

KRCC MOLECULAR PATHOLOGY TEST REQUEST FORM



Please send all samples to:

Synnovis Analytics Molecular Pathology Laboratory c/o Central Specimen Reception Blood Sciences Laboratory Ground Floor Bessemer Wing King's College Hospital, Denmark Hill London SE5 9RS

Tel: 020 3299 2265 Email: kch-tr.PND@nhs.net

All fields are mandatory. Illegible, unclear or incomplete forms will result in delays or rejection.

CONSENT STATEMENT: It is the referring clinician's responsibility to ensure that the patient/carer knows the purpose of the test and that the sample may be stored for future diagnostic testing. In signing this form the clinician has obtained consent for testing, storage and for the use of this sample and the information gathered from it to be shared with members of the donor's family through their health professionals (if appropriate). The patient should be advised that the sample may be used anonymously for quality assurance and training purposes. If the patient does not wish information to be shared please write this clearly in the clinical summary box.

ssurance and training purposes.	If the patie	ent does	not wish i	informa	ation to be	shai	red pleas	e wr	rite t	this clearly in th	ne clinical summary box.	, , , , , , , , , , , , , , , , , , , ,	
PATIENT DEMOGRAPHICS								PATIENT ETHNICITY					
First name:									٧	Vhite:	British ☐ Irish ☐ Any Oth	er White Background □	
Last name:							M		∕lixed:	White And Black Caribbean ☐ White And Black African ☐ White And Asian ☐			
DOB:	Gender: Male ☐ Female ☐ O					her 🗆	er Asian or		Asian or	Any Other Mixed Background ☐ Indian ☐ Pakistani ☐ Bangladeshi ☐			
NHS number:								Asian British:		Any Other Asian Background □			
Hospital no:	Family ref no:							Black or Caribbean ☐ African ☐ Any O Black British: Background ☐		ny Other Black			
Postcode:	Antenatal: Yes □ No □								Other Ethnic Groups: Chinese □ Any Other Ethnic Group □ (please specify:		· ·		
Non-NHSE funded (please attac	g details): □							_					
PATHOLOGY RESULTS Haematology indices									s	AMPLE REQUIF	REMENTS		
Iron / liver parameters Hb		HbF %						F	For haemoglobinopathy investigation: 2 x		2 x 4 ml EDTA blood		
Serum Iron	RBC			HbA2	%				-		ults (all other tests):	4 ml EDTA blood	
Serum TSat	MCV			Hb var	riant %				-	, ,		1 ml EDTA blood	
Serum Bilirubin	MCH			Absolu Reticu					-		Generation Sequencing:	3-5μg genomic DNA	
Serum	Platelets				locyte %				-	As DNA for all ot	ther tests:	1-5μg genomic DNA	
Ferritin Blood Film comments:	erritin			rictica	mocyte 70				С	ate sample col	lected:		
Reason for referral / fam	!!! . . . !! .		1 - 41-		ent affec		ı			·		_	
Molecular Tests - For Haemoglobinopathy referrals FBC and HPLC results MU. R93 Hb variant identification R93 Haemoglobinopathy investigations R93 Alpha thalassaemia R93 Beta thalassaemia Other (please state gene):							Pe provided R176 Gilbert's genotyping (TAs/6/7/8 repeat) R95 Hereditary haemochromatosis (HFE - C282Y and H63D variants) R191 Alpha-1-antitrypsin genotype (S and Z alleles) Thrombophilia genetic screen (please tick all that apply): FVL PT MTHFR R40/242/244 Familial variant testing ease state familial variant):						
For further details of each test please refer to the <u>Synnovis website</u> .													
Next Generation Sequencing - I	Please selec	t which	panel(s) a	re requ	uired								
 □ R91 Cytopenia (NOT Fanconi anaemia) □ Neutropenia □ Diamond-Blackfan anaemia □ Inherited bone marrow failure 				[□ R96 Iron metabolism d □ R168 Porphyria □ R169 Acute intermitte □ R170 Variegate porph 				mittent porphyria		☐ R92 Rare Anaemia ☐ Membranopathy ☐ Enzymopathy ☐ Haemoglobinopathy		
□ HLH □ R366 □ R405 □ R406 □					□ R366 Inl □ R405 He □ R406 Th	7 Inherited predisposition to AML 6 Inherited predisposition to ALL 5 Hereditary erythrocytosis 6 Thrombocythaemia 3 Sitosterolaemia				n to ALL	☐ Congenital dyserythropoietic anaemia ☐ Diamond-Blackfan anaemia ☐ Sideroblastic anaemia ☐ Haemolytic anaemia ☐ Sitosterolaemia ☐ Magaloblastic anaemia		
or details of genes in each subpanel, please refer to the <u>Synnovis website</u>											☐ Megaloblastic ana	iemia	
CLINICIAN DETAILS													
Requesting clinician / consultant								Responsible clinician / consultant (if different)					
Name: Hospital & department:								Name: Hospital & department:					
NHS email: Phone:							NHS en	IHS email: hone:					

Note: Please ensure the latest version of this request form is used, found on our website: www.southeastgenomics.nhs.uk