An investigation of proteins found in urine and blood to identify a combination of proteins which will detect bladder cancer

Submission date	Recruitment status	Prospectively registered
30/12/2014	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
06/02/2015	Completed	Results
Last Edited	Condition category	[] Individual participant data
31/10/2019	Cancer	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number ORECNI 80/04

Study information

Scientific Title

Evaluation and comparison of urine markers to investigate the development of a diagnostic index for bladder cancer

Study objectives

A combination of proteins secreted into urine and/or blood will be able to identify patients with bladder cancer

Ethics approval required

Old ethics approval format

Ethics approval(s)

Office for Research Ethics Committees Northern Ireland (80/04)

Study design

Observational case control study

Primary study design

Observational

Study type(s)

Diagnostic

Health condition(s) or problem(s) studied

Haematuria in bladder cancer

Interventions

This study recruited 200 patients who had reported blood in their urine to their doctor. Half of these patients had a final diagnosis of bladder cancer and the other half had other diagnoses. Blood and urine was taken from each patient. The levels of different proteins in the urine and blood were measured on anonymised blood and urine samples by a company called Randox Laboratories Ltd.

Intervention Type

Other

Primary outcome(s)

- 1. To compare specificities and sensitivities of the TCCB markers and generic cancer markers in predicting TCCB
- 3. To retain a bank of urine and blood samples if the need arise to assay further markers
- 4. To determine the best combination of factors for a prognostic index for TCCB

Key secondary outcome(s))

To compare biomarkers as prognostic indicators for TCCB

Completion date

31/10/2009

Eligibility

Kev inclusion criteria

- 1. Patients with haematuria who have undergone cystoscopy
- 2. Patients must be able to understand the study procedures and willing to give informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

- 1. Patients who have not had a flexible cytoscopy
- 2. Patients who did not present with haematuria
- 3. Patients with UTI destined to undergo TURB
- 4. Patients currently suffering from clinically evident alcoholism and or drug dependency

Date of first enrolment

13/11/2006

Date of final enrolment

22/04/2008

Locations

Countries of recruitment

United Kingdom

Northern Ireland

Study participating centre Belfast City Hospital

51 Lisburn Road Belfast United Kingdom BT9 7AB

Study participating centre Carigavon Area Hospital

68 Lurgan Road Portadown United Kingdom BT63 5QQ

Sponsor information

Organisation

Belfast City Hospital Trust

ROR

https://ror.org/02405mj67

Funder(s)

Funder type

Industry

Funder Name

Randox Laboratories Ltd (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration