



Accredited Medical Laboratory  
Reference No: 1934

Central Manchester and Manchester  
Children's University Hospitals



NHS Trust

**Contact**

**Molecular  
Genetics  
Genetic Medicine  
6th Floor  
St Mary's Hospital  
Oxford Road  
Manchester  
M13 9WL**

Tel: 0161 276 6122  
Fax: 0161 276 6606

**Samples:**

- Blood in EDTA
- Adults 10ml
- Children 5-10ml
- Neonates 1ml (minimum)
- Store at 4°C,
- Do not freeze
- Send by first class post
- Tumour samples – please contact laboratory for advice

**Request card available from**

[www.mangen.org.uk/](http://www.mangen.org.uk/)

**Neurofibromatosis type 2 (NF2) (OMIM#101000)**

**NF2 mutation screening (NCG referrals) – clinical pro-forma**

**Explanatory Notes:**

*This form should accompany all requests for mutation screening of the NF2 gene. It is not necessary to complete this form for follow on tests where a mutation has already been identified within a family. Please ensure the information you provide is as complete possible as this information will be used to determine eligibility for NCG funded testing and audit the effectiveness of the service. The referring clinician will be contacted for further information if criteria for NCG testing appear not to have been met.*

Patient Details	
Forename	<input type="text"/>
Surname	<input type="text"/>
Date of Birth <i>(dd/mm/yyyy)</i>	<input type="text"/>
NHS Number	<input type="text"/>
Address	<input type="text"/>
Local Ref No.	<input type="text"/>

Referrer Information	
Name	<input type="text"/>
Specialty	<input type="text"/>
Address	<input type="text"/>
Date of completion <i>(dd/mm/yyyy)</i>	<input type="text"/>

## Contact

Molecular  
Genetics  
Genetic Medicine  
6th Floor  
St Mary's Hospital  
Oxford Road  
Manchester  
M13 9WL

Tel: 0161 276 6122  
Fax: 0161 276 6606

### Samples:

- Blood in EDTA
- Adults 10ml
- Children 5-10ml
- Neonates 1ml (minimum)
- Store at 4°C,
- Do not freeze
- Send by first class post
- Tumour samples – please contact laboratory for advice

Request card  
available from

[www.mangen.org.uk/](http://www.mangen.org.uk/)

## Page 2 of 2

Clinical Summary	
Has the patient undergone NF2 mutation scanning previously?	Yes – Lymphocyte DNA <input type="checkbox"/>
	Yes – Tumour DNA <input type="checkbox"/>
	No <input type="checkbox"/>
	<i>If yes please provide details extent of screening e.g. SSCP of coding DNA &amp; MLPA and lab</i> <input type="text"/>
Does the patient have any of the following?	
Vestibular schwannoma	Yes <input type="checkbox"/> No <input type="checkbox"/>
	<i>If yes</i> bilateral <input type="checkbox"/> unilateral <input type="checkbox"/>
	age of onset <input type="text"/>
Meningioma	Yes <input type="checkbox"/> No <input type="checkbox"/>
	<i>If yes please give number</i> <input type="text"/>
	age of onset <input type="text"/>
Other Schwannoma	Yes <input type="checkbox"/> No <input type="checkbox"/>
	<i>If yes please indicate the site and number</i> <input type="text"/>
	age of onset <input type="text"/>
Juvenile posterior subcapsular lenticular opacities/ juvenile cortical cataract	Yes <input type="checkbox"/> No <input type="checkbox"/>
Family History	Yes <input type="checkbox"/> No <input type="checkbox"/>
Mononeuropathy	Yes <input type="checkbox"/> No <input type="checkbox"/>
Ependymoma	Yes <input type="checkbox"/> No <input type="checkbox"/>
	<i>If yes please give number</i> <input type="text"/>
	age of onset <input type="text"/>

**Thank you for your time and co-operation**