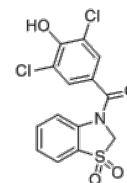


**Product Name** : Dotinurad  
**Cat. No.** : PC-60041  
**CAS No.** : 1285572-51-1  
**Molecular Formula** : C<sub>14</sub>H<sub>9</sub>Cl<sub>2</sub>NO<sub>4</sub>S  
**Molecular Weight** : 358.189  
**Target** : URAT1  
**Solubility** : 10 mM in DMSO



### Biological Activity

Dotinurad (FYU-981) is a potent and selective urate transporter 1 (**URAT1**) inhibitor with IC<sub>50</sub> of 37.2 nM.

Dotinurad weakly inhibited ATP-binding cassette subfamily G member 2 (ABCG2), organic anion transporter 1 (OAT1), and OAT3, with IC<sub>50</sub> values of 4.16, 4.08, and 1.32 μM, respectively, indicating higher selectivity for URAT1.

Dotinurad (1-30 mg/kg) decreased plasma urate levels and increased fractional excretion of urate (FEUA) in a dose-dependent manner in Cebus monkeys.

Dotinurad is a potent and selective urate reabsorption inhibitor is characterized by increased efficacy with decreasing plasma urate levels.

### References

- Tetsuya Taniguchi, et al. *J Pharmacol Exp Ther.* 2019 Oct;371(1):162-170.
- Hosoya T, et al. *Clin Exp Nephrol.* 2020 Mar;24(Suppl 1):53-61.

**Caution: Product has not been fully validated for medical applications. Lab Use Only!**

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