

# Dose-ranging study of SCH34117 in the treatment of patients with seasonal allergic rhinitis

<b>Submission date</b> 02/08/2002	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 02/08/2002	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 06/09/2007	<b>Condition category</b> Respiratory	<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**Protocol serial number**  
C98-001

## Study information

**Scientific Title**

**Study objectives**

Not provided at time of registration

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Randomised controlled trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

**Health condition(s) or problem(s) studied**

Seasonal allergic rhinitis

**Interventions**

Patients randomly assigned once-daily (QD) SCH34117 at doses of 2.5, 5, 7.5, 10, or 20 mg versus placebo for two weeks.

**Intervention Type**

Drug

**Phase**

Not Specified

**Drug/device/biological/vaccine name(s)**

SCH34117

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

01/06/1998

**Eligibility****Key inclusion criteria**

Subjects 12 years or older with a two-year documented history of seasonal allergic rhinitis and otherwise in good health

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/04/1998

**Date of final enrolment**

01/06/1998

## **Locations**

**Countries of recruitment**

United States of America

**Study participating centre**

**Medical officer**

Kenilworth

United States of America

NJ 07033-0530

## **Sponsor information**

**Organisation**

Schering-Plough Research Institute (SPRI) (USA)

**ROR**

<https://ror.org/02891sr49>

## **Funder(s)**

**Funder type**

Industry

**Funder Name**

Schering-Plough Research Institute (SPRI) (USA)

## Results and Publications

**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	Results	05/08/2002		Yes	No