## Retinoic acid-d6

Cat. No.: CAS No.: Molecular Formula: Molecular Weight: Target: Pathway:	HY-14649S3 2483831-72-5 C <sub>20</sub> H <sub>22</sub> D <sub>6</sub> O <sub>2</sub> 306.47 RAR/RXR; PPAR; Autophagy; Endogenous Metabolite Metabolic Enzyme/Protease; Vitamin D Related/Nuclear Receptor; Cell Cycle/DNA Damage; Autophagy	
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.	

BIOLOGICAL ACTIVITY		
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Description	Retinoic acid-d <sub>6</sub> is the deuterium labeled Retinoic acid[1]. Retinoic acid is a metabolite of vitamin A that plays important roles in cell growth, differentiation, and organogenesis. Retinoic acid is a natural agonist of RAR nuclear receptors, with IC50s of 14 nM for RAR $\alpha/\beta/\gamma$ . Retinoic acid bind to PPAR $\beta/\delta$ with Kd of 17 nM. Retinoic acid acts as an inhibitor of transcription factor Nrf2 through activation of retinoic acid receptor alpha[2][3][4][5][6][7].	
In Vitro	Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to affect the pharmacokinetic and metabolic profiles of drugs <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.	

## REFERENCES

[1]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. Ann Pharmacother. 2019 Feb;53(2):211-216.

[2]. Wu L, et al. Retinoid X Receptor Agonists Upregulate Genes Responsible for the Biosynthesis of All-Trans-Retinoic Acid in Human Epidermis. PLoS One. 2016 Apr 14;11(4):e0153556.;

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

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**Product** Data Sheet

