

Document No.	HAEM-W-844	Version No. Issue Date	2 20/10/2020
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## Haematological Malignancy Diagnostic Service

# Myeloid NGS Request Form

HMDS, Western General Hospital, Haematology/  
Biochemistry Combined Reception  
Immunophenotyping Laboratory  
Crewe Road, Edinburgh, EH4 2XUT

el. 0131-537-2374 email: [hmds.lothian@nhslothian.scot.nhs.uk](mailto:hmds.lothian@nhslothian.scot.nhs.uk)  
website: [www.edinburghlabmed.co.uk/Specialities/GeneScience/HMDS](http://www.edinburghlabmed.co.uk/Specialities/GeneScience/HMDS)

### REQUESTOR DETAILS:

Requesting Consultant/GP:			
Hospital/Site			
Contact telephone/page:		Destination for report:	

### PATIENT DETAILS:

Surname:		Forename:		
CHI (or Date of Birth):		Gender:	M / F	
Address :				
Specimen Ref. No.		Specimen type:	PB <input type="checkbox"/>	BM <input type="checkbox"/>
Sample collection date/time				

### Please tick a reason for testing:

- Intensive treatment-eligible AML patient
- Transplant eligible MDS patient
- Transplant eligible patient with PMF
- For differentiation of hypoplastic MDS/aplastic anaemia
- Atypical MPN (triple negative PMF phenotype, MDS/MPN overlap)
- Selected relapsed AML patient to provide therapeutic information
- DNA and RNA storage
- Other (please provide details)

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## Address for delivery:

HMDS, Western General  
Hospital,  
Haematology/Biochemistry  
Combined Reception  
Immunophenotyping  
Laboratory  
Crewe Road  
Edinburgh  
EH4 2XU

**Phone:** 0131-537-2374

**Internal:** 32374

Arrange for immediate transport to the laboratory.

It is your responsibility to ensure that samples are packaged to comply with the European Agreement concerning the International Carriage of Dangerous Goods by Road (ADR, 2017) at [https://www.unece.org/fileadmin/DAM/trans/danger/publi/adr/adr2017/ADR2017e\\_web.pdf](https://www.unece.org/fileadmin/DAM/trans/danger/publi/adr/adr2017/ADR2017e_web.pdf)

\*Fold along the dotted lines and place in sample bag with address facing outwards\*

## Consent

In accordance with the requirements of the Human Tissue (Scotland) Act 2006, **it is the responsibility of the referring clinician to ensure that appropriate informed consent has been obtained before any testing is undertaken.** The laboratory must be informed of any restrictions to this consent (e.g. storage of samples).

Unless otherwise informed, the laboratory assumes that all appropriate consent has been obtained from the patient for the tests requested and for storage of the derived DNA and RNA for future use both in assisting further testing (if required) and in the development of future diagnostic tests. If in doubt, contact a member of the HMDS team to discuss.

**Incomplete or illegible forms may cause delay or rejection of samples**

FOR LABORATORY USE ONLY			
Received by		Volume received	
Date/time received		Lab number	

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