PATIENT CONSENT FORM - PART III
(PET study)

Study title: A multicentre randomised clinical trial comparing rituximab with CHOP given 14 days and rituximab with CHOP given every 21 days for the treatment of patients with newly diagnosed diffuse large B cell non-Hodgkin's lymphoma

Substudy title: Blinded evaluation of prognostic value of FDG-PET after 2 cycles of chemotherapy in Diffuse Large B-cell Non-Hodgkin's Lymphoma

PATIENT CONSENT FORM PART III* (please read carefully)

Name of Researcher: _____

Please initial

I confirm that I have read and understand the information dated 18/07/08 (version 3.0) for the above study and have had an opportunity to ask questions.	
I understand that my participation in this study is voluntary and that I am free to withdraw at any time without giving a reason, without my medical care or legal rights being affected.	
I understand that the PET scan results will be anonymised, stored and analysed separately for the whole study. I understand that this will be linked to the trial data through the unique trial number and no other personal data will be held.	
I agree to take part in the PET study.	

_____	_____	_____
Name of Patient	Date	Signature
_____	_____	_____
Name of person taking consent (if different from researcher)	Date	Signature
_____	_____	_____
Name of Researcher	Date	Signature

* Three copies required: one each for the patient, researcher and hospital case notes