External radiotherapy versus expectancy in localized prostate cancer

Submission date	Recruitment status No longer recruiting	Prospectively registered		
08/08/2011		☐ Protocol		
Registration date 26/09/2011	Overall study status Completed Condition category	Statistical analysis plan		
		Results		
Last Edited		Individual participant data		
19/05/2017	Cancer	Record updated in last year		

Plain English summary of protocol

Background and study aims

Prostate cancer is a common cancer, where many patients have such a slowly growing tumor that it is uncertain if they need treatment. Other patients have very aggressive tumors, and in Sweden with 10000 newly diagnosed prostate cancer patients 2500 will die each year. No scientific studies had in 1986 shown that any of the treatment options for localized prostate cancer expectancy or radiotherapy or surgery was superior. Expectancy had been the treatment of choice in the beginnings of the 1980s. The aim of this study was to evaluate if radiotherapy could prolong progression free survival and reduce mortality in comparison to expectancy.

Who can participate?

Patients with prostate cancer under the age of 75

What does the study involve?

Participants are followed for the progression of their symptoms and are offered radiotherapy. The use of four-field EBRT, 64 Gy, 2.0 cm margin was increased during the beginning of 1980's, further technical improvements such as CT-based conformal radiotherapy was developed by the beginning of the 1990ies, dose was increased to 70 Gy with margins around 1-1.5 cm.

What are the possible benefits and risks of participating? Not provided at time of registration

Where is the study run from? Umeå University (Sweden)

When is study starting and how long is it expected to run for? April 1986 to January 1997

Who is funding the study?

- 1. Cancer Research Foundation (Sweden)
- 2. Lions Cancer Research Foundation, Umea University (Sweden)
- 3. Umeå University Hospital (Sweden)

Who is the main contact?
Professor Andrew Widmark
Anders.Widmark@onkologi.umu.se

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Version 1991 and 1995 04

Study information

Scientific Title

External radiotherapy versus expectancy in low-intermediate risk localized prostate cancer: a randomized trial

Study objectives

To investigate if radiotherapy will prolong progression free survival and reduce mortality in prostate cancer in early localized prostate cancer in comparison to active expectancy.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Study was part of the regular treatment program for localized prostate cancer in Northern Sweden (Ethics approval not necessary at that time [1996]).

Study design

Randomized open multi-center trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Prostate cancer

Interventions

Expectancy: Patients included: 107 out of (130) 160, Patients followed until symptomatic clinical progression, and then offered radiotherapy

Radiotherapy: Patients included: 107 out of (130) 160, Treated with 64-70 Gy, boxfield or conformal radiotherapy, and than follow up, until clinical progression and than castration treatment

Intervention Type

Other

Phase

Phase III

Primary outcome measure

- 1. Progression Free survival
- 2. Overall survival

Secondary outcome measures

Morbidity

Overall study start date

23/04/1986

Completion date

27/01/1997

Eligibility

Key inclusion criteria

- 1. Patients with verified localized prostate cancer, T1-T2b, pN0, Grade 1 and 2, M0
- 2. Age less than or equal to 75 years
- 3. Surgical obturatorious lymph node staging, with no signs of positive lymph nodes
- 4. No signs on metastasis after clinical investigation, bone scan and pulmonary X-ray
- 5. No previous treatment against their prostate cancer
- 6. No other known malign or benign disease that would shorten their lifes in relation to a healthy population
- 7. Patients approval of participation

Participant type(s)

Patient

Age group

Adult

Sex

Male

Target number of participants

260 (revised to 320)

Key exclusion criteria

Patients not willing to participate in the long follow up program

Date of first enrolment

23/04/1986

Date of final enrolment

27/01/1997

Locations

Countries of recruitment

Denmark

Norway

Sweden

Study participating centre Umea University

Umeå Sweden SE - 90187

Sponsor information

Organisation

Umeå University Hospital (Sweden)

Sponsor details

c/o Prof Anders Widmark Umeå Sweden SE - 901 87 +46 (0)90 785 2857 Anders.Widmark@onkologi.umu.se

Sponsor type

Hospital/treatment centre

Website

http://www.umu.se/english

ROR

https://ror.org/012k96e85

Funder(s)

Funder type

Government

Funder Name

Cancer Research Foundation (Sweden)

Alternative Name(s)

The Cancer Research Foundation, CRF

Funding Body Type

Government organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United States of America

Funder Name

Lions Cancer Research Foundation, Umeå University (Sweden)

Funder Name

Umeå university Hospital, Umeå (Sweden)

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?
Other publications		15/12/2001		Yes	No
Other publications		01/12/2009		Yes	No