061, 3028, 2983, 2890, 1667, 1600, 1494, 1450, 1285, 1192, 1142, 869, 701.

Rotamer *cis* **5b** (major diastereoisomer) ^1H NMR δ : 1.57–1.70 (2H, m, CH_2), 1.87–2.10 (2H, m, CH_2), 3.08–3.28 (2H, m, CH_2N), 3.74–3.82 (1H, m, NCH), 4.37 (1H, d, $J = 6.9$ Hz, $\text{CH}(\text{Ph})_2$), 5.11 (1H, dq, $J_{\text{H-F}} = 46.5$ Hz, $J_{\text{H-CF}_3} = 6.3$ Hz, CHF), 6.95–7.45 (m, 10H, Ph), ^{13}C NMR δ : 25.72 (CH_2), 39.19 (CH_2), 45.66 (CH_2N), 51.22 ($\text{CH}(\text{Ph})_2$), 61.10 (CH), 85.57 (dq, $J = 203.2$ Hz, $J = 33.6$ Hz, CHF), 120.52 (dq, $J = 282.1$ Hz, $J = 25.5$ Hz, CF_3), 128.41 (Ph), 128.84 (Ph), 129.56 (Ph), 144.72 (Ph), 159.53 (d, $J = 20.5$ Hz, $\text{C}=\text{O}$), ^{19}F NMR δ : –75.72 (3F, dd, $J_{\text{CF}_3-\text{F}} = 13.3$ Hz, $J_{\text{CF}_3-\text{H}} = 6.6$ Hz, CF_3), –200.68 (1F, dq, $J_{\text{F-H}} = 47.5$ Hz, $J_{\text{F-CF}_3} = 13.4$ Hz, F).

Rotamer *trans* **5b** (major diastereoisomer) ^1H NMR δ : 1.79–1.92 (2H, m, CH_2), 1.87–2.10 (2H, m, CH_2), 3.08–3.28 (2H, m, CH_2N), 3.74–3.82 (1H, m, NCH), 4.46 (1H, d, $J = 6.3$ Hz, $\text{CH}(\text{Ph})_2$), 5.05 (1H, dq, $J_{\text{H-F}} = 46.6$ Hz, $J_{\text{H-CF}_3} = 6.6$ Hz, CHF), 6.95–7.45 (m, 10H, Ph), ^{13}C NMR δ : 26.66 (CH_2), 39.34 (CH_2), 46.56 (CH_2N), 53.40 ($\text{CH}(\text{Ph})_2$), 61.10 (CH), 85.52 (dq, $J = 199.7$ Hz, $J = 33.5$ Hz, CHF), 120.94 (dq, $J = 282.8$ Hz, $J = 26.3$ Hz, CF_3), 128.45 (Ph), 129.24 (Ph), 129.30 (Ph), 145.52 (Ph), 161.18 (d, $J = 22.4$ Hz, $\text{C}=\text{O}$), ^{19}F NMR δ : –76.64 (3F, dd, $J_{\text{CF}_3-\text{F}} = 11.0$ Hz, $J_{\text{CF}_3-\text{H}} = 6.5$ Hz, CF_3), –202.67 (1F, dq, $J_{\text{F-H}} = 46.4$ Hz, $J_{\text{F-CF}_3} = 11.1$ Hz, F).

Rotamer *cis* **5b** (minor diastereoisomer) ^1H NMR δ : 1.22–1.40 (2H, m, CH_2), 1.79–1.92 (2H, m, CH_2), 3.40–3.64 (2H, m, CH_2N), 4.08–4.17 (1H, m, NCH), 4.57 (1H, d, $J = 5.5$ Hz, $\text{CH}(\text{Ph})_2$), 4.92 (1H, dq, $J_{\text{H-F}} = 45.4$ Hz, $J_{\text{H-CF}_3} = 5.5$ Hz, CHF), 6.95–7.45 (m, 10H, Ph), ^{13}C NMR δ : 26.90 (CH_2), 35.06 (CH_2), 46.58 (CH_2N), 52.25 ($\text{CH}(\text{Ph})_2$), 61.36 (CH), 85.64 (dq, $J = 201.2$ Hz, $J = 33.7$ Hz, CHF), 121.03 (dq, $J = 282.2$ Hz, $J = 26.0$ Hz, CF_3), 128.04 (Ph), 128.63 (Ph), 129.57 (Ph), 140.89 (Ph), 159.49 (d, $J = 19.3$ Hz, $\text{C}=\text{O}$), ^{19}F NMR δ : –75.74 (3F, dd, $J_{\text{CF}_3-\text{F}} = 12.6$ Hz, $J_{\text{CF}_3-\text{H}} = 5.9$ Hz, CF_3), –200.59 (1F, dq, $J_{\text{F-H}} = 45.7$ Hz, $J_{\text{F-CF}_3} = 12.4$ Hz, F).

Rotamer *trans* **5b** (minor diastereoisomer) ^1H NMR δ : 1.22–1.40 (2H, m, CH_2), 1.41–1.55 (2H, m, CH_2), 3.40–3.64 (2H, m, CH_2N), 4.00–4.08 (1H, m, NCH), 4.62 (1H, d, $J = 5.5$ Hz, $\text{CH}(\text{Ph})_2$), 4.93 (1H, dq, $J_{\text{H-F}} = 44.4$ Hz, $J_{\text{H-CF}_3} = 6.5$ Hz, CHF), 6.95–7.45 (m, 10H, Ph), ^{13}C NMR δ : 27.40 (CH_2), 35.26 (CH_2), 46.28 (CH_2N), 54.25 ($\text{CH}(\text{Ph})_2$), 61.26 (CH), 82.21 (dq, $J = 209.3$ Hz, $J = 33.7$ Hz, CHF), 120.57 (dq, $J = 280.0$ Hz, $J = 24.7$ Hz, CF_3), 128.44 (Ph), 128.93 (Ph), 130.17 (Ph), 140.93 (Ph), 161.20 (d, $J = 21.2$ Hz, $\text{C}=\text{O}$), ^{19}F NMR δ : –76.62 (3F, dd, $J_{\text{CF}_3-\text{F}} = 14.4$ Hz, $J_{\text{CF}_3-\text{H}} = 6.0$ Hz, CF_3), –203.73 (1F, dq, $J_{\text{F-H}} = 45.1$ Hz, $J_{\text{F-CF}_3} = 14.4$ Hz, F).

MS m/z (rel. int.) 264 (5), 198 (100), 165 (20), 129 (3), 101 (5), 77 (2), 70 (50).

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