Photodynamic versus white light-guided treatment of non-muscle invasive bladder cancer

Submission date	Recruitment status No longer recruiting	[X] Prospectively registered		
25/07/2014		[X] Protocol		
Registration date	Overall study status	Statistical analysis plan		
25/07/2014	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
26/04/2023	Cancer			

Plain English summary of protocol

https://www.cancerresearchuk.org/about-cancer/find-a-clinical-trial/a-trial-white-light-blue-light-surgery-people-bladder-cancer-photo

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

PHOTOdynamic versus white light-guided treatment of non-muscle invasive bladder cancer: randomised trial of clinical and cost-effectiveness

Acronym

PHOTO

Study objectives

Bladder cancer is a high priority area for research into clinical and cost-effective management and the findings from the PHOTO trial are likely to remain highly relevant and important to the needs of the NHS over the next 20 years, the expected life span of the equipment for the PDD technology. A further compelling reason for the study is the current piecemeal adoption of PDD within the NHS, resulting in variation in provision of PDD service. This gives further urgent need for better quality evidence to guide providers of bladder cancer services and the relevant practice guidance authorities to make early decisions around wholesale adoption or disinvestment in PDD technology.

Ethics approval required

Old ethics approval format

Ethics approval(s)

NRES Committee North East - Newcastle & North Tyneside 2, 16/07/2014, ref: 14/NE/1062

Study design

Randomised; Interventional; Design type: Treatment

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

Topic: Cancer; Subtopic: Bladder Cancer; Disease: Bladder (superficial)

Interventions

Randomisation will be undertaken centrally using either the secure web-based or the 24-hour Interactive Voice Response randomisation system at the Centre for Healthcare Randomised Trials (CHaRT) in Aberdeen, using minimisation by centre and gender, to allocate participants 1:1 to the control and experimental groups. The minimisation algorithm will incorporate a random element in order to prevent deterministic treatment allocation.

Guidance on treatment and follow up of NMIBC, developed by the European Association of Urology (EAU), recommend that patients who have high risk disease, or where there are indications that the TURT may not have been complete, should have a further resection of the original disease site 26 weeks after initial treatment. This reresection is intended to ensure that the initial resection was as complete as possible and to check that muscle invasive disease, which would require further radical treatment, has not been missed. Following initial or reresection, patients in both intermediate and high risk groups are recommended to receive further (adjuvant) intravesical treatment with the aim of reducing the risk of future recurrence and progression. Patients in the intermediate risk group are usually given further treatment with chemotherapy once a week for 6 weeks. Patients in the high risk group are treated with intravesical immunotherapy weekly for 6 weeks and may have further treatments every few months for up to 3 years. Patients will have 3 monthly cystoscopies to check for any sign of recurrence for 2 years, then 6 monthly to five years, after which they will continue to be followed up annually.

PHOTO follow-up schedule

All participants will be followed up according to EAU guidelines, with regular cystoscopies initially 3 monthly following surgery (either initial TURT or second TURT for those who have one). The PHOTO trial will collect the following information at the 3,6,9,12,18,24 and 36 month routine visits:

- 1. Outcome of cystoscopy
- 2. Adverse event information

Participants will be asked to complete quality of life and health service utilisation questionnaires 3, 6, 12, 18, 24 and 36 months post randomisation. Participants will also be asked to complete a patient costs questionnaire 30 months after randomisation. Questionnaires will be posted to patients homes by CHaRT.

Intervention Type

Other

Phase

Phase III

Primary outcome measure

- 1. To compare time to recurrence, for each of the two treatment strategies, with a principal point of interest at 3 years.
- 2. To evaluate cost effectiveness by the incremental cost for recurrence avoided and cost utility as the incremental cost per quality adjusted life year (QALY) gained at three years.

Secondary outcome measures

Secondary objectives will further explore clinical and cost effectiveness of photo dynamic surgery:

Clinical effectiveness:

- 1. Measure relative rate of disease progression at 3 years
- 2. Measure relative harms and safety
- 3. Measure health-related quality of life (HRQoL) and cancer specific survival

Overall study start date

01/09/2014

Completion date

23/06/2021

Eligibility

Key inclusion criteria

- 1. Adult men and women aged ≥ 16 years
- 2. First suspected diagnosis of bladder cancer
- 3. Visual/ultrasound/CT diagnosis of intermediate/high risk NMIBC
- 3.1. White light visual appearances of intermediate or high risk disease (=> 3cm, two or more tumours, or flat velvety erythematous changes alerting a clinical suspicion of CIS).
- 3.2. Suspicion of papillary bladder tumour > 3cm based on ultrasound or computerized tomography (CT) scanning (without hydronephrosis)
- 4. Written informed consent for participation prior to any study specific procedures
- 5. Willing to comply with lifestyle guidelines

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Planned Sample Size: 533; UK Sample Size: 533

Total final enrolment

538

Key exclusion criteria

- 1. Visual evidence of low risk NMIBC (solitary tumour < 3cm)
- 2. Visual evidence of MIBC on preliminary cystoscopy, i.e. nonpapillary or sessile mass (attached directly by its base without a stalk)
- 3. Imaging evidence of MIBC CT/USS (this includes the presence of hydronephrosis, which may be present despite clear imaging of MIBC in the bladder)
- 4. Upper tract (kidney or ureteric) tumours on imaging
- 5. Any other malignancy in the past 2 years (except: nonmelanomatous skin cancer cured by excision, adequately treated carcinoma in situ of the cervix, DCIS/LCIS of the breast or prostate cancer in patients who have a life expectancy of >5 years upon trial entry)
- 6. Evidence of metastases
- 7. Porphyria or known hypersensitivity to porphyrins

- 8. Known pregnancy (based on history and without formal testing, in keeping with day-to-day NHS practice of PDD use)
- 9. Any other conditions that in the Principal Investigators opinion would contraindicate protocol treatment
- 10. Unable to provide informed consent
- 11. Unable or unwilling to complete follow up schedule (including questionnaires)

Date of first enrolment

23/10/2014

Date of final enrolment

14/02/2018

Locations

Countries of recruitment

England

Scotland

United Kingdom

Wales

Study participating centre

Freeman Hospital

Freeman Road Newcastle upon Tyne United Kingdom NE7 7DN

Study participating centre Royal Devon and Exeter Hospital

Barrack Road Exeter United Kingdom EX2 5DW

Study participating centre **Churchill Hospital**

Old Road Oxford **United Kingdom**

OX3 7LE

Study participating centre Ninewells Hospital and Medical School Dundee United Kingdom DD2 1UB

Study participating centre
University College Hospital London
235 Euston Road
London
United Kingdom
NW1 2BU

Study participating centre Princess Of Wales Hospital Coity Road Bridgend United Kingdom CF31 1RQ

Study participating centre St Peter's Hospital Guildford Road Chertsey United Kingdom KT16 0PZ

Study participating centre Western General Hospital Crewe Road South Edinburgh United Kingdom EH4 2XU

Study participating centre Castle Hill Hospital Castle Road Cottingham United Kingdom HU16 5JQ

Study participating centre Basingstoke and North Hampshire Hospital

Aldermaston Road Basingstoke United Kingdom RG24 9NA

Study participating centre James Cook University Hospital

Marton Road Middlesborough United Kingdom TS4 3BW

Study participating centre Charing Cross Hospital

Fulham Palace Road London United Kingdom W6 8RF

Study participating centre Darent Valley Hospital

Darenth Wood Road Dartford United Kingdom DA2 8DA

Study participating centre Southampton General Hospital

Tremona Road Southampton United Kingdom SO16 6YD

Study participating centre St James's University Hospital

Beckett Street Leeds United Kingdom LS9 7TF

Study participating centre Morriston Hospital

Heol Maes Eglwys Swansea United Kingdom SA6 6NL

Study participating centre Royal Stoke University Hospital

Newcastle Road Stoke-on-Trent United Kingdom ST4 6QG

Study participating centre Salisbury District Hospital

Odstock Road Salisbury United Kingdom SP2 8BJ

Study participating centre Aberdeen Royal Infirmary

Foresterhill Aberdeen United Kingdom AB25 2ZN

Study participating centre Royal Derby Hospital

Uttoxeter Road Derby United Kingdom DE22 3NE

Study participating centre Barnet General Hospital

Wellhouse Lane Barnet United Kingdom EN5 3DJ

Study participating centre

Lister Hospital

Coreys Mill Lane Stevenage United Kingdom SG1 4AB

Sponsor information

Organisation

Newcastle upon Tyne Hospitals NHS Foundation Trust (UK)

Sponsor details

Northern Centre for Cancer Care Freeman Road High Heaton Newcastle upon Tyne England United Kingdom NE7 7DN

Sponsor type

Hospital/treatment centre

ROR

https://ror.org/05p40t847

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

Results and Publications

Publication and dissemination plan

The main trial results will be published in a peer-reviewed journal.

Intention to publish date

30/09/2022

Individual participant data (IPD) sharing plan

The current data sharing plans for the current study are unknown and will be made available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details protocol	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article		03/09/2019	05/10/2020	Yes	No
Results article		02/09/2022	27/09/2022	Yes	No
Results article		01/10/2022	28/10/2022	Yes	No
HRA research summary			28/06/2023	No	No