

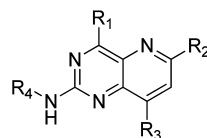
Pyridopyrimidines as Inhibitors of Hepatitis C Virus

Gerard Rosse*

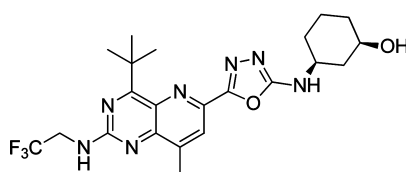
Structure Guided Chemistry, Dart Neuroscience LLC, 7473 Lusk Boulevard, San Diego, California 92121, United States, and Adjunct Associate Professor, Department of Pharmacology and Physiology, College of Medicine, Drexel University, New College Building, 245 North 15th Street, Philadelphia, Pennsylvania 19102, United States

Title: Pyridopyrimidines as Inhibitors of Hepatitis C Virus
Patent/Patent Application Number: WO 2013090840 A1
Publication date: June 20, 2013
Priority Application: US 2011-576284P
Priority date: December 15, 2011
Inventors: Jin, H.; Kim, C. U.; Li, J.
Assignee Company: Gilead Sciences, Inc., USA
Disease Area: Hepatic C Virus infection
Biological Target:
Summary: This application claims pyridopyrimidine analogues for the treatment and prevention of HCV infections.

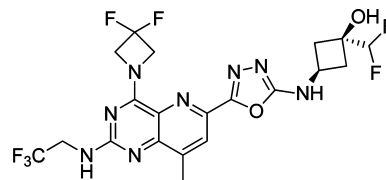
Important Compound Classes:



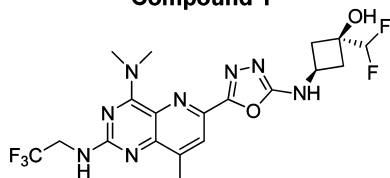
Key Structures:



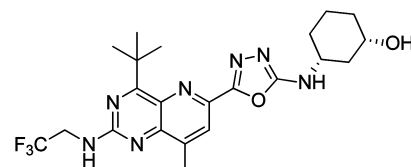
Compound 1



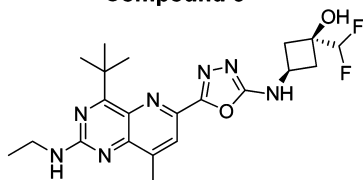
Compound 2



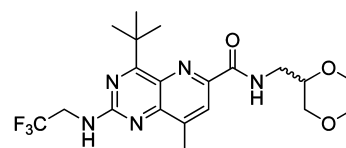
Compound 3



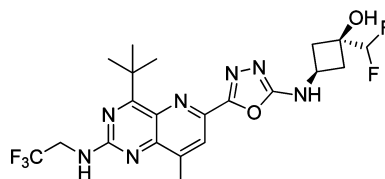
Compound 4



Compound 5



Compound 6



Compound 7

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Recent Review Articles:**Biological Assay:**

Effect of compounds was evaluated using HCV replicon assays developed from H77 (GT1a), Con1 (GT1b), or JHF-1 (GT2a) strain.

Pharmacological Data:

Seven compounds were tested for antiviral activity in HCV replicon assay. The unit for the EC₅₀ is not described.

Compound	HCV replicon 1A EC ₅₀	HCV replicon 2A EC ₅₀	HCV replicon 1B EC ₅₀
1	1295.5	0.684	5.566
2	4444.4	18.447	444.44
3	2991.7	2.495	42.367
4	94.726	0.165	1.075
5	1882.3	0.507	0.756
6	4444.4	3.879	4.502
7	300.77	0.092	0.461

Synthesis:

The synthesis of seven compounds is described.

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Notes

The authors declare no competing financial interest.