## **Product** Data Sheet

## Treprostinil-d7

Molecular Weight:

Cat. No.: HY-100441S3  $\label{eq:hybrid} \mbox{Molecular Formula:} \qquad \mbox{$C_{23}$H}_{27}\mbox{$D_7$O}_5$ 

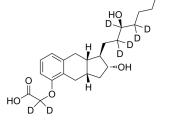
Target: Isotope-Labeled Compounds; Prostaglandin Receptor

Pathway: Others; GPCR/G Protein

**Storage:** Please store the product under the recommended conditions in the Certificate of

Analysis.

397.56



## **BIOLOGICAL ACTIVITY**

Description

Treprostinil-d7 (UT-15-d7) is a deuterated version of Treprostinil (HY-100441). Treprostinil is a highly potent DP1 and EP2 agonist with EC50s of 0.6 nM and 6.2 nM, respectively<sup>[1][2][3][4][5][6][7]</sup>.

## **REFERENCES**

- [1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. Ann Pharmacother. 2019;53(2):211-216.
- [2]. Whittle BJ, et al. Binding and activity of the prostacyclin receptor (IP) agonists, treprostinil and iloprost, at human prostanoid receptors: treprostinil is a potent DP1 and EP2 agonist. Biochem Pharmacol. 2012 Jul 1;84(1):68-75.
- [3]. Ferrantino M, et al. Inhaled treprostinil sodium for the treatment of pulmonary arterial hypertension. Expert Opin Pharmacother. 2011 Nov;12(16):2583-93.
- [4]. Ghonem N, et al. Treprostinil, a prostacyclin analog, ameliorates ischemia-reperfusion injury in rat orthotopic liver transplantation. Am J Transplant. 2011 Nov;11(11):2508-16.
- [5]. Smadja DM, et al. Treprostinil indirectly regulates endothelial colony forming cell angiogenic properties by increasing VEGF-A produced by mesenchymal stem cells. Thromb Haemost. 2015 Oct;114(4):735-47.
- [6]. Kazemi Z, et al. Repurposing Treprostinil for Enhancing Hematopoietic Progenitor Cell Transplantation. Mol Pharmacol. 2016 Jun;89(6):630-44.
- [7]. Nikam VS, et al. Treprostinil inhibits the recruitment of bone marrow-derived circulating fibrocytes in chronic hypoxic pulmonary hypertension. Eur Respir J. 2010 Dec;36(6):1302-14.

Caution: Product has not been fully validated for medical applications. For research use only.

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