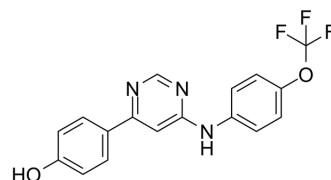


PROTAC BCR-ABL1 ligand 1

Cat. No.:	HY-130297
CAS No.:	2489876-34-6
Molecular Formula:	C ₁₇ H ₁₂ F ₃ N ₃ O ₂
Molecular Weight:	347.29
Target:	Bcr-Abl
Pathway:	Protein Tyrosine Kinase/RTK
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	PROTAC BCR-ABL1 ligand 1, compound GMB-475, is the ligand of PROTAC that allosterically targets BCR-ABL1 protein and recruits the E3 ligase Von Hippel-Lindau, resulting in ubiquitination and subsequent degradation of BCR-ABL1 ^[1] .								
In Vitro	<p>GMB-475 (0-10 μM) exhibits cell proliferation with IC₅₀ values of 1.11, 1.98, 0.37 μM for BCR-ABL1 WT, T315I, G250E cells, respectively^[1].</p> <p>Caution: Product has not been fully validated for medical applications. For research use only.</p> <p>GMB-475 (0-30 μM) induces the degradation of BCR-ABL1 and c-ABL1 with concomitant inhibition of downstream signaling via the STAT5 pathway, in a dose- and time-dependent fashion in the context of both K562 and Ba/F3 cells^[1].</p> <p>MCE has not independently confirmed the accuracy of these methods. They are for reference only.</p> <p>Western Blot Analysis^[1]</p> <table border="1"> <tr> <td>Cell Line:</td> <td>K562 and Ba/F3 cells</td> </tr> <tr> <td>Concentration:</td> <td>0-30 μM</td> </tr> <tr> <td>Incubation Time:</td> <td></td> </tr> <tr> <td>Result:</td> <td>Decreased p-STAT5 expression.</td> </tr> </table>	Cell Line:	K562 and Ba/F3 cells	Concentration:	0-30 μM	Incubation Time:		Result:	Decreased p-STAT5 expression.
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REFERENCES

[1]. Burslem GM, et al. Targeting BCR-ABL1 in Chronic Myeloid Leukemia by PROTAC-Mediated Targeted Protein Degradation. Cancer Res. 2019 Sep 15;79(18):4744-4753.