

Patient Details:

Patient's full name:

Sex: Male Female

Date of birth: / /

Hospital number:

Responsible clinician:

Hospital:

PD REHAB trial number:

AE Description:

Date event started:/...../.....

Date event ceased:/...../.....

Outcome: Fatal Recovered Continuing

Details of adverse event:

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Did the event require hospitalisation? No Yes No of days

Reason why you consider event to be intervention related:

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Name of person reporting:

Telephone Number:

Date:/...../.....