

**Responsible doctor** \* Country Code

Practice / hospital

Title / first name / last name

Street / number

CC\*      Postcode / city

Telephone

Fax



WM-3092-EN-006

Field for barcode

Referring laboratory

### Patient information

First name / last name

Date of birth (DD/MM/YYYY)

Singleton pregnancy     Twin pregnancy

**PrenaTest®** gestational week 9 + 0 to 32 + 1 p.m.

**Option 1**     Trisomy 21

**Option 2**     Trisomies 21/18/13

**Option 2 Plus**     Trisomies 21/18/13, rare autosomal aneuploidies (RAAs\*), Copy Number Variations (CNVs\*\*)

**Option 3**     Trisomies 21/18/13, gonosomal aneuploidies (X/Y)

**Option 3 Plus**     Trisomies 21/18/13, gonosomal aneuploidies (X/Y), RAAs\*, CNVs\*\*

\* RAAs: Examination of all autosomal chromosomes (Chr. 1 – 22) for monosomies & other trisomies  
\*\* CNVs: Examination of all autosomal chromosomes (Chr. 1 – 22) for partial (Micro-)deletions/-duplications (≥7Mb)

**Additional PrenaTest® option**

**22q11.2 microdeletion** –  / not for option 1. Please read the fact sheet!

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**Post analysis 22q11.2 microdeletion** –  / without new blood sample  
Based on PrenaTest® order not older than 3 months (not for option 1). Please read the fact sheet!      Prior barcode no.

Order by Fax: +49 (0) 89-2323735690

**NIPT-RhD** gestational week 11 + 0 p.m.

Non-invasive prenatal RhD genotyping from maternal blood for RhD negative women.

Please provide blood in 7.5 to 10 ml EDTA blood collection tube without separating gel

**Important: Patient MUST NOT be anti-D alloimmunized**

Please read the fact sheet!

**NIPT-RhD** –

Date of blood draw (DD/MM/YYYY)

Repeat blood sample (if 1st analysis failed)

Singleton pregnancy      Twin pregnancy

Gestational week      +      p.m.

Other Information (e.g. vanishing twin)

Body height      cm

Weight prior to pregnancy      kg

### Patient consent

The responsible doctor or the referring laboratory confirm by signing that the patient

- a) has received explanations and human genetic counseling in accordance with national legislation and has consented to the chosen genetic testing,
- b) has agreed to have her test result sent to the referring laboratory for forwarding to the responsible doctor,
- c) has agreed that her personal data and her blood sample may be transmitted to, processed and used by Eurofins LifeCodexx to carry out the genetic testing,
- d) is aware that she can withdraw her consent to the responsible doctor at any time. In that case the processing of the already collected personal data remains lawful.
- e) has taken note of the Notes on Data Protection (see reverse).

The patient consents to the storage and use of surplus examination material that is not identified by name for purposes of quality assurance, scientific research as well as the development of new diagnostic options

Yes     No

Result report will be provided in English language.

Place, Date	Signature / Stamp of the responsible doctor
	X
Place, Date	Signature / Stamp of the referring laboratory
	X

## Notes on Data Protection

1. Hereinafter we inform you about the collection of personal data in business relations. Personal data means any information relating to you as an identifiable natural person, e.g. name, address, e-mail address and payment details.
2. Responsible processor pursuant to Art. 4 para. 7 General Data Protection Regulation (GDPR) is Eurofins LifeCodexx GmbH, Lochhamer Straße 15, 82152 Planegg, Germany.
3. In order to provide the contractual services, personal data is collected, stored and, if necessary, passed on to third parties. Therefore the collection, storage and transmission are carried out for the purpose of fulfilling contractual or precontractual obligations and on the basis of Art. 6 para. 1 sentence 1 lit. b GDPR.  
If we ask for giving your consent to processing personal data, we obtain the consent for data processing on the legal basis of Art. 6 para. 1 lit. a GDPR.  
Failure to provide this information may result in that the contract cannot be concluded. A processing of personal data only takes place if you have given your prior consent or if permitted by law.
4. We maintain up-to-date technical measures to ensure the protection of personal data. These are adapted to the current state of the art.
5. You have the right to request information concerning the personal data stored with us at any time (Art. 15 GDPR). This also applies to the purpose of storage and the recipients or categories of recipients to whom the personal data is transmitted. In addition, you have the right to request correction and/or deletion and/or restriction of the data processing under the conditions of Art. 16 GDPR, Art. 17 GDPR and Art. 18. GDPR. Under the conditions of Art. 20 GDPR you can request a data transfer at any time.
6. If the purpose of storage has been omitted, personal data will be deleted or blocked immediately. A longer storage may take place if required by the European or national legislators, by EU regulations, laws or other regulations to which the controller is subject. Unless there is a need for further storage in order to conclude or fulfill a contract, blocking or deletion of the personal data takes place if a storage period prescribed by the mentioned standards expires.
7. All requests for information or objections to data processing should be directed by e-mail to our data protection officer at [Datenschutz@lifecodexx.com] or to the address mentioned above (2). For further information, please refer to the full text of the GDPR available on the internet at <http://eur-lex.europa.eu/eli/reg/2016/679> and our privacy statement, which is available on the internet at [www.lifecodexx.com](http://www.lifecodexx.com). Moreover you have the opportunity to complain to the responsible supervisory authority about data protection matters.

**Eurofins LifeCodexx GmbH**  
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