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Author(s): Kiss, Loránd; Nonn, Melinda; Sillanpää, Reijo; Fustero, Santos; Fülöp, Ferenc

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Efficient regio- and stereoselective access to novel fluorinated β-aminocyclohexanecarboxylates

Loránd Kiss¹, Melinda Nonn², Reijo Sillanpää³, Santos Fustero⁴ and Ferenc Fülöp*¹,²

Abstract
A regio- and stereoselective method has been developed for the synthesis of novel fluorinated 2-aminocyclohexanecarboxylic acid derivatives with the fluorine attached to position 4 of the ring. The synthesis starts from either cis- or trans-β-aminocyclohex-4-enecarboxylic acids and involves regio- and stereoselective transformation of the ring C–C double bond through iodooxazine formation and hydroxylation, followed by hydroxy–fluorine or oxo–fluorine exchange.

Introduction
Fluorine chemistry is an expanding area of research that has generated increasing interest in pharmaceutical and medicinal chemistry in recent years because of its considerable impact in drug discovery. There is currently extensive research activity in synthetic chemistry for the preparation of various biologically active fluorinated products [1-10].

Of special interest among such materials are the fluorinated amino acids, which in most cases exhibit higher bioactivities than the nonfluorinated counterparts. The fluorinated α- or acyclic β-amino acids have acquired significance as antibacterial or antifungal agents, enzyme inhibitors or as antitumoral compounds. Introduction of a fluorinated amino acid into a peptide may generate specific protein–ligand or protein–protein interactions, thereby determining thermal or metabolic stabilities, which is of great importance in peptide-based drug research [11-35]. These changes in properties may be more appreciable in the case of peptide oligomers formed from conformationally restricted fluorinated amino acids. Although cyclic β-amino acids are of major interest in pharmaceutical
chemistry and in peptide research [36-60], only a relatively small number of fluorinated derivatives of this class of compounds have been synthesized so far [61-70].

**Results and Discussion**

We recently developed a synthetic method for the regio- and stereoselective introduction of a fluorine atom onto the skeleton of a β-aminocyclohexanecarboxylic acid. The synthesis starts from the Boc-protected 2-aminocyclohex-4-enecarboxylic acid or 2-aminocyclohex-3-enecarboxylic acid and involves ring C–C bond transformation by regio- and stereoselective hydroxylation via iodosylactonization, followed by hydroxy–fluorine exchange. This protocol was applied to synthesize fluorinated β-aminocyclohexane scaffolds with the fluorine atom on either position 3 or 5 of the ring. Whereas the procedure is a convenient economical route to fluorinated cyclohexane or cyclohexene β-amino acids, it did not allow extension to the synthesis of similar derivatives with the fluorine atom on position 4.

During our work performed to fill this gap, we have developed a synthetic procedure for gaining access to fluorinated β-aminocyclohexanecarboxylic acids.

This synthesis starts from ethyl cis-2-aminocyclohex-4-enecarboxylate 1 [57] and follows two different strategies. One is based on regio- and stereoselective hydroxylation via iodo-oxazine formation, followed by fluorination, while the other includes stereoselective epoxidation and regioselective oxirane opening, followed by hydroxy–fluorine exchange. In the former protocol, amino ester 1 is treated with KI/I₂ in H₂O/CH₂Cl₂, which affords iodooxazinone derivative 2 stereo- and regioslectively (Scheme 1, Figure 1). Next, compound 2 is transformed to 3 by amide N-Boc protection with Boc₂O and 4-dimethylaminopyridine (DMAP) in THF. Removal of the iodine from the cyclohexane skeleton in 3 is accomplished under reductive conditions. On treatment with n-Bu₃SnH in the presence of a catalytic amount of azobisisobutyronitrile (AIBN) in dichloromethane under reflux, 3 undergoes deiodination to give ester 4 in 70% yield. Oxazinone 4 is then subjected to heterocycle ring opening with NaOEt in EtOH at 0 °C to furnish all-cis hydroxylated amino ester 5 with the hydroxy group on position 4 of the skeleton (Scheme 1, Figure 2).

During our work performed to fill this gap, we have developed a synthetic procedure for gaining access to fluorinated β-aminocyclohexanecarboxylic acids.

| Scheme 1: Synthesis of all-cis ethyl 4-hydroxylated β-aminocyclohexanecarboxylate 5. | Figure 1: ORTEP diagram of iodooxazinone 2. Thermal ellipsoids have been drawn at the 20 % probability level. |

1. **CO₂Et**
   - NH:Boc
   - KI, I₂, CH₂Cl₂
   - H₂O, rt, 18 h
   - 76%

2. **CO₂Et**
   - Iₗ
   - Boc₂O, DMAP
   - THF, 0 °C to rt
   - 22 h
   - 79%

3. **CO₂Et**
   - NaBH₄, EtOH
   - 70 °C, 2 h
   - 72%

4. **CO₂Et**
   - NaOEt, EtOH
   - 0 °C, 1 h
   - then H₂O
   - 61%
hydroxy function on position 4 (for analogous transformations, see reference [60]).

Hydroxylated amino ester 5 was next further used as a key compound for the synthesis of fluorinated target materials. A fluorine atom was introduced by hydroxy–fluorine exchange with bis(dimethoxyethylaminosulfur trifluoride) (Deoxo-Fluor®) reagent. The reaction was carried out under different experimental conditions, with variation of the temperature (−40 °C, 0 °C or 20 °C) and the solvent (toluene, CH₂Cl₂ or THF). Finally, it was found that hydroxylated amino ester 5 underwent inversion on reaction with a 50% Deoxo-Fluor toluene solution in CH₂Cl₂ at 0 °C [68,69] to give monofluorinated cyclohexane amino ester 7 in 32% yield (Scheme 2). This rather modest yield is attributed to the relatively large amount of elimination materials (40% overall). In continuation, the geminal difluorinated β-aminocyclohexanecarboxylic acid derivative with the fluorine atoms on position 4 was efficiently synthesized. Oxidation of the hydroxy group of amino ester 5 with pyridinium chlorochromate (PCC) in CH₂Cl₂ yielded the corresponding oxo-group-containing amino ester 8, which was then converted with Deoxo-Fluor in CH₂Cl₂ at 0 °C to the corresponding geminal difluoro amino ester 9 in good yield (Scheme 2).

The synthetic route presented above could be extended to the preparation of other 4-fluorinated cyclohexane amino acid derivatives, stereoisomers of 7 or 9. Ethyl trans-2-aminocyclohex-4-enecarboxylate 10 [57] was analogously transformed to its cis counterpart through regio- and stereoselective iodo-oxazine formation with KI/I₂ to give compound 11 (Scheme 3). N-Protection of 11, followed by reductive deiodination, proceeded via 12 (Figure 3) to afford ester 13. Opening of the heterocyclic ring with NaOEt in EtOH at 0 °C furnished 4-hydroxylated amino ester 14, a stereoisomer of 5 (Scheme 3, Figure 4).
Figure 3: ORTEP diagram of iodooxazinone derivative 12. Thermal ellipsoids have been drawn at the 20% probability level.

Figure 4: ORTEP diagram of hydroxylated amino ester 14. The water molecule oxygen atom O4 is situated on the twofold axis with a population parameter of 0.6. Thermal ellipsoids have been drawn at the 20% probability level.

Conclusion

In conclusion, a simple and convenient procedure has been developed for the introduction of one or two fluorine atoms onto the skeleton of either cis- or trans-β-aminocyclohexanecarboxylates. The synthetic concept involves regio- and stereoselective hydroxylation via iodooxazine formation, followed by hydroxy-fluorine or oxo-fluorine exchange.

Supporting Information

Supporting Information File 1
Experimental procedures and characterization of compounds.
[http://www.beilstein-journals.org/bjoc/content/supplementary/1860-5397-9-130-S1.pdf]

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References


55. Mansawat, W.; Vilaivan, C.; Balázs, Á.; Aitken, D. J.; Vilaivan, T. 
56. Berlicki, Ł.; Pitsl, L.; Weber, E.; Mándity, I. M.; Cabrele, C.; 
Martinek, T. A.; Fülop, F.; Reiser, O. Angew. Chem., Int. Ed. 2012, 51, 
2208. doi:10.1002/anie.201107702
2012, 8, 100. doi:10.3762/bjoc.8.10
doi:10.1016/j.tet.2010.03.030
60. Kiss, L.; Forró, É.; Martinek, T. A.; Bernáth, G.; De Kimpe, N.; Fülop, F. 
Tetrahedron 2008, 64, 5036. doi:10.1016/j.tet.2008.03.068
doi:10.1002/ejoc.201100032
Soloshonok, V.; Sorochinsky, A. Synthesis 2011, 3045. 
64. Fustero, S.; Sánchez-Roselló, M.; Aceña, J. L.; Fernández, B.; 
3414. doi:10.1021/jo900296d
2006, 8, 4633. doi:10.1021/ol061892w
doi:10.1016/S0960-894X(02)00958-7
4993. doi:10.1002/ejoc.201100583
69. Kiss, L.; Forró, É.; Fustero, S.; Fülop, F. Org. Biomol. Chem. 2011, 9, 
6528. doi:10.1039/c1ob05648d
70. Nonn, M.; Kiss, L.; Hänninen, M. M.; Sillanpää, R.; Fülop, F. 

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