## **Product** Data Sheet

## PKR activator 1

 $\begin{array}{lll} \textbf{Cat. No.:} & \textbf{HY-135883} \\ \textbf{CAS No.:} & 2283420\text{-}62\text{-}0 \\ \textbf{Molecular Formula:} & \textbf{C}_{16}\textbf{H}_{14}\textbf{N}_{8}\textbf{OS}_{2} \\ \end{array}$ 

Molecular Weight: 398.47

Target: Pyruvate Kinase

Pathway: Metabolic Enzyme/Protease

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

Analysis.

## **BIOLOGICAL ACTIVITY**

| Description               | PKR activator 1 is a potent pyruvate kinase-R (PKR) activator extracted from patent WO2019035865A1, compound E7-93 <sup>[1]</sup> .  |
|---------------------------|--|
| IC <sub>50</sub> & Target | Pyruvate kinase-R (PKR) <sup>[1]</sup>   |
| In Vitro                  | Pyruvate kinase (PK) is a metabolic enzyme that converts phosphoenolpyruvate to pyruvate during glycolysis. Four PK iso forms exist in mammals: the L and R iso forms (from the PKLR gene) are expressed in liver and red blood cells respectively, and the PKM gene encodes two splice variants, the MI isoform that is expressed in most adult tissues, and the M2 isoform that is expressed during embryonic development and in some adult tissues including the kidney and hematopoietic stem cells. Many tumor cells also express PKM2. This tetrameric allosterically regulated isoform is intrinsically designed to downregulate its activity, through post-translational modification, allosteric modulation by small molecule ligands including some amino acids, and by subunit dissociation (into the dimeric form), which results in partial inhibition of glycolysis at the last step <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only. |

## **REFERENCES**

[1]. Charles KUNG, et al. Pyruvate kinase modulators and use thereof. WO2019035865A1.

 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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