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| Clinical information sheetNeurofibromatosis (NF)NNFF International **NF1** Genetic Mutation Analysis Consortium Submission Form |
|  |
| Centrum Medische Genetica UZ Gent  |

FORM USE

Name of investigator (required field): Click and enter text.

Institution (required field): Click and enter text.

Please include phone number and/or e-mail address (one of them is required so that we may contact you if we have questions about your submission):

Investigator's e-mail address: Click and enter text.

Investigator's Phone Number: Click and enter text.

1. demographic information

**Mutation Database Number:** Click and enter text.

See above under Definition of General Terms. Enter this number if this is not the first submission for a given patient.

* 1. NNFFIDB Number: Click and enter text.

 *Clinical Database Participants Only*

See above under Definition of General Terms. If this patient is known to have been entered into the Clinical Database, please complete this information, otherwise leave it blank.

* 1. Sex

[ ]  Male

[ ]  Female

In the case of ambiguous gender, use the chromosomal definition of sex.

* 1. Date of Birth: Choose date.

If there is any uncertainty as to birth date, enter the date which is commonly used by the government on health records. Date of birth is used by the Central Database Centre as the primary variable to ensure that a patient or family member is only entered once into the Database. It is therefore essential that this information be accurate.

* 1. Date of Death

Enter exact date of death. If only year, or month and year, is known, enter that information and leave the other fields blank.

[ ]  Deceased, Date of Death: Choose date.

[ ]  Not applicable (still alive)

[ ]  Deceased, but date unknown

* 1. Maternal ethnic origins?

[ ]  Asian – Japanese [ ]  Latin American – Other

[ ]  Asian – Chinese [ ]  Native American – Other

[ ]  Asian – Indian Sub-continent [ ]  White

[ ]  Asian – Other [ ]  Other – Specify: Click and enter text.

[ ]  Black [ ]  Other – Unknown

* 1. Paternal ethnic origins?

[ ]  Asian – Japanese [ ]  Latin American – Other

[ ]  Asian – Chinese [ ]  Native American – Other

[ ]  Asian – Indian Sub-continent [ ]  White

[ ]  Asian – Other [ ]  Other – Specify: Click and enter text.

[ ]  Black [ ]  Other – Unknown

1. 1. N.I.H. Diagnostic Criteria

Please indicate which of the diagnostic criteria are met for NF1 in this patient. (Check all that apply)

[ ]  Six or more cafe au lait macules: Prepubertal > 5mm,Postpubertal > 15mm

[ ]  Two or more neurofibromas of any type or one plexiform neurofibroma

[ ]  Freckling in the axillary or inguinal regions

[ ]  Optic glioma

[ ]  Two or more Lisch nodules

[ ]  A distinct osseous lesion such as sphenoid dysplasia

[ ]  A first-degree relative with NF1 by the above criteria

* 1. PEDIGREE

Please use the space on this page to provide a pedigree using the symbols listed on the form. Omit names and local identification/lab numbers but include the Mutation Database Number and Clinical Database (NNFFID) Number where available.

The format for describing the pedigree should look similar to the following:

|  |  |  |  |
| --- | --- | --- | --- |
| Individual  | Gender | Nf-Affected | Tested For Mutation |
|  | ------ |  |  |
| Proband | Enter text. | Click and enter text. | Click and enter text. |
| Sibling 1  | Enter text. | Click and enter text. | Click and enter text. |
| Sibling 2  | Enter text. | Click and enter text. | Click and enter text. |
| etc. | Enter text. | Click and enter text. | Click and enter text. |
| Father | Enter text. | Click and enter text. | Click and enter text. |
| Mother | Enter text. | Click and enter text. | Click and enter text. |
| etc. | Enter text. | Click and enter text. | Click and enter text. |

Please enter your pedigree information in the following text box:

|  |
| --- |
| Click and enter text.  |

1. NF Signs and Symptoms
	1. Date of Exam: Choose date.

If exact date of latest exam is unknown complete the month and year fields as appropriate.

* 1. Superficial neurofibromas (check all types that are present)

[ ]  None

[ ]  Cutaneous

[ ]  Subcutaneous

[ ]  Pendulous

[ ]  Present – type unknown

[ ]  Unknown

Use the "Unknown" option if it is not known whether or not the patient has superficial neurofibromas.

A subcutaneous neurofibroma is located beneath the skin and will move little when the surrounding skin is pulled laterally. A cutaneous neurofibroma is located superficially in the skin and will move with the surrounding skin when the skin is pulled laterally. A pendulous neurofibroma is located above the skin surface and appears as a tag, or is connected to the body by a stalk.

Use the following space to record additional relevant information or to indicate any unusual findings such as pigmentation, atypical biopsy results or unusual distribution:

Click and enter text.

* 1. Plexiform neurofibromas (check all locations that apply)

[ ]  None [ ]  Trunk – ventral

[ ]  Orbit [ ]  Arm

[ ]  Face [ ]  Leg

[ ]  Head/Neck [ ]  Present – location unknown

[ ]  Trunk – dorsal [ ]  Unknown

If a plexiform lesion extends over more than one of the locations listed above, choose the area involved with the greatest portion of the neurofibroma for each plexiform.

Use the following space to record additional relevant information or to indicate any unusual findings such as pigmentation, atypical biopsy results, unusual location:

Click and enter text.

* 1. Internal neurofibromas

[ ]  Absent

[ ]  Present – asymptomatic

[ ]  Present – symptomatic

[ ]  Possible

[ ]  Unknown

Use the following space to note the location and size of internal neurofibromas and briefly describe any symptoms they are causing:

Click and enter text.

* 1. Neurological Problems

[ ]  Absent

[ ]  Present (please describe in following comment field)

[ ]  Unknown

Use the following space to record any neurological problems, whether or not they appear related to NF1:

Click and enter text.

* 1. Education completed

[ ]  Not applicable (too young for school) [ ]  Grade 10-high school completion

[ ]  Still in school at or above age level [ ]  Undergraduate degree

[ ]  Still in school behind age level [ ]  Higher degree

[ ]  Grade 5 or less [ ]  Unknown

[ ]  Grade 6-9

Educational level is at or above the age appropriate level if the child is in a regular school and is passing all subjects. Child is behind age appropriate level if he is in a special school for the learning disabled, needs substantial classroom assistance or is failing subjects or grades in a regular school.

Please use the space below to note any special circumstances which would explain early school leaving or failing grades in a child or adult. Examples include a child who has repeated a grade as a result of extensive hospitalizations or immigrants whose first language is not that of your school system.

Click and enter text.

* 1. Developmental examination

[ ]  Not Done

[ ]  Normal (provide I.Q. if known): Click and enter text.

[ ]  Abnormal (provide I.Q. if known): Click and enter text.

[ ]  Unknown

Accept results of any testing done by a qualified professional. Comment on areas of abnormality such as significantly delayed verbal or spacial abilities:

Click and enter text.

* 1. Cardiovascular disease

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Unknown

Comment on any clinical findings, ECG, angiogram, ultrasound abnormalities etc. and on any specific diagnosis such as aortic stenosis, ASD, hypertension, renal artery stenosis etc.

Click and enter text.

* 1. Skeletal anomalies

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Unknown

Comment on abnormalities such as pseudoarthrosis, congenitally bowed tibia, scoliosis, dysplastic vertebrae, dysplastic sphenoid wing, bony overgrowth etc.

Click and enter text.

* 1. Gastrointestinal complications

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Unknown

Do not include uncertain diagnoses. Do not include tumours under this item.

Comment on abnormalities such as pseudoarthrosis, congenitally bowed tibia, scoliosis, dysplastic vertebrae, dysplastic sphenoid wing, bony overgrowth etc.

Click and enter text.

* 1. Noonan phenotype

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Possible

[ ]  Unknown

Use "Possible" option if patient has borderline features of Noonan Syndrome. Use a comment to specify features, e.g. short stature, low set ears, midface hypoplasia, hypertelorism, ptosis, webbed neck, pulmonic stenosis, etc.

Click and enter text.

* 1. Other dysmorphic features

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Unknown

Use a comment to specify the dysmorphic feature, e.g. large ears, prominent jaw. Do not include the effects of plexiform neurofibromas or other overgrowth which have been described in items III.3 or III.9 above.

Click and enter text.

* 1. Neoplasms

[ ]  Absent

[ ]  Present (please describe with a comment)

[ ]  Unknown

Specify type of neoplasm e.g. pheochromocytoma, sarcoma, Schwannoma. Specify location, histology, age of onset, and course of any neoplasm with a comment.

Click and enter text.