Lonza

QC Testing Catalog 2016–2017





QC Insider™ e-Learning



Seminars and workshops are great ways to transfer knowledge but at the same time require significant dedication of time. Participants may also find that with pre-defined course content, their individual training needs are not fully met by these types of programs.

Our e-Learning Modules, accessible via our QC Insider™ Toolbox, are the solution. These modules are a suite of online, endotoxin-specific training courses available at your fingertips.

Benefit from:

- Learning at your own pace as your schedule permits
- No travel costs
- Creating customized training packages targeted to your training needs
- An integrated test and certification

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How to Use this Catalog

This section explains some tools and hints we have included in this catalog to help you find the information you need at a glance.

Icons

The icons below refer to additional product-related information, how to reach Customer Service or Scientific Support and additional purchase details.

- Great additions to our expanding products portfolio
- Additional product or technical information available online
- Contact our Customer Service or Scientific Support Representatives for additional information or ordering assistance
- Ordering information and instructions
- Product related scientific references
- Storage conditions

How to Order

There are several ways to order your products of choice with Lonza: via our online shop, e-mail, phone, fax or mail.

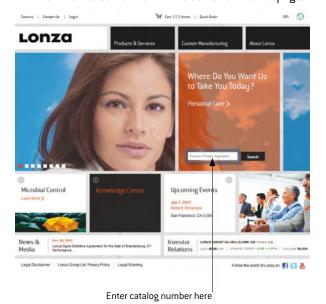
Online Ordering

Our website encompasses an extensive product portfolio. Along with the essential tools needed to manage your electronic purchases, you will find the process simple, secure and reliable. Our website offers:

- Highest security standards
- Ordering available 24 hours, 7 days a week
- Latest technical, application, and safety information for our products
- Custom pricing information (if you register with your customer number)
- Sales tax, shipping charges, and expected shipping dates displayed before order is placed
- Certificates of Analysis

To order, you need a valid credit card (MasterCard®, VISA® or American Express®) or purchase order, and a shipping address. Our system will always inform you of inventory availability as you add items to your shopping basket on the top center of the website.

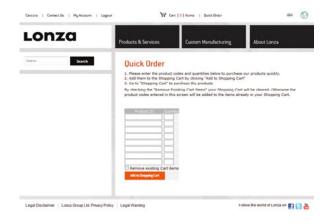
Go to: http://www.lonza.com to access our online shop. To find your product of interest, please enter the Catalog Number into the search field in the center of the webpage.



Ouick Order Form

It's easy and saves time.

If you already know the Catalog Numbers and quantities of the products you would like to purchase in our online shop, you may use the Quick Order Form (see screenshot) available at http://www.lonza.com/quickorder



Ordering via Ariba® or Other Third Party Providers

We support Ariba® and other third party purchasing systems through secure cXML communications. You have the choice of purchasing via a pricelist (CIF electronic catalog) or using a punch-out catalog system. We also accept and support EDI standards 850 (purchase order) and 810 (sales invoice), if your company conducts electronic transactions using EDI protocols.

Please contact us if you would like to purchase products using any of these types of electronic transmissions.

Direct Ordering via cXML

If you have developed your own internal purchasing system, you can link directly to our order entry systems via secure cXML communications protocols. Please contact us if you would like to discuss using this type of system to place your orders.

For prompt response to your e-commerce questions, please e-mail us at **ecommerce@lonza.com**.

How to Order

Continued

Ordering via E-mail, Phone, Fax or Mail

Please include all of the following information on all of your orders:

- Purchase order number
- Shipping address and billing address
- Contact name and telephone number
- Name and department of end user
- Quote or reserve lot information
- Catalog Number, size, quantity, and name of products ordered

To place an order with our Customer Service, please use any of the following convenient options:

North America

Hours: Monday through Friday, 8:00am to 6:00pm EST

E-mail: order.us@lonza.com Phone: 800 638 8174 Fax: 301 845 8338

Mail

Lonza Walkersville, Inc. Customer Service Department 8830 Biggs Ford Road Walkersville, MD 21793 USA

Europe

Hours: Monday through Friday, 8:30am to 5:30pm CET

E-mail: order.europe@lonza.com

Customer Service European Office Numbers

	Phone	Fax
Austria	0800 201 538	0800 201 536
Belgium	+32 87 321 611	+32 87 321 634
Denmark	808 83 159	808 83 178
France	0800 91 19 81	0800 91 19 80
Germany	0800 182 52 87	0800 182 52 78
Ireland	1 800 654 253	1 800 654 259
ltaly	800 789 888	800 789 889
Spain	900 963 298	900 963 299
Sweden	020 790 220	020 797 710
Switzerland	0800 83 86 20	0800 83 86 19
The Netherlands	0800 022 4525	0800 022 4942
United Kingdom	0808 2349 788	0808 2349 778

Mail

Lonza Verviers, S.p.r.l. Parc Industriel de Petit-Rechain 4800 Verviers, Belgium

Ordering Information — Outside the U.S. and Europe

	Phone	Fax	
Argentina	+54 34 8442 4343	+54 34 8442 4343	
	E-mail: argentina.info@	⊡lonza.com	
Australia	+61 3 9550 0883	+61 3 9550 0890	
	E-mail: bioscience.aust	ralia@lonza.com	
China	+ 86 21 6340 3488	+86 21 6340 3308	
	E-mail: lbs.china@lonz	a.com	
Brazil	+55 11 5641 3325	+55 11 2274 00 51	
	E-mail: contact.br@lon:	za.com	
India	+91 22 4342 4000	+91 22 4342 4050	
	E-mail: scientificsuppo	rt.india@lonza.com	
Japan	+81 3 5566 0612	+81 3 5566 0619	
	E-mail: scientific.support.jp@lonza.com		
Singapore	+65 6521 4379	+65 6521 4378	
	E-mail: bioscience.asia	@lonza.com	

If your country is not listed above, please check the current list of Lonza global distributors on our website:

www.lonza.com/dist

In the event that you do not find an authorized distributor for your country in the list, please contact:

Lonza Walkersville, Inc.

Customer Service Department 8830 Biggs Ford Road Walkersville, MD 21793 USA Phone: +1 301 898 7025 Fax: +1 301 845 8338

or

Lonza Verviers, S.p.r.l.

Verviers, Belgium

Phone: +32 87 321 611 Fax: +32 87 351 967

E-mail: info.europe@lonza.com

Terms and Conditions

Please visit our website to find the current terms and conditions:

www.lonza.com/termsandconditions

How to Get Scientific Support

Providing world-class technical support for our products is a top priority. Valuable information is available to you around the clock in our QC Insider™ Toolbox, which contains comprehensive support tools that help users perform the bacterial endotoxins test. The QC Insider™ Toolbox puts support right at your fingertips and is designed for novices and experts alike: www.lonza.com/qcinsider.

www.lonza.com/coa

To access Certificates of Analysis

Safety Data Sheets

They can be accessed via the specific product page and are available after log-in.

Our Scientific Support Representatives rely on years of laboratory experience to assist with product selection and help you maximize product performance. Do you need advice on which products suit your projects? Do you need support on a challenging experiment?

Get in Touch with Our Scientific Support Team by Phone or E-mail:

Scientific Support North America

Hours: Monday through Friday, 8:00am to 6:00pm EST

Phone: 800 521 0390 Fax: 301 845 8338

E-mail: scientific.support@lonza.com

Phone from outside the US: +1 301 898 7025

Scientific Support International

Hours: Monday through Friday; 9:00am and 6:00pm CET

Phone: +32 87 321 611

E-mail: scientific.support.eu@lonza.com

Trademarks and Patents

Trademarks of Lonza Group or its affiliates

Endotoxin Challenge Vials™ PyroTec™

Kinetic-QCL™ PyroWave™

Kinectic SmartStop™ QC Insider™

PyroGene™ QCL-1000™

PYROGENT™ SpeedFill™

PYROSPERSE™ WinKQCL™

Patents

Lonza Group or its affiliates are owner or licensee** of the following patents. Products described herein may be covered by one or more of these United States patents, by pending patent applications, or by corresponding patents or applications in other countries.

Various components of PyroGene™ are protected under the following patents: Israel 71,906; Japan 2,129,487 & 2,644,447; Korea 51,077; Mexico 164,250; Norway 173,944; New Zealand 208,259; Philippines 25,395; South Africa 843,914; Spain 532,825; Taiwan 50,740. Additional Patents Pending.

Trademarks from other companies

Ariba is a registered trademark of Ariba Inc. or its subsidiaries in the United States and other jurisdictions.

BD Falcon is a trademark of Becton, Dickinson and Company.

BioTek, ELx808, FLx800, Eon, 4-Zone and Synergy are trademarks of BioTek™ Instruments, Inc.

Eppendorf, Combtips and Biopur are registered trademarks of Eppendorf AG. Molecular Devices, and SpectraMax are registered trademarks and Gemini and VersaMax are trademarks of Molecular Devices, LLC.

Vortex-Genie is a registered trademark of Scientific Industries.

Tecan and EVOware are registered trademarks and Sunrise is a trademark of Tecan Group Ltd.

Active Directory and Windows are registered trademarks of Microsoft Corporation in the United States and other countries.

All other trademarks mentioned herein are either the intellectual property of Lonza or belong to other companies where their related products were used in the scope of research experiments or referenced scientifically otherwise.

QC Insider™ Toolbox

Endotoxin Expertise At Your Fingertips™

The QC Insider™ Toolbox has been designed for endotoxin testing novices as well as experts to provide endotoxin testing expertise at any level. The online portal contains a comprehensive offering of beginner and advanced support tools, a wide range of training resources, and a library of information that can be accessed at any time and from anywhere with internet access. The QC Insider™ Toolbox is organized into three categories so that users can easily navigate directly to the support tool they need.



The QC Insider™ Support offers oneon-one guidance, detailed information about software support, recertification and testing services, reader installation and maintenance, and workflow optimization.



The QC Insider™ Training contains self-directed training resources that will help users increase their endotoxin testing expertise, including a series of how-to videos that demonstrate different assay procedures.



The QC Insider™ Library consists of technical resources such as package inserts, quick guides, and technical tips that will help lead to success with endotoxin testing.



Become a QC Insider™ Expert today and ensure the support you need is always within reach.



Endotoxin Detection Assays



Endotoxin Detection Assays

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Endotoxin Detection: A Brief History

Ever since the pharmaceutical industry began manufacturing injectables, pyrogen detection tests have been an absolute necessity. Pyrogens are substances that can cause fever, shock, and even death if high levels are introduced into the body. Endotoxins are natural compounds found in the outer cell membrane of Gram-negative bacteria and are released upon cell lysis. Endotoxins are a type of pyrogen. Today, endotoxin detection tests are performed on raw materials, in-process materials, and for the final release of pharmaceutical and medical device products.

For most of the 20th century, the rabbit pyrogen test was the standard method of testing for pyrogenicity. This test, which takes approximately four hours, is accomplished by injecting the drug being analyzed into a rabbit's ear. If the animal develops a fever it confirms the presence of pyrogens.

The LAL (Limulus Amebocyte Lysate) test was commercially introduced in the 1970s. LAL is derived from the blood cells, or amebocytes, of the Atlantic horseshoe crab (Limulus polyphemus). LAL was developed into a test for endotoxin after Frederick Bang and Jack Levin observed that the amebocytes of the horseshoe crab contain a clotting agent that forms in the presence of Gram-negative bacteria. They recognized that this clotting agent could be used as a definitive way to test pharmaceutical drugs for the presence of Gram-negative bacteria and their endotoxins. In a notice published in the Federal Register on November 4, 1977, the FDA described conditions for the use of LAL as an end-product test for endotoxin in human biological products and medical devices. The FDA widely recognizes that the LAL test is much faster, more economical, and more efficient than the rabbit pyrogen test. In addition, the LAL test is less labor intensive than the rabbit test, which makes it possible to perform many tests in a single day.

To obtain the lysate required for the LAL test, horseshoe crabs are taken from the ocean floor and a small amount of their blood is drawn. The animals are then returned unharmed to the sea. The crab's blood cells, or amebocytes, are then separated and lysed to obtain the cellular proteins.

As LAL became the preferred endotoxin detection test, different methods were developed. Each method with its own unique benefits. For example, Gel Clot LAL (PYROGENT[™]) provides a simple positive/negative result and is mentioned in most pharmacopeial monographs as the official referee test. The kinetic turbidimetric LAL assay (PYROGENT[™]-5000) gives a quantitative result and offers an economical choice for water or large volume parenterals. The endpoint chromogenic LAL test (QCL-1000[™]) offers a quantitative result and exhibits less product interference than LAL methods utilizing the clotting protein. Detecting as low as 0.005 EU/mL, the kinetic chromogenic LAL assay (Kinetic-QCL[™]) provides the benefit of less product interference for proteins, vaccines, and other biologicals while also being our most sensitive assay.

Currently the FDA, the United States Pharmacopeia (USP), the European Pharmacopeia (EP), and the Japanese Pharmacopeia (JP) accept all of the above LAL methods, as do most individual country pharmacopeias.

More recently, Lonza scientists have developed a reliable and sustainable endotoxin detection test method that is not derived from horseshoe crab blood. The PyroGene™ Assay is based on the recombinantly expressed Factor C which is the first component in the LAL clotting cascade activated by endotoxin. It is specific for endotoxin and offers a reliable alternative for endotoxin release testing. The PyroGene™ Assay promises to reduce the dependence on animal-based endotoxin assays. In 2009, the FDA approved 510(K) applications that included the PyroGene™ Assay as the final release test. The latest FDA Guidance for Industry document on "Pyrogen and Endotoxins Testing: Questions and Answers" from 2012 accepts the use of PyroGene™ as an alternative method. Please refer to page 20 for further information.



Overview of LAL Testing Procedures

There are four basic types of assays, each of which is designed to perform a different aspect of LAL testing. Our WinKQCL™ Software supports all of these assay types and is the ideal tool to accompany your quantitative endotoxin assays. It offers a fully integrated solution for reporting and analyzing your endotoxin assay results.

Routine

A routine assay calculates the concentration of endotoxin in unknowns by comparison to the performance of a series of endotoxin standards. As part of a routine assay, the user has the option to include a Positive Product Control (PPC) as a monitor for product inhibition or enhancement. A PPC is a sample of product to which a known amount of endotoxin has been added. For quantitative assays, our WinKQCL™ Software automatically calculates the amount of endotoxin recovered in the PPC and compares it to the known amount of the endotoxin in the well to give the user a percentage of recovery.

Inhibition/Enhancement

The Limulus Amebocyte Lysate reaction is enzyme mediated and, as such, has an optimal pH range, specific salt concentrations, and divalent cation requirements. Occasionally, test samples may alter these optimal conditions to an extent that the lysate is rendered insensitive to endotoxin. Negative results with samples that inhibit the LAL test do not necessarily indicate the absence of endotoxin.

An inhibition/enhancement assay is designed to determine what level of product dilution or other treatment overcomes inhibition or enhancement. Each product dilution must be accompanied by a Positive Product Control (PPC). For quantitative assays, our WinKQCL™ Software calculates the amount of endotoxin recovered in the PPC for comparison to the known amount of endotoxin spike. In this manner it can be determined which product dilutions are non-inhibitory.

RSE/CSE

An RSE/CSE assay is designed to determine the potency of a Control Standard Endotoxin (CSE) in terms of the concentration units of the Reference Standard Endotoxin (RSE). The assay requires a single series of RSE dilutions and one or more sets of dilutions of the CSE. If you buy matched reagents, we have already performed this test for you. Our CSE is matched against the USP RSE. Matched CSE is either part of the kit or is available separately.

Initial Qualification

An Initial Qualification assay is required as part of the validation of the LAL assay and is also to be performed with each new lot of reagents. It serves to confirm reagent performance and assure reproducibility. In addition, it shows analyst qualification. For this assay, a series of endotoxin standards is prepared and tested in at least triplicate. To confirm sensitivity/linearity, the test result must meet regulatory requirements as defined by pharmacopeia. For gel clot assays, the determined end-point must fall between 2 λ and 0.5 λ of the labeled sensitivity. For the quantitative assays, the results are used to generate a standard curve which must have a correlation coefficient of $\geq |0.980|$. The Initial Qualification assay does not provide for the inclusion of any samples.

Endotoxin Detection Methods

Qualitative (Yes/No Answer)

Product: PYROGENT™ Gel Clot LAL Assay

- Method Visual inspection of gel formation
- Maximum sensitivity 0.03 EU/mL
- Instrument required A dry heat block or water bath

Benefits

Simple LAL test not requiring sophisticated instrumentation and software



Overview of Endotoxin Detection Methods

Continued

Quantitative

(Results calculated from standard curve)

Product: Kinetic-QCL™ Kinetic Chromogenic LAL Assay

- Method Kinetic measurement of color development
- Maximum sensitivity 0.005 EU/mL
- Instrument required Incubating absorbance reader

Benefits

- Our most sensitive LAL-based method
- Less sensitive to product inhibition
- Ideal for biological products such as vaccines and antibiotics

Product: QCL-1000™ Endpoint Chromogenic LAL Assay

- Method Endpoint measurement of color development
- Maximum sensitivity 0.1 EU/mL
- Instrument required Spectrophotometer or absorbance reader, dry heat block

Benefits

Results in 16 minutes

Product: PYROGENT™-5000 Kinetic Turbidimetric LAL Assay

- Method Kinetic measurement of turbidity development
- Maximum sensitivity 0.01 EU/mL
- Instrument required Incubating absorbance reader

Benefits

 Cost-effective method for water and large volume parenterals

Product: PyroGene™ rFC Endpoint Fluorescent Assay

- Method Endpoint measurement of fluorescence
- Maximum sensitivity 0.005 EU/mL
- Instrument required Incubating fluorescence reader

Benefits

- Elimination of false positive glucan reactions
- Less lot-to-lot variability
- Animal-free source and security of supply
- FDA acknowledged alternative to LAL

Kinetic Chromogenic LAL Assay Overview

The Kinetic-QCL™ Kinetic Chromogenic Assay is a quantitative, kinetic assay for the detection of Gramnegative bacterial endotoxin. A sample is mixed with the reconstituted LAL reagent in a 96-well plate and placed in an incubating absorbance plate reader that measures absorbance at 405 nm. The reaction is automatically monitored over time for the appearance of a yellow color.

In the presence of endotoxin, the lysate will begin to cleave the chromogenic substrate, causing the solution to become yellow. The time required for the change is inversely proportional to the amount of endotoxin present. The concentration in unknown samples can be calculated from a standard curve. Due to the nature of this assay, the Kinetic-QCL™ Assay is less impacted by inhibitory products that may interfere with the clotting mechanism in turbidimetric and gel clot assays. This feature, along with the sensitivity range of 0.005 to 50 EU/mL, makes this assay optimal for biological products such as vaccines and antibiotics.

Using our extensive experience and practical expertise with endotoxin detection and its regulatory requirements, Lonza has developed an integrated system to support quantitative endotoxin detection. Each system component has been validated and can be verified. This all leads to reliable, reproducible, and accurate quantitative results.

Each quantitative system incorporates three elements:

- Kinetic-QCL™ Kinetic Chromogenic LAL Assay
- WinKQCL™ Endotoxin Detection and Analysis Software
- Incubating Absorbance Plate Reader

These elements integrate seamlessly to meet your testing requirements, providing meaningful results that allow you to be confident in your critical decision-making.



Benefits

Sensitivity range from 0.005 to 50 EU/mL

Applications

Ideal for biological products such as vaccines and antibiotics

Requirements

- Incubating Absorbance Plate Reader
- WinKQCL™ Software
- LAL Reagent Water (for larger kits)
- Pyrogen-free Test Tubes
- LAL Reagent Grade Multi-well Plates

Kinetic-QCL™ Kinetic Chromogenic LAL Assay

The Kinetic-QCL™ Kinetic Chromogenic Assay kits contain co-lyophilized lysate/substrate and matched control standard endotoxin (Cat. No. 50-650U also contains LAL Reagent Water).

Bulk kit configurations are also available for those kits. Kinetic Chromogenic LAL and matched control standard endotoxin are packaged separately but should be ordered together. These bulk configurations are made to order, and therefore require a lead time.

Please contact Customer Service for more information.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/kqcl
- 2°C to 8°C



Benefits

- Sensitivity range from 0.005 to 50 EU/mL
- Less sensitive to product inhibition than assays requiring gel formation
- Available in 192-, 2040-, and 2400-test kit and bulk configurations

Ordering Information -Kinetic-QCL™ Kinetic Chromogenic LAL Assay

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
50-650U	50-650U	Kinetic-QCL™ Kinetic Chromogenic LAL Assay	8×24 tests/vial Lysate, 2 vials endotoxin, 3×30 mL vial LAL Reagent Water	192 tests	0.005 to 50
50-650NV	50-650NV	Kinetic-QCL™ Kinetic Chromogenic LAL Assay	85 × 24 tests/vial Lysate, 15 vials endotoxin	2,040 tests	0.005 to 50
50-650H	50-650H	Kinetic-QCL™ Kinetic Chromogenic LAL Assay	100 × 24 tests/vial Lysate, 10 vials endotoxin	2,400 tests	0.005 to 50
K50-643L	K50-643L	Kinetic-QCL™ Bulk Kinetic Chromogenic LAL Assay	25 × 24 tests/vial Lysate	600 tests	0.005 to 50
K50-643U	K50-643U	Kinetic-QCL™ Bulk Kinetic Chromogenic LAL Assay	100 × 24 tests/vial Lysate	2,400 tests	0.005 to 50

Control Standard Endotoxin for Kinetic-QCL™ Bulk Kinetic Chromogenic LAL

The Control Standard Endotoxin, derived from E. coli 055:B5, is referenced against the USP Reference Standard Endotoxin.

Ordering Information – Control Standard Endotoxin for Kinetic-QCL™ Bulk Kinetic Chromogenic LAL

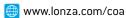
Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
E50-643L	E50-643L	Control Standard Endotoxin for Kinetic-QCL™ Bulk Kinetic Chromogenic LAL, <i>E. coli</i> Strain 055:B5	50 EU/mL	25 vials

Related Products	Page
WinKQCL™ Endotoxin Detection and Analysis Software	33
ELx808™ Incubating Absorbance Plate Reader	30
LAL Reagent Grade Multi-well Plates	37
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	
Tips and Reserviors	

QCL-1000™ Endpoint Chromogenic LAL Assay

The QCL-1000™ Endpoint Chromogenic LAL Assay is the most rapid of the LAL tests. This chromogenic LAL method is based on the formation of a yellow color and is measured photometrically at 405–410 nm. With the QCL-1000™ Assay, a multichannel pipette, a dry heat block and a 96-well plate, you can run 96 reaction wells at one time. This assay can also be run in tubes.

For your convenience, Certificates of Analysis are available online:





Benefits

- Less sensitive to product inhibition than assays requiring gel formation
- Sensitivity from 0.1 to 1 EU/mL
- Quantitative results in 16 minutes
- Flexible format use test tubes or 96-well plates
- Can be run with a simple spectrophotometer, no need for incubating absorbance plate reader



2°C to 8°C

Ordering Information – QCL-1000™ Endpoint Chromogenic LAL Assay

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
50-647U	50-647U	QCL-1000™ Endpoint Chromogenic LAL Assay	5×24 tests/vial Lysate, 1×1 mL vial endotoxin, 2×6.5 mL vial chromogenic substrate, 2×30 mL vial LAL Reagent Water	120 tests	0.1 to 1
50-648U	50-648U	QCL-1000™ Endpoint Chromogenic LAL Assay	5×60 tests/vial Lysate, 2 \times 1 mL vial endotoxin, 5×6.5 mL vial chromogenic substrate	300 tests	0.1 to 1

Ordering Information - Control Standard Endotoxin for QCL-1000™ Endpoint Chromogenic LAL

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
E50-640	E50-640	Control Standard Endotoxin for QCL:1000™ Endpoint Chromogenic LAL, <i>E coli</i> Strain 0111:B4	Endotoxin, 15 to 40 EU/mL	1 vial	n/a

Related Products	Page
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	36
LAL Reagent Grade Multi-well Plates	37
LAL Reagent Reservoirs	38
Eppendorf® 2–200 µL Biopur® Pipette tips	38
Eppendorf® 50–1000 μL Biopur® Pipette tips	38
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Kinetic Turbidimetric LAL Assay Overview

The PYROGENT™-5000 Assay is a quantitative, kinetic assay for the detection of Gram-negative bacterial endotoxin. A sample is mixed with the reconstituted LAL reagent in a 96-well plate and placed in an incubating absorbance plate reader that measures absorbance at 340 nm. The reaction is automatically monitored over time for the appearance of turbidity.

In the presence of endotoxin the lysate will begin to gel, causing the solution to become cloudy or turbid. The time required for the change is inversely proportional to the amount of endotoxin present. The concentration in unknown samples can be calculated from a standard curve.

The PYROGENT™-5000 Assay is perfect for laboratories needing to process a large number of samples. It is ideal for water samples, large volume parenterals, and water rinse from medical devices.

Using our extensive experience and technical expertise with endotoxin detection and its regulatory requirements, Lonza has developed an integrated system to support quantitative endotoxin detection. Each system component has been validated and can be verified. This all leads to reliable, reproducible, and accurate quantitative results.

Each quantitative system incorporates three elements:

- PYROGENT™-5000 Kinetic Turbidimetric LAL Assay
- WinKQCL™ Endotoxin Detection and Analysis Software
- Incubating Absorbance Plate Reader

These elements integrate seamlessly to meet your testing requirements, providing meaningful results that allow you to be confident in your critical decision-making.



Benefits

- Sensitivity range from 0.01 to 100 EU/mL
- Select from a wide range of kit sizes

Applications

Cost-effective method for water and large volume parenterals

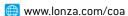
Requirements

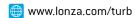
- Incubating Absorbance Plate Reader
- WinKQCL™ Software
- LAL Reagent Water
- Pyrogen-free Test Tubes
- LAL Reagent Grade Multi-well Plates

PYROGENT™-5000 Kinetic Turbidimetric LAL Assay

PYROGENT™-5000 kits contain turbidimetric lysate, reconstitution buffer for the lysate, and matched control standard endotoxin. Bulk kit configurations are available with the three assay components packaged separately. The kinetic turbidimetric LAL, reconstitution buffer, and matched Control Standard Endotoxin should be ordered together. These bulk configurations are made to order and therefore require a lead time. Please contact Customer Service for more information.

For your convenience, Certificates of Analysis are available online:









Benefits

- Sensitivity range from 0.01 to 100 EU/mL
- Available in 100-, 200-, 2,250-, and 4,500-test kit and bulk configurations

Ordering Information -PYROGENT™-5000 Kinetic Turbidimetric LAL Assay

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
N383	N383	PYROGENT™-5000 Kinetic Turbidimetric LAL Assay	2 × 50 tests/vial Lysate, 2 vials reconstitution buffer, 1 vial endotoxin	100 tests	0.01 to 100
N384	N384	PYROGENT™-5000 Kinetic Turbidimetric LAL Assay	2 × 100 tests/vial Lysate, 2 vials reconstitution buffer, 1 vial endotoxin	200 tests	0.01 to 100
N588	N588	PYROGENT™-5000 Kinetic Turbidimetric LAL Assay	45 × 50 tests/vial Lysate, 45 vials reconstitution buffer, 10 vials endotoxin	2,250 tests	0.01 to 100
N688	N688	PYROGENT™-5000 Kinetic Turbidimetric LAL Assay	${45\times100~\text{tests/vial Lysate, }45~\text{vials}}$ reconstitution buffer, 10 vials endotoxin	4,500 tests	0.01 to 100
T50-300L	T50-300L	PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL Assay	25 × 50 tests/vial Lysate	1,250 tests	0.01 to 100
T50-300U	T50-300U	PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL Assay	100 × 50 tests/vial Lysate	5,000 tests	0.01 to 100
T50-600L	T50-600L	PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL Assay	25 × 100 tests/vial Lysate	2,500 tests	0.01 to 100
T50-600U	T50-600U	PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL Assay	100 × 100 tests/vial Lysate	10,000 tests	0.01 to 100

Reconstitution Buffer for PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL

The reconstitution buffer is provided for rehydration of the PYROGENT™-5000 LAL Reagent.

Ordering Information - PYROGENT™-5000 Bulk Kinetic Turbidimetric Reconstitution Buffer

0				
Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
B50-300L	B50-300L	PYROGENT™-5000 Bulk Kinetic Turbidimetric Reconstitution Buffer	Reconstitution buffer for T50-300L	25 vials
B50-300U	B50-300U	PYROGENT™-5000 Bulk Kinetic Turbidimetric Reconstitution Buffer	Reconstitution buffer for T50-300U	100 vials
B50-600L	B50-600L	PYROGENT™-5000 Bulk Kinetic Turbidimetric Reconstitution Buffer	Reconstitution buffer for T50-600L	25 vials
B50-600U	B50-600U	PYROGENT™-5000 Bulk Kinetic Turbidimetric Reconstitution Buffer	Reconstitution buffer for T50-600U	100 vials

Control Standard Endotoxin for PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL

The Control Standard Endotoxin, derived from E. coli 055:B5, is referenced against the USP Reference Standard Endotoxin.

Ordering Information — Control Standard Endotoxin for PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity
7460L	7460L	Control Standard Endotoxin for PYROGENT™-5000 Bulk Kinetic Turbidimetric LAL, <i>E. coli</i> Strain 055:B5	100 EU/mL	25 vials	n/a

Related Products	Page
WinKQCL™ Endotoxin Detection and Analysis Software	33
ELx808™ Incubating Absorbance Plate Reader	30
LAL Reagent Grade Multi-well Plates	37
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	36

PyroGene™ Recombinant Factor C Assay

The PyroGene™ Recombinant Factor C Assay is an animal-free alternative to LAL that has been accepted by the FDA as an alternative method.* It is based on a recombinantly produced form of Factor C (rFC), the first component in the horseshoe crab clotting cascade. It is activated by endotoxin binding. The active moiety created then acts to cleave a synthetic substrate, which results in the release of a fluorophore. The reaction is run in a 96-well microplate and measured at time zero and after a one-hour incubation in a fluorescence microplate reader using excitation/emission wavelengths of 380/440 nm.

A global, multi-center study demonstrated that the recovery of endotoxin from water and other tested products using the PyroGene™ Recombinant Factor C Assay was comparable to that of LAL-based methods. The results of the assay validation were published in the Pharmacopeial Forum Vol. 36(1) [Jan. – Feb. 2010].

In June 2012, the FDA issued the document "Guidance for Industry - Pyrogen and Endotoxins Testing: Questions and Answers" which allows for the use of a recombinant Factor C based assay as an alternative to Limulus Amebocyte Lysate (LAL)-based assays. In July 2015, rFC became officially recognized by the European Pharmacopoeia (Ph. Eur.) as an alternative endotoxin detection methodology to the LAL and Rabbit Pyrogen Tests in the new draft of Chapter 5.1.10.

USP 28—NF 33 General Notices allows alternative methods if they provide advantages regarding accuracy, sensitivity, precision, selectivity, or adaptability to automation. However, to use these alternative methods for final release testing, one may need to validate the test method on their products as described in the general chapter "Validation of Compendial Procedures" < 1225> and it must be shown to give equivalent or better results.

Each quantitative system incorporates three elements:

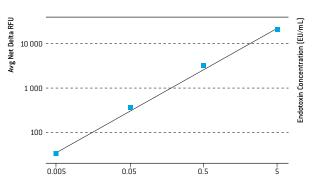
- PyroGene™ Recombinant Factor C Assay
- WinKQCL™ Endotoxin Detection and Analysis Software
- Fluorescence Plate Reader

These elements integrate seamlessly to meet your testing requirements, providing meaningful results that allow you to be confident in your critical decisions-making.

Applications

- Water testing
- In-process testing
- Final release testing
- Testing plant-based material





Standard curve illustrating assay range from 0.005 to 5 EU/mL

Benefits

- Sensitivity range from 0.005 to 5 EU/mL
- Higher endotoxin specificity
- Elimination of false positive glucan reactions
- Less lot-to-lot variability
- Animal-free
- Security of supply
- FDA acknowledged alternative to LAL

Requirements

- Incubating fluorescence reader
- WinKQCL™ Software
- Pyrogen-free test tubes
- LAL reagent grade multi-well plates
- LAL reagent water (for larger kits)

For your convenience, Certificates of Analysis are available online:



www.lonza.com/pyrogene

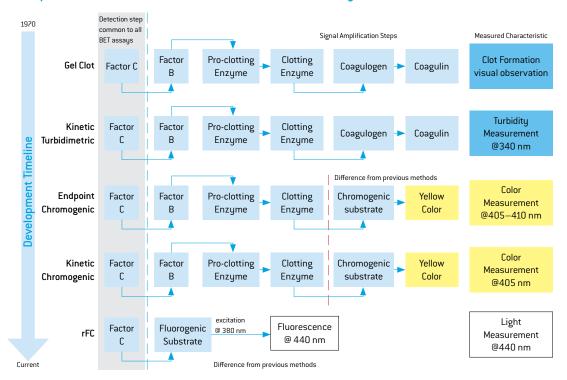
2°C to 8°C

* According to the FDA "Guidance for Industry - Pyrogen and Endotoxins Testing: Questions and Answers" document from June 2012, alternative assays should be validated as described in the USP General Chapter <1225>, "Validation of Compendial Procedures".

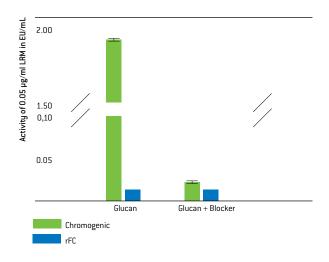
PyroGene™ Recombinant Factor C Assay

Continued

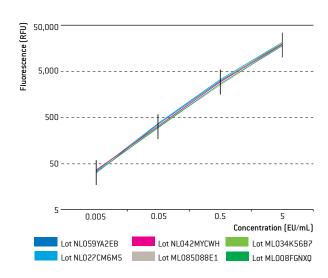
Comparison of Amplification Methods in Bacterial Endotoxin Detection Assays



rFC is the same binding protein operating in the LAL assay. The activated recombinant Factor C enzyme cleaves a substrate directly instead of activating another enzyme in a series (the LAL cascade). The substrate has a fluorescent tag, which gives a wide dynamic range with better resolution.



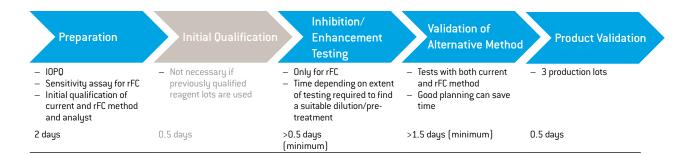
Comparison of reactivity towards glucans between kinetic chromogenic LAL and rFC. The false positive signal from the LAL assay is reduced in the presence of a glucan blocker. rFC does not detect any glucan activity; it is endotoxin-specific.



Endotoxin standard curves using 6 different lots of rFC. The log net fluorescence is proportional to the log endotoxin concentration and is linear in the 0.005 - 5 EU/mL range. Lot-to-lot standard curves exhibit excellent reproducibility.

PyroGene™ Validation Timeline

A possible validation scheme is outlined below. One validation can be accomplished in as little as 5 days, assuming that the product has been previously validated with a quantitative LAL method. The validation scheme is identical to that which would be needed for any LAL-based method with just the addition of one extra step — "Validation of alternative method". Lonza offers a full validation protocol that can be followed for your convenience. For further information, please contact Scientific Support or your local sales representative.



Ordering Information -PyroGene™ Recombinant Factor C Endpoint Fluorescent Assay

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
50-658U	50-658U	PyroGene™ Recombinant Factor C Endpoint Fluorescent Assay	2×96 tests/vial rFC enzyme solution, 2×6 mL vial fluorogenic substrate, 2×5 mL vial rFC assay buffer, 2 vials endotoxin, 2×30 mL vial LAL Reagent Water	192 tests	0.005 to 5
50-658NV	50-658NV	PyroGene™ Recombinant Factor C Endpoint Fluorescent Assay	30×96 tests/vial rFC enzyme solution, 30×6 mL/vial fluorogenic substrate, 30×5 mL/vial rFC assay buffer, 10 vials endotoxin	2,880 tests	0.005 to 5

Related Products		
Pyrogen-free Test Tubes		
LAL Reagent Grade Multi-well Plates		
LAL Reagent Reservoirs		
Eppendorf® 2–200 µL Biopur® Pipette tips		
Eppendorf® 50–1000 µL Biopur® Pipette tips		
PyroWave™ XM Fluorescence Plate Reader	31	

PYROGENT™ Gel Clot LAL Assay Overview

The PYROGENT™ Gel Clot LAL Assay is a qualitative LAL test for Gram-negative bacterial endotoxin. The gel clot assay is performed in tubes that are placed in a water bath or dry heat block at 37°C. After a one-hour incubation period, the tubes are inverted 180°. A firm clot that stays in the bottom of the tube indicates a positive reaction. If liquid flows down the side of the tube, the result is negative for endotoxin.

Like other enzymatic reactions, the LAL assay is pH dependent. The PYROGENT™ lysate formulation contains a buffer to help with these adjustments. As a result, many products will not require pH adjustments prior to testing.

PYROGENT™ Gel Clot LAL kits are available in two formats:

	Lysate	Matched Endotoxin
PYROGENT™ Gel Clot LAL		
PYRNGENT™ Plus Gel Clot I Al		



Benefits

- Easy-to-read qualitative results
- Simple LAL test not requiring sophisticated instrumentation and software
- Select from a wide range of kit sizes and sensitivities

Applications

- Water testing
- In-process testing
- Final release testing
- Testing plant-based material
- Testing acidic/basic material

Requirements

- A water bath or dry heat block
- LAL Reagent Water (LRW)
- Pyrogen-free test tubes

PYROGENT™ Gel Clot LAL Assay

PYROGENT™ Gel Clot LAL Assay standard kit sizes include 250 tests and 80 tests. Both the 250- and 80-test kits require depyrogenated 10×75 mm glass reaction tubes to run the assay.

In addition, kits of PYROGENT™ Gel Clot LAL Single Test Vials are available, which consist of 25 single use vials of lysate. Each single test vial serves as one reaction tube for an assay.

These kits do not include a matched control standard endotoxin. However, the standard can be purchased separately (Control Standard Endotoxin, page 27).

Benefits

- Sensitivities of 0.03, 0.06, 0.125, and 0.25 EU/mL available
- Easy-to-read qualitative results
- Also available as single test vials and as bulk kits



For your convenience, Certificates of Analysis are available online:

www.lonza.com/coa

www.lonza.com/gelclot

📜 2°C to 8°C

Ordering Information - PYROGENT™ Gel Clot LAL Assay (without endotoxin)

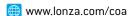
Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
N183-06	N183-06	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 16 tests/vial Lysate	80 tests	0.06
N183-125	N183-125	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 16 tests/vial Lysate	80 tests	0.125
N194-03	N194-03	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 50 tests/vial Lysate	250 tests	0.03
N194-06	N194-06	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 50 tests/vial Lysate	250 tests	0.06
N194-125	N194-125	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 50 tests/vial Lysate	250 tests	0.125
N184-25	N184-25	PYROGENT™ Gel Clot LAL Assay (without endotoxin)	5 × 50 tests/vial Lysate	250 tests	0.25
N189-06	N189-06	PYROGENT™ Gel Clot LAL Assay (without endotoxin) Single Test Vials	25 single tests/vial Lysate	25 vials	0.06
N189-125	N189-125	PYROGENT™ Gel Clot LAL Assay (without endotoxin) Single Test Vials	25 single tests/vial Lysate	25 vials	0.125
N189-25	N189-25	PYROGENT™ Gel Clot LAL Assay (without endotoxin) Single Test Vials	25 single tests/vial Lysate	25 vials	0.25

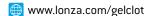
Related Products	Page
Control Standard Endotoxin for Gel Clot LAL	27
Bulk kits	
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	
Eppendorf® 2–200 μL Biopur® Pipette tips	
Eppendorf® 50–1000 μL Biopur® Pipette tips	38

PYROGENT™ Plus Gel Clot LAL Assay

The PYROGENT™ Plus Gel Clot LAL Assay combines PYROGENT™ LAL with a matched control standard endotoxin together in one kit box. Standard kit sizes include 4,000 tests, 200 tests, or 64 tests. The 4,000-, 200-, and 64-test kits require depyrogenated 10 × 75 mm glass reaction tubes to run the assay. In addition, kits of PYROGENT™ Plus Gel Clot LAL Single Test Vials are available that include 24 single-use test vials of lysate serving as the reaction tubes

For your convenience, the Certificate of Analysis documenting the FDA and USP required RSE/CSE correlation is available online:







Benefits

- Sensitivities of 0.03, 0.06, 0.125 and 0.25 EU/mL available
- No need to purchase CSE separately
- Also available as single test vials and as bulk kits



Ordering Information - PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
N283-06	N283-06	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 16 tests/vial Lysate, 1 vial endotoxin	64 tests	0.06
N283-125	N283-125	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 16 tests/vial Lysate, 1 vial endotoxin	64 tests	0.125
N294-03	N294-03	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 50 tests/vial Lysate, 1 vial endotoxin	200 tests	0.03
N294-06	N294-06	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 50 tests/vial Lysate, 1 vial endotoxin	200 tests	0.06
N294-125	N294-125	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 50 tests/vial Lysate, 1 vial endotoxin	200 tests	0.125
N284-25	N284-25	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin)	4 × 50 tests/vial Lysate, 1 vial endotoxin	200 tests	0.25
N494-03	N494-03	PYROGENT™ Plus Bulk Gel Clot LAL Assay (with endotoxin)	80 × 50 tests/vial Lysate, 20 vials endotoxin	4,000 tests	0.03
N494-06	N494-06	PYROGENT™ Plus Bulk Gel Clot LAL Assay (with endotoxin)	80 × 50 tests/vial Lysate, 20 vials endotoxin	4,000 tests	0.06
N494-125	N494-125	PYROGENT™ Plus Bulk Gel Clot LAL Assay (with endotoxin)	80 × 50 tests/vial Lysate, 20 vials endotoxin	4,000 tests	0.125
N288-25	N288-25	PYROGENT™ Plus Bulk Gel Clot LAL Assay (with endotoxin)	80 × 50 tests/vial Lysate, 20 vials endotoxin	4,000 tests	0.25
N289-06	N289-06	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin), Single Test Vials	24 single tests/vial Lysate, 1 × 1 mL vial endotoxin	24 tests	0.06
N289-125	N289-125	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin), Single Test Vials	24 single tests/vial Lysate, 1 × 1 mL vial endotoxin,	24 tests	0.125
N289-25	N289-25	PYROGENT™ Plus Gel Clot LAL Assay (with endotoxin), Single Test Vials	24 single tests/vial Lysate, 1 × 1 mL vial endotoxin	24 tests	0.25

Related Products	Page
Bulk kits	26
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	36
Eppendorf® 2–200 μL Biopur® Pipette tips	38
Eppendorf® 50–1000 μL Biopur® Pipette tips	38

PYROGENT™ Bulk Gel Clot LAL Assay

Bulk kit configurations of PYROGENT™ Gel Clot LAL are available for laboratories using large volumes of reagents. These configurations are made to order and production lead times are required. Please inquire with your sales representative for more information.

Benefits

- Bulk configurations for large volume use
- Bulk kits with and without endotoxin standard available
- Sensitivities of 0.03, 0.06, 0.125, and 0.25 EU/mL available

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- mww.lonza.com/gelclot
- 2°C to 8°C



Ordering Information -PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
E194L-03	E194L-03	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	25 × 50 tests/vial Lysate	1,250 tests	0.03
E194L-06	E194L-06	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	25 × 50 tests/vial Lysate	1,250 tests	0.06
E194L-125	E194L-125	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	25 × 50 tests/vial Lysate	1,250 tests	0.125
E209L-25	E209L-25	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	25 × 50 tests/vial Lysate	1,250 tests	0.25
F245U-06	F245U-06	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 16 tests/vial Lysate	1,600 tests	0.06
F245U-125	F245U-125	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 16 tests/vial Lysate	1,600 tests	0.125
E194U-03	E194U-03	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 50 tests/vial Lysate	5,000 tests	0.03
E194U-06	E194U-06	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 50 tests/vial Lysate	5,000 tests	0.06
E194U-125	E194U-125	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 50 tests/vial Lysate	5,000 tests	0.125
E209U-25	E209U-25	PYROGENT™ Bulk Gel Clot LAL Assay (without endotoxin)	100 × 50 tests/vial Lysate	5,000 tests	0.25

Related Products	Page
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	36
Eppendorf® 2–200 µL Biopur® Pipette tips	38
Eppendorf® 50–1000 μL Biopur® Pipette tips	38
Control Standard Endotoxin for PYROGENT™ Gel Clot LAL	27

Control Standard Endotoxin for PYROGENT™ Gel Clot LAL

Lonza's Control Standard Endotoxin is referenced against the USP Reference Standard Endotoxin.

Certificates of Analysis showing potency are available online:

www.lonza.com/coa





Ordering Information - Control Standard Endotoxin for PYROGENT™ Gel Clot LAL

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size	Sensitivity (EU/mL)
N186	N186	Control Standard Endotoxin for PYROGENT™ Gel Clot LAL Assays	Endotoxin, <i>E. coli</i> 055:B5	5 vials	n/a
7360L	7360L	Bulk Control Standard Endotoxin for PYROGENT™ Gel Clot LAL Assays	Endotoxin, E. coli 055:B5	25 vials	n/a

Related Products	Page
Gel Clot LAL Assays	24
LAL Reagent Water (LRW)	39
Pyrogen-free Test Tubes	36
Eppendorf® 2–200 μL Biopur® Pipette tips	38
Eppendorf® 50–1000 μL Biopur® Pipette tips	38

Instrumentation and Software



Instrumentation and Software

ELx808™ Incubating Absorbance Plate Reader	30
PyroWave™ XM Fluorescence Reader	31
PyroTec™ Liquid Handling System	32
WinKQCL™ Endotoxin Detection and Analysis Software	33

ELx808™ Incubating Absorbance Plate Reader

For Kinetic-QCL™ and PYROGENT™-5000 Kinetic LAL Assays

The ELx808™ Incubating Absorbance Plate Reader has been validated as part of our quantitative endotoxin detection systems. This 96-well microplate reader features 4 insulated incubating zones, providing well-to-well temperature uniformity available on the market. Utilizing interference filters instead of monochromators, the reader is optimized for the LAL-based endotoxin tests and comes pre-configured with the following interference filters: 340nm, 405nm, 450nm, 490nm, and 630nm. The ELx808™ Reader seamlessly interfaces with Lonza's WinKQCL™ Endotoxin Detection and Analysis Software.

For the kinetic and endpoint LAL assays, the ELx808™ Reader is highly reliable and easy to use. It features fully-automated kinetic reads of the 96-well plate via the WinKQCL™ Software. This reader can also be used as a standard spectrophotometer and results can be read from the QCL-1000™ Endpoint Chromogenic LAL Assay and ELISA Assays.

On-site service and preventive maintenance contracts are available to help ensure that your instrument is working properly.

Benefits

- Interference filters provide up to 5x the light energy of monochromator-based filters
- Excellent temperature uniformity
- Precise and accurate
- Cost effective filter-based reader



ELx808 [™] Absorbance Reader Specifications					
Wavelength Range	340 to 900 nm				
Filters Supplied	340, 405, 450, 490 and 630 nm				
Absorbance Range	0.000 to 4.000 0D @ 400 to 900 nm 0.000 to 3.000 0D @ 340 to 400 nm				
Temperature Control	4°C above ambient to 50°C				
Read Method	Kinetic or endpoint under WinKQCL™ Control				
Light Source	Tungsten halogen bulb				
Dimensions	16-inches deep \times 15.5-inches wide \times 8.75-inches high (40.6 cm \times 39.4 cm \times 22.2 cm)				
Weight	35 lb (15.9 kg)				

Ordering Information –ELx808™ Reader

Cat. No. NA	Cat. No. EU	Product Name	Product Description
25-315S	25-315\$	ELx808™ Reader	Incubating 8-channel absorbance plate reader
25-342	25-342	Stepped Neutral Density Plate*	For optical density validation across eight channels
7260522	7260522	BioTek™ Absorbance Test Plate	For alignment, repeatability, and accuracy validation
05105	LAL3400508	Replacement Bulb for ELx808™ Reader	
196171	75053	Computer Connection Cable for ELx808™ Reader	9 pin female to 25 pin female RS232 cable
196004	-	UPS-APC 1500VA	Uninterruptible power supply with LCD, 120 V output (US edition)
-	BE00196004	UPS-APC 1500VA	Uninterruptible power supply with LCD, 230 V output (European edition)
196005	196005	4-Port Serial PCI Card	Standard/low profile PCI card
25-361	25-361	USB to Serial Port Converter	10 cm cable with LED communication indicator lights

*Recertification service is available (see Recertification Services page 48)

PyroWave™ XM Fluorescence Reader

For the PyroGene™ rFC Endpoint Fluorescent Assay

The PyroWave™ XM Fluorescence Reader is a part of the quantitative endotoxin detection system that supports the PyroGene™ rFC Endpoint Fluorescent Assay. This reader replaces the Lonza FLx800™ LBS reader, bringing a new generation in fluorescence technology to users of the PyroGene™ Assay. Optimized specifically for Lonza's PyroGene™ Assay, this reader brings new and improved technology to the laboratory. The PyroWave™ XM Reader offers numerous enhancements in incubation, optics, automation compatibility, and overall reader maintenance and robustness. Controlled by Lonza's WinKQCL™ Endotoxin Detection and Analysis Software, version 5.2 and higher, Lonza delivers a high performance and easy to use system for users interested in an alternative to the horseshoe crabbased endotoxin detection test methods.

On-site service and preventive maintenance contracts are available to help ensure that your instrument is working properly.

Benefits

- Improved PyroGene™ Assay performance
- Reduced reader maintenance
- Automation compatible
- High performance optics
- 4-Zone™ Incubation at 37°C



PyroWave™ XM Fluores	cence Reader	Specification	ns		
Read capabilities	Fluorescence, Luminescence**, TRF**, FP**				
Read position	Top read				
Light source	Xenon flash	lamp			
Detection	High perforn	nance photo	multi-plier tu	be	
Fluorescence sensitivity	Fluorescein	@ 1 pM/well	in a 96-well p	olate	
Wavelength range	200 to 850 r	nm**			
Filters	One easy-to-swap filter cube with the following filter configuration:				
	Assay Type	Excitation Filter (nm)	Emission Filter (nm)	Dichroic Mirror (nm)	
	.	Filter (nm)			
	Туре	Filter (nm) 380/20	Filter (nm)	Mirror (nm)	
	Type PyroGene™ Fluorescein **Additional filte	Filter (nm) 380/20 485/20 r cubes and filter	Filter (nm) 440/30	Mirror (nm) 400 510 assed from reader	
	Type PyroGene™ Fluorescein **Additional filte	Filter (nm) 380/20 485/20 r cubes and filter r additional wave	Filter (nm) 440/30 528/20 s must be purches	Mirror (nm) 400 510 assed from reader	
Temperature control Power	Type PyroGene™ Fluorescein **Additional filte manufacturer fo ±0.2 °C at 37	Filter (nm) 380/20 485/20 r cubes and filter r additional wave	Filter (nm) 440/30 528/20 s must be purchallengths and read	Mirror (nm) 400 510 assed from reader	
	PyroGene** Fluorescein **Additional filte manufacturer fo ±0.2 °C at 37 100 – 240 v 39.1 cm W x	380/20 485/20 r cubes and filter r additional wave	Filter (nm) 440/30 528/20 s must be purchallengths and read 0 Hz 32.8 cm H	Mirror (nm) 400 510 assed from reader	

Ordering Information -PyroWave™ XM Reader

Cat. No. NA	Cat. No. EU	Product Name	Product Description
25-345\$	25-345\$	PyroWave™ XM Fluorescence Reader	Incubating Fluorescence Reader
196004		UPS-APC 1500VA	Uninterruptible power supply with LCD, 120 V output (US edition)
	BE00196004	UPS-APC 1500VA	Uninterruptible power supply with LCD, 230 V output (European edition)
196005	196005	4-Port Serial PCI Card	Standard/low profile PCI card
204511	7160013	Sodium Fluorescein Liquid Test Kit	For reader operational qualification and PM verification

Related Products	Page
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PyroGene™ rFC Endpoint Fluorescent Assay	20

PyroTec™ Liquid Handling System

The PyroTec™ Liquid Handling System is a robotic workstation to help automate endotoxin detection testing. The system includes a user-defined platform size to accommodate tips, reagent troughs and 96-well plates. The robotic arm picks up tips and dispenses samples and reagents to 96-well plates. Heating units can incubate plates prior to delivery into plate readers.

WinKQCL™ Software is fully integrated with the Tecan® Freedom EVOware® Software, allowing assay templates to be executed using robotic scripts for the PyroTec™ Liquid Handling System.

The PyroTec™ System can be tailored to your testing needs. Contact your local Lonza sales representative or scientific support for additional information.

On-site service and preventive maintenance contracts are available to help ensure that your instrument is working properly.



- Flexible platform to automate filling of assay plates
- Help high throughput labs manage their large daily sample requirements



Ordering Information - PyroTec™ Liquid Handling System

Cat. No. NA	Cat. No. EU	Product Name	Product Description	
25-601 25-601 PyroTec™ 200 Liquid Handling System Robotic workstation for filling nine 96-well plates at one time		Robotic workstation for filling nine 96-well plates at one time		
25-602	25-602	PyroTec™ 150 Liquid Handling System Robotic workstation for filling three 96-well plates at one time		
25-603	25-603	PyroTec™ 150 Positive ID System	stem Robotic barcode scanner	

Related Products	Page
WinKQCL™ Endotoxin Detection and Analysis Software	33
Kinetic-QCL™ Kinetic Chromogenic LAL Assay	
PYROGENT™-5000 Kinetic Turbidimetric LAL Assay	
PyroGene™ rFC Endpoint Fluorescent Assay	

WinKQCL™ Endotoxin Detection and Analysis Software

Quantitative methods such as the Kinetic-QCL™ Assay generate significant amounts of raw data that require careful analysis before reporting can take place. The WinKQCL™ Software offers a fully integrated solution for your quantitative endotoxin detection testing, data management, and reporting needs.

WinKQCL™ 5 Software meets 21 CFR Part 11 technical requirements for electronic records, signatures, audit trails and database archiving. Reader validation tests can be run from the WinKQCL™ Software and are stored in the same database. The built-in database backup and maintenance scheduler makes it easy to maintain the system.

Benefits

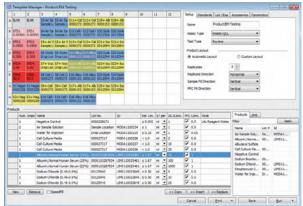
- Kinetic SmartStop™ monitoring feature to address split pair and other reaction conditions, without sacrificing time waiting for a fixed number of reads
- Enterprise level IT features including wide area network support, ability to work across time zones, application virtualization, Active Directory® integration, and data segregation by lab
- Bi-directional interface with 3rd party database systems
- Customizable endotoxin test reports
- Multi-language user interface: English, French, German, Italian, Japanese, Spanish, Portuguese, Simplified Chinese and Traditional Chinese

Applications

- For use with the PyroWave™ XM, ELx808™, and FLx800™ Readers. Extended reader integration now includes Molecular Devices® SpectraMax®, Gemini™ and VersaMax™ Readers; and the BioTek® Eon™ and Synergy™ 2 Readers. The software is also interfaced with the Tecan® Sunrise™ Reader
- Tecan® EVOware® interface integration for PyroTec™
 Liquid Handling System
- Supports all quantitative endotoxin detection assays from Lonza including QCL-1000™ Endpoint Chromogenic LAL, Kinetic-QCL™ Kinetic Chromogenic LAL, PYROGENT™-5000 Kinetic Turbidimetric LAL and PyroGene™ rFC Endpoint Fluorescent Assays
- Installation as a simple standalone system or as an interface with multiple robots and readers in multiple labs around the world, all storing data in a single database

The user-friendly and flexible Template Manager allows you to customize plate layout with a click of a mouse using the SpeedFill™ and drag 'n drop features. The interactive and enhanced trending tools provide meaningful results on demand, helping you easily detect drift and enabling you to make proactive decisions.





The endotoxin detection instruments and software from Lonza are available fully supported with Installation, Operational and Performance Qualification (IOPQ) manuals and a WinKQCL™ 5 Software Validation Package. In addition, a trained specialist from Lonza can perform the IOPQ of the complete system to help you with your system validation process. Please inquire with your local sales representative for further details.

WinKQCL™ Endotoxin Detection and Analysis Software

Continued

Ordering Information - WinKQCL™ Endotoxin Detection and Analysis Software

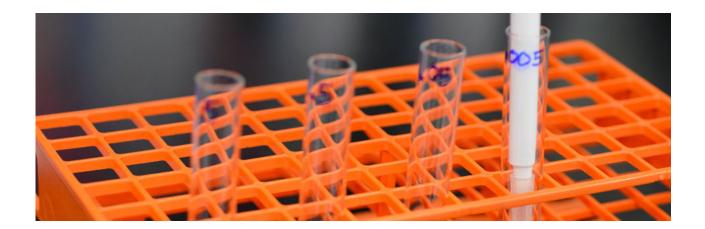
Cat. No. NA	Cat. No. EU	Product Name	Product Description	
25-501	25-501	WinKQCL™ 5 Software Package	Installation DVD, workgroup license, reader license	
25-502	25-502	WinKQCL™ 5 Workgroup License	Additional workgroup license	
25-503	25-503	WinKQCL™ 5 Reader License	Additional reader license	
25-504	25-504	WinKQCL™ 5 Qualification Manual	DVD containing IOPQ files for software and readers	
25-505	25-505		DVD containing software validation information	
25-339S	25-339S	System Qualification Service	IQ/OQ/PQ validation on site, labor only	
25-501SUP	25-501SUP	Annual WinKQCL™ Support for Cat. No. 25-501	Advanced IT/software support	
25-502SUP	25-502SUP	Annual WinKQCL™ Support for Cat. No. 25-502	Advanced IT/software support	
25-503SUP	25-503SUP	Annual WinKQCL™ Support for Cat. No. 25-503	Advanced IT/software support	

Ordering Information – WinKQCL™ Software Delivered Immediately by Email

Cat. No. NA	Cat. No. EU	Product Name	Product Description	
25-501E	25-501E	E-Delivered WinKQCL™ 5 Software Package	Installation files, workgroup license, reader license	
25-502E	25-502E	E-Delivered WinKQCL™ 5 Workgroup License	Additional workgroup license	
25-503E	25-503E	E-Delivered WinKQCL™ 5 Reader License	Additional reader license	
25-504E	25-504E	E-Delivered WinKQCL™ 5 Qualification Manual	IOPQ files for software and readers	
25-505E	25-505E	E-Delivered WinKQCL™ 5 Validation Package	File containing software validation information	

Related Products	Page	
Kinetic-QCL™ Kinetic Chromogenic LAL Assay	15	
PYROGENT™-5000 Kinetic Turbidimetric LAL Assay		
PyroGene™rFC Endpoint Fluorescent Assay		
ELx808™ Incubating Absorbance Reader		
PyroWave™ XM Incubating Fluorescence Reader		

Accessory Products



Accessory Products

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Introduction

In addition to the endotoxin detection kits, instruments, and software, Lonza offers many of the accessory items necessary to run endotoxin detection assays. Many of the items have been tested with the Kinetic-QCL™ Kinetic Chromogenic LAL Assay to help ensure their compatibility with our endotoxin detection methods. We also offer products such as the Endotoxin Challenge Vials™ to help with your oven depyrogenation validations.



Test Tubes

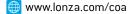
All test tubes are made from USP Type I flint borosilicate glass.

Both products N201 and N205 are recommended for use as reaction tubes in gel clot assays. N201 are provided with polypropylene screw caps. Product number N207 is recommended for dilution of endotoxin standards and test samples for all endotoxin detection assays.

Benefits

Certified to contain less than 0.005 EU/mL endotoxin

For your convenience, Certificates of Analysis are available online:



www.lonza.com/accessories



Ordering Information - Test Tubes

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
N207	N207	Pyrogen-free Test Tubes	Without caps, $13 \times 100 \text{ mm}$	30/foil pack
N201	N201	Pyrogen-free Test Tubes	With caps, 10×75 mm	50/box
N205	N205	Pyrogen-free Test Tubes	Without caps, 10 × 75 mm	50/foil pack

Sample Containers

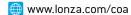
Sample containers are intended for transporting product samples for endotoxin analysis or sample storage. Proper container and storage conditions need to be validated for each individual sample.

Products 80-507L and 80-507U contain 10 mL glass vials with screw caps. Products BE2098 and BE2099 are plastic sample containers that offer greater capacity at a reduced cost

Benefits

Certified to contain less than 0.005 EU/mL endotoxin

For your convenience, Certificates of Analysis are available online:







Ordering Information - Sample Containers

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
80-507L	80-507L	Sample Containers	Depyrogenated, 10 mL glass bottle with screw cap	25/box
80-507U	80-507U	Sample Containers	Depyrogenated, 10 mL glass bottle with screw cap	100/box
	BE2098	Polypropylene Sample Containers	Endotoxin tested, 50 mL tubes	50/pack
	BE2099	Polystyrene Sample Containers	Endotoxin tested, 15 mL tubes	50/pack

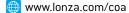
Plates

These 96-well plates can be used with the QCL-1000™ Endpoint Chromogenic LAL Assay, Kinetic-QCL™ Kinetic Chromogenic LAL Assay, PYROGENT™-5000 Kinetic Turbidimetric LAL Assay and PyroGene™ rFC Endpoint Fluorescent Assay. Each case contains individually wrapped plates.

Benefits

- Certified to contain less than 0.0005 EU/well endotoxin
- Certified for compatibility with the endotoxin detection assays
- Certified to be free from inhibition

For your convenience, Certificates of Analysis are available online:



www.lonza.com/accessories

Ordering Information - Plates

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
25-340	BE25-340	LAL Reagent Grade Multi-well Plates	96-well plates, endotoxin-tested (<0.0005 EU/well)	50/case

Pipette Tips

Pyrogen-free pipette tips are to be used when testing with any of our endotoxin detection systems. Eppendorf® Biopur® pipette tips are certified to contain <0.001 EU/mL endotoxin. The new design of the tips allows compatibility with different pipettors. Catalog Number 25-416 is for use with multi-channel pipettors. Products BE25-413 and BE25-414 are certified to contain <0.005 EU/mL endotoxin. They can be used with pipettes from different manufacturers. Eppendorf® Combitips® are for use with a multi-step pipette.

Benefits

- Endotoxin tested
- Broad offering for various pipette types

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories

Ordering Information - Pipette Tips

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
25-415	BE25-415	Eppendorf® 2–200 μL Biopur® Pipette Tips	< 0.001 EU/mL	5 trays/pkg; 96 tips/tray
25-416	BE25-416	Eppendorf® 2–300 μL Biopur® Pipette Tips	< 0.001 EU/mL, for multi-channel pipettors	5 trays/pkg; 96 tips/tray
5-417	BE25-417	Eppendorf® 50–1000 μL Biopur® Pipette Tips	< 0.001 EU/mL	5 trays/pkg; 96 tips/tray
	89634	Eppendorf® Combitips®, 0.5 mL	Single packed	100
	89650	Eppendorf® Combitips®, 2.5 mL	Single packed	100
	89669	Eppendorf® Combitips®, 5 mL	Single packed	100
	89677	Eppendorf® Combitips®, 10 mL	Single packed	100
	BE10035	Eppendorf® Pipette Tips, 2–200 μL	Endotoxin tested, single packed	100/box
	BE10051	Eppendorf® Pipette Tips, 50–1000 µL	Endotoxin tested, single packed	100/box
	BE25-413	LAL Reagent Grade Pipette Tips, 2–200 µL	<0.005 EU/mL	10 × 96 tips
	BE25-414	LAL Reagent Grade Pipette Tips, 50–1250 µL	<0.005 EU/mL	10 × 96 tips
	BE7521	Pipette BD Falcon™, 1 mL	Endotoxin tested, single packed	100
	BE7507	Pipette BD Falcon™, 2 mL	Endotoxin tested, single packed	100
	BE7543	Pipette BD Falcon™, 5 mL	Endotoxin tested, single packed	200
	BE7551	Pipette BD Falcon™, 10 mL	Endotoxin tested, single packed	200

Reservoirs

The LAL Reagent Reservoirs are for use with multi-channel pipettors when adding reagents to a 96-well plate. The reservoirs are provided in a zip closure bag enabling you to conveniently store unused reservoirs for later use.

For your convenience, Certificates of Analysis are available online:

www.lonza.com/coa



Benefits

- Certified to contain less than 0.005 EU/mL endotoxin

Ordering Information - LAL Reagent Reservoirs

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
190035	190035	LAL Reagent Reservoirs	<0.005 EU/mL	10/pack



Dry Heat Block, Inserts and Vortex Mixer

The dry heat block is used for incubation of LAL gel clot assays. The aluminum block with lid adaptor for a dry heat block allows a 96-well plate to be incubated at 37°C for use with QCL-1000™ Endpoint Chromogenic LAL Assay.





Ordering Information - Dry Heat Block, Inserts and Vortex Mixer

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
25-038A	25-038A	Aluminum Block with Lid	Heat block insert for 96-well plate	Each
	BEF3503	Aluminum Insert Block for Techne Dry Heat Block		For 20 tubes
	BEDB-2D	Techne Dry Heat Block	Digital, from 25°C to 100°C, requires 2 × BEF3503	For 40 tubes
	FDB03DD	Techne Dry Heat Block	Digital, from 25°C to 100°C, requires 3 × BEF3503	For 60 tubes
	BENP5051	Vortex Genie® 2		

LAL Reagent Water

LAL Reagent Water is recommended for reconstituting LAL reagents, as well as for the dilution of control standard endotoxin and test samples for endotoxin testing. LAL Reagent Water is equivalent to Water for Bacterial Endotoxins Test (BET). In addition to USP-required WFI tests, Lonza tests LAL Reagent Water for compatibility with our endotoxin detection assays.

Benefits

- Certified to contain less than 0.005 EU/mL endotoxin
- Available in a variety of sizes
- Certified for Positive Product Control Recovery within 75 to 150%
- 🤨 2°C to 8°C (W50-640)
- 15°C to 30°C (W50-100, W50-500, W50-1000)



For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories

Ordering Information - LAL Reagent Water

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
W50-640	W50-640	LAL Reagent Water	<0.005 EU/mL, 30 mL	1 bottle
W50-100	W50-100	LAL Reagent Water	<0.005 EU/mL, 100 mL	1 bottle
W50-500	W50-500	LAL Reagent Water	<0.005 EU/mL, 500 mL	1 bottle
W50-1000	W50-1000	LAL Reagent Water	<0.005 EU/mL, 1000 mL	1 bottle

β -G-Blocker

 β -D-glucans can produce false positive results in LAL assays. Some examples of glucan sources include yeast and cellulosic materials such as hemodialysis filters. Our β -G-Blocker may be used with any of our LAL assays.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories
- Benefits
- Certified to contain less than 0.005 EU/mL endotoxin
- Functionality tested to demonstrate a reduction of enhancement caused by β-D-glucans



Ordering Information – β -G-Blocker

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
N190	N190	β-G-Blocker	Glucan blocker, 5 mL/vial	5 vials

PYROSPERSE™ Dispersing Agent

PYROSPERSE™ Dispersing agent, helps eliminate endotoxin binding or masking in some samples — solving problems of inhibitory behavior. Examples of samples that may show endotoxin binding behavior include plasma protein fractions, electrolyte solutions, and lipid emulsions. The PYROSPERSE™ Dispersing Agent may be used with any of our LAL kits.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories
- Benefits
- Endotoxin and functionality tested



2°C to 30°C (unopened)

Ordering Information -PYROSPERSE™ Dispersing Agent

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
N188	N188	PYROSPERSE™ Dispersing Agent	5 mL/vial	5 vials

MgCl₂

MgCl₂ can be used as the sample diluent when attempting to overcome inhibitory chelation effects. Examples of samples which chelate divalent cations include heparin and EDTA. MgCl₂ may be used to prepare samples for any endotoxin detection assay.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories
- Benefits
- Certified to contain less than 0.005 EU/mL endotoxin
- 📒 2°C to 30°C (unopened)



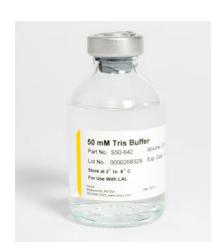
Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
S50-641	S50-641	MgCl₂ 10 mM Solution	30 mL/vial	1 vial

Tris Buffer

Tris Buffer can be used in place of water as the sample diluent for highly acidic or basic samples (for endotoxin testing, samples and sample dilutions should be between pH 6–8 after lysate addition). Tris Buffer may be used to prepare samples for any of our endotoxin detection assays.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories
- Benefits
- Certified to contain less than 0.005 EU/mL endotoxin
- Certified pH range from 7.0 to 7.4 @ 25°C
- Certified to ensure good buffering performance



10 mM MgCl₂ Solutio



Ordering Information - Tris Buffer 50 mM

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
S50-642	S50-642	Tris Buffer 50 mM	30 mL/vial	1 vial

Endotoxin and Endotoxin Challenge Vials™

Endotoxin (*E. coli*) Challenge Vials™ are for use in oven validation studies. Each vial contains >1,000 EU/vial. The vials may be tested using any of our endotoxin detection kits. Product 193783 contains high potency endotoxin and is intended for use in endotoxin removal system challenges, i.e. depyrogenation ovens, and other spiking studies. Each vial contains >1,250,000 EU/vial. E700 is the USP Reference Standard Endotoxin. Each vial contains 10,000 EU/vial.

For your convenience, Certificates of Analysis are available online:

- www.lonza.com/coa
- www.lonza.com/accessories
- Benefits
- Products 192568 and 193783 are devoid of fillers
- 192568 and 193783 are stored at 2°C to 8°C
- E700 storage conditions are -20°C





Ordering Information - Endotoxin and Endotoxin Challenge Vials™

Cat. No. NA	Cat. No. EU	Product Name	Product Description	Size
193783	193783	Endotoxin	> 1.25 million EU/vial	5 vials
192568	192568	Endotoxin Challenge Vials™	>1,000 EU/vial	25 vials
E700	E700	USP Reference Standard Endotoxin	10,000 EU/vial	1 vial

Testing Services



Testing Services

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QC Testing Solutions Services

Another means by which Lonza continues to strive to meet the needs of our QC microbiology customers is by providing complementary contract testing services for many of our QC microbiology related products.

QC Testing Solutions Services offered:

- Endotoxin Testing Services
- Recertification Services

QC Testing Solutions Services are presently operated out of two global Lonza facilities to provide our customers with worldwide access to our current service capabilities*:

- Lonza Walkersville, Maryland, USA
- Lonza Verviers, Belgium

Please contact your local sales representative or Customer Service for more information about where and how to submit samples to one of our service locations above.

Register for our QC Testing Solutions e-newsletter to stay up to date on the latest endotoxin detection products and services. Featured content includes technical tips, white papers and webinar events.

www.lonza.com/hsc

^{*}Not all testing services are offered in both locations.

Endotoxin Detection Testing Services

Lonza's Endotoxin Detection Testing Services offer you routine as well as customized endotoxin testing. Our facilities provide expertise in gel clot, kinetic chromogenic and kinetic turbidimetric LAL methods, as well as the rFC endpoint fluorescent assay.

USP, EP and FDA Compliant

All pharmaceutical grade endotoxin testing is performed in accordance with current regulatory documents such as the U.S. and European Pharmacopeia (USP and EP) Bacterial Endotoxins Test (BET), the 2012 U.S. Food and Drug Administration (USFDA) Guidance for Industry and the USP monograph for medical device testing. These documents originate from or are acceptable to the major Pharmacopeia and their regulatory authorities.

Whether you are from a university, research laboratory or a Quality Control laboratory in a major pharmaceutical company, Lonza has the capabilities you need for reliable, accurate and confidential results.

Benefits

- Market leader in endotoxin detection testing systems
- Technical expertise and full service reporting
- USP and EP compliant

STAT Testing Available*

For customers who may need faster turn-around times, Lonza offers an expedited testing service for preliminary screening, endotoxin determination, and the Endotoxin Challenge Vial™ test. Our STAT service delivers preliminary (unaudited) results in 4 days or less. STAT requests require prior approval from Lonza's Endotoxin Testing Services.

*STAT testing is only available in the US.

www.lonza.com/endotestserv

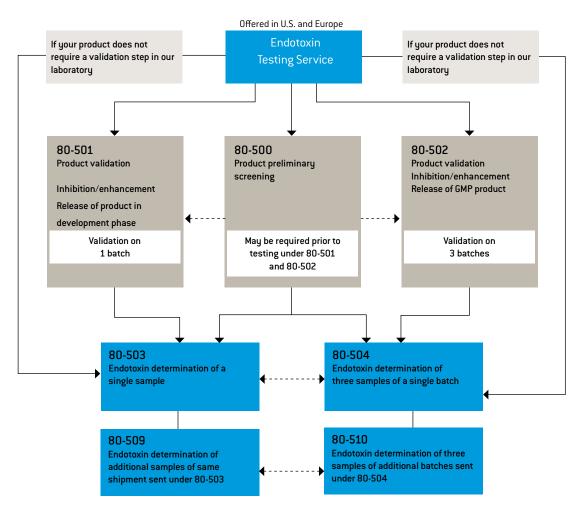


Figure 1. Endotoxin Testing Service.

Endotoxin Detection Testing Services

Continued

Pre-paid, Mail-in Anytime Testing Service

Available in Europe, the Endotoxin Testing Service offers a low cost mail-in anytime service for users that do not require testing according to pharmaceutical or medical device regulations.

This type of endotoxin testing is suitable for a wide range of non-standard tests such as:

- Samples from cell culture materials
- Samples of DNA plasmids
- Water samples from autoclaves or dialysis machines
- Dialysis water or fluids
- Research samples

Samples received using this service will be tested at the required dilutions indicated in the Pre-paid Endotoxin Testing Service diagram.

Benefits

- Low cost, convenient and quick endotoxin testing using Kinetic-QCL™ LAL Assay
- Quick, easy ordering process via web, phone, or fax
- Pre-addressed packaging for smooth delivery to Lonza's Endotoxin Testing Service

To utilize the Pre-paid Service, simply order one or more of the Rapid Endotest packs, place your sample in the sample containers provided, and mail the pre-addressed envelope back to Lonza.

When ordering the Pre-paid Service, the pre-paid endotest packs are included for shipment of the samples to the Endotoxin Testing Services. When you have a requirement for an endotoxin test, simply mail in your samples. Samples will be tested and results sent to the customer within 5 days of receipt.

www.lonza.com/endotestserv

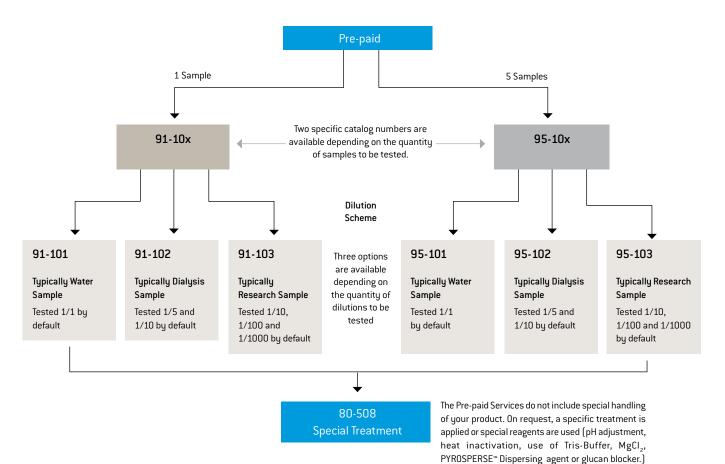


Figure 2. Pre-paid Testing Service, available in Europe. This service is not avilable in the US.

Endotoxin Detection Testing Services

Continued

Ordering Information - Endotoxin Detection Testing Services

Cat. No. NA	Cat. No. EU	Product Name	Product Description
80-500	BE80-500	Product Preliminary Screening	
80-501	BE80-501	Product Validation Inhibition/Enhancement	Release of product in development phase. Validation of one batch.
80-502	BE80-502	Product Validation Inhibition/Enhancement	Release of GMP product. Validation of three batches.
80-503	BE80-503	Endotoxin Determination of Single Sample	
80-509	BE80-509	Additional samples of same shipment sent under 80-503	
80-504	BE80-504	Endotoxin Determination of Three Samples of a Single Batch	
80-510	BE80-510	Endotoxin Determination of three samples of additional batches sent under 80-504	
80-508	BE80-508	Unusual sample treatment or handling charge per hour	
80-514	BE80-514	Exclusion of β-D-glucan activity testing	
80-535		Depyrogenation Endotoxin Challenge Vial™ Test	-
80-544	BE80-544	Sample container screening	
80-500S		STAT: Product preliminary screening of one batch	
80-503S		STAT: Endotoxin determination of single sample	
80-509S		STAT: Additional samples of same shipment sent under 80-503	
80-504S		STAT: Endotoxin determination of three samples of a single batch	
80-510S		STAT: Endotoxin determination of three samples of additional batches sent under 80-504	
80-514S		STAT: Exclusion of β-D-glucan activity testing	
80-535S		STAT: Depyrogenation Endotoxin Challenge Vial™ Test	

STAT Testing is not available in Europe.

Ordering Information - Pre-paid Endotoxin Detection Testing Services

Cat. No. NA	Cat. No. EU	Product Name	Product Description
	BE91-101	Rapid-Endotest Single Water Sample	
	BE91-102	Rapid-Endotest Single Dialysis Fluid Sample	
	BE91-103	Rapid-Endotest Single Research Sample	
	BE95-101	Rapid-Endotest Five Water Samples	
	BE95-102	Rapid-Endotest Five Dialysis Fluid Samples	
	BE95-103	Rapid-Endotest Five Research Samples	
	80-515-X1	15 Endotoxin Challenge Vials™	Endotoxin testing of baked vials with positive product controls. Price includes ECVs.
	80-515-X2	20 Endotoxin Challenge Vials™	Endotoxin testing of baked vials with positive product controls. Price includes ECVs.
	80-515-X3	25 Endotoxin Challenge Vials™	Endotoxin testing of baked vials with positive product controls. Price includes ECVs.

Please contact your local Customer Service Team to obtain general information and the appropriate sample submission forms.

U.S. Endotoxin Testing Services

E-mail: scientific.support@lonza.com

European Endotoxin Testing Services

E-mail: scientific.support.eu@lonza.com

Recertification Services

Recertification Service for Stepped Neutral Density Test Plates

The Stepped Neutral Density (SND) and Absorbance (Universal) Test plates provide the ELx808™ Microplate Reader user with the ability to manage risk and help assure the reader is functioning properly between the recommended six-month preventive maintenance visits.

The ELx808™ Readers are robust instruments designed to provide years of reliable performance for endotoxin detection and other absorbance-based assays. In following current Good Manufacturing Practices (cGMP), the reader's performance should be verified and documented at regular intervals.

In order to properly assess the reader performance, the plates used in the testing must be shown to be acceptable. Plate manufacturers recommend the SND plate be recertified at least once per year.

Benefits

- Manages risk and helps ensure optimal performance of the ELx808™ Readers
- cGMP compliant processes, including testing of multiple parameters
- NIST-traceable testing conducted in an ISO 9001 certified facility

Parameters tested

- Absorbance accuracy (linearity and slope)
- Precision (reproducibility of results)
- Wavelength accuracy
- Optical alignment
- Detailed physical inspection



Reports include

- Physical inspection observations
- Coefficient of variation (%CV) for each plate standard
- Slope, R-value and Y-intercept linearity data
- Reader and plate uncertainty values
- Approval by a Quality Assurance representative
- Recertification due date (default is 12 months)

Please refer to the table below for additional details and ordering information. For testing parameters, reports and pricing, please contact your local sales representative.



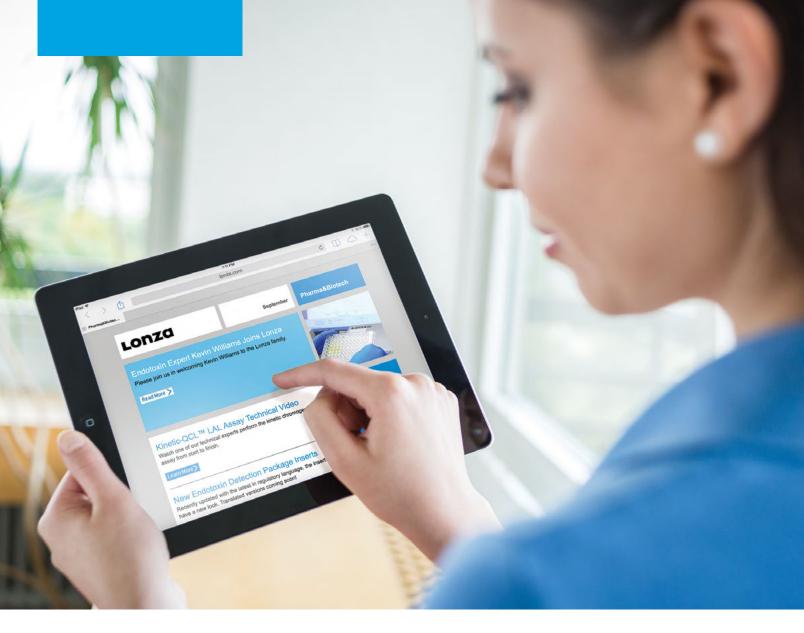
www.lonza.com/recert

Ordering Information – Recertification Services

Cat. No. NA	Cat. No. EU	Product Name	Product Description
85-011-SND	85-011-SND	Stepped Neutral Density Plate Recertification Service	Two filter wavelengths, 405 nm and 340 nm
85-012-SND	85-012-SND	Stepped Neutral Density Plate Recertification Service	Each additional filter wavelength
85-011BA	85-011BA	Stepped Neutral Density Plate Recertification Service	As Found and As Left testing at two filter wavelengths 405 nm and 340 nm
	85-013	Absorbance (Universal) Test Plate	Recertification Service*

SND plate recertification services offered for Lonza plate part number 25-342 only.

^{*}Recertification services for BioTek Instruments, Inc., plate part number 7260522 (Absorbance Test Plate) are only available in Europe.



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