# PRODUCT INFORMATION



Zanamivir-<sup>13</sup>C, <sup>15</sup>N<sub>2</sub> (hydrate)

Item No. 30733

5-(acetylamino)-4-[(amino-15N-imino-Formal Name:

<sup>15</sup>N-methyl-<sup>13</sup>C)amino]-2,6-anhydro-

3,4,5-trideoxy-D-glycero-D-ido-non-2-

enonic acid, hydrate

 $C_{11}[^{13}C]H_{20}N_2[^{15}N]_2O_7 \bullet XH_2O$ MF:

FW: 335.3 **Purity:** ≥98% Supplied as: A solid Storage: -20°C Stability: ≥4 years XH<sub>2</sub>O

Information represents the product specifications. Batch specific analytical results are provided on each certificate of analysis.

### **Laboratory Procedures**

Zanamivir-13C,15N2 (hydrate) is supplied as a solid. A stock solution may be made by dissolving the zanamivir-<sup>13</sup>C,<sup>15</sup>N<sub>2</sub> (hydrate) in water. Zanamivir-<sup>13</sup>C,<sup>15</sup>N<sub>2</sub> (hydrate) is slightly soluble in water. We do not recommend storing the aqueous solution for more than one day.

### Description

Zanamivir-13C,15N2 is intended for use as an internal standard for the quantification of zanamivir (Item No. 15123) by GC- or LC-MS. Zanamivir is a sialic acid analog that inhibits neuraminidase release of newly replicated influenza virus particles. It has been shown to selectively inhibit the growth of influenza A and B viruses in plaque reduction assays with  $IC_{50}$  values ranging from 5 to 14 nM and to directly inhibit influenza A and B virus neuraminidases with IC<sub>50</sub> values ranging from 0.6 to 7.9 nM in vitro. Intranasal zanamivir administration at 0.4 mg/kg twice daily reduces mortality and viral titers in lung homogenates of mice infected with influenza.

### Reference

1. Elliott, M. Zanamivir: From drug design to the clinic. Philos. Trans. R. Soc. Lond. B Biol Sci. 356(1416), 1885-1893 (2001).

WARNING
THIS PRODUCT IS FOR RESEARCH ONLY - NOT FOR HUMAN OR VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

This material should be considered hazardous until further information becomes available. Do not ingest, inhale, get in eyes, on skin, or on clothing. Wash thoroughly after handling. Before use, the user must review the complete Safety Data Sheet, which has been sent via email to your institution.

## WARRANTY AND LIMITATION OF REMEDY

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