

### CENTRUM MENSELIJKE ERFELIJKHEID

8-24990-92-00

# REQUEST FORM NON-INVASIVE PRENATAL TESTING (NIPT)





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Website: http://www.uzleuven.be/nl/centrum-menselijke-erfelijkheid/centrum-mense

IDENTIFICATION	MOTHER		CLINICAL	INFORMATION MOTHER	
PATIENT IDENTIFICATION  Last name: Address: Date of birth:  INSURANCE COMPANION INSURA	EAD-/HOS-nr. First name:  Gender:  KG1/KG2: Family relation  Ime institution:		Expected dat  Singleton pregnancy	□ MCMA pregnancy: kg	
INFORMED CONSENT			TO BE COMPLETED ON THE BACK PAGE		
	other (cfDNA tube [Streck/Roche]) sh		be delivered to t	he laboratory within 72 hours.  Date of blood sampling://201	
Maternal blood:	3470 ☐ 1 full cfDNA tube [white cap, Roche]			Time of blood sampling: h min	
INDICATION					
☐ Maternal age:  Specify: years of age			☐ Personal request of the patient		
☐ Family history: Specify: in		-		r relevant clinical information:  fy:	
CLINICIANS IN COPY Gynaecologist:	<b>1</b>		Family doctor:		
DATE OF REQUEST: / / 201			FETUS (FOR INTERNAL USE)		
REFERRING Dr:  I.D. nr.: N° INAMI: L  Signature		Last Addi Date INS Insu Stan	URANCE COMPAN rance in	First name:  Gender:	
				Department:	

## CONSENT FORM NON-INVASIVE PRENATAL TESTING (NIPT)



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#### **INFORMED CONSENT OF THE MOTHER**

- I have been informed about the possibilities and limitations of this test, as described in the information brochure. I have had the opportunity to ask additional information from my doctor.
- 2. I understand that this test is designed to detect trisomy 21, 18 and 13 as from 10 weeks of gestation. Other more appropriate testing may be required when there is an increased risk for certain other genetic disorders.
- 3. I have been informed that this test is very accurate, but not 100%. In case of a normal result, the probability that the baby would still have trisomy 21, 18 or 13 is very low, but cannot be completely excluded. An abnormal result should always be confirmed by invasive prenatal testing (preferably amniocentesis).
- 4. I have been informed that the result will be available within a maximum of 7 calendar days from blood sampling. I can consult my results in my online medical file through www.mynexuzhealth.be.
- 5. I understand that in less than 1,5% of the cases, the NIPT is inconclusive or fails. In this case, the NIPT can be repeated once on a second blood sample (at no additional cost).
- 6. Using NIPT, all chromosomes are analyzed. Therefore, in rare cases, NIPT can also detect other chromosomal abnormalities, such as a trisomy of another chromosome or a chromosome abnormality important for my own health or that of my baby. The Centre for Human Genetics or my gynaecologist will contact me should this be the case.
- 7. I understand that NIPT is reimbursed in Belgium. In that case, my personal cost for the laboratory test is € 8,68. In case of an increased allowance, the NIPT is free of charge. When I'm not a member of a Belgian service for public health insurance, I will be charged € 260.
- □ I confirm that during this pregnancy a combined test or NIPT has not already been performed and reimbursed by the public health insurance.
- ☐ I understand the above information and I agree that NIPT may be performed.