

Informed Consent Form

for genetic testing

Purpose of test (indications):						
Patient details:						
First name:			Medical Records No.:			
Surname:			Sex:		M F	
Date of birth	day mon	h year	PESEL: (for individuals without PESEL, enter type and number of other personal ID document)			
Contact details - address and telephone number: (for underage and incapacitated patients, full name and address of their legal guardian)						
Details of biological sample:						
Type of sample	blood	fibroblasts	trophoblast	an	nniotic fluid	other
suspicion / clinical diagnosis of:						
 In addition, I have been informed that: the test result may indicate that it is advisable to take biological samples from other family members (to increase diagnostic significance), the test result may help to establish whether my family members are carriers of a genetic disorder, the test result may help to establish the genetic risk for a particular disease in the family, including in the extended family (if practicable), in some cases the test result will be non-informative, the test will fail for technical reasons or the DNA will be compromised, making it necessary for the samples to be taken again, the test result may reveal an abnormality in the tested gene other than the abnormality which is significant for the suspected or diagnosed disorder specified above, if an underage patient becomes of age between the date of taking the sample and the date of presenting the test result, he or she will need to sign an additional Informed Consent Form before receiving the test result, I may withdraw my consent for storing isolated DNA and its use for scientific research at any time. 						
 I am aware that once the sample is taken from me for testing and I have signed this Informed Consent Form, the service commences (and starts generating expenses) and therefore my withdrawal (if any) from the test, although it will stop any test procedures, it will not entitle me to claim a refund of the fee for the test, whether in whole or in part. I consent to being informed of the risk of a pathological development in case of identifying any so-called unexpected abnormalities. I am aware that the service provider is under no obligation to inform me of the risk of such so-called unexpected abnormalities. 						
Full name of patient / legal guar consenting to sample collection		Signature of pat	ient / legal guardian*			Date
Signature of pl	hysician		Date			* delete as applicable