

## Substituted Benzofurans as Inhibitors of HCV NS5B Protein

## Gerard Rosse\*

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Title: Substituted Benzofurans as Inhibitors of HCV NS5B Protein

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Inventors: Yeung, K.-S.; Kadow, J. F.

Assignee Company: Bristol-Myers Squibb Company, USA

Disease Area: HCV infection Biological Target: HCV NSSB protein

Summary: This application claims a series of benzofuran analogues inhibits HCV NSSB protein and may provide a treatment against

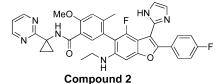
HCV infections.

Important Compound Classes:

$$R_1$$
 $R_2$ 
 $R_3$ 
 $R_4$ 
 $R_6$ 
 $R_5$ 

**Key Structures:** 

Compound 1



Compound 3

Biological Assay: An on-bead solid-phase homogeneous assay was used to asses NSSB inhibitors. Compound efficacy was evaluated using

HCV replicon luciferase assay.

Pharmacological Data:

|            | NS5B Inhibition    | HCV replicon assay |
|------------|--------------------|--------------------|
|            | $(IC_{50}, \mu M)$ | $(EC_{50}, \mu M)$ |
| Compound 1 | 0.030              | 0.030              |
| Compound 2 | NT                 | 0.0072             |
| Compound 3 | NT                 | 0.0090             |

Synthesis: Preparation of 3 compounds.

## AUTHOR INFORMATION

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Notes Special Issue: HCV Therapies

The authors declare no competing financial interest.

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