THIS FORM SHOUL	D BE COMPLETED BY	THE PRESCRIBING PHYSICIAN A	AND SENT WITH THE BLOOD SAMPLE	
eurofins Biomnis		Cystic fibros	Clinical Information Form Cystic fibrosis gene analysis (CFTR gene)	
	ON • Tel.: +33 (0)4 72		73 56 • E-mail: international@biomnis.com	
PATIENT DETAILS		CLINICIAN		
First(s) name(s):				
Surname:				
Date of birth:			City:	
TESTS REQUESTED				
Cystic Fibrosis CFTR-se	creening of most free	uent mutations		
		by New Generation Sequencing	g (NGS)	
FAMILY TREE			CONSULTATION CERTIFICATE	
			AND PATIENT CONSENT FORM	
			(Decree n° 2008-321 of 4th April 2008, amended on 27th May 2013)	
			I, the undersigned, Medical Doctor (MD), certify that I have fully informed my patient	
			Mr/Mrs/Miss of the information defined according to the article R.1131-4 of decree n°2008-321 dated 4 April 2008 of the French public health code and amended on 27 th May 2013 and that I have obtained written informed consent from my patient under the conditions specified in article R.1131-5. Signed in <i>(city)</i>	
	on of the mutations vary d	epending on the ethnic/geographical	Physician's signature	
			I, the undersigned,	
REASON BEHIND THE TEST REQUEST FOR A Suspected cystic fibrosis ENT disease: Respiratory disease: Digestive system disease: Pancreatic affection: Sweat test: NO YES, result (please indice			R1131-5 of the public health code and decree of 27th May 2013. Signed in (city) on	
□ Infertility		,	Patient signature	
Bilateral absence of the Please include the ultraso		□ NO □ YES		
☐ Medically assisted pro	ocreation			
Ovum donation				
Suspected cystic fibro LMP: Amniocentesis: N Digestive enzyme assay Please include the ultraso	☐ Date of c O ☐ YES y on amniotic fluid:			
Family investigation Heterozygote screen Familial mutation to be Please include the CFTR	ing of the family of a screened for:	patient with cystic fibrosis	a partner of heterozygous individual	