



Diagnostic Service Facility (DSF)

Laboratory of Neurogenetics • Institute Born-Bunge
University of Antwerp - CDE • Building V - Parking P4
Universiteitsplein 1 • B-2610 Antwerp
Tel.: +32 3 265 1020 • Fax: +32 3 265 1037 • E-mail: DSF@bornbunge.be
Website: <http://www.molgen.ua.ac.be/DNAdiagnostics>

DSF director: Christine Van Broeckhoven, PhD DSc

DSF REQUEST FORM GENETIC TESTING

Please mention name, first name and date of birth on the sample tubes. Only clearly identified blood samples and a corresponding request form are accepted. It is the responsibility of the physician to complete and sign this request form. It is important to provide sufficient clinical data to guide the genetic testing.

<p>PATIENT DATA (fill in completely):</p> <p>Family name + first name:</p> <p>Date of birth:</p> <p>Gender:</p> <p>Address:</p> <p>.....</p> <p>Your reference number:</p> <p>.....</p> <p>Residual material will be used for genetic scientific research. Please check the box if the patient does NOT agree <input type="checkbox"/></p>	<p>PHYSICIAN DATA (fill in completely):</p> <p>Family name + first name:</p> <p>Hospital/Institute:</p> <p>.....</p> <p>Address:</p> <p>.....</p> <p>Phone:</p> <p>E-mail¹:</p> <p>Request date:</p> <p>Signature:</p> <p>Copy of results to:</p>
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¹Please provide an e-mail address for all further communications and receiving results

SAMPLE

<input type="checkbox"/> Blood: date taken: :	<input type="checkbox"/> Other material ² , type:
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²Other material can only be accepted after consultation

REQUEST

<input type="checkbox"/> Confirmation/exclusion ³ clinical diagnosis	<input type="checkbox"/> Presymptomatic testing
<input type="checkbox"/> Carrier testing	

³Circle correct choice

CLINICAL INFORMATION

Clinical diagnosis:
Clinical information:
.....
.....
.....
Clinical report enclosed: Yes / No / Will be provided



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Name patient + date of birth:

FAMILY AND PEDIGREE INFORMATION

Have you already sent material to our laboratory: yes / no

If yes: Name: First name:
Date of birth: Relationship:

Gene/mutation known in family: yes / no

If yes: Mutation details:

Pedigree:

- isolated/sporadic patient: no (known) family history
- familial: positive family history
- inherited: (autosomal) dominant / (autosomal) recessive / X-linked (please circle the correct choice)
- parents are related: yes / no
If yes, specify:

Family history:

.....

.....

.....

.....

.....

.....

Draw pedigree and indicate the person(s) to be tested with an arrow:



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Name patient + date of birth:

GENETIC TESTS FOR NEURODEGENERATIVE BRAIN DISEASES

<input type="checkbox"/> Dementia <input type="checkbox"/> APOE <input type="checkbox"/> Alzheimer dementia <input type="checkbox"/> PSEN1 <input type="checkbox"/> APP <input type="checkbox"/> APP duplication <input type="checkbox"/> PSEN2 <input type="checkbox"/> Familial Alzheimer dementia <input type="checkbox"/> PSEN1 <input type="checkbox"/> APP <input type="checkbox"/> APP duplication <input type="checkbox"/> PSEN2 <input type="checkbox"/> MAPT <input type="checkbox"/> PRNP <input type="checkbox"/> Cerebral amyloid angiopathy <input type="checkbox"/> APP <input type="checkbox"/> APP duplication <input type="checkbox"/> APOE <input type="checkbox"/> Frontotemporal lobar degeneration <input type="checkbox"/> C9orf72 <input type="checkbox"/> GRN <input type="checkbox"/> GRN dosage <input type="checkbox"/> MAPT <input type="checkbox"/> MAPT dosage <input type="checkbox"/> VCP <input type="checkbox"/> CHMP2B	<input type="checkbox"/> Parkinson disease <input type="checkbox"/> LRRK2 <input type="checkbox"/> PARK2 <input type="checkbox"/> SNCA <input type="checkbox"/> PARK2 dosage <input type="checkbox"/> SNCA dosage <input type="checkbox"/> PINK1 <input type="checkbox"/> DJ1 <input type="checkbox"/> Amyotrophic lateral sclerosis <input type="checkbox"/> C9orf72 <input type="checkbox"/> SOD1 <input type="checkbox"/> TARDBP <input type="checkbox"/> FUS <input type="checkbox"/> VCP <input type="checkbox"/> Prion diseases (CJD/GSS/FFI) <input type="checkbox"/> PRNP <input type="checkbox"/> Other diseases/genes:
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SAMPLE:

- Mention name, first name, date of birth on each tube
- Per patient 2 x 10 ml Lithium Heparin blood⁴ is required
- Store samples on room temperature or at 4°C, do NOT freeze
- Samples can be sent by regular mail
- Blood samples should reach our lab within 48 hours (Friday before 2PM)
- **Urgent analysis can only be performed after consultation**

⁴Note: other material can only be accepted after consultation

SHIPMENT:

Diagnostic Service Facility (DSF)
University of Antwerp - CDE
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REQUEST FORMS:

By mail (DSF@bornbunge.be), on the DSF website (<http://www.molgen.ua.ac.be/DNAdiagnostics>)