| Patient Study Number: | Patient Initials: |
|-------------------------|-------------------|
| Patient DOB: | Site ID: |
| Principal Investigator: | Version: |

[Delete this line, then print on Trust headed paper]

PATIENT CONSENT FORM

Where witnessed consent is required please use the Witnessed Consent Form



Pressure UlceR Programme Of ReSEarch

Pressure Ulcer Risk Assessment Framework (PURAF) Field Test 1

Patient initial after each question

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1. I confirm that I have read and understand the information sheet dated 23.05.2012 (version 2) for the above study. I have had the opportunity to ask questions and have had these answered satisfactorily.

2. I understand that my participation is voluntary and that I am free to withdraw at any time without my medical care or legal rights being affected.

3. I understand that if I withdraw from the above study, the data already collected from me will be used in analysing the results of the study unless I specifically withdraw consent for this.

study report or other publication. This information will be confidentially destroyed at the end of the study.

6. I understand that information and results arising from this study may be used to develop new research.

7. I understand that a copy of this consent form will be passed to the Clinical Trials Research Unit, University of Leeds.

9. I agree to take part in the study.

Name of Patient

Date

Signature

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I have given written information and a verbal explanation to the person named above who has freely given their consent to participate.

| Name of Person taking consent Date | Signature |
|------------------------------------|-----------|
|------------------------------------|-----------|

1 copy for patient, 1 for patient records, 1 for CTRU; original stored in Investigator Site File