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Preface

Safety Precautions

The Catalyst Dx* Chemistry Analyzer weighs approximately 50 pounds (22 kg). It may require multiple people to lift the instrument.

The analyzer does not contain any user-serviceable components. DO NOT disassemble.

DO NOT stack other equipment or containers on top of the analyzer.

Keep analyzer away from sources of heat or flames.

DO NOT place or operate the analyzer near x-ray equipment, photocopiers, or other devices that generate static or magnetic fields.

PROTECT your equipment from damp conditions or wet weather.

Take care not to spill water or other fluids on the unit.

DO NOT use any of the following liquids, abrasives, or aerosol sprays on, inside, or near the analyzer, as they may damage the analyzer and may influence results:

- Organic solvents
- Ammonia-based cleaners
- Ink markers
- Sprays containing volatile liquids
- Insecticides
- Disinfectant
- Polish
- Room freshener
- Canned air

Line voltage for the Catalyst Dx analyzer is 100–240 V AC, 50–60 Hz. Be sure to plug all equipment into properly grounded electrical outlets.

Use only the power cable supplied.

Disconnect the power cable:

- If the cable becomes frayed or otherwise damaged.
- If anything is spilled onto the equipment.
- If your equipment is exposed to excessive moisture.
- If your equipment is dropped or the case has been damaged.
- If you suspect that your analyzer needs service or repair.
- Whenever you clean the case.

If the equipment is used in a manner other than specified, the protection provided by the equipment may be impaired.
International Symbol Descriptions

International symbols are often used on packaging to provide a pictorial representation of particular information related to the product (such as expiration date, temperature limitations, batch code, etc.). IDEXX Laboratories has adopted the use of international symbols on our analyzers, product boxes, labels, inserts, and manuals in an effort to provide our users with easy-to-read information.

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
</tr>
</thead>
</table>
| Use by | A utiliser avant  
Verwendbar bis  
Usare entro  
Usar antes de  
使用期限 |
| Batch code (Lot) | Code de lot (Lot)  
Chargenbezeichnung (Partie)  
Codice del lotto (partita)  
Código de lote (Lote)  
ロット番号 |
| Serial number | Numéro de série  
Seriennummer  
Numero di serie  
Número de serie  
シリアル番号 |
| Catalog number | Numéro catalogue  
Bestellnummer  
Numero di catalogo  
Número de catálogo  
製品番号 |
| Authorized Representative in the European Community | Représentant agréé pour la C.E.E.  
Autorisierte EG-Vertretung  
Rappresentante autorizzato nella Comunità Europea  
Representante autorizado en la Comunidad Europea  
EC内の正規販売代理店 |
| Manufacturer | Fabricant  
Hersteller  
Ditta produttrice  
Fabricante  
製造元 |
| Caution, consult accompanying documents | Attention, consulter les documents joints  
Achtung, Begleitdokumente beachten  
Attenzione, consultare la documentazione allegata  
Precaución, consultar la documentación adjunta  
注意、添付文書をご参照ください。 |
| Temperature limitation | Température limite  
Zulässiger Temperaturbereich  
Temperatura limite  
Limitación de temperatura  
保存温度（下限） |
| Upper limit of temperature | Limite supérieure de température  
Temperaturobergrenze  
Limite superiore di temperatura  
Limite superior de temperatura  
保存温度（上限） |
| Consult instructions for use | Consulter la notice d'utilisation  
Gebrauchsanweisung beachten  
Consultare le istruzioni per l'uso  
Consultar las instrucciones de uso  
取扱説明書をご参照ください。 |
| Keep away from sunlight | Conserver à l’abri de la lumière  
Vor direkter Sonneneinstrahlung schützen  
Mantener alejado de la luz solar  
Tenere lontano dalla luce diretta del sole  
遮光してください。 |
WEEE-Richtlinie 2002/96/EG  
Directiva 2002/96/CE RAEE  
Direttiva RAEE 2002/96/CE  
廃電気電子機器指令（WEEE Directive 2002/96/EC） |
| Biological risks | Risques biologiques  
Biogefährlich  
Rischi biologici  
Riesgos biológicos  
生物学的リスク |
| Do not reuse | Usage unique  
Nicht wiederverwenden  
No reutilizar  
Non riutilizzare  再利用しないでください。 |
<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="image" alt="Caution, hot surface" /></td>
<td>Caution, hot surface  &lt;br&gt; Attention, surface très chaude  &lt;br&gt; Precaución, superficie caliente  &lt;br&gt; Vorsicht, heiße Oberfläche  &lt;br&gt; Attenzione, superficie rovente  &lt;br&gt; 高温注意</td>
</tr>
<tr>
<td><img src="image" alt="Keep dry" /></td>
<td>Keep dry  &lt;br&gt; Conserver dans un endroit sec  &lt;br&gt; Mantener seco  &lt;br&gt; Vor Nässe schützen  &lt;br&gt; Tenere al riparo dall'umidità  &lt;br&gt; 湿らせないこと</td>
</tr>
<tr>
<td><img src="image" alt="This side up" /></td>
<td>This side up  &lt;br&gt; Haut  &lt;br&gt; Este lado hacia arriba  &lt;br&gt; Diese Seite nach oben  &lt;br&gt; Alto  &lt;br&gt; この面を上にする</td>
</tr>
<tr>
<td><img src="image" alt="Do not freeze" /></td>
<td>Do not freeze</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="image" alt="Electrostatic-sensitive device" /></td>
<td>Electrostatic-sensitive device  &lt;br&gt; Appareil sensible aux charges électrostatiques  &lt;br&gt; Dispositivo sensible a descargas electrostáticas  &lt;br&gt; Gerät ist sensibel auf elektrostatische Ladung  &lt;br&gt; Dispositivo sensibile alle scariche elettrostatiche  &lt;br&gt; 静電気の影響を受ける装置</td>
</tr>
<tr>
<td><img src="image" alt="Fragile" /></td>
<td>Fragile  &lt;br&gt; Frágil  &lt;br&gt; Zerbrechlich  &lt;br&gt; Fragile  &lt;br&gt; 取扱注意</td>
</tr>
<tr>
<td><img src="image" alt="Date of manufacture" /></td>
<td>Date of manufacture  &lt;br&gt; Date de production  &lt;br&gt; Fecha de producción  &lt;br&gt; Herstelldatum  &lt;br&gt; Data di produzione  &lt;br&gt; 製造年月日:</td>
</tr>
</tbody>
</table>

**Other Symbols**

<table>
<thead>
<tr>
<th>Symbol</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="image" alt="USB symbol" /></td>
<td>USB symbol</td>
</tr>
<tr>
<td><img src="image" alt="Ethernet/network symbol" /></td>
<td>Ethernet/network symbol</td>
</tr>
</tbody>
</table>
Getting Started

Introduction

Welcome to the Catalyst Dx* Chemistry Analyzer.

The Catalyst Dx analyzer’s flexible test menu allows you to monitor the health status of specific organs, recheck values over time, customize profiles by adding single tests to CLIPs, and test blood and urine at the same time to uncover early renal disease (for a complete list of the individual slides and CLIPs available, see pages 14–15). You can even run up to 25 tests on a single sample.

The analyzer’s touch-screen interface provides easy-to-follow instructions to help you navigate the system, enter patient data, specify testing information, and more.

The Catalyst Dx analyzer is for veterinary use only.

IDEXX VetLab* Station Connectivity

The Catalyst Dx analyzer is part of the IDEXX VetLab* suite of analyzers, all of which connect to the IDEXX VetLab Station (IDEXX’s laboratory information management system). Connecting multiple analyzers to the IDEXX VetLab Station helps you attain a comprehensive picture of your patient’s health, with the ability to view test results from multiple analyzers on a single report, determine disease progression with parameter-trending capabilities, and more.

By connecting the Catalyst Dx analyzer to the IDEXX VetLab Station, you can:

- Automatically review patients’ prior results on every printout for easy comparison.
- Improve client communications with illustrated diagnostic or treatment progress printouts.
- Link to expert descriptions and common causes of abnormal values.
- Print information to help explain the significance of results to your clients.
- Allow new staff to train independently.
- Learn proper protocols and tips for best techniques.
IDEXX Dry-Slide Technology

The Catalyst Dx analyzer uses dry-slide technology—the most accurate technology available for in-house testing. Dry-slide technology uses layers to minimize impurities for the most accurate results from even compromised samples.

![Diagram of the Catalyst Dx analyzer layers]

**How it Works**

There are several important steps that the analyzer performs in order to present the results of a sample. Once the slides and sample have been inserted into the analyzer, the Catalyst Dx analyzer incubates the slides and sample. Then, if using a Catalyst* whole blood separator, the plasma is separated from a whole blood sample. The sample is then accurately dispensed onto the slides, the analyzer measures the color development of the slide, and then all used materials are removed from the analyzer.

**Catalyst Dx Components**

**Note:** Some of the components on the Catalyst Dx analyzer have been redesigned since the analyzer was originally launched. The images in this section show the new hardware designs (for example, a single tip/diluent drawer).
sample drawer assembly
raise to gain access to the slide carousel and optics windows

center latch
press to release/lift the sample drawer assembly and access the incubator ring, carousel, optics windows, etc.

centrifuge
spins the whole blood separator to separate the whole blood

whole blood separator carriers
transport whole blood separators from the sample drawer to the centrifuge

optics windows
allow LED light to pass through in one direction and either reflectance or fluorescence to return to be analyzed

incubator ring
incubates the chemical reaction of the slide and the sample

underside of carousel
white reference tile calibrates the chemistry optics module

fan filter
should be cleaned quarterly (monthly if operated in dusty/dirty environment)

power switch
use to turn on/off the analyzer

power receptacle
power cable plugs in here

Ethernet port
use to connect the analyzer to the IDEXX VetLab Station
Installing the Catalyst Dx Analyzer

The Catalyst Dx analyzer works in conjunction with the IDEXX VetLab Station.

To Install the Catalyst Dx Analyzer

1. Before you unpack the analyzer, choose an optimum location for the instrument. The analyzer should be placed on a level surface with a minimum of 2 inches (5 cm) between the back of the analyzer and any wall. Choose a well-ventilated area away from obvious sources of heat, direct sunlight, cold, humidity, or vibrations. For optimum results, room temperature should be at 15°C–30°C (59°F–86°F) and relative humidity at 15%–75%.

   IMPORTANT: Ensure proper ventilation. The analyzer’s cooling vents are in the base. Leave at least a 2-inch (5-cm) clearance around the machine so that air can circulate on all sides.

2. Unpack the analyzer.

   IMPORTANT: The analyzer weighs approximately 50 pounds (22 kg). It may require multiple people to lift the instrument.

3. Remove the packaging foam located inside of the open maintenance access doors.

4. Verify the two black whole blood separator carriers on the top of the sample drawer assembly are seated properly (flat) and in the left and right positions.

5. Verify the white centrifuge sleeve is in place to the right of the sample drawers.

6. Close the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 36–37).

7. Fill the tip drawer with pipette tips.

8. Ensure the Catalyst Dx analyzer is switched off and then connect the power cable to the analyzer and to a properly grounded electrical outlet.

   IMPORTANT: Do not power on the Catalyst Dx analyzer. After connecting the power cable, you must then connect to the router and to the IDEXX VetLab Station (instructions follow).

To Install the IDEXX VetLab Station Connectivity Router

Note: If you already have a network router connected directly to the IDEXX VetLab Station computer, you can skip this section and move to the “To Connect the Catalyst Dx Analyzer to the IDEXX VetLab Station” section (on the next page).

1. Connect the AC power adapter to the power port on the back of the network router supplied by IDEXX Laboratories.

2. Plug the AC power adapter into an electrical outlet.

3. Connect one end of the Ethernet cable (provided with the router) into any available port on the router.

   IMPORTANT: Do not connect the IDEXX VetLab Station directly to the Internet port on the router.

4. Connect the other end of the Ethernet cable (from step 3) into the IDEXX VetLab Station computer’s Ethernet port, which is located near the center panel on the back of the computer.

   Note: If this Ethernet port on the IDEXX VetLab Station computer is currently being used to connect the IDEXX VetStat* Electrolyte and Blood Gas Analyzer (red Ethernet cable) to your IDEXX VetLab Station, disconnect the red Ethernet cable. Then complete step 4 above. The “To Reconnect the VetStat Analyzer” section, found on the next page, explains how to reconnect the VetStat analyzer after the router is installed.
To Connect the Catalyst Dx Analyzer to the IDEXX VetLab Station
1. Connect the Ethernet cable provided with the Catalyst Dx analyzer to the next available port on the back of the router.
   **IMPORTANT:** Do not connect the Catalyst Dx analyzer directly to the Internet port on the router.
2. Connect the other end of the Ethernet cable (from step 1) into the Ethernet port on the back of the Catalyst Dx analyzer.
3. Power on the IDEXX VetLab Station. Ensure all analyzer icons (except Catalyst Dx) are present with a “Ready” status. Then, power on the Catalyst Dx analyzer. Once the Catalyst Dx Home screen displays and its icon displays on the IDEXX VetLab Station Home screen, your connections are complete.
   **Note:** If the Catalyst Dx icon does not appear on the IDEXX VetLab Station Home screen within 3 minutes, contact IDEXX Technical Support for assistance.

To Reconnect the VetStat Analyzer
**IMPORTANT:** The direct connection between the VetStat analyzer and the IDEXX VetLab Station uses a standard Ethernet cable with a light-purple cross-over adapter. This light-purple cross-over adapter, which is located on one end of the cable, must be removed before reconnecting the Ethernet cable to the router.
1. Ensure the light-purple cross-over adapter has been removed from the VetStat Ethernet cable.
2. Connect one end of the VetStat Ethernet cable to the VetStat analyzer and the other end to the next available port on the router.
   **IMPORTANT:** Do not connect the VetStat analyzer directly to the Internet port on the router.
3. On the VetStat Home screen, tap **System Manager** and then tap **Setup**.
4. Use the alphanumeric keypad to enter the setup password, 207, and tap **OK**.
5. From the Setup screen, tap **Hardware** and then tap the **Ethernet** tab.
6. Select **Dynamic** as the network setting and then tap **OK** to save the change.
7. Turn off the VetStat analyzer by toggling the power switch to OFF. Then, turn on the analyzer. After the VetStat analyzer returns to the Home screen, the VetStat icon will appear at the bottom of the IDEXX VetLab Station Home screen.
   **Note:** If the VetStat icon does not appear on the IDEXX VetLab Station Home screen within 3 minutes, contact IDEXX Technical Support for assistance.

Powering On the Analyzer
To turn on the analyzer, press the power switch on the back of the analyzer. The analyzer may take 15–25 minutes to warm up. While warming up and performing a system check, the Catalyst Dx analyzer screen will display “IDEXX Laboratories,” the maintenance access doors will open, and sample drawers will slide out and back in. The analyzer is ready for use once the “Initializing” status message disappears on the Catalyst Dx Home screen.

**Note:** Ensure the IDEXX VetLab Station is powered on prior to powering on the analyzer. If the IDEXX VetLab Station is restarted while the analyzer is on, you may need to reboot your analyzer.
Shutting Down the Analyzer

To Shut Down the Catalyst Dx Analyzer
1. Tap **Tools**.
2. Tap **Shut Down** and then tap **Yes** to confirm that you want to shut down the analyzer.
3. When the analyzer indicates it is okay to do so, press the power switch on the back of the analyzer to power the analyzer off.

Printing Test Results

The Catalyst Dx analyzer is connected to the IDEXX VetLab Station. Therefore, you print your Catalyst Dx test results using the print settings on the IDEXX VetLab Station (compatible printer required). For more information on printing from the IDEXX VetLab Station, see the *IDEXX VetLab Station Operator’s Guide*.

Catalyst Dx Analyzer Consumables

The following consumables are available for use with the Catalyst Dx analyzer:

**CLIPs, Panels, and Slides**

You can run any IDEXX slide on any species; however, reference intervals may not always be provided (see footnotes for more information).

<table>
<thead>
<tr>
<th>Chemistry</th>
<th>Abbreviation</th>
<th>Chem 17 CLIP</th>
<th>Chem 15 CLIP</th>
<th>Chem 10 CLIP</th>
<th>Equine 15 CLIP</th>
<th>NSAID 6 CLIP</th>
<th>UPC Panel</th>
<th>Lyte 4 CLIP</th>
<th>QC CLIP</th>
<th>Individual Slides</th>
</tr>
</thead>
<tbody>
<tr>
<td>Albumin</td>
<td>ALB</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
<td></td>
<td></td>
<td></td>
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<td></td>
<td>✓</td>
</tr>
<tr>
<td>Alkaline Phosphatase</td>
<td>ALKP</td>
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<td>✓</td>
<td>✓</td>
<td>✓</td>
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<tr>
<td>Alanine Aminotransferase</td>
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<td>Amylase</td>
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<tr>
<td>Calcium</td>
<td>Ca</td>
<td>✓</td>
<td>✓</td>
<td>✓</td>
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<td></td>
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<tr>
<td>Cholesterol</td>
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<td>Chloride</td>
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<td>C-Reactive Protein²</td>
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<tr>
<td>Fructosamine⁴</td>
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<td>✓</td>
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<tr>
<td>Gamma-glutamyltransferase</td>
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<tr>
<td>Glucose</td>
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<td></td>
<td></td>
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</tr>
<tr>
<td>Potassium</td>
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<tr>
<td>Lactate</td>
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<tr>
<td>Lactate Dehydrogenase</td>
<td>LDH</td>
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<td></td>
<td>✓</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
### Other Consumables

- Catalyst* Sample Cups
- Lithium Heparin Whole Blood Separator
- Catalyst* Pipette Tips
- 300 µL Pipette
- 300 µL Pipette Tips
- Catalyst* PHBR Control
- Catalyst* Advanced Control
- VetTrol* Control Fluid
- UPRO Control Fluid
- Urine P:C Diluent
- Alcohol Prep Pads
- Optical Tissues

### Compatible Species

Species with specific reference intervals:

- Canine†
- Bovine
- Feline†
- Llama
- Equine†
- Sea Turtle

†Species-specific intervals are available for these species. All other species are qualified as “other.”
Groups of species with guideline reference intervals:

**Note:** Guideline reference intervals will vary because there is diversity within the species of these groups.

<table>
<thead>
<tr>
<th>Avian</th>
<th>Monkey</th>
<th>Rat</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ferret</td>
<td>Mouse</td>
<td>Sheep</td>
</tr>
<tr>
<td>Goat</td>
<td>Pig</td>
<td>Snake</td>
</tr>
<tr>
<td>Lizard</td>
<td>Rabbit</td>
<td>Tortoise</td>
</tr>
</tbody>
</table>
Using the Catalyst Dx* Analyzer

Overview
The Catalyst Dx analyzer is controlled via a touch-screen monitor on the front of the analyzer and by the IDEXX VetLab* Station.

Using the Touch Screen
To get the best results when using the touch screen:
• Do not rest your hand on the touch screen. The screen is sensitive to touch. Pressure from your hand prevents the touch screen from functioning properly.
• Tap the screen firmly.
• Never tap the touch screen with a sharp or abrasive object.

Analyzing Samples
The Catalyst Dx analyzer allows you to run up to 25 tests on a single sample. You can even load multiple patient samples at the same time.

Before you begin, please take note of the following:
• Frozen CLIPs/panels/slides can be run on the Catalyst Dx analyzer (no thawing required).
• Most CLIPs/slides should be loaded within 5 minutes of opening their foil packaging. The Lyte 4 CLIP should be loaded within 2 minutes of opening its foil packaging.
• If you are running a Lyte 4 CLIP or NH₃ slide, be sure to load it in the sample drawer before any other CLIPs or slides. If running both, NH₃ slides should always be loaded first.
• If you are running a UPC panel or a PHBR slide, do not load any other CLIPs or slides in the sample drawer.
• Only one test that requires a reagent pack can be processed in a single run. (For example, a total T4 test cannot be processed with a CRP test.)
• If you run a special slide without selecting the applicable special slides check box and/or you do not follow the on-screen instructions, your results will be flagged and you may receive inaccurate results.

To Run a Sample
1. Enter the patient information on the IDEXX VetLab Station (for more information, see the “Analyzing Samples” chapter of the IDEXX VetLab* Station Operator’s Guide).
2. Once the patient’s name appears in the Pending list on the Catalyst Dx Home screen, tap the patient’s name and then tap Select.
3. Select the Sample Type (whole blood, plasma, serum, urine, or other).
   Note: To learn which sample types can be run for a particular slide or CLIP, see the chart on page 28.
4. If you are running a special slide, select the applicable special slides check box.
5. Tap **Next**.

6. If you are running a UPC panel or PHBR slide, follow the on-screen instructions and then tap Next.

7. Load the sample in the sample drawer in either a whole blood separator (whole blood samples only) or a sample cup (plasma, serum, or urine samples only).

8. Open the foil packaging containing the CLIP(s)/slide(s) you are running.

9. Load the slides in the sample drawer.
   - If you are loading a Catalyst CLIP, snap open the CLIP handle and then use the handle to load the CLIP onto the sample drawer. Once the slides are secure in the sample drawer, pull the CLIP to detach the slides from the handle.
   - Note: Urine should be centrifuged prior to loading.

10. If you are running a TT4, CRP, or PHBR slide, load the reagent in the tip/diluent drawer(s).

11. Tap **Run**. The Catalyst Dx analyzer begins to process the patient sample automatically and transfers the results to the IDEXX VetLab Station once the run is complete.

12. If you loaded a TT4, CRP, UPC panel or a PHBR slide, remove and dispose of the sample/wash cups from the diluent drawer when prompted.

---

**Diluting Samples**

Dilutions should only be performed when a test value is outside the reportable range or when the sample contains interfering substances (e.g., medications) that cause a nonlinear or invalid result. The Catalyst Dx analyzer supports automated dilutions (the analyzer mixes the sample and diluent for you) and manual dilutions (you prepare the dilution outside of the analyzer). Select the appropriate option on the Identify Sample screen.

Remember the following important notes when diluting samples for analysis on the Catalyst Dx analyzer:

- Only dilute tests with results outside of the reportable range. Diluting tests with results in the normal range may produce invalid results.

- All chemistries should be analyzed first on the undiluted sample. Some analytes, such as GGT and total bilirubin, have low serum/plasma concentrations. These analytes may be diluted out even with the lowest dilution. Dilute the remaining sample and analyze any chemistries that were outside of the reportable range on the first analysis.

- Perform a dilution only when a test value is accompanied by a greater-than symbol (>) or when the analyzer informs you a dilution is necessary to receive accurate results.

- Use the proper diluent material for your sample type.
  - For whole blood, plasma, and serum samples, use normal saline.
  - IDEXX does not recommend manually diluting whole blood in a Catalyst whole blood separator—only dilute the separated plasma.
  - For urine, use Catalyst Urine P:C Diluent.

- Use an accurate measuring device, such as a calibrated pipette or syringe.
For best results, start with a 1:2 dilution (1 part sample to 1 part diluent)—do not exceed 10 parts diluent.

Do not dilute samples that are undergoing ammonia, phenobarbital, fructosamine, total T4, or electrolyte testing.

Do not dilute small samples to achieve a minimum sample volume. Such dilutions on normal analyte concentration cannot be read accurately. When dilution is needed to determine some analytes at very high concentration, the sample should be diluted manually.

You cannot perform two automated dilution runs at the same time, but you can perform an automated dilution run with a manual dilution run.

An automated dilution run will be canceled if:
- The diluent and tip drawer(s) is/are opened during the run.
- There is insufficient diluent/sample volume.
- There is an insufficient number of tips in the tip drawer.
- There are too many slides in the run.

Preparing Manual Dilutions

To Prepare a 1:2 Dilution
1. Accurately measure the desired amount of sample to be diluted and gently transfer it to a sample cup.
2. Accurately measure an equal amount of diluent and transfer it to the sample collected in step 1.
3. Thoroughly mix the sample and diluent.
4. Analyze the sample using the “To Run a Diluted Sample” instructions below.

To Prepare Dilutions Greater Than 1:2
If additional dilutions beyond 1:2 are necessary, always begin with the original, undiluted sample. Then, incrementally increase the parts diluent as indicated in the dilution chart (below).

Volumes are for example only. Parts Sample + Parts Diluent = Total Parts (Dilution Factor)

<table>
<thead>
<tr>
<th>Parts Sample</th>
<th>Parts Diluent</th>
<th>Total Parts (Dilution Factor)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 (100 μL)</td>
<td>0</td>
<td>1 (undiluted sample)</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>1 (100 μL)</td>
<td>2</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>2 (200 μL)</td>
<td>3</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>3 (300 μL)</td>
<td>4</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>4 (400 μL)</td>
<td>5</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>5 (500 μL)</td>
<td>6</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>6 (600 μL)</td>
<td>7</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>7 (700 μL)</td>
<td>8</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>8 (800 μL)</td>
<td>9</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>9 (900 μL)</td>
<td>10</td>
</tr>
<tr>
<td>1 (100 μL)</td>
<td>10 (1000 μL)</td>
<td>11</td>
</tr>
</tbody>
</table>
To Run a Diluted Sample

1. Enter the patient information on the IDEXX VetLab Station (for more information, see the “Analyzing Samples” chapter of the IDEXX VetLab* Station Operator’s Guide).

2. Once the patient’s name appears in the Pending list on the Catalyst Dx Home screen, tap the patient’s name and then tap Select.

3. Select the Sample Type (plasma, serum, urine, or other).

4. Select a dilution option (Automated or Manual). Then, use the up/down arrows to specify the desired dilution factor (total parts).
   Note: You cannot perform an automated dilution on CRP, NH3, PHBR, TT4, or FRU tests or whole blood samples.

5. Tap Next.

6. If you chose to have the analyzer dilute the sample for you (automated dilution), follow these steps:
   a. Open the tip and diluent drawer(s). Do not open the drawer(s) if there is an automated dilution run in process.
   b. Fill the tip drawer completely.
   c. Load an empty sample cup in the left circular cup holder.
   d. Load a sample cup containing 300 µL of diluent in the right circular cup holder (the sample cup should fit inside the holder easily).
   e. Close the tip and diluent drawer(s).
   f. Tap Next.

7. Load the sample in the sample drawer in either a whole blood separator (whole blood samples only) or a sample cup (plasma, serum, or urine samples only). The minimum sample volume varies based on the dilution factor and the number of slides that are being diluted (see table below).

<table>
<thead>
<tr>
<th>Parts Sample + Parts Diluent = Diluent Ratio</th>
<th>Maximum Number of Slides per Dilution</th>
<th>Minimum Sample Volume Serum, Plasma, or Urine</th>
<th>Whole Blood</th>
<th>Diluent Volume</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 + 1 = 1:2</td>
<td>5</td>
<td>155 µL</td>
<td>700 µL</td>
<td>300 µL</td>
</tr>
<tr>
<td>1 + 3 = 1:4</td>
<td>10</td>
<td>130 µL</td>
<td>700 µL</td>
<td>300 µL</td>
</tr>
<tr>
<td>1 + 9 = 1:10</td>
<td>10</td>
<td>100 µL</td>
<td>700 µL</td>
<td>300 µL</td>
</tr>
<tr>
<td>1 + 20 = 1:21</td>
<td>10</td>
<td>110 µL</td>
<td>700 µL</td>
<td>300 µL</td>
</tr>
</tbody>
</table>

8. Open the foil packaging containing the CLIP(s)/slide(s) you are running.

9. Load the slides in the sample drawer.

10. Tap Run. The Catalyst Dx analyzer begins to process the patient sample automatically.

11. Remove (and dispose of) the sample cups from the diluent drawer when prompted.
Viewing Test Results

Once a test is completed, you can view the test results on the Catalyst Dx analyzer or on the IDEXX VetLab Station.

To View the Test Results on the Catalyst Dx Analyzer

1. On the Catalyst Dx Home screen, tap the Results list.
2. Tap the patient whose test results you want to view.
   **Note:** If you do not see the patient’s name in the Results list, tap the page up ▲ and page down ▼ arrows to view additional patient names.
3. Tap View Results to display the Test Results screen.

To View the Test Results on the IDEXX VetLab Station

See the *IDEXX VetLab Station Operator’s Guide* for detailed instructions on viewing test results.

Canceling a Run That Is In Process

To cancel a run that is in process, tap the applicable patient in the In Process list (on the Home screen) and then tap Cancel Run. Then, tap Yes to confirm the cancellation. The analyzer cancels the run and ejects the slides into the waste drawer.

You can also cancel a run using the Edit In Process List feature in the Tools screen. For more information, see “To Delete a Patient from the In Process List” on page 23.

Removing a Sample from the Analyzer

You can remove a sample from the sample drawer when loading a new sample, by using the Sample Available notification in the In Process list (on the Home screen) or by using the Remove Sample option in the Tools screen.

To Remove a Sample Using the Home Screen

1. Tap the patient in the In Process list (on the Home screen) when the Sample Available notification displays
2. Tap Remove Sample. The sample drawer opens.
3. Remove the sample cup or whole blood separator from the sample drawer.
4. Tap OK to confirm the sample has been removed. The sample drawer closes.

To Remove a Sample Using the Tools Screen

There are two Remove Sample buttons in the Tools screen (one for the left sample drawer and one for the right sample drawer). When a sample cup or whole blood separator is detected in a sample drawer, the patient name associated with that sample is listed on the button (for example, “Remove Sample Fluffy”). When a sample cup or whole blood separator is not detected, the Remove Sample buttons are unavailable.

1. Tap Tools.
2. Tap Remove Sample <Patient Name>. The sample drawer opens and a confirmation message displays on the screen.
3. Remove the sample cup or whole blood separator from the sample drawer.
4. Tap OK to confirm the sample has been removed. The sample drawer closes.
Outside of Reportable Range Samples

Occasionally a test value may be outside the analyzer’s reportable range capability. The test value may be greater than (">" ) the reportable range, or interfering substances in the sample may be causing a nonlinear or invalid result. See the following chart for reportable ranges on individual chemistries. If a value is required, it will be necessary to dilute the sample and repeat the test.

<table>
<thead>
<tr>
<th>Chemistry</th>
<th>U.S. Units</th>
<th>S.I. Units</th>
<th>French Units</th>
</tr>
</thead>
<tbody>
<tr>
<td>ALB</td>
<td>0.1–6.0 g/dL</td>
<td>1–60 g/L</td>
<td>1–60 g/L</td>
</tr>
<tr>
<td>ALKP</td>
<td>10–2000 U/L</td>
<td>10–2000 U/L</td>
<td>10–2000 U/L</td>
</tr>
<tr>
<td>ALT</td>
<td>10–1000 U/L</td>
<td>10–1000 U/L</td>
<td>10–1000 U/L</td>
</tr>
<tr>
<td>AMYL</td>
<td>5–2500 U/L</td>
<td>5–2500 U/L</td>
<td>5–2500 U/L</td>
</tr>
<tr>
<td>AST</td>
<td>0–1083 U/L</td>
<td>0–1083 U/L</td>
<td>0–1083 U/L</td>
</tr>
<tr>
<td>BUN/UREA</td>
<td>2–130 mg/dL</td>
<td>0.6–46.4 mmol/L</td>
<td>0.034–2.730 g/L</td>
</tr>
<tr>
<td>Ca</td>
<td>1.0–16.0 mg/dL</td>
<td>0.25–4.00 mmol/L</td>
<td>10–160 mg/L</td>
</tr>
<tr>
<td>CHOL</td>
<td>6–520 mg/dL</td>
<td>0.16–13.44 mmol/L</td>
<td>0.06–5.20 g/L</td>
</tr>
<tr>
<td>CK</td>
<td>10–2036 U/L</td>
<td>10–2036 U/L</td>
<td>10–2036 U/L</td>
</tr>
<tr>
<td>CI</td>
<td>50–160 mmol/L</td>
<td>50–160 mmol/L</td>
<td>50–160 mmol/L</td>
</tr>
<tr>
<td>CREA</td>
<td>0.1–13.6 mg/dL</td>
<td>9–1202 µmol/L</td>
<td>1.0–136.0 mg/L</td>
</tr>
<tr>
<td>CRP</td>
<td>0.1–10.0 mg/dL</td>
<td>1.0–100.0 mg/L</td>
<td>1.0–100.0 mg/L</td>
</tr>
<tr>
<td>FRU</td>
<td>100–1000 µmol/L</td>
<td>100–1000 µmol/L</td>
<td>100–1000 µmol/L</td>
</tr>
<tr>
<td>GGT</td>
<td>0–952 U/L</td>
<td>0–952 U/L</td>
<td>0–952 U/L</td>
</tr>
<tr>
<td>GLU</td>
<td>10–686 mg/dL</td>
<td>0.56–38.11 mmol/L</td>
<td>0.10–6.86 g/L</td>
</tr>
<tr>
<td>K</td>
<td>0.8–10 mmol/L</td>
<td>0.8–10 mmol/L</td>
<td>0.8–10.0 mmol/L</td>
</tr>
<tr>
<td>LAC</td>
<td>0.50–12.00 mmol/L</td>
<td>0.50–12.00 mmol/L</td>
<td>0.50–12.00 mmol/L</td>
</tr>
<tr>
<td>LDH</td>
<td>50–2800 U/L</td>
<td>50–2800 U/L</td>
<td>50–2800 U/L</td>
</tr>
<tr>
<td>LIPA</td>
<td>10–6000 U/L</td>
<td>10–6000 U/L</td>
<td>10–6000 U/L</td>
</tr>
<tr>
<td>Mg</td>
<td>0.5–5.2 mg/dL</td>
<td>0.21–2.17 mmol/L</td>
<td>5.0–52.0 mg/L</td>
</tr>
<tr>
<td>Na</td>
<td>85–180 mmol/L</td>
<td>85–180 mmol/L</td>
<td>85–180 mmol/L</td>
</tr>
<tr>
<td>NH₃</td>
<td>0–950 µmol/L</td>
<td>0–950 µmol/L</td>
<td>0–950 µmol/L</td>
</tr>
<tr>
<td>PHOS</td>
<td>0.2–16.1 mg/dL</td>
<td>0.06–5.19 mmol/L</td>
<td>2.00–161.00 mg/L</td>
</tr>
<tr>
<td>TBIL</td>
<td>0.1–27.9 mg/dL</td>
<td>2–477 µmol/L</td>
<td>1.0–279.0 mg/L</td>
</tr>
<tr>
<td>TP</td>
<td>0.5–12.0 g/dL</td>
<td>5–120 g/L</td>
<td>5–120 g/L</td>
</tr>
<tr>
<td>TRIG</td>
<td>10–375 mg/dL</td>
<td>0.11–4.23 mmol/L</td>
<td>0.10–3.75 g/L</td>
</tr>
<tr>
<td>TT₄ (canine)</td>
<td>0.5–10.0 µg/dL</td>
<td>6.43–128.7 nmol/L</td>
<td>6.43–128.7 nmol/L</td>
</tr>
<tr>
<td>TT₄ (feline)</td>
<td>0.5–20.0 µg/dL</td>
<td>6.4–257.4 nmol/L</td>
<td>6.4–257.4 nmol/L</td>
</tr>
<tr>
<td>UCRE</td>
<td>6–350 mg/dL</td>
<td>0.06–3.50 g/L</td>
<td>0.06–3.50 g/L</td>
</tr>
<tr>
<td>UPRO</td>
<td>5–400 mg/dL</td>
<td>0.05–4.00 g/L</td>
<td>0.05–4.00 g/L</td>
</tr>
<tr>
<td>URIČ</td>
<td>0.1–20.0 mg/dL</td>
<td>6–1190 µmol/L</td>
<td>1–200 mg/L</td>
</tr>
</tbody>
</table>

1 µg/mL = 4.31 µmol/L

† Indicates sample types that should not be diluted.
Modifying the Settings on the Analyzer

Overview
Some of the Settings and Tools screen features allow you to customize the analyzer, such as selecting a time/date format and editing the In Process and Pending lists on the Home screen. This chapter describes how to use those features.

Changing the Language/Local Settings
Tapping the Language/Local option on the Settings screen allows you to modify the analyzer’s language, name format, unit system, time, and/or date.

Notes:
• This option is unavailable when the Catalyst Dx analyzer is processing a sample run.
• The analyzer will prompt you to restart it whenever the language/local settings are changed. You must restart the analyzer in order for the changes to take effect.

To Change the Language/Local Settings
1. Tap Settings on the Catalyst Dx Home screen.
2. Tap Language/Local.
3. Select the desired language from the Language drop-down list. When a language is chosen, the Unit System and Name Format default settings change.
4. If desired, select a different Name Format option (last name, first name or last name first name).
5. If desired, select a different Unit System option (US, SI, or French SI).
6. Tap Next.
7. If desired, update the time settings:
   a. Tap the arrows above or below the hour/minutes text boxes to increase or decrease the hours/minutes incrementally.
   b. Select the AM or PM option for your system time.
   c. Select a time format (hh:mm in 12-hour format or hh:mm in 24-hour format).
8. If desired, update the date settings:
   a. Select a date format (mm/dd/yyyy or dd/mm/yyyy). The left and right date fields (above the date format options) vary depending on the date format you choose. For example, if you choose the date format mm/dd/yyyy, the month field is the left-most field, the day field is the middle field, and the year field is the right-most field. If you choose dd/mm/yyyy, the day field is the left-most field, the month field is the middle field, and the year field is the right-most field.
   b. To change the month, tap the arrow above/below the current month selection to change the month incrementally.
   c. To change the day, tap the arrow above/below the day to increase/decrease the day incrementally.
   d. To change the year, tap the arrow above/below the year to increase/decrease the year incrementally.
9. Tap Save. When prompted, tap Yes to restart your analyzer and save the new settings.
Deleting a Patient from the Pending and In Process Lists

The Tools screen is available from the Home screen and provides options for editing the Pending and In Process lists. You can edit these lists by deleting a patient from the list.

To Delete a Patient from the Pending List

1. Tap **Tools**.
2. Tap **Edit Pending**.
3. Tap to select the patient you want to remove from the Pending list.
4. Tap **Delete** in the Delete from Pending box.

To Delete a Patient from the In Process List

You can also delete a patient from the In Process list by selecting the patient in the In Process list (on the Catalyst Dx Home screen) and then tapping **Stop Run** in the Home screen’s center display area.

1. Tap **Tools**.
2. Tap **Edit In Process**.
3. Tap to select the patient you want to remove from the In Process list.
4. Tap **Delete** in the Delete From In Process box. The slides are ejected into the waste drawer. Remove the sample using the instructions on page 20.

**Note:** The IDEXX VetLab* Station displays the “New Results” alert even though no results exist for the deleted patient run (this message only displays if you’ve selected to receive a message with new results in the New Results Alert tab on the IDEXX VetLab Station’s Settings screen).
Sample Preparation and Storage

Supported Sample Types for Catalyst CLIPs and Slides

The following sample types can be used with Catalyst CLIPs and slides:

<table>
<thead>
<tr>
<th>CLIPs/Slides</th>
<th>Abbreviation</th>
<th>Serum</th>
<th>Lithium</th>
<th>Heparin-Treated</th>
<th>Fluoride/Oxalate-Treated</th>
<th>Untreated Whole Blood [using the Catalyst* Lithium Whole Blood Separator]</th>
<th>Urine</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chem 17 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chem 15 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chem 10 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Equine 15 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>NSAID 6 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>UPC Panel</td>
<td>N/A</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>☑</td>
<td>✔</td>
</tr>
<tr>
<td>Lyte 4 CLIP</td>
<td>N/A</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Albumin</td>
<td>ALB</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alkaline Phosphatase</td>
<td>ALKP</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alanine Aminotransferase</td>
<td>ALT</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Amylase</td>
<td>AMYL</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Aspartate Aminotransferase</td>
<td>AST</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Blood Urea Nitrogen</td>
<td>BUN/UREA</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Calcium</td>
<td>Ca</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cholesterol</td>
<td>CHOL</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Creatine Kinase</td>
<td>CK</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Creatinine</td>
<td>CREA</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C-Reactive Protein</td>
<td>CRP</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Fructosamine (part number 99-0003341)</td>
<td>FRU</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Gamma-glutamyltransferase</td>
<td>GGT</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td>GLU</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lactate</td>
<td>LAC</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lactate Dehydrogenase</td>
<td>LDH</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lipase</td>
<td>LIPA</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Magnesium</td>
<td>Mg</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ammonia</td>
<td>NH₃</td>
<td>☑</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Phenobarbital</td>
<td>PHBR</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Inorganic Phosphate</td>
<td>PHOS</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total Bilirubin</td>
<td>TBIL</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Total Protein</td>
<td>TP</td>
<td>✔</td>
<td>✔</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Preparing Samples for Use on the Catalyst Dx Analyzer

You can run untreated whole blood, lithium heparinized whole blood, plasma, serum, and urine samples on the Catalyst Dx analyzer.

**IMPORTANT:** Do not use EDTA or sodium heparin for chemistry analysis.

### To Prepare an Untreated Whole Blood Sample
(Using a Lithium Heparin Whole Blood Separator)

1. Remove the green cap from the lithium heparin whole blood separator to prepare it for sample collection.

2. **Immediately** after sample collection (to avoid clotting), dispense 0.6–0.8 cc of untreated (no additive) whole blood into the lithium heparin whole blood separator using an untreated syringe with the needle removed.

   **Tip:** Use the fill line on the separator to ensure proper fill volume.

   **Note:** Heparinized samples can be used in the lithium heparin whole blood separator except when running feline AST, LDH, or CK. Double dosing may elevate the results for these assays in feline samples.

3. Gently swirl (do not invert or shake) the whole blood separator at least 5 times to mix the sample with the anticoagulant.

   **Caution:** Ensure that the cap is removed before loading the separator into the analyzer.

### To Prepare a Plasma Sample

1. Use the appropriate tube and collection device.

2. Draw the sample gently and transfer if necessary.

   **Note:** Be sure to use the correct blood-to-lithium heparin ratio.

3. Gently invert (do not shake) the sample for 30 seconds to mix.
4. Centrifuge the sample.

5. Use a transfer pipette (or the 300 µL pipette provided) to transfer the appropriate volume of sample to a Catalyst sample cup (ensure there are no bubbles in the sample cup). The volume needed varies depending on the number of slides being used in the run—for more information, see “Proper Sample Cup Volume” on pages 30–31.

To Prepare a Serum Sample

1. Use the appropriate tube and collection device.

2. Draw the sample gently and transfer if necessary.

3. Let the sample clot for a minimum of 20 minutes.

4. Centrifuge the sample.

5. Use a transfer pipette (or the 300 µL pipette provided) to transfer the appropriate volume of sample to a Catalyst sample cup (ensure there are no bubbles in the sample cup). The volume needed varies depending on the number of slides being used in the run—for more information, see “Proper Sample Cup Volume” (below).

To Prepare a Urine Sample

1. Obtain the sample through cystocentesis (recommended), catheter, or free-catch method.

2. Transfer the sample to a disposable sample tube.

3. Centrifuge the sample.

4. Use a transfer pipette (or the 300 µL pipette provided) to transfer the appropriate volume of supernatant urine to a Catalyst sample cup (ensure there are no bubbles in the sample cup). The volume needed varies depending on the number of slides being used in the run—for more information, see “Proper Sample Cup Volume” (below).
Proper Sample Cup Volume

The volume of plasma, serum, or urine sample required varies based on the number of slides being used in the run:

<table>
<thead>
<tr>
<th>Number of slides</th>
<th>Sample cup fill volume (µL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>60</td>
</tr>
<tr>
<td>2</td>
<td>70</td>
</tr>
<tr>
<td>3</td>
<td>80</td>
</tr>
<tr>
<td>4</td>
<td>90</td>
</tr>
<tr>
<td>5</td>
<td>100</td>
</tr>
<tr>
<td>6</td>
<td>110</td>
</tr>
<tr>
<td>7</td>
<td>120</td>
</tr>
<tr>
<td>8</td>
<td>130</td>
</tr>
<tr>
<td>9</td>
<td>190</td>
</tr>
<tr>
<td>10</td>
<td>200</td>
</tr>
<tr>
<td>11</td>
<td>210</td>
</tr>
<tr>
<td>12</td>
<td>220</td>
</tr>
<tr>
<td>13</td>
<td>230</td>
</tr>
<tr>
<td>14</td>
<td>240</td>
</tr>
<tr>
<td>15</td>
<td>250</td>
</tr>
<tr>
<td>16</td>
<td>260</td>
</tr>
<tr>
<td>17</td>
<td>270</td>
</tr>
<tr>
<td>18</td>
<td>280</td>
</tr>
</tbody>
</table>

Sample Inspection After Centrifugation

It is good practice to examine the sample carefully following centrifugation in a centrifuge and/or in the analyzer (by running a whole blood separator). If fibrin strands can be seen in the sample, they may interfere with sample pipetting. It may be necessary to rim the serum/plasma with a wooden stick, respin the sample, and proceed.

Various conditions, such as hemolysis, may affect results. You might also want to modify your test panel based on the following visual observations. Refer to the “Chemistry Descriptions” section on pages 47–66 for information about how each condition may affect specific chemistries.

**Note:** We recommend that after you have centrifuged a Catalyst whole blood separator that you inspect the sample for the conditions listed above.

**Hemolysis**

**Visual:** Sample has a transparent reddish hue ranging from pale pink to deep red.

**Indications:** Damage to red blood cells during sample preparation or intravascular hemolysis.

**Icterus**

**Visual:** Plasma has a transparent yellow to opaque brown color.

**Indications:** Obstructive or toxic liver disease, intravascular hemolysis.

**Lipemia**

**Visual:** Sample has a pale, milky appearance, possibly with floating fat globules.

**Indications:** Recent ingestion of a fatty meal or dysfunction in lipid metabolism.
Sample Storage

We recommend that you prepare and analyze samples immediately after collection for best results. However if storage is necessary, follow these sample storage and testing guidelines.

Storing Serum/Plasma

For storage, the serum or plasma must be separated and removed immediately from the blood cells. Do not attempt to pour off the sample.

- Using a transfer pipette, carefully transfer the serum or plasma to an untreated collection tube, taking care not to draw up any white or red blood cells.
- Cap the tube tightly to avoid contamination and evaporation. Avoid frothing at any stage as this damages the serum proteins.

If you cannot perform analysis within four hours of drawing and processing the sample, refrigerate it at 2°C–8°C (36°F–46°F). If you cannot perform analysis for more than 48 hours, you should freeze the serum/plasma at -18°C (0°F). Frozen serum/plasma can be stored for up to 1 month.

Notes:

- For additional information on the effects of delays in removing serum or plasma from the cells, see the “Chemistry Descriptions” section on pages 46–66.
- See the calcium (Ca), total bilirubin (TBIL), lactate dehydrogenase (LDH), ammonia (NH3), electrolytes (Na, K, Cl), and glucose (GLU) chemistry descriptions for additional special handling and storage requirements.
- IDEXX does not recommend freezing samples that will be used to run electrolytes or NH3.

Storing Whole Blood

Lithium heparinized whole blood samples should be analyzed immediately. Samples that are not analyzed within 30 minutes should be placed in a tube to be separated and stored.

Important: Do not store whole blood samples in whole blood separators.

Storing Urine

Urine should be tested within two hours. Do not store urine in the refrigerator for more than 24 hours. Urine should not be stored in the freezer.

Analysis of Stored Samples

For samples stored at 2°C–8°C (36°F–46°F) and at -18°C (0°F):

- Allow the samples to come to room temperature (19°C–27°C/66°F–81°F).
- Mix the samples gently, but thoroughly, by inversion. Do not shake.
- Centrifuge the samples to remove any fibrin particles (or urine sediment) that may have formed during storage.
- Analyze the samples immediately after centrifugation.
Quality Control

Overview

The purpose of quality control (QC) is to verify the integrity of your slides and also to verify that your Catalyst Dx* Analyzer is functioning properly.

You should run a QC test:

- When the analyzer is first installed.
- After cleaning the internal components of the analyzer.
- If the analyzer has been moved.
- To verify system performance.

Quality Control Materials

VetTrol* Control

In each box of VetTrol Control, there are four vials containing freeze-dried powder (dark brown bottle marked “VetTrol Control”) and four vials containing diluent (lighter bottles marked “Diluent for VetTrol”). The lot numbers for the diluent and the control are different and can be found on the product packaging.

For more information about VetTrol Control, see its package insert.

Storage

Control and diluent vials should be stored frozen (-18°C/0°F). Discard opened control vials within 24 hours. Expired or unwanted material should be discarded with other clinical waste.

Note: Do not store in the freezer door; only in the main freezer compartment.

Stability and Handling

For most chemistries, VetTrol Control can be used up to 24 hours after reconstitution when it is stored in the refrigerator and equilibrated to room temperature before running (do not leave at room temperature for more than two hours). For creatine kinase and ammonia values, VetTrol Control fluid should be used within two hours following reconstitution. Exposure to light will affect total bilirubin and creatine kinase results. Ammonia concentration will increase with time.

UPRO Control

In each box of UPRO Control, there are six vials containing the control fluid. The lot number can be found on the product packaging.

Storage

Control fluid should be refrigerated (2°C–8°C/36°F–46°F). Discard at the expiration date. Expired or unwanted material should be discarded with other clinical waste.

Stability and Handling

Use within 24 hours after opening (refrigerate when not in use).
Advanced Control
In each box of Advanced Control, there is one vial containing the control fluid. The lot number can be found on the product packaging.

Note: Each vial contains enough fluid for 2 runs, in the event a secondary run is necessary.

Storage
Store frozen until the expiration date, or store in the refrigerator for up to 5 days.

Stability and Handling
Once opened, Advanced Control cannot be stored and reused—discard remaining fluid after use.

PHBR Control
In each box of PHBR Control, there are six vials containing the control fluid. The lot number can be found on the product packaging.

Storage
Store frozen until the expiration date, or store in the refrigerator for up to 7 days.

Stability and Handling
Once thawed, PHBR Control cannot be stored and reused—discard remaining fluid after use.

Quality Control CLIPs and Slides
IDEXX recommends that you perform monthly quality control testing after you have cleaned the internal components of your analyzer. The convenient Catalyst QC CLIP contains all of the chemistry slides needed to perform this task. It is also recommended that you also perform a quality control for electrolytes using the Catalyst* Lyte 4 CLIP.

Run the QC CLIP and the Lyte 4 CLIP
Use the convenient QC CLIP and the Lyte 4 CLIP in conjunction with the VetTrol Control fluid to perform quality control on your Catalyst Dx analyzer. It is recommended that you wait at least 30 minutes after running any slides before running the QC CLIP.

OR

Run Individual Slides
You can use individual slides to create your own QC panel and perform a quality control test (one slide per group). If you want to use individual slides to run quality control, we recommend a minimum of one slide from each of the groups below.

Group 1
- NH₃

Group 2
- AMYL
- CHOL
- GLU
- LAC
- LIPA
- TBIL
- TP
- TRIG
Preparing Control Fluid

The instructions for preparing control fluid vary depending on the type of control you are preparing.

To Prepare VetTrol Control Fluid

1. Remove one diluent and one control vial from freezer. Allow 60–90 minutes for vials to acclimate to room temperature.
2. Slowly invert the diluent vial several times to thoroughly mix the contents. Do not shake.
3. Gently tap the control vial on the counter several times to dislodge any material adhering to the stopper.
4. Remove the seal and stopper from each vial just before adding the diluent to control. Do not leave the vials open.
5. Transfer exactly 3.0 mL of diluent to the control vial, using a clean, dry, Class A volumetric pipette or an equivalent automatic pipette. Discard the remaining diluent.

**IMPORTANT:** Measurement must be precise or results will be incorrect.

6. Replace the stopper on the control vial and hold it firmly in place. Gently invert the vial 6–10 times every 10 minutes for 1 hour (the use of a slow rocker is recommended). Do not shake. Reconstitution, with occasional inversion, will take 45–60 minutes. Visually verify that all freeze-dried material is dissolved before using.
7. Run quality control on the Catalyst Dx analyzer (see instructions on page 32).

To Prepare UPRO Control Fluid

1. Take one vial of UPRO Control out of the refrigerator and gently invert it 6–10 times to mix thoroughly.
2. Transfer 300 µL of UPRO Control into a Catalyst sample cup (to be loaded in the sample drawer).
3. Let the contents in the sample cups reach room temperature (approximately 10 minutes).
4. Run quality control on the analyzer (see instructions on page 32).
To Prepare Advanced Control Fluid
1. If the Advanced Control has been frozen, allow it to thaw for 30 minutes prior to use.
2. Invert the Advanced Control vial at least 5 times.
3. Transfer the contents of the Advanced Control vial to a Catalyst* sample cup.
4. Run quality control on the analyzer (see instructions on page 32).

To Prepare PHBR Control Fluid
1. Take one vial of PHBR Control out of the freezer and allow it to reach room temperature (approximately 60 minutes).
2. Once you have confirmed that there is no visible frozen material in the vial, gently invert it 6–10 times to mix thoroughly.
3. Transfer 300 μL of PHBR Control into a Catalyst* sample cup.
   
   **Note:** You will need one PHBR slide wash and one PHBR slide for the quality control procedure.

4. Run quality control on the analyzer (see instructions on page 32).

Running Quality Control

To Run Quality Control
1. Prepare the fluid using the steps on page 31.
2. Tap **Instruments** on the IDEXX VetLab Station Home screen. The Instruments screen displays.
3. Tap the **Catalyst Dx** tab.
4. Tap **Quality Control**.
5. Tap the quality control lot number you are using and tap **Run QC**.
6. Tap the QC information in the Pending list on the Catalyst Dx Home screen and then tap **Load**.
7. If you are running VetTrol Control, follow the on-screen instructions for loading the quality control materials and then tap **Run**.
   OR
   If you are running Advanced Control, follow these steps:
   a. Load the sample cup containing Advanced Control and an applicable slide into the sample drawer.
   b. Tap **Run**.
   OR
   If you are running PHBR Control, follow these steps:
   a. Open the tip/diluent drawer and load PHBR wash and fill it with tips.
   b. Close the drawer and tap **Next**.
   c. Load the sample cup containing 300 μL of PHBR Control and a PHBR slide into the sample drawer.
d. Tap **Run**.

OR

If you are running UPRO Control, follow these steps:

a. Load the sample cup containing 300 µL of UPRO Control and a UPRO slide (**do not load a UCRE slide**) into the sample drawer.

b. Tap **Run**.

8. Once the results are complete, you can view the results using these methods:

   - Tap **View** in the View Results message (this message displays by default when a run is complete) to access the Records: Test Results screen.

   - Tap the QC run in the Recent Test Results list on the IDEXX VetLab Station Home screen and then tap **View**.

   - Tap **Records** on the IDEXX VetLab Station Home screen, select the QC run you want to view, tap **View Records**, tap the desired test result, and then tap **View Results**.
Maintenance

Overview
In addition to performing monthly quality control checks on the Catalyst Dx* analyzer, it is recommended that you:

- Clean the analyzer internally and externally.
- Upgrade the software promptly.

Upgrading the Software
As new features and functionality are added to the Catalyst Dx analyzer, you will receive software upgrades from IDEXX. If you have SmartService* Solutions, the upgrade will be sent via your IDEXX VetLab* Station automatically. If you do not have SmartService* Solutions, you will receive your upgrade in the mail. Be sure to read the software notes contained with each new release.

Opening/Closing the Maintenance Access Doors
The maintenance access doors provide access to the internal components of the analyzer. You will need to open the maintenance access doors during the cleaning procedure, when clearing a slide jam, etc.

Note: The procedure for opening/closing the maintenance access doors varies depending on the configuration of your analyzer.

To Open the Maintenance Access Doors
1. Press up on the door panel below the touch screen.
2. If the area above the door panel has vertical plastic slats (see photo 2A below), push down firmly on the door panel. The maintenance access doors are released.
   OR
   If the area above the door panel has a metal handle (see photo 2B below), pull down on both sides of the metal handle above the door panel until you hear a click. The maintenance access doors are released.
3. Place a finger beneath the center of the maintenance access doors and push up until the doors lock in place.
To Close the Maintenance Access Doors

1. If the area above the door panel has vertical plastic slats (see photo 1A below), push and hold down on the door panel. The maintenance access doors close automatically. OR
   If the area above the door panel has a metal handle (see photo 1B below), pull down on both sides of the metal handle above the door panel until you hear a click. The maintenance access doors close automatically.

2. Press up on the door panel below the touch screen until it clicks.

Cleaning the Internal Components of the Analyzer

To ensure optimal performance of your analyzer, it is important that you clean the internal components (incubator ring, optics window, and carousel) monthly and before performing quality control.

It is recommended that you wear clean powder-free latex or nitrile gloves when cleaning the internal components of the analyzer. Wearing clean latex gloves helps to avoid smudges on the components and ensures an effective cleaning.

IMPORTANT:

- Never use canned air in or around the Catalyst Dx analyzer.
- Never use cleaning materials (such as alcohol cleaning wipes containing sodium bicarbonate) that leave a residue once the alcohol/solvent evaporates.

To Clean the Internal Components

1. Tap Tools.

2. Tap Clean Analyzer.

3. Open the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).

4. Remove the black whole blood separator carriers, any whole blood separators or sample cups from the sample drawer, and the white centrifuge shield. Then, clean the black carriers and white shield with an IDEXX-supported alcohol prep pad and return them to their positions.
5. Lift the sample drawer assembly by pressing the center latch and lifting up.

6. Remove the carousel:
   - If there is a wire handle in the middle of the carousel (see photos 6a below), remove the carousel by lifting the center wire handle in the middle of the carousel straight up.
   - If there is a plastic handle in the middle of the carousel (see photo 6b below), remove the carousel by lifting up with the handle.

7. Using an IDEXX-supported alcohol prep pad, wipe the incubator ring track in a counterclockwise direction (do not wipe the optics or ion windows at this time). Repeat this step at least three times using a new tissue for each wipe.

8. Clean the optics, ion windows, and reference tile on the carousel using the instructions in step 7.

9. Using a dry optical tissue, dry the optics, ion windows, and reference tile, ensuring all signs of dampness have evaporated from the cleaned components. If streaks or smudges remain, repeat the cleaning process.
10. Replace the carousel on the incubator ring track:
   - If there is a wire handle in the middle of the carousel, ensure it engages securely with the two carousel mounting posts (see photo 10a below). Then, lower the wire handle.
   - If there is a plastic handle in the middle of the carousel, position the front of the carousel below the railing on the incubator ring track and then press the carousel down so that it locks into place (see photo 10b below).

11. Lower the sample drawer assembly and ensure that it is locked into place.
12. Close the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).
13. On the Catalyst Dx touch screen, tap Done. The analyzer initializes.

**Cleaning the Fan Filter**

Clean the fan filter once quarterly in normal laboratory conditions. If the Catalyst Dx analyzer is operated in environmental conditions that are dusty or dirty, the fan filter may need to be cleaned on a monthly basis instead of quarterly.

**To Clean the Fan Filter**

1. Locate the fan filter on the back right side of the analyzer.
2. Gently pull up on the black plastic tab to move the filter upward. Then, hold both sides of the filter to remove it.
3. Vacuum the filter thoroughly.
4. Slide the filter back into place.
Cleaning the Centrifuge

Clean the centrifuge as needed to remove any residue from the whole blood separator.

To Clean the Centrifuge

1. Open the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).
2. Remove the whole blood separator carriers over the sample drawer stations and ensure that there are no sample cups or whole blood separators in the sample drawers.
3. Return the whole blood separator carriers to their positions.
4. Lift the sample drawer assembly by pressing the center latch and lifting up.
5. To the right of the slide loading stations, gently pull the tab up on the white centrifuge shield to remove it from the analyzer and then gently clean it with mild soap and water to remove the residue. Once it’s rinsed and dried thoroughly, replace it by aligning the notch on the shield to the recessed section on the centrifuge and gently press down. The shield is properly seated in the centrifuge when it is level and does not spin when attempting to turn.
6. Lower the sample drawer assembly. Push in on the center latch to ensure that it is locked into place.
7. Close the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).

Cleaning the Exterior of the Analyzer

Always disconnect the power cable before cleaning the analyzer.

Clean the outside of the analyzer with a damp (not wet) lint-free cloth. A mild liquid soap will remove grease. Do not use any of the following near the analyzer: organic solvents, ammonia-based cleaners, ink markers, canned air, sprays containing volatile liquids, insecticides, disinfectant, polish, or room freshener.

Care should be taken not to spill any samples, chemicals, cleaning agents, water, or other fluids on/in the analyzer.

**Note:** Dust and animal hair can lead to analyzer failures. Routinely dust off the analyzer with a damp cloth and dust around its location. Do not block the cooling vents under the analyzer by allowing paper, loose materials, or dust to accumulate.

**WARNING:** Never wipe the analyzer or its surroundings with ammonia based cleaning products. Avoid urine odors around analyzer. Ammonia in the atmosphere will falsely increase ammonia (NH₃) quality control and patient test results.

Cleaning the Screen

If the screen gets dirty, apply an antistatic screen cleaning agent (NOT ammonia-based) to a clean cloth or paper towel and wipe the screen. Do not spray the cleaner directly onto the screen as liquid can run inside the case and damage electrical circuits. Take care not to scratch the screen.
Emptying the Waste Drawer

It is essential that you empty the waste drawer when prompted by the analyzer. The analyzer will not operate when the waste drawer is full. Pull the waste drawer to remove it from the analyzer. After you have emptied and replaced the waste drawer, tap Yes to confirm that the drawer has been emptied.

IMPORTANT: The waste drawer should not be opened or removed during a run.
Troubleshooting

Differences in Results

With a Commercial Laboratory or Other Instrument
Reference ranges must be created for each analyte and each new instrument or method of analysis. Every commercial laboratory must establish its own species reference ranges for the equipment and methodology used. IDEXX is continually doing this work for you with every software release.

Comparing results from different laboratories that may be using different equipment or methods is imprecise at best. Any comparisons should be performed on the same sample that has been “split,” stored under like conditions, and tested at approximately the same time. Compare each result to the reference range stated by IDEXX or the commercial laboratory (as appropriate). Each result should have the same relationship to its method’s reference range. For instance, a sample giving a Catalyst Dx* result that is slightly below the Catalyst Dx analyzer’s normal range should give a laboratory result slightly below the laboratory’s normal range.

Status Messages
Status messages are displayed in two different locations on the analyzer. Some are reported in the center display area on the Catalyst Dx Home screen. Others appear in the status bar at the top of the screen. These messages provide information on the current state of the analyzer.

Note: If you cannot run a sample on the analyzer, be sure to check the center display area and status bar on the Home screen for helpful messages.

Home Screen Messages

<table>
<thead>
<tr>
<th>Icon</th>
<th>Message</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td><img src="Icon" alt="Clock with red background" /></td>
<td>Sample drawers are in use</td>
<td>The analyzer is processing samples from both sample drawers. You cannot process another patient’s sample at this time. When this icon disappears from the Home screen, a sample drawer has become available again for use (after approximately 2 minutes).</td>
</tr>
<tr>
<td><img src="Icon" alt="Waste basket with red background" /></td>
<td>Empty the waste drawer</td>
<td>The analyzer has determined that the maximum number of slides and/or tips is currently in the waste drawer. To prevent waste overflow, please empty the waste drawer. Once you have confirmed that it is empty, this icon will disappear and you can use the analyzer as needed.</td>
</tr>
</tbody>
</table>
### Add pipette tips

The analyzer has determined that there are not enough tips to process a sample on the analyzer. Open the tip drawer and fill it with up to 12 pipette tips. The icon will then disappear and you can use the analyzer as needed.

### Dilution drawer in use

The analyzer is currently performing an automated dilution and/or is processing a test that requires a reagent pack. You cannot perform another automated dilution, including a UPC ratio, or run another test that includes a reagent pack at this time.

You can process a patient’s sample that does not require an automated dilution or run a test that does not require a reagent pack while this icon is present.

**Important:** Do not open the tip/diluent drawer while an automated dilution or a test that requires a reagent pack is in process.

### TT4 run in process

The analyzer is currently running a PHBR slide or initializing. You must wait for this icon to disappear before running another patient sample.

**Important:** Do not open the tip/diluent drawer while a PHBR run is in process.

### Cleaning required

The analyzer must be cleaned before another sample can be processed. Once you clean the internal components of the analyzer successfully, this icon will disappear and you can use the analyzer as needed.

### Calibration required

The analyzer must be calibrated before another sample can be processed. Once the analyzer has been calibrated successfully, this icon will disappear and you can use the analyzer as needed. Please contact IDEXX Technical Support.

### Status Bar Messages

<table>
<thead>
<tr>
<th>This status message...</th>
<th>Indicates...</th>
</tr>
</thead>
<tbody>
<tr>
<td>Close Tip Drawer</td>
<td>The tip drawer is open.</td>
</tr>
<tr>
<td>Initialize Analyzer</td>
<td>The maintenance access doors are open.</td>
</tr>
<tr>
<td>Maintenance Required</td>
<td>The analyzer requires cleaning.</td>
</tr>
<tr>
<td>Instrument Maintenance</td>
<td>The analyzer is performing an automatic self-check to ensure optimal performance of the optics. (This message will display periodically.)</td>
</tr>
<tr>
<td>Initializing</td>
<td>The analyzer is preparing for a ready state.</td>
</tr>
</tbody>
</table>
This status message... Indicates...

| Diagnostic                     | A white reference slide has been placed in the analyzer for calibration. |
| Initialization Required       | An error has occurred with the analyzer. Please contact IDEXX Technical Support. |
| Processing Dilution: Please Wait | An automated dilution run and/or a test that requires a reagent pack is in process. You must wait for the automated dilution/reagent pack run to complete prior to starting another automated dilution/reagent pack run. |
| Not ready—PHBR run in process | The analyzer is currently running a PHBR slide. |

Removing a Slide Jam

If there is a slide jam inside of the Catalyst Dx analyzer, use the following procedure to remove the slides.

To Remove a Slide Jam

1. Open the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).
2. Remove any slides and sample from the sample drawer.
3. Lift the sample drawer assembly by pressing the center latch and lifting up.
4. Remove the carousel:
   - If there is a wire handle in the middle of the carousel (see photos 4a below), remove the carousel by lifting the center wire handle in the middle of the carousel straight up.
   - If there is a green plastic handle in the middle of the carousel (see photo 4b below), remove the carousel by lifting up with the handle.
5. Ensure all slides have been removed from the carousel.

6. Replace the carousel on the incubator ring track:
   - If there is a wire handle in the middle of the carousel, ensure it engages securely with the two carousel mounting posts (see photo 6a below). Then, lower the wire handle.
   - If there is a green plastic handle in the middle of the carousel, position the front of the carousel below the railing on the incubator ring track and then press the carousel down so that it locks into place (see photo 6b below).

7. Lower the sample drawer assembly and secure the latch.

8. Close the maintenance access doors (for detailed instructions, see “Opening/Closing the Maintenance Access Doors” on pages 34–35).

9. Initialize the analyzer.
Appendices

Chemistry Descriptions

Serving veterinarians throughout the world, IDEXX Laboratories understands that medical content, including interpretation of diagnostic results and medical protocols may vary from country to country. A medical review board has approved the content presented in this document.

IDEXX has more than 40 reference laboratories worldwide employing over 100 veterinarians. If you have any questions about the medical content or interpretation of results in this document, please contact IDEXX Laboratories.

Introduction to Biochemical Profiling

By performing appropriate biochemical tests on quality samples, you can obtain information that, when combined with patient history and clinical findings, should assist you in making an accurate diagnosis. Appropriate biochemical tests are also essential for monitoring and prognostication purposes once a diagnosis is achieved.

Single tests are helpful in particular circumstances, such as following the course of an identified disease or for monitoring the effect of therapy. However, many individual chemistry tests give information about different organ systems and should be used in combination with other tests (panels or profiles) to help characterize disease.

Alanine Aminotransferase (ALT)

For practical purposes, the enzyme alanine aminotransferase is specific to the liver in dogs and cats. It is found in the hepatocyte cytoplasm and may be released into the blood during both reversible and irreversible (cell necrosis) changes.

Principal Reason for Performing the Test

To investigate hepatocellular injury in dogs and cats.

Note: This test is not useful in the detection of liver disease in ruminants, horses, and pigs as the enzyme activity in the liver is very low. Even with severe liver disease in these species, the increase in activity is minimal.

Most Common Abnormality Indicated by the Test

Hepatocellular injury.

Sample Type and Precautions

Remove plasma or serum promptly from the cells or clot. Hemolyzed specimens should not be used because ALT contamination from red blood cells will occur. If plasma is being collected, use only lithium heparinized samples.

Complementary Tests

Alanine aminotransferase activity is usually determined in conjunction with other tests of hepatic function or damage.
**Reaction Sequence**

\[
\text{alanine + } \alpha\text{-ketoglutarate} \xrightleftharpoons{\text{ALT}}_{\text{P-5-P}} \text{ pyruvate + glutamate}
\]

\[
\text{pyruvate + NADH + } H^+ \xrightleftharpoons{\text{LDH}} \text{ lactate + NAD}^+
\]

**Albumin (ALB)**

Albumin forms the largest fraction of the total serum protein in the healthy animal. It is synthesized solely by the liver, has a relatively low molecular weight, and plays an important role in the transport of endogenous and exogenous compounds by binding with those compounds. Albumin also plays a major role related to osmoregulation.

**Principal Reasons for Performing the Test**

To investigate causes of hypoalbuminemia: protein-losing nephropathy, protein-losing enteropathy, as well as hepatic insufficiency (decreased production) and decreased absorption due to malabsorption (gastrointestinal disease) or malnutrition. In addition, it is helpful in characterizing the degree of dehydration with increases in serum albumin concentrations and it is commonly decreased with active inflammatory disease (negative acute phase reactant).

The test should not be performed in isolation because of its lack of specificity.

**Most Common Abnormalities Indicated by the Test**

Decreased albumin—inflammatory disease, protein-losing nephropathy and enteropathy, and decreased production (hepatic insufficiency).

Increased albumin—dehydration.

**Sample Type and Precautions**

Remove plasma or serum promptly from the cells or clot. Hemolysis may occur if the sample is not handled properly. Although dry-slide technology minimizes the interfering effect of mild-to-moderate hemolysis, marked hemolysis will cause an increased albumin value.

**Complementary Tests**

Albumin concentration is usually determined in conjunction with the measurement of total protein and other tests of renal and hepatic function. When albumin is measured with total protein, the total globulins will be calculated automatically and given with the results.

**Reaction Sequence**

\[
\text{albumin + bromocresol green (BCG) } \xrightarrow{\text{BCG-albumin complex}}
\]

**Alkaline Phosphatase (ALKP)**

The enzyme alkaline phosphatase is found in many body tissues. Highest levels are found in the kidney cortex, small intestinal mucosa, and osteoblasts. The enzyme is also present in the liver primarily located on the bile canalicular; thus an increase in ALKP may indicate cholestasis.

In cats and horses, the half-life of hepatic alkaline phosphatase is very short for ALKP and even shorter for other natural tissue sources of ALKP due to rapid renal excretion/metabolism. Sensitivity of the test in cats and horses is low. Since the nonhepatic sources of ALKP have relatively short half-lives compared to the hepatic source, a mild-to-modest increase in ALKP in these species can be a specific indicator of cholestasis.
**Principal Reason for Performing the Test**
As an indicator of hepatic and/or biliary disease.

**Most Common Abnormality Indicated by the Test**
Obstructive changes in the biliary system. A special consideration for interpreting ALKP changes in the dog is required because there are “induced” forms of ALKP due to glucocorticoids and other influences that are not associated with the natural tissue sources of ALKP. The nonhepatic sources of ALKP (bone, intestinal, placental) in the dog will only rarely be measured as high as threefold above the high end of the reference range because of their relative short half-lives compared to the induced and hepatic forms of ALKP. With both the induced and hepatic source (cholestasis) of ALKP, serum enzyme activities are commonly greater than the threefold increase; therefore, when a greater than threefold increase is noted in ALKP in the dog, either cholestasis or induced enzyme is suspected.

**Sample Type and Precautions**
Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Hemolyzed specimens should not be used because ALKP contamination from red blood cells will increase results while hemoglobin decreases results. Above normal total bilirubin levels may reduce ALKP results.

**Complementary Tests**
Alkaline phosphatase activity is usually determined in conjunction with other tests of hepatic function and damage.

**Reaction Sequence**
\[
p\text{-nitropheryl phosphate} + \text{Mg}^{2+} + \text{AMP} \xrightarrow{\text{ALKP}} p\text{-nitrophenol} + H_3PO_4
\]

**Ammonia (NH}_3\)**
Ammonia is the catabolic product of protein digestion and is extremely toxic. It is converted rapidly in the liver to urea which is eliminated from the body by the kidneys.

**Principal Reason for Performing the Test**
To evaluate hepatic function.

**Most Common Abnormality Indicated by the Test**
Increased ammonia—decreased hepatic functional mass or hepatic vascular shunt.

**Sample Type and Precautions**
Use only lithium heparinized samples.

Blood should be processed and centrifuged immediately following collection; for this reason, plasma is recommended as the sample of choice.

Ammonia measurements in either plasma or serum are significantly affected by environmental factors and/or the passage of time. **Minimal exposure of the sample to the air is essential.** All sample containers should be capped unless sample is being introduced or withdrawn. Do not attempt to measure ammonia in hemolyzed samples. Contamination from the red blood cells will invalidate the test.
Complementary Tests
Ammonia may be determined in isolation but more often in conjunction with other tests of hepatic damage or dysfunction, such as pre- and postprandial bile acids.

Reaction Sequence
\[ \text{NH}_4^+ + \text{bromophenol blue (ammonia indicator)} \rightarrow \text{blue dye} \]

Amylase (AMYL)
This section should be read in conjunction with the Lipase (LIPA) section.

The main source of serum amylase is the pancreas, although pathology of the liver and small intestine may result in significant elevations of this enzyme (above the reference range). Since amylase is cleared by the kidneys, renal pathology may also result in elevation of amylase independent of pancreatic disease.

Principal Reason for Performing the Test
As an indicator of pancreatic disease and potential acute pancreatitis.

Most Common Abnormality Indicted by the Test
Acute necrotizing pancreatitis.

Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. Hemolyzed specimens should not be used. Do not use oxalate, citrate, or EDTA anticoagulants. If plasma is being collected, use only lithium heparinized samples.

Blood samples should be taken within one day of the onset of symptoms suggesting acute pancreatitis.

Complementary Tests
Amylase and lipase are usually determined in conjunction with one another. Evaluation of a comprehensive chemistry profile including electrolytes is generally recommended because of secondary effects of acute pancreatitis. Specific pancreatic lipase should be considered in suspected cases of pancreatitis.

Reaction Sequence
\[ \text{dyed amylopectin} \xrightarrow{\text{amylase}} \text{dyed saccharides} \]

Aspartate Aminotransferase (AST)
The enzyme aspartate aminotransferase is present in large amounts in multiple tissues of dogs, cats, and many other animal species. Hepatocytes, cardiac muscle cells, and skeletal muscle cells have relatively high concentrations of AST. It is found in the cytoplasm and mitochondria of the cells and is released into the blood during cell injury. If no increase in ALT is seen in conjunction with an increased AST in the dog and cat, cardiac or skeletal muscle cell injury is most likely. For increased AST values with equine, bovine, and porcine samples, liver, cardiac, and skeletal muscle cell injury must be considered.
Principal Reason for Performing the Test
To investigate damage to liver, cardiac, or skeletal muscle.

Most Common Abnormalities Indicated by the Test
Dogs and cats—cardiac or skeletal muscle injury when ALT is not increased; liver, cardiac, or skeletal muscle injury if both ALT and AST are increased.
Horses, cows, and pigs—liver, cardiac, or skeletal muscle injury.

Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. Hemolyzed specimens should not be used because AST contamination from red blood cells will occur. EDTA and fluoride/oxalate should not be used as anticoagulants. If plasma is being collected, use only lithium heparinized samples.

Blood samples should be processed and centrifuged immediately after collection. Even slight hemolysis can cause marked increases in activity because of high intracellular concentrations of AST in red blood cells.

Complementary Tests
Aspartate aminotransferase activity is usually determined in conjunction with other tests of liver, cardiac, or skeletal muscle function or damage.

Reaction Sequence
\[
\text{aspartate} + \alpha\text{-ketoglutarate} \xrightarrow{\text{AST}} \text{oxaloacetate} + \text{glutamate}
\]
\[
\text{oxaloacetate} \xrightarrow{\text{oxaloacetate decarboxylase}} \text{pyruvate} + \text{CO}_2
\]
\[
\text{pyruvate} + \text{phosphate} + \text{O}_2 \xrightarrow{\text{pyruvate oxidase}} \text{hydrogen peroxide} + \text{acetylphosphate}
\]
\[
\text{hydrogen peroxide} + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye}
\]

Blood Urea Nitrogen (BUN)
The catabolism of proteins results in the production of ammonia, which is extremely toxic. Ammonia is converted to urea in the liver and eliminated from the body by glomerular filtration in the kidneys.

Principal Reason for Performing the Test
As an indicator of renal disease or pathologic conditions that result in bleeding into the gastrointestinal tract.

Most Common Abnormalities Indicated by the Test
Increased urea—prerenal, postrenal and renal azotemia with decreased glomerular filtration rate; high protein diet or bleeding into the gastrointestinal tract.
Decreased urea—decreased protein intake; hepatic insufficiency; diuresis.

Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples.

Blood should not be drawn for urea determination within six hours of a meal. Do not use sodium fluoride or EDTA as anticoagulant. Samples that contain hemoglobin increase urea nitrogen.
**Complementary Tests**

Urea concentration should usually be determined in conjunction with measurements of creatinine, inorganic phosphate, total protein, albumin, and a complete urinalysis. Urea concentration is influenced by high protein diet rather than creatinine.

**Reaction Sequence**

\[ \text{H}_2\text{NCONH}_2 + \text{H}_2\text{O} \xrightarrow{\text{urease}} 2\text{NH}_3 + \text{CO}_2 \]

\[ \text{NH}_3 + \text{ammonia indicator} \rightarrow \text{dye} \]

**Calcium (Ca)**

Calcium is an essential element that is involved in many body systems. These include the skeleton, enzyme activation, muscle metabolism, blood coagulation, and osmoregulation. In the blood, calcium exists in ionized and protein bound forms. Factors governing the total plasma, whole blood, or serum concentration are complex and include interaction with other chemical moieties, proteins, and hormones.

Calcium, phosphorus, and albumin metabolism are interdependent.

**Principal Reason for Performing the Test**

As an indicator of certain neoplasias, bone disease, parathyroid disease, eclampsia, and renal disease.

**Most Common Abnormalities Indicated by the Test**

Increased calcium—hypercalcemia of malignancy (due to tumor release of PTH-like substances), spurious.

Decreased calcium—potential renal failure with resultant hyperphosphatemia, dietary, spurious.

**Sample Type and Precautions**

Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples.

Centrifugation should take place quickly after the sample has been drawn. The sample should not be exposed to the air for long periods. Glassware must be scrupulously cleaned to avoid contamination by sources of calcium (e.g., detergents). Prolonged contact with the clot may lead to lowered calcium values due to dilution by red blood cell water.

Do not use tubes containing fluoride, oxalate, citrate, or EDTA as these agents will cause significant negative interference due to calcium chelation.

If analysis cannot be performed within four hours, the sample should be removed from the red blood cells and refrigerated in a tightly stoppered container at 2°C–8°C (36°F–46°F) for short-term storage (up to 24 hours). The sample should not be frozen. The sample must be allowed to reach room temperature before analysis.

**Complementary Tests**

Calcium should be determined in conjunction with measurements of inorganic phosphate, albumin, total protein, and glucose. Ionized calcium measurement will provide more specific information related to the physiologic form of calcium.
**Chloride (Cl)**
Chloride is the major anion, predominantly in the extracellular spaces, where it maintains cellular integrity by influencing osmotic pressure. Chloride determination is significant in monitoring acid-base balance and water balance.

**Principal Reason for Performing the Test**
Low chloride levels are usually found in severe vomiting or diarrhea, ulcerative colitis, severe burns, heat exhaustion, fever, and acute infections. Elevated values are found in dehydration, hyperventilation, anemia, and cardiac decompensation.

**Most Common Abnormalities Indicated by the Test**
Hyperchloremia: if elevated with sodium then the same cause of hypernatremia. Without a concurrent increase in sodium: hyperchloremic acidosis: GI or renal loss of HCO₃.<br>Hypochloremia: (without related change in sodium): upper GI tract loss (vomiting).

**Sample Type and Precautions**
Avoid hemolysis—sample should be run as soon as possible after serum or plasma is separated from the cells or clot. If plasma is being collected, use only lithium heparinized samples.

Do not freeze samples for use with the Catalyst Dx analyzer.

**Complementary Tests**
Sodium, potassium, and chloride should always be assayed together to determine electrolyte balance. If sodium, potassium, chloride, and bicarbonate are measured together, accurate assessment of metabolic acid-base physiology is possible.

**Reaction Sequence**
\[
\text{Chloride + fluorescent dye} \rightarrow \text{fluorescence change}
\]

**Cholesterol (CHOL)**
Serum cholesterol occurs predominantly at high concentration in the esterified form; the remainder is in the free form. Cholesterol is synthesized in the liver and other tissues and is also absorbed in the free form from the small intestine. It is esterified in the liver and is the precursor of steroid hormones.

Cholesterol is broken down in the liver to bile acids and eliminated via the bile duct.

**Principal Reason for Performing the Test**
May be a marker for cholestasis or endocrine disease such as hypothyroidism, hyperadrenocorticism, diabetes mellitus, as well as nephrotic syndrome.

**Most Common Abnormality Indicated by the Test**
Increased cholesterol—hypothyroidism, postprandial, nephrotic syndrome.

**Sample Type and Precautions**
Remove plasma or serum promptly from the cells or clot. Blood should not be drawn within 12 hours of a meal. If plasma is being collected, use only lithium heparinized samples.
Complementary Tests
Cholesterol measurements should not be performed in isolation but as part of a profile of tests to investigate endocrine, hepatic, and renal disease. If high cholesterol is found in the absence of diabetes, hepatic, or renal disease, hypothyroidism may be present. This can be evaluated by measuring thyroid function.

Reaction Sequence
\[
\text{lipoprotein} \xrightarrow{\text{surfactant}} \text{TX100} \rightarrow \text{cholesterol + cholesterol esters + proteins}
\]

\[
\text{cholesterol esters + H}_2\text{O} \xrightarrow{\text{cholesterol ester hydrolase}} \text{cholesterol + fatty acids}
\]

\[
\text{cholesterol + O}_2 \xrightarrow{\text{cholesterol oxidase}} \text{cholest-4-en-3-one + H}_2\text{O}_2
\]

\[
\text{H}_2\text{O}_2 + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye + 2H}_2\text{O}
\]

Creatine Kinase (CK)
Creatine kinase is found at high activity only in the cytoplasm of cardiac and skeletal muscle. This enzyme catalyzes the reversible phosphorylation of creatine by ATP to creatine phosphate and ADP. Creatine phosphate is the major source of high-energy phosphate used in muscle contraction.

Principal Reason for Performing the Test
To identify injury to skeletal or cardiac muscle.

Most Common Abnormality Indicated by the Test
Skeletal muscle lesions attributable to trauma or vigorous exercise.

Sample Type and Precautions
Samples must be processed and centrifuged immediately after drawing blood. Blood samples should be taken within six hours of a suspect lesion. It is important to determine that the patient has not been exercised vigorously during the 12 hours prior to sampling. This may cause marked increases in creatine kinase activity. Remove plasma or serum from the cells or clot. If plasma is being collected, use only lithium heparinized samples. EDTA and fluoride/oxalate will reduce CK results.

Complementary Tests
Creatine kinase determination provides a specific, sensitive indication of muscle cell damage. Aspartate aminotransferase and lactate dehydrogenase activities may also be measured but are less specific and show smaller corresponding increases when muscle damage is present.
Creatinine (CREA)
Creatinine is a degradation product of creatine in muscle metabolism. The daily production of creatinine is fairly constant and not influenced markedly by age, diet, exercise, or catabolism. Creatinine is eliminated from the body by glomerular filtration and tubular secretion in the kidneys.

Principal Reasons for Performing the Test
As an indicator of renal disease and/or an index of glomerular filtration rate.

Most Common Abnormality Indicated by the Test
Increased creatinine—prerenal, postrenal, and renal azotemia.

Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples.

Interfering substances, such as creatine, in a sample can affect the analyzer’s ability to accurately provide creatinine results. When the analyzer detects such an interfering substance, dilution of the sample may be required to obtain an accurate creatinine value.

Complementary Tests
A complete urinalysis with a refractometry specific gravity measurement is essential for proper interpretation of increases in creatinine. Creatinine determinations should usually be performed in conjunction with measurements of BUN, inorganic phosphate, total protein, and albumin. A complete blood count (CBC) can sometimes demonstrate changes such as nonregenerative anemia with chronic renal failure.

Reaction Sequence
\[
\text{Reaction Sequence} \\
creatine + H_2O \xrightarrow{\text{amidohydrolase}} \text{creatinine} \\
\text{creatinine} + H_2O \xrightarrow{\text{amidohydrolase}} \text{creatinine} + \text{sarcosine} + \text{urea} \\
\text{sarcosine} + O_2 + H_2O \xrightarrow{\text{sarcosine oxidase}} \text{glycine} + \text{formaldehyde} + H_2O_2 \\
H_2O_2 + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye} + 2H_2O
\]
C-Reactive Protein (CRP)
C-reactive protein (CRP) is the major acute phase protein released by the liver in response to systemic inflammation in selected species including the dog. The Catalyst CRP Test is a sandwich immunoassay using monoclonal antibodies conjugated to gold nanoparticles and latex particles for the measurement of CRP.

Principal Reason for Performing the Test
CRP is a highly sensitive biomarker of active systemic inflammation in the canine patient. CRP will help the veterinarian detect active inflammation early, characterize the severity of the inflammatory response, and closely monitor the resolution or progression of the inflammatory process following therapeutic intervention.

Most Common Abnormality Indicated by the Test
CRP will be significantly elevated in any condition where active, systemic inflammation is present. The increase in CRP correlates with the severity of the inflammation. An elevated CRP value may be seen with infectious and noninfectious inflammatory disease (i.e., pneumonia, pancreatitis, pyelonephritis, pyometra, septicemia, and pyothorax), immune-mediated disease (i.e., immune-mediated hemolytic anemia and polyarthritis), as well as inflammation associated with tissue injury as seen in major surgery.

Sample Type and Precautions
Samples acceptable for CRP measurement include canine serum, plasma, and whole blood (when using the Catalyst Lithium Heparin Whole Blood Separator). Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium-heparinized samples.

When testing CRP on patients with suspected severe systemic inflammation, manual dilutions of the sample may be performed to avoid repeat testing when CRP values are above 10.0 mg/dL (100.0 mg/L). The recommended dilution is one part serum or plasma in one part normal saline (0.9% saline). IDEXX recommends only diluting tests with results outside of the reportable range. Diluting tests with results in the normal range may produce invalid results. Automated dilutions are not available for CRP on the Catalyst Dx.

Note: Whole blood samples processed in the whole blood separator should not be diluted.

Please note that only one test that requires a reagent can be processed during each run (i.e., CRP, Total T4, PHBR).

Complementary Tests
CRP should be evaluated in conjunction with a comprehensive history, physical examination, complete blood count, complete biochemical profile, and urinalysis to provide a comprehensive database when suspecting systemic inflammation. If infection is suspected, detecting of the pathogen is needed to make a final diagnosis.

Please note that for runs with greater than 18 slides, CRP must be loaded within the first 18 slides.

Fructosamine (FRU)
Fructosamine is glycated albumin or other proteins. Its concentration is related to blood glucose concentration during the preceding 2 to 3 weeks.

Principal Reason for Performing the Test
Measurement of fructosamine concentration as part of the routine evaluation of a diabetic patient undergoing therapy. It provides information about the status of glycemic control during the 2–3 weeks prior to evaluation. In cats, fructosamine concentrations can be measured to identify if a stress response or diabetes mellitus is the reason for high blood glucose concentrations.
In addition, during management of diabetes in both canine and feline patients, fructosamine concentration is used to clarify discrepancies between the history and physical examination findings and serial blood glucose concentration measurements. It is also used to assess the effectiveness of therapy.

**Most Common Abnormality Indicated by the Test**
Increased fructosamine indicates lack of or inadequate glucose regulation due to diabetes mellitus. Fructosamine concentrations increase with poor glycemic control and decrease when glycemic control improves. Less common, a low fructosamine may indicate prolonged hypoglycemia.

**Sample Type and Precautions**
**IMPORTANT:** FRU is currently available in two different formulations and their supported sample types vary:

<table>
<thead>
<tr>
<th>Part Number</th>
<th>Supported Sample Type(s)</th>
</tr>
</thead>
<tbody>
<tr>
<td>99-0003341</td>
<td>Serum or lithium heparin-treated plasma (you can use the plasma generated from a Catalyst* Lithium Heparin Whole Blood Separator)</td>
</tr>
</tbody>
</table>

**Note:** Please refer to part number and labeling on the slide box for supported sample type information.

It is important to separate the sample from the red blood cells as promptly as possible.

Serum is preferred for fructosamine testing as customer experience shows that it more consistently provides good quality samples.

Examine the serum or plasma for hemolysis. Although IDEXX dry-slide technology dramatically reduces the effect of this interfering substance, marked hemolysis can result in inaccurate fructosamine results. Typically, marked hemolysis will lower the reported value on the Catalyst analyzers.

**Complementary Tests**
The fructosamine test should be interpreted in conjunction with a blood glucose curve as well as the history and physical examination findings. A concurrent urinalysis should also be performed to evaluate for the presence of glucose and ketone. A urine culture is recommended in newly diagnosed diabetics and in animals with poorly controlled diabetes. In addition, a complete blood count and chemistry panel may be indicated to assess overall health of patient, to assess for secondary effects of poorly controlled diabetes, or for evidence of insulin antagonist disease. Further testing should be performed as indicated.

**Reaction Sequence**
Fructosamine + NBT $\xrightarrow{\text{OH}^-}$ formazan dye (measured at 560 nm)

**Gamma-glutamyltransferase (GGT)**
The enzyme gamma-glutamyltransferase is membrane-bound. It is present in large quantities in the kidney medulla and cortex and to a lesser extent in the small intestinal mucosa and bile ductular epithelium.

Despite the high activity of gamma-glutamyltransferase in the kidney, renal disease does not result in high enzyme activity in the serum sample. GGT in the kidney is primarily related to tubular lining epithelial cells and the enzyme is localized to the apical portion of the cell. Pathologic changes in these tubular epithelial cells result in loss of GGT directly into the urine. Measurement of GGT in the urine can prove to be a sensitive indicator of tubular epithelial cell injury/nephrotoxicity.
**Principal Reason for Performing the Test**
As an indicator of cholestasis or gallbladder disease.

**Most Common Abnormality Indicated by the Test**
Increased GGT—cholestasis.

**Sample Type and Precautions**
Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Hemolyzed specimens should not be used. Do not use fluoride/oxalate as an anticoagulant.

**Complementary Tests**
Serum gamma-glutamyltransferase activity is usually determined in conjunction with other tests of hepatic function or damage.

**Reaction Sequence**

\[
\text{L-γ-glutamyl-p-nitroanilide} + \text{glycylglycine} & \xrightarrow{\text{GGT}} \text{p-nitroaniline} + \text{γ-glutamyl glycylglycine}
\]

**Glucose (GLU)**
Glucose is the principal source of energy in monogastric mammals. The circulating concentration in the healthy animal is maintained within narrow limits.

**Principal Reason for Performing the Test**
To investigate carbohydrate metabolism.

**Most Common Abnormality Indicated by the Test**
Increased glucose—diabetes mellitus; glucocorticoid influence; epinephrine influence.

**Sample Type and Precautions**
For glucose determinations, the animal should have been fasted for 5–8 hours before sampling. Hemolysis may affect glucose results.

*For plasma samples*: Use only lithium heparinized samples. When blood is collected in lithium heparin, it is important that the sample be centrifuged immediately after collection. In this anticoagulant, glycolysis occurs quite rapidly in the presence of red blood cells and the glucose concentration in the sample can diminish at up to 10% an hour at room temperature. Remove plasma promptly from the red blood cells. Hemolyzed specimens should not be used.

*For serum samples*: Do not centrifuge serum samples until clotting is complete. Samples must be centrifuged completely. Remove serum promptly from the clot to avoid metabolism of glucose by the cells. A maximum of 30 minutes between drawing and separation from the clot is recommended. Hemolyzed specimens should not be used.

**Complementary Tests**
When the patient is a diagnosed diabetic, glucose tests may be performed in isolation. It is, however, useful to perform other tests for renal and hepatic function and lipid metabolism to monitor secondary effects of poorly controlled diabetes. Because stress in companion animals, particularly cats, can significantly raise glucose above the reference range, a fructosamine level should be considered in suspected cases of diabetes mellitus. A concurrent urinalysis should also be performed to evaluate for the presence of glucose and ketones.
Reaction Sequence

\[ \beta-D\text{-}glucose + O_2 + H_2O \xrightarrow{\text{glucose oxidase}} D\text{-}gluconic acid + H_2O_2 \]

\[ 2H_2O_2 + 4\text{-}aminoantipyrine + 1,7\text{-}dihydroxynaphthalene \xrightarrow{\text{peroxidase}} \text{red dye} \]

**Inorganic Phosphate (PHOS)**
Phosphorus plays a major role as a metabolic intermediate and is a constituent of nucleic acids, phospholipids, and nucleotides. Phosphates are also important components of buffering systems within the body fluids. Phosphate and calcium are absorbed in the small intestine. Absorption is influenced by the presence of other minerals, nutrients, vitamins, and intestinal pH. Calcium and phosphorous metabolism are interdependent.

**Principal Reason for Performing the Test**
As a measure of glomerular filtration rate.

**Most Common Abnormality Indicated by the Test**
Increased inorganic phosphate—decreased glomerular filtration.

**Sample Type and Precautions**
Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Do not use oxalate, fluoride, citrate, or EDTA as anticoagulants. Blood samples must be processed and centrifuged as soon as possible after collection as phosphates are released quickly from the red blood cells. Hemolysis can result in marked increases in phosphate concentration.

**Complementary Tests**
Inorganic phosphate determination should be performed in conjunction with measurements of calcium, albumin, total protein, and glucose. If renal disease is suspected, BUN, creatinine, albumin, total protein, and a complete urinalysis should also be determined.

**Reaction Sequence**
inorganic phosphate + ammonium molybdate \( \xrightarrow{\text{pH 4.2}} \) ammonium phosphomolybdate complex

ammonium phosphomolybdate complex \( \xrightarrow{\text{p-methy lamino phenol sulfate}} \) heteropolymolybdate blue

**Lactate Dehydrogenase (LDH)**
The enzyme lactate dehydrogenase is present in large amounts in all organs and tissues (including red blood cells) of most animals. It is found in the cell cytoplasm and is released into the blood during reversible and irreversible (necrosis) cell injury. The test is not a specific or sensitive indicator of damage to any organ or tissue.

**Note:** The normal range of lactate dehydrogenase in the dog and cat is wide, as can be the intra-animal variation from day to day. Consequently, small increases in activity due to minimal organ damage are difficult to identify. The measurement of lactate dehydrogenase is a somewhat traditional test whose diagnostic value is limited in practice.
**Principal Reason for Performing the Test**  
To investigate damage to liver, cardiac or skeletal muscle.

**Most Common Abnormality Indicated by the Test**  
Increased activity is usually associated with hepatic parenchymal lesions.

**Sample Type and Precautions**  
Remove plasma or serum promptly from the cells or clot and analyze as soon as possible. If plasma is being collected, use only lithium heparinized samples. Fluoride/oxalate and EDTA should not be used as anticoagulants.

Hemolyzed specimens should not be used because LDH contamination from red blood cells will occur.

**Complementary Tests**  
Lactate dehydrogenase activity is usually determined in conjunction with other tests of liver, cardiac, or skeletal muscle function or damage.

**Reaction Sequence**

\[
\text{pyruvate} + \text{NADH} + \text{H}^+ \xrightarrow{\text{LDH}} \text{lactate} + \text{NAD}^+
\]

**Lactate (LAC)**  
Lactate is produced by anaerobic metabolism of glucose and its concentration depends on relative rates of production in muscle cells and erythrocytes and metabolism in the liver.

**Principal Reason for Performing the Test**  
Elevated lactate levels usually are caused by overproduction or under metabolism. They result from tissue hypoxia, diabetes mellitus, malignancies, ethanol or methanol ingestion, and metabolic acidosis.

**Most Common Abnormality Indicated by the Test**  
Hypoxia secondary to severe exercise, shock, hypovolemia, cardiac disease, pulmonary edema, and seizures.

**Sample Type and Precautions**  
Use lithium heparinized or Fl/oxalated samples. When using lithium heparinized samples, separate the plasma from the red cells within 5 minutes of collection.

**Complementary Tests**  
CBC, biochemical panel, complete urinalysis, and blood gas.

**Reaction Sequence**

\[
\text{L- (+)-lactic acid} + \text{O}_2 \xrightarrow{\text{lactate oxidase}} \text{pyruvate} + \text{H}_2\text{O}_2
\]

\[
2\text{H}_2\text{O}_2 + 4\text{-aminoantipyrine} + 1,7\text{-dihydroxynaphthalene} \xrightarrow{\text{peroxidase}} \text{red dye}
\]
Lipase (LIPA)

Lipase is secreted by the pancreas and to a lesser extent by the gastrointestinal mucosa. Lipase is a relatively sensitive indicator of pancreatic pathology (as compared to amylase). Generally a greater than threefold increase above the reference range is supportive of pancreatitis.

Principal Reason for Performing the Test
As an indicator of acute pancreatitis.

Most Common Abnormality Indicated by the Test
Acute pancreatitis.

Sample Type and Precautions
Blood samples should be taken within one day of the onset of symptoms suggesting acute pancreatitis. Promptly remove plasma or serum from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Do not use oxalate/fluoride, citrate, or EDTA anticoagulants. Lipemia and icterus may increase lipase results.

Complementary Tests
Lipase and amylase are usually determined in conjunction with tests of hepatic and pancreatic function or damage. Canine and feline pancreas-specific lipase tests should be performed in questionable cases.

Reaction Sequence

1-oleoyl-2,3-diacetylglcerol $\xrightarrow{\text{lipase, colipase pH 8.5}}$ 2,3-diacetylglcerol + oleic acid

2,3-diacetylglcerol $\xrightarrow{\text{diacetylase}}$ glycerol + acetic acid

glycerol + ATP $\xrightarrow{\text{glycerol kinase MgCl}_2}$ L-α-glyceroephosphate + ADP

L-α-glyceroephosphate + O$_2$ $\xrightarrow{\text{L-α-glyceroephosphate oxidase}}$ dihydroxyacetone phosphate + H$_2$O$_2$

H$_2$O$_2$ + leuco dye $\xrightarrow{\text{peroxidase}}$ dye + 2H$_2$O

Magnesium (Mg)

Magnesium plays an important intracellular role in the activation of enzymes including those responsible for many anabolic and catabolic processes. It is also involved in the formation and destruction of acetylcholine, which governs the transmission of electrical impulses at the neuromuscular junction. The adrenal, thyroid, and parathyroid glands appear to regulate serum magnesium concentration.

Principal Reason for Performing the Test
The importance of measuring serum magnesium concentration in dogs and cats has not been fully investigated. However, there have been reports of hypomagnesemia in dogs following the removal of the parathyroid gland.
Most Common Abnormalities Indicated by the Test
Increased magnesium—decreased glomerular filtration.
Decreased magnesium—parathyroid gland removal.

Sample Type and Precautions
Blood samples should be centrifuged immediately after collection as magnesium is released from hemolyzed erythrocytes and can give erroneously high magnesium results. Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Do not use oxalate/citrate or EDTA as anticoagulants. Blood collection tubes preserved with sodium fluoride cause lower results.

Complementary Tests
See tests listed under Endocrine Profile in the "Profile Selection" table on page 70.

Reaction Sequence

\[
\text{Mg}^{2+} + \text{Ca}^{2+} \xrightarrow{\text{chelator}} \text{Mg}^{2+} + \text{Ca}^{2+} - \text{chelator complex}
\]

\[
\text{Mg}^{2+} + \text{formazan dye derivative} \xrightarrow{\text{pH 9.75}} \text{Mg}^{2+} - \text{dye complex}
\]

Phenobarbital (PHBR)
Phenobarbital is a commonly used drug used to treat seizures in a variety of species. Phenobarbital levels should be evaluated during initial dosing and throughout treatment to ensure that the blood levels are within the targeted therapeutic range.

Principal Reasons for Performing the Test
Phenobarbital is a controlled barbiturate medication that is used to treat veterinary patients that have seizures. The dosage of phenobarbital needs to remain within a specific range to be effective. If the level is <10 µg/mL, there may not be a sufficient level of phenobarbital to prevent seizures. If the level >30 µg/mL in cats or >40 µg/mL in dogs, phenobarbital can be toxic and potentially life threatening.

In most patients, steady state is achieved after two to three weeks of consistent dosing with phenobarbital. Once steady state is achieved, timing of sample collection is not important in more than 90% of patients. However, there can be variability of the phenobarbital half-life in a small percentage of patients. Therefore, if toxicity is suspected, a peak sample (4–5 hours post-pill) may be helpful, and if breakthrough seizures are occurring and inadequate dosing is suspected, a trough level (collected immediately prior to the next dose) may be helpful.

Most Common Abnormalities Indicated by the Test
Over or under dosage of medication.

Sample Type and Precautions
Do not use separator tubes as contact with the gel may decrease levels.

Complementary Tests
CBC, full chemistry panel, urinalysis, bile acids (minimally 2 times per year).
Reaction Sequence

Potassium (K)
Potassium is the major cation of intracellular fluid, where it is the major buffer within the cell, facilitates nerve conduction and muscle function, and helps maintain osmotic pressure. Abnormally high or low potassium levels cause changes in muscle irritability, respiration, and myocardial function.

Principal Reasons for Performing the Test
High potassium (hyperkalemia) is usually found in urinary obstruction, renal failure, metabolic or respiratory acidosis, and hypoadrenocorticism as well as excessive hemolysis for horses, cattle, cats, and some breeds of dogs. Decreased values (hypokalemia) usually follow excessive salt loss through severe vomiting or diarrhea, inadequate intake, anorexia (especially cats), malabsorption, and severe burns.

Most Common Abnormalities Indicated by the Test
Hyperkalemia—renal failure, postrenal obstruction.
Hypokalemia—excessive loss of potassium.

Sample Type and Precautions
Remove plasma or serum promptly from cells or clot. If plasma is being collected, use only lithium heparinized samples. Avoid hemolysis.

Do not freeze samples for use with the Catalyst Dx analyzer.

Complementary Tests
Sodium, potassium, and chloride should always be assayed together to determine electrolyte balance. The additional measurement of bicarbonate will allow accurate assessment of metabolic acid-base physiology.
ACTH stimulation test for suspect cases of hypoadrenocorticism.

Reaction Sequence
Potassium + ionophore — fluorescent dye — fluorescence change

Sodium (Na)
Sodium is the major cation of extracellular fluid, where it maintains osmotic pressure, acid-base balance, and transmits nerve impulses. The body maintains total sodium content, and only slight changes are found even under pathologic conditions.
**Principal Reasons for Performing the Test**
To evaluate electrolyte status in conjunction with potassium and chloride levels.

Low sodium (hyponatremia) is usually caused by a relative excess of body water. Reduced levels may be due to low intake, loss through vomiting or diarrhea plus adequate water and inadequate salt replacement, salt-losing nephropathy, osmotic diuresis, metabolic acidosis, and various glandular conditions.

Elevated values (hypernatremia) usually follow water loss in excess of salt loss through profuse sweating, severe vomiting or diarrhea, inadequate water intake, and dehydration of renal sodium conservation in hyperaldosteronism.

**Most Common Abnormality Indicated by the Test**
Hypernatremia secondary to dehydration, gastrointestinal fluid loss (vomiting or diarrhea).

**Sample Type and Precautions**
Remove plasma or serum promptly from cells or clot. If plasma is collected, use only lithium heparinized samples. Avoid hemolysis.

Do not freeze samples for use with the Catalyst Dx analyzer.

**Complementary Tests**
Sodium, potassium, and chloride should always be assayed together to determine electrolyte balance. The additional measurement of bicarbonate will allow accurate assessment of metabolic acid-base physiology.

**Reaction Sequence**
Sodium + ionophore $\rightarrow$ fluorescent dye $\rightarrow$ fluorescence change

**Total Bilirubin (TBIL)**
Hemoglobin from degenerated erythrocytes is converted to bilirubin in the monocyte-macrophage system. Free unconjugated bilirubin is transported to the liver bound to albumin, where it is conjugated with glucuronic acid and eliminated in the bile. In obstructive liver disease, the concentration of conjugated bilirubin in the blood increases.

During intravascular or extravascular hemolysis, very large numbers of erythrocytes may be destroyed quickly and the conjugation mechanism in the liver may become overloaded so that high concentrations of unconjugated bilirubin are found in the blood. If the loss of hemoglobin and erythrocytes is very large, anoxia may occur. Hepatocyte dysfunction follows leading to cellular swelling, which occludes the bile canaliculi preventing the elimination of conjugated bilirubin. A concomitant rise in circulating conjugated bilirubin then occurs.

**Principal Reason for Performing the Test**
To detect hepatobiliary disease and excessive erythrocyte destruction.

**Note:** In healthy dogs and cats, the concentration of total bilirubin in the serum is very low. Visual inspection of the sample will frequently indicate whether bilirubin determination is necessary (serum and plasma only).

**Most Common Abnormality Indicated by the Test**
Increased bilirubin—cholestatic liver disease (conjugated bilirubin) and hepatic insufficiency (unconjugated bilirubin), hemolytic disease (unconjugated and possible conjugated bilirubin), and intrahepatic obstruction.
Sample Type and Precautions
Remove plasma or serum promptly from cells or clot. Samples should be analyzed immediately as bilirubin degrades rapidly in light. If immediate analysis is impossible, the sample must be kept in the dark and preferably at 4°C–8°C (36°F–40°F) in a refrigerator. Sample must be allowed to come to room temperature before analysis. If plasma is collected, use only lithium heparinized samples.

It is critical that samples be properly centrifuged. Otherwise, leukocytes and platelets may remain in suspension, even when red blood cells have been separated. Cellular material on the slide may cause significant positive error. Also, hemoglobin increases total bilirubin results, so avoid even moderately hemolyzed samples.

Complementary Tests
Total bilirubin should be determined with other tests of hepatic function or damage. Hematocrit should also be performed to eliminate or confirm the presence of hemolytic disease. Determination of urinary urobilinogen and bilirubin may also be useful.

Reaction Sequence

Total Protein (TP)
The serum total protein concentration comprises all the proteins found in the aqueous phase of the blood. In healthy animals, albumin is the major single component. The remaining proteins are the alpha, beta, and gamma globulins. The globulin concentration is determined by subtracting the albumin from the total protein.

Principal Reason for Performing the Test
Total protein measurement may provide useful information when used in combination with tests to investigate hepatic and renal function, the degree of hydration, protein-losing enteropathies, or gammopathies. The test is nonspecific and, if performed in isolation, will be unlikely to provide diagnostic information.

Most Common Abnormalities Indicated by the Test
Increased total protein—dehydration, inflammatory disease.
Decreased total protein—loss of proteins through blood loss and gastrointestinal loss, decreased albumin associated with protein-losing nephropathy and enteropathy, and decreased albumin associated with hepatic insufficiency and inflammatory disease.
Impaired renal and hepatic function, dehydration, and gastrointestinal lesions.

Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. If plasma is collected, use only lithium heparinized samples. Moderate-to-marked hemolysis can result in false high total protein concentration.

Results obtained from the analysis of plasma may be slightly higher than serum due to the fibrinogen that remains in the plasma.

Complementary Tests
Total protein concentration is usually determined in conjunction with the measurement of albumin and other tests of renal and hepatic function.
Reaction Sequence

protein + copper tartrate $\xrightarrow{\text{LiOH}}$ colored complex

**Total T₄ (TT₄)**
An enzyme-linked immunosorbent assay (ELISA) for the quantitative measurement of total T₄ (thyroxine) in canine and feline patients. With a total T₄ test, you can assess thyroid function, provide comprehensive one-visit screening for feline hyperthyroidism, presumptive canine hypothyroidism, as well as monitor response to treatment and adjust dosages immediately.

**Principal Reason for Performing the Test**
To screen, diagnose, and monitor thyroid disease. The measurement of total thyroxine helps veterinary practitioners to assess thyroidal function by measuring the bound and unbound thyroxine in the blood. Thyroxine is the principal hormone secreted by the thyroid gland and is critical to metabolic processes.

**Most Common Abnormality Indicated by the Test**

- **Hyperthyroidism**—an elevated TT₄ is consistent with hyperthyroidism. Naturally occurring hyperthyroidism is a common endocrine disorder in cats and rare in dogs.

- **Hypothyroidism**—a decreased TT₄ is consistent with but not necessarily definitively diagnostic of hypothyroidism. Naturally occurring hypothyroidism is a common endocrine disorder in dogs and rare in cats.

- **Nonthyroidal illness (NTI)**—nonthyroidal illness can affect TT₄ levels (and potentially other thyroid tests as well). Nonthyroidal illness can lower TT₄ levels, potentially into the hypothyroid range. The more severe the nonthyroidal illness, the greater the potential impact on TT₄ levels.

**Sample Type and Precautions**
For use with serum, plasma, and whole blood (when using the Catalyst Lithium Heparin Whole Blood Separator).

Remove plasma or serum promptly from the cells or clot. If plasma is being collected, use only lithium heparinized samples. Hemolyzed samples should not be used. Do not use fluoride/oxalate as an anticoagulant.

**Complementary Tests**
Total T₄ should be evaluated in conjunction with a comprehensive history, physical examination, CBC, complete biochemical profile, and urinalysis to provide a comprehensive database of information in the diagnosis or suspicion of thyroid disease.

In dogs with low or low normal T₄ results and with consistent clinical signs, evaluate free T₄ (fT₄) and endogenous thyroid-stimulating hormone (TSH) and possibly thyroglobulin autoantibodies (TgAA) to aid in confirming hypothyroidism.

Cats with consistent clinical signs and total T₄ (TT₄) values in the borderline high range (gray zone) may have early hyperthyroidism or a concurrent nonthyroidal illness (NTI). In these cases, consider a free T₄ (fT₄), a T₃ suppression test, or radionuclide thyroid imaging to aid in confirming the diagnosis.
Triglycerides (TRIG)
Triglycerides are usually present in the diet of dogs and cats, especially when the animals are fed table scraps. They are also synthesized in the liver, mainly from carbohydrates, to provide a secondary energy source and are stored in fatty tissue. Their hydrolysis to mono- and diglyceride glycerol and free fatty acids is catalyzed by pancreatic lipase.

Principal Reason for Performing the Test
To detect abnormalities in lipid metabolism.

Most Common Abnormality Indicated by the Test
Increased triglycerides—High-fat diet or abnormalities in fat metabolism.

Sample Type and Precautions
Blood should not be drawn within 12 hours of a meal.
Remove plasma or serum promptly from the cells or clot. If plasma is collected, use only lithium heparinized samples. Grossly lipemic specimens probably have very high triglycerides and should be diluted before analysis.

Complementary Tests
Triglycerides should not be measured in isolation. If the sample is turbid or milky, the test should be determined in conjunction with measurements of cholesterol and glucose, and hepatic and renal function tests. Also consider repeat sampling if the patient has not been fasted for 12 hours.

Reaction Sequence
\[
\text{lipoproteins} \xrightarrow{\text{surfactant}} \text{triglycerides} + \text{proteins} \\
\text{triglycerides} + \text{H}_2\text{O} \xrightarrow{\text{lipase}} \text{glycerol} + \text{fatty acids} \\
\text{glycerol} + \text{ATP} \xrightarrow{\text{glycerol kinase}} \text{L-}\alpha\text{-glycerophosphate} + \text{ADP} \\
\text{L-}\alpha\text{-glycerophosphate} + \text{O}_2 \xrightarrow{\text{L-}\alpha\text{-glycerol-phosphate oxidase}} \text{dihydroxyacetone phosphate} + \text{H}_2\text{O}_2 \\
\text{H}_2\text{O}_2 + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye} + 2\text{H}_2\text{O}
\]

Uric Acid (URIC)
Uric acid determinations are useful in avian patients and Dalmations dogs in place of urea determinations. In all dogs (except Dalmations) with diffuse hepatic disease, there is marked elevation of blood uric acid above the normal levels of <1 mg/dL.

Principal Reason for Performing the Test
As an indicator of the severity of renal disease in avian populations (and Dalmations).

Most Common Abnormality Indicated by the Test
Increased uric acid—prerenal, postrenal and renal azotemia associated with decreased glomerular filtration rate.
Sample Type and Precautions
Remove plasma or serum promptly from the cells or clot. If plasma is collected, use only lithium heparinized samples. Plasma collected from sodium fluoride, citrate, or EDTA preservative should not be used.

Complementary Tests
Creatinine, UCRE/CREA, UPRO.

Reaction Sequence

\[
2\text{H}_2\text{O} + \text{uric acid} \xrightarrow{\text{uricase}} \text{allantoin} + \text{H}_2\text{O} + \text{CO}_2
\]

\[
\text{H}_2\text{O}_2 + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye}
\]

Urine Creatinine (UCRE)
Urine creatinine is determined so that the concentration of electrolytes filtered or lost through the glomeruli or renal tubules, such as urinary protein or cortisol, can be quantitated, compared, and expressed as ratios with diagnostic significance.

Principal Reason for Performing the Test
To be performed with urine protein in order to determine the urine protein:creatinine ratio (UPC).

Most Common Abnormality Indicated by the Test
Proteinuria indicating early renal disease, protein-losing nephropathy.

Sample Type and Precautions
Urine, preferably collected through cystocentesis, collected in a clean container. An inactive urinary sediment should be demonstrated and urinary tract infection (UTI) via culture and sensitivity should be ruled out before performing, as UTI may mildly to moderately raise the UPC.

Complementary Tests
Complete urinalysis with culture and sensitivity. Serum chemistries such as creatinine, BUN, albumin, and globulin; CBC; and SNAP* 4Dx* Test.

Storage Information
Urine samples should be run within 2 hours of collection and can be stored in a refrigerator for up to 24 hours. DO NOT freeze urine samples.

Reaction Sequence

\[
\text{creatinine} + \text{H}_2\text{O} \xrightarrow{\text{creatinine amidohydrolase}} \text{creatine}
\]

\[
\text{creatinine} + \text{H}_2\text{O} \xrightarrow{\text{creatinine amidohydrolase}} \text{sarcosine} + \text{urea}
\]

\[
\text{sarcosine} + \text{O}_2 + \text{H}_2\text{O} \xrightarrow{\text{sarcosine oxidase}} \text{glycine} + \text{formaldehyde} + \text{H}_2\text{O}_2
\]

\[
\text{H}_2\text{O}_2 + \text{leuco dye} \xrightarrow{\text{peroxidase}} \text{dye} + 2\text{H}_2\text{O}
\]
Urine Protein (UPRO)
Urinary protein is determined and compared to the concentration of creatinine in order to assess the level of renal protein (glomeruli and tubular) loss to determine the urine protein:creatinine (UPC) ratio.

Principal Reason for Performing the Test
To be performed with urine creatinine in order to determine the urine protein:creatinine (UPC) ratio.

Most Common Abnormality Indicated by the Test
Proteinuria indicating early renal failure, protein-losing nephropathy.

Sample Type and Precautions
Urine, preferably collected through cystocentesis, collected in a clean container. An inactive urinary sediment should be demonstrated and urinary tract infection (UTI) via culture and sensitivity should be ruled out before performing as UTI may mildly to moderately raise the UPC.

Complementary Tests
Complete urinalysis with culture and sensitivity. Serum chemistries such as creatinine, BUN, albumin, and globulin.

CBC
SNAP 4Dx Test

Storage Information
Urine samples should be run within 2 hours of collection and can be stored in a refrigerator for up to 24 hours. DO NOT freeze urine samples.

Reaction Sequence

Medical Protocol Descriptions

Ammonia Protocol
Baseline ammonia levels should be assessed in animals with signs of hepatic encephalopathy or in patients suspected of having portosystemic shunts (PSS). Ammonia tolerance tests may be considered to evaluate for PSS where bile acids are not considered (for example, in Maltese dogs).

Ammonia tolerance test: A baseline sample is drawn after the patient has been fasted for 12 hours. Ammonium chloride (0.1 g/kg) by mouth via stomach tube or gelatin capsules. A second sample is drawn 30 minutes after ammonium chloride administration.

Note: Vomiting during the procedure will invalidate results.

Sample Requirements: 1 mL heparinized plasma, separated from RBCs. Do not use serum.

Storage/Stability: 48 hours—store plasma frozen

Interferences: Hemolysis, glucose levels over 600 mg/dL (33.33 mmol/L), high BUN values

Comments: Anticoagulated blood must be centrifuged immediately after collection. Separate plasma and place it in a glass container (RTT). Freeze immediately and keep frozen if not running sample immediately.

Note: Ammonia levels increase with time.
UPC Protocol

**Principle Reason for Performing Test:** To aid in the diagnosis of protein-losing nephropathies such as glomerulonephritis and amyloidosis and as an early marker of chronic renal failure.

**Includes:** Urine protein (UPRO), urine creatinine (UCRE), protein:creatinine (UPC) ratio

**Sample Requirements:** 2 mL urine in a sterile container

**Storage/Stability:** 48 hours at 2°C–8°C (36°F–46°F)

**Interferences:** Gross hematuria, pyuria.

**Complementary Tests:** Complete urinalysis with culture and sensitivity. Serum chemistries such as creatinine, BUN, albumin, globulin; CBC; SNAP* 4Dx* Test; and imaging studies.

**Interpretation:** Proteinuria requires proof of persistence and localization to prerenal, renal, or postrenal origins. Prove persistence of proteinuria by repeating the UPC ratio at least three times, a minimum of two weeks apart.

- Prerenal proteinuria is possible when a CBC and a biochemical profile detect hemolysis, hyperglobulinemia or evidence of muscle damage. Recommend investigation and management for the underlying cause.

- Postrenal proteinuria is caused by urogenital tract diseases, hematuria, or pyuria. Repeat the test with a cystocentesis sample or evaluate urine sediment for hemorrhage or inflammation. Consider a urine culture. Recommend investigation and management for the underlying cause.

- Renal proteinuria: evaluate in the face of azotemia.

**Nonazotemic, persistent, renal proteinuria (dogs and cats):**

- UPC <0.5 = within reference range
- UPC 0.5–1.0 = questionable, repeat at appropriate range
- UPC 1.0–2.0 = excessive proteinuria; recommend investigation for underlying systemic diseases
- UPC ≥2.0 = excessive proteinuria; recommend investigation for underlying systemic diseases and medical management

**Azotemic, persistent, renal proteinuria (dogs):**

- UPC <0.5 = warrant monitoring and investigation
- UPC ≥0.5 = excessive proteinuria; recommend investigation for underlying systemic diseases and medical management

**Azotemic, persistent, renal proteinuria (cats):**

- UPC <0.4 = warrant monitoring and investigation
- UPC ≥0.4 = excessive proteinuria; recommend investigation for underlying systemic diseases and medical management
History and physical exam

CBC, biochemical profile, urinalysis (examine urine sediment and specific gravity)

- Active sediment
- Pollakiuria
- Azotemia

≠ Active sediment
± Pollakiuria
± Azotemia

+ Active sediment
+ Pollakiuria
- Azotemia

Prerenal
If prerenal ruled out
Renal
If postrenal ruled out
Postrenal

+ Hemoglobinuria
+ Myoglobinuria
+ Bence Jones proteinuria (myeloma)

IDEXX Urine P:C Ratio

+ Azotemia
- Azotemia

<0.5 canine <0.4 feline
Within reference range
Investigate azotemia

≥0.5 canine ≥ 0.4 feline
Within reference range
Monitor, investigate, treat

≥0.5 canine and feline
Within reference range
Repeat IDEXX UPC ratio in two weeks

<0.5 canine and feline
Monitor

≥0.5 to <1.0 canine and feline
Monitor
Investigate potential causes of immune-mediated glomerulonephritis or interstitial nephritis

≥1.0 to <2.0 canine and feline
Monitor, investigate

≥2.0 canine and feline
Monitor, investigate, treat

Work up for proteinuria

+ Urinary tract infection
+ Stones
+ Tumors
Total T₄ Protocols

**Canine hypothyroidism suspected**

**Common clinical signs in dogs**
- Obesity
- Skin disease
- Lethargy
- Mental dullness
- Exercise/Cold intolerance

**Initial database**
- Total T₄
- CBC
- Chemistry with electrolytes
- Complete urinalysis

**Low T₄ with NTI**
- <1.0 µg/dL
  - (<13.0 nmol/L)
  - Address NTI

**Low T₄**
- <1.0 µg/dL
  - (<13.0 nmol/L)
  - Hypothyroidism unlikely

**Low Normal T₄**
- 1.0–2.0 µg/dL
  - (13.0–26.0 nmol/L)

**Normal T₄**
- 2.0–4.0 µg/dL
  - (26.0–51.0 nmol/L)
  - Hypothyroidism unlikely

**Low T₄ ± high TSH ± positive TgAA**
- Hypothyroidism likely
  - Clinical trial

**Normal T₄ and TSH negative TgAA**
- Hypothyroidism unlikely
  - Repeat testing in 4–6 weeks if hypothyroidism still suspected

CBC = Complete blood count

*Note:* 1 µg/dL is equal to 12.87 nmol/L. A result that falls within the low normal range of the assay should be considered ambiguous.
Feline hyperthyroidism suspected

Common clinical signs in cats
- Weight loss
- Hyperactivity
- Polyphagia
- Palpable goiter
- Unkempt coat

Initial database
- Total T₄
- CBC
- Chemistry with electrolytes
- Complete urinalysis

Low T₄
<0.8 µg/dL
(<10.0 nmol/L)
- Euthyroid sick or iatrogenic

Normal T₄
0.8–4.7 µg/dL
(10.0–30.0 nmol/L)
- Normal T₄ (gray zone)
  2.3–4.7 µg/dL
  (30.0–60.0 nmol/L)
  - Low or normal fT₄
  - Hyperthyroidism unlikely†

High T₄
>4.7 µg/dL
(>60.0 nmol/L)
- High fT₄
- Hyperthyroidism likely

†If strong suspicion of hyperthyroidism still exists, consider retesting in 4–6 weeks or a technetium scan.

Note: 1 µg/dL is equal to 12.87 nmol/L. A result that falls within the gray zone of the assay should be considered ambiguous.

CBC = Complete blood count

Common clinical signs in cats
- Weight loss
- Hyperactivity
- Polyphagia
- Palpable goiter
- Unkempt coat

Initial database
- Total T₄
- CBC
- Chemistry with electrolytes
- Complete urinalysis

Low T₄
<0.8 µg/dL
(<10.0 nmol/L)
- Euthyroid sick or iatrogenic

Normal T₄
0.8–4.7 µg/dL
(10.0–30.0 nmol/L)
- Normal T₄ (gray zone)
  2.3–4.7 µg/dL
  (30.0–60.0 nmol/L)
  - Low or normal fT₄
  - Hyperthyroidism unlikely†

High T₄
>4.7 µg/dL
(>60.0 nmol/L)
- High fT₄
- Hyperthyroidism likely

†If strong suspicion of hyperthyroidism still exists, consider retesting in 4–6 weeks or a technetium scan.

Note: 1 µg/dL is equal to 12.87 nmol/L. A result that falls within the gray zone of the assay should be considered ambiguous.
### Profile Selection

Suggested CLIPs/profiles that will aid in the identification of abnormalities in tissues, organs, and metabolic systems in the most economical way are:

<table>
<thead>
<tr>
<th>Circumstances for the choice of the profile</th>
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</thead>
<tbody>
<tr>
<td>A diagnosis is not evident on clinical examination but one or more lesions are suspected.</td>
<td>ALB</td>
<td>ALKP</td>
<td>ALT</td>
<td>AMYL</td>
<td>AST</td>
<td>BUN/UREA</td>
<td>Ca²⁺</td>
<td>CHOL</td>
<td>CK</td>
<td>CREA</td>
<td>CRP</td>
<td>FRU</td>
<td>GGT</td>
<td>GLU</td>
<td>LDH</td>
<td>LIPA</td>
<td>Mg²⁺</td>
<td>NH₃</td>
<td>PHBR</td>
<td>PHOS</td>
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<tr>
<td>A smaller version of the General/Geriatric profile.</td>
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<tr>
<td>As a general precaution for any anesthetic candidate.</td>
<td>Chem 17 CLIP</td>
<td>Chem 15 CLIP</td>
<td>Chem 10 CLIP</td>
<td>Gastrointestinal profile</td>
<td>Cardiac profile</td>
<td>Endocrine profile</td>
<td>Hepatic profile</td>
<td>Lipid profile</td>
<td>Pancreatic profile</td>
<td>Renal profile</td>
<td>Young preanesthetic profile</td>
<td>Seizures</td>
<td>Diabetes</td>
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<td>Gastrointestinal diseases are suspected.</td>
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<td>Cardiac disease is suspected.</td>
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<td>Endocrine disease is suspected. Results may indicate that specific hormone tests are required.</td>
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<td>Hepatic damage is suspected.</td>
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<td>The investigation of fat metabolism in hypo-thyroidism, obesity, or lipemic samples.</td>
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<td>Acute pancreatitis is suspected.</td>
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<td>Renal disease is suspected.</td>
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</table>

### Notes:

1. **CK:** Samples should be taken within 6 hours of a suspected lesion.
2. **AMYL/LIPA:** Samples should be taken within 1 day of the onset of pancreatitis symptoms.
Technical Specifications

Dimensions
Width: 14 inches (35.56 cm)
Depth: 16.25 inches (41.28 cm)
Height: 17.25 inches (43.82 cm)
Weight: approximately 50 pounds (22 kg)

Power Supply
Input: 100–240 V AC, 50–60 Hz, 3.5 Amps
Power Supply Protection: IPX0

Input/Output Connections
There are four Input/Output connections for the Catalyst Dx analyzer. Three are on the rear of the instrument (power connection, Ethernet port for connection to IDEXX VetLab* Station, and a USB port) and one is accessible when the waste drawer is removed (USB port).

Operating Conditions

<table>
<thead>
<tr>
<th></th>
<th>Operating</th>
<th>Storage</th>
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</thead>
<tbody>
<tr>
<td>Temperature</td>
<td>15°C–30°C</td>
<td>5°C–38°C</td>
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<tr>
<td></td>
<td>(59°F–86°F)</td>
<td>(41°F–100°F)</td>
</tr>
<tr>
<td>Relative Humidity</td>
<td>15%–75%</td>
<td>20%–85%</td>
</tr>
</tbody>
</table>
IDEXX Technical Support Contact Information

IDEXX Sales Representative: ________________________________
Telephone/Voice Mail: ________________________________

United States: 1-800-248-2483
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Belgium: 32 (0)27 00 64 38
Brazil: 0800-777-7027
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Ireland: 353 (0)1 562 1211
Italy: 39 02 87 10 36 76
Japan: 0120-71-4921