

Institut für Humangenetik I Im Neuenheimer Feld 366 I 69120 Heidelberg

Institute of Human Genetics

Prof. Dr. med. Christian Schaaf

Medical Director

Genetic Outpatient Clinic

phone: +49 (0)6221 56-5087 fax: +49 (0)6221 56-5080

Diagnostics Laboratories

fax: +49 (0)6221 56-5091 Molecular Genetics: +49 (0)6221 56-32484 Cytogenetics and Molecular Cytogenetics: +49 (0)6221-56-36879

Patient / Person being tested	
Family name, first name Date of birth	自然學學
Address	
I have been informed about the significance and consequences of the planned genetic analyses and I have had sufficient time for questions and reflection. I have received a patient information sheet (see QR code). I have no further questions. I am aware that I can revoke my consent at any time.	
With my signature, I consent / confirm consent on behalf of my relatives / the person for whom I have custody, to the collection of the necessary blood / tissue samples and agree that the findings may be stored in the UKHD patient data system as an aid to diagnosis:	
In the course of the planned examination(s), the analysis /analyses may reveal genetic alterations that are not directly related to the indication of the analysis. Such incidental findings may be medically relevant – or possibly become relevant later in life – however, receiving a genetic test result may also cause distress, be burdensome and /	□ yes
or have implications for your life and future.	□ no
As a person capable of giving consent, I would like to be informed about incidental findings concerning myself / the person in my custody.	
Specificities when performing genetic testing on children:	□ yes
I would like to be informed about incidental findings that are of potential clinical significance for my child during childhood / adolescence.	□ no
Incidental findings will always be reported - at the discretion of the competent physician - if withholding or failure to act upon them would result in harm to the child.	3
I would also like to be informed about incidental findings that will only be of clinical significance for my child during adulthood (adult-onset conditions)	□ yes □ no
The GenDG stipulates that sample material should be destroyed when no longer required for the testing for which it was requested.	☐ yes
I consent to the storage of the sample material and its use for results verification, future genetic analyses of myself and within the context of my family and for quality assurance.	□ no
Surplus material is an important source for quality assurance and for scientific purposes; it is kept encoded, which makes it impossible for unauthorized individuals to attribute the sample to you / your relatives / the person in your custody.	□ yes
I consent to the use of remaining sample material to aid medical teaching and research.	
I allow that the medical and genetic data that has been collected from me / my relatives / the person for whom I have custody, may be used for scientific purposes in a (partially) coded form and under anonymized conditions be published in scientific journals.	☐ yes ☐ no
The GenDG stipulates that results of genetic analyses ought to be destroyed after 10 years. However, this data could become important for you / your child / the person in your custody and other family members in the future.	☐ yes
I agree to the storage of genetic data and analysis results beyond the legally defined period.	□ no
	•
Place, date Signature of patient / person to be examined / legal representative	
Name of Treating Physician Signature of Treating Physician	

Informed consent to genetic testing according to the Genetic Diagnostics Act (GenDG)