

NATIONAL HAEMOGLOBINOPATHY REFERENCE LABORATORY
REQUEST FORM: OXFORD RED CELL PANEL (ORCP)

PATIENT DETAILS (please fill in or attach addressograph)

Surname:	First Name:
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Gender:	D.O.B:	Post Code:	Patient reference No:
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NHS No:	Ethnicity:
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SUSPECTED DIAGNOSIS:

This is the PROBAND: <input type="checkbox"/>	This is a FAMILY MEMBER: <input type="checkbox"/> Proband name: _____ Relationship to Proband: _____ This individual is: AFFECTED <input type="checkbox"/> UNAFFECTED <input type="checkbox"/>	Proband Date of Birth: _____
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REQUESTER DETAILS

Requesting Clinician:	Contact details (email & phone no):
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Address for report:	Address for invoice:
	Same as for report: <input type="checkbox"/>

TO AID INTERPRETATION OF RESULTS PLEASE PROVIDE US WITH THE FOLLOWING DETAILS:

RBC (red blood cell count, x10 ¹² /L)	WBC (white blood cell count: x10 ⁹ /L)
HGB (haemoglobin, g/L)	NEUTROPHILS (x10 ⁹ /L)
MCV (mean corpuscular volume, fl)	LYMPHOCYTES (x10 ⁹ /L)
MCH (mean corpuscular haemoglobin, pg)	BILIRUBIN (umol/L)
RETICULOCYTES (x10 ¹² /L & %)	SGOT/AST (Serum glutamic oxaloacetic transaminase, IU/L)
PLATELETS (x10 ⁹ /L)	SGPT/ALT (Serum glutamic pyruvic transaminase, IU/L)
FERRITIN (mlg/L)	LDH (lactate dehydrogenase: IU/L)
TRANSFERRIN (%)	DAT (Direct Antiglobulin Test/Coombs Test)
	POS/NEG/ not tested

BLOOD SMEAR RESULTS:

BONE MARROW SMEAR RESULTS (please also supply date sample analysed):

TO AID INTERPRETATION OF RESULTS PLEASE PROVIDE THE FOLLOWING CLINICAL INFORMATION:

ANAEMIA ONSET AND TYPE (circle all that apply)	Fetus/Infant/Child/Adult Onset Acute Chronic Transfusion dependent	JAUNDICE (circle all that apply)	Prolonged neonatal Intermittent Chronic
SPLENOMEGALY	Y / N	HEPATOMEGALY	Y / N
PANCREATIC INSUFFICIENCY	Y / N	GALLSTONES	Y / N
DYSMORPHIC FACIES	Y / N	SKELETAL, LIMB OR DIGIT ABNORMALITIES	Y / N
DEVELOPMENTAL DELAY /LEARNING DIFFICULTIES	Y / N	ANY OTHER ORGAN ABNORMALITIES	Y / N
SHORT STATURE/ FAILURE TO THRIVE	Y / N	FREQUENT INFECTIONS	Y / N
FAMILY HISTORY (provide relationship to patient)	Anaemia Other relevant features:	CONSANGUINITY	Y / N
ANY OTHER RELEVANT DETAILS: eg pertinent treatment details, results of other investigations such as enzyme levels, other non-haematological finding, history of bone marrow transplant, etc)			

Genes included on the panel are as follows:

ABCB7, ALAS2, ALDOA, ANK1, C15orf41, CDAN1, ENO1, EPB41, EPB42, G6PD, GATA1, GATA2, GCLC, GPI, GPX1, GSR, GSS, HK1, KIF23, KLF1, LPIN2, NT5C3A, PFKM, PGK1, PIEZO1, PKLR, RHAG, RPL5, RPL9, RPL11, RPL26, RPL27, RPL35A, RPS7, RPS10, RPS17, RPS19, RPS24, RPS26, RPS27, RPS29, SBDS, SEC23B, SLC2A1, SLC4A1, SLC11A2, SLC25A38, SPTA1, SPTB, TMPRSS6, TPI1.

Variants identified during this analysis will be reported if there is reasonable evidence supporting clinical significance.

This may include mutations associated with non-anaemia phenotypes or carrier status with reproductive implications.

Sample type (circle): DNA or EDTA blood

Date sampled:.....

Please send samples to the address provided towards the foot of the form.

Consent has been obtained for the DNA/RNA of this sample to be stored and used in research/development projects that have been granted ethical approval (please delete as appropriate):

Yes / No **Signed**.....

Clinician:..... **Date**.....

Labelling Standards:

Please label samples with the **patient's**: Name, Date Of Birth, NHS Number (or Hospital Number for non-UK referrals). A minimum of 2 identifiers must be provided or the sample cannot be accepted for testing.

Address: Molecular Haematology, Level 4, John Radcliffe Hospital, Headington, Oxford, OX3 9DU

Sample reception: 01865 572769 Sec: 01865 572826 Fax: 01865 572775

Email : oxford.molecularhaem@nhs.net Web site: <http://www.oxford-translational-molecular-diagnostics.org.uk/>

Further copies of this form can be downloaded from: www.oxford-translational-molecular-diagnostics.org.uk/