



Maryland Department of Health  
Office of Health Care Quality  
Laboratory Licensing Programs  
7120 Samuel Morse Drive  
Second Floor  
Columbia, Maryland 21046  
Phone: 410.402.8025 Fax: 410.402.8213

## Instructions for Completion of State Compliance Application

Changes to your current State laboratory license must be submitted on the Laboratory Licensing Change Form. Forms can be downloaded on our website

It is important that you fill out this application completely, including signatures where required. Original (ink) signatures are required on all initial applications and must be mailed or hand delivered to our office (address listed above). If the application is incomplete, it will delay the licensing process. Initial applications are not accepted by fax or email.

Please allow 3-4 weeks for permit processing and mailing

There is no fee for this licensure.

If you have any questions, please call the Laboratory Licensing Division at (410) 402-8025.

**\*\*\*Important\*\*\***

**\*\*\*Before submitting your application, please review the checklist on the last page.\*\*\***

<b>Maryland Department of Health</b> <b>Laboratory Licensing Programs</b> <b>Office of Health Care Quality</b>	<b>Date/Amount Paid</b>	Office use only
	<b>Invoice #</b>	Office use only
	<b>Check #</b>	Office use only
	<b>State Permit #</b>	Applicant, if known please enter
	<b>CLIA #</b>	Applicant, if known please enter

## State Compliance Application

☐ Initial Application

☐ Reinstatement

<b>I. Laboratory Information</b>			
<b>Type of Laboratory</b> <input type="checkbox"/> <b>Physician Office</b> <input type="checkbox"/> <b>Point of Care</b> <input type="checkbox"/> <b>Independent/Reference</b> <input type="checkbox"/> <b>Hospital</b>			
Laboratory Practice/ Entity Name		Contact Person Name/Phone Number	
Address, City, State and Zip Code	Email Address	Fax	
Mailing address if different from above			
<b>II. Director Information</b>			
Laboratory Director Name	Degree	Full Time	Part Time (hours/week)
Certification by American Specialty Board (Name, Date, Number)		State Medical License Number	
<b>III. Laboratory Supervisor/Consulting Supervisor/Manager Information</b>			
Name	Degree	Full Time	Part Time (hours/week)
Certification by American Specialty Board (Name, Date, Number)			

## IV. Schedule A - General Permit

\*\*\* If you are only performing tests on Excepted list, Schedule B, do not use this section\*\*\*

<b>Chemistry</b>  <input type="checkbox"/> Routine <input type="checkbox"/> Blood Gas <input type="checkbox"/> Endocrinology <input type="checkbox"/> Toxicology: Drugs of Abuse <input type="checkbox"/> Toxicology: Therapeutic <input type="checkbox"/> Toxicology: Heavy Metals <input type="checkbox"/> Radioimmunoassay	<b>Genetics</b>  <input type="checkbox"/> Routine <input type="checkbox"/> Molecular <input type="checkbox"/> Cytogenetics	<b>Forensic Toxicology</b>  <input type="checkbox"/> Toxicology: Job Related	<b>Microbiology</b>  <input type="checkbox"/> Bacteriology <input type="checkbox"/> Parasitology <input type="checkbox"/> Mycology <input type="checkbox"/> Mycobacteriology <input type="checkbox"/> Virology	<b>Health Awareness *</b>  <input type="checkbox"/> Cholesterol/Lipids <input type="checkbox"/> Glucose Finger Stick <input type="checkbox"/> Hemoglobin A1c  <small>* performed at health fairs not routine chemistry lab *must be CLIA waived</small>
<b>Immunohematology/ Blood Bank</b>  <input type="checkbox"/> ABO/Rh/Non Trans- fusion/Transplant <input type="checkbox"/> ABO/Rh <input type="checkbox"/> Antibody Detection <input type="checkbox"/> Antibody Identification <input type="checkbox"/> Compatibility Testing	<b>Hematology</b>  <input type="checkbox"/> Routine <input type="checkbox"/> Coagulation <input type="checkbox"/> CLIA Waived CBC (Sysmex)	<b>Molecular Biology</b>  <input type="checkbox"/> Nucleic Acid Probes <input type="checkbox"/> PCR Amplifications <input type="checkbox"/> Recombinant Nucleic Acid Techniques	<b>Pathology</b>  <input type="checkbox"/> Histopathology <input type="checkbox"/> Dermatopathology <input type="checkbox"/> Oral Pathology <input type="checkbox"/> Cytology-GYN <input type="checkbox"/> Cytology-Non- GYN	<b>Immunology</b>  <input type="checkbox"/> General Immunology <input type="checkbox"/> Syphilis Serology <input type="checkbox"/> Histocompatibility

## V. Schedule B - Excepted Tests \*

\* Note: Not all tests excepted by Maryland regulations are waived by CLIA. You can check the test categories for CLIA at <http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfclia/search.cfm>

<b>Chemistry</b> <input type="checkbox"/> CLIA waived blood lipid analysis for cholesterol, HDL, LDL, and triglycerides. <input type="checkbox"/> Dipstick Glucose <span style="float: right;">BNP <input type="checkbox"/></span> <input type="checkbox"/> Dipstick Urinalysis <span style="float: right;">Microscopic Urinalysis <input type="checkbox"/></span> <input type="checkbox"/> Dipstick Microalbumin & creatinine, urine <input type="checkbox"/> Fructosamine (whole blood) <input type="checkbox"/> <input type="checkbox"/> Glucose (FDA Home Device) <input type="checkbox"/> Hemoglobin A1c (Glycohemoglobin) <input type="checkbox"/> Waived Whole Blood Lead Testing <input type="checkbox"/> CLIA Waived Urine Drug Screen	<b>Hematology</b>  <input type="checkbox"/> Fern Test <input type="checkbox"/> Hematocrit <input type="checkbox"/> Hemoglobin <input type="checkbox"/> Nitrazine Test <input type="checkbox"/> Semen analysis, qualitative <input type="checkbox"/> Sick Cell Testing <input type="checkbox"/> CLIA Waived PT/INR
<b>Immunology</b>  <input type="checkbox"/> Bladder marker, H-related protein, qualitative <input type="checkbox"/> H.Pylori (whole blood) <input type="checkbox"/> Heterophyle AG (whole blood) <input type="checkbox"/> Mono Slide Test <input type="checkbox"/> NMP Bladder Marker, qualitative <input type="checkbox"/> Rheumatoid Factor <input type="checkbox"/> Urine Pregnancy Test	<b>Microbiology</b>  <input type="checkbox"/> Dermatophyte Screen <span style="float: right;">Trichomonas vaginalisantigen <input type="checkbox"/></span> <input type="checkbox"/> Bacterial Sialidase <input type="checkbox"/> Gram Stain <span style="float: right;">Adenovirus antigen eye fluid <input type="checkbox"/></span> <input type="checkbox"/> Group A Strep Screen (non-culture) <input type="checkbox"/> Influenza Antigen (nasal or throat swab) <input type="checkbox"/> KOH Preparation <input type="checkbox"/> Occult Blood <input type="checkbox"/> Occult Blood, gastric <input type="checkbox"/> Pinworm Prep <input type="checkbox"/> Urine Colony Count (no ID) <input type="checkbox"/> Wet Mount

## VI. Mandatory, You Must List Testing Instrumentation and Test Kits Used in the Laboratory

\*\*\*Please also include test discipline/subdiscipline (e.g. Chemistry-Routine) if using Schedule A\*\*\*


## VII. Proficiency Testing

☐ I am not enrolled

☐ I am enrolled (complete below)

Name of Company

Discipline


## VIII. Ownership Information

### A. Type of Entity

☐ Sole Proprietorship   ☐ Partnership   ☐ Corporation   ☐ Unincorporated Association  
☐ Other (Specify) \_\_\_\_\_

B. This section is MANDATORY, application will be returned if left blank. Social Security Number is unacceptable  
Attention- Laboratories not located in Maryland, the EIN must match what you have on file in the CMS CLIA database. Only include one EIN Number below, not several please.

Name	Address	EIN Federal Tax ID

### IX. Attestation

I certify that the information provided in this application is true and complete, understanding that any knowing and willful false statement or representation, or failure to fully and accurately disclose the requested information in this application, may be prosecuted under applicable federal or State laws, may lead to a denial, suspension or revocation of the medical laboratory license for this entity, or could result in termination of participation in State or federal reimbursement programs. I further understand that compliance with State laws may not assure compliance with federal laws.

\_\_\_\_\_  
Signature of Laboratory Director

\_\_\_\_\_  
Date

**For Informational Purposes Only**  
**Examples of Testing for Schedule A- General Permit (Do Not Circle)**

**Chemistry**

Alkaline Phosphatase  
Amylase  
B-HCG (quantitative)  
Blood Lead  
CK-MB  
Digoxin  
Iron  
Lipase  
Phenytoin  
T4-Free  
Troponin  
TSH  
Vitamin D

**Genetics**

Chromosome Analysis  
FISH Studies (Neoplastic and Congenital)  
Fragile X Screen  
Gaucher Disease (*GBA*) 8 Mutations  
Tay-Sachs (*HEXA*) 7 Mutations  
Y Chromosome Deletions

**Forensic Toxicology**

Job Related Alcohol  
Job Related Drugs of Abuse

**Microbiology**

AFB Smear  
Bacterial Culture  
Blood Culture  
CSF Bacterial Antigen  
Fungus/Yeast Culture  
Ova and Parasite  
Sensitivity Testing  
Viral Culture

**Hematology**

APTT  
CBC  
Differential  
Fetal Hemoglobin  
Fibrinogen  
INR  
Prothrombin Time  
Reticulocyte Count  
Sedimentation Rate

**Molecular Biology**

Adenovirus PCR  
BD Affirm Probe Test  
Chlamydia PCR  
EBV PCR  
HCV Genotyping  
HIV Drug Resistance Genotyping  
HIV Viral Load

**Pathology**

Dermatopathology  
Fine Needle Aspirations  
Grossing  
Histopathology  
Oral Pathology  
Other Cytology  
Pap Smear Interpretations

**Immunology**

Anti-Nuclear Antibody  
Epstein Barr Antibodies  
GM1 Antibody  
Hepatitis B Surface Antibody  
Hepatitis B Surface Antigen  
Herpes Antibody  
HIV Antibody  
Lyme Antibody  
Non Transplant Related Histocompatibility

To prevent a delay in processing your application please check to make sure all of the following are included:

- ☐ Completed application with each section completely filled out
- ☐ Signature of Laboratory Director must match the name in section II of application
- ☐ If the status of your CLIA certificate is changing, a completed CMS 116 form must be submitted
- ☐ Director Qualifications
  - ☐ Copy of CV, Diploma (highest degree), ECFMG (if applicable), board certification for MD or PhD (if applicable)
- ☐ Technical Supervisor Qualifications (for the discipline of HISTOLOGY)
  - ☐ Copy of American Pathology Board certification in Anatomical Pathology
  - ☐ Copy of Maryland (Board of Physicians) license to practice medicine
- ☐ Genetics Testing
  - ☐ Copy of Technical Supervisor's diploma (must be MD, DO or PhD), board certification from the American Board of Medical Genetics or 4 years of verified (not self-generated) experience in clinical genetics and CV
  - ☐ Copy of Test Menu
  - ☐ Copy of a Validation Study of one test (includes a summary and raw data)
  - ☐ Letter from Director documenting that the lab does not perform "Direct to Consumer" testing
- ☐ Certificate of Accreditation Laboratories
  - ☐ Copy of enrollment verification from the designated accrediting organization

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### Applicants Located in Maryland

- ☐ Completed CLIA application in agreement with State application
- ☐ Copy of Director's Maryland (Board of Physicians) license to practice medicine
- ☐ For High Complexity Laboratories: Documentation of training, education and previous experience that meets CLIA Sec. 493.1443: Standard: Laboratory Director Qualifications
- ☐ For Moderate Complexity Laboratories: Board Certification or Documentation of 20 CME from approved programs for Medical Director that meets CLIA Sec. 493.1405
- ☐ Documentation of licensure as a practitioner seeking a Letter of Exception (midwife, nurse practitioner, etc.)

### Applicants Located Out of State

- ☐ Copy of CLIA certificate and State Laboratory License, if applicable
- ☐ Copy of most recent survey, which includes cited deficiencies and corrective actions
- ☐ Copy of Director's State license to practice medicine from the State where the laboratory is located
- ☐ Documentation of training, education and previous experience that meets CLIA Sec. 493.1443: Standard: Laboratory Director Qualifications
- ☐ Proof of most recent participation in annual GYN cytology proficiency testing