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A concise and sequential synthesis of the nitroimidazooxazole based drug, Delamanid and related compounds[†]

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A concise, protection-group free and sequential route has been developed for the synthesis of the nitroimidazole based FDA-approved multi-drug resistant anti-tuberculosis drug, Delamanid and anti-leishmanial lead candidate VL-2098. The synthesis required chiral epoxides (**11** and **17**) as key intermediates. The chiral epoxide **11** was synthesised by sequential reaction cascades *viz.*, allylation, selective *N*-arylation, Mitsunobu etherification, Sharpless asymmetric dihydroxylation and epoxidation, which do not require any special/dry reaction conditions. The steps involved towards the synthesis of epoxide also worked nicely in gram scales. After the synthesis of epoxide **11**, the synthesis of Delamanid was achieved by reaction with 2-bromo-4-nitroimidazole **12** with an overall yield of 27%. Similarly, anti-leishmanial lead candidate VL-2098 was also synthesized in an overall yield of 36%.

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Imidazoles are present in a wide variety of biologically relevant molecules exhibiting diverse pharmaceutical properties. Specifically, nitroimidazole containing compounds are active therapeutic agents against a wide variety of protozoan, bacterial (anaerobic) and leishmanial infections of humans and animals.1 The notable examples of drugs are fexindazole,2 metronidazole,³ benznidazole,⁴ tinidazole,⁵ etc. Nitroimidazole was also well explored in the area of tuberculosis (TB) drug discovery (the structure of nitroimidazole containing drugs and lead compounds shown in Fig. 1).6-9 CGI-17341 (I, developed by Ciba-Geigy, India) represents one of the very earliest outcomes but was not continued because of mutagenicity.6 Continued efforts to overcome the mutagenic liability led to the discovery of Delamanid (II, OPC-67683, a blockbuster against multi-drug resistant-TB, approved in 2014 by the EU)⁷ and Pretomanid^{8a} (III, PA-824, against multi-drug-resistant-MTB, approved in 2019 by the USFDA).¹⁰ Our involvement in the area of the TB drug discovery program and nitroimidazole chemistry^{8b,c} motivated us to develop a new concise and improved strategy towards the synthesis of Delamanid. The first synthesis of Delamanid was done by Tsubouchi et al. in 2004 (Otsuka Pharmaceutical Ltd. WO2004033463A1) which involved 16 steps (details shown in Fig. S1 of ESI[†]).^{11,12} The synthesis involves protection-deprotection strategy and made the process lengthy. Later on, in 2011, Otsuka had developed another concise route

for Delamanid with improved yield (upper half of Fig. 2, details shown in Fig. S2 of ESI†). This method involved the Sharpless epoxidation of 2-methyl allylalcohol followed by ring opening with 4-bromophenol. Then coupling with 4(4-trifluoromethoxy phenyl)piperidine fragment under palladium catalyzed conditions, which in turn was synthesized in 2–5 steps.¹³ Considering the importance of Delamanid, It could be better if more concise route was developed. In this regard, here we devised a route which involves sequential addition of fragments which not only avoid protection and deprotection but also provides the advantages of avoidance of dry conditions and costly intermediates. The lower half of Fig. 2 represents the strategy of the present method used for the synthesis of Delamanid, which involve the sequential coupling with following cascade *viz.*, allylation, selective *N*-arylation, Mitsunobu ether formation,



Fig. 1 Nitroimidazole containing drugs and leads.

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Fig. 2 Previous and current approaches

Sharpless dihydroxylation, epoxidation, ring opening and cyclization using inexpensive and easily available starting materials.

The present synthesis started with 2-methylallyl chloride as first starting material and its selection is because of following reasons (i) as protecting group, (ii) inexpensive (33\$ per 100 mL, Sigma) in comparison to 2-methylallyl alcohol (766\$ per 100 mL, Sigma) which is used in earlier reported method and (iii) provide double bond functionality to generate epoxide *via* Sharpless dihydroxylation approach, which operates under open atmosphere conditions and avoids the anhydrous and dry environment as required for Sharpless epoxidation, used in the previous reported methods.^{7,11b,12}

Result and discussion

The synthesis started with allylation of 4-iodophenol 2. The 2methylallyl chloride 1 (or 3-chloro-2-methylprop-1-ene) reacted with 4-iodophenol 2 in the presence of potassium carbonate as base in DMF at 60 °C to furnish 1-iodo-4-((2-methylallyl)oxy) benzene 3 in 97% yield (Scheme 1).

The second step involves the C-N bond formation between 1iodo-4-((2-methylallyl)oxy)benzene 3 and 4-hydroxypiperidine 4. Here, our aim was to develop a condition for selective C-N bond formation in the presence of free hydroxyl group. In the literature, copper catalysis has been used for selective C-N bond formation,¹⁴ however, no such attempts were made in the synthesis of Delamanid, which on the other hand was achieved by Pd-catalysis.^{10,12-14} We started with copper catalysis and various attempts were made (details provided in Scheme 2, entries 1-10). The optimization started by using CuI as catalyst in the presence of different ligands as well as solvents. The reaction with CuI (10 mol%), phenanthroline (10 mol%) and Cs_2CO_3 (2 equiv.) gave the desired *N*-arylated product 5 in a yield of 55% without any O-arylated product 5' (entry 1). When the same reaction was performed without any base, it gives 53% yield of 5 (entry 2). In the next attempt, reaction was performed by using bipyridine as ligand which gave 5 in a yield of 60% (entry 3). After that, pyrrolidine was attempted which gave poor yield of 35% of 5 (entry 4). Then, combination of CuI (10 mol%) and L-proline (10 mol%) was tried at room temperature which gave the yield of desired N-arylated product 5 in a yield of 80% (entry 6). The reaction with CuI (10 mol%) and L-proline (10 mol%) at 80 °C furnished the N-arylated product 5 and Oarylated product 5' in a yield of 40% and 20% respectively (entry 7). Then varying the molar ratio of ligand (10 : 20%) at 80 °C did not show any improvement and desired product 5 was obtained in a yield of 48% along with 25% of O-arylated product 5' (entry 8). In further refinement experiments (entries 9-11), different molar ratio of ligand and catalyst was tried, where CuI (20 mol%) and L-proline (40 mL%) gave highest yield of 5 (85%). After optimization, the synthesis of desired N-aryalted product 5 was performed at 100 mmol scale which worked nicely and delivered the required product 5 in a yield of 81%.

The third step involves the ether bond formation between 1-(4-((2-methylallyl)oxy)phenyl)piperidin-4-ol 5 and 4-trifluoromethoxyphenol 7. In the first instance, the ether formation was tried*via*mesylation of hydroxyl group followed bynucleophilic substitution with 4-trifluoromethoxyphenol 7. Themesylation of 1-(4-((2-methylallyl)oxy)phenyl)piperidin-4-ol 5was carried out by treating with mesyl chloride in presence of



Scheme 1 Allylation. Reagent and conditions: (i) compound 1 (100 mmol), compound 2 (200.0 mmol), K_2CO_3 (200.0 mmol), DMF, 60 °C, 97%.

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conditions 5' Catalvst Yield entrv Ligand Base (mol%) Cul (mol%) 1 10 Phenanthroline(10) 55 Cs₂CO₃ 2 10 Phenanthroline(10) 53 3 10 Bipyridine(10) 60 ____ 4 10 Pyrrolidine(10) 35 5 10 L-Proline(10) 80 6 10 L-Proline(10) Cs₂CO₃ 80 70 40(20)^d 10 L-Proline(10) ____ 48 (25)^d 8^c 10 L-Proline(20) 9 20 L-Proline(30) 74 10 20 L-Proline(40) 85 11 40 L-Proline(80) 78

Scheme 2 C–N bond formation. ^aReagents and conditions: compound 3 (0.5 mmol), compound 4 (1 mmol), DMF, 10–24 h; ^bisolated yields; ^creaction performed at 80 °C; ^dyield of *O*-arylated product 5'.



Scheme 3 Nucleophilic substitution for ether formation. Reagent and conditions: (i) compound 5 (4 mmol), mesyl chloride (4.4 mmol), Et_3N (6 mmol), DCM, 0 °C to rt; (ii) compound 6 (0.31 mmol), phenol 7 (0.31 mmol), base (0.62), 2–12 h; ^aisolated yield.

triethylamine which afforded the *O*-mesylated product **6** in the yield of 93%. The coupling of *O*-mestlated product **6** with 4-trifluoromethoxyphenol 7 have been tried by using different

solvents (*N*,*N*-diemthylformamide, toluene) and bases (Cs₂CO₃, K₂CO₃, NaH) combination, but none gave satisfactorily yields of coupled product **8** (Scheme 3, entries 1–5). Then, we switch over to Mitsunobu reaction conditions using triphenyl phosphine (PPh₃) and diethyl azodicarboxylate (DEAD) for ether formation. Again various attempts have been made under different solvents (Scheme 4).¹⁴ By screening different solvents, THF as solvent and 2.0 equivalents of PPh₃ and DEAD gave best result towards the formation of required key intermediate **8** in a yield of 70%. Taking the best conditions, the ether formation was tried in 20 mmol scale which afforded the key product **8** in a yield of 72%. Here, we have provided a new approach for the ether formation. The use of environmental friendly version of Mitsunobu (polymeric or catalytic variants) will be tried during upscaling.¹⁵

The next step involved the generation of enantiopure epoxide which in present case was achieved via Sharpless asymmetric dihydroxylation using AD mix-a and AD mix-B.16 The optimization started with the reaction of 1-(4-((2-methylallyl)oxy)phenyl)-4-(4-trifluoromethoxy) phenoxy) piperidine 8 with AD mix- α at 20 °C which gave diol product 9 in a yield of 84% with an ee (enantiomeric excess) of 82%. Decrease in the temperature conditions from 20 °C to 0 °C improved the enantioselctivity where the reaction at 0 °C gave diol product 9 in a yield of 86% with ee of 94% (HPLC detail provided in ESI[†]). The synthesized diol 9 has shown an optical rotation of $\left[\alpha\right]_{D}^{25}$ -6.65°. In order to confirm the R-configuration of diol 9, the same has also been synthesized by reported method (detail provided in Fig. S3 of ESI[†]), which is in well agreement with our synthesized diol 9.9,17 In the further exploration, when the reaction was attempted with AD mix-β, the diol product was obtained with yield of 89% having ee of 94% (and shown opposite optical rotation $\left[\alpha\right]_{D}^{25}$ +5° and in agreement with literature reported S-configured diol. The present approach provides the opportunity to synthesize both the enantiomers which can be exploited in hit to lead optimization strategy of medicinal chemistry program. As the present strategy required R-isomer, our next steps are focused with R-isomer. The temperature played an important role in this reaction to get the better enantioselctivity. The Sharpless asymmetric dihydroxylation conditions operated under open atmospheric conditions, which provided another advantage of present method over the existing method.



Scheme 4 Mitsunobu reaction for ether formation. ^aReagents and conditions: compound 5 (0.8 mmol), compound 7 (0.8 mmol), PPh₃ (1.14 mmol), DEAD (0.97 mmol), rt, 24 h; ^bisolated yield.

$HO \longrightarrow OCF_3 \xrightarrow{Conditions^a} HO \longrightarrow OCF_3 \xrightarrow{OC} OCF_3$							
entry	Reagents	Temp.	Time	Yield	ee (%)	[α]D	
		°C	h				
1	AD mix-α	20	6.0	84	82		
2	AD mix-α	10	6.0	87	94		
3	AD mix-α	0	6.0	86	94(R)	-6.65°	
4	AD mix-β	0	6.0	89	94(S)	+5.0°	

Scheme 5 Sharpless asymmetric dihydroxylation. ^aReagent and conditions: compound 8 (2.0 mmol), AD mix- α or β (2.8 g, 1.4 g for 1 mmol of alkene), *tert*-butanol : water (1 : 1).

The diol R-(-)-9 was converted into a key intermediate epoxide 11 *via* mesylation and ring formation as per the conditions provided in Scheme 6.

The 2-bromo-4-nitroimidazole **12**, another key fragment and coupling partner was synthesized from 4-nitroimidazole (detail shown in Fig. S4 of ESI[†]). After the successful synthesis of both the key intermediates *viz.*, epoxide **11** and 2-bromo-4-nitroimidazole **12**, the final attempt towards the suitable coupling conditions were made (details given Table S1 of ESI[†]) where DIPEA (act as base as well as solvent) gave the coupled product **13** which subsequent treated with Cs_2CO_3 as base successfully furnished the final construct Delamanid **II** with an overall yield of 63% (Scheme 7).

After finishing this, we further extended the present approach to VL-2098 (**VII**) which is another nitroimidazole based lead reported against visceral leishmaniasis.^{1d} As per the literature, two routes are reported for its synthesis. The first synthesis was reported by Tsubouchi *et al.* (2004) which involved 7 steps (details shown in Fig. S5 of ESI[†]).^{11,18} Later on, in 2017, Pati *et. al.* developed another alternative route with an overall yield of 19% (details shown in Fig. S6 of ESI[†]) (Fig. 3).⁹

To synthesized VL-2098, 2-methylallyl chloride **1** (or 3-chloro-2-methylprop-1-ene) reacted with 4-trifluoromethoxy phenol 7 under basic conditions to furnished 1-((2-methylallyl)oxy)-4-



Scheme 6 Epoxide formation. Reagent and conditions: (i) compound R-(-)-9 (3 mmol), mesyl chloride (3.6 mmol), Et₃N (6 mmol), DCM, 0 °C to rt; (ii) compound **10** (0.31 mmol), DBU (0.62 mmol), rt, 2 h, yield 82%.



Scheme 7 Final coupling. Reagent and conditions: (i) compound 12 (3 mmol), compound 11 (3 mmol), DIPEA, 115 °C, yield 90%; (ii) compound 13 (1 mmol), Cs_2CO_3 (2 mmol), DMF, 50 °C, 2 h, yield 70%.

(trifluoromethoxy)benzene **14** in 97% yield. The next step involved the Sharpless asymmetric dihydroxylation using AD mix- α .¹⁶ The reaction started with the 1-((2-methylallyl)oxy)-4-(trifluoromethoxy)benzene **14** with AD mix- α at 20 °C which gave diol product *R*-(+)-15 with an yield of 88%. The synthesized diol has shown an optical rotation of $[\alpha]_{D}^{25}$ +8.83°. The diol *R*-(+)-15 was converted into a key intermediate epoxide **17** *via*



Fig. 3 Previous and current approaches for VL-2098.



Scheme 8 Synthesis of VL-2098. Reagent and conditions: (i) Compound 7 (28 mmol), K_2CO_3 (56 mmol), DMF, $60-65 \degree C$, 2 h, 97%; (ii); compound 14 (4.3 mmol), AD-mix α (3 mmol), t-BuOH-H₂O (1 : 1), rt, 2 h, 88%; (iii) compound *R*-(+)-15 (3.7 mmol), MsCl (2.2 mmol), Et₃N (4 mmol), EtOAc, <15 °C - rt, 2 h; (iv) DBU (4 mmol), EtOAc, rt, 2 h, 89%; (v) compound 17 (1 mmol), DIPEA,110 °C, 2 h; (vi) Cs₂CO₃ (2 mmol), DMF, 50 °C, 2 h, 48%.

mesylation as per the conditions provided in Scheme 8. The final step involved the coupling of key fragment epoxide 17 with 2-bromo-4-nitroimidazole 12 to get the desired product VL-2098 (VII) with an optical rotation of $[\alpha]_D^{25}$ +7° ($[\alpha]_D^{25}$ +8.445°).⁹ This new approach has an overall yield of 36%.

In summary, we have developed an alternative concise route for the synthesis of nitroimidazole based drug molecule Delamanid and lead candidate VL-2098 with an overall yield of 27% and 36% respectively. The present strategy avoids the use of protecting groups and expensive reagents. The present strategy also provides the opportunity towards the generation of opposite enantiomers and manipulation of functionalities/ groups, which is presently being pursued by our group to generate best-in-class nitroimidazole based TB drug free of hERG liability and better drug-like properties.

Experimental section

General information

All product mixtures were analysed by thin layer chromatography silica TLC plates. UV-active compounds were detected with a UV lamp ($\lambda = 254$ nm). For column chromatography, silica gel (60–120 mesh) was used as stationary phase. ¹H NMR, ¹³C and DEPT NMR spectra were recorded on 400 MHz spectrometer using CDCl₃, DMSO solvent. ESI-MS and HRMS spectra were recorded on LC-MS/MS and HRMS-6540-UHD machines.

Synthesis of Delamanid

Step 1: Allylation

Procedure for the synthesis of the 1-iodo-4-((2-methylallyl)oxy) benzene (3). The 4-iodophenol 2 (21.9 g, 100 mmol, 1 equiv.) and potassium carbonate (K_2CO_3) (200 mmol, 2 equiv.) were taken in *N*,*N*-dimethylformamide (DMF) and then methallyl chloride 1* (200 mmol, 1.3 equiv.) was added to the reaction mixture and heated to 60 °C for a period of 2 h. Upon cooling the reaction mixture, it was diluted with EtOAc and washed twice with H₂O and brine. Then organic layer was evaporated under vacuum and purified by column chromatography on a silica gel (60–120) with hexane/ethyl acetate (9 : 1) as the eluent to get the desired product **3** as oily liquid. (26.5 g, yield 97%). TLC $R_f = 0.5$ (20%

EtOAc : hexane) ¹H NMR (400 MHz, CDCl₃) δ 7.56–7.54 (d, J = 8.0 Hz, 2H), 6.72–6.70 (d, J = 8.0 Hz, 2H), 5.08 (s, 1H), 5.00 (s, 1H), 4.41 (s, 2H), 1.83 (s, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 158.74, 140.53, 138.27, 117.35, 113.07, 83.03, 71.87, 19.48; GC-MS(EI) m/z (relative intensity) 274 (M⁺, 100%).

#The care has to be taken while handling methallyl chloride as it is a reported to have mutagenic properties.

Step 2: Selective N-arylation

Synthesis of the 1-(4-((2-methylallyl)oxy)phenyl)piperidin-4-ol (5) (Scheme 2, entry 10). An oven-dried round bottom flask was charged with copper iodide (19 mg, 0.1 mmol) and L-proline (23 mg, 0.4 mmol). The flask was evacuated with N_2 atmosphere & then fitted with a rubber septum. DMF (2 mL) was next added, as well as the aryl iodide 3 (136 mg, 0.5 mmol), which was added at this stage. The resulting blue solution was stirred for 10 min before adding 4-hydroxypiperidine 4 (202 mg, 2 mmol). Upon completion, sodium hydroxide (1 M aqueous solution) was added to the reaction mixture and extracted twice with ethyl acetate. Anhydrous sodium sulphate was added to the organic layer, filtered and concentrated to get the desired pure product as solid 5 (104 mg, yield 85%).

Scale up reaction

An oven-dried flask was charged with copper iodide (3.8 g, 20 mmol) and L-proline (4.6 g, 40 mmol). The flask was evacuated with N₂ atmosphere & then fitted with a rubber septum. DMF (120 mL) was next added, as well as the aryl iodide 3 (27 g, 100 mmol), which was added at this stage. The resulting blue solution was stirred for 10 min before adding 4-hydroxypiperidine 4 (40 g, 400 mmol). Upon completion, sodium hydroxide (1 M aqueous solution) was added to the reaction mixture and extracted twice with ethyl acetate. Anhydrous sodium sulphate was added to the organic layer, filtered and concentrated to get the desired pure product as solid 5 (19.8 g, yield 81%). TLC $R_{\rm f} = 0.5$ (40% EtOAc : hexane).

¹H NMR (400 MHz, CDCl₃) δ 6.84–6.76 (m, 4H), 5.00 (s, 1H), 4.89 (s, 1H), 4.31 (s, 2H), 3.76–3.72 (m, 1H), 3.34–3.31 (m, 2H), 2.78–2.71 (m, 2H), 1.96–1.92 (m, 2H), 1.75 (s, 3H), 1.68–1.60 (m, 2H); ¹³C NMR (101 MHz, CDCl₃): δ (ppm) = 153.01, 145.92, 141.32, 118.69, 115.49, 112.43, 72.19, 48.94, 34.47, 29.70, 19.28; HRMS (ESI⁺) calcd for C₁₅H₂₁NO₂ 248.1651 ([M + H]⁺), found 248.1621.

Step 3: Mitsunobu ether formation

Synthesis of 1-(4-((2-methylallyl)oxy)phenyl)-4-(4-(trifluoromethoxy)phenoxy)piperidine (8) (Scheme 4, entry 2). Toa solution of 4-hydroxypiperidine derivative 5 (200 mg, 0.8mmol) in THF (3 mL) was added PPh₃ (420 mg, 1.6 mmol) andfurther a solution of 4-trifluoromethoxyphenol 7 (144 mg, 0.8mmol) and DEAD (324 mg, 1.6 mmol) was added slowly dropwise below 5 °C, and stirred at room temperature for 24 hours.The reaction mixture was allowed to concentrate and aqueoussodium hydroxide solution was added, followed by extractionwith EtOAc. Then anhydrous sodium sulfate was added to theorganic layer and concentrated under reduced pressure andpurified by silica gel column chromatography (eluent; hexane : EtOAc = 4 : 1 (v/v)) to obtain 1-(4-((2-methylallyl)oxy) phenyl)-4-(4-(trifluoromethoxy)phenoxy)piperidine as light brown solid material **8** (231 mg, yield 70%).

Scale up reaction

To a solution of 4-hydroxypiperidine derivative 5 (5 g, 20 mmol) in THF (30 mL) was added PPh₃ (10.5 g, 40 mmol) and further a solution of 4-trifluoromethoxyphenol 7 (3.56 g, 20 mmol) and DEAD (8.1 g, 40 mmol) was added slowly dropwise below 5 °C, and stirred at room temperature for 24 hours. The reaction mixture was allowed to concentrate and aqueous sodium hydroxide solution was added, followed by extraction with EtOAc. Then anhydrous sodium sulfate was added to the organic layer and concentrated under reduced pressure and purified by silica gel column chromatography (eluent; hexane : EtOAc = 4:1 (v/v)) to obtain 1-(4-((2-methylallyl)oxy))phenyl)-4-(4-(trifluoromethoxy)phenoxy)piperidine as light brown solid material 8 (5.93 g, yield 72%). TLC $R_{\rm f} = 0.5$ (5%) EtOAc : hexane).

¹H NMR (400 MHz, CDCl₃) δ 7.17–7.14 (d, J = 12.0 Hz, 2H), 6.96–6.87 (m, 6H), 5.10 (s, 1H), 4.99 (s, 1H), 4.40 (s, 3H), 3.41– 3.36 (m, 2H), 3.03–2.97 (m, 2H), 2.16–2.11 (m, 2H), 2.01–1.93 (m, 2H), 1.84 (s, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 155.97, 153.16, 145.86, 142.84, 141.32, 122.48, 119.33 (q, $J_{C-F} = 256.54$ Hz), 118.71, 116.93, 115.53, 112.52, 72.96, 72.30, 48.27, 30.77, 19.41; HRMS (ESI⁺) calcd for C₂₂H₂₄F₃NO₃ 408.1787 ([M + H]⁺), found 408.1796.

Step 4: Sharpless asymmetric dihydroxylation

Synthesis of (R)-2-methyl-3-(4-(4-(4-(trifluoromethoxy)phenoxy) piperidin-1-yl)phenoxy)propane-1,2-diol $(\mathbf{R} \cdot (-) \cdot \mathbf{9})$ (Scheme 5, entry 3). In a round bottom flask, AD-mix-α (12 g) was taken in tertbutyl alcohol : water (1 : 1, 30 mL) and stir at room temperature to dissolve the AD-mix- α . The mixture was cooled to 0 °C and then added 1-(4-((2-methylallyl) oxy)phenyl)-4-(4-(trifluoromethoxy) phenoxy) piperidine 8 (5 g, 12.28 mmol) and again stir vigorously at 0 °C for 6 h and then 1.93 g (15.3 mmol) of sodium sulphite was added. The reaction mixture was allowed to stir at room temperature for 1 h and diluted with 20 mL of DCM and 40 mL of water and extracted with portions of DCM. The anhydrous MgSO4 was added to the organic phase, filtered and concentrated to afford the crude oil. The residue was purified by chromatography on a silica gel column with gradient elution with 4:1 (hexanes/ ethyl acetate) as brownish gummy material R-(-)-9. Yield (4.7 g, 86%). Enantiomeric excess was determined by chiral HPLC (chiral pack ADH normal phase column, 4.6 \times 250 mm, 5 μ m; enantiomeric excess (ee): 94%). TLC $R_f = 0.5$ (70% EtOAc : hexane). $\left[\alpha\right]_{\mathrm{D}}^{25}$ -6.65° (c 1, chloroform).

¹H NMR (400 MHz, CDCl₃) δ 7.08–7.06 (d, J = 8.0 Hz, 2H), 6.87– 6.82 (m, 4H), 6.80–6.76 (m, 2H), 4.37–4.32 (m, 1H), 3.84–3.78 (dd, J = 12.0 Hz, 1H), 3.66–3.63 (d, J = 12.0 Hz, 1H), 3.51–3.48 (d, J = 12.0 Hz, 1H), 3.33–3.27 (m, 2H), 2.95–2.89 (m, 2H), 2.06–2.01 (m, 2H), 1.90–1.84 (m, 2H), 1.21 (s, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 155.89, 152.65, 146.28, 142.86, 122.46, 119.33(q, $J_{C-F} = 260.58$ Hz), 118.72, 116.93, 115.39, 73.53, 72.91, 72.01, 67.83, 48.14, 30.60, 21.49; HRMS (ESI⁺) calcd for C₂₂H₂₇F₃NO₅ 442.1841 ([M + H]⁺), found 442.1843.

Synthesis of (S)-2-methyl-3-(4-(4-(4-(trifluoromethoxy)phenoxy) piperidin-1-yl)phenoxy)propane-1,2-diol (S-(+)-9) (Scheme 5, entry 4). In a round-bottomed flask equipped with magnetic stirring bar, AD-mix- β (12 g) was taken in *tert*-butyl alcohol : water (1 : 1, 30 mL) and stir at room temperature to dissolve the AD-mix-β. The mixture was cooled to 0 °C and then added the 1-(4-((2-methylallyl) oxy)phenyl)-4-(4-(trifluoromethoxy)phenoxy)piperidine 8 (5 g, 12.28 mmol) and again stir vigorously at 0 °C for 6 h and then 1.93 g (15.3 mmol) of sodium sulphite was added. The reaction mixture was allowed to stir at room temperature for 1 h and diluted with 20 mL of DCM and 40 mL of water and the reaction mixture was extracted with portions of DCM. The anhydrous MgSO4 was added to the organic phase, filtered and concentrated to afford the crude oil. The residue was purified by chromatography on a silica gel column with gradient elution with 4:1 hexanes/ethyl acetate as brownish gummy material S-(+)-9. Yield (4.8 g, 89%). Enantiomeric excess was determined by using chiral HPLC (chiral pack ADH, normal phase column, 4.6 \times 250 mm, 5 μ m; enantiomeric excess (ee): 95%). $[\alpha]_{D}^{25}$ +5° (*c* 1, chloroform).

¹H NMR (400 MHz, CDCl₃) δ 7.14 (d, J = 8.0 Hz 2H), 6.93–6.97 (m, 4H), 6.87–6.90 (d, J = 12.0 Hz, 2H), 4.41 (m, 1H), 3.87 (d, J = 12.0 Hz, 1H), 3.95 (d, J = 12.0 Hz, 1H), 3.74 (d, J = 12.0 Hz, 1H), 3.60 (d, J = 12.0 Hz, 1H), 3.38 (m, 2H), 3.0 (m, 2H), 2.1 (m, 2H), 1.99 (m, 2H), 1.30 (s, 3H).

Step 5: Epoxidation

Synthesis of (R)-2-hydroxy-2-methyl-3-(4-(4-(4-(trifluoromethoxy) phenoxy)piperidin-1-yl)phenoxy)propyl methanesulfonate 10 and (R)-1-(4-((2-methyloxiran-2-yl)methoxy)phenyl)-4-(4-trifluoromethoxy)phenoxy) (11) (Scheme 6). In a 100 mL round-bottomed flask, (R)-2methyl-3-(4-(4-(trifluoromethoxy)phenoxy) piperidin-1-yl) phenoxy)propane-1,2-diol R-(-)-9 (0.9 g, 2 mmol) was taken into DCM solvent and add triethylamine (0.570 mL, 4 mmol) followed by the addition of methanesulfonyl chloride (0.170 mL, 2.2 mmol) dropwise at 0 °C and stirred at room temperature for a period of 2 h. Then solvent was evaporated and extract with ethyl acetate and aqueous layer. The organic solvent was evaporated to afford the mesylated product 10 as gummy material and used as such for the next reaction. In the next step, mesylated compounds 10 was dissolved in ethyl acetate and DBU (0.596 mL, 4 mmol) was added and stirred the reaction mixture for a period of 2 h and then water was added to the reaction mixture and extract with ethyl acetate and evaporate the organic layer under rotary evaporator followed by purification over 100-200 silica gel and ethyl acetate : hexane (40:60) as eluent to afford the compound **11** as brown gummy liquid. (0.7 g, yield 82%).

¹H NMR (400 MHz, CDCl₃) δ 7.15–7.13 (d, J = 8 Hz, 2H), 6.94– 6.91 (dd, J = 4 Hz, 4H), 6.88–6.86 (d, J = 8 Hz, 2H), 4.44–4.38 (m, 1H), 4.00–3.90 (dd, J = 16 Hz, 2H), 3.40–3.35 (m, 2H), 3.02–2.96 (m, 2H), 2.86–2.85 (d, J = 4.0 Hz, 1H), 2.72–2.71 (d, J = 4 Hz, 1H), 2.14–2.09 (m, 2H), 1.99–1.92 (m, 2H), 1.48 (s, 3H); ¹³C NMR (101 MHz, CDCl₃) δ 156.02, 152.94, 146.28, 142.86, 122.46, 119.40 (q, $J_{C-F} = 256.54$ Hz), 118.66, 116.97, 115.53, 73.01, 72.20, 55.63, 51.97, 48.14, 30.77, 18.49; HRMS (ESI⁺) calcd for C₂₂H₂₄F₃NO₄ 424.1730 ([M + H]⁺, found 424.1733.

Step 6: Ring opening and cyclisation

Paper

Synthesis of 1-(2-bromo-4-nitro-1H-imidazol-1-yl)-2-methyl-3-(4-(4-(trifluoromethoxy)phenoxy)piperidin-1-yl)phenoxy)propan-2ol (13) (Scheme 7). A mixture of epoxide 11 (424 mg, 1 mmol), 2bromo-4-nitro-1H-imidazole 12 (190 mg, 1 mmol), and DIPEA (2.0 mL) were taken into a sealed tube and heated to 115 °C for a period of 12 h. Upon cooling the reaction mixture, dichloromethane was added and evaporated to dryness under reduced pressure and then residue was purified on silica gel 100–200 and then eluted with 0.5% EtOAc/CH₂Cl₂ gave the desired compound 13 (552 mg) in a yield of 90%.

¹H NMR (400 MHz, CDCl₃) δ 8.11 (s, 1H), 7.16–7.14 (d, J = 8.0 Hz, 2H), 6.95–6.91 (m, 4H), 6.83–6.81 (d, J = 8.0 Hz, 2H), 4.43 (m, 1H), 4.31–4.28 (d, J = 16 Hz, 1H), 4.20–4.16 (d, J = 16 Hz, 1H), 3.87–3.79 (m, 2H), 3.42–3.35 (m, 2H), 3.02 (m, 2H), 2.17–2.06 (m, 2H), 2.00–1.91 (m, 2H), 1.33 (s, 3H). Mass (m/z): 615 (M + H)⁺.

Synthesis of 2-methyl-6-nitro-2-((4-(4-(4-(trifluoromethoxy)phenoxy)piperidin-1-yl)phenoxy)methyl)-2,3-dihydroimidazo[2,1-b]oxazoles (II, Delamanid, Scheme 7). To the stirred solution of compound 13 (307 mg, 0.5 mol) in anhydrous DMF, caesium carbonate was taken and stirred the reaction mixture for 2 h at 50 °C and then upon completion of reaction, add water and extract with ethyl acetate. Then evaporate the solvent and purified the reaction mixture with silica gel 100-200 eluting with 40% ethyl acetate: dichloromethane gave compound (Delamanid II) as white solid (185 mg, 70%). Mp 195–196 °C. ¹H NMR (CDCl₃) δ 7.58 (s, 1H), 7.17-7.15 (d, J = 8.0 Hz, 2H), 6.84-6.91 (m, 4H), 6.81-6.78 (d, *J* = 12.0 Hz, 2H), 4.53–4.50 (d, *J* = 12.0 Hz, 1H), 4.46–4.41 (1H, m), 4.21–4.18 (d, J = 10.2 Hz, 1H), 4.07–4.04 (dd, J = 12.0 Hz, 2H), 3.41-3.35 (m, 2H), 3.04-2.98 (m, 2H), 2.14-2.09 (m, 2H), 1.99-1.93 (m, 2H), 1.78 (s, 3H); ¹⁹F NMR (CDCl₃, 376 MHz): δ –58.32; ¹³C NMR (101 MHz, CDCl₃) δ 155.92, 151.71, 146.89, 122.50, 119.29 (q, $J_{\rm C-F} = 256.54$ Hz) 118.56, 116.84, 115.57, 112.54, 93.24, 72.70, 72.23, 51.43, 47.86, 30.57, 23.16; $[\alpha]_{\rm D}^{25}$ -12° (c 1, chloroform) $[[\alpha]_D^{25} - 9.9^\circ];^7$ HRMS (ESI⁺) calcd for $C_{25}H_{25}F_3N_4O_6$ 535.180 ([M + H]⁺, found 535.179.

Synthesis of 2,5-dibromo-4-nitro-1H-imidazole S_2 (Fig. S4 of ESI†). In a three necked round bottom flask (3 L), 4-nitroimidazole S_1 was taken into water (400 mL) and stir the mixture to temperature below 5 °C. Then sodium bicarbonate (86 g, 1 mol) was added followed by the slow addition of bromine (160 g, 1.0 mmol) to reaction mixture over a period of 4.0 h by keeping the temperature below 0 °C and then stir it at 65 °C for 6.0 h. After completion of reaction, pH of the reaction mixture was adjusted with aqueous hydrochloric acid (6.0 N, 50 mL) to pH 2–3 and stirred it for 1.0 h below 5 °C and filtered. The compound S_2 obtained as a pale-yellow solid (104 g, 88%). Mp 229–231 °C.

¹H NMR: (400 MHz, DMSO-d₆) δ 7.98 (s, 1H); mass (*m*/*z*): 271.4 (M + H)⁺.

Synthesis of 2-bromo-4-nitro-1H-imidazole (12) (Fig. S4 of ESI⁺). In a 3 L capacity round bottom flask, 2,5-dibromo-4-nitro-1H-imidazole, S_2 (100 g, 378 mmol) was taken into glacial acetic acid (1.2 L) and stirred it at room temperature. Then potassium iodide (94 g, 567 mmol), and sodium sulfite (71 g, 567 mmol) was added to the reaction mixture and heated to 120–125 °C for 16 h. Upon completion, glacial acetic acid was distill off and water (500 mL) was added to the reaction mixture and stirred for 10 min, followed by the addition of aqueous sodium bisulfite solution (5%) to quench excess iodide. The reaction mixture was extracted with ethyl acetate (3×250 mL) and washed with water (2×250 mL), dried over magnesium sulphate, and concentrated to get 2-bromo-4-nitroimidazole, **12** as a yellow solid (72 g, 65%). Mp 240 °C.

 ^{1}H NMR: (400 MHz, DMSO-d_6) δ 14.14 (bs, 1H), 8.41 (s, 1H); HRMS (ESI^+) calcd for $C_{3}H_{3}BrN_{3}O_{2}$ 191.9409 ([M + H]^+, found 191.9402.

Synthesis of VL-2098

Step 1: Allylation

Procedure for the synthesis of the 1-((2-methylallyl)oxy)-4-(trifluoromethoxy)benzene (14) (Scheme 8). The starting material 4-trifluoromethoxyphenol 7 (5 g, 28 mmol, 1 equiv.) and K₂CO₃ (7.8 g, 56 mmol, 2 equiv.) were taken in DMF and then methallyl chloride 1 (3 mL, 36.4 mmol, 1.3 equiv.) was added and heated the reaction mixture to 60 °C for a period of 2 h. Upon cooling the reaction mixture, it was diluted with EtOAc and washed twice with H₂O and brine. Then organic layer was evaporated under vacuum and purified by column chromatography on a silica gel (60-120) with hexane/ethyl acetate (9:1) as the eluent to get the desired product (6.32 g, yield 97%). ¹H NMR (400 MHz, $CDCl_3$) δ 7.12 (d, J = 8.5 Hz, 2H), 6.89 (d, J = 8.5 Hz, 2H), 5.09 (s, 1H), 5.00 (s, 1H), 4.40 (s, 2H), 1.82 (s, 3H). ¹⁹F NMR (CDCl₃, 376 MHz): δ -58.32. ¹³C NMR (101 MHz, CDCl₃) δ 157.40, 142.87, 140.55, 124.52, 122.32, 121.98, 119.44 (q, J_{C-F} = 256.54 Hz), 115.54, 112.88, 72.06, 19.14. GC-MS(EI) m/z (relative intensity) 232 (M⁺, 100%).

Step 2: Sharpless asymmetric dihydroxylation

Synthesis (*R*)-2-methyl-3-(4-(trifluoromethoxy)phenoxy) of propane-1,2-diol R-(+)-15 (Scheme 8). In a 100 mL roundbottomed flask AD-mix- α (4 g) was taken in *tert*-butyl alcohol-: water (1 : 1, 30 mL) and stir at room temperature to dissolve the AD-mix-α. Add 1-((2-methylallyl)oxy)-4-(trifluoromethoxy) benzene 14 (1 g, 4.31 mmol) at 0 °C and stir it for a period of 6-12 h and then sodium sulphite (1.93 g, 15.3 mmol) was added. The reaction mixture was allowed to stir at room temperature for 1 h and diluted with DCM (20 mL) and water (40 mL) and extracted with DCM. The combined organic phase was dried over anhydrous MgSO₄, filtered and concentrated to get crude oil. The residue was purified through column chromatography on a silica gel (100-200 mesh size) with gradient elution with 4:1 hexane/ethyl acetate as white solid material R-(+)-15. Yield 1.0 g (88%). $[\alpha]_{D}^{25}$ +8.83°.

¹H NMR (400 MHz, CDCl₃) δ 7.14 (d, J = 8.8 Hz, 2H), 6.90 (d, J = 9.0 Hz, 2H), 3.90 (q, J = 8.9 Hz, 2H), 3.71 (dd, J = 11.0 Hz, 1H), 3.57 (dd, J = 11.0 Hz, 1H), 2.82 (s, 1H), 2.44 (s, 1H), 1.29 (s, 3H). LC-MS (ESI⁺): m/z calcd for (M–OH)⁺ 249.20.

Step 3: Epoxidation

Synthesis of (R)-2-hydroxy-2-methyl-3-(4-(trifluoromethoxy) phenoxy)propyl methanesulfonate **16** and (R)-1-(4-((2methyloxiran-2-yl)methoxy)phenyl)-4-(4-trifluoromethoxy) phenoxy) (**17**) (Scheme 8). In a 100 mL round-bottomed flask, (R)-2-methyl-3-(4-(trifluoromethoxy)phenoxy)propane-1,2-diol **R**-(+)-**15** (0.9 g, 3.7 mmol) in DCM solvent was taken triethylamine (0.570 mL, 4 mmol) and then methanesulfonyl chloride (0.170

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mL, 2.2 mmol) was added dropwise at 0 °C and stir the reaction mixture at room temperature for a period of 2 h. Then solvent was evaporated and extract with ethyl acetate and aqueous layer. Then evaporation of organic phase gave the mesylated product **16** as gummy material and used as such for the next reaction. In the next step, mesylated compounds **16** was dissolved in ethyl acetate and DBU (0.596 mL, 4 mmol) was added and stirred the reaction mixture for a period of 2 h and then water was added to the reaction mixture and extract with ethyl acetate and evaporate the organic layer under rotary evaporator followed by purification over 100–200 silica gel and ethyl acetate : hexane (40 : 60) as eluent to afford the compound **17** as yellow clear liquid (0.75 g) with the yield of 89%.

¹H NMR (400 MHz, CDCl₃) δ 7.10 (d, J = 8.7 Hz, 2H), 6.87 (d, J = 8.8 Hz, 2H), 4.00 (d, J = 10.5 Hz, 1H), 3.87 (d, J = 10.5 Hz, 1H), 2.82 (d, J = 4.6 Hz, 1H), 2.69 (d, J = 4.6 Hz, 1H), 1.44 (s, 3H). LC-MS (ESI⁺): m/z calcd for (M + H)⁺ 249.20.

Step 4: Ring opening and cyclization

Synthesis of (R)-2-methyl-6-nitro-2-((4-(trifluoromethoxy)phenoxy)methyl)-2,3-dihydroimidazo[2,1-b]oxazole (VL-2098, VII) (Scheme 8). A mixture of epoxide 17 (248 mg, 1 mmol), 2-bromo-4-nitro-1*H*-imidazole (12, 190 mg, 1 mmol), and DIPEA (2.0 mL) were taken in a sealed tube and heated at 115 °C for a period of 2 h. Upon cooling the reaction mixture, dichloromethane was added and evaporated to get crude product and used for the next reaction. To the stirred solution of crude product in anhydrous DMF was added caesium carbonate and stirred the reaction mixture for 2 h at 50 °C and then upon completion of reaction, add water and extract the reaction mixture with ethyl acetate. Then evaporate the solvent and purified the reaction mixture with silica gel 100–200 eluting with 40% ethyl acetate : dichloromethane gave compound **VII** as white solid (172 mg, 48%). $[\alpha]_{D}^{25} + 7^{\circ}$ (*c* 1, chloroform) $[[\alpha]_{D}^{25} + 8.445^{\circ}]$.

¹H NMR (400 MHz, CDCl₃) δ 7.53 (s, 1H), 7.11 (d, J = 8.8 Hz, 2H), 6.81 (d, J = 9.1 Hz, 2H), 4.46 (d, J = 10.3 Hz, 1H), 4.19 (d, J = 10.1 Hz, 1H), 4.05 (dd, J = 10.0, 8.0 Hz, 2H), 1.75 (s, 3H). LC-MS (ESI⁺): m/z calcd for (M + H)⁺ 360.20.

Conflicts of interest

There are no conflicts to declare.

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