

Laboratorium Klinische Genetica

Consent form for genomic diagnostic testing (gene set/WES/WGS)

| First name and surname | X-number: |
|--|--|
| | |
| Street and house number | * For trio-sequencing and/or patients under the age of 16, please fill in the details of parents and/or legal |
| ZIP code and address Date of birth | guardians. |
| Date of birth PLAKEN | For trio-sequencing, one completed consent form is sufficient, provided both parents' co-sign. |
| Gender m / f | |
| BSN | |
| Parent 1 / legal guardian* | |
| First name and surname | |
| Street and house number STICKER | |
| ZIP code and address | |
| Date of birth | |
| BSN | |
| Parent 2 / legal guardian* | 7 |
| | |
| First name and surname | |
| Street and house number | |
| Date of birth DIAKKEN | |
| BSN | |
| and/or whole exome sequencing (WES) and/or whole genome sequencing | |
| Incidental findings | |
| During consultation It was explained that incidental findings are reported | as follows: |
| The analysis state as a discrete control of the con | or monitoring is possible. |
| The predisposition to a disorder will be reported if medical treatment of | |
| The predisposition to a disorder will be reported if medical treatment of the predisposition to a disorder will not be reported if, our actual infor | |
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| The predisposition to a disorder will not be reported if, our actual infor | mation suggests that no medical treatment or |
| The predisposition to a disorder will not be reported if, our actual infor monitoring is possible. | mation suggests that no medical treatment or |
| The predisposition to a disorder will not be reported if, our actual infor monitoring is possible. If an incidental finding has been identified that presents a high risk of a be reported. | mation suggests that no medical treatment or a disorder not for me but for my (unborn)child, this will |
| The predisposition to a disorder will not be reported if, our actual informonitoring is possible. If an incidental finding has been identified that presents a high risk of a be reported. Only to be completed by the applicant in the event of nonconformity from | mation suggests that no medical treatment or a disorder not for me but for my (unborn)child, this will the above agreements |
| The predisposition to a disorder will not be reported if, our actual informonitoring is possible. If an incidental finding has been identified that presents a high risk of be reported. Only to be completed by the applicant in the event of nonconformity from the fother agreements were made during the consultation about reporting in the consultation | mation suggests that no medical treatment or a disorder not for me but for my (unborn)child, this will the above agreements acidental findings, please list them below: |
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Future contact

It is possible that new data and information of importance to you could become available in the future. I can indicate if I wish to be informed about this or not.

Make your choice:

The Clinical Genetics Department may contact me in the future regarding new data or information which becomes available. The Clinical Genetics Department may **not** contact me in the future about new data or information which becomes available.

General and signatures

| I have been informed both verbally and in writing about extensive genomic diagnostic testing. I fully understand that I have the option to change or withdraw my consent at any time. | |
|---|---|
| Sign here | |
| Patient's name* | Patient's signature* |
| Name of parent 1 / legal guardian* | Signature of parent 1 / legal guardian* |
| Name of parent 2 / legal guardian* | Signature of parent 2 / legal guardian* |
| Date | |

* Patients under the age of 12 are not required to sign in person: the signature of both parents/legal guardians will suffice. For patients between the ages of 12 and 16, both parents/legal guardians co-sign with the patient if possible. A signature form both parents is required for trio sequencing.

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