

**Requisition Form**

Ship Samples, photocopy of Insurance card(s) (Front and Back) and copy of this form to:

Genoscientific
Durham Center,
2 Ethel Road, Suite 203 C, Edison, NJ 08817
Tel: (732) 662-5543
Fax: (732) 662-5544

Patient Information**Patient Name,** _____

Address, _____

City, _____ State, _____ Zip, _____

Collection Information

Collection Date, _____

Check at least one Box below for a personalized medicine management report:

Specimen Type: ☐ Blood, ☐ Buccal Swab ☐ Urine, ☐ Other**Swabs rubbed firmly in each cheek and under the gum****Patient Consent**

I request and authorize the CLIA accredited laboratory to perform the below designated test(s) on the DNA sample provided by me. My signature below constitutes my acknowledgement that have read the Patient Information Form which outlines the benefits and limitations of this testing which have been explained to my satisfaction by a qualified health professional.

Patient Signature, _____**Physician Information****Physician Name,** _____

Practice, _____

Office Phone, _____ Fax Number, _____

Physician Signature, _____**Insurance Information**

Patient Insurance Information: Include photocopy of insurance card(s) (both sides) OR complete Insurance section below.

Medicare Number _____ Relation Ship to Insured(optional) _____

Primary Insurance	ID Number	Group Number
Secondary Insurance	ID Number	Group Number

Advance Beneficiary Notice Instructions

All tests on this request form are subject to coverage limitations by Medicare and may require that an advance Beneficiary Notice (ABN) be signed by the patient prior to obtaining the specimen. When ordered tests are likely to be denied by Medicare, please complete a separate ABN with the patient's signature and date; submitting it with this requisition.

<input type="checkbox"/> Cardiac Panel	<input type="checkbox"/> Women's Health	<input type="checkbox"/> Hem-Pathology Molecular
<input type="checkbox"/> Prothrombin (Factor II) G20210A Mutation <input type="checkbox"/> Factor V Leiden Mutation <input type="checkbox"/> Factor V Leiden Mutation _HR <input type="checkbox"/> Factor XIII A V34L <input type="checkbox"/> MTHFR 677 and 1298 Mutations <input type="checkbox"/> Beta Blocker CYP 2D6 <input type="checkbox"/> CYP 2c19(Plavix) <input type="checkbox"/> CYP2C9 (Includes VKORC1) <input type="checkbox"/> ApoE C112R R158C <input type="checkbox"/> ApoB-100 R350Q <input type="checkbox"/> IL-6 G-174C <input type="checkbox"/> PAI 1 Drug Metabolism Panel. <input type="checkbox"/> Beta Blocker CYP 2D6 <input type="checkbox"/> CYP 2c19(Plavix) <input type="checkbox"/> CYP2C9 (Includes VKORC1) <input type="checkbox"/> CYP 3A4 and A5 Mental health Panel. <input type="checkbox"/> Beta Blocker CYP 2D6 <input type="checkbox"/> CYP 2c19(Plavix) <input type="checkbox"/> CYP2C9 (Includes VKORC1) <input type="checkbox"/> MTHFR 2 mutations <input type="checkbox"/> 5HT2C HLA Typing <input type="checkbox"/> HLA Dq6 and DQ8 <input type="checkbox"/> HLA B27 DNA Typing <input type="checkbox"/> Celiac Diseases DNA testing	<input type="checkbox"/> HPV-HR and genotyping by Real-Time PCR <hr/> <input type="checkbox"/> C. trachomatis <input type="checkbox"/> N. gonorrhoeae <input type="checkbox"/> Trichomonas Vaginitis <input type="checkbox"/> Herpes Simplex Viruses 1 and 2. <input type="checkbox"/> BD Affirm™ (Candida, Gardnerella, and Trichomoniasis) <input type="checkbox"/> Candida albicans <input type="checkbox"/> Gardnerella vaginalis <input type="checkbox"/> Candida glabrata <input type="checkbox"/> M. genitalium <input type="checkbox"/> M. hominis <input type="checkbox"/> U. urealyticum 16S <input type="checkbox"/> T. gondii <hr/> <input type="checkbox"/> Hepatitis C Viral Load Quantitative <input type="checkbox"/> Hepatitis C Genotype <input type="checkbox"/> Hepatitis B Viral Load Quantitative <hr/> <input type="checkbox"/> Cystic fibrosis mutations panel (liquid-based Pap specimens) 39 or 60 Mutations	<input type="checkbox"/> B-Cell clonality <input type="checkbox"/> BCR/ABL1 kinase mutation analysis <input type="checkbox"/> P210 BCR/ABL1 mutation analysis, quantitative <input type="checkbox"/> P190 BCR/ABL1 mutation analysis, quantitative <input type="checkbox"/> CEBPA mutation analysis <input type="checkbox"/> JAK2 mutation analysis, qualitative/quantitative <input type="checkbox"/> MPL W515/K mutation analysis <input type="checkbox"/> FML/RARA fusion protein <input type="checkbox"/> T-Cell clonality <input type="checkbox"/> IgVH hypermutation analysis <hr/> Leukemia Translocation Panel <input type="checkbox"/> AML (8;21) <input type="checkbox"/> ALL (9;22) <input type="checkbox"/> ALL (1;19) <input type="checkbox"/> APL (15;17) <input type="checkbox"/> CML (9;22) <input type="checkbox"/> Inv 16 <input type="checkbox"/> MLL <hr/> <input type="checkbox"/> Oncology. <input type="checkbox"/> KRAS-BRAF <input type="checkbox"/> BRAF <input type="checkbox"/> EGFR Other oncogenic assays
<input type="checkbox"/> Icd-9	<input type="checkbox"/> ICD-9	<input type="checkbox"/> ICD-9