

Aminoquinoline Derivatives as HCV Inhibitors

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Title:	Aminoquinoline derivatives as HCV inhibitors			
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Priority Application:	US 2011-576278P	Priority date:	Dec 15, 2011	
Inventors:	Cai, Z. R.; Du, Z.; Ji, M.; Jin, H.; Kim, C. U.; Li, J.; Phillips, B. W.; Pyun, HJ.; Saugier, J. H.			
Assignee Company:	Gilead Sciences, Inc., USA			
Disease Area:	HCV infection	Biological Target:		
Summary:	The application claims new aminoquinoline derivatives that may provide a treatment against HCV infections.			
Important Compound Classes:		R^{6} N R^{2}		

Key Structures:



R⁴ R³

Biological Assay: Pharmacological Data: The HCV replicon assay was developed using cell line HCV subgenomic RNA derived from GT1a, GT1b, and GT2a. Over 180 compounds were tested for antiviral activity in HCV replicon assay. The unit for the EC₅₀ is not described.

	HCV replicon	HCV replicon	HCV replicon
	assay – GT1a	assay – GT1b	assay – GT2a
	(EC_{50})	(EC_{50})	(EC ₅₀)
Compound 4	0.060	0.076	9.423
Compound 5	0.071	0.221	130.71
Compound 25	0.028	0.064	25.750
Compound 105	0.951	1.306	753.840

Synthesis:

Over 180 compounds were prepared.

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Notes

The authors declare no competing financial interest.

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