

# Natural plants for the management of nausea and vomiting

<b>Submission date</b> 29/09/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 29/09/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
<b>Last Edited</b> 30/09/2014	<b>Condition category</b> Signs and Symptoms	<input type="checkbox"/> Individual participant data <input type="checkbox"/> 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

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**  
N0284173983

## Study information

**Scientific Title**

**Study objectives**

The purpose is to assess whether two specific herbs (ginger and peppermint) in capsule form are effective treatments for the management of nausea and vomiting caused by chemotherapy.

**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

**Secondary study design**

Randomised controlled trial

**Study setting(s)**

Not specified

**Study type(s)**

Treatment

**Participant information sheet****Health condition(s) or problem(s) studied**

Signs and symptoms: nausea and vomiting

**Interventions**

Patients will be randomised into groups, some will be given the peppermint or ginger and some will be given a placebo.

**Intervention Type**

Other

**Phase**

Not Applicable

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

15/03/2005

**Completion date**

15/03/2006

## Eligibility

**Key inclusion criteria**

Not provided at time of registration

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

Patients will be recruited from clinic

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

15/03/2005

**Date of final enrolment**

15/03/2006

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

8 Eileen Grove West

Manchester

United Kingdom

M14 5NW

## Sponsor information

**Organisation**

**Sponsor details**

The Department of Health, Richmond House, 79 Whitehall  
London  
United Kingdom  
SW1A 2NL  
+44 (0)20 7307 2622  
dhmail@doh.gsi.org.uk

**Sponsor type**

Government

**Website**

<http://www.dh.gov.uk/Home/fs/en>

**Funder(s)****Funder type**

Government

**Funder Name**

Wrightington, Wigan and Leigh NHS Trust (UK)

**Funder Name**

NHS R&D Support Funding (UK)

**Results and Publications****Publication and dissemination plan**

Not provided at time of registration

**Intention to publish date****Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration