## Healthlink: Information for GPs or consultants referring patients to the Department of Clinical Genetics (DCG), Children's Health Ireland (CHI), at Crumlin, D12 N512

General points to aid referral	We accept referrals of adults and children
	Include patient's mobile number
	When referring a couple or more than one family member
	together, include names and dobs of each individual
	If family member previously attended DCG include Genetic
	Family reference number (often a 5-digit number, PEDxxxxx)
	Enclose copy of genetic test(s) as appropriate
	Enclose copy of post mortem results as appropriate
REFERRAL REASON	INFORMATION REQUIRED
Referring a pregnant patient for	Do <b>NOT</b> send the referral via Healthlink
prenatal advice	Email the referral as an attachment to <u>prenatal.GC@olchc.ie</u>
	Prenatal referrals must be marked URGENT
	Any problems with above, contact DCG at 01 409 6739 or the
	Consultant Clinical Geneticist on-call via switchboard 01 409
	6100
	Always include the patient's LMP
Referral for <i>predictive</i> genetic	Include our family reference/pedigree number
testing (general and cancer)	If family member seen elsewhere enclose copy of their genetic
(Referring an unaffected	result or details (name/dob/genetic centre attended) so that
individual with a genetically	report can be obtained. We cannot offer predictive testing
confirmed condition in a family	without a copy of the genetic result
member)	State the relationship between your patient and the affected
	family member with the genetically confirmed condition
Referral for diagnostic genetic	Briefly describe the phenotype
testing (general)	List any previous diagnoses
	Enclose copies of any genetic tests and other relevant tests
	Outline the family history
Referring a child with	Please refer to a paediatrician in the first instance for basic
developmental delay or autism	genetic investigations.
	The paediatrician will refer on to genetics if necessary
Referral for diagnostic genetic	State the age at diagnosis of any cancer
testing (cancer)	Outline the family history of cancer stating the relative's age at
	cancer diagnosis, type of cancer, whether alive or dead and
Fruther information link	relationship (including side of family) to your patient
Further information link	a family history questionnaire (FHQ) completed by the patient,
	may be enclosed with the referral, which is available at:
	https://www.olchc.ie/Services/Departments-A-Z/Department-of-
	Clinical-Genetics/Information-Leaflets-Forms/Genetics-Family-
Deferred of confloated nations	History-Cancer-Questionnaire.pdf
Referral of <i>unaffected</i> patient concerned about risk of cancer	Outline the family history of cancer stating the relatives ages at
due to family history	cancer diagnosis, type of cancer, whether alive or dead and relationship (including side of family) to your patient
due to failing history	
	Please note: only 5-10% of cancers are genetic (early age of onset, multiple cancers in an individual, same cancer in family,
Further information link	cancers associated with a cancer syndrome, unusual/rare cancers
	or cancers coming down the generations may suggest a genetic
	cause)
	There needs to be a living affected family member to initiate
	testing in (provided testing criteria are met when we assess the
	family history)
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Referring a patient for advice regarding a genetic test result Carrier testing for Cystic Fibrosis	<ul> <li>We may not be able to offer genetic testing but may offer surveillance advice.</li> <li>FHQ completed by the patient may be enclosed with the referral, see link above. We do not accept FHQ without a referral attached.</li> <li>Referrals will be rejected if a copy of the genetic test report is not attached</li> <li>This is done directly by GPs.</li> <li>If you wish to arrange carrier testing for Cystic fibrosis for your patient, further information can be found here: <a href="https://www.olchc.ie/Services/Departments-A-Z/Department-of-Clinical-Genetics/Information-Leaflets-Forms/Cystic-Fibrosis-">https://www.olchc.ie/Services/Departments-A-Z/Department-of-Clinical-Genetics/Information-Leaflets-Forms/Cystic-Fibrosis-</a></li> </ul>
	<u>Genetic-Testing-Questionnaire.pdf</u>
Diagnostic or carrier testing for Haemochromatosis	<ul> <li>We do not accept referrals for haemochromatosis.</li> <li>Hemochromatosis is a very common disorder in Ireland and only a small percentage of individuals who carry specific variants in both copies of the gene develop the disease.</li> <li>Further information can be found in our handbook in lieu of appointment. Please see link Rare Disease Handbook FINAL</li> </ul>
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Diagnostic and carrier testing for haematological conditions	<ul> <li>Please refer to a haematologist, not a geneticist, particularly for common haematological conditions e.g. sickle cell disease, thalassaemia.</li> </ul>
Referring a patient for advice regarding Direct To Consumer (DTC) testing	<ul> <li>We do not recommend that the public organise DTC testing for diagnostic purposes but recognise that this increasingly occurs</li> <li>Advice should be sought from the genetic counsellors employed by the DTC company</li> <li>Patients can be referred when common genetic conditions are identified (e.g. cystic fibrosis or a known cancer predisposition gene).</li> <li>Your patient should be aware that we may need to repeat testing in a clinically accredited laboratory following appropriate counselling</li> <li>Most other DCT results have limited clinical utility and we are likely to reject the referral and let you know why. We will be unable to accept referrals for carriers of very rare conditions.</li> <li>BEWARE the patient with a strong personal or family history suggestive of a genetic condition who claims to a have a 'normal' DTC test result, such patients may warrant genetics referral</li> </ul>

If you have queries concerning any of the above or a question not covered here please telephone 01 409 6739 or email <u>clinical.genetics@olchc.ie</u> Do not use this email for referrals