A DIVISION OF GENESIS CLINICAL DIAGNOSTICS
A MEMBER OF GENESIS BIOTECHNOLOGY GROUP

MEDICAL DIAGNOSTIC LABORATORIES
2439 Kuser Road • Hamilton, NJ 08690-3303
(609) 570-1000 • Fax (609) 245-7665
Toll Free (877) 269-0090
www.mdlab.com

CAP ACCREDITED
COLLEGE of AMERICAN PATHOLOGISTS

Family Practice 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:

Assigned Sex at Birth (Required): ☐ Female ☐ Male | Date of Birth (Required): | Patient ID#:

Phone Number: ☐ Cell Phone ☐ Home Phone

Race: ☐ Alaska Native or American Indian ☐ Asian ☐ Black or African American ☐ Multiracial ☐ Native Hawaiian or other Pacific Islander ☐ Other race ☐ White ☐ Does not wish to disclose ☐ Not provided

Ethnicity: ☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown

Gender Identity: ☐ Male ☐ Female ☐ Gender nonconforming ☐ Transgender male-to-female ☐ Transgender female-to-male ☐ Does not wish to disclose ☐ Not provided ☐ Not applicable

Sexual Orientation: ☐ Bisexual ☐ Straight ☐ Gay or Lesbian ☐ Something else ☐ Does not wish to disclose ☐ Not provided ☐ Not applicable

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

Billing Type: ☐ Patient ☐ Insurance ☐ Client | Relation (Required): ☐ Self ☐ Spouse ☐ Dependant

Insured's Name (if not patient):

Insured's SS#: | Insured's DOB:

Primary Insurance Carrier: | Medicare, Medicaid or Policy ID#:

Claims Address:

Employer/Group Name: | Group#:

Specimen Information

Date Collected (Required):

Specimen Type: Specimen Source:

☐ ***One*Swab®**: _____
☐ **Blood**
☐ **Saliva**
☐ ***Uro*Swab®**

Was ***Uro*Swab®** collected from a catheterized patient? ☐ Yes

Sexually Transmitted Infections - *OneSwab®*

Common ICD10 codes (required):
N89.8 ☐ Other specified noninflammatory disorders of vagina
Z20.2 ☐ Contact with and (suspected) exposure to infections with a predominantly sexual mode of transmission
N76.4 ☐ Abscess of vulva
☐ Other:_____

121 ☐ Leukorrhea Panel Includes-
105 ☐ *Chlamydia trachomatis* (*Reflex to antibiotic resistance by Molecular Analysis),
167 ☐ *Neisseria gonorrhoeae* (*Reflex to antibiotic resistance by Molecular Analysis)
111 ☐ *Trichomonas vaginalis* (*Reflex to **M** resistance),
129 ☐ *Mycoplasma genitalium* (*Reflex to antibiotic resistance by Molecular Analysis)

115 ☐ Genital Ulcer Disease Panel Includes -
122 ☐ *Haemophilus ducreyi* 126 ☐ Herpes subtype (HSV-1, HSV-2) 110 ☐ *Treponema pallidum* (syphilis)

Vaginitis & Vaginosis - *OneSwab®*

Common ICD10 codes (required):
N76.0 ☐ Acute vaginitis
N89.8 ☐ Other specified noninflammatory disorders of vagina
R10.2 ☐ Pelvic and perineal pain
☐ Other:_____

759 ☐ Bacterial Vaginosis (BV) Panel with Lactobacillus Profiling by qPCR Includes -
Fannyhessea vaginae (*Atopobium vaginae*), BVAB1, BVAB2, BVAB3, *Bacteroides fragilis, Bifidobacterium breve, Megasphaera* Type 1 & 2, *Gardnerella vaginalis, Mobiluncus curtisii, M. mulieris, Prevotella bivia, Sneathia sanguinegens, Streptococcus anginosus*

182 ☐ Aerobic Vaginitis (AV) Panel Includes -
127 ☐ Group B Streptococcus (GBS) 141 ☐ *Escherichia coli*
184 ☐ *Staphylococcus aureus* 153 ☐ *Enterococcus faecalis*

560 ☐ Candida Vaginitis Panel Includes -
551 ☐ *Candida albicans* 559 ☐ *Candida glabrata*
558 ☐ *Candida parapsilosis* 557 ☐ *Candida tropicalis*
566 ☐ *Candida krusei*

134 ☐ Urogenital Mycoplasma & Ureaplasma Panel
Includes - 129 ☐ *Mycoplasma genitalium* (ᵠReflex to **AZ** & **FL** resistance by Pyrosequencing) 130 ☐ *Mycoplasma hominis*, 320 ☐ *Ureaplasma urealyticum* (*Reflex to antibiotic resistance by Molecular Analysis)

Intestinal Pathogens - *OneSwab®* Loose Stool, Rectal Swab, No Lubricant

Common ICD10 codes (required):
R19.7 ☐ Diarrhea, unspecified
R10.9 ☐ Unspecified abdominal pain
R11.2 ☐ Nausea with vomiting, unspecified
☐ Other: _____

365 ☐ *Campylobacter jejuni*
162 ☐ *Clostridium difficile* (Toxins A, B)
371 ☐ *Cryptosporidium parvum*
372 ☐ *Entamoeba histolytica*
168 ☐ *E. coli* (O157 and Shiga toxin)
370 ☐ *Giardia intestinalis*
310 ☐ *Helicobacter pylori*
274 ☐ Human Rotavirus A
158 ☐ *Listeria monocytogenes*
272 ☐ Norwalk virus (Norovirus)
160 ☐ Salmonella
161 ☐ Shigella

Refer to the back for antibiotic abbreviation key.

Other Tests/Panels:

ICD10 codes (required):

For a full menu of testing, please visit www.mdlab

Skin & Soft Tissue Infections - *OneSwab®*

Common ICD10 codes (required):
A49.9 ☐ Bacterial infection, unspecified
B99.9 ☐ Unspecified infectious disease (opportunistic)
B99.8 ☐ Other infectious disease
☐ Other: _____

366 ☐ Skin & Soft Tissue Infections (SSTI) Panel Includes -125 ☐ *Bacteroides fragilis,* 153 ☐ *Enterococcus faecalis,* 141 ☐ *Escherichia coli,* 1112 ☐ Group A Streptococcus, 127 ☐ Group B Streptococcus (GBS), 727 ☐ *Klebsiella oxytoca,* 728 ☐ *K. pneumoniae,* 1118 ☐ MRSA: Methicillin Resistant and Methicillin Susceptible (MSSA) *Staphylococcus aureus* by Conventional PCR, 1119 ☐ CA-MRSA: Community-Associated MRSA. Panton-Valentine Leukocidin (PVL) DNA**(Type IV MRSA + #1118 Req.) [Community Associated MRSA = Type IV MRSA+ and PVL+], 362 ☐ Prevotella species Group 1 (*P. bivia, P. disiens, P. intermedia, P. melaninogenica*), 363 ☐ Prevotella species Group 2 (*P. corporis, P. albensis*), 146 ☐ *Proteus mirabilis,* 174 ☐ *Pseudomonas aeruginosa,* 368 ☐ *Fusobacterium* species, 553 ☐ *Aspergillus fumigatus*

367 ☐ SSTI Panel Antibiotic Resistance Includes - [*E. faecalis*, *E. coli*, GAS, GBS, *K. oxytoca, K. pneumoniae, P. mirabilis, P. aeruginosa,* MRSA: **AC**, **AM** (for *E. faecalis*), **C**, **CL**, **D**, **TS**, **CP**, **CF**, **PT**, **I**, **G**] (153, 141, 1112, 127, 727, 728, 146, 174, or 1118 Req.) When panel is ordered and individual tests are not selected, all 9 will be performed & billed)

286 ☐ Dermatologic Viruses Panel Includes - HSV-1 & HSV-2, HPV, 219 ☐ Human herpesvirus-6 (HHV-6) Variants A & B 263 ☐ HHV-7 128 ☐ Molluscum contagiosum virus (MCV) 215 ☐ Varicella-zoster virus (VZV)

593 ☐ *Candidozyma auris*

285 ☐ Monkeypox virus (Non-variola Orthopoxvirus)

UTI & STI - *UroSwab®*

Common ICD10 codes (required):
N39.0 ☐ Urinary tract infection, site not specified
R30.0 ☐ Dysuria
R30.1 ☐ Vesical tenesmus
☐ Other: _____

176 ☐ Urinary Pathogens Antibiotic Resistance* Includes - 141 ☐ *E. coli,* 727 ☐ *K. oxytoca,* 728 ☐ *K. pneumoniae,* 146 ☐ *P. mirabilis*-**AC, C, TS, N, CP, F** 153 ☐ *E. faecalis,* 154 ☐ *E. faecium*- **A, N, CP, F, D, L** 174 ☐ *P. aeruginosa* - **CF, PT, I, A, G**
*(141, 153, 154, 728, 727, 146 or 174 Req. When panel is ordered and individual tests are not selected, all 7 will be performed & billed)

Common ICD10 codes (required):
Z20.2 ☐ Contact with and (suspected) exposure to infections with a predominantly sexual mode of transmission
R36.9 ☐ Urethral discharge, unspecified
R30.9 ☐ Painful micturition, unspecified
☐ Other: _____

SEXUALLY TRANSMITTED INFECTIONS Applicable for adolescent females who are not candidates for pelvic exams.

121 ☐ Leukorrhea Panel Includes -105 ☐ *C. trachomatis* (*Reflex to antibiotic resistance by Molecular Analysis), 167 ☐ *N. gonorrhoeae,* 111 ☐ *T. vaginalis* (*Reflex to **M** resistance), 129 ☐ *M. genitalium* (*Reflex to antibiotic resistance by Molecular Analysis)

109 ☐ *N. gonorrhoeae & *C. trachomatis****

110 ☐ *Treponema pallidum* (syphilis)

Pharmacogenomics- *whole blood or saliva*

ICD10 codes (required):

Note: This testing requires a signed Patient Informed Consent/Insurance Acknowledgment Form

3407 ☐ Depressive Disorder & Major Depressive Disorder Includes- Amitriptyline, Antidepressants (gen), Antipsychotics (gen), Aripiprazole, Bupropion, Citalopram, Clomipramine, Desipramine, Desvenlafaxine, Diazepam, Doxepin, Duloxetine, Escitalopram, Fluoxetine, Fluvoxamine, Imipramine, Maprotiline, Milnacipran, Mirtazapine, Nortriptyline, Olanzapine, Opipramol, Paroxetine, Quetiapine, Sertraline, SSRIs (gen), Trimipramine, Venlafaxine, Vilazodone, Vortioxetine (ABCB1, ADRA2A, ANKK1, COMT, CYP1A2, CYP2B6, CYP2C19, CYP2C9, CYP2D6, CYP3A4, CYP3A5, DRD4, GABRP, GRIK4, HTR2A, HTR2C, MTHFR, UGT1A1)

3102 ☐ Statins Includes- Atorvastatin, Fluvastatin, Lovastatin, Pitavastatin, Pravastatin, Rosuvastatin, Simvastatin (ABCB1, ABCG2, APOE, CYP2C9, CYP2D6, CYP3A4, CYP3A5, KIF6, MTHFR, SLCO1B1)

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

My signature below acknowledges the patient has been informed about the purpose, limitation and possible risks of genetic testing. The patient has been given the opportunity to ask questions about this consent and seek outside genetic counseling.

If the genetic 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's signed informed consent 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: _____

HPV Testing - *OneSwab® or ThinPrep®*

Common ICD10 codes (required):
Z01.411 ☐ Encounter for gynecological examination (general) (routine) with abnormal findings
Z01.419 ☐ Encounter for gynecological examination (general) (routine) without abnormal findings
Z11.51 ☐ Encounter for screening for human papillomavirus (HPV)
☐ Other:_____

739 ☐ HPV Type-Detect® 4.0 by Multiplex Real-Time PCR (Includes HPV Types:16, 18, 45, 58, 35, 51, 52, 56, 39, 31, 33, 59, 68)

795 ☐ HPV Type-Detect® 4.0 (High Risk Subtypes only, reflex to Vaginal Dysbiosis)

Antibiotic Abbreviations Key

A = aztreonam **AC**= amoxicillin-clavulanic acid, **AP** = ampicillin, **AZ** = azithromycin, **CC** = ceftriaxone/cefixime, **C** = cephalothin (cephalexin), **CF** = cefepime, **CP** = ciprofloxacin, **CL** = clindamycin, **D** = doxycycline, **F** = fosfomycin, **FL** = fluoroquinolone **G** = gentamicin, **I** = imipenum, **L** = linezolid, **M** = metronidazole **N** = nitrofurantoin, **PT** = piperacillin-tazobactam, **TS** = trimethoprim-sulfamethoxazole.

Medical Necessity Guidelines:

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. MDL provides practitioners with the flexibility to choose appropriate individual tests for each specimen to assure that the convenience of ordering panels does not impede them from ordering tests/panels that are medically necessary. All tests listed in panels may be ordered individually using this test requisition form. 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.

** This test can only be performed when the test in parenthesis is positive. All tests performed will be billed.

OneSwab® & *UroSwab®* are registered in the USPTO.

Test by Real-Time PCR unless otherwise specified.

Specimen Collection Platform		TAT*	Stability	Test Additions¥	Specimen Collection
OneSwab®		24 - 72 hours	7 days	30 days to add tests	1. Collect specimen with the sterile swab provided. 2. Insert swab into the transport media, break off swab handle and tightly re-secure the cap on the transport media vial.
UroSwab®		24 - 72 hours	4 days	14 days to add tests	1. Have patient collect a urine specimen in a collection cup. 2. Dip the sponge swab into collection cup to absorb the urine. 3. Tightly re-secure the cap on the vial.
Blood	Yellow top tube (ACD solution A)	3 - 5 days	48 Hours	30 days to add tests	1. In accordance with the standard operating procedure of your facility, collect blood in two yellow top (ACD solution A) tubes. 2. Allow the tubes to fill properly to ensure the proper blood to anticoagulant ratio. 3. Invert gently several times to mix and prevent clot formation. Do not shake the tubes. Do not centrifuge.
Saliva		5 - 10 days	48 Hours	--	• Vigorously rinse your mouth with clean water 5 minutes prior to specimen collection (30 minutes prior is ideal). • After rinsing, do not brush your teeth, use mouthwash, eat, drink, chew gum or smoke prior to sample collection. 1. Begin collecting your sample by allowing saliva to pool in your mouth. Then spit into the wide funnel of the tube allowing saliva to collect in the upper chamber of the tube. Fill the tube until the amount of saliva (not bubbles) reaches the fill line as shown. 2. Once filled, unscrew the funnel allowing the saliva to flow into the lower chamber of the tube containing the stabilizing solution. Discard the funnel. 3. Use the blue cap to close the tube tightly. Shake the capped tube for 5 seconds.

** Up to 72 hours with reflex/antiobiotic resistance testing* *¥ Pending QC review for sufficient specimen volume*

Specimen Packaging:

1. Label every vial with a minimum of 2 patient identifiers including the patient's name and date of birth. Be sure the name on the vial is written exactly the same way as on the test requisition form.
2. Place the vial into the Styrofoam/Cardboard container. You can fit up to 3 vials in one container.
3. Place the Styrofoam/Cardboard container into the central pocket of a biohazard bag containing absorbent material.
4. Place a completed test requisition form for each vial in the front pocket of the biohazard bag.
5. Place the biohazard bag into the prepaid Lab pack Envelope that has a preaddressed airbill attached. One envelope will accommodate 6-7 containers. Package as many containers in one Labpack as possible.
6. Be sure to seal the Lab pack by removing the plastic from the top of the adhesive.

Specimen Pick-up:

- If you have a specimen pick-up for a local courier in the NJ, PA, DE, D.C., MD, VA, KS, MO or Phoenix, AZ area, please call (877) 205-0005 no later than 2 hours prior to the closing of your facility.
- If you have a specimen pick-up, please call your sales representative or MDL customer service at 877.269.0090 no later than 2 hours prior to the closing of your facility.

Helpful Hints Checklist

Please review these helpful hints to reduce specimen discrepancies and enhance turnaround time.

Verify Patient Name - did you:
- ✓ attach the correct demographics sheet?
- ✓ write the patient's name on the requisition form?

Patient Name Matches on Vial & Requisition Form- did you:
- ✓ make sure names on vial and requisition form match?
- ✓ list the patients married or maiden name?
- ✓ list a nickname by mistake?

Verify Date of Collection- did you:
- ✓ write the correct year?
- ✓ write the correct month?
- ✓ list the date of birth instead?

Verify Tests- did you:
- ✓ clearly mark each box?
- ✓ order tests appropriate for the specimen type?

No Tests Ordered- did you:
- ✓ mark the boxes for the tests/panels ordered?

Supply Orders:

Easily place supply orders online by visiting our website:

http://www.mdlab.com/physicians/supplies/#

Supply orders may also be placed by contacting our Client Services department toll free 877.269.0090 or by fax 609.570.1050. Supply requests are processed and shipped on a daily basis. Please allow 3 to 5 business days for delivery, depending upon your location.

MDL Contact Information	TOLL FREE	FAX
GBS Hotline *24 hours - 7 days a week* *Group B Strep & HSV results only*	877.MDL.GBS7 877.635.4277	
Quality Control Department *For Technical Assistance*	877.269.0090	609.245.7665
Client Services *General Questions, Results*	877.269.0090	609.570.1050
Client Services *Billing Questions*	877.333.9233	609.245.7683