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## DATASHEET

tatM2NX

### Product overview

|                               |   |
|-------------------------------|---|
| <b>Name</b>                   | tatM2NX   |
| <b>Cat No</b>                 | HB7349  |
| <b>Biological description</b> | Novel, potent and cell permeable TRPM2 antagonist ( $IC_{50} = 396nM$ ). Prevents ligand binding and TRPM2 activation. Inhibits over 90% of human TRPM2 channel currents at concentrations as low as 2 $\mu M$ . Shows neuroprotective effects in animal models of focal and global ischemia. Active <i>in vivo</i> . |
| <b>Biological action</b>      | Antagonist  |
| <b>Purity</b>                 | >93%  |
| <b>Description</b>            | TRPM2 antagonist. Cell permeable.   |

### Solubility & Handling

|                                      |   |
|--------------------------------------|---|
| <b>Storage instructions</b>          | -20 °C  |
| <b>Solubility overview</b>           | Soluble in aqueous buffer   |
| <b>Storage of solutions</b>          | Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.                  |
| <b>Shipping Conditions Important</b> | Stable for <b>ambient temperature</b> shipping. Follow storage instructions on receipt.<br>This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use |

### Chemical Data

|                                |  |
|--------------------------------|--|
| <b>Molecular Weight</b>        | 4354.17  |
| <b>Molecular Formula</b>       | $C_{190}H_{323}N_{71}O_{45}S$  |
| <b>Sequence (one letter)</b>   | YGRKKRRQRRRGSRPEGM LPRK LKRVLRQEFWW  |
| <b>Sequence (three letter)</b> | H-Tyr-Gly-Arg-Lys-Lys-Arg-Arg-Gln-Arg-Arg-Arg-Gly-Ser-Arg-Glu-Pro-Gly-Glu-Met-Leu-Pro-Arg-Lys-Leu-Lys-Arg-Val-Leu-Arg-Gln-Glu-Phe-Trp-Val-OH |

### References

#### Extended therapeutic window of a novel peptide inhibitor of TRPM2 channels following focal cerebral ischemia.

Shimizu T et al (2016) Experimental neurology 283  
**PubMedID** [27317297](#)

#### Characterization and Optimization of the Novel Transient Receptor Potential Melastatin 2 Antagonist tatM2NX.

Cruz-Torres I et al (2020) Molecular pharmacology 97  
**PubMedID** [31772034](#)

