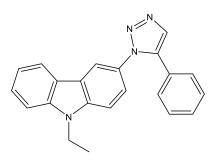


## Catalog # 10-4434 MBQ-167

CAS# 2097938-73-1 1-(9-Ethyl-9H-carbazol-3-yl)-5-phenyl-1H-1,2,3-triazole; 9-Ethyl-3-(5-phenylcarbazol-1-yl)carbazole Lot # FBA8084



MBQ-167 is a dual inhibitor of the Rho GTPases Rac (IC<sub>50</sub> = 103 nM) and Cdc42 (IC<sub>50</sub> = 78 nM).<sup>1</sup> It inhibits breast cancer cell migration, viability, and mammosphere formation. MBQ-167 induced polarity loss resulting in 95% cell rounding and detachment from the substratum in metastatic MDA-MB-231 cells and was active in GFP-HER2-BM, MDA-MB-468, and Hs578t breast cancer cells as well as Mia-PaCa-2 pancreatic cancer cells, SKOV3 ovarian cancer cells, AGS and NCI-N87 gastric cancer cells, and SH-SY5Y neuroblastoma cells. Non-cancerous mammary epithelial MCF10A and epithelial breast cancer MCF-7 cells were resistant to MBQ-167. MBQ-167 inhibited viability and induced apoptosis in gefitinib and lapatinib resistant SKBR3 breast cancer cells.<sup>2</sup> MBQ-167 inhibited triple negative breast cancer tumor growth and lung metastasis in a mouse model.<sup>3</sup> Active *in vivo*.

- 1) Humphries-Bickley et al. (2017) Characterization of a Dual Rac/Cdc42 Inhibitor MBQ-167 in Metastatic Cancer; Mol.Cancer Ther. **16** 805
- 2) Borrero-Garcia et al. (2021) Rac inhibition as a novel therapeutic strategy for EGFR/HER2 targeted therapy resistant breast cancer; BMC Cancer 21 652
- Cruz-Collazo et al. (2021) Efficacy of Rac and Cdc42 inhibitor MBQ-167 in triple negative breast cancer; Mol. Cancer Ther. 20 2420

## PHYSICAL DATA

Molecular Weight:	338.41
Molecular Formula:	C <sub>22</sub> H <sub>18</sub> N <sub>4</sub>
Purity:	>98% (HPLC)
Solubility:	DMSO (>25 mg/ml)
Physical Description:	Off-white solid
Storage and Stability:	Store as supplied at -20° for up to one year from the date of purchase. Solutions in
	DMSO may be stored at -20°C for up to 2 months

Materials provided by Focus Biomolecules are for laboratory research use only and are not intended for human or veterinary applications.

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