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**Data Sheet** 

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Product Name	:	PF-07328948
Cat. No.	:	PC-23451
CAS No.	:	2936625-34-0
Molecular Formula	:	C <sub>16</sub> H <sub>8</sub> F <sub>4</sub> O <sub>3</sub> S
Molecular Weight	:	356.29
Target	:	Other Targets
Solubility	:	10 mM in DMSO



CAS: 2936625-34-0

## **Biological Activity**

PF-07328948 is the first potent, selective branched-chain ketoacid dehydrogenase kinase (BDK or BCKDK) with in vitro IC50 of 15 nM, and cellular IC50 of 46 nM in human skeletal muscle BDK activity assay.

PF-07328948 is not only a BDK inhibitor but also BDK degrader.

PF-07328948 shows a high degree of selectivity over a number of enzymes, receptors, and ion channels, as well as excellent kinase selectivity.

PF-07328948 binds to the BDK allosteric pocket and does not contain a traditional ATP-binding site motif.

PF-07328948 binds to BDK with SPR Kd of 4.8 nM.

PF-07328948 also destabilizes the interaction between BDK and E2.

PF-07328948 also shows high species potency against mouse BDK IC50 = 41 nM, rat BDK IC50 = 24 nM, and dog BDK IC50

= 24 nM. lowered plasma biomarkers, including BCAAs and branched-chain ketoacids (BCKAs) in vivo with enhanced pharmacodynamic effect upon chronic dosing due to BDK degradation.

## References

Filipski KJ, et al. J Med Chem. 2024 Nov 19. doi: 10.1021/acs.jmedchem.4c02230.

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

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