

Specimen Collection and Handling

With DLO, you're good to GO

At DIO, we understand how critical each specimen and test result can be in managing your patients' health. We take all possible care to maintain specimen integrity from the moment it is picked up through test completion. Specimens are picked up, packaged, tracked and delivered directly to the laboratory by our reliable, efficient Route Service Representatives (RSRs), thereby minimizing the need for follow-up, thus freeing time for your staff.

About this section

This section will ac quaint you with DIO's specimen handling process.

Electronic resources for testing and specimen collection

- Quest Diagnostics T est Direct ory
- IntelliTest Manag er[™]
- Virtual Test Guide on dlolab .com

Specimen Handling and Transport Overview

Blood Specimen Collection

Microbiology Specimen Collection

Cytology Specimen Collection

For additional assistance with test ordering, please contact DLO's Customer Support Center at 800.891.2917, option 2.

Specimen Collection and Handling

Quality results depend on quality specimens

Quality results begin with the manner in which specimens are collected and prepared for testing. With a comprehensive menu of more than 3,500 tests, DIO and Quest Diagnostics perform testing on a wide range of sample types. Properly collecting and preparing patient specimens ensur es y ou get the results y ou need to care for your patients.

Specimen Collection Requirements

Refer to the digital tools explained below for expanded instructions on patient preparation and laboratory specimen collection procedures for individual tests.

Test Directory

Information on all tests off ered through DLO/ Quest

DLO's Virtual Test Guide

Complete test and specimen c ollection guide for frequently ordered and DLO specific t ests

dlolab.com/virtual-test-guide or dlolab.com/vtg

Directory of Services

Testing and specimen c ollection information with helpful explanations for standard Quest policies and pr ocedures

Intellitest Manager™

Online tool to access new test information, test updat es and changes

intellitestmanager.com

Ouanum™ Solutions

View specimen c ollection requirements at time of or der proces sing

Specimen requirements include information such as specimen volume collection and transport containers as well as transport temperature.

Adequate specimen volume must submitted for analysis. The volume listed is enough for initial analysis as well as for any confirmatory tests that must be performed. If an inadequate specimen is submitted, we may not be able to perform the initial test or required confirmatory procedures.

Patient Preparation

Many tests require that the patient be prepared in some specific way to ensure useful results. Please refer to the digital tools previously explained or call Customer Services for clarification of any patient preparation that might be needed.

A fasting specimen is preferred for the majority of tests performed on serum, plasma, or whole blood. Non-fasting specimens often contain fat particles that can interfere with many analytical procedures.

Supplies

Specimen collection devices supplied by DLO are to be used only for the collection of specimens for processing by DLO. Supplies are not to be used to store or dispose of biological materials, including sharp instruments, or for any activity not connected with the collection of specimens for processing by

Specimens collected and/or transported in expired collection or transport devices may be rejected. Routinely check to ensure your supplies are not outdated.

Health and Safety Precautions

Specimens should be handled in a safe manner and according to applicable legal requirements or guidance. Information on safe specimen handling may be obtained from the US Occupational Safety and Health Administration (OSHA) and the Centers for Disease C ontrol and Prevention (CDC).

Additional Details and Instructions

- Specimen leak age or contamination of collection device
- Specimens should ne ver be frozen in glass tubes
- No needles or other sharps in the package which could cause injury or pathog enic exposure

DLO reserves the right to refuse to accept any transports that pose a safety hazard to its employees.



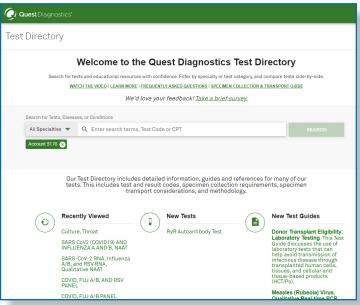
Quest Test Directory

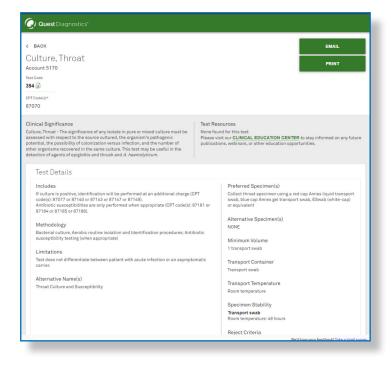
Improved access to Quest's latest testing information

Quest Diagnostics' Test Directory is a valuable resource available to providers and their staff for testing information. Links to the Test Direct ory can be found on the dlolab.com homepage, in the Tools and Resources for Providers, and throughout the Virtual Test Guide.

On your first visit to the Test Directory, select "Click Here" under "Ordering from a Quest Diagnostics affiliate" and then select DIO to save your service area. Any sear ches will now test information available for DIO.







Quest's Test Directory provides comprehensive information for all tests available through DLO/Quest.

- Search engine dedicat ed to the Test Center
- DIO/ Quest t est name
- Test c ode(s)
- Billing CPT ode(s)
- Additional testing or reflex criteria
- Methodology
- Limitations
- Reference ranges
- Clinical significance
- Link to FAQs, algorithms, test reference material orrelated articles
- Preferred and ac ceptable alt ernative specimens
- Links to related sections of quest diagnostics. com

Intellitest Manager™

Easily manage the test changes that are most important to you

IntelliTest Manager is a flexible online tool that provides best-in-class features for accessing new test information, test updates and changes based on specific account utilization. Clients can simply visit **intellitestmanager.com** and log in with their client number and 5-digit zip code.

Get the test update data you need in the format you want with IntelliTest Manager

Features

Filter by the utilization of multiple accounts

Browse and perform keyword search across all updated tests

Manage recipients of email notifications about lab updates

Export information in the product-specific format specified by your EMR or LIS wendor

Filter and browse tests by specific client utilization

Customize the view by selecting and hiding data fields

Sort information based on the following: new tests, CPT code, specimen requirements, transport temperature, specimen stability, reference range or methodology

View test chang e documents online, 2 4/7

See t est updat e history with effective date range

See detailed information for updated test(s), including specimen requirements and effective dates

Export and download list of all updated tests to Excel and PDF

Update notification a vailable by email

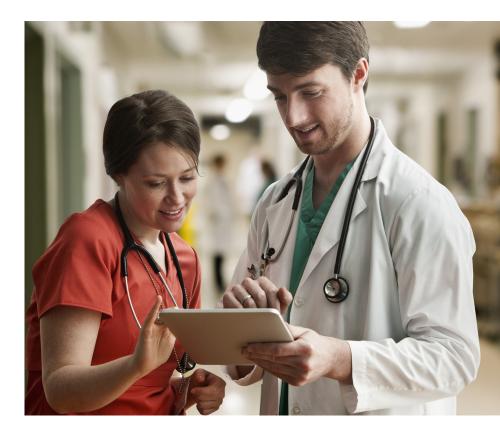
Interface mapping information provided, including LOINC

See pricing messages for price matching due to test code changes

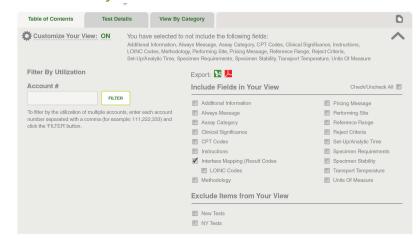
Browse and print new test off erings

For questions or support:

Email intellitestmanag er@quest diagnostics.com, call 1.800 .697.930 2, Option 1, then 6, or ask your DIO Account Executive



Customize your view



Virtual Test Guide

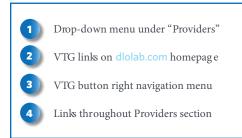
DLO's premium laboratory testing reference tool

An innovative online, no-cost solution to providing testing information with specimen guideline and visual collection guides for Oklahoma's healthcare providers.

DIO's Virtual Test Guide (VTG) on dlolab.com features test information, specimen collection specifics and a visual collection guide for individually select ed frequently used tests and tests with a history of collection and/or submission difficulties. The Virtual Test Guide homepage can be ac cessed se veral different ways.

Easily accessed through your phone, tablet or computer at dlolab.com/vtg.



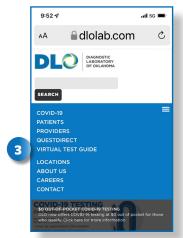




OVID-19 TESTIN

PATIENT BILL PAY

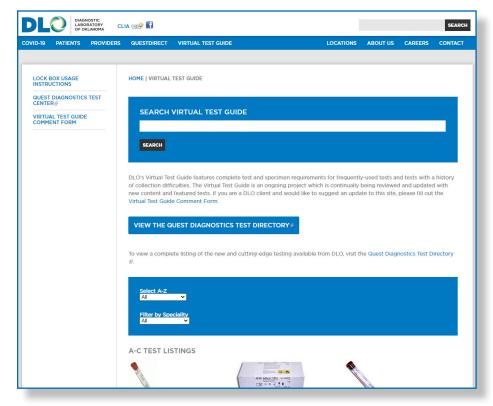
LOCATIONS



Virtual Test Guide

Gain complete test information in one location

Finding the information you need has never been easier.



VTG Search Engine is dedicated to search key words, disease state, tests names, tests numbers and specimen collection devices within the guide.

Tests are listed alphabetically on the VTG homepage, according to test name.

Filters allow tests to be sorted according to specific specialties and/or alphabetic ranges

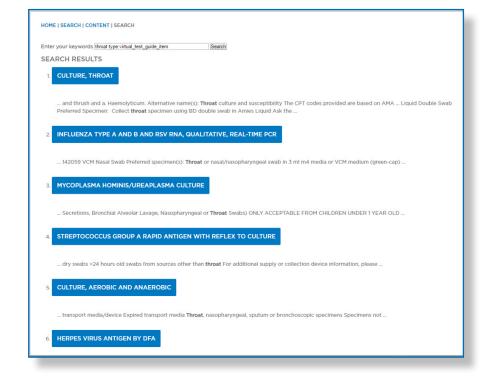
Specimens must be segregated according to temperature while being stored for transport. Lock Box Usage instructions clarify what is needed to protect the integrity of each specimen until pickup by a DIO RSR.

Quest's Test Directory contains information on all of the more than 3,500 tests available through DLO and Quest Diagnostics.

Can find a test? Have a comment you want to share? **Virtual Test Guide Comment Form** sends all submissions to DIO's VTG Team for review and response.

Providing accurate results every time.

VTG search results will list all tests which contain any part of the submitted search criteria. The tests are listed according to the percentage of matching criteria.



Virtual Test Guide

The only place to go for all your test information



All information you need to make the best decision for your patients.

DIO/ Quest t est name

Collection device(s) photo

Test c ode(s)

Billing CPT ode(s)

Additional testing or reflex criteria

Methodology

Limitations

Reference ranges

Clinical significance

Linkto FAQs, algorithms, test reference material or related articles

Alternative test names

Complete listing of the specimen(s) with collection device(s)

Easy to understand instructions and clinical explanations

Collection device with DLO supply order number

Additional test c odes, when a vailable

Preferred specimen

Collection instructions with illustrations, when available

Specimen transport and storage

Transport container

Transport temperature

Specimen stabilit y

Rejection crit eria

Specialties as sociated with test

Click any "Virtual Test Guide" link to return to the VTG homepage.

"HOME" link at the top of any page within dlolab.com will lead to the site's homepage.

Information on related tests or disease states not listed in the VTG can be found at the Quest Diagnostics' Test Directory.

Quality testing starts with proper specimen preparation

Properly collecting and preparing patient blood specimens can minimize errors or inaccurate results and reduce test delays or cancellations. Common examples of inac curate results and err or messages include, but are not limited to:

- Falsely ele vated potas siums
- Falsely decr eased gluc oses
- Falsely ele vated lactat e dehydrogenase le vels
- "Specimen received unspun" comment on reports
- "Quantity not sufficient" (e.g., QNS) comments on reports
- "Red Blood Cells present in specimen" c omment on report

Order of Draw

Inorder for a blood specimen to be appropriate for testing, it must be drawn in a specific order. The following "Order of Draw" procedure must be followed to ensure a suitable blood specimen is obtained and to avoid cross contamination of specimens with additives from a previous tube or c ontainer.

	>					
Blood Cultures	Citrate Tube	Separator	Heparin Tube	EDTA	Sodium Fluoride	Citrate ACD
Varies	Light Blue	Serum Tube Gel - Red/ Gray or Gold	Invert 8-10 times Royal Blu	Lavender, Tan or	Gray	Yellow
Invert 8-10 times	Invert 3-4 times			Royal Blue Invert 8-10 times	Invert 8-10 times	Invert 8-10 times
		No Gel - Red				
		Gently Invert 5 times				

- Allow the SST and red top serum tubes to clot for a minimum of 30 minutes, but no longer than 45 minutes, before centrifugation.
- Tubes should be allo wed to clot in a vertical position (e.g., in a test tube r ack) at room temperature unless otherwise not ed.
- If your centrifuge is a swing bucket centrifuge, spin the SST and serum tubes f or 15 minutes at 2, 200 RP M.

For serum or plasma specimens, draw a sufficient volume of whole blood to obtain the required serum or plasma volume after centrifugation (approximately 2 ½ times mor e whole blood).

For serum, gently invert the tube eight times after filling; allow the blood to clot for at least 30 minutes in a vertical position and separate by centrifugation.

- 10 minutes for horizontal spin centrifuges
- 15 minutes for fixed head c entrifuges

For plasma and whole blood, completely fill the tube to eliminate dilution from the anticoagulant or preservative; immediately mix the blood by gently and thoroughly inverting the tube ten times. Separate plasma by centrifugation. Transfer plasma to a plastic tube and label the tube as "plasma."



Proper Phlebotomy Techniques

This chart shows the various tube tops used during the collection of DLO lab specimens, including the additive, number of inversions and order of draw.

Order of Draw and Number of Inversions are for specimens drawn in plastic tubes only.

COLLECT IN THIS ORDER



8-10



Citrate

Gel, serum

Tube must be filled completely. Note: When using a winged blood collection set for venipuncture and a coagulation (citrate) tube is the first specimen to be drawn, a discard tube should be drawn first. The discard tube must be used to fill the blood collection set tubing's "dead space" with blood, but the discard tube does not need to be completely filled. This important step will ensure maintenance of the proper blood-to additive ratio of the blood specimen.







Do not use gel tubes for toxicology or





No gel, Serum 5





Heparin 8







EDTA 8



EDTA

8





Sodium Fluoride (Glucose)

Tubes with other additives



Citrate SCD

Last tube drawn

Please properly fill and separate all specimens.

The information on this chart is valid as of November 17, 2021 and is subject to change without notice. ©2021 Diagnostic Laboratory of Oklahoma, L.L.C. All rights reserved.

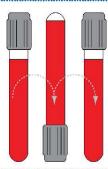
BD Vacutainer® Plus Plastic Citrate Tube Draw Volume Guide

Sufficient volume achieved if blood drawn falls above minimum fill indicator. For blood transfer, do not fill above illustrated dashed maximum line.

Note: The quantity of blood drawn into evacuated tubes varies with altitude, ambient temperature, barometric pressure, tube age, venous pressure and filling technique



* According to CLSI guideline, Dec. 2003, Doc. H1-A5, Vol. 23, No. 33.





Clot 30 minutes

Allow blood to clot for 30-45 minutes in a vertical position



Spin 10 minutes

Centrifuge at full speed (between 1100 & 1300g) for 10 minutes for swing-head units or 15 minutes for fixed angle units (balance tube in centrifuge).

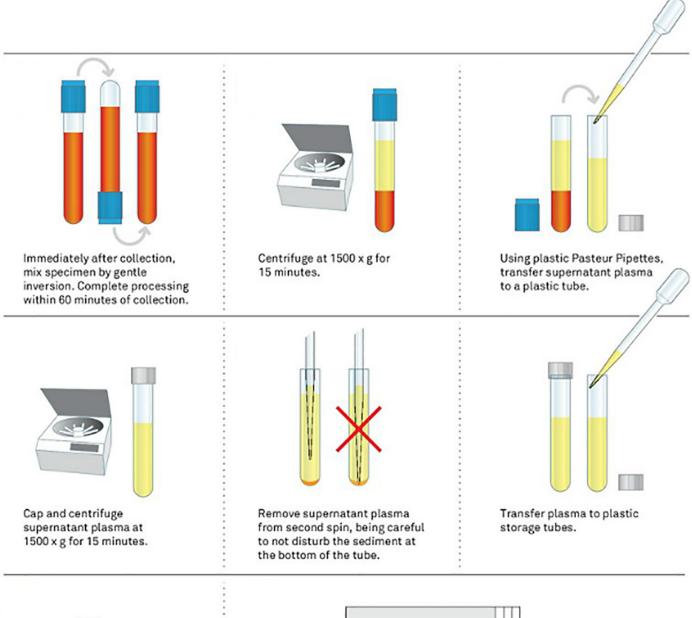


Fill Transport Vial Using a pipet, move the

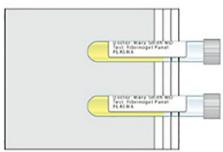
serum to a clear transport vial. Label with Specimen Type and two patient identifiers.

Coagulation Testing

Preparing Platelet - Poor Plasma for Coagulation Testing







Factors That Compromise Specimens

Specimen integrity is imperative to achieve quality test results.

Your care, skill, and knowledge when preparing the patient and specimen are essential to the provision of the highest quality standards for testing and services.

The following guidelines highlight the various factors that can compromise patient's specimens and as says.

Hemolysis

Hemolysis occurs when the erythrocytes are ruptured and release their contents into the serum or plasma. The hemolyzed serum or plasma will look light pink to bright red. Hemolysis, even in small amounts, may alter test results markedly, particularly potas sium and LDH. Grossly or moderately hemolyzed specimens may be rejected.

Causes of hemolysis include:

- Small needle used t o collect specimen
- Difficult phlebotomy
- Placing red top tubes in the r efrigerator without allowing 30 minut es at r oom temperature for complete clotting
- Vigorous shaking of specimens
- Storing specimens in excessive heat or in a refrigerator that is too cold

Quantity Not Sufficient (QNS)

Each as say requires a minimum amount of specimen required to perform the test accurately. If we do not receive enough of a specimen to meet the minimum volume requirements, we will not perform the test. For serum or plasma specimens, please draw more than the amount request ed in our specimen requirements (for example, 2 ½ times more than the request ed volume). If you suspect a specimen will be QNS, list tests in or der of priority.

Lipemia

Excessive lipids in the blood produce a cloudy or milky specimen. Moderately to grossly lipemic specimens may invalidate many test results. Lipemic specimens may be the result of a recent meal prior to the blood collection. Follow the general rules of fasting before a blood specimen is obtained (e.g., the patient should have nothing to eat or drink, except water, for 8-12 hours prior to the draw).

Hyperbilirubinemia

Icteric serum or plasma will appear dark to bright yellow. Icterus may affect some results. To ensure quality we may request another specimen be collected for analysis.

Specimens collected with outdated supplies

Please check routinely to ensure that your supplies are not outdated. All specimens received in expired collection or transport tubes will be r eject ed by DIO.



Specimen Handling and Transport

Quality testing starts with proper specimen preparation.

Urine Collection

Urine collections require providing specific instructions to the patient. Clean catch patient instructions and 24-hour collection instructions are detailed in DIO's Virtual Test Guide and/or Quest Diagnostics Directory of Services. For 24-hour test collections, total urine volume must be included on both the labor atory order and the urine aliquot submitt ed for analysis.

Tissue Collection

Ensure that tissue specimens are covered completely in 10% formalin. For further details about collection and preparation of tissue specimens, r efer to the Virtual Test Guide, Quest Diagnostics digital Dir ectory of Services or Test Directory.

Collection (other)

Comprehensive collection procedures for trace elements, cultures, and toxicology specimens can be found in the Virtual Test Guide, digital Directory of Services or Test Directory

Labeling

Each specimen container must be labeled with appropriate patient identification in order to be test ed. Specimens with missing patient identification will not be test ed. If a significant discrepancy is noted with the patient information provided on the specimen and the laboratory order, your facility will be contacted for clarification. Specimens for HIV testing, blood bank (immunohematology) testing and other sensitive tests that are inconsistently labeled will not be test ed.

Storage

All specimens must be stored at the appropriate temperature prior to transport to the laboratory for testing. Refer to the Virtual Test Guide, Directory of Services or online Test Directory for information on specimen st orage temperatures.

- Storage temperatures are defined as:
- Ambient/Room Temperature (15 30 degr ees C)
- Refrigerated (2 10 degr ees C)
- Frozen (-20 degr ees C or c older)

Minimum Volume Requirements

Test volumes listed in the Virtual Test Guide, Quest Diagnostics Directory of Services or online Test Directory allow for multiple test determinations. The minimum volume allows for a single test including instrument dead volume. Adequate specimen volume for each test request ed should be submitted to DIO to avoid delays in processing and to expedite turnaround time. Prioritizing tests for low volume (short) specimens: Specimens with questionably small sample volumes can have the tests prioritized on the test order form. Minimum testing requirements are available by calling DIO customer services.

Collection Supplies

Specimens collected and/or transported in expired collection or transport devices will be rejected by DIO. Please routinely check to ensure your supplies are not out dated.

Additional Details an Instructions

- Specimen c ollection and handling should always take place using Universal Precautions.
- Specimens should ne ver be frozen in glass tubes.
- Needles or s yringes should ne ver be submitt ed to the laboratory.

Specimen Handling and Transport

Microbiology Collection Devices







































The information on this chart is valid as of November 17, 2021 and is subject to change without notice.

Cytology Specifications

Comprehensive testing from one vial, one specimen

Image-guided Pap with age-based screening protocols— DLO SMART Codes

SMART test codes are comprised of Imaged Pap testing with HPV and additional STI tests, appropriate for her age, based on professional guidelines.*

Send in the specimen using either ThinPrep® or SurePath™, and provide the patient's date of birth

Image-guided Pap with age-based screening protocols						
Test Offerings	ThinPrep	SurePath				
Image-Guided Pap with Age-Based Scr eening Protocols	91384	91384				
Image-Guided Pap with Age-Based Scr eening, Plus CT/NG	91385	91385				
Image-Guided Pap with Age-Based Scr eening, Plus CT/NG/T rich	91386	91386				

^{*}Ask your Quest Diagnostics representative, or visit quest diagnostics. com/smartcodes, for a full explanation of the use of SMART Codes, as well as to see the most current professional cervical cancer and STI screening guidelines.

Additional Testing Options

		1					
Reflex and co-testing options for Pap and HPV (based on ACOG guidelines)							
	•	Test Offerings	ThinPrep w/ Imaging	SurePath w/ Imaging			
	Age 21-29	Pap	58315	18810			
		Pap (reflexes to HPV if ASCUS	909 34	18811			
Cytology e very 3 years for		Pap (refle xes to HPV if ASCUS) and CT/NG ¹	91912	18817			
women 21-29; co-testing	Age 30- 65	Pap	58315	18810			
(Pap and HPV combined) for		Pap & HPV	909 33	18813			
women 30-65		Pap & HPV and CT/NG ¹	91339	18828			
		Pap & HPV mRNA E6/E7, reflex HPV 16,18/45 Pap has to be neg, HPV has to be detected, then reflex to 16, 18/45	91414	18829			
CT/NG, C. trachomatis/N gonorrhoeae RNA							

Out-of-the-vial tests						
Test Offerings	ThinPrep	SurePath				
СТ	11361	11361				
NG	11362	11362				
CT/NG	11363	11363				
Trichomonas vaginalis		905 21				
	0056.0	00560				
	9088 7	92203				
HPV Genotypes 16, 18/45	91826					
HPV Reflex to Genotypes 16, 18/45	909 42	92211				





1. For patients with risk factors for sexually transmitted infections.

Specimen Handling and Transport

Proper Specimen Identification

The College of American Pathologists (CAP), DLO's laboratory accrediting agency, requires that all specimens submitted for testing must have two patient identifiers located on the specimen container upon submission.

Specimen labels

All specimens should be labeled at the time of collection with at least two patient identifiers that must also appear on the requisition.

Examples of patient identifiers are as follows:

The patient's name (full last name, then full fir st name or initial) or a unique ID code is always required.

The sec ond patient identifier may be one of the f ollowing:

- Date of birth (month/date/year)
- Other unique patient identifier that is also on the t est requisition, e.g., hospital or offic e ID code or file number
- DIO requisition number or specimen bar code label
- Other barcode labels can be used if the barcode matches the unique identifiers on the printed requisition (the barcode does not need to be human readable)

NOTE: Location-based identifier s are *NOT* acceptable, e.g., hospital r oom number or street address

Each specimen c ontainer must have a secur ely affixed label with the f ollowing information:

- the patient's name writt en exactly as it appear s on the test requisition (e.g. Doe, Jane)
- a sec ond patient identifier as not ed above
- your account number
- date of collection

Additional Instructions

If the label is hand-written, use a ballpoint pen—do not use a

If glass slides are submitted, use a pencil for labeling the **frosted end** — two identifiers are preferred although patient's name alone is ac ceptable

If labeling a sample that is intended to be frozen, secure the label with transparent tape.

When using an electronically-generated DLO test requisition, place the label lengthwise on the tube.

When submitting a specimen in a container other than the tube used to draw the sample (e.g., transfer vials), indicate specimen **type on the label** (e.g., serum, plasma, urine, et c.).

When submitting specimens for microbiological testing (e.g., cultures, bacterial antigen, microscopic examination) the nature and anatomic source of the sample and the specific organism(s) to be detected, if any, should be specified.

Quanum will automatically print specimen labels with submitted orders. Improper labeling of patient specimens may result in test cancellations.

Specimen Transport Preparation

DIAGNOSTIC LABORATORY OF OKLAHOMA

Information Regarding Requisiton Folding

DLO is dedicated to delivering accurate results on-time, every time. Our goal is to be accountable for your patient's specimen from the time we pick it up until the time it is tested and resulted.

Specimen tracking is the tool that allows us to capture individual patient demographics for each specimen you entrust in our care.

By folding the requisiton in a manner that allows us to scan the barcode, we are able to capture each patient's information. This allows us to track the status of our patient's specimen at any given time until the results are in your hands.



Print Requisition



Folded requisition option



Requisition inserted in sleeve outside the bag with barcode on clear side.



dlolab.com/pickup

Introducing online pickup, a new service from DLO

We know you're busy. That's why we now offer convenient online scheduling for specimen pickups. It's part of our commitment to creating new and better ways to support you in the important work of caring for patients.

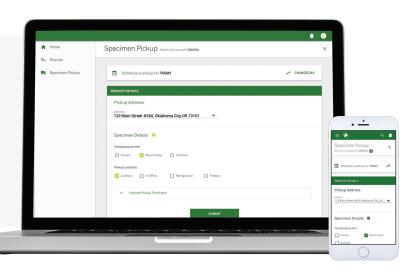
Just click, enter, and click. DONE.

Scheduling a Diagnostic Laboratory of Oklahoma (DLO) specimen pickup may now be done online through our Quanum™ for Healthcare Professionals website. Simply go online to dlolab.com/pickup to log in, enter specimen details, schedule the pickup, and submit. That's all it takes. You'll receive confirmation, and a DLO courier will pick up your specimen(s).

dlolab.com/pickup—quick and convenient

No more phone calls or waiting to speak with the right person. Online pickup is easy to use and available anytime to fit into your workflow and help you streamline operations.

- Simple 4-click ordering
- **Schedule**, modify, or cancel a request electronically
- **Receive confirmation** that the pickup was successfully scheduled
- Available 24/7—no need to wait for a dispatcher during peak call times
- Available from your PC, tablet, or smartphone
- **1-click option** for frequently requested pickups





Ready to try online pickup? It's easy.

dlolab.com/pickup

Signing up for online pickup is a simple, 1-time process:

Go to dlolab.com/pickup, click the "Log In" button and log in.

Login

Ummare

Passered

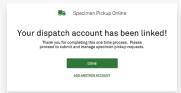
On the next screen select **SPECIMEN PICKUP**.



Follow the instructions to enter your dispatch account information and to confirm pickup address.†



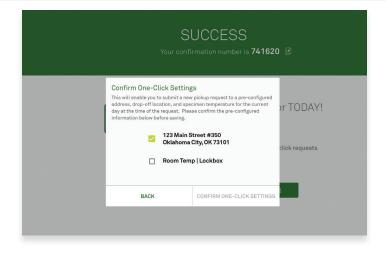
Click **SUBMIT** to confirm you're ready to schedule your first pickup!



Have a frequent pickup need? Try 1-click.

Online pickup gives you the option to save your pickup order information, so you can save even more time. Just choose the **1-click option**, enter the information, and you're always just 1 click away from more efficient scheduling.

At DLO, our goal is to help you simplify practice operations so you have more time to focus on enhancing the patient experiences that can lead to healthier outcomes.



If you need assistance, have DLO lock box questions, or want to contact DLO Logistics directly to schedule a specimen pickup, call **1.800.891.2917**, option **3.**



For questions about Quanum for Healthcare Professionals or Online Specimen Pickup, contact the DLO client services at 1.800.891.2917.



800.891.2917 • dlolab.com

^{*} You can log in to the site using your existing Quanum/Care360 credentials or follow the steps on the screen to verify your Quest Diagnostics account. If you're having trouble logging in, call 1.800.891.2917 for support. † Please note: DLO cannot pick up from P.O. box addresses.

Lock Box Usage Instructions

Help to ensure proper specimen transport!

Ambient, refrigerated, and frozen specimens MUST be segregated.

If you need additional lock box capacity to allow for complete segregation. **Call 1.800.891.2917**, **Option 3**, to notify DLO if you need additional lock box capacity to properly srore specimens.

Ambient

Place ambient specimens farthest away from any cold packs.

If there are specimen tubes requiring different temperature states within a single patient sample collection, place each specimen tube in a separate specimen bag with a copy of the ordering requisition. This will facilitate those tubes being placed in the correct areas of the lock box.

DO NOT put ambient specimens on top of refrigerated specimens.

Refrigerated

When placing a refrigerated specimen inside the lock box, use a cold pack and place refrigerated specimen directly on the cold pack either by:

- 1) rubber banding specim to the cold pack
- placing specimen in separate bag with the cold pack

DO NOT put refrigerated specimens in the frozen tote bag.

Frozen

Frozen specimens should be placed inside a frozen tote container to allow them to remain frozen while in the lock box. Make sure the specimen is completely frozen before placing it inside the frozen container.

Remember to take the frozen tote out of the lock box in the morning to refreeze the gelpacks before reusing. Freeze only the gel packs, not the entire tote.





