

The clinical management of chemotherapy-induced nausea and vomiting with adjuvant Progressive Muscle Relaxation Training (PMRT) and imagery techniques in breast cancer patients

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
10/10/2002	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
10/10/2002	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
05/10/2007	Cancer	

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr A Molassiotis

Contact details

Nethersole School of Nursing
Chinese University of Hong Kong

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Hong Kong

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Additional identifiers

Protocol serial number

811006

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

Breast cancer

Interventions

PMRT 1 hour prior to chemotherapy administration and daily thereafter for another 5 days (total of 6 PMRT sessions). Each session lasts for 25 minutes followed by 5 minutes of imagery techniques.

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

01/01/2001

Eligibility

Key inclusion criteria

1. Patients diagnosed with breast cancer and have no exposure to cytotoxic chemotherapeutic agent prior to the study (chemotherapy-naive)
2. Females between the age of 18 - 70 years

3. Clinical indication to receive adriamycin with cyclophosphamide (AC) as adjuvant or salvage chemotherapy for treatment of breast cancer
4. Able to read and write Chinese
5. Agree to sign the informed consent

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

01/01/1999

Date of final enrolment

01/01/2001

Locations

Countries of recruitment

Hong Kong

Study participating centre

Nethersole School of Nursing

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Hong Kong

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Sponsor information

Organisation

Hong Kong Health Services Research Fund (China)

ROR

<https://ror.org/03qh32912>

Funder(s)

Funder type

Government

Funder Name

Hong Kong Health Services Research Fund (Hong Kong)

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?
Results article	Results	01/04/2002		Yes	No