

Patient Registration Form

Centre Number	Patient Number	
Hospital Name		
NOTE: Fill only section 1 – 3 if samples are be	eing sent to UK Biocentre with the sample information sheets for the first	
	ng sent to UK Biocentre do not require this form to be filled. ent sample is preferable, but if not possible any other time point when the patient	
has adequate neutr	rophil count (>1.0 x 10 ⁹ /L) is acceptable	
1 - Patient Information		
Sex Initials	Male Female Male	1.01
Date of Birth		1.02
		1.03
Hospital Number		1.04
NHS/CHI Number		1.05
Postcode		1.06
Ethnicity		1.07
Is the patient on a clinical trial?	Yes No	1.08
Trial name	Trial number	1.09
Trial name other		1.10
2 - Consent		
Who is giving consent? Has consent been given for the following statemen	Patient Parent Guardian N	2.01
Samples to be stored in the VIVO Biobank?	<u> </u>	2.03
Samples to be used in research worldwide?	Sample originally taken	2.04
DNA testing to be performed on samples?Samples to be used in animal research?	using deferred consent	2.05
Additional comments		2.07
Date consent obtained	1 1	2.08
I confirm consent has been obtained for the	ne VIVO Biobank and a copy retained in the patient's medical notes.	
Name		2.09
Job Title		2.10
Date	1 1	2.11
Person confirming consent		2.12
3 – Diagnosis and Treatment Information		
Diagnosis (please select most appropriate, leave b	olank if diagnosis not yet known & add other diagnoses in other diagnosis b	ox)
Main Category - Main Diagnosis	- Sub-Type - Anatomical Location - Additional Information	
-		3.01
E.g. Bone - Osteosarcoma	- High Risk - Femur - MYCN Amplified	
Other diagnosis (not on drop-down)		3.02
Date of diagnosis	/ / /	3.03
Stage (not leukaemia)		3.04
Grading (not leukaemia)		3.05
Immunophenotype (Leukaemia only)		3.06
Other immunophenotype (do not abbreviate)		3.07
Genetic information (if available)		3.08
GTAB Number (if available)		3.09
Whole Genome Sequencing (WGS) done for this par	tient? Y N	3.10