# PRODUCT INFORMATION



## Ravidasvir (hydrochloride)

Item No. 37794

CAS Registry No.: 1303533-81-4

Formal Name: N-[(1S)-1-[[(2S)-2-[5-[6-[2-[(2S)-1-

> [(2S)-2-[(methoxycarbonyl)amino]-3methyl-1-oxobutyl]-2-pyrrolidinyl]-1Hbenzimidazol-6-yl]-2-naphthalenyl]-1Himidazol-2-yl]-1-pyrrolidinyl]carbonyl]-2methylpropyl]-carbamic acid, methyl ester,

dihydrochloride

Synonym: **PPI-668** 

MF: C<sub>42</sub>H<sub>50</sub>N<sub>8</sub>O<sub>6</sub> • 2HCl

FW: 835.8 **Purity:** >98%

UV/Vis.:  $\lambda_{\text{max}}$ : 267, 321 nm

Supplied as: A solid Storage: -20°C Stability: ≥4 years

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

### **Laboratory Procedures**

Ravidasvir (hydrochloride) is supplied as a solid. A stock solution may be made by dissolving the ravidasvir (hydrochloride) in the solvent of choice, which should be purged with an inert gas. Ravidasvir (hydrochloride) is soluble in organic solvents such as ethanol, DMSO, and dimethyl formamide (DMF). The solubility of ravidasvir (hydrochloride) in ethanol is approximately 1 mg/ml and approximately 5 mg/ml in DMSO and DMF.

### Description

Ravidasvir is an inhibitor of hepatitis C virus (HCV) non-structural protein 5A (NS5A).<sup>1</sup> It inhibits HCV replication in Huh7 replicon cells containing NS5A from the HCV genotypes 1a, -1b, and 3a with EC<sub>50</sub> values of 0.12, 0.01, and 1.14 nM, respectively.

### Reference

1. Zhong, M., Peng, E., Huang, N., et al. Discovery of ravidasvir (PPI-668) as a potent pan-genotypic HCV NS5A inhibitor. Bioorg. Med. Chem. Lett. 26(18), 4508-4512 (2016).

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.

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