

Request form for genetic examination - gamete donor

Personal data of the patient (label):		Referring physician:	
Name and surname: ID number: Date of birth: Insurance: self-payer Gender: male female Address: Diagnosis (MKN):		 (name, specialisation, identification number, workplace, stamp, signature)	
Primary sample: peripheral blood (5 ml of non-coagulating blood)		Other material:	
☐ in K ₃ EDTA (<i>molecular genetics</i>) ☐ in Li-Heparin (<i>cytogenetics, karyotype</i>)		☐ isolated DNA from: (<i>molecular genetics</i>)	
Date and time of collection:		Date and time of indication (if different from the collection date):	
Clinical data: (to be completed by the referring physician)		☐ STATIM	
Required examinations:			
☐ Karyotype ☐ CarrierTest - preconception panel of donor candidate (Carrier status for recessive mutations in genes <i>GJB2</i>, <i>CFTR</i>, <i>SMN1</i>, <i>COL4A5</i>, <i>OTC</i>, <i>DMD</i>, <i>GLA</i>, <i>MTM1</i>, <i>IL2RG</i>, <i>NBN</i>, <i>ATM</i>; response to FSH stimulation; HLA-C typing) ☐ Additional evaluation of selected genes for matching: (Can be requested only if the donor has undergone a CarrierTest using the WES method at GNTlabs.) List the individual genes separated by a semicolon: ☐ Genetic testing of genes (1 to 10 genes) in a donor without previous CarrierTest using the WES method at GNTlabs. List the individual genes separated by a semicolon:			
Separate tests (sample not tested using CarrierTest)			
Fragile X syndrome – detection of CGG repeat expansion in the <i>FMR1</i> gene Cystic fibrosis – 50 mutations + Tn variants of IVS8 in the <i>CFTR</i> gene Spinal muscular atrophy – determination of copy number of exon 7 and 8 in the <i>SMN1</i> gene AR deafness – detection of 35delG <i>GJB2</i> mutation HLA-C DNA Test - HLA-C typing (C1/C2) Response to hormonal stimulation – Ser680Asn polymorphism in the <i>FSHR</i> gene		Thrombophilic mutations ☐ Leiden mutation (G1691A) in the <i>F5</i> gene ☐ G20210A <i>F2</i> mutation (Prothrombin) ☐ C677T polymorphism in the <i>MTHFR</i> gene ☐ A1298C polymorphism in the <i>MTHFR</i> gene	
Informed consent* - examined person			
AGREES with the examination of the sample with the use of the sample for research with sample storage		DISAGREES with sample storage	
*) By submitting the request, the referring physician confirms that the patient or legal representative has signed the Informed Consent, which is either stored in the patient's documentation or attached to this request.			
Examination conducted by: GENNET, Ltd., GENNET Laboratories, Pekařská 635/6, 158 00 Praha 5 - Jinonice, Tel: 226 231 691			
Laboratory records:			
Date and time of sample/request reception:		Sample/request accepted by:	

