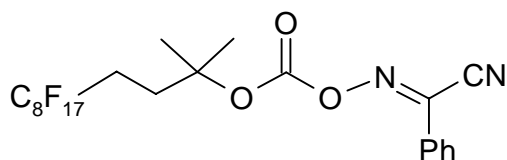
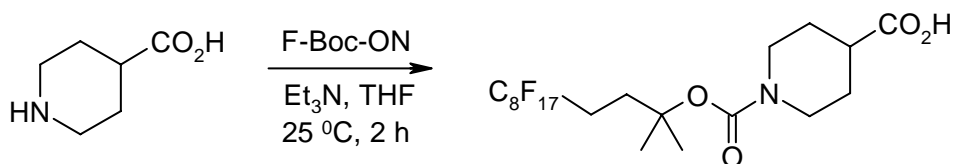


F017003

 2-[2-(1H,1H,2H,2H-Perfluorodecyl)
 isopropoxycarbonyloxyimino]-2-phenylacetonitrile

Chemical Formula:	C ₂₂ H ₁₅ F ₁₇ N ₂ O ₃
Formula Weight:	678.34
CAS Number:	356056-15-0
Appearance:	White, free-flowing solid
Soluble in:	Dichloromethane, chloroform, THF, ether, toluene, and many other organic solvents
Stability:	Stable in air

Why Fluorous? 2-(tert-Butoxycarbonyloxyimino)-2-phenylacetonitrile (Boc-ON) has been widely used for protection of amino groups in peptide synthesis and multistep organic synthesis of small molecules.^{1,2} F17-Boc-ON is a fluorous equivalent which has comparable utility to the normal Boc-ON. Protection of the amino group with F-Boc-ON and deprotection are achieved under standard reaction conditions, with the advantage that products containing the F-Boc group can easily be separated from organic reagents, reactants or products by performing a quick fluorous solid phase extraction (F-SPE) over FluoroFlash[®] silica gel.³

TYPICAL PROTECTION:¹

Materials Needed
Quantity

Amine	29 mg, 1.2 mmol
F17-Boc-ON	88 mg, 1.3 mmol
Et ₃ N	20 mg, 0.2 mmol
THF (dry)	10 mL
FluoroFlash [®] SPE cartridge	1 x 5 g
MeOH:H ₂ O, 80:20	20 mL
MeOH	20 mL

Stepwise Procedure:

- 1) Mix F-Boc-ON, amine, and Et₃N in THF;
- 2) Stir 2 h at room temperature;
- 3) Concentrate reaction mixture to < 0.3 mL;
- 4) Charge to F-SPE cartridge;
- 5) Elute cartridge with MeOH:H₂O ;
- 6) Elute cartridge with MeOH and concentrate eluent to give 93 mg, 96% yield of product.

TYPICAL DEPROTECTION:¹
Materials Needed
Quantity

F-Boc Protected Amine	0.05 mmol
TFA:CH ₂ Cl ₂ , 1:1	1.0 mL
FluoroFlash [®] SPE cartridge	1 x 2 g
MeOH:H ₂ O, 80:20	8 mL

Stepwise Procedure:

- 1) Add TFA:CH₂Cl₂ to F-Boc protected amine;
- 2) Stir 2.5 h at room temperature;
- 3) Concentrate reaction mixture to < 0.15 mL;
- 4) Charge to F-SPE cartridge;
- 5) Elute cartridge with MeOH:H₂O and concentrate eluent to give the deprotected product. Typical yields are 90-100%.

Insider Tips:

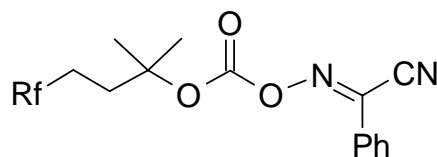
- The F-Boc protected amine is in the second SPE fraction of MeOH, while the deprotected product is in the first fraction of MeOH:H₂O.
- Loading solvents and volumes affect the reliability of the F-SPE in predictable ways. Be sure to read the application note on "Fluorous Solid Phase Extractions" if you are new to this technique.⁴
- The spent F-SPE cartridge can be regenerated by washing with THF, then reconditioning according to the above application note.
- Another method for deprotection that can be used is heating the F-Boc amine with HCl:MeOH (1:3) for 16 h at 65 °C.

ADDITIONAL OPTIONS:

F17-Boc has appropriate fluorine content for tagging of diverse organic molecules in combination with F-SPE. The F-Boc-ON homologs are useful for fluorous mixture synthesis (FMS).⁵

Additional F-BOC-ON Homologs
 Available:

Rf	Catalog No.
C ₃ F ₇	F007003
C ₄ F ₉	F009003
C ₆ F ₁₃	F013003
C ₁₀ F ₂₁	F021003



REFERENCES:

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- 3) Curran, D. P. *Synlett* **2001**, 1488. Curran, D. P. 'Separations with Fluorous Silica Gel and Related Materials', in *Handbook of Fluorous Chemistry*, Wiley-VCH: Weinheim, 2004; pp 101-127.
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- 5) Zhang, W. *Arkivoc* **2004** (i), 101-109.
- 6) Zhang, W. Tempest, P. "Highly Efficient Microwave-Assisted Fluorous Ugi and Post-Condensation Reactions for Benzimidazoles and Quinoxalinones" *Tetrahedron Lett.* **2004**, 45, 6757-6760.
- 7) Sreeman K. Mamidyala and Steven M. Firestine "Fluorous synthesis of minor groove binding agents related to distamycin" *Tetrahedron Lett.* **2006**, 47, 7431-7434.