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**DESCRIPTION**

*Antiviral Research* publishes research reports, short communications, review articles and commentaries on the control of viral infections in humans and animals. Its scope encompasses: antiviral drugs, antibodies and host-response modifiers, including their synthesis, *in vitro* and *in vivo* testing and mechanisms of action; new or improved vaccines against viral infections of humans; assessments of drug and vaccine safety; evolution of drug- or vaccine-resistant viruses and the development of effective countermeasures; identification and validation of new drug targets; laboratory animal models of viral diseases; pathogenesis of viral diseases and mechanisms of viral evasion of host immune responses.

We encourage the submission of manuscripts describing the activity of well-defined chemical compounds for the treatment of viral diseases of humans and vertebrate animals. We also publish manuscripts on the protective activity of vaccines for humans, but we do not accept research reports on veterinary vaccines. All papers must include a sufficiently detailed description of methods to permit other investigators to replicate the experiments. Authors reporting the antiviral effect of a novel small-molecule drug must reveal its chemical structure. Claims of *in vitro* or *in vivo* efficacy of a drug or vaccine must be supported by appropriate statistical analysis.

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*Antiviral Research* is an official publication of the International Society for Antiviral Research ([http://www.isar-icar.com](http://www.isar-icar.com)).

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AUDIENCE

Virologists, clinicians, veterinarians, medicinal chemists, researchers in the pharmaceutical industry and members of regulatory authorities.

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The scope of AVR is limited to viral diseases of humans and vertebrate animals. We do not accept reports on viral diseases of plants or of insects, crustaceans or other invertebrates. We discourage the submission of manuscripts reporting the antiviral activity of unpurified natural products, or of partially purified substances of natural origin for which a mechanism of action has not been determined. Manuscripts claiming an antiviral effect of homeopathic products or other highly diluted preparations, or which fail to clearly identify the biological ingredient or molecule responsible for the antiviral activity of an experimental therapy, will not be considered for publication. Articles describing antiseptics with broad-spectrum antimicrobial activity will not be accepted. We also discourage submission of in silico docking studies or other computer-based predictions of antiviral activity that are not supported by data from biological assays.

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- One U1i inhibitor revealed high potency in various transient transfection assays.
- This potency is due to a tandem target site in the HIV-1 RNA.
- U1i combined with a lentiviral gene therapy approach needs optimization.

**From "Orthopoxvirus targets for the development of new antiviral agents"**
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**From "An engineered inhibitor RNA that efficiently interferes with hepatitis C virus translation and replication"**
We analyse the anti-HCV potential of the RNA molecule HH363-24.
It cleaves the HCV genome and binds to the essential domain IIId of the IRES region.
We analyse the anti-HCV potential of the RNA molecule HH363-24.
The inhibitor interferes with viral translation and replication in cell culture.

From "Polyfunctional CD8+ T cells are associated with the vaccination-induced control of a novel recombinant influenza virus expressing an HCV epitope"
Lipopeptide vaccination elicits polyfunctional CD8+ T cells at multiple organ sites.
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Polyfunctional CD8+ T cells are associated with control of the recombinant HCV influenza virus.

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