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Immunosuppression for progressive membranous nephropathy

Submission date	Recruitment status	[_] Prospectively registered
25/10/2000	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
25/10/2000	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
01/02/2013	Urological and Genital Diseases	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
G9721265

Study information

Scientific Title

Study objectives

To test whether immunosuppression alters the natural history of progressive membranous glomerulonephritis and to compare two types of immunosuppressive treatment: cyclosporin and prednisolone/chlorambucil

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Multi-centre randomised controlled trial

Primary study design
Interventional

Study type(s)
Not Specified

Health condition(s) or problem(s) studied
Progressive membranous glomerulonephritis

Interventions
Randomisation will be between three groups:
1. One receiving supportive treatment only
2. One receiving 12 months treatment with cyclosporin
3. One receiving six months treatment with a combination of prednisolone and chlorambucil

Intervention Type
Drug

Phase
Not Specified

Drug/device/biological/vaccine name(s)
Cyclosporin and prednisolone/chlorambucil

Primary outcome(s)
Glomerular filtration rate with a further decline of 20% being an end-point

Key secondary outcome(s))
Proteinurea and adverse effects

Completion date
31/03/2009

Eligibility

Key inclusion criteria

1. Idiopathic membranous nephropathy with a 20% decline in glomerular filtration rate
2. Patients aged between 18 and 75

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 years

Sex
All

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
15/01/1999

Date of final enrolment
31/03/2009

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Academic Renal Unit
Bristol
United Kingdom
BS10 5NB

Sponsor information

Organisation
University of Bristol (UK)

ROR
https://ror.org/0524sp257

Funder(s)

Funder type
Research council

Funder Name
Medical Research Council (MRC) (UK)

Alternative Name(s)
Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory Council, MRC

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United Kingdom

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	02/03/2013		Yes	No