

INSTITUT PERUBATAN DAN PERGIGIAN TERMAJU (IPPT) **UNIVERSITI SAINS MALAYSIA (USM) Advanced Diagnostic Laboratory**





No. Tel: 04- 5622293

MOLECULAR DIAGNOSTIC REQUEST FORM			
PATIENT INFORMATION:	For Laboratory Use Only		
Name:	_		
I/C No.:			
Age: PID: Chinasa	1		
Gender: Male Female Ethnicity: Malay Chinese If female; Pregnant? Indian Others:			
Yes, Weeks: No			
Ward / Clinic:			
Date of Sampling: Date Sent:	TYPE OF TEST REQUEST:		
Clinical Diagnosis:	Sex-determining Y Region (SRY) gene Southeast Asian Ovalocytosis (SAO)*		
Parental consanguinity: YES NO	Microdeletion Y (MDY) Screening		
	Molecular Alpha Thalassaemia (MAT)*		
CLINICAL SUMMARY/ FAMILY HISTORY (include family pedigree):	DNA extraction		
	Indication of test: NEW CASE		
	CASCADE SCREENING		
	Information of index case		
	Name:		
	IC no:		
	Lab no: Result:		
	OTHERS (please specify)		
	Official Stamp of Requesting Doctor:		
	Name, Signature & Date		
	Please provide contact detail for return of result		
	(Name, Organization Email, Contact No):		

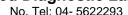
*Note:

- All paeditarics (≤12y/o) samples must be accompanied with parents' samples.
- SAO test request must be accompanied with a copy of recent FBP (< 3 months) result.

 MAT test request must be accompanied with a copy of recent FBP (< 3 months) result and Hb Analysis result.



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INFORMED CONSENT FOR DNA TESTS

Patient Name:	Patient ID:		
I understand the following:			
This test is specific for			
- A BOSITIVE regult is an indication that I may be predigneded to	o or have the enecific discose	or condition	Further tection

- A POSITIVE result is an indication that I may be predisposed to or have the specific disease, or condition. Further testing
 may be needed to confirm the diagnosis.
- There is a chance that I will have this genetic condition but that the genetic test results will be **NEGATIVE**. Due to limitations in the technology and incomplete knowledge of genes, some changes in DNA or protein products that cause disease, may not be detected by the test.
- There may be a possibility that the laboratory findings will be **UNINTERPRETABLE** or of unknown significance. In rare circumstances, findings may be suggestive of a condition different than the diagnosis that was originally considered.
- 1. The potential benefit of this test is **to confirm the diagnosis of the condition** and to determine which other family members may be carriers or have increased risk of having the defective gene.
- 2. Erroneous results and incorrect interpretation may occur because of rare variation in the DNA of the individual, rare technical error, mis-incorporation of DNA bases by the enzyme used to perform the test, sample misidentification, sample contamination, primer site mutations and general laboratory error.
- 3. Accurate interpretation of the DNA test result depends on **correct information** about the **clinical diagnosis** and about the **biological relationships** within the family.
- 4. DNA testing may reveal **non-paternity**, meaning that the stated father is not the biological father.
- 5. The tests offered are considered to be **the best available at this time**. If technology improves and more mutations (gene defects) are detectable in future, I authorize the Laboratory to re-analyse, at the Laboratory's option, any remaining DNA for the same disease without being informed. If the sample is insufficient, my doctor may ask me for a fresh sample. There may be additional fees for such tests.
- 6. In order to help me understand the test results, the results will be reported to me only through a doctor or genetic counsellor.
- 7. This report shall be used **ONLY** for clinical interpretation.
- 8. This report shall **NOT** be used for any forensic purposes or is **NOT VALID** for forensic interpretation.
- 9. This report shall NOT be used in any courts of law or in legal matters and is NOT VALID for legal interpretation.
- 10. Residual unused sample and their derivative **MIGHT BE** used for educational and research purposes subject to rules and regulation of USM.

For prenatal testing, the following also apply:

- 1. This DNA test will determine the status of the fetus for this disease ONLY.
- 2. Besides rare DNA variation and the technical error, erroneous results may also arise from maternal contamination of the fetal sample.

WRITTEN CONSENT:

- 1. A biologic specimen (blood, tissue, amniotic fluid or chorionic villi) will be collected for DNA tests for the above condition.
- 2. After DNA testing is completed, a small amount of my DNA may be made anonymous and used for medical education, quality control or research. Since the samples have been anonymised, any research results obtained cannot be reported to me. I understand that any biologic specimens obtained for the purpose of this genetic testing become the exclusive property of AMDI's laboratories. After the specific test(s) requested have been completed, the laboratory may dispose, retain, or use the de-identified specimen(s) for test validation or education; i.e publication into journals. I understand that my identity will be protected.
- DNA results are strictly CONFIDENTIAL and will not be released to anyone including my relatives/ other than my doctors without my consent.
- 4. Some individuals who have chosen to have predictive DNA testing and been found to carry the gene leading to the disease have experienced discrimination (insurance, employment and social).

To be completed by the:		
PATIENT/ PARENTS/ LEGAL GUARDIAN	DOCTOR/ COUNSELLOR	
I have read and received a copy of this consent form. I understand the information provided in this document and I have had the opportunity to ask questions about testing, the procedure and the associated risks, benefits and limitations. I agree to have genetic testing and accept the risks and limitations.	I have fully explained the nature of the requested test(s) to the patient/ parent/ legal guardian	
SIGNATURE : NAME & IC NO.:	SIGNATURE & OFFICIAL STAMP: NAME:	
DATE:	DATE:	