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Prospective pilot study: evaluation of a new dressing for patients treated in peritoneal dialysis

Submission date	Recruitment status	[_] Prospectively registered
23/08/2006	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
30/10/2006	Completed	[_] Results
Last Edited	**Condition category**	[_] Individual participant data
19/09/2007	Urological and Genital Diseases	[_] Record updated in last year

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
1445

Study information

Scientific Title

Study objectives

Evaluation of a new Peritoneal Dialysis (PD) dressing.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Submitted to CPP Grenoble (reference number: 06-GAMB-1), will receive verdict on 20th September 2006.

Primary study design
Observational

Study design
Monocentric, open, comparative study

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Chronic renal failure

Interventions
Evaluation of dressing advantages via questionnaires

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
Dressing advantages are evaluated via questionnaires, which have to be filled in at specific time schedule. Comfort and safety evaluation will be performed by the patient at the end of each period of product use (end of PD Immo use and end of usual dressing use).

Key secondary outcome(s)
Handling is also evaluated via the patient questionnaire.
The incidence of Adverse Events (AE) will be followed during the study period.

Completion date
25/11/2006

Eligibility

Key inclusion criteria
1. Patients with well healed exit site
2. Patients treated for at least three months in peritoneal dialysis irrespective of the treatment mode

3. Patients with healthy skin judged by investigator
4. Patients aged 18 years and older
5. Patients having signed a written consent (informed consent) to participate in the study

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Adult

Lower age limit
18 Years

Sex
All

Key exclusion criteria
1. Patients treated in hemodialysis
2. Patients with diagnosed abdominal pain not related with PD-solution
3. Peritonitis within one month prior to the study
4. Presence of exit-site, tunnel infection
5. Patients with ongoing peritonitis
6. Patients participating in other studies during the period of this study
7. Patients practising swimming
8. Patients under guardianship
9. Pregnancy, lactation

Date of first enrolment
25/09/2006

Date of final enrolment
25/11/2006

Locations

Countries of recruitment
France

Study participating centre
AGDUC
Grenoble
France
38 700

Sponsor information

Organisation
Gambro Industries (France)

ROR
https://ror.org/01mgtdr23

Funder(s)

Funder type
Industry

Funder Name
Gambro Industries (France)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration