

How to designate PT providers and products

Updated November 2025

The proficiency testing (PT) products listed meet the New York State (NYS) PT requirements. The Category Specific Help link provides information on NYS mandated analytes that require PT participation. Please consult the PT provider's website and catalog to determine if you are designating an appropriate product for your methodology and instrumentation, and to find additional important details.

Please contact the PT Administration Group by email at CLEPPT@health.ny.gov with any questions regarding PT product enrollment and/or designation.

Note that you can save your work at any time. The information you entered will still be available if you need to come back to the page, even after logging out.

Login to eCLEP through the Health Commerce System (HCS) home page.

PLEASE LOGIN TO BEGIN USING THE HEALTH COMMERCE SYSTEM (HCS)



User ID

Password

Forgot Your [User ID](#) or [Password](#) ☐ Remember User ID

LOGIN

Don't Have An Account? [Sign Up Here](#)

On the HCS home page, under **My Applications**, select **eCLEP**.

My Applications

Acronyms & Abbreviations	
eCLEP	i
Emergency Contacts	
ServNY	i

Refresh My Applications List
↻

On the **Welcome to e-CLEP** home page, under **Proficiency Testing**, select **PT Designations**.



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New York State Department of Health

eCLEP
electronic Clinical Laboratory Evaluation Program

home

Welcome to e-CLEP
This site contains a collection of tools and resources to assist you in meeting the requirements of the New York State Clinical Laboratory Reference System.
For general information and guidance, please refer to the [Wadsworth Center Public Website](#).

Date	Priority	Message
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Permit Materials
Laboratory Reapplication
Laboratory Changes

Proficiency Testing
PT Designations
PT Documents

Gross Annual Receipts
Reporting

LDT Approval
Test approval materials for permitted and non-permitted laboratories

Survey
Tools and information related to laboratory surveys

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BSAR

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Alternately, on the **Proficiency Testing (PT) – Home** page, select **PT Designations**.

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home > proficiency testing

Select Facility

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[PT Home](#)

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PFI: Name:

Proficiency Testing (PT) - Home

Use the links below to designate PT providers or to view PT related documents.

- **PT Designations**, when open, is used to notify CLEP of both the PT Provider(s) and product(s) the laboratory intends to enroll with for all tests/analytes offered that are either listed in CMS 42 CFR 493 subpart I (CLIA subpart I) OR defined by NYS as requiring PT (collectively, NYS mandated PT analytes).
- **PT Documents** include important time-sensitive information for your laboratory including verification of PT enrollment and PT participation issues, and PT performance.

[PT Designations](#)

[PT Documents](#)

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Either choice will bring you to the **Proficiency Testing (PT) – Designations** home page.

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home > proficiency testing > pt designations

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Proficiency Testing (PT) - Designations

Use the links at the left to begin or continue the designation process, view instructions, or find additional help regarding specific PT products.

Laboratories seeking or holding a New York State (NYS) clinical laboratory permit through the Clinical Laboratory Evaluation Program (CLEP) must enroll in proficiency testing (PT) as defined by NYS (NYS mandated PT). NYS mandated PT includes all tests/analytes offered by the laboratory that are either listed in CMS 42 CFR 493 subpart I (CLIA subpart I) OR defined by NYS as requiring PT.

In addition, each laboratory must designate the PT products that will be used to satisfy these PT requirements. Laboratories currently holding a NYS CLEP permit or those in applied status for a permit must designate these PT products via this website. This process applies ONLY to your laboratory's PRIMARY test method.

This website lists tests that require PT (a.k.a NYS mandated PT analytes) and prescreened PT products offered by the CMS-approved PT Providers that have been screened by CLEP to identify those that meet New York State PT requirements for NYS mandated PT analytes. **Note that ONLY those PT products listed are acceptable.**

For each test, your laboratory must indicate whether the test is offered on NYS specimens and, if so, indicate a PT Provider and product. You must click on the links at the left side of the page to progress through the process. You must click "submit" on the final page before the notification period ends.

- Failure to designate and submit your planned enrollment may result in citation under NYS Clinical Laboratory Standard of Practice **Proficiency Testing Standard of Practice 1 (PT S1): Enrollment, Department Notification and Participation.**
- Laboratories are required to authorize their PT Providers to submit PT results to NYS. **Be sure to provide the PT Provider with the NYS PFI number.** If your PT Provider requests a contact name and fax number, please indicate Beverly Rauch at fax number (518) 408-8666. **Failure to authorize release of PT results to NYS may result in citation under NYS Clinical Laboratory Standard of Practice **Proficiency Testing Standard of Practice 2 (PT S2): Authorized Release of Proficiency Testing Results.****
- CLEP will compare your planned enrollment to the PT scores we receive from the PT Providers in January. **Failure to participate in the PT products selected here may result in citation under NYS Clinical Laboratory Standard of Practice **Proficiency Testing Standard of Practice 1 (PT S1): Enrollment, Department Notification and Participation.****
- Laboratories must participate with their chosen provider(s) for the entire calendar year before designating a different program.

NOTE: The laboratory must comply with **Proficiency Testing Standards of Practice 3 (PT S3): Alternative to Proficiency Testing** for all tests not subject to NYS PT requirements (e.g., those not indicated as a NYS mandated PT analyte).

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Only categories your laboratory holds or has requested (is in applied status for) that contain analytes requiring PT participation will appear in the drop-down menu on the page labeled “Indicate tests Offered on NYS Specimens”.

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Designation Steps

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- Step 4. Submit designations

PFI:
Name:

Indicate Tests Offered on NYS Specimens

PT designation period is closed.

Laboratories applying for or holding a New York State (NYS) clinical laboratory permit through the Clinical Laboratory Evaluation Program (CLEP) must enroll in proficiency testing (PT) as defined by NYS (NYS mandated PT). NYS mandated PT includes all tests/analytes offered by the laboratory that are either listed in CMS 42 CFR 493 subpart I (CLIA subpart I) OR defined by NYS as requiring PT.

Laboratories offering these tests on NYS specimens must designate which PT provider and product they will use to satisfy these requirements for the upcoming calendar year. For a list of NYS-acceptable PT providers and surveys, please visit:
<https://www.wadsworth.org/regulatory/clep/pt/provider-search>.

Categories Requiring PT: ▼

Ther. Sub. Mon./Quant. Tox.
Toxicology - Blood Lead-Comprehensive

Help/Instructions
See the Category Specific

[Category Specific Help](#)

Show 40 ▼ entries

Search:

Name	Test Status
No Data	
Name	Test Status

Showing 0 to 0 of 0 entries (filtered from 16 total entries)
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You must save your work for EACH permit category by clicking the 'Save' button at the bottom of the page before choosing a different permit category or clicking a different link. Any unsaved data will be lost.

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home > proficiency testing > test selection Select Facility

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PFI: **Name:**

Indicate Tests Offered on NYS Specimens

PT designation period is open from 11/03/2022 08:00 till 11/30/2022 17:00.

Laboratories applying for or holding a New York State (NYS) clinical laboratory permit through the Clinical Laboratory Evaluation Program (CLEP) must enroll in proficiency testing (PT) as defined by NYS (NYS mandated PT). NYS mandated PT includes all tests/analytes offered by the laboratory that are either listed in CMS 42 CFR 493 subpart I (CLIA subpart I) OR defined by NYS as requiring PT.

Laboratories offering these tests on NYS specimens must designate which PT provider and product they will use to satisfy these requirements for the upcoming calendar year. For a list of NYS-acceptable PT providers and surveys, please visit:
<https://www.wadsworth.org/regulatory/clep/pt/provider-search>.

Categories Requiring PT: Toxicology - Blood Lead-Comprehensive ▼

Help/Instructions—

Toxicology - Blood Lead-Comprehensive

- Refer to Category Specific Help for additional information
- Laboratories are required to enroll in a program(s) that includes:
 - a minimum of five samples per testing event.
 - three shipments per year
 - samples for all CLIA subpart I analytes as listed in the Category Specific Help document.

[Category Specific Help](#)

Show 40 ▼ entries Search:

Name	Test Status
blood lead	Test Offered ▼
Name	Test Status

Showing 1 to 1 of 1 entries (filtered from 16 total entries)

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Step 2. Designate PT provider and product

Click on “Step 2. Designate PT provider and product”

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PFI: Name:

Designate PT provider and product

Next, please choose a PT provider and PT product. Approved PT products must include at least 5 samples provided 3 times per year (except for Mycobacteriology, which is 2 times per year). This requirement also applies to laboratories offering these tests using waived kits/devices. For a list of NYS-acceptable PT providers and products, please visit <https://www.wadsworth.org/regulatory/clep/pt/provider-search>.

Categories Requiring PT: Toxicology - Blood Lead-Comprehensive ▼

Show 40 ▼ entries

Search:

PT Designations - PRIMARY METHOD

Test Name	Provider	Product
blood lead	WI State Laboratory of Hy	Blood Lead - 01080 ▼
Test Name	American Proficiency Institute College of American Pathologists WI State Laboratory of Hygiene	Product

Showing 1 to 1 of 1 entries 1 row selected

*Your PT Provider will not be saved unless you have also chosen a product.

Save Clear

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Select a category from the drop-down menu.

- The message 'no data' means that your laboratory has selected "Test Not Offered" for all tests that require PT participation in this permit category.
- Select your PT provider from the 'Provider' drop-down menu for each "Test Offered" from Step 1.
- The 'Product' drop-down menu will automatically populate with appropriate product(s) offered by that provider.
- Select the PT product in which you have enrolled or intend to enroll for each "Test Offered".
- Click the 'Save' button for each permit category before selecting another category or leaving this page.

NOTES

- Designate PT ONLY for your primary method. Your laboratory may enroll in additional PT for secondary methods for these tests. Do not designate them here.
 - While you may order and receive additional PT samples of your designated PT Product to assess secondary methods, it is critical that the additional materials and secondary method(s) be tested AFTER you have submitted results for the PT event AND the PT event deadline has passed.
- PT for waived testing performed in a comprehensive clinical laboratory regulated by CLEP must comply with CLIA requirements (generally, five PT samples per event and three events per year). PT products that DO NOT meet these criteria are omitted from the PT Designation drop-down menus.
- Enrolling in PT products that are not listed in the PT Designation drop-down menus may result in citations under New York State Clinical Laboratory Standard **Proficiency Testing Standard of Practice (PT S1) Enrollment, Department Notification and Participation**. PT Products that are not listed in the PT Designation drop-down menus have not been approved to meet PT requirements by CLEP.

Repeat Steps 1 and 2 for all permit categories

Click on “Step 3. View designations”

This page summarizes the information you designated and any records on PT provider and/or product on file.

- Provider and product designations include both those from previous year(s) and those modified in Step 2. For modified records, both additions (new “Test Offered”) and deletions (“Test Not Offered”), are denoted by a ‘+’ in the chart.
- Please review and make any necessary changes before submitting. Changes can be made by returning to Steps 1 and 2.
- Print this page for your records.
- Alternately, use the “Download” buttons to save a csv file to your computer.

Step 4. Submit Designations

Click on “Step 4. Submit Designations”

- If there are tests listed on this page, you either did not select a test status in Step 1 or you did not designate a PT provider and/or product in Step 2, or both. Return to the steps to enter the required information.
- When there are no tests listed on this page, read the **Attestation** statement, and then check the box to indicate your agreement.
- Once you click the ‘Submit’ button to complete the PT designation process you will no longer be able to make changes.

The screenshot displays the Wadsworth Center eCLEP (electronic Clinical Laboratory Evaluation Program) interface. The header includes the Wadsworth Center logo and the eCLEP title. A navigation bar contains tabs for Permit Materials, Proficiency Testing (selected), Gross Annual Receipts, LDT Approval, Survey, Blood Resources, and Tools. A left sidebar lists various links, with 'Step 4. Submit designations' highlighted in a red box. The main content area is titled 'Submit designations' and shows a message that the PT designation period is closed. It lists tests under the category 'Mycology': Antigen ID of fungi, Culture ID of fungi, and Molecular ID of fungi. Below this is an 'Attestation' section with a statement to be read and a checkbox for agreement. A comment box is provided at the bottom, and a 'Submit' button is visible.

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Submit designations

PFI: Name:
PT designation period is closed.

Additional information is required for all tests listed below. You must complete the information required in order to submit designations. Review steps 1 and 2 to assure that all information is included.

Category	Test Name
Mycology	Antigen ID of fungi
	Culture ID of fungi
	Molecular ID of fungi

Attestation
Please read the following attestation carefully. If you agree, signify by clicking the checkbox (* required), then click 'Submit'.
I understand that signing and submitting this record in this fashion is the legal equivalent of having placed my handwritten signature on the submitted record and this affirmation. I understand and agree that by electronically signing and submitting this record in this fashion I am affirming to the truth of the information contained therein.
I, the laboratory director or delegated submitter, attest that my laboratory has enrolled or will enroll in the PT surveys designated here. I understand that these designations are binding for the coming calendar year and that failure to enroll and participate in these surveys for these analytes may result in regulatory sanctions. I further understand that satisfactory PT performance is required to maintain my laboratory's CLEP permit.
☐ I have read, and agree with, the above attestation.

Comment
If you would like to provide a comment, please do so below: (500 characters max) Characters Remaining: 500

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