

# Intranasal zolmitriptan is effective and well tolerated in acute cluster headache

<b>Submission date</b> 14/02/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 23/02/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 06/08/2008	<b>Condition category</b> Signs and Symptoms	<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**  
ZINCH-1

## Study information

**Scientific Title**

**Acronym**  
ZINCH

**Study objectives**

Zolmitriptan intranasal is more effective than placebo in the treatment of acute cluster headache.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Ethics approval received from the National Hospital for Neurology and Neurosurgery Ethics Committee on the 8th May 2003 (ref: 02/N031).

**Study design**

Randomised, placebo-controlled, double-blind cross-over trial

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Cluster headache

**Interventions**

Zolmitriptan nasal spray versus placebo.

**Intervention Type**

Drug

**Phase**

Not Specified

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

Zolmitriptan

**Primary outcome(s)**

Proportion of patients who have taken active drug with headache relief at 30 minutes.

**Key secondary outcome(s))**

Pain free at 30 minutes.

**Completion date**

01/01/2005

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

Cluster headache with attacks longer than 45 minutes.

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

Contraindications to zolmitriptan in participating countries.

**Date of first enrolment**

01/09/2003

**Date of final enrolment**

01/01/2005

## Locations

**Countries of recruitment**

United Kingdom

England

Germany

Italy

**Study participating centre**

**Institute of Neurology**

London

United Kingdom

WC1N 3BG

## Sponsor information

**Organisation**

AstraZeneca (UK)

**ROR**

<https://ror.org/04r9x1a08>

# Funder(s)

## Funder type

Industry

## Funder Name

AstraZeneca (UK)

## Alternative Name(s)

AstraZeneca PLC, Pearl Therapeutics, AZ

## Funding Body Type

Government organisation

## Funding Body Subtype

For-profit companies (industry)

## Location

United Kingdom

# Results and Publications

## Individual participant data (IPD) sharing plan

### IPD sharing plan summary

## Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	Results	01/11/2006		Yes	No