



A DIVISION OF

GENESIS DIAGNOSTICS

MEDICAL DIAGNOSTIC LABORATORIES, L.L.C.

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Core OB/GYN Test Requisition Form

Ordering Physician/Laboratory

(Required: Include the ordering physician's first & last name, NPI, practice name, complete address, phone number and fax number.)

Physician to receive additional result report:

Physician's Signature:

Date:

Patient Information (Please Print)

Name (Last, First) (Required):

In Care of:

Patient Address:

City:

State:

Zip:

Gender (Required): Female Male

Date of Birth (Required):

Patient SS#:

Ethnicity†:

Phone Number:

Email:

Billing Information (Please include a copy of the front & back of card.)

- Patient Billing
Insurance Billing
Path Lab/Hospital
Physician Account

Relation (Required): Self Spouse Dependant

Diagnosis Codes (Required): Please provide ALL applicable diagnosis codes.

Primary Insurance Carrier:

Insured's Name (if not patient):

Insured's SS#:

Insured's DOB:

Claims Address:

Medicare, Medicaid or Policy ID#:

Employer/Group Name:

Group#:

Specimen Information

Date Collected (Required):

Specimen Source:

Pharmacogenomics Test Selection

Whole Blood or Buccal Swab - *Informed Consent form must accompany specimen.

- 3101 Antiplatelet Agents-Aspirin, Cilostazol, Clopidogrel, Prasugrel, Ticagrelor
3102 Statins-Atorvastatin, Fluvastatin, Lovastatin, Pitavastatin, Pravastatin, Rosuvastatin, Simvastatin
3104 Thrombophilia-Susceptibility to Factor II, Factor V Leiden
3105 Calcium Channel Blockers-Amlodipine, Nifedipine
3106 Beta Blockers-Bufuralol, Carvedilol, Metoprolol, Propranolol, Talinolol, Timolol
3201 Pain Management-General-Alfentanil, Codeine, Fentanyl, Hydrocodone, Ketamine, Loroxicam, Methadone, Morphine, Opioids, Oxycodone, Sumatriptan, Tramadol
3303 Antimetabolites-Cytarabine, Fludarabine, Mercaptopurine, Methotrexate, Siilbinin
3306 Platinum Derivatives-Carboplatin, Cisplatin, Oxaliplatin, Platinum Compounds
3310 Uracil Derivatives-Capecitabine, Fluorouracil, Folfox, Folox, Leucovorin, Tegafur, Xelox
3404 Anxiety, Insomnia, Severe Agitation - Bupropion, Dexmedetomidine, Duloxetine, Escitalopram, Lorazepam, Midazolam, Oxazepam, Venlafaxine
3407 Depressive Disorder and Major Depressive Disorder-Agomelatine, Amitriptyline, Antidepressants
3603 Type-II Diabetes - Repaglinide, Sulfoureas, Urea Derivatives
3605 Inflammation-Anti-inflammatories
3704 Contraception-Oral Contraceptives

Confirmation of Informed Consent and Medical Necessity for Pharmacogenomic/Genetic Testing

My signature below certifies that I am a licensed medical professional or his/her representative or a genetic counselor authorized to order genetic testing. My signature further acknowledges the patient has been supplied information regarding genetic testing and has been informed about the purpose, limitation and possible risks.

In the event that the patient's health insurance plan determines for the test(s) I checked above that some of the genes that I requested for analysis are not covered, I understand that Medical Diagnostic Laboratories, L.L.C. shall perform and result the test(s) for the genes I selected, then submit the claim to the patient's health insurance plan for the testing of the genes covered under the patient's plan.

If the testing is covered by the patient's health plan and the out-of-pocket expense is less than \$150.00, testing will proceed without further delay or additional contact. The patient has given consent for genetic testing to be performed and the signed consent form is being provided with this requisition. I confirm that this testing is medically necessary for the specified patient and that these results will be used in the medical management and treatment decisions for this patient.

Medical Professional Signature (Req.): Date:

Pathology Test Selection

Date Collected (Required):

Specimen Source:

Anatomic Source (Required): Cervix/Endocervix Vagina Vaginal Cuff Other:

Date of LMP:

Previous Results: ASCUS LGSIL CIN 1 CIN 2 CIN 3

Date of Last Pap:

Normal LGSIL CIN 1 CIN 2 CIN 3

Pathology Test Requisition with completed clinical information must accompany specimen

Table with columns for ThinPrep, Liquid Pap only, and Liquid Pap with HPV Options (Ages 21 and older, Ages 30 and older)

HPV Reflex Test:

714 HPV Type-Detect 3.0 by Next Generation Sequencing High Risk Subtypes Only (Includes HPV Subtypes: 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59, 66, 68)

Infectious Disease Test Selection

ThinPrep or OneSwab (female - cervical, vaginal, rectal, lesion) (male - lesion, [urethral - special male swab required])

Testing by Real-Time PCR unless otherwise specified. To order panel components individually, select tests beneath the panel.

- 182 Aerobic Vaginitis (AV) Panel PCR (GBS, S. aureus, E.coli, E. faecalis)
127 Group B Streptococcus (GBS) Is the patient pregnant? Yes No
184 Staphylococcus aureus
141 Escherichia coli
153 Enterococcus faecalis
166 Bacterial Vaginosis (BV) Panel PCR [A. vaginae, BVAB2, G. vaginalis, Megasphaera species (Types 1&2)]
142 Atopobium vaginae
164 Bacterial Vaginosis Associated Bacterium 2 (BVAB2)
132 Gardnerella vaginalis
165 Megasphaera species (Type 1 and Type 2)
560 Candida Vaginitis Panel (C. albicans, C. glabrata, C. parapsilosis, C. tropicalis)
551 Candida albicans
559 Candida glabrata
558 Candida parapsilosis
557 Candida tropicalis
127 Group B Streptococcus (GBS) Is the patient pregnant? Yes No
137 Group B Streptococcus (GBS) Antibiotic Resistance by PCR (OneSwab only)
126 Herpes subtype (HSV-1, HSV-2)
121 Leukorrhea Panel (N. gonorrhoeae*, C. trachomatis**, T. vaginalis*)
105 Chlamydia trachomatis (**Reflex to azithromycin resistance by Pyrosequencing)
167 Neisseria gonorrhoeae (*Reflex to antibiotic resistance by Molecular Analysis)
111 Trichomonas vaginalis (*Reflex to metronidazole resistance)
129 Mycoplasma genitalium (*Reflex to azithromycin & fluoroquinolone resistance by Pyrosequencing)
110 Treponema pallidum (syphilis)
320 Ureaplasma urealyticum (†Reflex to fluoroquinolone resistance by Pyrosequencing)

MALE & FEMALE SEXUALLY TRANSMITTED INFECTIONS

UroSwab (Urine specimens only) by Real-Time PCR unless otherwise specified

- 105 Chlamydia trachomatis (**Reflex to azithromycin resistance by Pyrosequencing)
121 Leukorrhea Panel (N. gonorrhoeae*, C. trachomatis**, T. vaginalis*)
129 Mycoplasma genitalium (*Reflex to azithromycin & fluoroquinolone resistance by Pyrosequencing)
167 Neisseria gonorrhoeae (*Reflex to antibiotic resistance by Molecular Analysis)
109 N. gonorrhoeae* & C. trachomatis**
110 Treponema pallidum (syphilis)
111 Trichomonas vaginalis (*Reflex to metronidazole resistance)

Applicable for adolescent females who are not candidates for pelvic exams.

BRCAcare Test Selection

*BRCAcare Testing - Whole Blood or Mouthwash

To order panel components individually, select tests beneath the panel.

BRCAcare Test Requisition and Informed Consent forms must accompany specimen

- 1235 Breast Cancer High Risk Extended Panel Plus: 14 genes (BRCA1, BRCA2, CDH1, PTEN, TP53, STK11, ATM, CHEK2, PALB2, BARD1, BRIP1, MUTYH, RAD51C, RAD51D) by Gene Sequencing with BRCA1/2 Deletion/Duplication Analysis
1279 Lynch Syndrome Gene Panel: 5 Genes (EPCAM, MLH1, MSH2, MSH6, PMS2) by Gene Sequencing with Deletion/Duplication Analysis
1221 BRCA1/2: Comprehensive BRCA Analysis by Gene Sequencing with Deletion/Duplication Analysis

Genetic Carrier Screening Test Selection

Whole Blood (ACD Solution A), OneSwab, ThinPrep or Mouthwash

To order panel components individually, select tests beneath the panel.

Genetic Carrier Screening Test Requisition with completed clinical information must accompany specimen.

- 1231 Cystic Fibrosis Core Test (23 major CFTR variants approved by ACOG/ACMG)
1232 Cystic Fibrosis Comprehensive Test by Next Generation Sequencing (191 variants of the CFTR gene, including 23 major variants approved by ACOG/ACMG)
1274 Genetic Carrier Screening Panel (3 genes) includes:
1231 Cystic Fibrosis Core Test (23 major CFTR variants approved by ACOG/ACMG) (CFTR)
1272 Fragile X Syndrome (FMR1)
1273 Spinal Muscular Atrophy (SMN1)

Other Tests/Panels:

To view all of MDL's Test Requisition forms, please visit www.mdlab.com/forms/

Commonly Used ICD-10 Diagnosis Codes for Medical Diagnostic Laboratories, L.L.C. Tests

Please indicate your Diagnosis Code selection on the front of this test requisition in the designated spaces under "Billing Information – Diagnosis Codes (Required)".

This is a general, non-comprehensive guide for use by the healthcare provider to assist in the assignment of a diagnosis code to the laboratory testing ordered. The healthcare clinician must only order tests determined to be medically necessary for the diagnosis and treatment of the patient.

Pharmacogenomics Testing:

ICD-10	Description
Cardiology	
T45.615A	Adverse effect thrombolytic drugs, initial encounter
T46.1X5A	Adverse effect of calcium-channel blockers, initial encounter
T44.7X5A	Adverse effect of beta-adrenoreceptor, initial encounter
Pain Management	
T40.4X5A	Adverse effect of other synthetic narcotic, initial encounter
T48.1X5A	Adverse effect of skeletal muscle relaxants, initial encounter
Psychiatric Disorders (Including Addiction)	
T43.505A	Adverse effect of unspecified antipsychotics and neuroleptics, initial encounter
T43.205A	Adverse effect of unspecified antidepressants, initial encounter
Immunology / Immune Modulation	
T39.395A	Adverse effect of other nonsteroidal anti-inflammatory drugs [NSAID], initial encounter
Oral Contraceptives	
T38.4X5A	Adverse effect of oral contraceptives, initial encounter
Other	
T88.7XXA	Unspecified adverse effect of drug or medicament, initial encounter
T88.7XXD	Unspecified adverse effect of drug or medicament, subsequent encounter
Z79.899	Other long term (current) drug therapy

Infectious Disease Testing:

ICD-10	Description
Sexually Transmitted Diseases	
A63.8	Other specified predominantly sexually transmitted diseases
A64	Unspecified sexually transmitted disease
Z11.3	Encounter for screening for infections with a predominantly sexual mode of transmission
Z20.2	Contact with and (suspected) exposure to infections with a predominantly sexual mode of transmission
Z20.9	Contact with and (suspected) exposure to unspecified communicable disease
Z77.9	Other contact with and (suspected) exposures hazardous to health
A49.3	Mycoplasma infection, unspecified site
A49.9	Bacterial infection, unspecified
B37.3	Candidiasis of vulva and vagina
B37.9	Candidiasis, unspecified
A56.02	Chlamydia trachomatis infection of lower genitourinary sites
A74.89	Other chlamydial diseases
A74.9	Chlamydial infection, unspecified
A60.04	Herpesviral vulvovaginitis
A63.0	Anogenital (venereal) warts
B00.89	Other herpes viral infection
B07.9	Viral wart, unspecified
A49.3	Mycoplasma infection, unspecified site
A59.01	Trichomonal vulvovaginitis
A59.9	Trichomoniasis, unspecified
Vaginitis	
N76.0	Acute vaginitis
N76.1	Subacute and chronic vaginitis
Cervicovaginal Disorders	
N72	Inflammatory disease of cervix uteri
N73.9	Female pelvic inflammatory disease, unspecified
N84.1	Polyp of cervix uteri
N87.0	Mild cervical dysplasia
Z87.410	Personal history of cervical dysplasia
N89.8	Other specified noninflammatory disorders of vagina
N88.9	Non-inflammatory disorder of cervix uteri, unspecified
Z12.4	Encounter for screening for malignant neoplasm of cervix
Z12.72	Encounter for screening for malignant neoplasm of vagina
Z12.89	Encounter for screening for malignant neoplasm of other sites
Human Papillomavirus (HPV)	
R87.610	Atypical squamous cells of undetermined significance on cytologic smear of cervix (ASC-US)
R87.611	Atypical squamous cells cannot exclude high grade squamous intraepithelial lesion on cytologic smear of cervix (ASC-H)
R87.612	Low grade squamous intraepithelial lesion on cytologic smear of cervix (LGSIL)
R87.613	High grade squamous intraepithelial lesion smear of cervix (HGSIL)
R87.619	Unspecified abnormal cytological finding in specimens from cervix uteri
R87.810	Cervical high risk human papillomavirus (HPV) DNA test positive
R87.820	Cervical low risk human papillomavirus (HPV) DNA test positive
Z11.51	Encounter for screening for human papillomavirus (HPV)
Menstruation and Bleeding	
N92.5	Other specified irregular menstruation
N93.8	Other specified abnormal uterine and vaginal bleeding
N94.89	Other specified conditions associated with female genital organs and menstrual cycle
N95.0	Postmenopausal bleeding
N95.2	Postmenopausal atrophic vaginitis
R10.2	Pelvic and perineal pain
Pregnancy and Group B Streptococcus	
Z34.80	Encounter for supervision of other normal pregnancy, unspecified trimester
Z34.90	Encounter for supervision of normal pregnancy, unspecified, unspecified trimester
Z36	Encounter for antenatal screening of mother
O99.820	Streptococcus B carrier state complicating pregnancy
O99.824	Streptococcus B carrier state complicating childbirth
O99.825	Streptococcus B carrier state complicating the puerperium
Z22.330	Carrier of Group B streptococcus
Genetic Carrier Screening	
Cystic Fibrosis	
E84.0	Cystic fibrosis with pulmonary manifestations
E84.11	Meconium ileus in cystic fibrosis
E84.19	Cystic fibrosis with other intestinal manifestations
E84.8	Cystic fibrosis with other manifestations
E84.9	Cystic fibrosis, unspecified
Z14.1	Cystic fibrosis carrier
Fragile X Syndrome	
Q99.2	Fragile X chromosome
Spinal Muscular Atrophy	
G12.9	Spinal muscular atrophy, unspecified
G12.1	Other inherited spinal muscular atrophy

For BRCA testing ICD-10 codes, please refer to the back of the Genetic Test Requisition Form.

For Pathology testing ICD-10 codes, please refer to the back of the Comprehensive Test Requisition Form.

* Reflex to antibiotic resistance by Molecular Analysis. ♦ Reflex to metronidazole resistance by Real-Time PCR. ** Reflex to azithromycin resistance by Pyrosequencing. † Reflex to azithromycin & fluoroquinolone resistance.
 ‡ Reflex to fluoroquinolone resistance by Pyrosequencing). *** This test can only be performed when the test in parenthesis is positive. All tests performed will be billed.
 OneSwab® UroSwab®, X-Plate Technology® and HPV Type-Detect® are registered in the USPTO. ThinPrep® is a trademark of Hologic, Inc.

Physicians must only order tests that they have determined are medically necessary for the diagnosis and treatment of a patient. MDL offers individual tests, as well as a limited number of customized panels. If you choose to order a panel, please make certain that each and every test is medically necessary. If you check off a panel as your choice, MDL understands that the physician has determined that all of the component tests are medically necessary, and will perform, report and bill for all such component tests.