TEST REQUISITION FORM

PATIENT DETAILS	eurofins Genoma
Protocol no. (internal use only):	GeneSafe™
Surname	
Date of birth Place of birth	ORDERING LABORATORY / CLINICIAN
VAT no.	
Address:	
Date of blood withdraw	Name / Stamp
Gynecologist name:	PATIENT MEDICAL HISTORY
Address:	
Phone no.:	
E-mail:	
PREGNANCY HISTORY	INDICATION FOR TESTING
Patient current weight Kg Patient height Gestational age at draw + days Gestational age calculated by: Ultrasound;	□ Advanced maternal age; □ Advanced paternal age; □ Partner carrier of a genetic disorder: □ Male □ Female □ Specify disorder: □ Specify gene and mutation: □ Parental anxiety (low-risk) □ Abnormal ultrasound (describe): □ Other indication □ None
TYPF	OF TEST
ТҮРЕ	OF TEST
TYPE GeneSafe™	☐ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete)
O TH	☐ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo +
GeneSafe™ GeneSAFE™ Inherited □ GeneSAFE™ de novo □ GeneSAFE™ Complete (Inherited + de novo)	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) *** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome);
GeneSafe™ Inherited ☐ GeneSAFE™ de novo☐ GeneSAFE™ Complete (Inherited + de novo) Is it a redraw? ☐ Yes; ☐ NO CHECK LIST Please check if you provided the following information:	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome).
GeneSafe™ Inherited ☐ GeneSAFE™ de novo ☐ GeneSAFE™ Complete (Inherited + de novo) Is it a redraw? ☐ Yes; ☐ NO CHECK LIST Please check if you provided the following information: ☐ Patient's details;	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome).
GeneSafe™ Inherited	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES □ PHYSICIAN / LABORATORY
GeneSAFE™ Inherited	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES Name / Stamp
GeneSafe™ Inherited	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES □ PHYSICIAN / LABORATORY Name / Stamp Courier Mail; □ Courier Courier
GeneSAFETM Inherited GeneSAFETM de novo GeneSAFETM Complete (Inherited + de novo) Is it a redraw? Yes; NO CHECK LIST Please check if you provided the following information: Patient's details; Ordering Laboratory / Clinician; Pregnancy history Type of Test If wish to know the fetal gender;	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES □ PHYSICIAN / LABORATORY □ E-mail; □ On-Line; □ Mail; □ Courier □ PATIENT
GeneSAFE™ Inherited GeneSAFE™ de novo GeneSAFE™ Complete (Inherited + de novo) Is it a redraw? Yes; NO CHECK LIST Please check if you provided the following information: Patient's details; Ordering Laboratory / Clinician; Pregnancy history Type of Test If wish to know the fetal gender; If it is a redraw;	☐ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) ☐ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1936 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES ☐ PHYSICIAN / LABORATORY Name / Stamp ☐ E-mail; ☐ On-Line; ☐ Mail; ☐ Courier ☐ PATIENT ☐ E-mail; address
GeneSAFE™ Inherited	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES □ PHYSICIAN / LABORATORY □ E-mail; □ On-Line; □ Mail; □ Courier □ PATIENT □ □ E-mail; address □ □ On-Line; □ By Phone, no.: □
GeneSAFE™ Inherited	☐ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) ☐ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1936 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES ☐ PHYSICIAN / LABORATORY Name / Stamp ☐ E-mail; ☐ On-Line; ☐ Mail; ☐ Courier ☐ PATIENT ☐ E-mail; address
GeneSAFE™ Inherited	□ PrenatalSAFE®COMPLETE (PrenatalSAFE® Karyo + GeneSAFE™ Complete) □ PrenatalSAFE® COMPLETE Plus (PrenatalSAFE® Karyo Plus** + GeneSAFE™ Complete) ** This option includes in addition screening for the following syndromes: 22q11.2 syndrome (DiGeorge syndrome, Velocardialfacial syndrome); 1p36 deletion syndrome; Angelman syndrome (15q11.2 deletion syndrome); Prader-Willi syndrome (15q11.2 deletion syndrome); Cri du Chat syndrome (5p- deletion syndrome); Wolf-Hirschhorn syndrome (4p- deletion syndrome); Langer-Giedion syndrome (8q24 deletion syndrome); Jacobsen syndrome (11q23 deletion syndrome); Smith-Magenis syndrome (17p11.2 deletion syndrome). REPORTING PREFERENCES □ PHYSICIAN / LABORATORY □ PATIENT □ □ E-mail; □ On-Line; □ Mail; □ Courier □ PATIENT □ □ F-mail; address □ □ On-Line; □ By Phone, no.: □ □ In order to activate the on-line reporting option, you need to







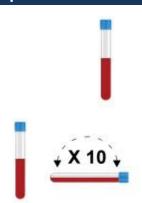
Test Submission Instructions

Informed consent and test requisition form

IMPORTANT: Fill in all required Test Requisition Form information to avoid delays and ensure timely reporting.

To ensure acceptance of your patient's specimen for testing, please verify that the **informed consent** has been signed from the patient and it has been enclosed with samples:

Sample collection instruction:



- Take the 10ml collection tube from the PrenatalSafe® Test Shipper Kit.
- Write the blood **collection date** in the specimen information section of the test requisition form.
- Write the patient's full name and date of birth on the collection tube label.
- Fill the collection tube <u>almost completely</u> with whole blood.
- Invert the collection tube 10 times.

Store collected blood at **room temperature** until ready for shipment. **Blood should never be**frozen!

Sample Packaging:

IMPORTANT: Always store kits at **room temperature**.

- Place the filled and properly labeled collection tube into the **PrenatalSafe**® shipper kit box. Only one patient sample per box.
- Place the completed **test requisition form** and **informed consent** into the shipper kit box, at the side.
- Put sample tubes inside the sponge and both inside the biohazard envelope. Close the box.
- Place shipper kit box inside of courier pack and seal.
- If you are shipping more than one sample, place as many as possible collection tubes into one shipper kit.
- If you are shipping more than one shipper kit, place as many as possible into one courier pack.
- Adhere the **courier airbill** pouch to the outside of the courier pack. Insert the airbill into the pouch.
- Call courier to arrange specimen pickup.
- Ship specimens, preferably the **same day** as collected. Specimens must be received by Genoma **within 5 days** of collection date. Genoma receives specimens Monday through Saturday.