

Clinical Information Form

Genetic characterisation of male infertility

INTERNATIONAL DIVISION

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Customer number:

REFERRING PHYSICIAN

First name(s): Last name:
 Address:
 Postcode: [][][][][] City:
 Country: E-mail:
 Tel.: [][][][][][][][][] Fax: [][][][][][][][][]

Physician's stamp

PATIENT (INDEX CASE)

First name(s): Last name:
 Name at birth:
 Date of birth: [D][D][M][M][Y][Y][Y][Y]
 Address:
 Postcode: [][][][][] City:
 Country:
 Date of sample collection: [D][D][M][M][Y][Y][Y][Y]

OTHER PHYSICIANS INVOLVED IN THE PATIENT'S TREATMENT (andrologist, endocrinologist, reproductive biologist, gynaecologist)

First name(s): Last name:
 Address:
 Postcode: [][][][][] City:
 Country: E-mail:
 Tel.: [][][][][][][][][] Fax: [][][][][][][][][]

TYPE OF ANOMALY

Obstructive azoospermia (excretory) Severe oligozoospermia
 Non-obstructive azoospermia (secretory) Cryptozoospermia
 Other (contact us):

FAMILY HISTORY

Infertile blood relatives YES NO
If yes, specify relationship (brother/sister, uncle/aunt/ cousin, specify whether paternal or maternal side).

Consanguinity YES NO

Geographical origin*:
 (*The frequency of genetic variants differs according to the ethnic/geographical origins of the patient)

Family tree

CONSENT FOR TESTING OF AN INDIVIDUAL'S GENETIC CHARACTERISTICS

(In accordance with Articles R.1131-4 R.1131-5 and the Code of Public Health).

I the undersigned
 born on [D][D][M][M][Y][Y][Y][Y]
 acknowledge that I have been informed by Dr:

on the tests of genetic characteristics that will be performed in order to:

- confirm or reject the diagnosis of a genetic disease related to my symptoms;
- confirm or reject the pre-symptom diagnosis of a genetic disease; identify healthy carrier status (screening for heterozygote or chromosomal rearrangement);
- evaluate my genetic susceptibility to a disease or to a drug treatment.

► To this end, I agree:

- to a biological sample being obtained from me
- to a biological sample being taken from my child or an adult under my guardianship
- to a sample being taken from my foetus.

I have been informed the results of the examination of genetic characteristics will be presented to me by the above physician in the context of an individual consultation. If examination reveals results other than those expected, the above-named physician will determine further investigations during an individual consultation.

► If part of the sample remains unused after examination,

- I agree to its use, if appropriate, for scientific research purposes. In this case, all the medical data about me will be protected by complete anonymisation. As a result of this I am aware that these scientific studies will neither benefit me nor put me at risk.

Signed in
 on [date] [D][D][M][M][Y][Y][Y][Y]

Signature of adult patient or legal guardian of the child or legal guardian of an adult under guardianship:

CERTIFICATE OF CONSULTATION

(Decree no. 2008-321 of 4 April 2008 – enacted May 27, 2013).

I the undersigned
 a physician, in accordance with Articles R.1131-4 and R. 1131-5 of the Code of Public Health, certify that I conducted a consultation with the patient named below on this date to provide him/her with information on investigated disease characteristics, means for detecting such mutations and the options for prevention and treatment.

Signed in
 on [date] [D][D][M][M][Y][Y][Y][Y]

Signature of the physician:

SAMPLE REQUIREMENTS

- 5 ml of EDTA whole blood at room temperature (refrigerate sample if transport > 48H) **and/or** a tube of extracted DNA at room temperature (minimum volume 50µL and minimum quantity 1µg)

CLINICAL AND BIOLOGICAL ASSESSMENT

Clinical details

Age years Weight..... kg
Height cm
Testicular volume: ml

Primary/secondary infertility,

Specify the duration:

- Bilateral cryptorchidism YES NO
Epididymitis (*uni- or bilateral*) YES NO
Acute torsion of the testicle YES NO
Testicular trauma YES NO
Orchitis YES NO
Chemoradiotherapy YES NO
Exposure to toxic substances YES NO
Systemic diseases YES NO
Kidney failure YES NO
Absence of vas deferens YES NO

Please specify:

- Epididymis abnormality (absence, dilatation) YES NO
Delayed onset of puberty, hypogonadism YES NO
Scrotal/endorectal ultrasound YES NO

if yes, attach report

- External genital organ anomalies YES NO
Morphological anomalies YES NO
Abnormal hair YES NO
Gynaecomastia YES NO

Biological examination

Semen analysis

Volume pH.....
No. of sperm No. of round cells.....

Analysis of hormone levels

FSH:
LH:
Testosterone: total
Inhibin B:
Seminal plasma markers:
Alpha-glucosidase:
L-carnitine:
Fructose
Citric acid
Acid phosphatase

GENETIC TESTS ALREADY PERFORMED

Karyotype performed YES NO

Abnormal result (*please specify*): *Please enclose the report*

Testing for microdeletions of AZF regions YES NO

Abnormal result (*please specify*): *Please enclose the report*

Testing for mutations of the CFTR gene YES NO

Abnormal result (*please specify*): *Please enclose the report*

Chromosomal microarray analysis (CMA) / Array CGH YES NO

Abnormal result (*please specify*): *Please enclose the report*

List of documents to be included with the request

- Consent and signed genetic counselling form
- Semen analysis (preferably two)
- Analysis of hormone levels
- Report of genetic analyses performed
- Copy of the pathological anatomy report, if a testicular/epididymal biopsy was performed
- Surgery report (urologist)