

## NIH AIDS Reagent Program

20301 Century Boulevard Building 6, Suite 200 Germantown, MD 20874 USA

Phone: 240 686 4740 Fax: 301 515 4015 aidsreagent.org

## **DATA SHEET**

Raltegravir (ISENTRESS/MK-0518) Reagent:

Catalog Number: 11680

Lot Number: 080169

Release Category: Е

**Provided:** 2 mg

 $\label{eq:n-constraint} $$N-[(4-Fluorophenyl)methyl]-1,6-dihydro-5-hydroxy-1-methyl-2-[1-methyl-1-[(5-methyl-1,3,4-oxadiazol-2-yl)carbonyl]amino]ethyl]-6-oxo-4-$ **Chemical Name:** 

pyrimidinecarboxamide monopotassium salt

**Empirical** 

Formula:

C20H20FN6O5K.

**HPLC Purity:** 99.85%

Molecular Weight:

482.52

**CAS Num:** 871038-72-1

**Purity:** 99.85%

Solubility: Raltegravir is soluble in water. It is slightly soluble in methanol and ethanol. It is insoluble

in isopropanol.

ALL RECIPIENTS OF THIS MATERIAL MUST COMPLY WITH ALL APPLICABLE BIOLOGICAL, CHEMICAL, AND/OR RADIOCHEMICAL SAFETY STANDARDS INCLUDING SPECIAL PRACTICES, EQUIPMENT, FACILITIES, AND REGULATIONS. NOT FOR USE IN HUMANS.

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Raltegravir is an HIV-1 integrase inhibitor. Inhibition of the viral integrase prevents insertion of HIV DNA into the human DNA genome thus blocking the ability of HIV-1 to replicate. Warning: Reagent can cause severe eye and respiratory irritation. Toxic to aquatic organisms.

Recommended Storage:

Room temperature. Once resuspended, working aliquots can be stored at -20°C.

**Contributor:** 

Merck & Company, Inc.

NOTE:

Acknowledgment for publications should read "The following reagent was obtained through the NIH AIDS Reagent Program, Division of AIDS, NIAID, NIH: Raltegravir from Merck &

Company, Inc (Cat # 11680)."

This compound is restricted for "research purposes only" and is limited to 2.0 mg

per requester. Not available for release to commercial organizations.

Recipient agrees that the reagent (Raltegravir) use is permitted only as a standard for in vitro and/or studies in animals for inhibition of HIV replication.

**Last Updated** 

July 24, 2018

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