



REFERRAL FOR AMBULATORY LABORATORY INVESTIGATIONS

Date			Number of analysis				
Refferal valid until			6 months				
CITO:							
	Gender: Woman	1an □	Date of birth (dd.mm.yyyy.): Identification Number:				
	ivame, oumanic		<u>-</u>				
	Address:		Phone:				
PATIENT	E-mail: Signature: * with my signature, I confirm acceptance to genetic testing * with my signature, I confirm acceptance that encrypted results of the testing will be sent to my e-mail						
	Basic Dg code _	week		code			
		,					
	Doctor:	Specialty cod	Specialty code or the title:				
8	ID:						
SENDER	Authority:		Phone, e-mail,	, where to repo	ort:		
MATERIAL							
A - blood B - buccal epithelium C - FFPE or fresh tissue D - missed abortion material and mother blood							

No.	Genetic Testing	Material	Price, €
1.	Karyotyping	Α	200
2.	Molecular Diagnosis of Trombophilia	A/B	100
3.	Non-invasive determination of fetal Rhesus factor	Α	168
4.	Molecular Testing of the AZF region of the Y chromosome	A/B	200
5.	PAI determination	A/B	80
6.	Determination of the RHD gene zygosity status	A/B	110
7.	Testing for the Fragile X chromosome syndrome	A/B	285
8.	Determination of the most common two CFTR gene mutations that causes cystic fibrosis	A/B	85
9.	CFTR mutation panel (determination of 68 mutations)	A/B	260
10.	FSH/LHC GR	A/B	90
11.	HLA typing	Α	250

Notes:

No.	Genetic Testing	Material	Price, €	No.	Genetic Testing	Material	Price, €
12.	DNA contamination test	Α	110 24.		Exome – Reanalysis – Hereditary recessive disease		160
13.	KIR typing	Α			panel		
14.	Sequencing (1 reaction)	A/B	200	25.	Exome – Reanalysis – Most common hereditary cancer gene panel: 33 genes	-	160
15.	Sequencing (1 reaction for 2 samples)	A/B	221	26.	Exome – Reanalysis – Genetic predisposition of embryonic	-	160
16.	Non-invasive prenatal testing (NIPT)	Α	315		development abnormalities		
17.	Non-invasive prenatal testing (NIPT+)	Α	400	27.	Exome – Reanalysis – The Early Menopause Panel	-	160
18.	Molecular testing of the CYP21A2 gene (congenital adrenal hyperplasia)	Α	495	28.	Exome – Reanalysis – Disorders of Spermatogenesis Panel	-	160
19.	Exome – Hereditary recessive disease panel	Α	650	29.	Skin Diagnostic DNA Test	A/B	160
20.	Exome – Most common hereditary cancer gene panel: 33 genes	Α	650	30.	Metabolism DNA Test	A/B	160
	Exome – Genetic			31.	Vitamins metabolism DNA Test	A/B	160
21.	predisposition of embryonic development abnormalities 650		32.	Sports achievements DNA test	A/B	160	
22.	Exome –The Early Menopause Panel	Α	650	33.	Mentality (way of thinking) DNA test	A/B	160
23.	Exome -Disorders of Spermatogenesis Panel	Α	650	34.	Determination of BRCA1 and BRCA2 hereditary mutations	A/B	200

Notes:

No.	Performed after prior agreement	Material	Price, €	No. Performed after prior agreement		Material	Price, €
35.	Determination of the 3 most common mutations that cause hereditary hemochromatosis		200	40.	Genetic testing for celiac disease	A/B	126
	nereditary hemochiomatosis			41.	Lactose intolerance test	A/B	200
36.	Determination of Alpha 1-antitrypsingene mutations (Z and S)	A/B	200	42.	Determination of large BRCA1/BRCA2 deletions/ insertions with MLPA	A/B	420
37.	Determination of resistance against HIV-1 infection	A/B	A/B 95		Hereditary cancer NGS panel (111 genes)	A/B	1310
38.	Spinal muscular atrophy (SMA) carrier testing with MLPA (1 patient)	Α	285	44.	Superior Onko	С	905
39.	SMA carrier testing with MLPA (pair)	Α	325				

No.	Missed abortion				Material	Price, €			
45.	Genetic testing of missed abortion materia		D	330					
46.	Genetic testing of missed abortion materi		С	370					
Spontaneously terminated pregnancy date and place:									
Pregnancy week:			How pregnancy was terminated: Medication Instrumental						
Number of missed abortion (including current):			Number of pregnancies: Number of births:						
Infect	ion investigations Yes No No	Type	of sample: Chorion 🗌	Paraffin blo	ck 🗌				
Comple	eted by the laboratory								
Samp	le identification number		Date and time of sampl	e registration in	the labora	tory			
The s	ample meets the criteria Yes	No 🗌							

Notes:





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