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IBS-C Study A2306

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

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
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.

Study design
Randomised controlled trial

Primary study design
Interventional

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