# **PRODUCT** INFORMATION

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## Dotinurad

Item No. 41014

| CAS Registry No.:<br>Formal Name:  | 1285572-51-1<br>(3,5-dichloro-4-hydroxyphenyl)(1,1-dioxido-<br>3(2H)-benzothiazolyl)-methanone |         |
|--|--|---------|
| Synonym:   | FYU-981  | × \ /~  |
| MF:  | C <sub>14</sub> H <sub>9</sub> Cl <sub>2</sub> NO <sub>4</sub> S                               |         |
| FW:  | 358.2  | О       |
| Purity:  | ≥98%   |         |
| Supplied as:   | A solid  | \<br>Cl |
| Storage:   | -20°C  | 01      |
| Stability:   | ≥4 years   |         |
| Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis. |  |         |

#### Laboratory Procedures

Dotinurad is supplied as a solid. A stock solution may be made by dissolving the dotinurad in the solvent of choice, which should be purged with an inert gas. Dotinurad is sparingly soluble (1-10 mg/ml) in DMSO.

### Description

Dotinurad is a urate transporter 1 (URAT1) antagonist (IC<sub>50</sub> = 0.0372  $\mu$ M).<sup>1</sup> It is selective for URAT1 over ATP-binding cassette subfamily G member 2 (ABCG2), organic anion transporter 1 (OAT1), and OAT3 (IC<sub>50</sub>s = 4.16, 4.08, and 1.32 µM, respectively). In vivo, dotinurad (30 mg/kg) decreases plasma levels of urate and increases urine levels of urate in Cebus monkeys. Formulations containing dotinurad have been used in the treatment of hyperuricemia and gout.

#### Reference

1. Taniguchi, T., Ashizawa, N., Matsumoto, K., et al. Pharmacological evaluation of dotinurad, a selective urate reabsorption inhibitor. J. Pharmacol. Exp. Ther. 371(1), 162-170 (2019).

WARNING THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

#### SAFETY DATA

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

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