

IBS-C Study A2306

Submission date 12/08/2003	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 11/09/2003	Overall study status Completed	☐ Protocol
Last Edited 20/02/2008	Condition category Digestive System	☐ Statistical analysis plan
		☐ Results
		☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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07936

Additional identifiers

Protocol serial number

CHTF919A2306

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.

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Irritable bowel syndrome (IBS) with constipation

Interventions

Not provided at time of registration.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration.

Key secondary outcome(s)

Not provided at time of registration.

Completion date

30/04/2003

Eligibility**Key inclusion criteria**

Females, 18 to 65 years, with IBS-C (Irritable Bowel Syndrome with constipation)

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 Years

Sex

Female

Key exclusion criteria

Not provided at time of registration.

Date of first enrolment

01/04/2002

Date of final enrolment

30/04/2003

Locations**Countries of recruitment**

United States of America

Study participating centre

Novartis Pharmaceuticals

East Hanover

United States of America

07936

Sponsor information**Organisation**

Novartis Pharmaceuticals Corporation

ROR

<https://ror.org/028fhxy95>

Funder(s)**Funder type**

Industry

Funder Name

Sponsored by Novartis Pharmaceuticals Corporation (<http://click.atdmt.com/GTO/go/thbmbnov00700895gto/direct/01/>)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration