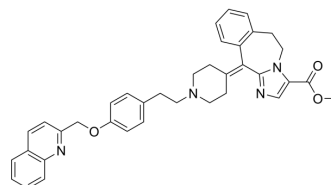


## Laniquidar

Cat. No.:	HY-132189
CAS No.:	197509-46-9
Molecular Formula:	C <sub>37</sub> H <sub>36</sub> N <sub>4</sub> O <sub>3</sub>
Molecular Weight:	584.71
Target:	P-glycoprotein
Pathway:	Membrane Transporter/Ion Channel
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

Description	Laniquidar (R101933) is a noncompetitive, third generation P-glycoprotein (P-gp) inhibitor with an IC <sub>50</sub> of 0.51 μM. Laniquidar can be used for modulating multidrug resistance transporters <sup>[1]</sup> . Laniquidar can also be used for studying acute myeloid leukemia (AML) and myelodysplastic syndrome (MDS) <sup>[2]</sup> . Laniquidar has limited oral bioavailability <sup>[3]</sup> .
IC <sub>50</sub> & Target	IC <sub>50</sub> : 0.51 μM (P-glycoprotein) <sup>[1]</sup>

### REFERENCES

- [1]. Luurtsema G, et al. Evaluation of [<sup>11</sup>C]laniquidar as a tracer of P-glycoprotein: radiosynthesis and biodistribution in rats. Nucl Med Biol. 2009 Aug;36(6):643-9.
- [2]. Ross DD. Modulation of drug resistance transporters as a strategy for treating myelodysplastic syndrome. Best Pract Res Clin Haematol. 2004 Dec;17(4):641-51.
- [3]. Fox E, et al. Tariquidar (XR9576): a P-glycoprotein drug efflux pump inhibitor. Expert Rev Anticancer Ther. 2007 Apr;7(4):447-59.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA