

HERA: A randomised three-arm multi-centre comparison of 1 year and 2-years of Herceptin® versus no Herceptin® in women with HER2-positive primary breast cancer who have completed adjuvant chemotherapy

Submission date 15/10/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 15/10/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 22/10/2018	Condition category Cancer	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT00045032

Secondary identifying numbers

N/A

Study information

Scientific Title

HERA: A randomised three-arm multi-centre comparison of 1 year and 2-years of Herceptin® versus no Herceptin® in women with HER2-positive primary breast cancer who have completed adjuvant chemotherapy

Acronym

HERA

Study objectives

Added 08/09/09:

Monoclonal antibodies such as trastuzumab can locate tumor cells and either kill them or deliver tumor-killing substances to them without harming normal cells. It is not yet known whether trastuzumab is effective in treating primary breast cancer in women who have completed adjuvant chemotherapy.

Primary objectives:

1. Compare the disease-free survival of women with HER2-positive primary breast cancer treated with trastuzumab (Herceptin®) for 1 year vs trastuzumab for 2 years vs standard supportive care.
2. Compare the overall survival of patients treated with these regimens.
3. Compare the relapse-free survival of patients treated with these regimens.
4. Compare the distant disease-free survival of patients treated with these regimens.
5. Compare the incidence of cardiac dysfunction in patients treated with these regimens.
6. Evaluate the safety and tolerability of these regimens in these patients.

Secondary objectives:

1. Compare time to recurrence in patients treated with these regimens.
2. Compare time to distant recurrence in patients treated with these regimens.
3. Compare outcomes, in terms of disease-free survival, overall survival, recurrence-free survival, distant disease-free survival, time to recurrence, time to distant recurrence, cardiac safety, and overall safety, in patients treated with trastuzumab for 1 year vs 2 years.

Please note that as of 08/09/09 this record has been extensively updated. All updates can be found under the relevant field with the above update date.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Multicentre randomised open label controlled parallel group trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

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

Breast cancer

Interventions

Current information as of 08/09/09:

This is a randomised, open-label, multicentre study. Patients are stratified according to nodal status (any nodal status and prior neoadjuvant chemotherapy vs no positive nodes and no prior neoadjuvant chemotherapy vs 1-3 positive nodes and no prior neoadjuvant chemotherapy vs 4 or more positive nodes and no prior neoadjuvant chemotherapy), prior adjuvant chemotherapy regimen (no anthracyclines or taxanes vs anthracyclines only vs anthracyclines and taxanes), receptor status and endocrine therapy (negative vs positive and no prior endocrine therapy vs positive and prior endocrine therapy), age (18 to 34 vs 35 to 49 vs 50 to 59 vs 60 and over), and participating center. Patients are randomized to 1 of 3 treatment arms.

1. Arm I: Patients receive trastuzumab (Herceptin®) IV over 1.5 hours on day 1. Courses repeat every 3 weeks for up to 1 year in the absence of disease progression or unacceptable toxicity.
2. Arm II: Patients receive trastuzumab as in arm I. Courses repeat every 3 weeks for up to 2 years in the absence of disease progression or unacceptable toxicity.
3. Arm III: Patients receive no trastuzumab. Patients may later receive trastuzumab as in arm I or arm II. Patients are followed every 3 months for 2 years, every 6 months for 3 years, and then annually thereafter.

Initial information at time of registration

1. Patients are randomised to receive Herceptin® every 3 weeks for 1 or 2 years
2. No further treatment

Intervention Type

Drug

Phase

Phase III

Drug/device/biological/vaccine name(s)

Trastuzumab (Herceptin®)

Primary outcome measure

Added 08/09/09:

1. Disease-free survival
2. Relapse-free survival
3. Distant disease-free survival
4. Incidence of cardiac dysfunction
5. Safety and tolerability

Secondary outcome measures

Added 08/09/09:

1. Overall survival
2. Time to recurrence
3. Time to distant recurrence

Overall study start date

01/03/2002

Completion date

01/12/2004

Eligibility

Key inclusion criteria

1. Females aged ≥ 18 years
2. Eastern Cooperative Oncology Group (ECOG) performance status ≤ 1
3. Non-metastatic operable primary invasive adenocarcinoma of the breast that is histologically confirmed, adequately excised and axillary node positive or negative
4. Known hormone receptor status
5. Completion of at least 3 months of an approved (neo-) adjuvant chemotherapy regimen
6. Baseline left ventricular ejection fraction (LVEF) $\geq 55\%$
7. Completion of radiotherapy for any patients undergoing radiotherapy
8. Overexpression of HER2 in the invasive component of the primary tumour
9. Completion of all necessary baseline lab and radiological investigations
10. Signed written informed consent

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Female

Target number of participants

3192

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/03/2002

Date of final enrolment

01/12/2004

Locations**Countries of recruitment**

Argentina

Australia

Austria

Belgium

Brazil

Canada

Chile

China

Colombia

Croatia

Denmark

England

France

Germany

Greece

Guatemala

Hong Kong

Hungary

Ireland

Israel

Italy

Japan

Netherlands

Poland

Portugal

Russian Federation

Singapore

South Africa

Spain

Sweden

Switzerland

Thailand

United Kingdom

Study participating centre
UKCCCR Register Co-ordinator
London
United Kingdom
NW1 2DA

Sponsor information

Organisation

Roche Products Limited (UK)

Sponsor details

P.O. Box 8
Welwyn Garden City, Hertfordshire
United Kingdom
AL7 3AY

Sponsor type

Industry

Website

<http://www.roche.com>

ROR

<https://ror.org/024tgbv41>

Funder(s)

Funder type

Industry

Funder Name

Roche Products Ltd (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Plain English results				No	Yes
Results article	results	20/10/2005		Yes	No
Results article	results	06/01/2007		Yes	No
Results article	results on adverse cardiac effects	01/09/2007		Yes	No
Results article	results	01/06/2008		Yes	No
Results article	results	20/06/2009		Yes	No
Results article	results	20/07/2010		Yes	No