



DMP 777

Catalog No: tcsc0355

| Available Sizes |
|---|
| Size: 5mg |
| Size: 10mg |
| Size: 25mg |
| Size: 50mg |
| Specifications |
| CAS No: 157341-41-8 |
| Formula: C ₃₁ H ₄₀ N ₄ O ₆ |
| Pathway: Metabolic Enzyme/Protease |
| Target: Elastase |
| Purity / Grade: >98% |
| Solubility: DMSO : ≥ 50 mg/mL (88.55 mM) |
| Alternative Names: L-694458 |
| Observed Molecular Weight: 564.67 |





Product Description

DMP 777 is a potent, selective, and orally active human leukocyte elastase (HLE) inhibitor.

In Vivo: DMP-777-treated rats show a marked decrease in H/K-ATPase staining parietal cells. DMP-777-induced loss of parietal cells is significantly ameliorated with coadministration of omeprazole. DMP-777-treated animals demonstrates marked foveolar hyperplasia in the fundus with prominent expansion of diastase-resistant, PAS-positive surface mucous cells. When DMP-777 is coadministered

with omprazole, there is a significant decrease in BrdUpositive S-phase cells compared with rats thatreceive DMP-777 alone^[1]. After oral dosing of monkeys at 40 mg/kg with DMP-777 the only stereoisomer detected in the post-dose plasma samples is the starting material DMP-777, and no inversion of the configuration at positions \'a\' and \'b\' of DMP-777 has occurred in vivo^[2]. Mist1-/- mice treated with DMP-777 show fewer chief cell to SPEM transitions. Mist1-/- mice treated with L635 demonstrates significantly fewer proliferative SPEM cells compared to control mice^[3].

All products are for RESEARCH USE ONLY. Not for diagnostic & therapeutic purposes!