



LABORATORY DIAGNOSTICS

CATALOGUE

Doing things differently in medical diagnostics

UNA HEALTH

ABOUT US

Una Health Ltd is a leading UK distributor of diagnostics for the laboratory and point of care. Founded in 2009, we are an independent company with one key aim – to reshape the patient journey and improve overall healthcare.

We believe that better healthcare is about harnessing existing technology to elevate outcomes as well as utilising cutting-edge, innovative diagnostics. We supply specialist, innovative and appropriate healthcare solutions, backed by a team with exceptional levels of experience, knowledge and support.

Collaborating with global suppliers, we strive to fulfil the diverse needs of our customers. Our pioneering approach to customer care is rooted in principles of fairness, flexibility, and transparency.

Our comprehensive solutions span various healthcare areas, including Community Care, Diabetes Management, Cardiovascular, Oncology, Paediatrics, Respiratory, Urinalysis, AMR and laboratory diagnostics.

The exceptional customer support and comprehensive in-person or e-learning training (CPD accredited) differentiate us in the industry. Our products participate in numerous EQA schemes, ensuring the highest standards of quality and performance.

WHAT MAKES US DIFFERENT?

Our aim is not only to provide innovative, cost-effective pathology and point-of-care diagnostics to the UK healthcare sector, but also to explore new ways in which our products can be used to improve efficiency and patient outcomes. Our approach to customer care is equally ground-breaking, with fairness, flexibility and transparency at its heart.



We innovate the way we approach diagnostics with a focus on challenging traditional pathways.



We're adaptable, accessible and flexible to our customer needs; with specialist professionals to add value to your business.



Our friendly and positive attitude means we look for solutions to 'how we can' rather than 'why we can't'.



Communication is key here at Una and we work closely with customers and suppliers alike.



We have our customers at our heart and are small enough to care yet big enough to cope.

OUR PARTNERS

We take pride in our commitment to seamlessly connect with global suppliers, fostering partnerships and networks that deliver exceptional products and services and fulfil the diverse needs of our customers.



NG Biotech is an innovative biotech company based in France, developing and manufacturing novel in vitro diagnostic tools for Therapy Monitoring at the Point of Care. Founded by pioneers of the rapid test industry, NG Biotech has developed a proprietary patented immunoassay platform enabling the development of qualitative and multiplex point-of-care diagnostics.



TECHLAB® is a market leader of innovative, rapid, non-invasive diagnostic tests for gastrointestinal diseases. The company designs, develops, and manufactures enteric diagnostics that are distributed worldwide and hold ISO 13485 certification, MDSAP, and FDA registration.



Fortress Diagnostics® is a multi-award-winning global provider of in vitro diagnostics (IVDs). Fortress develop, manufacture and support an extensive portfolio of clinical diagnostic tests in the United Kingdom, providing solutions to immunology, haematology and serological laboratories in hospitals, medical centres, clinics, blood banks and research institutions.



Nal von Minden is a German company developing and manufacturing in-vitro diagnostics. 4 decades of research, development and experience have gone into their products and ensure their unique quality. Nal von Minden were among the first to offer COVID-19 antigen tests. More than 300 million tests have been sold since then.



KOVA International is a leading manufacturer of quality controls and consumables for the clinical diagnostic laboratory. The company's primary focus lies in producing urinalysis controls available in both lyophilised and ready-to-use liquid forms. With a US manufacturing base, the KOVA® brand is certified to ISO 13485:2003 and it is cGMP qualified.



Quantimetrix® designs, develops, and manufactures laboratory-quality products at their headquarters in California. Leaders in the field of liquid-stable quality control products, Quantimetrix pioneered these innovations more than 40 years ago. Their portfolio improves the efficiency and reliability of laboratory testing and patient care.



AUDIT® MicroControls™, a US-based manufacturer, offers a complete line of multi-level materials including serum and urine-based matrices. Audit MicroControls' calibration verification/linearity and quality control ranges cover areas including blood gas, cardiac, diabetes, general chemistry, immunoassay, therapeutic drug monitoring, urinalysis.

LABORATORY DIAGNOSTICS

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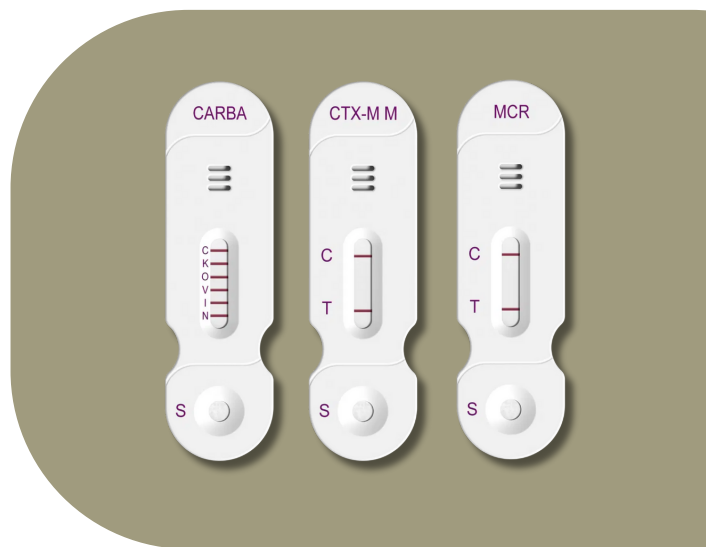


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NG BIOTECH LABORATOIRES

AMR RANGE



The rapid diagnostic tests from NG Biotech have been developed in collaboration with the French CEA, enabling fast, easy and accurate detection of antimicrobial resistance (AMR) in (multi) drug-resistant bacteria (MDR). These cutting-edge antimicrobial resistance tests enable rapid detection in just 15 minutes. Using patented technology for superior sensitivity and sensitivity, they provide results you can trust, verified by over 100 scientific publications.



FAST

Reduces wait times:
results in only 15 minutes



COMPREHENSIVE

Detects multiple mechanisms



EASY TO PERFORM

AMR detection direct from
bacterial cultures



ANTIMICROBIAL STEWARDSHIP

Ensures appropriate use of
tailored antibiotics

HOW IT WORKS

The NG-Test AMR rapid test is a visual, multiplex lateral flow immunochromatographic assay, developed using patented technology. Each single-use cassette offers rapid, qualitative detection and differentiation of different antimicrobial resistance mechanisms among non-susceptible colonies of Gram-negative bacteria.

The patented new multi-layer, multi-conjugate lateral flow platform utilised across the NG AMR product range increases the capacity of key biomarkers, whilst maintaining the high quality of detection. The NG-Test AMR platform is based on lateral flow immunochromatographic principles.

HOW IT HELPS

Rapid detection of antimicrobial resistant bacteria is paramount to a patient receiving appropriate treatment in a timely manner and is necessary to inform infection prevention actions. The NG Biotech lateral flow kits offer equivalent results to molecular detection methods in a fraction of the time, with no need for additional equipment and associated maintenance costs. Hands on time to prepare the test is minimal and results are available to read in 15 minutes.

CARBAPENEMASE DETECTION

NG-TEST® CARBA-5



NG-Test CARBA-5 is a visual multiplex immunochromatographic (lateral flow) qualitative assay for the detection and differentiation of the five most common carbapenemase families (KPC, OXA-48-like, VIM, IMP and NDM) from carbapenem non-susceptible pure bacterial colonies of Enterobacterales (including *Escherichia coli* and *Klebsiella pneumoniae*) and *Pseudomonas aeruginosa*. Results in 15 minutes.

Sensitivity: 100% Specificity: 100%

NG-Test® CARBA 5 detects the following variants:

- Type NDM: NDM-1 -2 -3 -4 -5 -6 -7 -8 -9 -11 -19
- Type KPC: KPC-1 -2 -3 -4 -5 -6 -7 -12 -14 -23 -28 -39
- Type IMP: IMP-1 -2 -4 -5 -6 -7 -8 -10 -11 -13 -14 -15 -16 -18 -19 -22 -26 -29 -31 -37 -39 -46 -47 -56 -58 -61 -63 -71 -79
- Type VIM: VIM-1 -2 -4 -5 -6 -19 -23 -26 -27 -31 -39 -46 -51 -52 -54 -56 -58 -59
- OXA-48-like: OXA-48 -162 -181 -204 -232 -244 -245 -370 -436 -484 -515 -517 -519 -535 -793

NGB-CAR-S23-021

20 tests/box

ESBL

NG-TEST® CTX-M MULTI



NG-Test CTX-M Multi detects the presence of the 5 major groups in the CTX-M-type enzymes of extended-spectrum beta-lactamases (ESBLs) produced by Enterobacteriaceae, from a bacterial colony. The Rapid Test detects enzymes belonging to CTX-M Groups 1, 2, 8, 9 and 25 including their most clinically relevant variants in the same cassette. Results in 15 minutes.

Sensitivity: 100% Specificity: 100%

NG-Test® CTX-M detects the following variants:

- Group 1: CTX-M-1 -3 -10 -15 -32 -37 -55 -57 -71 -82 -101 -182
- Group 2: CTX-M-2
- Group 8: CTX-M-8
- Group 9: CTX-M-9 -13 -14 -17 -18 -19 -24 -27 -38 -65 -93
- Group 25: CTX-M-94, -100

NGB-CTM-S23-016

20 tests/box

COLISTIN RESISTANCE

NG-TEST® MCR-1



NG-Test MCR-1 detects the presence of the MCR-1 enzyme responsible for Polymyxin E (colistin) resistance in Gram Negative bacteria, from a bacterial colony, in less than 15 minutes.

Sensitivity: 100% Specificity: 100%

NGB-MCR-S23-016

20 tests/box

ENTERIC DISEASES



TEHLAB®

TEHLAB® is a market leader of innovative, rapid, non-invasive diagnostic tests for gastrointestinal diseases. The TEHLAB® kits offer rapid laboratory detection of a range of faecal antigens and are the gold standard enzyme immunoassay (EIA) tests used to determine the common causes of diarrhoeal illness in different clinical scenarios.

Infectious causes:

- ***Clostridioides difficile*** GDH (glutamate dehydrogenase) and/or toxin A/B detection
- **Foodborne illness** – Shiga toxin-producing *Escherichia coli* (STEC), *Campylobacter* species, *Clostridium perfringens* enterotoxin detection
- **Faecal parasite detection** - *Giardia lamblia*, *Cryptosporidium*, *Entamoeba histolytica*
- ***Helicobacter pylori*** – a cause of gastritis, gastric ulcers or gastric cancer

Non-infectious causes:

- **Faecal lactoferrin** – for accurate differentiation between inflammatory bowel disease (IBD) and irritable bowel syndrome (IBS)

The TEHLAB® range of kits largely comprise two formats: CHEK™ and QUIK CHEK™.



CHEK™

96-well plate-based enzyme immunoassay (EIA)

- 96-well plate-based format
- Suitable for screening large numbers of samples
- Results within 2 hours
- Simple procedure
- Automatable
- Highly standardised



QUIK CHEK™

Single test membrane EIA technology in a cassette

- Direct faecal specimen testing in a rapid assay format
- Individual device
- Membrane bound EIA technology
- Suitable for smaller numbers of samples or for 'out-of-workflow' testing
- Results within 30 minutes
- Easy to interpret
- No equipment needed
- Highly specific and sensitive

CLOSTRIDIoidES DIFFICILE

Panel of in vitro diagnostics for detecting *C. difficile* and its toxins in faecal specimens from patients suspected of having the disease.

C. DIFF QUIK CHEK COMPLETE®	T30525C/T30550C	25 or 50 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: Toxins A&B GDH antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 30 min 			
C. DIFF QUIK CHEK®	30390	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: GDH antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 30 min 			
TOX A/B QUIK CHEK®	30394	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: Toxins A&B • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 30 min 			
C. DIFF CHEK™ - 60	TL5025/T5025B	96 tests	Microplate
<ul style="list-style-type: none"> • Analyte(s) Detected: GDH antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: < 1 hr, or 30 min (rapid format) 			
C. DIFFICILE TOX A/B II™	T5015/T5015B	96 tests	Microplate
<ul style="list-style-type: none"> • Analyte(s) Detected: Toxins A&B • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: < 1 hr, or 30 min (rapid format) 			
C. difficile Toxin/Antitoxin Kit	T5000	300-650 tests	Tissue Culture
<ul style="list-style-type: none"> • Analyte(s) Detected: Toxin B • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: 18 hrs 			
C. DIFFICILE TOX-B TEST	T5003	96 tests	Tissue Culture
<ul style="list-style-type: none"> • Analyte(s) Detected: Toxin B • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: 24-48 hrs 			

FOODBORNE PATHOGENS

Shiga toxin-producing *Escherichia coli* (STEC), *Campylobacter* species, *Clostridium perfringens* enterotoxin detection.

CAMPYLOBACTER CHEK™	T31096	96 tests	Microplate
<ul style="list-style-type: none"> • Analyte(s) Detected: C. jejuni, C. coli, C. lari, & C. upsaliensis • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: 1 hr 			
CAMPYLOBACTER QUIK CHEK™	T31025	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: C. jejuni, C. coli, C. lari, & C. upsaliensis • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 30 min 			
SHIGA TOXIN QUIK CHEK™	T30625	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: Shiga Toxin 1 and Shiga Toxin 2 • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S / Broth (GN or MAC) /Plate Culture (SMAC,CT-SMAC,CHROMagar@O157) • Time to Result: < 30 min 			
SHIGA TOXIN CHEK™	T30696	96 tests	Microplate
<ul style="list-style-type: none"> • Analyte(s) Detected: Shiga Toxin 1 and Shiga Toxin 2 • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S / Broth (GN or MAC) /Plate Culture (SMAC,CT-SMAC,CHROMagar@O157) • Time to Result: 50 min, or 20 min (rapid format) 			
Clostridium Perfringens Enterotoxin Test	T5006	96 tests	Microplate
<ul style="list-style-type: none"> • Analyte(s) Detected: Clostridium perfringens Enterotoxin • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: < 2.5 hrs 			

H. PYLORI

Diagnostic assays for detecting *H. pylori* in faecal specimens.

H. PYLORI CHEK™	T5051	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: Helicobacter pylori stool antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) up to 96 hours / Cary Blair / C&S • Time to Result: 1 hr 			
H. PYLORI QUIK CHEK™	30925	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: Helicobacter pylori stool antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) up to 96 hours / Cary Blair / C&S • Time to Result: < 30 min 			

PROTOZOAN PARASITES

Diagnostic tests for common intestinal parasites: Giardia, Cryptosporidium, Entamoeba histolytica.

TRI-COMBO PARASITE SCREEN	T30408	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: Giardia cyst antigen Cryptosporidium oocyst antigen E. histolytica antigen (adhesin) • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 2 hrs at RT 			
GIARDIA/CRYPTOSPORIDIUM CHEK®	30401	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: Giardia cyst antigen Cryptosporidium oocyst antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / 10% Formalin / SAF • Time to Result: < 2 hrs at RT 			
GIARDIA II™	PT5012	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: Giardia cyst antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / 10% Formalin / SAF • Time to Result: < 2 hrs at RT 			
CRYPTOSPORIDIUM II™	30406	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: Cryptosporidium oocyst antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / 10% Formalin / SAF / Cary Blair / C&S • Time to Result: < 2 hrs at RT 			
E. HISTOLYTICA II™	T5017	96 tests	Microplate ELISA
<ul style="list-style-type: none"> • Analyte(s) Detected: E. histolytica antigen (adhesin) • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) • Time to Result: < 2 hrs at RT 			
GIARDIA/CRYPTOSPORIDIUM QUIK CHEK™	T30407	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: Giardia cyst antigen Cryptosporidium oocyst antigen • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / 10% Formalin / SAF / Cary Blair / C&S • Time to Result: < 30 min at RT 			
E. HISTOLYTICA QUIK CHEK™	T30409	25 tests	Rapid EIA
<ul style="list-style-type: none"> • Analyte(s) Detected: E. histolytica antigen (adhesin) • Faecal Sample Type: Fresh & frozen faecal sample (unpreserved) / Cary Blair / C&S • Time to Result: < 30 min at RT 			

FAECAL LEUKOCYTE SCREEN

The LEUKO EZ VUE® is an immunochromatographic test for the qualitative detection of elevated levels of faecal lactoferrin, in liquid, semi-liquid, and solid faecal specimens.

LEUKO EZ VUE®

T30355

25 tests

Lateral Flow Qualitative

- **Analyte(s) Detected:** Lactoferrin—marker for faecal leukocytes & indicator for intestinal inflammation that can be caused by enteric infections
- **Faecal Sample Type:** Fresh samples (undiluted, unpreserved) stored between 2°C - 8°C or room temperature for up to 2 weeks, then stored frozen
- **Time to Result:** 10 min

INTESTINAL INFLAMMATION

Diagnostic tests for detection of lactoferrin and other markers of inflammation in the bowels.

LACTOFERRIN SCAN®

T5009

96 tests

Microplate ELISA Quantitative

- **Analyte(s) Detected:** Lactoferrin—marker for faecal leukocytes & indicator for intestinal inflammation; aids in the diagnosis of IBD/IBS populations
- **Faecal Sample Type:** Fresh samples (undiluted, unpreserved) stored between 2°C - 8°C or room temperature for up to 2 weeks, then stored frozen
- **Time to Result:** approx 75 min

LACTOFERRIN CHEK®

T5008

96 tests

Microplate ELISA Qualitative

- **Analyte(s) Detected:** Lactoferrin—marker for faecal leukocytes & indicator for intestinal inflammation; aids in the diagnosis of IBD/IBS populations
- **Faecal Sample Type:** Fresh samples (undiluted, unpreserved) stored between 2°C - 8°C or room temperature for up to 2 weeks, then stored frozen
- **Time to Result:** approx 75 min

LACTOFERRIN EZ VUE®

T5018

25 tests

Lateral Flow Qualitative

- **Analyte(s) Detected:** Lactoferrin—marker for faecal leukocytes & indicator for intestinal inflammation; aids in the diagnosis of IBD/IBS populations
- **Faecal Sample Type:** Fresh samples (undiluted, unpreserved) stored between 2°C - 8°C or room temperature for up to 2 weeks, then stored frozen
- **Time to Result:** 10 min

ASCA-CHEK™

T5016

96 tests

Microplate ELISA Qualitative

- **Analyte(s) Detected:** Detects anti-*S. cerevisiae* antibodies; aids in the diagnosis of Crohn's disease
- **Faecal Sample Type:** Fresh and frozen faecal samples (unpreserved; faecal specimens should be frozen if not tested within 48 hours) / Serum (freeze if not tested within 7 days)
- **Time to Result:** approx 75 min

FORTRESS® DIAGNOSTICS

Fortress Diagnostics® are a multi award-winning global provider of in vitro diagnostics (IVDs). Fortress develop, manufacture and support an extensive portfolio of clinical diagnostic tests in the United Kingdom. Our ongoing relationship means we are able to provide highly accurate medical testing solutions to immunology, haematology and serological laboratories in hospitals, medical centres, clinics, blood banks and research institutions.

FORTRESS® DIAGNOSTICS

HELICOBACTER PYLORI

The Fortress Diagnostics Helicobacter pylori kits are intended for use in the quantitative determination of Anti H pylori specific antibodies of IgA, IgG and IgM type in human serum or plasma by Microplate Enzyme Immunoassay.

Helicobacter Pylori, IgA	BXE0672A	96 tests	ELISA
Helicobacter Pylori, IgG	BXE0673A	96 tests	ELISA
Helicobacter Pylori, IgM	BXE0674A	96 tests	ELISA

FORTRESS® DIAGNOSTICS

SEXUALLY TRANSMITTED DISEASES



Fortress Diagnostics® are a multi award-winning global provider of in vitro diagnostics (IVDs). Fortress develop, manufacture and support an extensive portfolio of clinical diagnostic tests in the United Kingdom. Our ongoing relationship means we are able to provide highly accurate medical testing solutions to immunology, haematology and serological laboratories in hospitals, medical centres, clinics, blood banks and research institutions.



RELIABLE

A trusted provider in the UK



RESULTS

High sensitivity & specificity



METHOD

Lateral flow and EIA kits available



QUALITY

High-quality testing methods

HOW IT WORKS

All Fortress Diagnostics products are highly accurate and are available with a long shelf life. Suitable for preliminary or emergency medical screening for use in medical facilities with limited resources and laboratories with low tests throughput.

- High quality & easy-to-use.
- Quick results – 10 minutes to 2 hours – providing timely treatment interventions.
- Little or no additional equipment required.
- Possibility to store at room temperature for extended length of time.
- All CE marked

Apart from the methods presented in this section, we also provide rapid tests for sexually transmitted diseases, please refer to the Rapid Tests section of this catalogue.

SYPHILIS TESTING SOLUTIONS

Treponema Pallidum (TP) is the causative agent of the venereal disease Syphilis. TP is a spirochete bacterium with an outer envelope and a cytoplasmic membrane. Relatively little is known about the organism in comparison with other bacterial pathogens. According to the Center for Disease Control (CDC), the number of cases of Syphilis infection has markedly increased since 1985. Some studies have reported a substantial epidemiological correlation between the acquisition and transmission of the HIV virus and Syphilis.

HAEMAGGLUTINATION METHOD

Sensitive and specific indirect haemagglutination tests for the detection of antibodies to Treponema Pallidum.

TPHA (with +ve and -ve controls)	SYTP0100	100 tests	Haemagglutination test
TPHA (with +ve and -ve controls)	SYTP0200	200 tests	Haemagglutination test

SEROLOGICAL METHOD

Rapid Plasma Reagin or RPR Card test is a non-treponemal method for the serological detection of syphilis. The antigen – a particulate carbon suspension coated with lipid complexes – agglutinates in the presence of serum reagins. Reagins are antibodies present in the sera of syphilitic patients. Visible agglutination in the form of black clumps which can be viewed macroscopically, indicates the presence of such antibodies in the sample tested.

RPR Positive & Negative Control, Pipette/Stirrers, Test Cards, Dispensing Bottle and Needle	SYRPR050	50 tests	Non-treponemal method for serological detection
RPR Positive & Negative Control, Pipette/Stirrers, Test Cards, Dispensing Bottle and Needle	SYRPR100	100 tests	Non-treponemal method for serological detection
RPR Positive & Negative Control, Pipette/Stirrers, Test Cards, Dispensing Bottle and Needle	SYRPR500	500 tests	Non-treponemal method for serological detection
RPR (Reagent only)	SYCA0002	2ml (100T)	Reagent
RPR (Reagent only)	SYCA0005	5ml (250T)	Reagent
RPR (Reagent only)	SYCA0010	10ml (500T)	Reagent
RPR (Reagent only)	SYCA0100	100ml (5000T)	Reagent
RPR (Reagent only)	SYCA1000	1000ml (50000T)	Reagent

CONTROLS & CALIBRATORS

The Syphilis Control set is designed for the validation of the Fortress range of Syphilis Test Kits. It is recommended that a positive and negative control be included with each run of tests carried out.

Syphilis Control (Positive & Negative)	SYPN0010	2 x 5 x 1ml	Control
Syphilis Control Panel Level 1 - 6	BXC0806A	6 x 1 x 0.5ml	Control panel

SYPHILIS ELISA

The detection of anti-Treponema Pallidum (anti-TP) antibodies is achieved by antigen sandwich enzyme linked immunosorbent assay, where the microwells are coated with recombinant Treponema pallidum antigens expressed in E.coli. In vitro diagnostic kit for the detection of antibodies to Treponema pallidum in human serum or plasma. It's intended for use in the screening of blood donors and to aid in the diagnosis and management of clinical conditions of syphilis.

Syphilis	BXE0995A	96T	ELISA
Syphilis	BXE0995C	480T	ELISA

SEXUALLY TRANSMITTED DISEASES

HERPES SIMPLEX VIRUS (HSV)

The Fortress Herpes Simplex Virus kits are enzyme immunoassays intended for the qualitative detection of IgG and IgM antibodies to HSV-I, HSV-II and HSV-1/2 in human serum or plasma. It is intended for screening and as an aid in the diagnosis of possible HSV infection.

Herpes Simplex Virus (HSV-1/2)IgG	BXE0621A	96T	ELISA
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FORTRESS® DIAGNOSTICS

HEPATITIS

Fortress Diagnostics® are a multi award-winning global provider of in vitro diagnostics (IVDs). Fortress develop, manufacture and support an extensive portfolio of clinical diagnostic tests in the United Kingdom. Our ongoing relationship means we are able to provide highly accurate medical testing solutions to immunology, haematology and serological laboratories in hospitals, medical centres, clinics, blood banks and research institutions.

HEPATITIS SCREENING

HEPATITIS E

The Fortress Diagnostics Hepatitis E (HEV) kits are enzyme linked immunosorbent assays (ELISA) intended for the qualitative detection of Hepatitis E virus antigen, in human or plasma specimens. It is intended for use in clinical laboratories for the diagnosis and management of patients related to infection with hepatitis E Virus.

HEV Ab	BXE0745A	96T	ELISA
HEV Ag	BXE0903A	96T	ELISA
HEV IgG	BXE0901A	96T	ELISA
HEV IgM	BXE0902A	96T	ELISA

Apart from the methods presented in this section, we also provide rapid tests for hepatitis screening, please refer to the Rapid Tests section of this catalogue.

RAPID TESTS



FORTRESS® DIAGNOSTICS

Our rapid diagnostic tests range includes suitable for preliminary or emergency medical screening for use in medical facilities with limited resources and laboratories with low number of tests per day.



RELIABLE

A trusted provider in the UK



RESULTS

High sensitivity & specificity



METHOD

Lateral flow and EIA kits available



QUALITY

High-quality testing methods

DENGUE FEVER

Dengue viruses, transmitted by the *Aedes aegypti* and *Aedes albopictus* mosquitoes, are widely distributed throughout the tropical and subtropical areas of the world. Fortress Diagnostics have a Dengue NS1 Antigen Rapid Test Device for the detection of dengue virus NS1 in antigen in whole blood, serum or plasma and a Dengue IgG.IgM rapid test device for the qualitative detection of IgG and IgM antibodies to dengue virus.

Dengue IgG/IgM NS1 Test Device	DMNS1020	20T	Whole Blood/Serum/Plasma
Dengue IgG/IgM Test Device	DNGMC020	20T	Serum/Plasma/Whole Blood, Cut off 500ng/ml
Dengue Fever IgG/IgM Antibody Test	DNGMC040	40T	Whole Blood/Serum/Plasma

FAECAL OCCULT BLOOD TEST

The Fortress Diagnostics Faecal Occult Blood (FOB) One Step Test Device (faeces sample) is a rapid test to qualitatively detect low levels of faecal occult blood in faeces. The test uses double antibody sandwich assay to selectively detect as low as 50 ng/mL of hemoglobin or 6 µg hemoglobin/g faeces. In addition, unlike the guaiac assays, the accuracy of the test is not affected by the diet of the patients.

Faecal Occult Blood Test Device	FOBC0020	20T	Cut off 30 ng/ml - (Faeces)
Faecal Occult Blood Test Device	FOBC0040	40T	Cut off 30 ng/ml - (Faeces)

HEPATITIS

Fortress Diagnostics offers lateral flow chromatographic immunoassays for the qualitative detection of Hepatitis B surface antigen (HBsAg), anti-HBs and Hepatitis C antibodies in human whole blood, serum or plasma.

HBsAg Rapid Test	HBSCWB20	20T	Whole Blood/Serum/Plasma
HBsAg Rapid Test	HBSCWB40	40T	Whole Blood/Serum/Plasma
HBsAg Rapid Test	HBSWB050	50T	Whole Blood/Serum/Plasma
HBsAg Rapid Test	HBSWB100	100T	Whole Blood/Serum/Plasma
Anti-HBs Rapid Test	HBSAB050	50T	Whole Blood/Serum/Plasma
Anti-HBs Rapid Test	HBSAB100	100T	Whole Blood/Serum/Plasma
HCV Rapid Test	HCVC0020	20T	Whole Blood/Serum/Plasma
HCV Rapid Test	HCVC0040	40T	Whole Blood/Serum/Plasma
HCV Rapid Test	HCVS0050	50T	Whole Blood/Serum/Plasma
HCV Rapid Test	HCVS0100	100T	Whole Blood/Serum/Plasma

GASTROINTESTINAL

HELICOBACTER PYLORI

The Fortress Diagnostics H.pylori Ab Rapid Test is a sandwich lateral flow chromatographic immunoassay for the qualitative detection of antibodies (IgG, IgM, and IgA) anti-Helicobacter pylori in human whole blood, serum or plasma.

Helicobacter Pylori Test Strip	HPS00050	50T	Serum/Plasma
Helicobacter Pylori Ag	HCVC0040	40T	Whole Blood/Serum/Plasma
Helicobacter Pylori Ag	HCVS0050	50T	Whole Blood/Serum/Plasma

SALMONELLA TYPHI

S. typhi Antigen	STAGC040	40T	Serum/Plasma/Faeces
S. typhi Antigen	TPC00020	20T	Serum/Plasma/Faeces

SEXUALLY TRANSMITTED DISEASES

SYPHILIS

The Fortress Diagnostics Syphilis Ab Rapid Test Cassette (Whole Blood/Serum/Plasma) is a rapid, serological, immunochromatographic assay for the qualitative detection of antibodies (IgG, IgM and IgA) to Treponema Pallidum (TP) in human whole blood, serum or plasma.

Syphilis Device	TPC00040	40T	Serum/Plasma/Whole Blood
Syphilis Test Strips	TPS00050	50T	Serum/Plasma/Whole Blood
Syphilis Test Strips	TPS00100	100T	Serum/Plasma/Whole Blood

GONORRHEA

Gonorrhoea	GONC0020	20T	Swab
Gonorrhoea	GONC0040	40T	Swab

PREGNANCY TESTING

The Fortress Diagnostics hCG One Step Pregnancy Test Device (Urine/Serum) is a rapid chromatographic immunoassay for the qualitative detection of human chorionic gonadotropin in urine or serum to aid in the early detection of pregnancy.

hCG Test Device	HCGCSU20	20T	Sensitivity 25IU/ml - Serum/Urine, including disposable pipette
hCG Test Device	HCGC0040	40T	Sensitivity 25IU/ml - Urine Only, including disposable pipette
hCG Test Device	HCGC0050	50T	Sensitivity 25IU/ml - Urine Only, including disposable pipette
hCG Test Strip	HCGSU100	100T	Sensitivity 25IU/ml - Serum/Urine, including disposable pipette
hCG Test Strips (Urine Only)	HCGS0050	50T	Sensitivity 25IU/ml - Urine -2.5mm
hCG Test Strips (Urine Only)	HCGS0100	100T	Sensitivity 25IU/ml - Urine -2.5mm
HCG Rapid Test	HCGCSU20	20T	Serum/Urine
HCG Rapid Test	HCGCSU40	40T	Serum/Urine

MALARIA

The Fortress Diagnostics Malaria Pf Ag Rapid Test is a rapid lateral flow chromatographic immunoassay for the detection of Malaria P.falciparum specific histidine rich protein-2 (Pf HRP-II) in human blood specimen as an aid in the diagnosis of Malaria infection. This test is intended for In-Vitro Diagnostic use only.

Malaria Device P.falciparum	PFC00020	20T	Serum/Plasma/Whole Blood, with Test Tube & Buffer
Malaria Device P.falciparum	PFC00040	40T	Serum/Plasma/Whole Blood, with Test Tube & Buffer
Malaria P.falciparum/P. Vivax (Pan Malaria Device)	PVC00020	20T	Serum/Plasma/Whole Blood, with Test Tube & Buffer
Malaria P.falciparum/P. Vivax (Pan Malaria Device)	PVC00040	40T	Serum/Plasma/Whole Blood, with Test Tube & Buffer

MULTI-DRUG

The Fortress Diagnostics One Step Multi-Drug Screen Test Dipcard is a lateral flow chromatographic immunoassay for the qualitative detection of multiple drugs and drug metabolites in urine at the following cut-off concentrations in urine. The speed and sensitivity of immunoassays have made them the most widely accepted method to screen urine for multiple drugs of abuse.

Multi-drug - 6 Parameter Device	DOAC0006	20T	Test Device for multi-drug detection - Urine. Many combinations available
Multi-drug - 10 Parameter Device	DOAC0010	20T	Test Device for multi-drug detection- Urine- COC, AMP, MET, THC, MTD, MDMA, OPI, PCP, BAR, BZO
Multi-drug - 12 Parameter Device	DOAC0012	20T	Test Device for multi-drug detection - Urine - COC, AMP, TCA, MOP, MET, THC, MTD, MDMA, OPI, PCP, BAR, BZO

TROPONIN

The Fortress Diagnostics cTnI One Step Troponin I Test Device (Whole Blood/Serum/Plasma) is a simple test that utilises a combination of anti-cTnI antibody-coated particles and capture reagent to selectively detect cTnI in whole blood, serum or plasma. The minimum detection level is 0.5 ng/mL.

Troponin-I Device	TNIC0020	20T	Serum/Plasma/Whole Blood, Cut off 1 ng/ml
Troponin-I Device	TNIC0040	40T	Serum/Plasma/Whole Blood, Cut off 1 ng/ml
CK-MB, Troponin-I, Myoglobin	CTMC0020	20T	Whole Blood, Cut off CK-MB 5ng/ml, Tni-I 1ng/ml, Myo 80ng/ml

TUBERCULOSIS

The Fortress Diagnostics Tuberculosis IgG/IgM Rapid Test Device (Whole Blood/Serum/Plasma) is a membrane-based screening test for the rapid detection of IgM anti-Mycobacterium Tuberculosis and IgG anti-Mycobacterium Tuberculosis in human whole blood, serum or plasma. This innovative rapid screening test is based on lateral flow immunochromatography and is among the easiest point of care (POC) assay diagnostics.

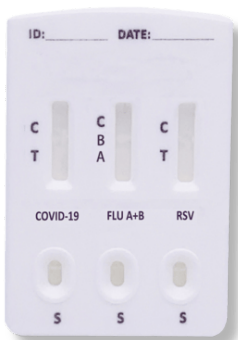
Tuberculosis Device	TBC00020	20T	Whole Blood / Serum / Plasma
Tuberculosis Device	TBC00040	40T	Whole Blood / Serum / Plasma

COVID-19

Covid-19 Total Ab Device	COVID010	10T	Whole Blood / Serum / Plasma
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ALLTEST

COVID-19/INFLUENZA A+B/RSV



Prepare for the winter season with the Beright SARS-CoV-2/Influenza A+B/RSV Antigen Combo Rapid Lateral Flow Test, a CTDA approved rapid chromatographic immunoassay for the qualitative detection of COVID-19, Influenza A, Influenza B and Respiratory Syncytial Virus (RSV). Save time and money with the detection of the four main respiratory viruses in a single test.

- SARS-CoV-2: Sensitivity: 97.0% Specificity: 99.0%
- Flu A: Sensitivity: 95.0% Specificity: 99.1%
- Flu B: Sensitivity: 92.9% Specificity: 99.1%
- RSV: Sensitivity: 94.3% Specificity: 96.2%

SARS-CoV-2/Influenza A+B/RSV Antigen Combo Test (from Beright)	ISIR-535	20T	Nasal or Nasopharyngeal Sample
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RAPID TESTS

NAL VON MINDEN



The Nal Von Minden GmbH NADAL® rapid tests cover a wide range of clinical conditions for both point-of-care settings and laboratories, where quick and accurate diagnosis can make all the difference. They are suitable in emergency medical screening or medical facilities with limited resources and laboratories with low test throughput. Whether diagnosing an infectious disease or cancer, right through to a heart attack, the Nal Von Minden rapid tests deliver certainty and peace of mind.

All tests are CE marked, have high sensitivity and specificity, confirm to IVDR regulations and are available with a long shelf life. They further expand Una Health's solutions to cover almost every branch of medicine.



FAST

Reduces wait times; results available in minutes



RESULTS

High sensitivity & specificity



COMPREHENSIVE

Extensive range of tests



FLEXIBILITY

Suitable for almost every branch of medicine

HOW IT WORKS

Our complete range of rapid tests are designed to give quick and reliable answers on the spot. Early diagnosis allows treatment to be introduced faster and generally reduces the duration and severity of illnesses. In addition, accurate and timely diagnosis can help reduce unnecessary use of antibiotics and supports antimicrobial stewardship alongside reducing the need for costly, unnecessary and uncomfortable lab examinations.

INFECTIOUS DISEASES

ESCHERICHIA COLI O157

The NADAL® E. coli O157 Test is a rapid chromatographic immunoassay for the qualitative detection of Escherichia coli (E. coli) O157 antigens in human faecal specimens. The test is intended for use as an aid in the diagnosis of an E. coli infection and is designed for professional use only.

E.Coli O157 Rapid Tests - Cassette

501006

10T

Faecal specimens

LEGIONELLA

The NADAL® Legionella Test is an in-vitro rapid chromatographic lateral flow immunoassay for the qualitative detection of Legionella pneumophila (L. pneumophila) serogroup 1 antigen in urine specimens from patients with symptoms of pneumonia. The NADAL® Legionella Test is intended to be used in conjunction with culture and other methods as an aid in the presumptive diagnosis of Legionella infection (Legionnaires' disease) caused by L. pneumophila serogroup 1.

Legionella rapid test

552020

10T

Urine

HEPATITIS

The NADAL® HAV IgG/IgM Test is a lateral flow chromatographic immunoassay for the simultaneous detection and differentiation of anti-hepatitis A virus (HAV) IgG and IgM in human serum, plasma or whole blood. The test is intended to be used by professionals as a screening test and as an aid in the diagnosis of a HAV infection. Any reactive result with the NADAL® HAV IgG/IgM Test must be confirmed with alternative testing method(s) and clinical findings.

HAV IgG/IgM test cassette

622070N-30

30T

Serum, Whole Blood, Plasma

The Hepatitis B Virus Surface Antibody Test is a rapid chromatographic immunoassay for the qualitative detection of antibodies to Hepatitis B virus surface antigen (HBsAg) in human whole blood, serum, plasma.

INFO® anti-HBs (HBsAb) (CE1434) test cassette

622091

40T

Serum, Whole Blood, Plasma

GROUP B STREP

The NADAL® Strep B Test is a rapid, visual immunoassay for the qualitative, presumptive detection of group B Streptococcus (GBS) antigens in specimens collected from vaginal or rectal swab, as well as swabs taken from the ear or throat in newborns

NADAL® Strep B Test

232001

20T

Vaginal, rectal, ear and throat swabs

TRICHOMONAS VAGINALIS

The NADAL® Trichomonas vaginalis Test is a simple, one step chromatographic immunoassay for the rapid, qualitative detection of Trichomonas vaginalis in vaginal swab specimens.

This test is intended for use as an aid in the diagnosis of trichomoniasis and designed for professional use only.

NADAL® Trichomonas vaginalis, test cassette

840003N-10

10T

Vaginal swabs

MYCOBACTERIUM TUBERCULOSIS

The NADAL® Tuberculosis IgG/IgM Test is a lateral flow chromatographic immunoassay for the simultaneous detection and differentiation of anti-Mycobacterium Tuberculosis (M.Tb) IgM and IgG in human serum, plasma or whole blood. The test is intended to be used as a screening test and as an aid in the diagnosis of infection with M.Tb. Any reactive specimen with the NADAL® Tuberculosis IgG/IgM Test should be confirmed with alternative testing method(s) and clinical findings.

NADAL® Tuberculosis IgG/IgM, test cassette

322003N-30

30T

Human serum, plasma or whole blood

STREPTOCOCCUS PNEUMONIAE

The NADAL® S. pneumoniae Test is a qualitative rapid assay which is intended to be used for the detection of Streptococcus pneumoniae antigen in urine without any dilution and as an aid in the diagnosis of pneumonia, meningitis and otitis media.

Streptococcus pneumoniae Rapid Test

572004N-10

10T

Urine

CARDIAC RELATED

The NADAL® Cardiac Combo Test is a rapid visual immunoassay for the qualitative presumptive detection of human Myoglobin, CK-MB and cardiac Troponin I in whole blood, serum or plasma. The test is intended for use as an aid in the diagnosis of myocardial infarction (MI).

Cardiac Combo Cassette tests (Myoglobin, CK-MB and cardiac Troponin I)

282003

5T

Serum, Whole Blood, Plasma

D-DIMER

The NADAL® D-Dimer Test is used for the qualitative detection of D-Dimer in human whole blood and plasma. The test is used as an aid in the assessment and evaluation of patients with suspected disseminated intravascular coagulation (DIC), deep vein thrombosis (DVT) and pulmonary embolism (PE).

Nadal® D-Dimer Test

351006N-10

10T

Whole Blood, Plasma

Nadal® D-Dimer Test

351006N-05

5T

Whole Blood, Plasma

CALPROTECTIN

The NADAL® Calprotectin Test is a rapid chromatographic immunoassay (non-invasive assay) for the qualitative detection of calprotectin (hCP) in human faecal specimens. The test is intended for use as an aid in the diagnosis of inflammatory gastrointestinal disorders and is designed for professional use only.

NADAL® Calprotectin, test cassette

1212001

10T

Faeces

CRP

The NADAL® CRP Test is a lateral flow chromatographic immunoassay for the semiquantitative detection of C-reactive protein (CRP) in human whole blood, serum or plasma specimens. The test is intended for use as an aid in the diagnosis of bacterial infectious diseases and inflammatory processes. Due to a large number of possible symptoms, the test is not restricted to a defined target patient group and can generally be used to differentiate between bacterial and viral infections or to assess the presence or severity of inflammatory processes. The test procedure is not automated and requires no special training or qualification. The NADAL® CRP Test is designed for professional use only.

NADAL® CRP, test cassette

311801N-20

20T

Serum, Whole Blood, Plasma

PROSTATE SPECIFIC ANTIGEN

The NADAL® PSA Test is a rapid visual immunoassay for the semiquantitative presumptive detection of prostate-specific antigen (PSA) in human serum, plasma or whole blood specimens with a cut-off of 4 ng/mL. The test detects total PSA (tPSA). The NADAL® PSA Test is intended for use as an aid in the diagnosis of prostate cancer by professional users as elevated PSA levels frequently indicate an increased risk of prostate carcinomas.

NADAL® PSA, test cassette

602003

20T

Serum, Whole Blood, Plasma

CHLAMYDIA

The NADAL® Chlamydia Test (swab/urine) is a rapid visual immunoassay for the qualitative presumptive detection of Chlamydia trachomatis in female cervical swab, male urethral swab and male urine specimens. This test is intended for use as an aid in the diagnosis of Chlamydia infection. The NADAL® Chlamydia Test is designed for professional use only.

NADAL® Chlamydia (CE0197), test cassette

212007

20T

Urine, Urethral Swab, Cervical Swab

ENTEROVIRUS

The NADAL® Enterovirus Test is a rapid chromatographic immunoassay for the qualitative detection of Enterovirus antigens (VP1 peptide) in human faecal specimens. The test is intended for use as an aid in the diagnosis of Enterovirus infection and is designed for professional use only.

NADAL® Enterovirus, test cassette

1222001

10T

Faeces

ANTI-HERPES SIMPLEX VIRUS

The NADAL® HSV 2 IgG/IgM Test is a lateral flow chromatographic immunoassay for the qualitative detection and differentiation of anti-herpes simplex virus-2 (HSV-2) IgG and IgM in human serum, plasma or whole blood. The test is intended to be used by professionals as a screening test and as an aid in the diagnosis of infection with HSV-2. Any reactive result with the NADAL® HSV 2 IgG/IgM Test must be confirmed with alternative testing method(s) and clinical findings.

NADAL® HSV 2 IgG/IgM, test cassette

1130005N-10

10T

Serum, Whole Blood, Plasma

The NADAL® HSV 1 IgG/IgM Test is a lateral flow chromatographic immunoassay for the qualitative detection and differentiation of anti-herpes simplex virus-1 (HSV-1) IgG and IgM in human serum, plasma or whole blood. The test is intended to be used by professionals as a screening test and as an aid in the diagnosis of infection with HSV-1. Any reactive result with the NADAL® HSV-1 IgG/IgM Test must be confirmed with alternative testing method(s) and clinical findings.

NADAL® HSV 1 IgG/IgM, test cassette

1130004N-10

10T

Serum, Whole Blood, Plasma

MONONUCLEOSIS

The NADAL® Mononucleosis Test is a fast, chromatographic immunoassay for the qualitative detection of heterophile antibodies against Epstein-Barr-Virus (EBV) in human whole blood, serum or plasma. The test is intended as an aid in the diagnosis of infectious mononucleosis (IM) and is for professional use only.

NADAL® Mononucleosis test

252003N-20

20T

Serum, Whole Blood, Plasma

SALMONELLA

The NADAL® Salmonella spp. Test is a rapid chromatographic immunoassay for the qualitative detection of Salmonella antigens in human faecal specimens. The test is intended for use as an aid in the diagnosis of salmonellosis. The test is designed for professional use only.

NADAL® Salmonella spp., test cassette

1242001

10T

Faeces

FORTRESS® DIAGNOSTICS

FEBRILE ANTIGENS



Febrile Antigens are stained bacterial antigen suspensions used to identify and measure antibodies, following infection. When the test serum sample is mixed with a febrile antigen, the solution will agglutinate if antibodies are present, thus indicating infection is present. The Fortress Diagnostics' range of febrile antigens can be used either as a screening test or as a confirmatory test.

BRUCELLA

The Fortress Diagnostics Brucella agglutination test is a serological test for Brucellosis. Specific antibodies to the Brucella species are detectable a few weeks after exposure. Specific antibodies to Brucella antigens if present in serum will react with the antigen suspension to produce an agglutination reaction. No agglutination indicates the absence of the specific antigens.

Brucella Abortus	FEBBAB05	1 x 5ml	5ml Stained Antigen Suspension
Brucella Abortus	FEBBAB1L	1000ml	1000ml Stained Antigen Suspension
Brucella Melitensis	FEBBME05	1 x 5ml	5ml Stained Antigen Suspension
Brucella Melitensis	FEBBME1L	1000ml	1000ml Stained Antigen Suspension
Brucella Abortus, Melitensis + Positive Control	FEBAMP05	2 x 1 x 5ml	Stained Antigen Suspension plus 1x0.5ml Positive control

FEBRILE ANTIGEN (WIDAL)

The Fortress Diagnostics Febrile Antigen (Widal) test is a common agglutination test employed in the serological diagnosis of enteric fever and helps to detect the presence of Salmonella antigens in a patient's serum. Antibodies in serum produced in response to exposure to Salmonella organisms will agglutinate bacterial suspension which carries homologous antigens.

Febrile Antigen Kit (Widal)	FEBNC100	8 x 5ml	Stained Salmonella Antigens, 100T per Antigen WITHOUT CONTROLS
Febrile Antigen Kit (Widal)	FEBWC100	8 x 5ml/2 x 1ml	Stained Salmonella Antigens, 100T per Antigen WITH CONTROLS
Febrile Negative Control	FEBNCO01	1 x 1ml	1ml polyvalent negative
Febrile Positive Control	FEBPCO01	1 x 1ml	1ml polyvalent positive

PROTEUS

Antibodies produced against rickettsial antigen cross reacts with OX19 and OX2 strains of *Proteus vulgaris* and OXK strains of *Proteus mirabilis*. The Fortress Diagnostics Proteus OX19 stained febrile antigen suspension can be used to identify and quantitate specific antibodies in human sera following infection with certain Rickettsiae pathogens. *Proteus* OX19 reacts strongly with the sera of patients with typhus group rickettsiae and rocky mountain of spotted fever.

Brucella Abortus	FEBBAB05	1 x 5ml	5ml Stained Antigen Suspension
Brucella Abortus	FEBBAB1L	1000ml	1000ml Stained Antigen Suspension
Brucella Melitensis	FEBBME05	1 x 5ml	5ml Stained Antigen Suspension

ROSE BENGAL

The Fortress Diagnostics Febrile Antigen (Widal) test is a common agglutination test employed in the serological diagnosis of enteric fever and helps to detect the presence of *Salmonella* antigens in a patient's serum. Antibodies in serum produced in response to exposure to *Salmonella* organisms will agglutinate bacterial suspension which carries homologous antigens.

Febrile Antigen Kit (Widal)	FEBNC100	8 x 5ml	Stained Salmonella Antigens, 100T per Antigen WITHOUT CONTROLS
Febrile Antigen Kit (Widal)	FEBWC100	8 x 5ml/2 x 1ml	Stained Salmonella Antigens, 100T per Antigen WITH CONTROLS
Febrile Negative Control	FEBNCO01	1 x 1ml	1ml polyvalent negative
Febrile Positive Control	FEBPCO01	1 x 1ml	1ml polyvalent positive

SALMONELLA

The Fortress Diagnostics have developed *Salmonella* Paratyphi tests for *Salmonella* Paratyphi A-H, A-O, B-H, B-O, C-H and C-O. The Febrile Antigen Widal test is a common agglutination test employed in the serological diagnosis of enteric fever and helps to detect the presence of *Salmonella* antigens in a patient's serum.

Salmonella paratyphi A-H	FEBSAH05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella paratyphi A-O	FEBSAO05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella paratyphi B-H	FEBSBH05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella paratyphi B-O	FEBSCO05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella paratyphi C-H	FEBSCH05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella paratyphi C-O	FEBSCO05	1 x 5ml	5ml Stained Antigen Suspension

WEIL-FELIX

The Fortress Diagnostics Febrile Antigens Weli-Felix test provides rapid detection and semi-quantitation of serum antibodies developed during the acute stage of disease caused by rickettsial infection. The antigens agglutinate in the presence of the homologous antibodies in the sample tested.

Weil-Felix	FEWF0025	5 x 5ml	"5ml vials of Brucella Abortus, Brucella Melitensis, Proteus OX19, Proteus OX2 and Proteus OXK"
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SALMONELLA

The Fortress Diagnostics Salmonella Typhi Widal test is a common agglutination test employed in the serological diagnosis of enteric fever and helps to detect the presence of Salmonella antigens in a patient's serum. Patients infected with Salmonella produce antibodies against the antigens of the organism. Antibodies in serum produced in response to exposure to Salmonella organisms will agglutinate bacterial suspension which carries homologous antigens.

Salmonella typhi H	FEBSTH05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella typhi H	FEBSTH1L	1000 ml	1000ml Stained Antigen Suspension
Salmonella typhi O	FEBSTO05	1 x 5ml	5ml Stained Antigen Suspension
Salmonella typhi O	FEBSTO1L	1000 ml	1000ml Stained Antigen Suspension
Salmonella typhi O , H +Positive Control	FEBSOH05	2 x 1 x 5ml	Stained Antigen Suspension plus 1x0.5ml Positive control"



FORTRESS® DIAGNOSTICS

IMMUNOASSAYS (ELISA)

Fortress Diagnostics have developed a range of Enzyme Linked Immunoabsorbent Assays (ELISA) and Chemiluminescence immunoassays (CLIA) tests as an effective and highly sensitive range of diagnostics solutions.



RELIABLE

A trusted provider in the UK



RESULTS

Highly accurate



COMPREHENSIVE

Wide range of tests available



GOLD STANDARD

Gold standard assay technique

HORMONE LEVEL TESTING

17α-OH Progesterone	BXE0996A	96T	Human Serum or Plasma
Anti-Mullerian Hormone (AMH)	BXE0999A	96T	Human Serum
DHEA-S	BXE0998A	96T	Human Serum
Estradiol	BXE0860A	96T	Human Serum or plasma
Estradiol	BXE0860B	192T	Human Serum or plasma
Free Testosterone	BXE0997A	96T	Human Serum or plasma
Testosterone	BXE0862A	96T	Human Serum or plasma
Testosterone	BXE0862B	192T	Human Serum or plasma
Inhibin-A	BXE0615A	96T	Human Serum or plasma
Inhibin-B	BXE0618A	96T	Human Serum or plasma

Luteinizing hormone (LH)	BXE0651A	96T	Human Serum
Luteinizing hormone (LH)	BXE0651B	192T	Human Serum
Luteinizing hormone (LH)	BXE0651C	480T	Human Serum
Human Growth Hormone (HGH)	BXE0991A	96T	Human Serum
Prolactin	BXE0671A	96T	Human Serum
Prolactin	BXE0671B	192T	Human Serum
Prolactin	BXE0671C	480T	Human Serum
FSH	BXE0631A	96T	Human Serum
FSH	BXE0631B	192T	Human Serum
FSH	BXE0631C	480T	Human Serum
TSH	BXE0681A	96T	Human Serum
TSH	BXE0681B	192T	Human Serum
TSH	BXE0681C	480T	Human Serum
FT3	BXE0731A	96T	Human Serum
FT4	BXE0721A	96T	Human Serum
T3	BXE0701A	96T	Human Serum or plasma
T3	BXE0701B	192T	Human Serum or plasma
T3	BXE0701C	480T	Human Serum or plasma
T3 STREPTAVIDIN	BXE0703A	96T	Human Serum or plasma
T3 STREPTAVIDIN	BXE0703B	192T	Human Serum or plasma
T3-Uptake	BXE0702A	96T	Human Serum or plasma
T3-Uptake	BXE0702B	192T	Human Serum or plasma
T4	BXE0711A	96T	Human Serum or plasma
T4	BXE0711B	192T	Human Serum or plasma
T4	BXE0711C	480T	Human Serum or plasma
T4 STREPTAVIDIN	BXE0712A	96T	Human Serum or plasma
T4 STREPTAVIDIN	BXE0712B	192T	Human Serum or plasma

TUMOR MARKERS

Alpha-Fetoprotein (AFP)	BXE0801A	96T	Human serum
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Alpha-Fetoprotein (AFP)	BXE0801B	96T	Human serum
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Alpha-Fetoprotein (AFP) is a glycoprotein with a molecular weight of 70 kDa. AFP is normally produced during fetal development by the hepatocytes, yolk sac and to a lesser extent by the gastrointestinal tract. Serum concentrations reach a peak level of up to 10 mg/ml at twelve weeks of gestation. This peak level gradually decreases to less than 25 ng/ml after one year of postpartum. Thereafter, the levels reduce further to less than 10 ng/ml. Elevated levels of AFP are found in patients with primary hepatoma and yolk sac-derived germ tumors. AFP is the most useful marker for the diagnosis and management of hepatocellular carcinoma.

CEA	BXE0811A	96T	Human serum
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CEA is the first of the so-called carcinoembryonic proteins and is the most widely used marker for gastrointestinal cancer. Although CEA is primarily associated with colorectal cancers, other malignancies that can cause elevated levels of CEA include breast, lung, stomach, pancreas, ovary and other organs. The Fortress Diagnostics CEA can be used for quantitative determination of carcinoembryonic antigen (CEA) concentration in human serum by a microplate immunoenzymometric assay.

MCM5	BXE0950A	96T	Human urine
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Mcm5 is a novel marker for early and sensitive determination of Prostate Carcinoma. Conventional testing for Mcm5 is accomplished by the determination of PSA and fPSA which are marked by low sensitivity and specificity. The Fortress Diagnostics Mcm5 Elisa is intended for quantitative determination of Mcm5 in human urine.

CA125	BXE0821A	96T	high concentrations are linked to ovarian cancer and a range of benign and malignant diseases
CA15-3	BXE0881A	96T	considered one of the first prognostic factors for breast cancer
CA19.9	BXE0841A	96T	linked to gastrointestinal cancer

The Fortress Diagnostics Cancer Antigen kits are intended for use in the quantitative determination of a range of cancer antigens.

ENDOCRINES

Anti-Tg	BXE0895A	96T	Serum or plasma
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Anti-TPO	BXE0896A	96T	Serum or plasma
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Cortisol	BXE0831A	96T	Serum or plasma
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Ferritin	BXE0891A	96T	Serum
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DIABETES RELATED

C-Peptide	BXE0616A	96T	Serum
Insulin	BXE0616A	96T	Serum

INFLAMMATION

CRP (High Sensitivity, quantitative)	BXE0617A	96T	Serum or plasma
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COVID-19

Covid-19 Total Antibody	BXE0882A	96T	ELISA
Covid-19 Igm Antibody	BXE0883A	96T	ELISA

CARDIAC RELATED

Digoxin	BXE0992A	96T	Human serum or plasma
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Digoxin - The clinical usefulness of the measurement of serum digoxin (DIG) is due to its low therapeutic ratio - a very small difference exists between therapeutic and toxic tissue levels. In addition, individuals may vary in their response to digoxin with an apparent increase in susceptibility to toxicity with age. The action of digoxin is to increase the force and velocity of the myocardial contraction. This is necessary in the treatment of congestive heart failure and arrhythmias such as atrial fibrillation and atrial flutter.

The Fortress Diagnostics Digoxin kit is intended for the quantitative determination of digoxin concentration in human serum or plasma by a microplate enzyme immunoassay.

TROPONIN-I	BXE0864A	96T	Human serum
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The cardiac-specific isoform of Troponin-I (cTnI) has been known as a marker of heart damage and myocardial cell death, due to myocardial infarction, for just over 10 years. Troponin-I (cTnI, 24kDA) is the inhibitory subunit of the Troponin complex of striated muscle. Most of the Troponin (I & T) proteins are located within the contractile apparatus of the striated muscle.

The Fortress Diagnostics Troponin-I kit is a colorimetric enzyme-linked immunosorbent assay (ELISA) for the quantitative determination of circulating Troponin-I concentrations in human serum.

GASTROINTESTINAL

Helicobacter Pylori, IgA	BXE0672A	96T	Human serum or plasma
Helicobacter Pylori, IgG	BXE0673A	96T	Human serum or plasma
Helicobacter Pylori, IgM	BXE0674A	96T	Human serum or plasma

ALLERGENS

IgE	BXE0641A	96T	Human serum
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MALARIA

Malaria (Antigen)	BXE0691A	96T	Whole blood
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VITAMIN D

Vitamin D	BXE0111A	96T	Human Serum or plasma
Vitamin D (Fast - 90 Mins)	BXE0112A	96T	Human Serum or plasma

FERTILITY AND NEONATAL

Free β hCG	BXE0872A	96T	Human Serum
hCG	BXE0871A	96T	Human serum
hCG	BXE0871B	192T	Human serum
hCG Rapid	BXE0873A	96T	Human serum
hCG Rapid	BXE0873B	192T	Human serum
hCG Rapid Extended Range	BXE0874A	96T	Human serum
hCG Rapid Extended Range	BXE0874C	480T	Human serum
Neonatal T4	BXE0863A	96T	Human whole blood
Neonatal TSH	BXE0682A	96T	Human whole blood
PAPP-A (Pregnancy associated plasma protein A)	BXE0888A	96T	Human serum or plasma
Progesterone	BXE0661A	96T	Human serum or plasma
Progesterone	BXE0661B	192T	Human serum or plasma
Unconjugated Estriol (UE3)	BXE0865A	96T	Human serum or plasma
Patient Cards for NTSH & NT4	BXEPC100	100	6 specimen per paper card



FORTRESS® DIAGNOSTICS

CLINICAL CHEMISTRY REAGENTS

Fortress Diagnostics manufactures, develops and supplies an extensive range of high-quality clinical chemistry reagents. The reagents are manufactured to the highest quality standards in an ISO 13485:2016-certified laboratory, ensuring excellent stability, accuracy and results you can trust.

WHY CHOOSE FORTRESS DIAGNOSTICS CLINICAL CHEMISTRY REAGENTS?

- Compatible with market leading analysers
- Controls available (see our QC and calibrators section)
- Flexibility: Lyophilised and liquid formats available, as well as a variety of kit sizes, to suit different laboratory sizes and needs
- Excellent stability, to ensure cost-effectiveness

Acid Phosphatase (Lyo.)	BXC0401A	R1: 1 x 65ml R2: 6 x 10ml R3: 3 x 10ml R4: 1 x 5ml
Acid Phosphatase (Lyo.)	BXC0401B	R1: 1 x 105ml R2: 10 x 10ml R3: 5 x 10ml R4: 1 x 5ml
Acid Phosphatase (Lyo.)	BXC0401C	R1: 1 x 35ml R2: 10 x 3ml R3: 5 x 3ml R4: 1 x 5ml

Methodology: Colorimetric- Fast Red
Form: Lyo
Sample type: Serum
Sample Volume: 200 µl
Measuring Range: 1-90 U/L
Storage: 2°C - 8°C
Working Stability: 5 days at 2-8°C protected from light

ADA (L.S)	BXC0208A	R1: 1 x 50ml R2: 1 x 5ml R3: 1 x 10ml R4: 1 x 5ml
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Methodology: Enzymatic UV
Form: LS
Sample type: Serum, Heparinized Plasma
Sample Volume: 25 µl
Measuring Range: 0 U/l - 210 U/l
Storage: 2°C - 8°C
Working Stability: 30 days at 2-8°C

Aldolase (Lyo.)

BXC0391A

R1: 5 x 20ml R2: 2 x 1ml R3: 1 x 0.5ml

Methodology: GDH/TIM/LDH (UV)

Form: Lyo.

Sample type: Serum, Heparinized Plasma, EDTA Plasma

Sample Volume: 200µl

Measuring Range: 1.5 – 100 U/l

Storage: 2°C - 8°C

Working Stability: 2 weeks at 2-8°C

Albumin

BXC0221A

R1: 6 x 60ml R4: 1 x 5ml

Albumin (Monoliquid)

BXC0222A

R1: 2 x 60ml R4: 1 x 5ml

Albumin (Monoliquid)

BXC0222B

R1: 6 x 60ml R4: 1 x 5ml

Methodology: Bromocresol Green (BCG) Concentrate

Form: Concentrate

Sample type: Serum, plasma

Sample Volume: Macro: 10µl Micro: 5µl

Measuring Range: 1.76 g/dl - 60 g/l

Storage: 15°C - 25°C

Working Stability: 12 weeks at 15°C - 25°C

Alkaline Phosphatase (L.S) Colorimetric

BXC0183A

R1: 1 x 50ml R2: 2 x 125ml R4: 1 x 5ml

Alkaline Phosphatase (L.S) AMP (IFCC)

BXC0184A

R1: 5 x 25ml R2: 1 x 25ml

Alkaline Phosphatase (L.S) AMP (IFCC)

BXC0184B

R1: 5 x 50ml R2: 1 x 50ml

Alkaline Phosphatase (L.S) AMP (IFCC)

BXC0184C

R1: 5 x 100ml R2: 1 x 100ml

Alkaline Phosphatase (L.S) DEA (DGKC)

BXC0185A

R1: 5 x 25ml R2: 1 x 25ml

Alkaline Phosphatase (L.S) DEA (DGKC)

BXC0185B

R1: 5 x 50ml R2: 1 x 50ml

Alkaline Phosphatase (L.S) DEA (DGKC)

BXC0185C

R1: 5 x 100ml R2: 1 x 100ml

Alkaline Phosphatase (L.S) DEA (DGKC) 4+1

BXC0187A

R1: 1 x 100ml R2: 1 x 25ml

Alkaline Phosphatase (Lyo.) DEA (DGKC)

BXC0181A

R1: 5 x 20ml R2: 5 x 20ml

Alkaline Phosphatase (Lyo.) DEA (DGKC)

BXC0181C

R1: 1 x 105ml R2: 10 x 10ml

Alkaline Phosphatase (Lyo.) DEA (DGKC)

BXC0181D

R1: 1 x 65ml R2: 20 x 3ml

Alkaline Phosphatase (Lyo.) AMP (IFCC)

BXC0182A

R1: 1 x 105ml R2: 5 x 20ml

Alkaline Phosphatase (Lyo.) AMP (IFCC)

BXC0182D

R1: 1 x 105ml R2: 10 x 10ml

Methodology: Colorimetric, AMP (IFCC) or DEA (DGKC)

Sample type: Human serum or plasma

Storage: 2°C - 8°C

Alpha-1-Acid Glycoprotein (L.S)	BXC0890A	R1: 1 x 20ml R2: 1 x 4ml
Alpha-1-Antitrypsin (L.S)	BXC0190A	R1: 1 x 20ml R2: 1 x 4ml
Alpha-1-Antitrypsin (L.S)	BXC0190B	R1: 2 x 20ml R2: 1 x 8ml
Alpha-1-Microglobulin (L.S)	BXC0888A	R1: 1 x 20ml R2: 1 x 4ml
Alpha-1-Microglobulin (L.S)	BXC0888B	R1: 2 x 20ml R2: 1 x 8ml
Alpha-2-Macroglobulin (L.S)	BXC0889A	R1: 1 x 20ml R2: 1 x 4ml

Methodology: Immunoturbidimetric (Without Sample Pre-dilution)

ALT (GPT) (L.S) Colorimetric (Standard)	BXC0212A	R1: 1 x 125ml R2: 1 x 125ml R4: 1 x 5ml R6: 1 x 100ml
ALT (GPT) (L.S) IFCC	BXC0213A	R1: 1 x 100ml R2: 1 x 20ml
ALT (GPT) (L.S) IFCC	BXC0213D	R1: 2 x 100ml R2: 2 x 20ml
ALT (GPT) (L.S) IFCC	BXC0213F	R1: 5 x 100ml R2: 1 x 100ml
ALT (GPT) (L.S) Colorimetric (Table)	BXC0214A	R1: 1 x 100ml R2: 1 x 100ml
ALT (GPT) (L.S) DGKC 4+1	BXC0215A	R1: 1 x 100ml R2: 1 x 25ml
ALT (GPT) (L.S) DGKC 4+1	BXC0215B	R1: 2 x 100ml R2: 2 x 25ml
ALT (GPT) (Lyo.) IFCC	BXC0211B	R1: 2 x 105ml R2: 10 x 20ml
ALT (GPT) (Lyo.) IFCC	BXC0211C	R1: 1 x 105ml R2: 10 x 10ml
ALT (GPT) (Lyo.) IFCC	BXC0211D	R1: 5 x 50ml R2: 5 x 50ml
ALT (GPT) (Lyo.) IFCC	BXC0211E	R1: 1 x 45ml R2: 20 x 2ml
ALT (GPT) (Monoliquid) UV	BXC0127A	R1: 1 x 100ml

Methodology: Kinetic UV

Sample Type: Human serum, plasma

Storage: 2°C - 8°C

Ammonia (Monoliquid) GLDH	BXC0376A	R1: 10 x 10ml R4: 1 x 1ml Controls: 2 x 1ml
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Methodology: Enzymatic UV - Dea (DGKC) - Lyophilised

Form: Mono

Sample Type: Plasma

Sample Volume: 1000 ul

Measuring Range: 9 - 1700 ug/dl

Storage: 2°C - 8°C

Working Stability: 5 days after opening at +2-8°C

Amylase (L.S) EPS-pNPG7 IFCC	BXC0563A	R1: 1 x 25ml R2: 1 x 5ml
Amylase (L.S) EPS-pNPG7 IFCC	BXC0563B	R1: 2 x 25ml R2: 2 x 5ml
Amylase (L.S) EPS-pNPG7 IFCC	BXC0563D	R1: 1 x 100ml R2: 1 x 20ml
Amylase (Lyo.) Benzylidene Blocked-pNPG7	BXC0561A	R1: 1 x 30ml R2: 5 x 5ml
Amylase (Monoliquid) CNPG3	BXC0562A	R1: 6 x 10ml
Amylase (Monoliquid) CNPG3	BXC0562B	R1: 10 x 10ml
Amylase, Pancreatic (L.S) EPS-pNPG7 IFCC	BXC0564A	R1: 5 x 10ml R2: 5 x 2ml

Sample Type: Human serum, plasma & urine

Storage: 2°C - 8°C

Angiotensin Converting Enzyme (ACE) (Monoliquid)	BXC0176A	R1: 5 x 10ml R4: 1 x 1ml
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Methodology: Enzymatic UV (Liquid Stable)

Form: Mono

Sample Type: Serum/Heparinised Plasma

Sample Volume: 100 µl

Measuring Range: 5-170 U/l

Storage: 2°C - 8°C

Working Stability: 30 days @ +2-8°C

AST (GOT) (L.S) Colorimetric (Standard)	BXC0202A	R1: 1 x 125ml R2: 1 x 125ml R4: 1 x 5ml R6: 1 x 100ml
AST (GOT) (L.S) IFCC	BXC0203A	R1: 1 x 100ml R2: 1 x 20ml
AST (GOT) (L.S) IFCC	BXC0203D	R1: 2 x 100ml R2: 2 x 20ml
AST (GOT) (L.S) IFCC	BXC0203F	R1: 5 x 100ml R2: 1 x 100ml
AST (GOT) (L.S) Colorimetric (Table)	BXC0204A	R1: 1 x 100ml R2: 1 x 100ml
AST (GOT) (L.S)	BXC0205A	R1: 1 x 100ml R2: 1 x 25ml
AST (GOT) (Lyo.) IFCC	BXC0201B	R1: 2 x 105ml R2: 10 x 20ml
AST (GOT) (Lyo.) IFCC	BXC0201C	R1: 1 x 105ml R2: 10 x 10ml
AST (GOT) (Lyo.) IFCC	BXC0201D	R1: 5 x 50ml R2: 5 x 50ml
AST (GOT) (Lyo.) IFCC	BXC0201E	R1: 1 x 45ml R2: 20 x 2ml
AST (GOT) (Monoliquid) UV	BXC0128A	R1: 1 x 100ml

Sample Type: Human serum, plasma

Sample Volume: 100 µl

Storage: 2°C - 8°C

Anti Thrombin-III (L.S) **BXC0429A** **R1: 2 x 20ml R2: 1 x 8ml**

Methodology: Immunoturbidimetric (Without Sample Pre-dilution)

Form: LS

Sample Type: Serum or plasma

Sample Volume: 2µl

Measuring Range: 3-80 mg/dl (0.03-0.80 g/l)

Storage: 2°C - 8°C

Working Stability: 4 weeks at 2-8°C

Apolipoprotein A1 (L.S) **BXC0411A** **R1: 2 x 20ml R2: 1 x 8ml**

Apolipoprotein B (L.S) **BXC0412A** **R1: 2 x 20ml R2: 1 x 8ml**

Methodology: Immunoturbidimetric (Without Sample Pre-dilution)

Form: LS

Sample Type: Serum or plasma

Sample Volume: 2µl

Storage: 2°C - 8°C

Working Stability: 30 days at + 2-8°C

ASO (L.S) **BXC0501A** **R1: 1 x 40ml R2: 1 x 10ml R4: 1 x 1ml**

ASO (L.S) **BXC0501B** **R1: 2 x 40ml R2: 2 x 10ml R4: 1 x 1ml**

ASO (L.S) **BXC0501C** **R1: 4 x 40ml R2: 4 x 10ml R4: 1 x 1ml**

Methodology: Turbidimetric Single Point

Form: LS

Sample Type: Serum

Sample Volume: 10 µl

Measuring Range: 12.5-800 U/l

Storage: 2°C - 8°C

Working Stability: 28 Days at +2-8°C

Access Fluid (Monoliquid) **BXC0161B** **1 x 1000ml**

Methodology: For use as a wetting agent for Technicon RA series of Analysers.

Form: Mono

Storage: At room temperature up to 25°C

Bile Acids (Lyo.) **BXC0581A** **R1: 1 x 65ml R2: 1 x 20ml R3: 6 x 10ml**

Methodology: Enzymatic Colorimetric

Sample Type: Serum or plasma

Sample Volume: 4 µl

Storage: 2°C - 8°C

Bilirubin (L.S) DIRECT	BXC0191A	R1: 2 x 50ml R2: 1 x 20ml
Bilirubin (L.S) DIRECT	BXC0191B	R1: 2 x 100ml R2: 1 x 40ml
Bilirubin (L.S) TOTAL	BXC0192A	R1: 2 x 50ml R2: 1 x 25ml
Bilirubin (L.S) TOTAL	BXC0192B	R1: 2 x 100ml R2: 1 x 50ml
Bilirubin (L.S) TOTAL & DIRECT MANUAL	BXC0193A	R1: 1 x 100ml R2: 1 x 100ml R3: 1 x 50ml / 1 x 10ml
Bilirubin (L.S) TOTAL & DIRECT MANUAL	BXC0193B	R1: 2 x 100ml R2: 1 x 100ml R3: 1 x 100ml / 1 x 20ml
Bilirubin Direct (L.S) Vanadate Oxidation	BXC0104A	R1: 2 x 70ml R2: 2 x 18ml
Bilirubin Total (L.S) Vanadate Oxidation	BXC0105A	R1: 2 x 70ml R2: 2 x 18ml

Sample Type: Vanadate Oxidation and Colorimetric Methods available

Form: LS

Sample Type: Serum or plasma

Storage: 15-25°C

C1 Esterase Inhibitor (L.S)	BXC0170A	R1: 1 x 20ml R2: 1 x 4ml
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Methodology: Immunoturbidimetric (Without Sample Pre-dilution)

Form: LS

Sample Type: Human serum or plasma

Sample Volume: 2 µl

Measuring Range: 0.10-106 mg/dl

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

Calcium (L.S) AMP/CPC	BXC0291A	R1: 1 x 125ml R2: 1 x 125ml R3: 1 x 12.5ml R4: 1 x 5ml
Calcium (Monoliquid) Arsenazo	BXC0292A	R1: 2 x 125ml R4: 1 x 5ml

Methodology: Colorimetric

Form: LS

Sample Type: Serum, Plasma, Urine

Sample Volume: 25 µl

Measuring Range: AMP/CPC 0.12-5.50mmol/l (0.5-22mg/dl); Arsenazo 0.05-4.8mmol/l (0.2-19 mg/dl)

Storage: +2-8°C

Working Stability: 28 Days @ +2-8°C

Ceruloplasmin (L.S)	BXC0765A	R1: 1 x 20ml R2: 1 x 4ml
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Methodology: Immunoturbidimetric (Without Sample Pre-dilution)

Form: LS

Sample Type: Human serum or plasma

Sample Volume: 2 µl

Measuring Range: 10-280 mg/dl (0.10-2.80g/l)

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

Chloride (Monoliquid)	BXC0281A	R1: 2 x 60ml STD L:1 x 5ml STD H:1 x 5ml
<p>Methodology: Colorimetric (Thiocyanate) Form: Mono Sample Type: Serum/Plasma Sample Volume: 10 µl Measuring Range: Varies with each assay Storage: 15- 25°C Working Stability: Up to expiry when stored at +15°C- 25°C</p>		

Cholesterol (Lyo.)	BXC0262A	R1: 2 x 105ml R2: 10 x 20ml R4: 1 x 5ml
Cholesterol (Lyo.)	BXC0262B	R1: 10 x 50ml R2: 10 x 50ml R4: 1 x 5ml
Cholesterol (Monoliquid)	BXC0261A	R1: 2 x 60ml R4: 1 x 5ml
Cholesterol (Monoliquid)	BXC0261B	R1: 6 x 60ml R4: 1 x 5ml
Cholesterol (Monoliquid)	BXC0261C	R1: 8 x 250ml R4: 1 x 5ml
Cholesterol (Monoliquid)	BXC0261D	R1: 4 x 250ml R4: 1 x 5ml
<p>Form: Mono/Lyo Sample Type: Serum/Heparinised Plasma Sample Volume: 10 µl Storage: +2-8°C</p>		

Cholinesterase (L.S)	BXC0801A	R1: 5 x 20 R2: 2 x 10ml
<p>Methodology: DGKC Form: LS Sample Type: Serum/Plasma Sample Volume: 20 µl Measuring Range: 35-25,000 U/l Storage: +2-8°C Working Stability: 24 Hrs @ 15-25°C, 5 days @ +2-8°C</p>		

Citrate (Urinary) (L.S)	BXC0120A	R1: 2x10ml; R2: 1x2ml; R3: 2x10ml; 1x5ml, R4: 2x3ml
<p>Methodology: Enzymatic Colorimetric Form: LS Sample Type: Human urine Sample Volume: 20 µl Measuring Range: 0-1600 mg/l (0.03-8.34mmol/l) Storage: +2-8°C Working Stability: 3 days @ +2-8°C</p>		

CK-MB (L.S)	BXC0452A	R1: 1 x 25ml R2: 1 x 5ml
CK-MB (L.S)	BXC0452B	R1: 2 x 25ml R2: 2 x 5ml
CK-MB (L.S)	BXC0452C	R1: 10 x 3ml R2: 1 x 6ml
CK-MB (Lyo.)	BXC0451A	R1: 1 x 35ml R2: 10 x 2.5ml R3:1 x 2ml
CK-MB (Lyo.)	BXC0451B	R1: 1 x 105ml R2: 10 x 10ml R3:1 x 2ml
CK-MB (Lyo.)	BXC0451C	R1: 1 x 15ml R2: 5 x 2.5ml R3:1 x 2ml
CK-MB 4+1 (L.S)	BXC0458A	R1: 1 x 20ml R2: 1 x 4ml
CK-MB 4+1 (L.S)	BXC0458A	R1: 2 x 20ml R2: 1 x 8ml

Methodology: Enzymatic UV

Form: LS

Sample Type: Serum or plasma

Measuring Range: 5-1000 U/l

Storage: +2-8°C

CK-nac (L.S)	BXC0252A	R1: 2 x 25ml R2: 2 x 5ml
CK-nac (L.S)	BXC0252B	R1: 2 x 50ml R2: 2 x 10ml
CK-nac (L.S)	BXC0252C	R1: 6 x 10ml R2: 1 x 12ml
CK-nac (Lyo.)	BXC0251B	R1: 1 x 105ml R2: 10 x 10ml
CK-nac (Lyo.)	BXC0251D	R1: 1 x 35ml R2: 10 x 2.5ml
CK-nac 4+1 (L.S)	BXC0253A	R1: 2 x 20ml R2: 1 x 10ml
CK-nac 4+1 (L.S)	BXC0253B	R1: 2 x 40ml R2: 1 x 20ml

Methodology: Enzymatic UV

Form: LS/Lyo

Sample Type: Serum or plasma

Measuring Range: 2-2300 U/l

Storage: +2-8°C

CO2 (Bicarbonate) (Monoliquid)	BXC0152A	R1: 10 x 5ml R4: 1 x 1ml
CO2 (Bicarbonate) (Monoliquid)	BXC0152B	R1: 10 x 10ml R4: 1 x 1ml

Methodology: Enzymatic

Form: LS

Sample Type: Serum

Sample Volume: 10 µl

Measuring Range: 1.12-50 mmol/l

Storage: +2-8°C

Working Stability: 7 days @ +2-8°C after opening

Complement C3 (L.S)	BXC0851A	R1: 2 x 20ml R2: 1 x 8ml
Complement C4 (L.S)	BXC0861A	R1: 2 x 20ml R2: 1 x 8ml
Methodology: Immunoturbidmetric Form: LS Sample Type: Human serum or plasma Sample Volume: 2 µl Measuring Range: Varies according to assay Storage: +2-8°C Working Stability: 4 weeks @ +2-8°C		

Copper (Monoliquid)	BXC0341A	R1: 2 x 25ml R4: 1 x 5ml
Copper (Urinary) (L.S)	BXC0342A	R1: 2 x 25ml R2: 1 x 10ml R4: 1 x 5ml
Copper Urinary (Monoliquid)	BXC0343A	R1: 2 x 25ml R4: 1 x 5ml
Methodology: Colorimetric Sample Type: Serum or plasma Measuring Range: Varies according to assay Storage: +2-8°C Working Stability: 28 days @ +2-8°C		

Creatinine (L.S)	BXC0111A	R1: 1 x 120ml R2: 1 x 120ml R4: 1 x 5ml
Creatinine (L.S)	BXC0111B	R1 3 x 60ml R2 3 x 60ml R4 1 x 5ml
Creatinine (L.S)	BXC0112A	R1: 1 x 120ml R2: 1 x 120ml R4: 1 x 35ml
Creatinine (L.S)	BXC0117A	R1: 1 x 120ml R2: 1 x 120ml R4: 1 x 5ml
Creatinine (Lyo.)	BXC0113A	R1: 4 x 25ml R2: 4 x 25ml R3: 2 x 10ml R4: 1 x 2ml
Creatinine (Lyo.)	BXC0113B	R1: 8 x 25ml R2: 8 x 25ml R3: 4 x 10ml R4: 1 x 2ml
Creatinine TCA Concentrate	BXC0114A	R1: 1 x 120ml
Methodology: Jaffe without Deproteinization/ Enzymatic, UV Sample Type: Serum/plasma and urine		

CRP (L.S)	BXC0382A	R1: 1 x 40ml R2: 1 x 10ml R4: 1 x 1ml
CRP (L.S)	BXC0382B	R1: 2 x 40ml R2: 2 x 10ml R4: 1 x 1ml
CRP (L.S)	BXC0382C	R1: 4 x 40ml R2: 4 x 10ml R4: 1 x 1ml
CRP (L.S)	BXC0384A	R1: 2 x 20ml R2: 1 x 8ml R4: 1 x 1ml
CRP (ULTRA SENSITIVE) (L.S)	BXC0383A	R1: 1 x 16ml R2: 1 x 4ml R4: 1 x 200ul
Methodology: Immunoturbidmetric/ Turbidmetric Form: LS Sample Type: Human serum or plasma Storage: +2-8°C		

Cystatin-C (L.S)

BXC0777A

R1: 1 x 20ml R2: 1 x 5ml

Methodology: Latex Enhanced Turbidmetric

Form: LS

Sample Type: Serum

Sample Volume: 3 µl

Measuring Range: 1-10 mg/ml

Storage: +2-8°C

Working Stability: 30 days @ +2-8°C

D3-Hydroxybutyrate (L.S)

BXC0542A

R1: 1 x 50ml R2: 1 x 8ml R4: 1 x 5ml

D3-Hydroxybutyrate (Lyo.)

BXC0541A

R1: 1 x 50ml R2: 5 x 10ml R4: 1 x 5ml

Methodology: Enzymatic

Sample Type: Serum or plasma

Sample Volume: 25 µl

Measuring Range: 1.12-50 mmol/l

Storage: +2-8°C

D-Dimer (L.S)

BXC0787A

R1: 1 x 20ml R2: 1 x 4ml

D-Dimer (L.S)

BXC0787B

R1: 1 x 20ml R2: 1 x 8ml

Methodology: Turbilatex

Form: LS

Sample Type: Serum

Sample Volume: 18 µl

Measuring Range: 0.2-10 ug/ml

Storage: +2-8°C

Working Stability: 4 weeks at 2-8°C

Drabkins Solution (For use with BXC0551A) (L.S)

BXC0552A

R1: 4 x 200ml

Form: LS

Sample Type: Whole blood

Sample Volume: 50 µl

Storage: Room temperature up to 25°C

Working Stability: 6 months @ Room Temperature up to 25°C

Ethanol (L.S)

BXC0491A

R1: 3 x 20ml R2: 2 x 5ml R4: 1 x 2ml

Methodology: Enzymatic UV

Form: LS

Sample Type: Serum/Plasma/Urine

Sample Volume: 8 µl

Measuring Range: 0 - 300 mg/dl

Storage: +2-8°C

Working Stability: 28 days at 2°C - 8°C

Ferritin (L.S)

BXC0441A

R1: 1 x 20ml R2: 1 x 5ml R4: 1 x 1ml

Methodology: Turbidimetric

Form: LS

Sample Type: Serum or plasma

Sample Volume: 90 µl

Measuring Range: 5.0-600 ug/l

Storage: 2-8°C / DO NOT FREEZE

Working Stability: Stable up to expiry date

Fibrinogen (L.S)

BXC0442A

R1: 2 x 20ml R2: 1 x 8ml

Methodology: Immunoturbidimetry

Form: LS

Sample Type: Plasma

Sample Volume: 50 ul

Measuring Range: 100-900 mg/dl

Storage: 2°C - 8°C

Working Stability: 28 days at 2-8°C

Fructosamine (L.S)

BXC0591A

R1: 3 x 6ml R2: 3 x 14ml

Methodology: Colorimetric-Enzymatic

Form: LS

Sample Type: Human serum or plasma

Sample Volume: 50 µl

Measuring Range: 10-1000 umol/l

Storage: 2°C - 8°C

Working Stability: 28 days @ +2-8°C

GGT (L.S)

BXC0362A

R1: 1 x 60ml R2: 1 x 12ml

GGT (L.S)

BXC0362D

R1: 2 x 60ml R2: 2 x 12ml

GGT (Lyo.)

BXC0361A

R1: 1 x 60ml R2: 5 x 10ml

Methodology: Carboxy

Sample Type: Serum or plasma

Measuring Range: 1-1200 U/l

Storage: +2-8°C

Glucose (L.S)	BXC0103A	R1: 5 x 25ml R2: 1 x 25ml R4: 1 x 5ml
Glucose (Lyo.)	BXC0102A	R1: 4 x 250ml R2: 4 x 250ml R4: 1 x 5ml
Glucose (Lyo.)	BXC0102B	R1: 10 x 500ml R2: 10 x 500ml R4: 1 x 5ml
Glucose (Monoliquid)	BXC0101A	R1: 6 x 60ml R4: 1 x 5ml
Glucose (Monoliquid)	BXC0101B	R1: 6 x 125ml R4: 1 x 5ml
Glucose (Monoliquid)	BXC0101C	R1: 2 x 500ml R4: 1 x 5ml
Glucose (Monoliquid)	BXC0101D	R1: 4 x 250ml R4: 1 x 5ml
Glucose (Monoliquid)	BXC0101E	R1: 10 x 500ml R4: 1 x 5ml

Methodology: Enzymatic

Sample Type: Blood, CSF, Urine (BXC0103 only)/Human serum or plasma

Storage: +2-8°C

Working Stability: 28 days @ +2-8°C

Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0571A	R1: 2 x 50ml R2: 1 x 2ml R3:1 x 2ml R4: 1 x 20ml
Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0574A	25 TESTS
Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0574B	50 TESTS
Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0574C	100 TESTS
Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0574D	500 TESTS
Glucose-6-Phosphate Dehydrogenase (G6PDH) (Lyo.)	BXC0574E	800 TESTS

Methodology: UV

Form: Lyo

Sample Type: Whole blood

Sample Volume: 10ul

Measuring range: 150-1250 U/l (BXC0571A only)

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

Glutathione Peroxidase (Lyo.)	BXC0551A	R1: 1 x 70ml R2: 5 x 10ml R3:1 x 1ml R6:2 x 200ml
Glutathione Reductase (Lyo.)	BXC0853A	R1: 1 x 50ml R2: 5 x 5ml R3: 5 x 3ml
GLYCOHAEMOGLOBIN (L.S)	BXC0650A	R1: 1 x 60ml R2: 1 x 10ml R4: 1 x 1ml
GLYCOHAEMOGLOBIN (L.S)	BXC0650B	R1: 3 x 100ml R2: 1 x 50ml R4: 1 x 1ml

Methodology: Enzymatic UV

Sample Type: Serum, plasma or erythrocytes

Storage: +2-8°C

Haemoglobin (Monoliquid)	BXC0482A	R1: 1 x 1000ml
Haemoglobin (Monoliquid)	BXC0482B	R1: 3 x 1000ml
GLYCOHAEMOGLOBIN (L.S)	BXC0650A	R1: 1 x 60ml R2: 1 x 10ml R4: 1 x 1ml
GLYCOHAEMOGLOBIN (L.S)	BXC0650B	R1: 3 x 100ml R2: 1 x 50ml R4: 1 x 1ml

Methodology: Colorimetric Method

Form: Conc

Sample Type: Human blood

Sample Volume: 20 µl

Measuring Range: 0.03-20 g/dl

Storage: +2-8°C

Working Stability: Undiluted RGT stable up to exp date shown when stored @ + 2-8°C: diluted @ room temp

Haptoglobin (L.S)	BXC0496A	R1: 2 x 20ml R2: 1 x 8ml
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Methodology: Immuno-turbidimetric

Form: LS

Sample Type: Human serum or plasma

Measuring Range: 20-570 mg/dl (0.20-5.70 g/l)

Storage: 2°C - 8°C

Working Stability: 4 weeks @ +2-8°C

HbA1c	BXC0670A	R1: 1 x 30ml R2: 1 x 10ml R3: 2 x 120ml R4: 2 x 0.5ml
HbA1c	BXC0668A	2 x 1 x 0.5ml
HbA1c	BXC0669A	2 x 1 x 0.5ml
HbA1c	BXC0671A	25 Tests
HbA1c	BXC0671B	10 Tests
HbA1c	BXC0671C	100 Tests
HbA1c	BXC0672A	R1: 1 x 7ml R2: 1 x 3ml R3: 1 x 4.5ml R4: 1 x 15ml
HbA1c	BXC0672B	R1: 1 x 21ml R2: 1 x 9ml R3: 1 x 12.5ml R4: 1 x 45ml

Methodology: Modified Enzymatic

Form: Conc

Sample Type: Human blood

Sample Volume: 6 µl

Measuring Range: 3-15%

Storage: 2°C - 8°C

Working Stability: 4 weeks @ 2-8°C

HDL Cholesterol (L.S)	BXC0421A	R1: 3 x 10ml R2: 1 x 10ml
HDL Cholesterol (L.S)	BXC0421B	R1: 6 x 30ml R2: 3 x 20ml
HDL Cholesterol (L.S)	BXC0421C	R1: 4 x 70ml R2: 2 x 50ml
HDL Cholesterol (L.S)	BXC0421D	R1: 3 x 10ml R2: 1 x 10ml R4: 1 x 1ml
HDL Cholesterol (L.S)	BXC0421E	R1: 6 x 30ml R2: 3 x 20ml R4: 1 x 1ml
HDL Cholesterol (L.S)	BXC0421F	R1: 4 x 70ml R2: 2 x 50ml R4: 1 x 1ml
HDL Cholesterol (L.S)	BXC0421G	R1: 4 x 125ml R2: 2 x 85ml R4: 1 x 1ml
HDL Cholesterol (Monoliquid)	BXC0422A	R1: 2 x 60ml R4: 1 x 5ml

Methodology: Enzymatic

Sample Type: Serum or heparinised plasma

Homocysteine (L.S)	BXC0690A	R1: 1 x 16ml R2: 1 x 4ml
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Methodology: Enzymatic UV

Form: LS

Sample Type: Serum or plasma

Sample Volume: 52 µl

Measuring Range: 0.5-50 umol/l

Storage: +2-8°C

Working Stability: 5 days at 2-8°C

IgA (L.S)	BXC0701A	R1: 2 x 20ml R2: 1 x 8ml
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Methodology: Immunoturbidmetric

Form: LS

Sample Type: Human serum or plasma

Sample Volume: 2 µl

Measuring Range: 50-800 mg/dl (0.5-8.00 g/l)

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

IgE (L.S)	BXC0751A	R1: 2 x 20ml R2: 1 x 8ml
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Methodology: Immunoturbidmetric

Form: LS

Sample Type: Human serum or plasma

Sample Volume: 35 µl

Measuring Range: 5.7-1500 IU/ml (16.4-2400 ng/ml)

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

IgG (L.S)	BXC0721A	R1: 2 x 20ml R2: 1 x 8ml
Methodology: Immunoturbidmetric Form: LS Sample Type: Human serum or plasma Sample Volume: 2 µl Measuring Range: 300-3100 mg/dl Storage: +2-8°C Working Stability: 4 weeks @ 2-8°C		
IgM (L.S)	BXC0731A	R1: 2 x 20ml R2: 1 x 8ml
Methodology: Immunoturbidmetric Form: LS Sample Type: Human serum or plasma Sample Volume: 2 µl Measuring Range: 25-650 mg/dl (0.25-6.5 g/l) Storage: +2-8°C Working Stability: 4 weeks @ 2-8°C		
Iron (L.S)	BXC0232A	R1 2 x 125ml R2 1 x 65ml
Iron (L.S)	BXC0235A	R1: 2 x 50ml R2: 2 x 10ml R4: 1 x 5ml
Iron (L.S)	BXC0235B	R1: 4 x 50ml R2: 2 x 20ml R4: 1 x 5ml
Iron (Monoliquid)	BXC0236A	R1: 2 x 50ml R4: 1 x 5ml
Iron/UIBC (L.S)	BXC0234A	A:1 x 100ml B:2 x 15ml C:1 x 10ml D:1 x 10ml E:1 x 10ml F:1 x 100ml G:1 x 5ml
Sample Type: Human serum or plasma Storage: +2-8°C Working Stability: 4 weeks @ 2-8°C		
Kappa Light Chains (L.S)	BXC0331A	R1: 2 x 20ml R2: 1 x 8ml
Methodology: Immunoturbidmetric Form: LS Sample Type: Serum or plasma Sample Volume: 2 µl Measuring Range: 25-900 mg/dl (0.25-9.00 g/l) Storage: +2-8°C Working Stability: Onboard stability 28 days at 2-8°C		

Lactate (Lyo.)	BXC0621A	R1: 1 x 105ml R2: 16 x 6ml R4: 1 x 5ml
Lactate (Monoliquid)	BXC0622A	R1: 2 x 50ml R4: 1 x 5ml
Lactate Dehydrogenase (LD) (L.S)	BXC0242A	R1: 5 x 20ml R2: 1 x 20ml
Lactate Dehydrogenase (LD) (L.S)	BXC0242B	R1: 10 x 10ml R2: 2 x 10ml
Lactate Dehydrogenase (LD) (L.S)	BXC0243A	R1: 5 x 20ml R2: 1 x 20ml
Lactate Dehydrogenase (LD) (L.S)	BXC0243B	R1: 10 x 10ml R2: 2 x 10ml

Methodology: Enzymatic Colorimetric

Sample Type: Plasma and CSF

Sample Volume: 10 µl

Storage: 2°C - 8°C

Lambda Light Chains (L.S)	BXC0381A	R1: 2 x 20ml R2: 1 x 8ml
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Methodology: Immunoturbidmetric

Form: LS

Sample Type: Serum or plasma

Sample Volume: 2 µl

Measuring Range: 30-500 mg/dl : 0.30-5.00 g/l

Storage: +2-8°C

Working Stability: 4 weeks @ 2-8°C

LDL Cholesterol (L.S)	BXC0431A	R1: 3 x 10ml R2: 1 x 10ml
LDL Cholesterol (L.S)	BXC0431B	R1: 6 x 30ml R2: 3 x 20ml
LDL Cholesterol (L.S)	BXC0431C	R1: 4 x 70ml R2: 2 x 50ml
LDL Cholesterol (L.S)	BXC0431D	R1: 3 x 10ml R2: 1 x 10ml R4: 1 x 1ml
LDL Cholesterol (L.S)	BXC0431E	R1: 6 x 30ml R2: 3 x 20ml R4: 1 x 1ml
LDL Cholesterol (L.S)	BXC0431F	R1: 4 x 70ml R2: 2 x 50ml R4: 1 x 1ml
LDL Cholesterol (Monoliquid)	BXC0432A	R1: 2 x 60ml R4: 1 x 5ml

Methodology: Enzymatic Colorimetric

Sample Type: Serum & plasma

Sample Volume: 5 µl

Storage: 2°C - 8°C

Lipase (L.S)

BXC0511A

R1: 2 x 10ml R2: 1 x 10ml R4: 1 x 1ml

Lipase (UV) (Lyo.)

BXC0512A

R1: 1 x 35ml R2: 10 x 2.5ml R4: 3 x 1ml

Methodology: Enzymatic Colorimetric

Form: LS

Sample Type: Serum or plasma

Sample Volume: 30 µl

Measuring Range: 2.43-548 U/l

Storage: +2-8°C

Working Stability: 31 days @ +2-8°C

Lithium (L.S)

BXC0125A

R1: 1 x 20ml R2: 1 x 10ml

Methodology: Enzymatic

Magnesium (L.S) Calmagite

BXC0351A

R1: 1 x 60ml R2: 1 x 60ml R4: 1 x 5ml

Magnesium (L.S) Calmagite

BXC0351B

R1: 2 x 20ml R2: 2 x 20ml R4: 1 x 5ml

Magnesium (Monoliquid) Xylidyl Blue

BXC0352A

R1: 2 x 60ml R4: 1 x 5ml

Magnesium (Monoliquid) Xylidyl Blue

BXC0352B

R1: 6 x 60ml R4: 1 x 5ml

Magnesium (Monoliquid) Arsenazo

BXC0353A

R1: 2 x 60ml R4: 1 x 5ml

Sample Type: Serum, plasma or urine

Storage: 15-25°C

Working Stability: 7 days @ +15-25°C (Magnesium calmagite); Up to expiry when stored @ +15-25°C (Magnesium Xylidyl Blue & Arsenazo)

Microalbumin (L.S)

BXC0471A

R1: 1 x 50ml R2: 1 x 10ml R4: 5 x 1ml(C1-C5)

Methodology: Immunoturbidmetric

Form: LS

Sample Type: Urine

Sample Volume: 60 µl

Measuring Range: 0.25-200 mg/l

Storage: +2-8°C

Working Stability: up to expiry when stored at +2-8°C

Oxalate (Urinary) (L.S)

BXC0129A

40 TESTS

Methodology: Immunoturbidmetric

Form: LS

Sample Type: Human urine

Sample Volume: 25 µl

Measuring Range: 0.003-2 mmol/l

Storage: +2-8°C

Working Stability: up to expiry when stored @+15-25°C

Phosphorus (Inorganic) (L.S)	BXC0301A	R1: 2 x 30ml R2: 2 x 70ml R4: 1 x 5ml
Phosphorus (Inorganic) (Monoliquid)	BXC0302A	R1: 5 x 20ml R4: 1 x 5ml
Methodology: Molybdate (UV) Sample Type: Serum or urine (BXC0302 only) Sample Volume: 10 µl Measuring Range: 0.25-6.5 mmol/l Storage: 15°C - 25°C		

Potassium (L.S)	BXC0132A	R1: 1 x 30ml R2: 1 x 30ml R3: 1 x 60ml R4: 1 x 5ml
Potassium (L.S)	BXC0135A	R1: 4 x 20ml R2: 2 x 10ml R4: 2 x 5ml
Potassium (Monoliquid)	BXC0138A	R1: 5 x 20ml R4: 1 x 5ml
Potassium (Monoliquid)	BXC0138B	R1: 10 x 20ml R4: 1 x 5ml
Sample Type: Serum Storage: 2-8°C		

Pre-Albumin (L.S)	BXC0449A	R1: 2 x 20ml R4: 1 x 8ml
Methodology: Immunoturbidimetric Form: LS Sample Type: Serum and plasma Sample Volume: 2 µl Measuring Range: 3-80 mg/dl (0.07-0.80 g/l) Storage: +2-8°C Working Stability: 4 weeks at 2-8°C		

RF (L.S)	BXC0611A	R1: 1 x 40ml R2: 1 x 10ml R4: 1 x 2ml
RF (L.S)	BXC0611B	R1: 2 x 40ml R2: 2 x 10ml R4: 1 x 2ml
RF (L.S)	BXC0611C	R1: 4 x 40ml R2: 4 x 10ml R4: 1 x 2ml
Methodology: Turbidimetric Form: LS Sample Type: Human serum or plasma Sample Volume: 7 µl Measuring Range: 6-160 IU/ml Storage: 2-8°C Working Stability: Stable up to expiry date when stored at +2-8°C		

Sodium (L.S)	BXC0142A	R1: 2 x 20ml R2: 2 x 10ml R4: 2 x 5ml
Sodium (Monoliquid)	BXC0146A	R1: 5 x 20ml R4: 2 x 5ml
Methodology: Enzymatic Sample Type: Human serum Storage: 2-8°C		

TIBC (L.S)	BXC0233A	R1: 1 x 100ml R2: 1 x 20g
TIBC (L.S)	BXC0237A	R1: 1 x 50ml R2: 1 x 15ml
<p>Methodology: Biuret (Colorimetric & ready to use) Form: LS Sample Type: Human serum or plasma Sample Volume: 20 µl (TIBC Direct)</p>		
Total Antioxidant Status (TAS) (L.S)	BXC0553A	R1: 1 x 50ml R2: 1 x 10ml STD1 1 x 1ml STD2 1 x 1ml
<p>Methodology: Enzymatic colorimetric Form: LS Sample Type: Serum and plasma Sample Volume: 50 µl Measuring Range: Linear up to 2.7 mmol/l Storage: +2-8°C Working Stability: Opened reagents - 28 days @ + 2-8°C</p>		
Total Lipid (Colorimetric) (L.S)	BXC0263A	R1: 2 x 100ml R4: 1 x 10ml
<p>Methodology: Colorimetric Form: LS Sample Type: Serum Sample Volume: 10 µl Measuring Range: 3-1300 mg/dl Storage: Room temperature up to 25°C Working Stability: up to expiry when stored at RT up to +25°C</p>		
Total Protein (L.S)	BXC0171A	R1: 4 x 125ml R2: 1 x 50ml R4: 1 x 5ml
Total Protein (Monoliquid)	BXC0173A	R1: 2 x 60ml R4: 1 x 5ml
Total Protein (Monoliquid)	BXC0173B	R1: 6 x 60ml R4: 1 x 5ml
<p>Methodology: Biuret (Colorimetric & ready to use) Sample Type: Human serum or plasma Sample Volume: 20 µl Measuring Range: 6-160 IU/ml Storage: Room temperature up to 25°C</p>		
Transferrin (L.S)	BXC0741A	R1: 2 x 20ml R2: 1 x 8ml
<p>Methodology: Immunoturbidimetric Form: LS Sample Type: Human serum or plasma Sample Volume: 2 µl Measuring Range: 15-850 mg/dl Storage: +2-8°C Working Stability: up to expiry when stored at +2-8°C</p>		

Triglycerides (Lyo.)	BXC0272A	R1: 1 x 105ml R2: 10 x 10ml R4: 1 x 5ml
Triglycerides (Lyo.)	BXC0272B	R1: 2 x 105ml R2: 10 x 20ml R4: 1 x 5ml
Triglycerides (Lyo.)	BXC0272C	R1: 4 x 50ml R2: 4 x 50ml R4: 1 x 5ml
Triglycerides (Monoliquid)	BXC0271A	R1: 2 x 60ml R4: 1 x 5ml
Triglycerides (Monoliquid)	BXC0271B	R1: 4 x 60ml R4: 1 x 5ml
Triglycerides (Monoliquid)	BXC0271C	R1: 5 x 20ml R4: 1 x 5ml
Triglycerides (Monoliquid)	BXC0271D	R1: 12 x 60ml R4: 1 x 5ml

Methodology: Enzymatic
Sample Type: Human serum or plasma
Sample Volume: 10 µl
Measuring Range: 3-1000 mg/dl (0.05-11.4 mmol/l)
Storage: 2°C - 8°C

Troponin-I (L.S)	BXC0469A	R1: 1 x 40ml R2: 1 x 10ml
Troponin-I (L.S)	BXC0469B	R1: 1 x 80ml R2: 1 x 20ml

Methodology: Latex Enhanced Turbidimetry
Form: LS
Sample Type: Serum or plasma
Sample Volume: 25 µl
Measuring Range: 0.28-10 ng/ml
Storage: +2-8°C
Working Stability: Onboard stability 28 days @ + 2-8°C

Urea (L.S)	BXC0122A	R1: 2 x 125ml R2: 1 x 50ml R3:2 x 6.5ml R4: 1 x 5ml
Urea (L.S)	BXC0122B	R1: 4 x 125ml R2: 2 x 50ml R3:4 x 6.5ml R4: 1 x 5ml
Urea (L.S)	BXC0123A	R1: 1 x 100ml R2: 1 x 20ml R4: 1 x 5ml
Urea (L.S)	BXC0123B	R1: 3 x 100ml R2: 1 x 60ml R4: 1 x 5ml
Urea (L.S)	BXC0123C	R1: 5 x 100ml R2: 1 x 100ml R4: 1 x 5ml
Urea (L.S)	BXC0124A	R1: 1 x 100ml R2: 1 x 25ml R4: 1 x 5ml
Urea (Lyo.)	BXC0121A	R1: 1 x 105ml R2: 5 x 20ml R4: 1 x 5ml
Urea (Lyo.)	BXC0121B	R1: 2 x 105ml R2: 10 x 20ml R4: 1 x 5ml
Urea (Monoliquid)	BXC0126A	R1: 1 x 100ml R4: 1 x 5ml

Methodology: Enzymatic
Sample Type: Human serum or plasma
Sample Volume: 10 µl
Storage: 2°C - 8°C

Uric Acid (L.S)	BXC0602A	R1: 4 x 50ml R2: 1 x 50ml R4: 1 x 5ml
Uric Acid (L.S)	BXC0602B	R1: 4 x 20ml R2: 1 x 20ml R4: 1 x 5ml
Uric Acid (Lyo.)	BXC0601A	1: 1 x 105ml R2: 5 x 20ml R4: 1 x 5ml
Uric Acid (Lyo.)	BXC0601B	R1: 2 x 105ml R2: 10 x 20ml R4: 1 x 5ml
Uric Acid (Monoliquid)	BXC0603A	R1: 5 x 20ml R4: 1 x 5ml
Uric Acid (Monoliquid)	BXC0603B	R1: 1 x 100ml R4: 1 x 5ml
Uric Acid (Monoliquid)	BXC0603C	R1: 5 x 100ml R4: 1 x 5ml

Methodology: Enzymatic

Sample Type: Human serum, plasma or urine

Storage: 2°C - 8°C

Urinary & CSF Protein (L.S)	BXC0174A	R1: 1 x 50ml R2: 1 x 10ml R4: 1 x 5 x 1ml
Urinary & CSF Protein (Monoliquid)	BXC0172A	R1: 2 x 60ml R4: 1 x 5ml
Urinary & CSF Protein (Monoliquid)	BXC0172B	R1: 4 x 60ml R4: 1 x 5ml

Methodology: Colorimetric

Sample Type: Urine and CSF fluid

Sample Volume: 30 µl

VITAMIN-D (L.S)	BXC0472A	R1: 1 x 9ml R2: 1 x 18ml R3: 1 x 9ml R4: 1 x 18ml CAL: 5 x 1ml
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Methodology: Enzymatic

Form: LS

Sample Type: Serum and plasma

Sample Volume: 3 µl

Measuring Range: 7.6- 147.8 ng/ml

Storage: 2°C - 8°C

Working Stability: Up to expiry when stored at 2-8°C

Zinc (L.S)	BXC0461A	R1: 2 x 50ml R2: 1 x 25ml R3: 1 x 25ml R4: 1 x 10ml
Zinc (Monoliquid)	BXC0462A	R1: 2 x 40ml R4: 1 x 5ml

Methodology: Latex Enhanced Turbidimetry

Form: LS

Sample Type: Serum or plasma

Sample Volume: 3 µl

Measuring Range: 7.6- 147.8 ng/ml

Storage: +2-8°C

Working Stability: Up to expiry when stored at 2-8°C

FORTRESS® DIAGNOSTICS

AQUEOUS STANDARDS



Fortress Diagnostics are a multi-award-winning global provider of in vitro diagnostics (IVDs). They develop, manufacture and support an extensive portfolio of clinical diagnostic tests and biochemistry solutions in the United Kingdom.

Albumin	BXCSTD01	45 g/l (4.5g/dL)	10 x 5ml
Calcium	BXCSTD02	Referenced against NIST 909b (2.5mmol/l/10mg/dL)	10 x 5ml
Chloride	BXCSTD03	Referenced against NIST 909b (75mmol/l)	10 x 5ml
Chloride	BXCSTD04	Referenced against NIST 909b (100mmol/l)	10 x 5ml
Cholesterol	BXCSTD05	Referenced against NIST 909b (5.17mmol/l/200mg/dL)	10 x 5ml
Creatinine	BXCSTD06	Referenced against NIST 909b (177 µmol/l/2 mg/dL)	10 x 5ml
Creatinine/Glucose/Urea Standard	BXCSTD24	Referenced against NIST 965a / 909b (5.55 mmol/l/100mg/dL/ 177 µmol/l / 8.33mmol/l)	10 x 5ml
Glucose	BXCSTD07	Referenced against NIST 965a (5.55 mmol/l/100mg/dL)	10 x 5ml
Haemoglobin Standard	BXCSTD08	For Drabkin's Method (18g/dL)	5 x 10 ml
Haemoglobin Standard Set	BXCSTD09	For Drabkin's Method (8, 10, 12, 15 & 18g/dL)	5 x 1 x 10ml
Iron	BXCSTD10	35.8 µmol/l (0.2 mg/dL)	10 x 5ml
Iron	BXCSTD11	107.4 µmol/l (0.6 mg/dL)	10 x 5ml
Lactate	BXCSTD12	4.44mmol/l (40mg/dL)	10 x 5ml

Magnesium	BXCSTD13	Referenced against NIST 909b (1.0 mmol/12.43mg/dL)	10 x 5ml
Phosphorus	BXCSTD14	1.61 mmol/l (5 mg/dL)	10 x 5ml
Potassium	BXCSTD15	Referenced against NIST 909b (0.5 mmol/l)	10 x 5ml
Potassium	BXCSTD16	Referenced against NIST 909b (5 mmol/l)	10 x 5ml
Pyruvate	BXCSTD17	2 mmol/l	10 x 5ml
Sodium	BXCSTD18	Referenced against NIST 909b (150.0 mmol/l150 mEq/l)	10 x 5ml
Total Protein	BXCSTD19	Referenced against NIST 927d (60g/l (6.0 g/dL)	10 x 5ml
Triglycerides	BXCSTD20	Referenced against NIST 909b (2.28 mmol/1200mg/dL)	10 x 5ml
Urea	BXCSTD21	Referenced against NIST 909b (8.33 mmol/150mg/dL)	10 x 5ml
Uric Acid	BXCSTD22	Referenced against NIST 909b (595 μ mol/110mg/dL)	10 x 5ml
Urinary/CSF Protein Standard	BXCSTD23	Referenced against NIST 927d (1g/l)	10 x 5ml

FORTRESS® DIAGNOSTICS

BLOOD GROUPING



Blood grouping reagents are used for the identification of blood types to ensure total transfusional compatibility between receivers and donors.

There are 43 blood group systems containing 345 antigens recognised by the International Society of Blood Transfusion however the most important blood groups in transfusion are the ABO blood group system and the RH blood group system.

Fortress Diagnostics offers both traditional methods and new techniques to identify the major and rare blood grouping types. The high-quality blood grouping reagents and gel cards are accurate, easy to use, competitively priced, and conveniently offered in different sizes in order to meet the needs of our distributors, blood banks, clinics, hospital laboratories and transfusion centres across the world.

AHG (Coomb's)	BGAG0010	Anti Human Globulin. A blend of rabbit Anti-Human IgG and Monoclonal Anti-Human c3d	1x10ml
AHG (Coomb's)	BGAG001L	Anti Human Globulin. A blend of rabbit Anti-Human IgG and Monoclonal Anti-Human c3d	1000ml
AHG (Coomb's)	BGAG1010	Anti Human Globulin. A blend of rabbit Anti-Human IgG and Monoclonal Anti-Human c3d	10x10ml
AHG (Coomb's)	BGAG2210	Anti Human Globulin. A blend of rabbit Anti-Human IgG and Monoclonal Anti-Human c3d	221x10ml

The Fortress Anti-Human Globulin (AHG) reagent is a blend of rabbit Anti-Human IgG and Monoclonal Anti-Human c3d and is available in various sizes.

Anti-c	BGLC0005	Monoclonal	1 x 5ml
Anti-C	BGBC0005	Monoclonal	1 x 5ml

The Fortress Diagnostics monoclonal IgM Anti- Rh blood grouping reagent contains human monoclonal antibodies diluted with sodium chloride, bovine albumin and macromolecular potentiators. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-CDE	BGCDE010	Polyclonal	1 x 10ml
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The Fortress Diagnostics Anti-C reagent is for the in vitro detection and identification of the human C blood group antigen by direct agglutination.

Anti-Cw	BGCW0002	Polyclonal	1 x 2ml
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The Fortress Diagnostics Anti-Cw blood grouping reagent contains human monoclonal IgM antibody, diluted in a phosphate buffer containing sodium chloride, bovine albumin and a preservative. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-e	BGLE0005	Monoclonal	1 x 5ml
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Anti-E	BGBE0005	Monoclonal	1 x 5ml
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The Fortress Diagnostics Anti-E reagent is for the in vitro detection and identification of the human E blood group antigen by direct agglutination.

Anti-H	BGBH0002	Lectin	1 x 2ml
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The Fortress Diagnostics Anti-H blood grouping reagent is prepared from a Ulex Europaeus seed extract, diluted with a sodium chloride solution containing bovine albumin. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-Lea	BGLEA002	Monoclonal	1 x 2ml
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Anti-Leb	BGLEB002	Monoclonal	1 x 2ml
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Fortress Diagnostics manufacture Anti LEa and Anti-LEb blood grouping reagents for the in vitro detection and identification of the human LEa and LEb blood group antigens by direct agglutination.

Anti-Lua	BGLUA002	Polyclonal	1 x 2ml
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Anti-Lub	BGLUB002	Polyclonal	1 x 2ml
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The Fortress Diagnostics Anti-Lua and Anti-Lub kits are Polyclonal Blood Grouping Reagent available in 1 x 2ml kits.

Anti-M	BGM00002	Polyclonal	1 x 2ml
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The Fortress Diagnostics Anti- M blood grouping reagent contains a murine monoclonal IgG antibody diluted in buffer containing sodium chloride, bovine albumin and a preservative. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-N	BGN00002	Monoclonal	1 x 2ml
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The Fortress Diagnostics Anti-N Lectin blood grouping reagent is prepared from an extract of Vicia unijugaleaves, diluted with a sodium chloride solution containing bovine albumin. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-P1	BGP10002	Monoclonal	1 x 2ml
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The Fortress Diagnostics Anti-P1 blood grouping reagent contains mouse monoclonal IgM antibodies prepared from the cell line, Clone 650, diluted in a solution containing sodium chloride and bovine albumin. This reagent is supplied ready for use and does not require any further dilution or addition.

Anti-S	BGBS0002	Monoclonal	1 x 2ml
Anti-s	BGLS0002	Monoclonal	1 x 2ml

The Fortress Diagnostics Anti-S and Anti-s reagents consist of human monoclonal IgG antibodies diluted in phosphate buffer containing sodium chloride and bovine albumin. The reagent requires no further dilution or addition for recommended techniques.

Low Ionic Strength Solution	BGLS001L	Ready to Use Low Ionic Strength Solution	1000ml
Low Ionic Strength Solution	BGLS1010	Ready to Use Low Ionic Strength Solution	10x10ml
Low Ionic Strength Solution	BGLS2210	Ready to Use Low Ionic Strength Solution	221x10ml
Low Ionic Strength Solution	BGLSS010	Ready to Use Low Ionic Strength Solution	1 x 10ml

The Fortress Diagnostics L.I.S.S. Ready for Use is a low ionic strength solution which is iso-osmotic with human red blood cells. The solution contains glycine, phosphate buffer and 0.03M sodium chloride.

Serological Albumin 22%	BG22001L	Caprylate Free. Free from stabilisers and non-specific agglutinins	1000ml
Serological Albumin 22%	BG221010	Caprylate Free. Free from stabilisers and non-specific agglutinins	10x10ml
Serological Albumin 22%	BGAL2210	Caprylate Free. Free from stabilisers and non-specific agglutinins	1 x 10ml
Serological Albumin 30%	BG30001L	Caprylate Free. Free from stabilizers and non-specific agglutinins	1000ml
Serological Albumin 30%	BG301010	Caprylate Free. Free from stabilisers and non-specific agglutinins	10x10ml
Serological Albumin 30%	BGAL3010	Caprylate Free. Free from stabilisers and non-specific agglutinins	1 x 10ml

The Fortress Diagnostics Serological Albumin 22% and 30% are prepared from bovine serum albumin and buffered saline. No artificial avidity enhancers or high molecular weight agglutination potentiators are added to the serological albumin solution. This reagent is supplied ready for use and does not require any further dilution or addition.

FORTRESS® DIAGNOSTICS

HAEMATOLOGY REAGENTS



Haematology involves the study, diagnosis and treatment of blood and bone marrow disorders. Fortress Diagnostics has developed a range of haematology reagents, controls and calibrators for use in detecting blood related disorders.

CELLDIFF+ CLEAN	HAEM042A	1 Litre
CELLDIFF+ CLEAN	HAEM042B	5 Litres
CELLDIFF+ DILUENT	HAEM034A	20 Litres
CELLDIFF+ ENHANCED CLEANER	HAEM033A	100 mls. Daily & weekly Maintenance Solution
CELLDIFF+ LYSE	HAEM043A	1 Litre
CELLDIFF-3+ Reagent Starter Pack	FORTSP01	5 Litres Diluent; 1 Litre Cleaner; 100mls Lyse.

Fortress Celldiff + products available include Celldiff +Clean, Celldiff + Enhanced Cleaner, Celldiff + Lyse and Celldiff -3+ Reagent Starter Pack.

Haematology Triple Pack Box	HAEM1000	20 Litres Diluent; 1 Litre Lyse; 5 Litres Cleaner
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The Fortress Haematology Triple Pack Box contains diluent, clean and lyse reagent for 1000 tests on the Fortress CELLDIFF-3+ haematology analyser.

Proteolyse	HAEM035A	50mls
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The Fortress CELLDIFF Proteolyse kit is intended for use as an enhanced cleaning solution for clot removal and aperture cleaning in the CELLDIFF fluid path.

FORTRESS® DIAGNOSTICS

HAEMOSTASIS COAGULATION REAGENTS



Haemostasis is the natural process in which blood flow slows and a clot forms to prevent blood loss when an injury occurs, with haemo meaning blood, and stasis meaning stopping. During haemostasis, blood changes from a fluid liquid to a gelatinous state.

Fortress Diagnostics manufactures, develops and supplies an extensive range of high quality haemostasis coagulation reagents.

WHY CHOOSE FORTRESS HAEMOSTASIS COAGULATION REAGENTS?

- Compatible with market leading analysers
- Controls available (see our QC and calibrators section)
- Flexibility: Lyophilised and liquid formats available, as well as a variety of kit sizes, to suit different laboratory sizes and needs
- Excellent stability, to ensure cost-effectiveness

APTT Reagent	COAG103A	Liquid Stable	1 x 3ml
APTT Reagent	COAG103B	Liquid Stable	5 x 3ml
APTT Reagent	COAG103C	Liquid Stable	1 x 10ml
APTT Reagent with Calcium Chloride	COAG110A	Liquid Stable	6 x 3ml; 2 x 10ml

The activated partial thromboplastin time (APTT) is used as a general screening test for the detection of coagulation abnormalities in the intrinsic pathway. The APTT is sensitive to deficiencies or abnormalities of factors VIII, IX, XI, XII, X and II, prekallikrein, high molecular weight kininogen (HMWK), and fibrinogen. APTT is also sensitive to inhibitors of blood coagulation such as lupus inhibitor and fibrin/fibrinogen degradation products. The APTT is the most widely used method for monitoring intravenous heparin anticoagulation therapy.

The Fortress Diagnostics APTT reagent is an in vitro diagnostic assay intended for use in determining activated partial thromboplastin time (APTT) and coagulation factor assays that are based on a modified APTT.

Calcium Chloride (0.02M)	COAG104A	Ready To Use for APTT	1 x 10ml
Calcium Chloride (0.02M)	COAG104B	Ready To Use for APTT	5 x 10ml

The Fortress Diagnostics APTT reagent is an in vitro diagnostic assay intended for use in determining activated partial thromboplastin time (APTT) and coagulation factor assays that are based on a modified APTT.

D-Dimer Latex	COAG115C	D-Dimer Latex Assay	25T
D-Dimer Latex	COAG115D	D-Dimer Latex Assay	50T
D-Dimer Latex	COAG115E	D-Dimer Latex Assay	100T

D-Dimer Latex Test is intended for the rapid qualitative or semi-quantitative evaluation of circulating derivatives of cross-linked fibrin degradation products (XL-FDP) in human plasma.

FACTOR II ASSAY KIT	COAG129A	5 x 1ml; 1 x 60ml
FACTOR IX ASSAY KIT	COAG127A	5 x 1ml; 1 x 60ml
FACTOR V ASSAY KIT	COAG128A	5 x 1ml; 1 x 60ml
FACTOR VII ASSAY KIT	COAG124A	5 x 1ml; 1 x 60ml
FACTOR VIII ASSAY KIT	COAG122A	5 x 1ml; 1 x 60ml
FACTOR X ASSAY KIT	COAG123A	5 x 1ml; 1 x 60ml
FACTOR XI ASSAY KIT	COAG133A	5 x 1ml; 1 x 60ml
FACTOR XII ASSAY KIT	COAG126A	5 x 1ml; 1 x 60ml

The Fortress Diagnostics Factor Deficient Substrate Plasma Assays are intended for the quantitative determination of the respective factor in patients suspected of having a congenital or acquired deficiency of this coagulation protein.

Fibrinogen Degradation Products (FDP)	COAG114A	Latex Assay	60 tests
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The Fortress Diagnostics Fibrinogen reagent is an in vitro diagnostic assay intended for quantitative determination of fibrinogen in plasma.

Fibrinogen Liquid Stable	COAG116A	Quantitative Estimation of Fibrinogen	1 x 10ml Buffer; 1 x 2ml Bovine Thrombin; 1 x 1ml Calibrator
Fibrinogen Liquid Stable	COAG116B	Quantitative Estimation of Fibrinogen	1 x 20ml Buffer; 2 x 2ml Bovine Thrombin; 1 x 1ml Calibrator
Fibrinogen Lyophilised	COAG105A	Quantitative Estimation of Fibrinogen	20 tests
Fibrinogen Lyophilised	COAG105B	Quantitative Estimation of Fibrinogen	60 tests
Fibrinogen Lyophilised	COAG105C	Quantitative Estimation of Fibrinogen	100 tests

The Fortress Diagnostics Fibrinogen assays are intended for use in the determination of fibrinogen in plasma.

HAEMOSTASIS CONTROL NORMAL	COAG131A	Including Factor Assays	1 ml
HAEMOSTASIS CONTROL PATHOLOGICAL	COAG132A	Including Factor Assays	1 ml

The Fortress Diagnostics Haemostasis Control is intended for use as a quality control plasma to monitor the precision of laboratory testing procedures for the analytes listed in this package insert.

Kaolin Clotting Time	COAG121A	For the Determination of Kaolin Clotting Time	R1: 1 x 2ml R2: 1 x 2ml
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The Fortress Diagnostics Kaolin Clotting Time reagent for the detection of Lupus anticoagulants.

Plasma Control Level II	COAG109B	For Coagulation Series only	5 x 1ml
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The Fortress Diagnostics Plasma Control Level 2 is intended for checking Precision and Accuracy for Coagulation assays PT, APTT and Fibrinogen.

Protein C	COAG118A	Quantitative Determination of Protein C in Human Plasma	40 TESTS
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Protein S	COAG119A	Quantitative Determination of Protein S in Human Plasma	40 TESTS
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The Fortress Diagnostics Protein C and S assays are intended for the determination of Protein C/S activity in human plasma.

PT High Sensitivity	COAG101A	Liquid Stable, ISI ~1.15	5 x 1ml
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PT High Sensitivity	COAG101B	Liquid Stable, ISI ~1.15	5 x 5ml
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PT High Sensitivity	COAG101C	Liquid Stable, ISI ~1.15	10 x 5ml
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PT High Sensitivity	COAG101D	Liquid Stable, ISI ~1.15	6 x 8ml
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PT High Sensitivity	COAG101E	Liquid Stable, ISI ~1.15	6 x 5ml
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PT High Sensitivity	COAG101F	Liquid Stable, ISI ~1.15	20 x 5ml
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PT High Sensitivity	COAG101G	Liquid Stable, ISI ~1.15	20 x 8ml
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PT Low Sensitivity	COAG102A	Liquid Stable, ISI ~1.5	1 x 5ml
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PT Low Sensitivity	COAG102B	Liquid Stable, ISI ~1.5	5 x 5ml
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PT Low Sensitivity	COAG117A	Liquid Stable, ISI ~1.8	1 x 5ml
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PT Low Sensitivity	COAG117B	Liquid Stable, ISI ~1.8	5 x 5ml
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The Fortress Diagnostics PT reagent is an in vitro diagnostic assay intended for use in performing the one stage prothrombin time (PT) test and assays which are based on a modified prothrombin time.

Thrombin Time Reagent	COAG107A	For Qualitative estimation of Fibrinogen	5 x 1ml
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The Fortress Diagnostics Thrombin Time (Bovine) is an in vitro diagnostic assay intended for qualitative determination of fibrinogen in plasma.

FORTRESS® DIAGNOSTICS

LATEX SEROLOGY



Latex agglutination testing, also called latex serology, is widely used as a laboratory method to identify certain antibodies and antigens. The test uses a variety of sample types including blood, urine, saliva and cerebrospinal fluid, depending on what type of sample is needed. Fortress Diagnostics manufactures, develops and supplies an extensive range of high-quality latex serology reagents.

ASO	5ml Latex	LXASO005	1 x 5ml
ASO	1000ml Latex	LXASO01L	25,000T
ASO	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXASO025	25T
ASO	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXASO050	50T
ASO	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXASO100	100T
ASO	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXASO150	150T

The Fortress Diagnostics ASO-Latex agglutination is the rapid and simple test for the qualitative and semi-quantitative measurements of antibodies to Anti-Streptolysin-O (ASO) in human serum.

When the latex reagent is mixed with a serum containing ASO, agglutination occurs. In acute streptococcal infection the toxic immunogenic exoenzyme Streptolysin-O (ASO) is produced in response to Streptolysin O antigens liberated by haemolytic streptococci of groups A, C and G.

Infectious Mononucleosis	LXIM00020	20T
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The Fortress Diagnostics Infectious Mononucleosis (IM) test is a rapid slide test for the qualitative detection of heterophile antibodies to IM in human serum. The reagent agglutinates when mixed with serum containing the heterophile antibodies.

CRP	5ml Latex	LXCRP005	1 x 5ml
CRP	1000ml Latex	LXCRP01L	25,000T
CRP	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXCRP025	25T
CRP	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXCRP050	50T
CRP	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXCRP100	100T
CRP	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXCRP150	150T

The Fortress Diagnostics CRP Latex test is a rapid slide agglutination test for the qualitative and semi-quantitative detection of C-Reactive Protein in serum. The reagent containing particles coated with specific anti-human C-Reactive protein antibodies, agglutinates in the presence of CRP in the patient's serum.

hCG White Latex	1000ml Latex	LXHCW01L	25,000T
hCG White Latex	Sensitivity 200IU/ml - White Latex. Kit contains: Latex reagent, +ve and -ve controls, pipettes/stirrers and test	LXHCW025	25T
hCG White Latex	Sensitivity 200IU/ml - White Latex. Kit contains: Latex reagent, +ve and -ve controls, pipettes/stirrers and test	LXHCW050	50T
hCG White Latex	Sensitivity 200IU/ml - White Latex. Kit contains: Latex reagent, +ve and -ve controls, pipettes/stirrers and test	LXHCW100	100T

The Fortress hCG Latex test is a rapid slide agglutination procedure, developed for the direct detection of hCG in urine. The presence or absence of a visible agglutination indicates the presence or absence of hCG in the sample tested.

RF	5ml Latex	LXRF0005	1 x 5ml
RF	1000ml Latex	LXRF001L	25,000T
RF	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXRF0025	25T
RF	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXRF0050	50T
RF	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXRF0100	100T
RF	Latex Reagent, +ve and -ve Controls, Pipette/Stirrers & Re-usable Test Slide	LXRF0150	150T

The Fortress Diagnostics RF-Latex test is a rapid slide agglutination test for the qualitative and semi-quantitative detection of Rheumatoid Factors in serum. Latex suspension coated with human gamma-globulin, agglutinates in the presence of rheumatoid factors in the patient serum.

Rheumatoid factors are antibodies directed against antigenic sites in the Fc fragment of human and animal IgG. Their frequent occurrence in rheumatoid arthritis makes them useful for diagnosis and monitoring of the disease, although RF is also found in a number of rheumatic disorders such as Sjogren's syndrome and Lupus erythematosus (SLE).

FORTRESS® DIAGNOSTICS

QUALITY CONTROLS AND CALIBRATORS



Fortress Diagnostics' quality control range Seraqual is utilised in laboratories across the world for accurate results you can rely on. The Seraqual controls & calibrators are designed and manufactured in-house to the highest quality standards. Kits are supplied with methodology and instrumentation-related comparisons.

WHY CHOOSE FORTRESS DIAGNOSTICS QUALITY CONTROLS?

- Manufactured in-house for full control and traceability.
- Extensive and ever-growing range available.
- Custom quality control service available.
- Highly compatible, designed to mimic human samples with minimal matrix effects being observed.
- Offer a wide parameter range allowing more analyte coverage single control.

ADA CALIBRATOR	BXC0209A	1 x 5ml
ADA CONTROL LEVEL-1	BXC0210A	1 x 5ml
ADA CONTROL LEVEL-2	BXC0211A	1 x 5ml

The Fortress Diagnostics ADA Calibrator is intended to be used with the Fortress ADA Liquid stable kit. The calibrator is used for standardising Fortress ADA Liquid stable kit

Alcohol, Ammonia, Carbonate Calibrator	BXC0492A	1 x 2ml
Alcohol, Ammonia, Carbonate Control Level 1	BXC0493A	1 x 2ml
Alcohol, Ammonia, Carbonate Control Level 2	BXC0494A	1 x 2ml

The Fortress Diagnostics alcohol/ammonia/carbonate control is intended to be used with Fortress alcohol, ammonia and carbonate kits. The control can be used for assessing accuracy and precision of the assays.

Aldolase Calibrator	BXC0394A	3 x 1ml
Aldolase Control Normal	BXC0392A	3 x 1ml
Aldolase Control Elevated	BXC0393A	3 x 1ml

The Fortress Diagnostics Aldolase Control Level 1 is intended to be used with the Fortress Aldolase Kit.

Alpha-1-Antitrypsin Calibrator	BXC0711A	1 x 2ml
Alpha-1-Antitrypsin Control L1 & L2	BXC0712A	2 x 1 x 2ml

The Fortress Diagnostics Alpha-1-Antitrypsin Calibration and Control (levels 1 & 2) are intended for use in the control and calibration of Alpha-1-Antitrypsin assays.

Ammonia Calibrator	BXC0373A	3 x 2ml
Ammonia Control Low	BXC0374A	3 x 2ml
Ammonia Control High	BXC0375A	3 x 2ml

The Fortress Diagnostics Ammonia control (low & high) and calibrator kits are intended for use in assessing the accuracy and precision of Ammonia Liquid stable kits.

Angiotensin Converting Enzyme (ACE) Calibrator	BXC0177A	3 x 1ml
Angiotensin Converting Enzyme (ACE) Control Level 1	BXC0178A	3 x 1ml
Angiotensin Converting Enzyme (ACE) Control Level 2	BXC0179A	3 x 1ml

The Fortress Diagnostics Angiotensin Converting Enzyme (ACE) Calibrator is intended to be used as a calibration material for the ACE assay.

APO A1/B CALIBRATOR SERIES	BXC0413A	1 x 6 x 1ml
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The Fortress Diagnostics Apolipoprotein (APO) A1 & Apolipoprotein B Calibrator is intended for calibrating the Fortress Apolipoprotein A1 and B assays.

ASO (Single Point) Calibrator	BXC0323A	1 x 1ml
ASO (Single Point) Calibrator	BXC0323B	3 x 1ml
ASO, CRP, RF CONTROL	BXC0645A	3 x 1ml

The Fortress Diagnostics ASO, CRP and RF controls are intended for use as an assayed quality control material to monitor the consistency of performance of laboratory test procedures associated with determination and monitoring of the clinical status. The product is human serum-based, liquid stable control, stabilised with preservatives.

Bilirubin Calibrator	BXC0319A	3 x 1ml
Bilirubin Calibrator, Liquid Stable	BXC0303A	2 x 1ml
Bilirubin Control, Liquid Stable	BXC0306A	1 x 1ml
Bilirubin Control, Liquid Stable	BXC0306B	3 x 1ml
Bilirubin Control, Lyo.	BXC0318A	1 x 1ml
Bilirubin Control, Lyo.	BXC0318B	5 x 1ml
Bilirubin Control, Liquid Stable	BXC0307A	1 x 1ml
Bilirubin Control, Liquid Stable	BXC0307B	3 x 1ml
Bilirubin Control Set, Liquid Stable	BXC0304A	2 x 1 x 1ml
Bilirubin Control Set, Paediatric, Liquid Stable	BXC0305A	2 x 1 x 1ml

The Fortress Diagnostics Bilirubin Liquid Stable control is ready to use and stable up to the expiry date when stored at 2-8°C. Once opened the control is stable for 6 months when stored at 2-8°C and kept in the dark.

Blood Gas & Electrolyte Control Level 1	BXC0108A	1 x 2ml
Blood Gas & Electrolyte Control Level 1	BXC0108D	10 x 2ml
Blood Gas & Electrolyte Control Level 2	BXC0108B	1 x 2ml
Blood Gas & Electrolyte Control Level 2	BXC0108E	10 x 2ml
Blood Gas & Electrolyte Control Level 3	BXC0108C	1 x 2ml
Blood Gas & Electrolyte Control Level 3	BXC0108F	10 x 2ml

The Fortress Diagnostics Blood Gas and Electrolyte controls are intended for use in the precision and accuracy assessment of Blood Gas, Electrolyte and Metabolite assays.

Cardiac Control Set	BXC0455C	2 x 1 x 2ml
Cardiac Control Set	BXC0456C	2 x 1 x 1ml
Cardiac Control-Level-I	BXC0455A	5 x 2ml
Cardiac Control-Level-I	BXC0456A	5 x 1ml
Cardiac Control-Level-II	BXC0455B	5 x 2ml
Cardiac Control-Level-II	BXC0456B	5 x 1ml

Cardiac quality controls are used to make sure that diagnostic tests for cardiac assessment, as well as analytical systems, are performing properly. Cardiac markers with Fortress Diagnostics kits include CK, CKMB, Tni, TnT, hsCRP, digitoxin, Homocysteine, BPB, NT-ProBNP.

Calibration Serum	BXC0321K	1 x 3ml
Calibration Serum	BXC0321L	5 x 3ml
Calibration Serum	BXC0321M	10 x 3ml

The Fortress Diagnostics Blood Gas and Electrolyte controls are intended for use in the precision and accuracy assessment of Blood Gas, Electrolyte and Metabolite assays.

Citrate (Urinary) Calibrator (200mg/L)	BXC0136A	2 x 1ml
Citrate (Urinary) Control L1 (50mg/L)	BXC0137A	2 x 1ml
Citrate (Urinary) Control L2 (600mg/L)	BXC0139A	2 x 1ml

The Fortress Diagnostics Citrate Control is intended for use in assessing the precision and accuracy of the Citrate Liquid stable assay.

CK / CKMB Calibrator	BXC0454A	1 x 2ml
CK / CKMB Calibrator	BXC0454B	5 x 2ml
CK / CKMB Control Level-1	BXC0453A	1 x 2ml
CK / CKMB Control Level-1	BXC0453B	5 x 2ml
CK / CKMB Control Level-2	BXC0459A	1 x 2ml
CK / CKMB Control Level-2	BXC0459B	5 x 2ml

The Fortress Diagnostics CK-CKMB Controls and Calibrators are intended for checking accuracy and precision of CK MB and CK NAC on automated and semi-automated analysers. All expected values are provided for guidance only until each laboratory establishes its own values and SD for its methods.

CO2 (Bicarbonate) Calibrator	BXC0155A	3 x 2ml
CO2 (Bicarbonate) Control Low	BXC0156A	3 x 2ml
CO2 (Bicarbonate) Control High	BXC0157A	3 x 2ml

The Fortress Diagnostics CO2 Bicarbonate Control High is intended for checking the accuracy and the precision of the Bicarbonate Kit.

CSF Control Level I	BXC0673A	1 x 1ml
CSF Control Level II	BXC0673B	1 x 1ml

The Fortress Diagnostics CSF Control Level 1 & Level 2 kits are intended for checking accuracy and precision of laboratory methods for: Chloride, Glucose, Lactate, Microalbumin, Sodium, Urinary Protein, IGG

CRP (Multi Point) Calibrator	BXC0324B	1 x 6 x 1ml
CRP (Single Point) Calibrator	BXC0324A	3 x 1ml
CRP (ULTRA SENSITIVE) Low Control	BXC0325A	3 x 1ml
CRP (ULTRA SENSITIVE) Standard Set	BXC0327A	5 x 1 x 0.5ml
CRP Control Level 1	BXC0326A	5 x 1ml
CRP Control Level 2	BXC0326B	5 x 1ml

The Fortress Diagnostics CRP control kits are intended for use as an assayed quality control material to monitor the consistency of performance of laboratory test procedures associated with determination and monitoring of the clinical status. The product is human serum based, liquid stable control, stabilised with preservatives.

Cyanmethaemoglobin Standard Set (8,10,12,15,18 g/dL)	BXC0483A	5 x 1 x 10ml
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The Fortress Diagnostics Cyanmethaemoglobin Standard Set is intended for use in assessing the precision and accuracy of Cyanmethaemoglobin assays.

Cystatin C Calibrator Set Level 1 - 6	BXC0334A	6 x 1ml
Cystatin C Control Level 1	BXC0333A	2 x 1ml
Cystatin C Control Level 2	BXC0333B	2 x 1ml

The Fortress Diagnostics Cystatin C control and calibrator sets are intended for use with the Fortress Latex Enhanced Cystatin C Assay.

D3-Hydroxybutyrate Control, Level 1	BXC0543A	6 x 1ml
D3-Hydroxybutyrate Control, Level 2	BXC0543B	2 x 1ml

The Fortress Diagnostics D3HBD Control is intended to be used in assessing the accuracy and precision of D3 Hydroxybutyrate assays.

D-DIMER Immunoturbidimetric Calibrator Set (L1-L5)	BXC0788A	5 x 1ml
D-DIMER Immunoturbidimetric Control Set (L1,L2)	BXC0789A	2 x 1ml

The Fortress Diagnostics D-Dimer Calibrator is intended for use in the calibration of the Fortress D-Dimer Assay.

Drugs of Abuse Control Level 1	BXC0784A	10 x 10ml
Drugs of Abuse Control Level 2	BXC0785A	10 x 10ml
Drugs of Abuse Control Level 3	BXC0786A	10 x 10ml

The Fortress Diagnostics Drugs of Abuse control is intended for use as an in vitro diagnostic tool to monitor the performance and precision of laboratory urine toxicology screening procedures.

Electrolytes Calibrator	BXC0140A	5 x 5ml
Electrolytes Control Low (Na, K, Cl)	BXC0143A	5 x 5ml
Electrolytes Control Normal (Na, K, Cl)	BXC0144A	5 x 5ml
Electrolytes Control High (Na, K, Cl)	BXC0145A	5 x 5ml

The Fortress Diagnostics Electrolyte Control Level 1-3 (low, normal & high) kits are intended for use on ISE systems for accuracy and precision checks.

ESR Control Level 1	BXC0631A	2 x 10ml
ESR Control Level 2	BXC0632A	2 x 10ml

The Fortress Diagnostics Erythrocyte Sedimentation Rate (ESR) Control kits are intended for use as a quality control tool to measure the precision of ESR laboratory testing procedures.

FERRITIN CONTROL LEVEL-1	BXC0443A	1 x 1ml
FERRITIN CONTROL LEVEL-2	BXC0444A	1 x 1ml

The Fortress Diagnostics Ferritin Control (Levels 1 & 2) and calibration kits are intended for use as an assayed quality control material to monitor the performance consistency of Ferritin assays. The controls are human serum based, liquid stable control and stabilised with preservatives.

Fructosamine Calibrator	BXC0592A	1 x 1ml
Fructosamine Control Low	BXC0593A	3 x 1ml
Fructosamine Control High	BXC0594A	3 x 1ml

The Fortress Diagnostics Fructosamine controls (levels 1 & 2) are intended for use as a Quality Control method for assessing the precision and accuracy of Fructosamine Assays.

G-6-PDH Control Deficient	BXC0572A	3 x 0.5ml
G-6-PDH Control Normal	BXC0573A	3 x 0.5ml

The Fortress Diagnostics G-6-PDH control (deficient & normal) kits are intended for use as an accuracy and precision control for G-6-PDH testing.

Glutathione Peroxidase Control	BXC0556A	5 x 1ml
Glutathione Reductase Control	BXC0460A	5 x 1ml

The Fortress Diagnostics Glutathione control (Peroxidase & Reductase) controls are intended for use in the analysis of precision and accuracy of the Fortress Glutathione Peroxidase kit.

Glycerol Control	BXC0278A	2 x 5ml
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The Fortress Diagnostics Glycerol Control kit is intended for use in assessing the precision and accuracy of glycerol assays.

HAEMATOLOGY CALIBRATOR (for 3-part Differential Analysers)	HAEMC005	3 x 3ml
HAEMATOLOGY CONTROL LOW	HAEMC001	3 x 3ml
HAEMATOLOGY CONTROL NORMAL	HAEMC002	3 x 3ml
HAEMATOLOGY CONTROL HIGH	HAEMC003	3 x 3ml
HAEMATOLOGY CONTROL, 3-LEVELS	HAEMC004	3 x 3 x 3ml
HAEMATOLOGY CONTROL, 3- LEVELS	HAEMC313	3 x 1 x 3ml
HAEMATOLOGY CONTROL, 3- LEVELS	HAEMC004	3 x 3 x 3ml

The Fortress Diagnostics Haematology Controls are intended for use in monitoring the accuracy and precision of blood cell counts on Haematology analysers.

HbA1c (Control Set 2 Levels)	BXC0675A	2 x 2 x 0.5ml
HbA1c Control Low	BXC0676A	2 x 0.5ml
HbA1c Control High	BXC0677A	2 x 0.5ml
HbA1c Calibrator Series 2 Levels	BXC0678A	2 x 1 x 0.5ml
HbA1c (HPLC) Control Low	BXC0779A	2 x 0.5ml
HbA1c (HPLC) Control High	BXC0780A	2 x 0.5ml

The Fortress Diagnostics HbA1c Control kits are intended for use as an assayed quality control material to monitor the precision of laboratory testing procedures for multiple analytes. The HbA1c Calibrator kits are intended for the preparation of a standard curve for calibration of HbA1c assays.

hCG Serum Control Positive	BXC0681A	1 x 1ml
hCG Serum Control Negative	BXC0682A	1 x 1ml
hCG Serum Control Positive	BXC0683A	1 x 5ml

The Fortress hCG serum control is intended for use in assessing the precision and accuracy of hCG assays.

HDL/LDL Calibrator	BXC0315B	1 x 3ml
HDL/LDL Calibrator	BXC0315C	5 x 3ml
HDL/LDL Calibrator	BXC0315D	5 x 1ml

The Fortress Diagnostics HDL/LDL calibrator is intended for use with the Fortress HDL and LDL Cholesterol assays.

Homocysteine Calibrator 5 Levels	BXC0691A	5 x 1 x 1ml
Homocysteine Calibrator 2 Levels	BXC0691B	2 x 1 x 1ml
Homocysteine Control 2 Levels	BXC0692A	2 x 1 x 1ml

Fortress Diagnostics have Homocysteine control and calibrator sets available. The Homocysteine Control Set is intended for checking the accuracy and precision of the Homocysteine assay and the Calibrator Set is intended for calibrating the Homocysteine assay.

Human Assayed Control (Normal)	BXC0312A	10 x 5ml
Human Assayed Control (Elevated)	BXC0312B	10 x 5ml
Human Assayed Control (Normal)	BXC0312C	5 x 5ml
Human Assayed Control (Elevated)	BXC0312D	5 x 5ml
Human Assayed Control (Normal)	BXC0312E	1 x 5ml
Human Assayed Control (Elevated)	BXC0312F	1 x 5ml

The Fortress Diagnostics Human Assayed Controls are lyophilised human serum based controls, intended for laboratory use. These are used to monitor the precision and accuracy of substrates, electrolytes and enzymes testing procedures.

Human Precision Control (Normal)	BXC0314A	10 x 5ml
Human Precision Control (Elevated)	BXC0314B	10 x 5ml
Human Precision Control (Normal)	BXC0314C	5 x 5ml
Human Precision Control (Elevated)	BXC0314D	5 x 5ml
Human Precision Control (Normal)	BXC0314E	1 x 5ml
Human Precision Control (Elevated)	BXC0314F	1 x 5ml

The Fortress Diagnostics human precision controls are lyophilised human serum based, intended for use in the laboratory as precision controls. These are ideal tools for testing reproducibility.

Immunoassay Control Level I	BXC0363A	4 x 3ml
Immunoassay Control Level II	BXC0363B	4 x 3ml
Immunoassay Control Level III	BXC0363C	4 x 3ml
Immunoassay Tri-Level Control	BXC0363D	3 x 4 x 3ml

The Fortress Diagnostics Immunoassay controls are intended for use as an assayed quality control material to monitor the consistency of laboratory performance test procedures associated with determination and monitoring of clinical status. The controls are human based, lyophilized controls and can be used with RIA, EIA, ELISA or FIA methods.

Lipase Calibrator	BXC0510A	3 x 1ml
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The Fortress Diagnostics Lipase Calibrator is intended for use as a calibrator for Fortress Lipase assays.

Lipid Calibrator	BXC0317A	3 x 1ml
Lipid Control Elevated	BXC0316A	3 x 1ml
Lipid Control Normal	BXC0330A	3 x 1ml

Fortress Diagnostics have 2 Lipid controls and a Lipid calibrator available in lyophilized format. The parameters covered by control 1 include: Normal, Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL. The parameters covered by control 2 include: Elevated, Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL. The parameters covered by this calibrator include: Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL.

Lipoprotein (a) Calibrator High	BXC0134A	1 x 1ml
Lipoprotein (a) Control Low	BXC0131A	1 x 1ml
Lipoprotein (a) Control High	BXC0133A	1 x 1ml

The Fortress Diagnostics Lipoprotein calibrator and control kits are intended for use with the Fortress Lipoprotein assay.

Lithium Calibrator (3000 mmol/L)	BXC0106A	2 x 1ml
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The Fortress Diagnostics Lithium calibrator is intended for use in the calibration of Lithium assays.

Microalbumin Calibrator Series	BXC0329A	5 x 1ml
Microalbumin Control	BXC0328A	2 x 1ml
Microalbumin Control	BXC0328B	5 x 1ml

Fortress Diagnostics have Microalbumin calibrator and control kits available. The calibrator Liquid Stable is for the calibration of quantitative immunoturbidimetric assays for Microalbumin (Albumin in Urine) and the control kits is intended for monitoring the performance of quantitative immunoturbidimetric assays for microalbumin (albumin in urine). Multiple levels of control are available to allow monitoring of the test system's reliability.

NTProBNP Calibrator	BXC0335A	1 x 1ml
NTProBNP Control Set Level 1 - 6	BXC0336A	1 x 6 x 1ml

The Fortress Diagnostics NTProBNP calibrator and control series is designed for use in assessing the precision and accuracy of NTProBNP assays.

Oxalate (Urinary) Calibrator 0.5 mmol/L	BXC0147A	1 x 5ml
Oxalate (Urinary) Control L1 (0.2mmol/L)	BXC0148A	1 x 5ml
Oxalate (Urinary) Control L2 (0.7mmol/L)	BXC0149A	1 x 5ml

The Fortress Diagnostic calibrator and control sets are intended for use in assessing the accuracy and precision of Oxalate (Urinary) assays.

Paediatric Control**BXC0807A****3 x 1ml**

The Fortress Diagnostics Paediatric control is intended for use as an assayed quality control serum to monitor the precision of multiple analytes including; Bilirubin, Calcium, Chloride, Glucose, Magnesium, Potassium, Sodium and Phosphorous Inorganic.

PAPP-A & fβhCG Controls Level 1**BXC0337A****1 x 0.5ml****PAPP-A & fβhCG Controls Level 2****BXC0338A****1 x 0.5ml****PAPP-A & fβhCG Controls Level 3****BXC0339A****1 x 0.5ml****PAPP-A Control****BXC0240A****1 x 0.5ml**

Pregnancy associated plasma protein A (PAPPA) is another metalloproteinase, originally discovered as a glycoprotein in pregnant women, produced by the syncytiotrophoblasts of the placenta. The Fortress Diagnostics PAPP-A control levels are intended for use in assessing the precision and accuracy of PAPP-A assays.

Plasma Calibrator**COAG125A****5 x 1ml****Plasma Control Level I****COAG108A****1 x 1ml****Plasma Control Level I****COAG108B****5 x 1ml****Plasma Control Level I & II****COAG112A****2 x 5 x 1ml****Plasma Control Level II****COAG109A****1 x 1ml****Plasma Control Level II****COAG109B****5 x 1ml**

The Fortress Diagnostics Plasma control level kits are intended for assessing the precision and accuracy in Coagulation Assays, PT, APTT and Fibrinogen.

Protein Control Level 1**BXC0641A****1 x 2ml****Protein Control Level 2****BXC0642A****1 x 2ml****Protein Control Set Level 1 and 2****BXC0643A****2 x 1 x 2ml**

The Fortress Diagnostics Protein Controls are intended for use as an assayed quality control material to monitor the consistency of performance of laboratory test procedures associated with determination and monitoring of the clinical status. This product is human serum based, liquid stable control, stabilised with preservatives.

Retinol Binding Protein Control**BXC0995A****1 x 1ml**

Retinol-binding proteins are a family of proteins with diverse functions. They are carrier proteins that bind retinol, the assessment of retinol-binding protein is used to determine visceral protein mass in health-related nutritional studies. The Fortress Diagnostics retinol binding protein control is a liquid stable control available in 1x1ml kit size.

RF (Multi-Point) Calibrator	BXC0325B	3 x 2ml
RF (Multi-Point) Calibrator	BXC0612A	1 x 2ml

The Fortress Diagnostics Rheumatoid Factor (RF) Mutlipoint Calibrator is intended for the calibration of the Fortress Diagnostics RF Latex Immunoturbidmetry assay. The calibrator is made from human serum base and is presented as a ready to use material that is stable when stored at 2-8°C.

Specific Protein Calibrator	BXC0644A	1 x 2ml
Specific Protein Calibrator Set	BXC0646A	1 x 6 x 1ml

The Fortress Diagnostics Specific Protein calibrator is intended for use in the preparation of reference curves for the quantitative determination of various proteins in human serum with protein assays. The calibrator set is prepared for human serum.

Superoxide Dismutase Control	BXC0433A	5 x 1ml
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The Fortress Diagnostics Superoxide Dismutase control is intended for determining the accuracy and precision control of the Fortress Superoxide Dismutase Kit.

Syphilis Control Panel	BXC0806A	6 x 1 x 0.5ml
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The Fortress Diagnostics Syphilis control panel is intended for checking the accuracy and precision of syphilis assays.

Thalassaemia (α & β) Control Level 1	BXC0665A	2 x 0.5ml
Thalassaemia (α & β) Control Level 2	BXC0665B	2 x 0.5ml

The Fortress Diagnostics Thalassaemia control is available for use on all major systems & Methods including HPLC & Immunoassay , HbA2, HbF, HbS

Therapeutic Drug Monitoring Level I	BXC0781A	5 x 5ml
Therapeutic Drug Monitoring Level II	BXC0782A	5 x 5ml
Therapeutic Drug Monitoring Level III	BXC0783A	5 x 5ml

The Fortress Diagnostics Therapeutic Drug Monitoring Control (TDM) kit is intended for use as an assayed quality control serum to monitor the precision of laboratory testing procedures for a range of analytes; for a full list, please contact us directly.

Total Antioxidant Status (TAS) Control	BXC0554A	10 x 5ml
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The Fortress Total Antioxidant Status control and calibrator kits (TAS) are intended for assessing the accuracy, precision and calibration of the Fortress Total Antioxidant Status Kit.

Troponin-I Calibrator Set	BXC0473A	6 x 1 x 1ml
Troponin-I Control Set	BXC0470A	2 x 1 x 1ml
Troponin-T High Sensitivity Control	BXC0457A	3 x 1ml

The Fortress Diagnostics Troponin-T High Sensitivity Control is available for use on all major systems.

Urine Assayed Control Level I	BXC0661A	10 x 10ml
Urine Assayed Control Level II	BXC0661B	10 x 10ml

The Fortress Diagnostics Urine Assayed Control (levels 1 & 2) kits are intended for in vitro diagnostic use in the quality control of urine on clinical chemistry systems.

Urine Precision Control Level I	BXC0662A	10 x 10ml
Urine Precision Control Level II	BXC0662B	10 x 10ml

The Fortress Diagnostics Urine Precision control (levels 1 & 2) kits are intended for in vitro diagnostic use in the quality control of urine on clinical chemistry systems.

Urine Strip Control Level I & II	BXC0663A	2 x 3 x 12ml
Urine Strip Control Level I	BXC0663D	3 x 12ml
Urine Strip Control Level II	BXC0663E	3 x 12ml

The Fortress Diagnostics Urine Strip control (levels 1 & 2) kits are intended for use as a tool to assay the precision of Urine and hCG Strip assays.

VITAMIN-D Calibrator Set	BXC0475A	5 x 1ml
VITAMIN-D Control Set	BXC0474A	2 x 1ml

The Fortress Diagnostics Vitamin-D Calibrator and Control kits are intended for use with Vitamin D assays.

Zinc & Copper Calibrator (Lyo.)	BXC0463A	2 x 1ml
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The Fortress Diagnostics Zinc & Copper calibrator is a lyophilized kit intended for use with Zinc & Copper Assays.

AUDIT® MICROCONTROLS™

QUALITY CONTROLS AND CALIBRATION VERIFICATION/LINEARITY



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CALIBRATION VERIFICATION/LINEARITY

GENERAL CHEMISTRY

Linearity LQ Ammonia/Ethanol	K712M-5	5 x 2ml
Analytes: Ammonia, Ethanol		
Linearity FD General Chemistry	K701M-5	10 x 5ml
Analytes: Albumin, Alkaline Phosphatase, ALT (SGPT), Amylase, AST (SGOT), Bilirubin (Total and Direct), BUN, Calcium, Chloride, Cholesterol, CO ₂ , Creatine Kinase (CK), Creatinine, Gamma-GT, Glucose, HDL Cholesterol, Iron, Lactate, LDH, LDL Cholesterol, Lipase, Magnesium, Osmolality, Phosphorus, Potassium, Sodium, TIBC, Total Protein, Triglycerides, Uric Acid		
Linearity LQ ISE	K726M-5	5 x 5ml
Analytes: Chloride, Potassium, Sodium		

Linearity FD Lipids	K709M-5	5 x 2ml
Analytes: Apo A, Apo B, Cholesterol, HDL Cholesterol, LDL Cholesterol, Triglycerides		
Linearity FD UIBC	K732M-5	5 x 1ml
Analytes: UIBC		
Linearity FD General Chemistry (for Beckman AU)	K820M-5	10 x 5ml
Analytes: Albumin, Alkaline Phosphatase, ALT (SGPT), Amylase, AST (SGOT), Bilirubin (Total and Direct), BUN, Calcium, Chloride, Cholesterol, CO ₂ , Creatine Kinase (CK), Creatinine, Gamma-GT, Glucose, HDL Cholesterol, Iron, Lactate, LDH, LDL Cholesterol, Lipase, Magnesium, Phosphorus, Potassium, Sodium, Total Protein, Triglycerides, Uric Acid		
Linearity LQ Ammonia/Ethanol (for Roche Systems)	K881M-5	5 x 2ml
Analytes: Ammonia, Ethanol		
Linearity FD Bilirubin (for Roche Systems)	K886M-5	5 x 1ml
Analytes: Bilirubin (Total and Direct)		
Linearity FD Lipids (for Roche Systems)	K882M-5	5 x 2ml
Analytes: Apo A, Apo B, Cholesterol, HDL Cholesterol, LDL Cholesterol, Triglycerides		
Linearity FD General Chemistry (for Roche Systems)	K880M-5	10 x 5ml
Analytes: Albumin, Alkaline Phosphatase, ALT (SGPT), Amylase, AST (SGOT), Bilirubin (Total), BUN, Calcium, Chloride, CO ₂ , Creatine Kinase (CK), Creatinine, Gamma-GT, Glucose, Iron, Lactate, LDH, Lipase, Magnesium, Phosphorus, Potassium, Sodium, Total Protein, Uric Acid		
CARDIAC		
Linearity FD BNP (High)	K713M-5	5 x 1ml
Analytes: BNP-32, NT-ProBNP		
Linearity FD BNP (Low)	K708M-5	5 x 1ml
Analytes: BNP-32, NT-ProBNP		
Linearity FD BNP (Extended)	K737M-5	5 x 1ml
Analytes: BNP-32, NT-ProBNP		
Linearity FD Cardiac Markers	K710M-5	5 x 1ml
Analytes: CK-MB, Myoglobin, Troponin I		

Linearity FD D-Dimer	K717M-5	5 x 1ml
Analytes: D-Dimer		
Linearity LQ Homocysteine	K704M-5	5 x 1ml
Analytes: Homocysteine		
Linearity LQ hs-CRP	K705M-5	5 x 1ml
Analytes: hs-CRP		
Linearity FD D-Dimer (PATHFAST)	K883M-5	5 x 1ml
Analytes: D-Dimer		
Linearity FD BNP (RAMP)	K888M-5	5 x 1ml
Analytes: NT-ProBNP		
Linearity FD Cardiac (RAMP)	K887M-5	5 x 1ml
Analytes: CK-MB, Myoglobin, Troponin I		
Linearity FD BNP (Siemens Centaur)	K803M-5	5 x 1ml
Analytes: BNP-32		
Linearity FD Cardiac Markers (Siemens Centaur)	K804M-5	5 x 1ml
Analytes: CK-MB, Myoglobin, Troponin I		
Linearity LQ Homocysteine (Siemens Centaur)	K801M-5	5 x 1ml
Analytes: Homocysteine		
Linearity FD BNP (Siemens Stratus)	K892M-5	5 x 1ml
Analytes: NT-Pro BNP		
Linearity FD Cardiac (Siemens Stratus)	K891M-5	5 x 1ml
Analytes: CK-MB, Myoglobin, Troponin I		
Linearity FD D-Dimer (Siemens Stratus)	K893M-5	5 x 1ml
Analytes: D-Dimer		

IMMUNOASSAY

Linearity LQ ASO	K721M-5	5 x 1ml
Analytes: ASO		
Linearity FD Immunoassay	K714M-5	10 x 5ml
Analytes: Set 1: Cortisol, Digoxin, Estradiol, Ferritin, Folate, FSH, hCG, LH, Progesterone, Prolactin, Testosterone, Total PSA, Total T3, Total T4, TSH, Vitamin B12 Set 2: Free T3, Free T4		
Linearity LQ RF/CRP	K720M-5	5 x 1ml
Analytes: CRP, RF		
Linearity FD Tumor Markers	K719M-5	10 x 1ml
Analytes: Set 1: AFP, CA 125, CEA, Total PSA, CA 19-9, CA 27/29 Set 2: CA 15-3, Free PSA		
Linearity LQ Vitamin D	K729M-5	5 x 2ml
Analytes: Vitamin D		
Linearity FD Immunoassay (Abbott Architect i Series)	K831M-5	10 x 5ml
Analytes: Set 1: Ferritin, Folate, Free PSA, FSH, LH, Progesterone, Prolactin, Testosterone, Total PSA, Total T3, Total T4, TSH, Vitamin B12 Set 2: Free T3, Free T4		
Linearity FD Procalcitonin (bioMérieux VIDAS and miniVIDAS)	K841M-5	5 x 1ml
Method Val FD Procalcitonin (bioMérieux VIDAS and miniVIDAS)	K842M-25	25 x 1ml
Analytes: Procalcitonin		
Linearity LQ Vitamin D (Roche Systems)	K885M-5	5 x 2ml
Analytes: Vitamin D		
Linearity FD Anemia (Siemens Centaur)	K809M-5	5 x 3ml
Analytes: Cortisol, Ferritin, Folate, Vitamin B12		
Linearity FD Fertility (Siemens Centaur)	K808M-5	10 x 3ml
Analytes: Set 1: Estradiol, FSH, hCG, LH, Progesterone, Prolactin Set 2: Testosterone		

Linearity FD Thyroid (Siemens Centaur)	K807M-5	10 x 3ml
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Analytes:
Set 1: Total T3, Total T4, TSH
Set 2: Free T3, Free T4

Linearity FD Tumor Markers (Siemens Centaur)	K805M-5	10 x 1ml
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Analytes:
Set 1: AFP, CA 125, CA 19-9, CA 27-29 (BR), CEA, PSA
Set 2: CA 15-3

Cal Ver LQ T-Uptake (Tosoh AIA)	K851M-5	5 x 1ml
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Analytes: T-Uptake

IMMUNOLOGY

Linearity LQ Protein	K702M-5	5 x 2ml
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Analytes: Alpha-1-Antitrypsin, Complement C3, Complement C4, Immunoglobulin A, Immunoglobulin G, Immunoglobulin M, Total Protein, Transferrin

DIABETES

Linearity LQ Beta-Hydroxybutyric Acid	K728M-5	5 x 1ml
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Analytes: Beta-Hydroxybutyric Acid

Linearity FD Glycohemoglobin A1c	K703M-5	5 x 0.5ml
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Analytes: Glycohemoglobin A1c

Linearity LQ Special Diabetes	K730M-5	5 x 2ml
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Analytes: C-Peptide, Fluctosamine, Insulin

URINE CHEMISTRY

Linearity LQ Microalbumin/Microprotein	K722M-5	5 x 2ml
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Analytes: Microalbumin, Microprotein

Linearity FD Urine/Fluids Chemistry	K723M-5	10 x 3ml
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Analytes:
Set 1: Osmolality, Phosphorus, Urea Nitrogen, Uric Acid
Set 2: Amylase, Calcium, Chloride, Creatinine, Glucose, Magnesium, Microalbumin, Microprotein, Potassium, Sodium

THERAPEUTIC DRUG MONITORING

Linearity FD TDM

K707M-5

5 x 5ml

Analytes: Acetaminophen, Carbamazepine, Digoxin, Gentamicin, Lithium, Phenobarbital, Phenytoin, Quinidine, Salicylate, Theophylline, Tobramycin, Valproic Acid, Vancomycin

Linearity FD TDM (Siemens Centaur)

K802M-5

5 x 5ml

Analytes: Carbamazepine, Digoxin, Gentamicin, Phenobarbital, Phenytoin, Theophylline, Valproic Acid, Vancomycin

BLOOD GAS

Cal Ver LQ Blood Gas

K727M-5

15 x 2ml

Analytes: BUN, Chloride, Creatinine, Glucose, Ionised Calcium, Lactate, pCO₂, pH, pO₂, Potassium, Sodium

DAILY QUALITY CONTROL

GENERAL CHEMISTRY

Control LQ Ammonia/Ethanol (2 Levels)

K062M-6

6 x 3ml

Analytes: Ammonia, Ethanol

Control FD Assayed Chemistry, Level 1

K0311-10

10 x 5ml

Analytes: Albumin, Alkaline Phosphatase, ALT (SGPT), Amylase, AST (SGOT), Bilirubin (Total and Direct), BUN, Calcium, Chloride, Cholesterol, CO₂, Creatine Kinase (CK), Creatinine, Gamma-GT, Glucose, HDL Cholesterol, Iron, Lactate, LDH, LDL Cholesterol, Lipase, Magnesium, Osmolality, Phosphorus, Potassium, Sodium, TIBC, Total Protein, Triglycerides, UIBC, Uric Acid

Control FD Assayed Chemistry, Level 2

K0312-10

10 x 5ml

Analytes: Albumin, Alkaline Phosphatase, ALT (SGPT), Amylase, AST (SGOT), Bilirubin (Total and Direct), BUN, Calcium, Chloride, Cholesterol, CO₂, Creatine Kinase (CK), Creatinine, Gamma-GT, Glucose, HDL Cholesterol, Iron, Lactate, LDH, LDL Cholesterol, Lipase, Magnesium, Osmolality, Phosphorus, Potassium, Sodium, TIBC, Total Protein, Triglycerides, UIBC, Uric Acid

CARDIAC

Control FD BNP (2 Levels)

K023M-6

6 x 1ml

Analytes: BNP-32, NT-ProBNP

Control FD Cardiac Markers (3 Levels)

K026M-6

6 x 2ml

Analytes: CK-MB, Myoglobin, Troponin I

Control FD D-Dimer (3 Levels)	K072M-6	6 x 1ml
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Analytes: D-Dimer

Control LQ Homocysteine (2 Levels)	K020M-6	6 x 2ml
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Analytes: Homocysteine

Control LQ hs-CRP (2 Levels)	K021M-6	6 x 2ml
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Analytes: hs-CRP

IMMUNOASSAY

Control FD Immunoassay (3 Levels)	K063M-6	6 x 3ml
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Analytes: Cortisol, Digoxin, Estradiol, Ferritin, Folate, Free T3, Free T4, FSH, hCG, LH, Progesterone, Prolactin, Testosterone, Total PSA, Total T3, Total T4, TSH, Vitamin B12

Control FD Procalcitonin (2 Levels)	K071M-6	6 x 2ml
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Analytes: Procalcitonin

Control LQ RF/CRP (3 Levels)	K073M-6	6 x 2ml
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Analytes: CRP, RF

Control FD Tumor Markers (2 Levels)	K070M-6	6 x 2ml
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Analytes: AFP, CA125, CA15-3, CEA, Free PSA, Total PSA

Control LQ Vitamin D (2 Levels)	K075M-6	6 x 2ml
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Analytes: Vitamin D

IMMUNOLOGY

Control LQ Serum Protein (2 Levels)	K074M-6	6 x 2ml
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Analytes: Albumin, Alpha1-Acid Glycoprotein, Alpha-1-Antitrypsin, Alpha2-Macroglobulin, Antithrombin, Complement C3, Complement C4, Haptoglobin, Immunoglobulin A, Immunoglobulin E, Immunoglobulin G, Immunoglobulin M, Prealbumin, Total Protein, Transferrin

Control LQ Spinal Fluid (2 Levels)	K060M-6	6 x 3ml
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Analytes: Chloride, Glucose, Immunoglobulin A, Immunoglobulin G, Immunoglobulin M, Lactate, LDH, Microalbumin, Microprotein, Sodium

DIABETES

Control FD Glycohemoglobin A1c (2 Levels)

K061M-8

8 x 0.5ml

Analytes: Glycohemoglobin A1c

Control LQ Glycohemoglobin A1c (2 Levels)

K067M-8

8 x 0.5ml

Analytes: Glycohemoglobin A1c

URINE CHEMISTRY

Control FD Urine Chemistry, Level 1

K0661-10

10 x 3ml

Analytes:

Set 1: Phosphorus, Urea Nitrogen, Uric Acid

Set 2: Amylase, Calcium, Chloride, Creatinine, Glucose, Magnesium, Microalbumin, Microprotein, Osmolality, Potassium, Sodium

Control FD Urine Chemistry, Level 2

K0662-10

10 x 3ml

Analytes:

Set 1: Phosphorus, Urea Nitrogen, Uric Acid

Set 2: Amylase, Calcium, Chloride, Creatinine, Glucose, Magnesium, Microalbumin, Microprotein, Osmolality, Potassium, Sodium

Control DROP LQ Urine Dipstick (2 Levels)

K064M-10

10 x 5ml

Control DROP LQ Urine Dipstick (2 Levels)

K065M-4

4 x 25ml

Analytes: Acetest, Bilirubin, Blood, Clinitest, Glucose, hCG, ketones, Leukocytes, Microalbumin, Nitrite, pH, Protein, Specific Gravity, Urobilinogen

THERAPEUTIC DRUG MONITORING

Control FD TDM

K068M-6

6 x 5ml

Analytes: Acetaminophen, Carbamazepine, Digoxin, Gentamicin, Lithium, Phenobarbital, Phenytoin, Quinidine, Theophylline, Tobramycin, Valproic Acid, Vancomycin

KOVA INTERNATIONAL

QUALITY CONTROLS AND CALIBRATORS



KOVA International is a leading manufacturer of quality controls and consumables for the clinical diagnostic laboratory industry primarily catering to the global urinalysis market. With a US manufacturing base, the KOVA® brand is certified to ISO 13485:2003 and is cGMP qualified. The company's primary focus lies in producing urinalysis controls available in both lyophilised and ready-to-use liquid forms.

KOVA-Trol® and KOVA Liqua-Trol® are intended for use in the clinical laboratory as controls for macroscopic and microscopic examination as well as physiochemical and chemical analysis of urine specimens. KOVA-Trol®, a human urine-based control with simulated leukocytes, is used as a daily control for the physical, chemical and microscopic examination of the urinalysis laboratory. KOVA-Trol® is available in three levels: High Abnormal (Level I), Low Abnormal (Level II), and Normal (Level III). Liqua-Trol is available in two levels, Normal and Abnormal, with or without microscopics.

KOVA LIQUA-TROL®

KOVA Liqua-Trol® is a ready-to-use liquid control with urobilinogen, saving sample preparation time and reducing inconsistencies and errors. The KOVA Liqua-Trol® control uses esterase to mimic a patient sample.

KOVA Liqua-Trol® is available in two levels, Normal and Abnormal, with or without microscopics. The 15 mL size configuration for easy-to-use chemical testing and the 120 mL size, for larger microscopic testing.

Values are assigned for visual and instrument reading on all major systems. With proper storage (2-8°C), Liqua-Trol has a proven shelf-life of up to 27 months from date of manufacture, or 30 days at room temperature.

- Reduces time and eliminates reconstitution errors
- Assures consistency from bottle to bottle
- Available with or without microscopics

KOVA® Liqua-Trol® Level I (Abnormal) and Level II (Normal w/ HCG)	87112E	Urine	4 x 25ml
KOVA Liqua-Trol® Level II (Normal) w/ hCG and microscopics	87122E	Urine	2 x 120ml
KOVA Liqua-Trol® Level II (Normal) w/ hCG and microscopics	87123E	Urine	4 x 120ml
KOVA® Liqua-Trol® Level I (Abnormal) w/ Microscopics	87176E	Urine	2 x 120ml
KOVA® Liqua-Trol® Level I (Abnormal) w/ Microscopics	87177E	Urine	4 x 120ml

KOVA-TROL®

The KOVA-Trol® line of lyophilised, human, urine-based controls provides complete quality control for the physical, chemical and microscopic examination of urine specimens. KOVA-Trol® is available in three levels to monitor the entire decision range: High Abnormal (with or without urobilinogen), Low Abnormal and Normal. Values are assigned for visual and instrument reading on all major systems. KOVA-Trol® provides you with the maximum quality control information for urinalysis testing.

- Stable human urine control (freeze-dried) for complete quality control of physical-chemical and microscopic examination of urine specimens
- Available in three levels to monitor the entire decision ranges for reagent strip chemistries
- Red and white cells included for QC of microscopic analysis
- hCG positive in KOVA-Trol® Level III (Normal)
- KOVA-Trol® Level II (Low Abnormal) and KOVA-Trol® III (Normal) have microalbumin and creatinine value assignments

STABILITY

- Store at 2 – 8 °C, protect from light
- Unreconstituted shelf life is 27 months from date of manufacture
- All constituents are stable from day 1 of reconstitution until day 7
- 1 month frozen stability for strip testing and hCG

KOVA-Trol® I: High Abnormal w/o Urobilinogen	87329E	Urine	4 X 15ml
KOVA-Trol® I: High Abnormal w/o Urobilinogen	87325E	Urine	4 x 60ml
KOVA-Trol® I: High Abnormal w/o Urobilinogen	87426E	Urine	8 x 60ml
KOVA-Trol® I: High Abnormal w/ Urobilinogen	87332E	Urine	4 X 60ml
KOVA-Trol® I: High Abnormal w/ Urobilinogen	87533E	Urine	8 x 60ml
KOVA-Trol® I: High Abnormal w/ Urobilinogen	87334E	Urine	4 x 15ml
KOVA-Trol® II: Low Abnormal	87130E	Urine	4 x 15ml
KOVA-Trol® II: Low Abnormal	87428E	Urine	8 x 60ml
KOVA-Trol® III: Normal w/ hCG	87331E	Urine	4 X 15ml
KOVA-Trol® III: Normal w/ hCG	87327E	Urine	4 x 60ml
KOVA-Trol® III: Normal w/ hCG	87528E	Urine	8 x 60ml

QUANTIMETRIX

QUALITY CONTROLS



Quantimetrix designs, develops, and manufactures laboratory-quality products at their headquarters in Redondo Beach, California. Leaders in the field of liquid-stable quality control products, Quantimetrix pioneered these innovations more than 40 years ago. Their portfolio improves the efficiency and reliability of laboratory testing and patient care.

DipperPOCT Control – 2 Level Set	1400-01	62 x 1.5ml
DipperPOCT Control – 2 Level Set	1400-02	20 x 1.5ml
Analytes: Bilirubin, Blood, Creatinine, Glucose, hCG, Ketones, Leukocytes, Microalbumin, Nitrite, pH, Protein, Specific Gravity, Urobilinogen		



Dipper POCT® Urinalysis Dipstick Control is a revolutionary single-use liquid control with long stability. Made with a simulated human urine matrix, Dipper POCT's stability exceeds that of all other urinalysis controls, currently on the market, formulated with native ketones. Perfect for use in every testing environment including central labs, reference labs, nursing stations and doctor's offices.

- 3 years refrigerated shelf life from date of manufacture
- 3 months room temperature stability
- Full dipstick immersion simulates patient testing
- Minimised risk of contamination
- Designed for use with most urinalysis reagent strips

Dropper Plus POC Urine Dipstick Control Set	1440-04	10 x 5ml
Dropper Plus POC Urine Dipstick Control Set	1440-06	2 x 5ml
Analytes: Bilirubin, Blood, Creatinine, Glucose, hCG, Ketones, Leukocytes, Microalbumin, Nitrite, pH, Protein, Specific Gravity, Urobilinogen		



Dropper® Plus POC Urinalysis Dipstick Control eliminates storage problems and gives you maximum portability by providing one month of room temperature stability – perfect for sites without refrigeration. Easy-to-use dropper bottles make dispensing simple. Dropper Plus POC Urinalysis Dipstick Control is designed for use with most urinalysis reagent strips. It can also be used for confirmatory tests and refractometry.

- Liquid, made with human urine
- 1-month open vial stability at room temperature for maximum portability
- 18-month shelf life & open vial stability from date of manufacture when stored at 2°C–8°C





GET IN TOUCH

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