QUALITY CONTROLS CATALOGUE

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2021 2021	
United Kingdom Fortress Diagnostics LTD	
Fortress Diagnostics	



How To Order

What information do we need?

Please quote your:

- Address
- Telephone Number
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- **Product Description**
- Catalogue Number
- Quantity Required
- 10 Any Specific Delivery Instructions

Where should you send your quote?

Please send your quote to us using one of the following options:

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Through Our Website fortressdiagnostics.com/contact-us

Over the Phone +44 (0)28 9448 7676

By Email



Advancing Global Diagnostics

supplying an extensive portfolio of products to over 100 countries worldwide.

We have been providing our distributors and end users with high quality products for over 20 years. The objective of Fortress Diagnostics is to provide the highest level of diagnostic solutions, globally, through continued investment in research and development.

Why choose Fortress Diagnostics?



Our proven product performance, matched with our selection of reputable distribution partners, has been instrumental in our continued success and growth. Built on a culture of quality and performance, we are committed to providing our customers with a reliable service and results they can trust.



Our ISO 13485:2016 certification for the design, development, manufacturing and distribution of in vitro diagnostics (IVD) medical reagents and instrumentation endorses our product range and ensures that the highest quality standards are constantly maintained.



Through our global distribution network, we provide highly accurate medical testing solutions to immunology, haematology and serological laboratories in hospitals, medical centres, clinics, blood banks and research institutions in over 100 international markets.



Hosting an in-house Research and Development team, we are committed to creating the next innovative IVD solutions to exceed our customer's requirments and advance the healthcare industry. We reinvest over 30% of our turnover in R&D every year.

Fortress Quality Controls & Calibrators

Quality controls and calibrators play a vital role in clinical diagnostics by ensuring the accuracy and reliability of test results. Quality controls are designed to monitor the performance of diagnostic assays and reagents, helping to detect any variations or issues that may arise during testing. This ongoing assessment ensures that laboratories maintain high standards and can identify problems before they affect patient care. Calibrators, on the other hand. are used to establish the measurement accuracy of diagnostic instruments by providing a known reference point against which test results can be compared.

Together, these elements contribute to standardisation and consistency in laboratory testing, fostering confidence in the results provided to healthcare professionals. Ultimately, robust quality control and calibration processes are essential for delivering precise diagnoses, guiding effective treatment decisions, and ensuring patient safety in clinical practice.

Fortress Quality Controls (Seraqual) are supplied with instrument and method specific assigned values. We can also provide guality control solutions, tailor-made to customer specification and requirements, enabling laboratories to be more efficient and reduce costs. We also work with many universities, research institutions and laboratories to develop quality control and calibration solutions for unique and novel biomarkers.

The Seragual Portfolio

Fortress Diagnostics Seragual Controls and Calibrators are available for:

Antioxidants **Aqueous Standards Blood Gas & Electrolytes** Cardiac **Clinical Chemistry** Coagulation/Haemostasis Diabetes

Drugs of Abuse Haematology Immunoassay/ Infectious Immunology/Proteins Lipids Maternal Screening Neonatal Screening **Paediatric Screening**

Fortress Diagnostics | Quality Controls Catalogue

Fortress Diagnostics is a well-established, multi award-winning IVD manufacturer based in the United Kingdom.

Speciality Controls Thalassaemia **Therapeutic Drugs** Torch Paediatric Urine Xanthochromia

Why Choose Fortress Quality Controls & Calibrators?



Customer Support

Fortress Diagnostics employ a highly-skilled team of technical support staff to assist customers with all their needs and technical questions. We are always on-hand to help.



As well as offering an extensive range of controls, Fortress Diagnostics has the ability to manufacture custom quality control products to meet any customer requirements providing a fully flexible solution.



Cost Saving

Our Quality Controls Range have a wide-ranging analyte content, enabling laboratories to consolidate their quality control purchases and therefore benefit from cost savings.





costs.

Long Shelf Life

Flexible



Accurate

Manufactured by Fortress Diagnostics in our ISO 13485:2016 accredited labs, Seraqual products are designed to provide highly accurate target, values mimicking human samples, with minimal matrix effects being observed.





Reliable

Our fully in-house logistics team enables us to ship orders globally and our customer support team are available to assist with any labs' special requirements or custom orders to ensure customer satisfaction is always exceeded.







Consolidation

Traceability

directives.

Fortress Quality Controls offer a wide parameter range allowing more analyte coverage with a single control.



Third Party Controls Suitable For Global Market

We manufacture a wide range of third party controls which are valueassigned, without bias, towards any methods or instrumentation, enabling labs to meet their ISO accreditations through accurate and robust training.



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Value Assignment The Fortress value assignment involves six European Reference

Laboratories, (chosen on the basis of the analytes performed in these

Compatibility

Fortress Controls are made from human derived material, with minimal matrix effects being observed.



High Quality

Seragual products are manufactured without the use of preservatives or stabilisers, utlising human-based material, which have undergone stringent quality checks for HIV, HBsAg, HCV, HTLV, and Syphilis for FDA approved methods. Products of animal origin are not added in the Fortress controls, therefore eliminating the lack of specificity in antibody-based tests.





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Fortress Diagnostics offer quality controls in two different formats - lyopholised (stable for up to 4 years) and liquid (stable with a shelf life of up to 18 months) - enabling labs to utilize lot supply and save

Fortress Controls and Calibrators are CE marked and fully traceable to the reference material according to ISO 17511 and ISO 18153

laboratories), each of which performs five independant runs.











Find Out More





Website fortressdiagnostics.com

The Fortress Diagnostics website details our full product range, allowing users to add products of interest to a "Quote Request" which is sent directly to our sales team. We regularly post blogs, event information, resources and company updates.

Quality Control Inserts Portal

insert.fortressdiagnostics.com

Fortress Diagnostics are excited to launch the new Fortress QC Inserts Portal. The portal allows customers to easily search our database of quality control inserts by product name, catalogue number or batch number in our easy-to-use filter tool. Inserts can be viewed online or downloaded in PDF format.





Brochures & Catalogues

fortressdiagnostics.com/resources

All Fortress Diagnostics brochures and catalogues can be viewed and downloaded on the Fortress Resource Hub in PDF format.

Videos

youtube.com/@fortressdiagnostics

Subscribe to our YouTube channel to explore information videos, tutorial guides for Seraqual 365, interviews for World Awareness Days and more.





Social Media

Explore and follow our social media pages across **Instagram**, **LinkedIn**, **Facebook** and **Twitter**. We regularly post updates on product launches, events Fortress are exhibiting at, news from the world of diagnostics and company developments.

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Antioxidant Controls

Antioxidant quality controls in diagnostics are crucial for ensuring the accuracy and reliability of tests that measure oxidative stress and antioxidant levels in biological samples. These quality controls are used to validate the performance of diagnostic assays, ensuring that the test results are consistent and accurate. Positive controls, which contain known quantities of antioxidants, help confirm that the assay is working correctly, while negative controls, without antioxidants, ensure there is no false detection of antioxidant activity.

By using antioxidant controls, laboratories can detect any potential issues with the assay, such as reagent degradation or equipment malfunction. In addition to positive and negative controls, diagnostic assays may also use blank and solvent controls to account for any potential interference from other substances or experimental factors. Consistent use of antioxidant quality controls helps to standardize the measurement of oxidative stress biomarkers, ensuring that results are comparable across different laboratories and studies. These controls are critical in clinical diagnostics, where accurate measurements of antioxidant levels can be used for disease risk assessment, monitoring therapeutic responses, and guiding treatment decisions, particularly in conditions like cardiovascular diseases, diabetes, and neurodegenerative disorders.

The Fortress Seraqual Antioxidant Quality Controls are lyophilised for enhanced stability, offer excellent accuracy and precision and cover a range of key antioxidants in clinical practice. The controls are supplied with instrument and method specific assigned values.

			8	6
Lyophilised	Liquid Frozen	Liquid Stable	100% Human Serum	Assayed Target Values Provided
ilutathione Peroxidase C	ontrol			
 This control is presente Reconstituted stability 100% Human serum ba 		e to expiry date at 2-8 °C.		
Description		Size	Cat No.	Туре
Glutathione Peroxidase Control		5 x 1 ml	BXC0556/	A Lyo
	librator & Control eductase Controls are intended Is are supplied in a Lypholised f			
	7 day at 2-8°C or 3 days at 25 °C		-8 L.	
Description		Size	Cat No.	Туре
Glutathione Reductase Calibrat	or	5 x 1 ml	BXC0999	A Lyo
Glutathione Reductase Control		5 x 1 ml	BXC0460	A Lyo
Superoxide Dismutase Co	ntrol			
 The Control is presente Reconstituted stability 100% Human serum ba 		o expiry date at 2-8 °C.		
Description		Size	Cat No.	Туре
Superoxide Dismutase Control		5 x 1 ml	BXC0433	A Lyo
Total Antioxidant Status (Calibrator & Control			0
 100% Human serum ba Reconstituted stability 				
Description		Size	Cat No.	Туре
Total Antioxidant Status (TAS) (Calibrator	1 x 1 ml	BXC0555	A Lyo
Total Antioxidant Status (TAS) (Control	5 x 5 ml	BXC0554	A Lyo



Aqueous Standards

Aqueous standards in diagnostics are reference solutions used to calibrate and validate diagnostic assays, ensuring accurate measurement of analytes in biological fluids such as blood, urine, and saliva. These standards consist of known concentrations of target molecules dissolved in water, closely mimicking the physiological environment.

By using aqueous standards, laboratories can calibrate instruments and establish a baseline for comparison with patient samples. This allows for precise detection and quantification of biomarkers, essential for diagnosing diseases and monitoring health conditions. Aqueous standards are especially important in ensuring consistency across different diagnostic platforms and laboratories. They help identify any variation or drift in instrument performance over time and confirm that the reagents used in tests are functioning properly. In clinical diagnostics, the use of well-prepared aqueous standards ensures that test results are reliable and reproducible, which is critical for making informed medical decisions, particularly in areas such as metabolic monitoring, hormone assays, and infection diagnostics.

The Fortress Aqueous Standards are available in a liquid stable format for convenience and ease-of-use, with a long shelf life, and are traceable to reference standard material.

Liquid

Stable















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Aqueous Standards

Liquid stable (ready-to-use). •

Long shelf life.

Traceable to reference standard material.

Description	Method	Size	Cat No.	Туре
Albumin	45 g/L (4.5g/dL)	10 x 5 ml	BXCSTD01	LS
Calcium	Referenced against NIST 909b (2.5mmol/L 10mg/dL)	10 x 5 ml	BXCSTD02	LS
Chloride	Referenced against NIST 909b (75mmol/L)	10 x 5 ml	BXCSTD03	LS
Chloride	Referenced against NIST 909b (100mmol/L)	10 x 5 ml	BXCSTD04	LS
Cholestrol	Referenced against NIST 909b(5.17mmol/L 200mg/dL)	10 x 5 ml	BXCSTD05	LS
Creatinine	Referenced against NIST 909b (177 µmol/L 2 mg/dL)	10 x 5 ml	BXCSTD06	LS
Creatine/Glucose/Urea Standard	Referenced against NIST 965a / 909b (5.55 mmol/L 100mg/dL/ 177 µmol/L / 8.33mmol/L)	10 x 5 ml	BXCSTD24	LS
Glucose	Referenced against NIST 965a (5.55 mmol/L 100mg/dL)	10 x 5 ml	BXCSTD07	LS
Glucose-Fructose Standard		5 x 1 ml	BXCSTD26	LS
Haemoglobin Standard	For Drabkin's Method (18g/dL)	5 x 10 ml	BXCSTD08	LS
Haemoglobin Standard Set	For Drabkin's Method (8, 10, 12, 15 & 18g/dL)	5 x 1 x 10 ml	BXCSTD09	LS
Iron	35.8 µmol/L (0.2 mg/dL)	10 x 5 ml	BXCSTD10	LS
Iron	107.4 µmol/L (0.6 mg/dL)	10 x 5 ml	BXCSTD11	LS
Lactate	4.44mmol/L (40mg/dL)	10 x 5 ml	BXCSTD12	LS
Magnesium	Referenced against NIST 909b (1.0 mmol/l2.43mg/dL)	10 x 5 ml	BXCSTD13	LS
Phosphorus	1.61 mmol/L (5 mg/dL)	10 x 5 ml	BXCSTD14	LS
Potassium	Referenced against NIST 909b (0.5 mmol/L)	10 x 5 ml	BXCSTD15	LS
Potassium	Referenced against NIST 909b (5 mmol/L)	10 x 5 ml	BXCSTD16	LS
Pyruvate	2 mmol/L	10 x 5 ml	BXCSTD17	LS
Sodium	Referenced against NIST 909b (150.0 mmol/L 150 mEq/L)	10 x 5 ml	BXCSTD18	LS
Total Protein	Referenced against NIST 927d (60g/L (6.0 g/dL)	10 x 5 ml	BXCSTD19	LS
Triglycerides	Referenced against NIST 909b (2.28 mmol/L 200mg/dL)	10 x 5 ml	BXCSTD20	LS
Urea	Referenced against NIST 909b (8.33 mmol/L 50mg/dL)	10 x 5 ml	BXCSTD21	LS
Uric Acid	Referenced against NIST 909b (595 µmol/L 10mg/dL)	10 x 5 ml	BXCSTD22	LS
Urinary Copper Standard	Referenced against NIST-SE 3114 (1.6mol/L)	10 x 5 ml	BXCSTD25	LS
Urinary/CSF Protein Standard	Referenced against NIST 927d (1g/L)	10 x 5ml	BXCSTD23	LS



Quality Controls Catalogue | Fortress Diagnostics

SECTION 3 BLOOD GAS & ELECTROLYTES

Blood Gas & Electrolyte Controls

Blood gas and electrolyte controls in diagnostics are essential for ensuring the accuracy and reliability of tests that measure critical parameters like pH, oxygen (PaO2), carbon dioxide (PaCO2), and key electrolytes such as sodium, potassium, chloride, and bicarbonate. These controls, typically pre-prepared solutions with known concentrations, are used to calibrate and validate diagnostic equipment, such as blood gas analyzers.

By regularly running these controls, laboratories can detect any inaccuracies in the testing process, ensuring that instruments and reagents are functioning correctly and that test results are precise. In clinical settings, the proper use of blood gas and electrolyte controls helps prevent diagnostic errors, which are crucial in high-stakes situations such as critical care and emergency medicine. Any variation in these tests can significantly impact patient management, from adjusting ventilation settings to correcting electrolyte imbalances.

The Fortress Blood Gas & Electrolytes Quality Controls contain values for ten parameters, with values assigned by European Reference Laboratories, covering pH, pCO2, pO2 and electrolytes. Available in a liquid stable format for convenience and ease-of-use. The controls are supplied with instrument and method specific assigned values.



Blood Gas & Electrolyte Controls

- Liquid stable controls with a shelf life of 1 year when stored, unopened at 2-8 °C.
- Values are assigned by European reference laboratories.
- Store at 2-8 °C. Do not freeze or expose to temperatures greater than 30 °C.
 When opened, analyse immediately for Blood Gas parameters and use within 1 hour for electrolytes.

Analytes						
Calcium Chloride Glucose	Lactate Potassium Sodium	рСО2 рН рО2		Total CO2		
Description		Size	Cat No.	Туре		
Blood Gas & Electrolyte Control (Level 1)		1 x 2 ml	BXC0108A	LS		
Blood Gas & Electrolyte Control (Level 1)		10 x 2 ml	BXC0108D	LS		
Blood Gas & Electrolyte Control (Level 2)		1 x 2 ml	BXC0108B	LS		
Blood Gas & Electrolyte Control (Level 2)		10 x 2 ml	BXC0108E	LS		
Blood Gas & Electrolyte Control (Level 3)		1 x 2 ml	BXC0108C	LS		
Blood Gas & Electrolyte Control (Level 3)		10 x 2 ml	BXC0108F	LS		
Blood Gas & Electrolyte Control (Tri-Level)		30 x 2 ml	BXC0108G	LS		

Electrolyte Controls

• Liquid stable format with a shelf life of 2 years when stored at 2-8 °C.

Values are assigned by European reference Laboratories.

Store at 2-8 °C. Do not freeze or expose to temperatures greater than 30 °C. When opened the analytes are stable for a period of 15 days when stored at 2-8 °C.

Analytes

Chloride (ISE & Colorimetry)	Potassium (ISE & Co	olorimetry)	рН		Sodium (ISE & Colorimet
Description	Details	Size		Cat No.	Туре
Electrolytes Calibrator		5 x 5 ml		BXC0140A	LS
Electrolytes Control (Level 1)	Low (Na, K, Cl)	5 x 5 ml		BXC0143A	LS
Electrolytes Control (Level 2)	Normal (Na, K, Cl)	5 x 5 ml		BXC0144A	LS
Electrolytes Control (Level 3)	High (Na, K, Cl)	5 x 5 ml		BXC0145A	LS



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Blood Grouping Controls

Blood grouping controls in diagnostics are vital for ensuring the accuracy of blood type determination, which is essential for safe blood transfusions, organ transplants, and pregnancy management. These controls are standardised samples with known blood types (A, B, AB, O, and Rh factor) used to verify the performance of blood typing assays and equipment. By including both positive and negative controls in the testing process, laboratories can ensure that reagents are reacting correctly with antigens and antibodies, reducing the risk of errors in identifying a patient's blood group.

Using blood grouping controls regularly helps maintain the reliability of diagnostic tests, preventing potentially lifethreatening transfusion reactions caused by incorrect blood typing. It also ensures consistency across different testing environments, whether in routine clinical laboratories or emergency settings. These controls allow for early detection of any technical issues with reagents, equipment, or the procedure itself, ensuring accurate and dependable blood group determination, which is critical for patient safety.

The Fortress Seraqual ABO-Rhesus Blood Grouping Quality Control Set is available in a liquid stable format for convenience and ease-of-use and covers all major blood types. The controls are supplied with instrument and method specific assigned values.



AB0-Rhesus Blood Grouping Control Set

- Ready-to-Use
- Covers all major blood types.
- Total Blood Kit.

Analytes						
A AB	B O	Rhesus Rh	Kell	Phenotype		
Description	Details	Size	Cat No.	Туре		
AB0-Rhesus Blood Grouping Control Set	For use Rh-reagents and Anti-K Monoclonal	4 x 4 ml	BXC0070A	LS		
Monoclonal Anti-D Negative Control		1 x 10 ml	BDNC0010	LS		
Precise Weak Anti-D Control	Sensitivity Control for AHG	1 x 5 ml	BGWD0005	LS		



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Cardiac Controls

Cardiac controls are essential in diagnostics for ensuring the accuracy and reliability of tests that measure key cardiac biomarkers, such as troponin and B-type natriuretic peptide (BNP), which are critical for diagnosing heart attacks, heart failure, and other cardiovascular conditions. These controls, which contain known concentrations of these biomarkers, help validate the performance of diagnostic assays, ensuring that testing equipment and reagents are working properly. By using these controls, laboratories can identify potential issues early, minimising the risk of inaccurate test results that could lead to misdiagnosis or inappropriate treatment.

The role of cardiac controls is particularly crucial in emergency and critical care settings, where timely and precise biomarker measurements are vital for guiding immediate medical interventions. Reliable cardiac diagnostics help physicians make informed decisions about treatment strategies, such as initiating therapies for acute coronary syndromes or adjusting treatments for chronic heart conditions. Regular use of cardiac controls ensures consistency across tests, supporting better patient outcomes by enabling accurate diagnosis and monitoring of cardiovascular health.

The Fortress Seraqual Cardiac Quality Controls have been designed to cover several cardiac analytes. Available in a lyophilised format for enhanced stability and manufactured from 100% human based serum to negate matrix effect. The controls are supplied with instrument and method specific assigned values.



Cardiac Controls

- Values are assigned by European reference laboratories.
- Lyophilised format stable up to expiry at 2-8 °C.
- Reconstituted stability of 5 days at 2-8 °C or 4 weeks at -20 °C.
- 100% Human serum base.

Analytes					
Homocysteine CK Total	CK MB Activity CK MB Mass	Digoxin Troponin I		Troponin T Myoglobin	
Description	Details		Size	Cat No.	Туре
Cardiac Control (Level 1)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 2 ml	BXC0455A	Lyo.
Cardiac Control (Level 1)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 1 ml	BXC0455B	Lyo.
Cardiac Control (Level 2)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 2 ml	BXC0456A	Lyo.
Cardiac Control (Level 2)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 1 ml	BXC0456B	Lyo.
Cardiac Control (Level 3)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 2 ml	BXC0464A	Lyo.
Cardiac Control (Level 3)	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	5 x 1 ml	BXC0464B	Lyo.
Cardiac Control Set	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	3 x 1 x 2ml	BXC0450A	Lyo.
Cardiac Control Set	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	3 x 1 x 1ml	BXC0450B	Lyo.
Cardiac Tri Level Control	CK Total, CKMB Activity, CKMB Mass, Tni,TnT, Dig Myoglobin	goxin, Homocysteine,	3 x 1 x 0.5ml	BXC0881A	Lyo.

CK/CK MB Calibrators & Controls

- Lyophilised format stable up to expiry at 2-8 °C.
- Values are assigned by European Reference Laboratories.
- 100% Human serum base.
- Reconstituted stability 5 days at 2-8 °C, 8 hours at +25 °C and 4 weeks at -20 °C.

Analytes						
	CK Total	CK MB Activit	у			
Description		Size	Cat No.	Туре		
CK/CK MB Calibrator		1 x 2 ml	BXC0454A	Lyo.		
CK/CK MB Calibrator		5 x 2 ml	BXC0454B	Lyo.		
CK/CK MB Control (Level 1)		1 x 2 ml	BXC0453A	Lyo.		
CK/CK MB Control (Level 1)		5 x 2 ml	BXC0453B	Lyo.		
CK/CK MB Control (Level 1)		10 x 2 ml	BXC0453C	Lyo.		
CK/CK MB Control (Level 2)		1 x 2 ml	BXC0459A	Lyo.		
CK/CK MB Control (Level 2)		5 x 2 ml	BXC0459B	Lyo.		
CK/CK MB Control (Level 2)		10 x 2 ml	BXC0459C	Lyo.		

D-DIMER Calibrators & Controls

- Lyophilised format stable up to expiry at 2-8°C.
- Prepared from human serum
- Open vial stability of 30 days, at 2-8°C.

Description

D-Dimer Calibrator Set

D-Dimer Control Set (Level 1 & Level 2)

Homocysteine Calibrators & Controls

- Fortress Homocysteine controls are intended for monitoring accuracy and precision.
- The Calibrators & controls are provided in a Lyophilised format stable up to expiry at 2-8 °C.
- 100% Human serum based.

Description

Homocysteine Calibrator (2 Levels)

Homocysteine Control (2 Levels)

Myoglobin Calibrators & Controls

- Lyophilised to achieve long shelf life.
- Reconstituted stability of 30 days at 2-8°C and 12 months at -20°C.

Description

Myoglobin Calibrator Series

Myoglobin Control

NTProBNP Calibrators & Controls

•	Lyophilised format stable to expiry at 2-8 °C.
	Prepared from human serum

- Prepared from human serum. Reconstituted stability of 8 hours at 2-8°C.

Description	Details	Size	Cat No.	Туре
NTProBNP Calibrator	Reconstitued Stability 8 hours at 2-8°C	1 x 1 ml	BXC0335A	Lyo.
NTProBNP Control Series (6 Levels)	Reconstitued Stability 8 hours at 2-8°C	6 x 1 x 1 ml	BXC0336A	Lyo.
BNP Calibrator Level 1	human Serum Base, Reconstituted Stability 8 hours at 2-8°C	1 x 0.5ml	BXC0455S	Lyo.
BNP Calibrator Level 2	human Serum Base, Reconstituted Stability 8 hours at 2-8°C	1 x 0.5ml	BXC0456S	Lyo.

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CK	MB	Activity	



Si	ze	Cat No.	Туре
5 :	x 1 x1ml	BXC0788A	Lyo.
2 :	x 1 x1ml	BXC0789A	Lyo.

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Size	Cat No.	Туре
2 x 1 x 1ml	BXC0691B	Lyo.
2 x 1 x 1 ml	BXC0692A	Lyo.

Size	Cat No.	Туре
5 x 1 x 0.5ml	BXC0486A	Lyo.
1 x 0.5ml	BXC0487A	Lyo.
		80

SECTION 5 | Cardiac

Troponin-T (High Sensitivity) Controls

- Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Cut-off value of 13.5ng/L for automated systems. Suitable for use in point-of-care testing. Open vial stability of 1 week, at 2-8°C. .
- .

Troponin-T High Sensitivity Control	For use on all major systems	3 x 1 ml	BXC0457A	15
Description	Details	Size	Cat No.	Туре

Troponin-I Controls & Calibrators

Lyophilised for long shelf life, stable to expiry at 2-8 °C.
Reconstituted stability of 5 days at 2-8°C.

Description	Details	Size	Cat No.	Туре
Troponin-I Control Set	For use on all major systems	2 x 1 x 1 ml	BXC0470A	Lyo.
Troponin-I Calibrator Set	For use on all major systems	6 x 1 x 1 ml	BXC0473A	Lyo.





Clinical Chemistry Controls

Clinical chemistry controls are vital for ensuring the accuracy, reliability, and validity of laboratory tests that analyze bodily fluids, such as blood and urine. These controls, which include both positive and negative samples with known concentrations of analytes, help validate the performance of diagnostic assays and ensure that testing equipment and reagents are functioning correctly. By routinely using these controls, laboratories can detect any inconsistencies or errors in the testing process, thereby minimising the risk of false positives or negatives that could lead to misdiagnosis or inappropriate treatment.

The role of clinical chemistry controls is particularly important in maintaining quality assurance in laboratory settings. They help ensure that the results are consistent and reproducible across different tests and patient samples, supporting accurate clinical decision-making. In critical situations, such as emergency medicine or monitoring chronic conditions, reliable laboratory data is essential for guiding treatment plans and assessing patient outcomes. Overall, clinical chemistry controls are integral to enhancing the overall quality of healthcare by providing dependable diagnostic information that informs patient management and care strategies.

The Fortress clinical chemistry range of controls are suitable for a range of clinical chemistry analysers, offering flexibility for your laboratory. The controls are supplied with instrument and method specific assigned values.



Featured Product

Human Assaved Controls



Multi-Analyte Human Assayed Controls are essential tools in clinical laboratories for ensuring the accuracy and reliability of diagnostic tests. These controls contain a blend of multiple analytes—such as hormones, proteins, and other biomarkers—mimicking human samples.

They are used to validate and monitor the performance of assays, which are tests designed to measure the presence or concentration of specific substances in a sample. By incorporating a range of analytes, these controls help laboratories simultaneously assess the precision and accuracy of various diagnostic assays under a single test run. This comprehensive approach not only enhances the efficiency of quality control procedures but also ensures consistency in test results, ultimately contributing to better patient care by maintaining high standards in laboratory diagnostics.

Why Choose Fortress Human Assayed Controls?



Human serum based: Same matrix as samples.



Lyophilised format: Fortress Human Assayed Controls are stable to expiry at 2-8ºC.



Custom levels tailored to specific requirments available: Adaptable to perfectly suit unique experimental conditions, analytical needs, and regulatory standards.



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0.2µ filtered: Helps remove particulate contaminants and potential microbial contaminants, leading to more accurate and reproducible outcomes in various assays.

Method based values with reference traceability provided.

Reconstituted stability of 7 days at 2-8°C, or 4 weeks at -20°C.

	Substrates
Entymor	Albumin
Enzymes Alkaline Phosphatase	APO A1
•	APO B
ALT (GPT) Amvlase	Bile Acids
5	Bilirubin Direct
AST (GOT) Cholinesterase	Bilirubin Total
	Calcium
CK (Total) Gamma GT	Cholesterol
GLDH	Соррег
HBDH	Creatinine
LDH	D3- Hyrdroxybutyrate
	Glucose
Lipase Pancreatic Amylase	HDL Cholesterol
Total Acid Phosphatase	IGA
Prostatic Acid-Phosphatase	IGG
Total Amylase	IGM
TOLAL AITIYLASE	Iron
	Iron/TIBC

		-		
Enzymes Alkaline Phosphatase ALT (GPT) Amylase AST (GOT) Cholinesterase CK (Total) Gamma GT GLDH HBDH LDH Lipase Pancreatic Amylase Total Acid Phosphatase Prostatic Acid-Phosphatase Total Amylase	Substrates Albumin APO A1 APO B Bile Acids Bilirubin Direct Bilirubin Total Calcium Cholesterol Copper Creatinine D3- Hyrdroxybutyrate Glucose HDL Cholesterol IGA IGG IGM Iron	Inorganic Phosphorous LDL Cholesterol NEFA Osmolality Total Protein Transferrin Triglycerides UIBC Urea Uric Acid Zinc Electrolytes Bicarbonate Chloride Lithium Magnesium Potassium Sodium	۱ F T	munoassays Cortisol Folate FT3 FT4 Total T3 Total T4 Total PSA TSH /itamin B12 Drugs Digoxin Gentamicin 'aracetamol Salicylate 'heophyline Tobramycin
Description		Size	Cat No.	Туре
Human Assayed Control (Level 1)		10 x 5ml	BXC0312A	Lyo.
Human Assayed Control (Level 2)		10 x 5ml	BXC0312B	Lyo.
Human Assayed Control (Level 1)		5 x 5ml	BXC0312C	Lyo.
Human Assayed Control (Level 2)		5 x 5ml	BXC0312D	Lyo.
Human Assayed Control (Level 1)		1 x 5ml	BXC0312E	Lyo.

GenChem UltraOC TR

Alkaline Phosphatase Albumin Alkaline Phosphatase Alpha-1-Acid Glycoprotein Alpha-1-Antitrypsin Alpha-Fetoprotein ALT (GPT) Amylase (Pancreatic) Amylase (Total) Apo-A1 ADO-B AST (GOT) Beta-2-Macroglobulin Bicarbonate Bile Acids Bilirubin, Direct Bilirubin, Total C-Reactive Protein Ceruloplasmin Calcium

Carcinoembryonic Antigen (CEA) Chloride Cholesterol Cholinesterase CK-NAC Complement C3 Complement C4 Copper Cortisol Creatinine D3-Hydroxybutyrate DHEA-S Ethanol (alcohol) Ferritin Folate Free T3 Free T4 FSH Gamma-GT GLDH Glucose

Description

Liquid Assayed Chemistry Control (Level-1)

Liquid Assayed Chemistry Control (Level-2)

Liquid Assayed Chemistry Control (Level-3)

Analytes

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)	a-HBDH Iron Haptoglobin HDL Cholesterol (Direct) IgA IgE IgG LgM Lactate LDH LACTATE DEHYDROGENASE LDL Cholesterol Direct Leutinising Hormone (LH) Lipase Lipoprotein (a) Lithium Magnesium Myoglobin Osmolality Phosphate Inorganic Potassium Prealbumin Progesterone	Prolactin Protein, Total PSA Total Sodium T Uptake Testosterone TSH TIBC Total Beta hCG Total T3 Total T4 Transferrin Triglycerides Troponin T UIBC Urea Uric Acid Vitamin B12 Zinc

Size	Cat No.	Туре
6 x 5 ml	BXC0365A	LS (Frozen).
6 x 5 ml	BXC0365B	LS (Frozen)
6 x 5 ml	BXC0365C	LS (Frozen)

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Calibration Serum

• Approximately 40 parameters are value assigned by reference laboratories in Europe.

- Method based values with reference traceability provided.
- Lyophilised product stable to expiry at 2-8 °C.
 Reconstituted stability of 7 days at 2-8 °C, or 4 weeks at -20 °C.

	7.	latyces		
Enzymes ALP ALT (GPT) Total Amylase AST (GOT) Cholinesterase CK NAC Gamma GT GLDH HBDH LDH LDH Lipase	Pancreatic Amylase Prostatic Phosphatase Substrates Albumin Bicarbonate Bile Acids Bilirubin Direct Bilirubin Total Calcium Cholesterol Copper	Creatinine D3- Hydroxybutyrate Glucose HDL Cholesterol Iron Iron/TIBC Inorganic Phosphorous Lactate LDL Cholesterol Total Protein Triglycerides Urea	Uric Acid Zinc Electrolytes Chloride Lithium Magnesium Potassium Osmolality Sodium	
Description		Size	Cat No.	Туре
Calibration Serum		1 x 3 ml	BXC0321K	Lyo.
Calibration Serum		5 x 3 ml	BXC0321L	Lyo.
Calibration Serum		10 x 3 ml	BXC0321M	Lyo.
Calibration Serum Normal		3 x 5 ml	BXC0321A	Lyo.
Calibration Serum Normal		5 x 5 ml	BXC0321B	Lyo.
Calibration Serum Normal		10 x 5 ml	BXC0321C	Lyo.
Calibration Serum Elevated		3 x 5 ml	BXC0321D	Lyo.
Calibration Serum Elevated		5 x 5 ml	BXC0321E	Lyo.
Calibration Serum Elevated		10 x 5 ml	BXC0321F	Lyo.

Analytes

ADA Calibrators & Controls

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• Liquid Stable version (Ready-to-use) stable to expiry at 2-8 °C & Open vial stability of 30 days, at 2-8°C.

Lyophilised version reconstituted stability of 30 days at 2-8°C.
 Open vial stability of 30 days, at 2-8°C.

Description	Size	Cat No.	Туре
ADA Calibrator	1 x 1 ml	BXC0209A	LS
ADA Control (Level 1)	1 x 1 ml	BXC0210A	LS
ADA Control (Level 2)	1 x 1 ml	BXC0216A	LS
ADA Calibrator	1 x 1 ml	BXC0217A	Lyo.
ADA Control (Level 1)	1 x 1 ml	BXC0218A	Lyo.
ADA Control (Level 2)	1 x 1 ml	BXC0219A	Lyo.

Alcohol, Ammonia, Carbonate Calibrators & Controls

	Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.
•	Open vial stability of 24 hours at 2-8°C.

Description	Size	Cat No.	Туре
Alcohol, Ammonia, Carbonate Calibrator	1 x 2 ml	BXC0492A	LS
Alcohol, Ammonia, Carbonate Control (Level 1)	1 x 2 ml	BXC0493A	LS
Alcohol, Ammonia, Carbonate Control (Level 2)	1 x 2 ml	BXC0494A	LS

Aldolase Calibrators & Controls

• Lyophilised product with a 3 year shelf life when stored at 2-8 °C.

Reconstituted stability of 5 days at 2-8°C.

Values assigned by reference laboratories in Europe

Description	Size	Cat No.	Туре
Aldolase Calibrator	3 x 1 ml	BXC0394A	Lyo.
Aldolase Control Elevated	3 x 1 ml	BXC0393A	Lyo.
Aldolase Control Normal	3 x 1 ml	BXC0392A	Lyo.

Quality Controls Catalogue | Fortress Diagnostics

Alpha-1-Acid Glycoprotein Calibrators & Controls

- Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.
- Open vial stability of 30 days, at 2-8°C.

Description

Alpha-1-Acid Glycoprotein Calibrator

Alpha-1-Acid Glycoprotein Control

Alpha-1-Antitrypsin Calibrators & Controls

•	Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.
•	Open vial stability of 30 days, at 2-8°C.

Description	Details
Alpha-1-Antitrypsin Calibrator	Immunoturbidimetric

Alpha-1-Antitrypsin Control

(Level 1 & Level 2)

Alpha-1-Microglobulin Calibrator

Alpha-1-Microglobulin Control

(Level 1 & Level 2)

Aluminium Calibrators & Controls

Liquid Stable (Ready-to-Use) stable to expiry at 2-8 °C.
 Open vial stability of 30 days, at 2-8°C.

Description

Aluminium Control Level 1

Aluminium Control Level 2

Aluminium Control Level 3

Ammonia Controls & Calibrators

- Liquid stable Calibrator and Control with a 2 year shelf life.
- Values are assigned in house using Enzymatic UV method.
- Storage at 2-8 °C.
- One month open vial stability at 2-8 °C.

Description

Ammonia Calibrator

Ammonia Control High

Ammonia Control Low

Angiotensin Converting Enzyme (ACE) Controls & Calib

Lyophilised for long shelf life, stable to expiry at 2-8 °C.
 Reconstituted stability of 7 days at 2-8°C.

Description

Angiotensin Converting Enzyme (ACE) Calibrator Angiotensin Converting Enzyme (ACE) Control (Level 1)

Angiotensin Converting Enzyme (ACE) Control (Level 2)

Anti-CCP Controls & Calibrators

- Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.
 When vial opened it must be used immediately, store at 2-8°C.
- Description

Anti-CCP Calibrator

Anti-CCP Calibrator Series

Anti-CCP Control Set

Fortress Diagnostics | Quality Controls Catalogue

Clinical Chemistry | SECTION 6

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Size	Cat No.	Туре
2 x 1 ml	BXC0892A	LS
2 x 1 ml	BXC0891A	LS

Size	Cat No.	Туре
1 x 2 ml	BXC0711A	LS
2 x 1 x 2ml	BXC0712A	LS
1 x 1 ml	BXC0887A	LS
2 x 1 x 1ml	BXC0889A	LS

Size	Cat No.	Туре	
1 x 1 ml	BXC0696A	LS	
1 x 1 ml	BXC0697A	LS	
1 x 1 ml	BXC0698A	LS	

	Size	Cat No.	Туре
-	3 x 2 ml	BXC0373A	LS
3	3 x 2 ml	BXC0375A	LS
	3 x 2 ml	BXC0374A	LS
bratoi	rs		

Size	Cat No.	Туре	
3 x 1 ml	BXC0177A	Lyo.	
3 x 1 ml	BXC0178A	Lyo.	
3 x 1 ml	BXC0179A	Lyo.	

с.			
	Size	Cat No.	Туре
	1 x 1 ml	BXC0379D	LS
	6 x 1 x 1ml	BXC0379B	LS
	2 x 1 x 1ml	BXC0379C	LS

 Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. When vial opened it must be used immediately, store at 2-8°C. 				
Description	Size	Cat No.	Туре	
Anti-Mullerian Hormone (AMH) Calibrators & Control Set	Calibrator Set: 2 x 1x0.5ml ; Control Set: 2x1x0.5ml	BXC0999A	LS	
APO A1/B Calibrator Series				(
Description	Size	Cat No.	Туре	
APO A1/B Calibrator Series	6 x 1 x 1 ml	BXC0413A	LS.	
ASO & RF Control				(
Description	Size	Cat No.	Туре	
ASO & RF Control	3 x 1ml	BXC0640A	LS.	
 Values are assigned by in house methods and instruments 100% human material designed for matrix conformity. The liquid stable ASO Single point calibrator has a shelf life of 2 ye Once opened the calibrator is stable for a period of 30 days when 		8 °C.		
Description	Size	Cat No.	Туре	
ASO (Single Point) Calibrator	1 x 1 ml	BXC0323A	LS	
ASO (Single Point) Calibrator	3 x 1 ml	BXC0323B	LS	
ASO, CRP, RF Control				
 This control is presented in a liquid stable format and is stable up The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up 	cored without contamination at 2-8	°C.		
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description	ored without contamination at 2-8 p to expiry at 2-8 °C. Size	Cat No.	Туре	
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description	cored without contamination at 2-8 p to expiry at 2-8 °C.		Type LS	
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control	ored without contamination at 2-8 p to expiry at 2-8 °C. Size	Cat No.		
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control Reta-2- Microglobulin Calibrators & Controls	ored without contamination at 2-8 p to expiry at 2-8 °C. Size	Cat No.		
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control Description Description	cored without contamination at 2-8 p to expiry at 2-8 °C. Size 3 x 1 ml	Cat No. BXC0645A	LS	
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control eta-2- Microglobulin Calibrators & Controls Description Beta-2- Microglobulin Calibrator	cored without contamination at 2-8 p to expiry at 2-8 °C. Size 3 x 1 ml Size	Cat No. BXC0645A Cat No.	LS Type	(
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control Beta-2- Microglobulin Calibrators & Controls Beta-2- Microglobulin Calibrator Beta-2- Microglobulin Calibrator	cored without contamination at 2-8 p to expiry at 2-8 °C. Size 3 x 1 ml Size 1 x 1 ml	Cat No. BXC0645A Cat No. BXC0647A	LS Type LS	
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description	cored without contamination at 2-8 p to expiry at 2-8 °C. Size 3 x 1 ml Size 1 x 1 ml	Cat No. BXC0645A Cat No. BXC0647A	LS Type LS	
 The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity. Once opened the control is stable for a period of 30 days when st This material is presented in a liquid stable format and is stable up Description ASO, CRP, RF Control Beta-2- Microglobulin Calibrators & Controls Description Beta-2- Microglobulin Calibrator Beta-2- Microglobulin Calibrator Beta-2- Microglobulin Control (Level 1 & Level 2) Bile Acid Controls & Calibrators Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. 	cored without contamination at 2-8 p to expiry at 2-8 °C. Size 3 x 1 ml Size 1 x 1 ml	Cat No. BXC0645A Cat No. BXC0647A	LS Type LS	(

2 x 1 x 1 ml

2 x 1 x 1 ml

Blood Alcohol Controls & Calibrators

Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. . When vial opened it must be used immediately, store at 2-8°C.

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Description	Size	Cat No.	Туре
Blood Alcohol Calibrator	1 x 2 ml	BXC0488A	LS
Blood Alcohol Control (Level 1)	1 x 2 ml	BXC0489A	LS
Blood Alcohol Control (Level 2)	1 x 2 ml	BXC0490A	LS
	TAE IIK	Briconsort	20

Bilirubin Controls & Calibrators

- Human Serum based Controls simulating the human serum matrix.
- •
- . •
- Storage at 2-8 °C.
- Reconstituted stability of 5 days at 2-8 °C.
- Values for Total and Direct Bilirubin. .
- Ideal for laboratories measuring Neonatal Bilirubin. Open vial stability for liquid 3 months.

Description	Size	Cat No.	Туре
Bilirubin Calibrator	3 x 1ml	BXC0319A	Lyo.
Bilirubin Control (Elevated)	1 x 1ml	BXC0318A	Lyo.
Bilirubin Control (Elevated)	5 x 1ml	BXC0318B	Lyo.
Bilirubin Calibrator	2 x 1ml	BXC0303A	LS
Bilirubin Control Set	2 x 1 x 1ml	BXC0304A	LS
Bilirubin Control Set, Paediatric	2 x 1 x 1ml	BXC0305A	LS
Bilirubin Control (Normal)	1 x 1ml	BXC0306A	LS
Bilirubin Control (Normal)	3 x 1ml	BXC0306B	LS
Bilirubin Control (Elevated)	1 x 1 ml	BXC0307A	LS
Bilirubin Control (Elevated)	3 x 1 ml	BXC0307B	LS

Citrate (Urinary) Calibrators & Controls

The controls are presented in a Liquid format and are stable up to expiry at 2-8 °C. • . Open vial stability of 30 days, at 2-8°C.

Description

Citrate (Urinary) Calibrator [200mg/L]

Citrate (Urinary) Control (Level 1) [50mg/L]

Citrate (Urinary) Control (Level 2) [600mg/L]

CO₂ (Bicarbonate) Controls & Calibrators

- Values are assigned by in house methods using Enzymatic CO2 kits.
- The CO2 (Bicarbonate) calibrator is intended for calibrating the Fortress CO2 (Bicarbonate) kit.
- CO2 (Bicarbonate) Calibrator & Controls are Liquid Stable.
- Stable to expiry at 2-8 °C.
 Five hours open vial stability at 2-8 °C.

Description

- CO2 (Bicarbonate) Calibrator
- CO2 (Bicarbonate) Control Low

CO2 (Bicarbonate) Control High

LS

LS

BXC0583A

BXC0584A

Bile Acid Control (Level 1)

Bile Acid Control (Level 2)

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Values assigned by reference laboratories in Europe. Typical levels: Total Bilirubin around 350 umol/l (around 20 mg/dl) & Direct Bilirubin around 100 umol/l (around 6.0 mg/dl). Lyophilised product and Liquid Stable version ready-to-use stable up to expiry at 2-8 °C.

Size	Cat No.	Туре	
2 x 1 ml	BXC0136A	LS	
2 x 1 ml	BXC0137A	LS	
2 x 1 ml	BXC0139A	LS	

Size		Cat No.	Туре
3 x 2 m	l	BXC0155A	LS
3 x 2 m	l	BXC0156A	LS
3 x 2 m	l	BXC0157A	LS

CSF Controls

• Human CSF based controls.

- Method based values with reference traceability provided.
- Liquid stable (Ready-to-Use) Stable to expiry at 2-8 °C.
- Once opened, the control is stable for a period of 30 days when stored at 2-8 °C without contamination.

Analytes		
Electrolytes Chloride Lactate Sodium	Substrates Albumin Glucose	
Size	Cat No. T	јуре
1 x 1 ml	BXC0673A L	.S
1 x 1 ml	BXC0673B L	.5
	Electrolytes Chloride Lactate Sodium Size 1 x 1 ml	Electrolytes Substrates Chloride Albumin Lactate Glucose Sodium Cat No. T 1 x 1 ml BXC0673A L

CSF Diff Control

			-
Description	Size	Cat No.	Туре
CSF Diff Control (Level 1)	5 x 2 ml	BXC0080A	LS.
CSF Diff Control (Level 2)	5 x 2 ml	BXC0081A	LS.
CSF Diff Control (Level 3)	5 x 2 ml	BXC0082A	LS.

CRP Controls & Calibrators

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· Fortress CRP controls are intended for monitoring accuracy and precision of CRP turbidimetric, Immunoturbidimetric and Nephelometric assays. • This control is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.

• The values are assigned in European Reference Laboratories. 100% Human serum designed for matrix conformity.

• Once opened the control is stable for a period of 30 days when stored without contamination at 2-8 °C.

Description	Size	Cat No.	Туре
CRP (Single Point) Calibrator	3 x 1 ml	BXC0324A	LS
CRP (Multi Point) Calibrator	6 x 1 x 1 ml	BXC0324B	LS
CRP (Ultra Sensitive) Low Control	3 x 1 ml	BXC0325A	LS
CRP (Ultra Sensitive) Standard Set	5 x 1 x 0.5 ml	BXC0327A	LS
CRP Control (Level 1)	5 x 1 ml	BXC0326A	LS
CRP Control (Level 2)	5 x 1 ml	BXC0326B	LS
CRP Control Set	2 x 1 x 1 ml	BXC0326C	LS

Cyanmethaemoglobin Standard Set

• Typical concentrations 8,10,12,15,18 g/dL.

• The standard set comprises of 5 standards.

• Values are assigned using in house methods.

• Store the standard set at 2-8 °C. The Standard set is stable up to expiry when stored unopened at 2-8 °C.

• Once opened the standard set is stable for a period of 1 month when stored at 2-8°C without contamination. • Do not freeze

Description	Size	Cat No.	Туре
Cyanmethaemoglobin Standard Set	5 x 1 x 10 ml	BXC0483A	LS

Cystatin C Controls & Calibrators

• Values are assigned using in house methods.

• These controls and calibrators are supplied in liquid stable format with a shelf life of 2 years when stored at 2-8 °C. • When opened and stored at 2-8 °C, without contamination, the controls and calibrators are stable for a period of 1 month.

Mix gently before use.

• The calibrator set and controls are liquid stable. • Human serum based to ensure matrix compatibility.

• Do not freeze.

Description

Cystatin C Calibrator Set (6 Levels)

Cystatin C Control (Level 1)

Cystatin C Control (Level 2)

D3-Hydroxybutyrate Controls & Calibrators

Lyophylised product stable to expiry at 2-8 °C.

Reconstituted Stability of 7 Days at 2-8°C.

Description

D3-Hydroxybutyrate Calibrator

D3-Hydroxybutyrate Control (Level 1)

D3-Hydroxybutyrate Control (Level 2)

D-Dimer Calibrators & Controls

- Liquid Stable (Ready-to-use)
- Prepared from human serum.
- Open vial stability of 30 days, at 2-8°C.

Description

D-Dimer Calibrator Set (5 Levels)

D-Dimer Control Set (Level 1 & Level 2)

ESR Controls

 Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. • Open vial stability of 30 days, at 2-8°C.

Description

ESR Control (Level 1)

ESR Control (Level 2)

Ferritin Controls & Calibrators

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. • Open vial stability of 30 days, at 2-8°C.

Description

Ferritin	Calibrator	
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Ferritin Calibrator Set

Ferritin Control (Level 1)

Ferritin Control (Level 2)

Fructosamine Controls & Calibrators

• Values are assigned using both in house and reference laboratory values. • 100% Human Serum designed for matrix conformity.

Description

Fructosamine Calibrator

Fructosamine Control (Level 1)

Fructosamine Control (Level 2)

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Size	Cat No.	Туре	
6 x 1 ml	BXC0334A	LS	
2 x 1 ml	BXC0333A	LS	
2 x 1 ml	BXC0333B	LS	
			0

Size	Cat No.	Туре
1 x 3 ml	BXC0544A	Lyo
1 x 5 ml	BXC0543A	Lyo
1 x 5 ml	BXC0543B	Lyo
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Size	Cat No.	Туре
5 x 1 ml	BXC0788A	LS
5 x 1 ml	BXC0789A	LS

Size	Cat No.	Туре	
2 x 10 ml	BXC0631A	LS	
2 x 10 ml	BXC0632A	LS	
			0

1 x 1 ml BXC0445A LS 6 x 1 x 1 ml BXC0445B LS 3 x 1 ml BXC0443A LS	Size	Cat No.	Туре
	1 x 1 ml	BXC0445A	LS
3 x 1 ml BXC0443A LS	6 x 1 x 1 ml	BXC0445B	LS
	3 x 1 ml	BXC0443A	LS
3 x 1 ml BXC0444A LS	3 x 1 ml	BXC0444A	LS

	Size	Cat No.	Туре
	1 x 1 ml	BXC0592A	LS
Low	3 x 1 ml	BXC0593A	LS
High	3 x 1 ml	BXC0594A	LS

G-6-PDH Controls

- Values assigned by reference Laboratories in Europe.
- Stable to expiry at 2-8 °C.
- Reconstituted stability of 5 days at 2-8 °C.
- Lypholised for enhanced stability.
- Typical Levels:
 - G-6-PDH Deficient Control : 100 200 U/l G-6-PDH Normal Control: 900 - 1400 U/l

Description	Size	Cat No.	Туре
G-6-PDH Control Normal	3 x 1.0ml	BXC0573B	Lyo.
G-6-PDH Control Intermediate	1 x 1 ml	BXC0572B	Lyo.
G-6-PDH Control Deficient	3 x 1.0ml	BXC0572B	Lyo.

Glucose-Fructose Control

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Description	Size	Cat No.	Туре	
Glucose-Fructose Control (Level 1)	TBC	BXC0099A	LS	
Glucose-Fructose Control (Level 2)	TBC	BXC0099B	LS	

Glycerol Control

Size	Cat No.	Туре	
2 x 5 ml	BXC0278A	LS	

HDL/LDL Calibrators

Fortress HDL/LDL Cholesterol Calibrator is presented in a Lyophilised format for enhanced stability. Stable up to expiry at 2-8 °C.
 Once opened and reconstituted the control is stable for 7 days at 2-8 °C, or 30 days at -20 °C.
 100% human material designed for matrix conformity.

	Analytes		_
HDL Cholesterol (Direct)	LDL Cholesterol (Direct)		
Description	Size	Cat No. Type	
HDL/LDL Calibrator	5 x 3 ml	BXC0315C Lyo.	
HDL/LDL Calibrator	3 x 1 ml	BXC0315D LS.	
HDL/LDL Calibrator	1 x 1 ml	BXC0315E LS.	
HDL/LDL Calibrator	3 x 1 ml	BXC0315F Lyo.	

IgE Calibrators & Controls

- Lyophilised product stable to expiry at 2-8 °C.
- Reconstituted stability of 7 days at 2-8°C.
- Reconstituted stability of 30 days at -20°C.

Description	Size	Cat No.	Туре
IgE Calibrator	1 x 1 ml	BXC0752A	Lyo.
IgE Control Set (Level 1 & Level 2)	2 x 2 x 1 ml	BXC0753A	Lyo.

Immunoglobulin Calibrator

* Liquid Stable (Ready-to-use) stable to expiry at 2-8 $^\circ\mathrm{C}.$

Open vial stability of 30 days, at 2-8°C.
Prepared from Human Serum.

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AgI	IgG	IgM	l
Description	Size	Cat No.	Туре
mmunoglobulin Calibrator	3 x 1 ml	BXC0040A	LS
Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.			
 Pase Calibrator Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C. 	Size	Cat No.	Туре
 Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C. 	Size 3 x 1 ml	Cat No. BXC0510A	Type LS

- Liquid stable controls and calibrators stable up to expiry at 2-8 °C.
 Once opened the calibrator and controls are stable for a period of 30 days when stored at 2-8 °C.
 Lipoprotein Low and High controls typically have the following the levels:
- Lipoprotein Control Low around 20mg/dl.
- Lipoprotein Control High around 50mg/dl.

Do not freeze.

Description

Lipoprotein (a) Calibrator

Lipoprotein (a) Control Low

Lipoprotein (a) Control High

Lithium Calibrator

- Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.
- Open vial stability of 30 days, at 2-8°C.

Description

Lithium Calibrator (3000 mmol/L)

Lithium Calibrator

Methanol Calibrators & Controls

Description Methanol Calibrator Methanol Control (Level 1) Methanol Control (Level 2)

Microalbumin Calibrators & Controls

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. • Open vial stability of 30 days, at 2-8°C.

Description

Microalbumin Calibrator Series

Microalbumin Control (Level 1)

Microalbumin Control (Level 2)

Fortress Diagnostics | Quality Controls Catalogue

Clinical Chemistry | SECTION 6

Size	Cat No.	Туре	
1 x 1 ml	BXC0134A	LS	
1 x 1 ml	BXC0131A	LS	
1 x 1 ml	BXC0133A	LS	

Size	Cat No.	Туре
2 x 1 ml	BXC0106A	LS
1 x 2 x 1 ml	BXC0141A	LS

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Size	Cat No.	Туре
1 x 2ml	BXC0497A	LS
1 x 2ml	BXC0498A	LS
1 x 2ml	BXC0499A	LS

Size	Cat No.	Туре	
5 x 1 ml	BXC0329A	LS	
2 x 1 ml	BXC0328A	LS	
5 x 1 ml	BXC0328B	LS	

Myoglobin Calibrators & Controls

 Lyophilised product stable to expiry at 2-8 °C. Reconstituted stability of 30 days at 2-8°C. 			
Description	Size	Cat No.	Туре
Myoglobin Calibrator Series	1 x 1 x 0.5ml	BXC0486A	Lyo.
Myoqlobin Control	1 x 0.5ml	BXC0487A	Lvo.

Oxalate Calibrators & Controls

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.

 Open via 	al stability of 30 days, at 2-8°C.	
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Description	Size	Cat No.	Туре
Oxalate (Urinary) Calibrator [0.5 mmol/L]	1 x 5ml	BXC0147A	LS
Oxalate (Urinary) Control (Level 1) [0.2 mmol/L]	1 x 5ml	BXC0148A	LS
Oxalate (Urinary) Control (Level 2) [0.7 mmol/L]	1 x 5ml	BXC0149A	LS

RF (Multi-Point) Calibrators

- This Calibrator is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.
 The Calibrator is presented as a single level calibrator which can be diluted as necessary for calibrating RF Turbidimetric Assays.
 Values are assigned by in house methods and instruments.

- 100% human material designed for matrix conformity.
 Once opened the control is stable for a period of 30 days. when stored without contamination at 2-8 °C.

Description	Size	Cat No.	Туре
RF (Multi-Point) Calibrator	1 x 2ml	BXC0612A	LS
RF (Multi-Point) Calibrator	3 x 2ml	BXC0612B	LS

TIBC Calibrator

Zinc & Copper Calibrator

Lyophilised product stable to expiry at 2-8 °C.
Reconstituted stability of 30 days at 2-8°C.

Description	Size	Cat No.	Туре
Zinc & Copper Calibrator	2 x 1 ml	BXC0463A	Lyo.





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Туре

Lyo.



THROMBO

SECTION 7 Coagulation / Haemostasis

Coagulation / Haemostasis Controls

Coagulation and haemostasis controls are essential in diagnostics for accurately assessing a patient's ability to form blood clots and maintain haemostasis. These controls include standardized samples with known coagulation properties used to validate laboratory tests that measure parameters like prothrombin time (PT), activated partial thromboplastin time (aPTT), and platelet counts. By employing these controls, laboratories can ensure the accuracy and reliability of test results, which are critical for diagnosing bleeding disorders, monitoring anticoagulant therapy, and guiding treatment decisions in various clinical scenarios, such as surgery or trauma care.

The importance of coagulation and haemostasis controls extends to maintaining quality assurance within laboratory settings, allowing for consistent and reproducible results across different testing environments. Reliable coagulation diagnostics are crucial for preventing complications related to abnormal bleeding or thrombosis, ultimately improving patient outcomes. Regular use of these controls helps identify any technical issues with reagents or instrumentation, ensuring that healthcare providers have accurate information to make informed decisions regarding patient management and treatment strategies.

The Fortress Haemostasis Plasma Controls are provided in a lyophilised format for enhanced stability and are 100% human serum based to negate matrix effects. The controls are supplied with instrument and method specific assigned values.



Plasma Calibrator & Controls

For Coagulation Series only, PT, aPTT, Fibrinogen, Thrombin Time

• The calibrator and controls are supplied in lyophilised format and stable to expiry when stored at 2-8 °C.

- Once opened and reconstituted, the calibrators and controls are stable for a period of 24 hours at 2-8 °C.
- 100% Human plasma to mimic human samples.
- Do not Freeze

Description	Size	Cat No.	Туре
Plasma Calibrator	5 x 1 ml	COAG125A	Lyo.
Plasma Control (Level 1)	1 x 1 ml	COAG108A	Lyo.
Plasma Control (Level 1)	5 x 1 ml	COAG108B	Lyo.
Plasma Control (Level 2)	1 x 1 ml	COAG109A	Lyo.
Plasma Control (Level 2)	5 x 1 ml	COAG109B	Lyo.
Plasma Control Set (Level 1 & Level 2)	2 x 5 x 1 ml	COAG112A	Lyo.



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CUSTOMISED QUALITY CONTROLS

SECTION 8

Customised Quality Controls

Fortress Diagnostics has extensive experience and expertise in the production of customised quality controls for EQA programmes, and manufaturing custom made QC solutions for laboratories throughout the global market.

Over 200 analytes can be customized in the required matrix, with concentrations, levels and volumes. The following list of quality controls can be customised by Fortress Diagnostics:

- Antioxidant Controls Clinical Chemistry Cardiac Controls Haemostasis Controls **Diabetes Controls** Immunoassay Controls
- Immunology / Specific Protein Controls Lipids Tumor Markers TDM and DOA **Urine Assayed Urine Strip Controls**

Why Customise Fortress Controls?



Cost Effective

Fortress Diagnostics operate throughout the global market. Over 200 analytes can be customized in the required matrix, with concentrations, levels and volumes, leading to cost savings.



Customised

Quality

The customised advantage helps in getting the laboratory's requirement in analytes and levels in one control. High usage controls can be manufactured in higher volumes.



Choice of Matrix Controls are available in different matrices, like Serum, Plasma, Whole Blood, Urine and aqueous material.

All controls are CE marked and manufactured according to the ISO13485 specifications.



Fast Lead Time

The total order, testing and manufacturing time is 60 days. This cuts the ordering of customised control to half the time that is required by other companies to manufacture customised controls.











Diabetes

Diabetes controls are crucial in diagnostics for ensuring the accuracy and reliability of tests that measure blood glucose levels and other related biomarkers, enabling timely diagnosis, effective monitoring of glycemic control, and informed treatment decisions to manage diabetes and prevent complications.

The Fortress Diabetes Controls are lyophilised for enhanced stability, and are ideal for both clinical and research use. The controls are supplied with instrument and method specific assigned values.



Fructosamine Controls & Calibrators

• Values are assigned using both in house and reference laboratory values.

- 100% Human Serum designed for matrix conformity.
- Lyophilised Controls and Calibrators stable to expiry at 2-8 °C.
 Reconstituted stability of 28 days at -20 °C.

Description	Size	Cat No.	Туре
Fructosamine Calibrator	1 x 1 ml	BXC0592A	LS
Fructosamine Control (Level 1)	3 x 1 ml	BXC0593A	LS
Fructosamine Control (Level 2)	3 x 1 ml	BXC0594A	LS

HbA1c Calibrators

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- 100% human blood designed for matrix conformity.
- The Calibrator is supplied in a Lyophilised format with a 2 year shelf life when stored at 2-8 °C.
- Reconstituted stability of 5 days at 2-8 °C.
- HbA1c Calibrator Set: Typical levels:
- HbA1c Calibrator Level 1: 5-6%
- HbA1c Calibrator Level 2: 11-13% • Do not freeze.

Description		Size	Cat No.	Туре
HbA1c Calibrator Series	For use with BXC0670	2 x 1 x 0.5 ml	BXC0668A	Lyo.
HbA1c Calibrator Set (2 Levels)		2 x 1 x 0.5 ml	BXC0678A	Lyo.
HbA1c (HPLC) Calibrator Set (2 Levels)	For HPLC Method Only	2 x 1 x 0.25 ml	BXC0780A	Lyo.

HbA1c Controls

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• Controls are supplied in a Lyophilised format stable to expiry at 2-8 °C.

The controls are value assigned in European Reference Laboratories.
 The Control carries values for methods such as Immunoturbidimetry, Micro-column, HPLC and Enzymatic HbA1c & point of care devices.

• Values are provided for both NGSP DCCT and IFCC methodologies.

Description		Size	Cat No.	Туре
HbA1c (HPLC) Control Set (2 Levels)	For HPLC Method Only	2 x 1 x 0.5 ml	BXC0779A	Lyo.
HbA1c Control High		2 x 0.5 ml	BXC0677A	Lyo.
HbA1c Control Low		2 x 0.5 ml	BXC0676A	Lyo.
HbA1c Control Set		2 x 2 x 0.5 ml	BXC0675A	Lyo.
HbA1c Control Set (2 Levels)	For use with BXC0670	2 x 1 x 0.5 ml	BXC0669A	Lyo.

Thalassaemia (Alpha & Beta) Control



For use on all major systems and methods including HPLC, Immunoassay, HbA2, HbF, HbS.

Designed for monitoring accuracy and precision of Haemoglobin Variants measurements associated with Thalassaemia.

• The controls are supplied in a Lyophilised format stable up to expiry at 2-8 °C.

• 100% Human blood.

Description	Size	Cat No.	Туре
Thalassaemia (Alpha & Beta) Control (Level 1)	2 x 0.5 ml	BXC0665A	Lyo.
Thalassaemia (Alpha & Beta) Control (Level 2)	2 x 0.5 ml	BXC0665B	Lyo.



Quality Controls Catalogue | Fortress Diagnostics

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Drugs of Abuse

Drugs of abuse controls are vital in diagnostics for accurately detecting and monitoring the use of illicit substances, prescription medications, and other psychoactive drugs in patients. These controls consist of standardized samples with known concentrations of specific drugs or their metabolites, used to validate the performance of drug testing assays, such as immunoassays and gas chromatography. By implementing these controls, laboratories ensure the reliability and accuracy of test results, which are crucial for effective patient management in various settings, including addiction treatment, pain management, and legal cases.

The role of drugs of abuse controls extends beyond accuracy; they are essential for maintaining quality assurance in drug testing laboratories. Regular use of these controls helps identify potential errors, such as reagent degradation or instrument malfunction, thus preventing false positives or negatives that could impact patient care or legal outcomes. Moreover, reliable drug testing plays a critical role in public health initiatives, workplace drug screening, and monitoring compliance with treatment programs, ultimately contributing to safer and more effective healthcare practices.

The Fortress Drugs of Abuse Controls, are available in a liquid stable (ready-to-use) format for convenience and easeof-use, offer an independent assessment of performance for Drugs of Abuse Immunoassays, and are suitable for semiquantitative and quantitative use. The controls are supplied with instrument and method specific assigned values.



Drugs of Abuse Controls

• Fortress Drugs of Abuse controls are intended for checking assay performance of Drugs of Abuse Immunoassays.

These controls are urine based and have concentrations 25% above and below the SAMHSA (Substance Abuse And Mental Health Services Administration) recommended cut off levels.

Liquid stable controls (ready-to-use).

• Suitable for semi-quantitative and quantitative use.

• 2 years shelf life.

30 days open vial stability when stored at 2-8 °C.

Typical Values in Drugs of Abuse Control Levels

Analyte	DOA Level (ng/ml)	DOA Level 2 (ng/ml)	DOA Level 3 (ng/ml)
Amphetamines	600	900	1350
Barbiturates	100	400	1000
Benzodiazepines	102	250	600
Cannabinoids	50	90	150
Cocaine	300	275	400
LSD	0.05	0.1	0.2
Methadone	200	300	800
Opiates	1000	1500	6000
Phencyclidine	15	25	100
Propoxyphene	60	100	-
тса	50	90	180

	Anal	ytes	
Amphetamines Barbiturates Benzodiazepines	Cannabinoids Cocaine TCA	LSD Methadone Opiates	Phencyclidine Propoxyphene

Description

 Drugs of Abuse Control (Level 1)

 Drugs of Abuse Control (Level 1)

 Drugs of Abuse Control (Level 2)

 Drugs of Abuse Control (Level 2)

 Drugs of Abuse Control (Level 3)

 Drugs of Abuse Control (Level 3)

Drugs of Abuse | SECTION 10

Size	Cat No.	Туре
10 x 10 ml	BXC0784A	LS
10 x 3 ml	BXC0784B	LS
10 x 10 ml	BXC0785A	LS
10 x 3 ml	BXC0785B	LS
10 x 10 ml	BXC0786A	LS
10 x 3 ml	BXC0786B	LS





Haematology

Hematology controls are crucial in diagnostics for ensuring the accuracy and reliability of tests that analyse blood components, including red and white blood cells, platelets, and hemoglobin levels. These controls consist of standardised samples with known cell counts and characteristics, used to validate laboratory instruments and procedures for tests like complete blood counts (CBC) and coagulation profiles. By implementing these controls, laboratories can detect any discrepancies in test results, which is essential for diagnosing conditions such as anemia, leukemia, and other blood disorders, as well as for monitoring patient response to treatment.

The role of hematology controls extends to maintaining quality assurance within laboratory settings, allowing for consistent and reproducible results across different testing environments. Reliable hematology diagnostics are vital for guiding clinical decision-making and ensuring appropriate patient management. Regular use of these controls helps identify potential issues with reagents or equipment, thereby minimising the risk of misdiagnosis or inappropriate treatment. Overall, hematology controls play a significant role in enhancing the quality of healthcare by providing accurate diagnostic information that informs treatment strategies and improves patient outcomes.

The Fortress Haematology Controls combines 14 analytes, covering the most commonly tested blood profiles in a single control. With 3-part differential technology an unbiased, independent assessment of analytical performance. The controls are supplied with instrument and method specific assigned values.



Haematology Controls & Calibrators

• Fortress Haematology calibrators are intended for calibrating 3 part differential analysers.

- The controls are intended for monitoring accuracy and precision in 3 part differential analysers.
- The product is supplied as a liquid stable material with storage at 2-8 °C.
- When opened, this product is stable for a period of 21 days, when stored at 2-8 °C.

Values Available for the Following Analysers:

Abbott Cell Dyn 1600/1700/1800, ABX Micros 60, 80/80XL, Coulter AcT 5 Diff, Coulter ACT Diff/ACT Diff 3, Coulter MD Series, Coulter STKS/MAXM/ HMX, Coulter T890/T660/T540, Danam DATACELL 16CP/I1800/18/18MS/18MS Plus, Siemens ADVIA 120/2120/1210i, Sysmex XE2100/XE5000/ XT1800i/XT2000i/XS1000i, Sysmex KX21/K1000, ERMA PCE210, Merck Medonic CA620, Mindray BC1800/BC3000/BC-3000CT/BC3200/BC3000 Plus/ BC2600/BC2800, Nihon Kohden CELLTAC Alpha MEK 6318, MEK 6410K, Fortress CellDiff 3, Swelab AC920, Medonic CA610, Abacus Junior.

Analytes					
Haemoglobin HCT MCH MCHC	MCV MPV Platelets RDW CV	RDW SD Total RBC Total WBC Granulocytes		mphocytes Mid Cells	
Description		Size	Cat No.	Туре	
Haematology Calibrator		3 x 3 ml	HAEMC005	LS	
Haematology Control (Level 1)		3 x 3 ml	HAEMC001	LS	
Haematology Control (Level 2)		3 x 3 ml	HAEMC002	LS	
Haematology Control (Level 3)		3 x 3 ml	HAEMC003	LS	
Haematology Control (3 Levels)		3 x 3 x 3 ml	HAEMC004	LS	





Immunoassay

Immunoassay controls are essential in diagnostics for ensuring the accuracy and reliability of tests that detect and quantify specific proteins, hormones, drugs, and antibodies in biological samples. These controls consist of standardised samples with known concentrations of analytes, which are used to validate the performance of immunoassay methods such as enzyme-linked immunosorbent assays (ELISA) and radioimmunoassays. By incorporating these controls, laboratories can monitor assay precision, sensitivity, and specificity, thus minimising the risk of false-positive or false-negative results that could lead to incorrect diagnoses or inappropriate treatment decisions.

The role of immunoassay controls extends to maintaining quality assurance in laboratory operations, ensuring that test results are consistent and reproducible across different assays and testing platforms. Reliable immunoassay diagnostics are critical for guiding clinical decisions, such as monitoring hormone levels, diagnosing infectious diseases, and screening for autoimmune disorders. Regular use of these controls helps identify any technical issues with reagents, instrumentation, or assay protocols, ultimately supporting accurate and timely patient management. Overall, immunoassay controls significantly enhance the quality of healthcare by providing trustworthy diagnostic information that informs treatment strategies and improves patient outcomes.

The Fortress Immunoassay Controls Range have been designed to cover a variety of the most comprehensive markers, including: thyroid hormones, fertility hormones, tumour markers, growth deficiency, allergy, diabetes, anaemia and steroids. All Fortress controls in the Immunoassay range are manufactured from 100% human serum to negate matrix effects. The controls are supplied with instrument and method specific assigned values.



Immunoassay Controls

Fortress Immunoassay Controls are intended for monitoring accuracy and precision.

• Controls are provided in a Lyophilised format stable up to expiry at 2-8 °C.

Reconstituted stability of 7 days at 2-8 °C or 4 week at -20 °C.

· Values are assigned in European reference laboratories.

28 analytes.

Analytes						
Thyroid Hormones Total T3 Total T4 Free T3 Free T4 TSH T-uptake Fertility Hormones LH FSH Prolactin HCG	Tumor Markers CEA AFP PSA fPSA Cardiac Markers Digoxin Growth Deficiency Human Growth Hormone	Allergy IgE Diabetes C-peptide Insulin Anaemia Ferritin	1 Prog C Unconju Prog Test	eroids 7-OH lesterone ortisol gated Estriol lesterone osterone HEA-S		
Description		Size	Cat No.	Туре		
Immunoassay Control (Level 1)		4 x 3 ml	BXC0363A	Lyo.		
Immunoassay Control (Level 2)		4 x 3 ml	BXC0363B	Lyo.		
Immunoassay Control (Level 3)		4 x 3 ml	BXC0363C	Lyo.		
Immunoassay Tri-Level Control		3 x 4 x 3 ml	BXC0363D	Lyo.		

Immunoassay Speciality

• Lypholised stable to expiry at 2-8 °C

100% Human serum

Reconstituted stability of 7 days at 2-8 °C or 28 days stored at -20 °C.

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1-25 (OH) Vitamin D Anti TG Anti TP

Description

Immunoassay Speciality (Level 1)

Immunoassay Speciality (Level 2)

Immunoassay Speciality (Level 3)

Immunoassay Speciality Tri-Level Control

Anti-Mullerian Hormone (AMH) Calibrators & Controls

Lyophilised for long shelf life, stable to expiry at 2-8 °C.

- Reconstituted stability of 7 days at 2-8°C.
- Compatable with all AMH assays.

Description

Anti-Mullerian Hormone (AMH) Calibrators & Controls

hCG Serum Control

- Fortress hCG Serum controls are intended for monitoring the accurate
- Controls are supplied as lyophilised materials stable up to expiry at 2-8 °C.
- The controls are available in two levels positive and negative
- Value assignment of the Positive Controls is performed in European reference laboratories.
- 100% Human serum based for matrix conformity.

Description		Size	Cat No.	Туре
hCG Serum Control	Positive	1 x 1 ml	BXC0681A	LS.
hCG Serum Control	Negative	1 x 1 ml	BXC0682A	LS.
hCG Serum Control	Positive	1 x 5 ml	BXC0683A	LS.

H-Pylori Control

 Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C.

Description

H-Pylori Control (2 Levels)

Tumour Marker Controls

For use on all major systems & methods including Immunoassay Instruments

- Fortress Tumour Marker controls are multi analyte controls designed for monitoring accuracy.
- The controls are supplied in a Lyophilised format for enhanced stability. • 100% human serum.
- Reconstituted stability of 10 days at 2-8 °C and 30 days at 20 °C.
- Stable up to expiry at 2-8 °C.

Size Cat No. Type	Description
5 x 2 ml BXC0792A Lyo.	Tumour Marker Control (Level 1)
5 x 2 ml BXC0792B Lyo.	Tumour Marker Control (Level 2)
2 x 1 x 2ml BXC0793A Lyo	Tumour Marker Control SET (2-Levels)
5 x 2 ml BXC0792B I	Tumour Marker Control (Level 2)

Tumour Marker Control (Level 1)
Tumour Marker Control (Level 2)
Tumour Marker Control SET (2-Levels)

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Immunoassay / Infectious Diseases SECTION 12

Analy	tes		
C Pept IgF- Insul	1	Oesteocalci PTH Calcitonin	
	Size	Cat No.	Туре
	1 x 2 ml	BXC0364A	Lyo.
	1 x 2 ml	BXC0364B	Lyo.
	1 x 2 ml	BXC0364C	Lyo.
	3 x 1 x 2 ml	BXC0364D	Lyo.
S			<u></u>
	Size	Cat No.	Туре
	Cal A: 1 x 0.5ml; Cal B-G: 6 x 0.5ml; Ctl L1: 1 x 0.5ml; Ctl L2: 1 x 0.5ml	BXC0999A	Lyo.
			0 8 4
ccuracy	and precision of hCG assays in both o	quantitative and qua	alitative formats.

Positive control has a hCG concentration of >/= 20 lU/ml, making it ideal for sensitivity testing of gualitative assays.

Size	Cat No.	Туре
2 x 1 x 1 ml	BXC0684A	LS

Analytes

SECTION 12 Immunoassay / Infectious Diseases

Vitamin-D Calibrators & Controls

• Lyophilised product stable to expiry at 2-8 °C.

 Reconstituted stability of 30 days at 2-8° 	c.	
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Description	Size	Cat No.	Туре
Vitamin-D Calibrator Set (5 Levels)	5 x 1 x 1ml	BXC0475A	LS.
Vitamin-D Control Set (2 levels)	2 x 1 x 1ml	BXC0474A	LS.

Infectious Diseases

Infectious diseases controls are crucial in diagnostics for ensuring the accuracy and reliability of tests that identify and quantify pathogens, such as bacteria, viruses, fungi, and parasites, in biological samples. These controls consist of standardised samples containing known concentrations of specific pathogens or their genetic material, which are used to validate the performance of diagnostic assays like polymerase chain reaction (PCR) tests and enzyme immunoassays. By incorporating these controls, laboratories can monitor the sensitivity and specificity of tests, thereby reducing the risk of false-negative or false-positive results that could impact patient management and public health responses.

The role of infectious diseases controls extends to maintaining quality assurance in laboratory settings, ensuring consistent and reproducible results across different testing environments and methodologies. Reliable diagnostics for infectious diseases are essential for timely treatment, outbreak control, and infection prevention strategies. Regular use of these controls helps identify potential issues with reagents, equipment, or testing protocols, which is critical for maintaining the integrity of diagnostic results. Overall, infectious diseases controls significantly enhance the quality of healthcare by providing accurate diagnostic information that informs treatment decisions and public health initiatives, ultimately improving patient outcomes and community health safety.

The Fortress Infectious Diseases Controls are available in a liquid stable format for convenience and ease-of-use, being made from 100% human serum to negate matrix effects. The controls are supplied with instrument and method specific assigned values.



COVID-19 Anitgen Controls

Description	Size	Cat No.	Туре
COVID-19 Anitgen Control Pair	2 x 1 x 0.5ml	COVIDNPC	LS

HBsAb Control Panel

- 100% Human serum from single donors.
- The panel is supplied in 5 levels.

• Open vial stability of 60 days when stored at 2-8 °C without contamination.

• HBsAg Controls have the following concentrations; L1 0.2IU/ml, L2 0.5IU/ml, L3 1.0IU/ml, L4 2.0IU/ml, L5 4.0IU/ml.

Description	Size	Cat No.	Туре
HBsAb Control Panel (2 Levels)	2 x 1 x 0.5ml	BXC0805A	LS
HBsAb Control Panel (5 Levels)	5 x 1 x 0.5ml	BXC0800A	LS

HCV Antibody Control Panel

• The control panel is supplied as a liquid panel stable up to expiry at 2-8 °C.

 100% Human serum • The panel is supplied in 4 levels.

• Open vial stability: 30 days at 2-8 °C without contamination.

Description	Size	Cat No.	Туре
HCV Antibody Control Panel (4 Levels)	4 x 1 x 0.5ml	BXC0804A	LS

HE4 Description HE4 (Level 1) HE4 (Level 2)

HIV-I Antibody Control Panel

- The control panel is supplied as a liquid panel stable up to expiry a
- 100% Human serum • The panel is supplied in 4 levels.
- Open vial stability: 30 days at 2-8 °C without contamination.

Description

HIV-I Antibody Control Panel (4 Levels)

HIV-I Antigen (p24) Control Panel

- The Control Panel is supplied as a liquid panel stable up to expiry 100% Human serum from single donors.
- The panel is supplied in 4 levels.
- Open vial stability: 30 days when stored at 2-8 °C without contamination
- Level 1 1.25U/m
- , 2.50U/ml Level 2
- 5.00U/ml Level 3
- Level 4 10.00U/ml

Description

HIV-I Antigen (p24) Control Panel (4 Levels)

IFN-y Control

Description

IFN-y Control

Procalcitonin Control

Description

Procalcitonin Control Set (2 Levels)

Protein Control

- This control is presented in a liquid stable format and is stable up
- Values are assigned by reference laboratories based in Europe.
- Values for Immunoglobulin, Inflammatory proteins, Complements 100% Human material designed for matrix conformity.
- Once opened the control is stable for a period of 30 days when st

Description

Protein Control (Level 1)

Protein Control (Level 2)

Protein Control Set (Level 3)

Protein Control Set (Level 1 , Level 2 & Level 3)

Retinol Binding Protein Control

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C.

Description

Retinol Binding Protein Control

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Size	Cat No.	Туре	
1 x 0.25ml	BXC0425A	Lyo.	
1 x 0.25ml	BXC0425B	Lyo.	
		80	
/ at 2-8 °C.			
Size	Cat No.	Туре	
4 x 1 x 1ml	BXC0802A	LS	
		80	
y at 2-8 °C.			
ination.			
Size	Cat No.	Туре	
4 x 1 x 0.5 ml	BXC0803A	LS	
		<u></u>	
Size	Cat No.	Туре	
3 x 1 ml	BXC0702A	Lyo.	
		<u>_</u>	
Size	Cat No.	Туре	
2x1x1ml	BXC0590B	Lyo.	
	2	_,	
		0	
p to expiry when stored at 2-8 °C.			
ts, Carrier and Storage proteins are availab	le in the control		
	te in the control.		
stored without contamination at 2-8 °C.	Cable	Trees	
Size	Cat No.	Туре	
1 x 1 ml	BXC0638A	LS	
1 x 1 ml 1 x 1 ml	BXC0639A BXC0640A	LS	
3 x 1 x 1ml	BXC0640A BXC0641A	LS	
	DAC004TA	LJ	
Size	Cat No.	Туре	
1 x 1 ml	BXC0995A	LS	

SECTION 12 | Immunoassay / Infectious Diseases

 The Control Panel is supplied as a liquid panel stable 100% Human serum from single donors. The panel is supplied in 6 levels. Open vial stability: 30 days when stored at 2-8 °C with 				
escription	Size	Cat No.	Туре	
/philis Control Panel (6 Levels)	6 x 1 x 0.5ml	BXC0806A	LS	
rch Controls Negative				

Description	Size	Cat No.	Туре
TORCH Control Negative	3 x 1 ml	BXC0799A	LS

Torch Controls Positive

The Fortress TORCH Control Positive is intended for use in monitoring precision of TORCH Assays.
The control is supplied in a liquid stable format.
The control is stable up to expiry at 2-8 °C.
Values are assigned in European Reference Laboratories.
TORCH-IgM controls are supplied individually to prevent cross-reactivity.
TORCH-IgG controls are supplied together / vial.

Analytes			
IgG Toxoplasma Gondii IgG	HSV -2 IgG VZV IgG	IgM Toxoplasma Gondii IgM	HSV-2 IgM
Rubella IgG	EBV IgG	Rubella IgM	Syphilis RPR
CMV IgG HSV-1 IgG	Syphilis IgG	CMV IgM HSV-1 IgM	

Description	Size	Cat No.	Туре
TORCH IgG Control Positive	3 x 1 ml	BXC0799B	LS
TORCH IgM Control Positive	5 x 1 x 1 ml	BXC0799C	LS

Xanthochromia Quality Control

• Lyophilised format for convenience and longer shelf life.

• Human Based Material.

True Third Party control providing unbiased assessment of performance.
Open reconstituted Vial stability of 2 days at 2° to 8°C. Stable for 1 year when stored unopened at 2° to 8°C.

Description	Size	Cat No.	Туре
Xanthochromia Positive Control	3 x 1 ml	BXC0355A	Lyo



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PROTEINS

Immunology Controls

Immunology and protein controls are essential in diagnostics for ensuring the accuracy and reliability of tests that assess various immune responses and protein levels in biological samples. These controls consist of standardised samples with known concentrations of specific proteins or antibodies, utilised to validate the performance of assays such as enzyme-linked immunosorbent assays (ELISA) and Western blots. By incorporating these controls, laboratories can evaluate assay precision, sensitivity, and specificity, thus reducing the risk of incorrect results that could lead to misdiagnoses or inappropriate treatment decisions.

Immunology and protein controls play a key role in maintaining quality assurance in laboratory processes, ensuring consistent and reproducible test results across various assays. Reliable diagnostics are crucial for diagnosing autoimmune disorders, allergies, and infections, as well as monitoring treatment responses. Regular use of these controls helps identify potential issues with reagents or equipment, facilitating accurate and timely patient management. Overall, they significantly enhance healthcare quality by providing dependable diagnostic information that informs clinical decisions and improves patient outcomes.

The majority of the Fortress Immunology/Protein Controls are available in a liquid stable format for convenience and ease-of-use and are supplied with instrument and method specific assigned values.



Lyophilised for long shelf life, stable to expiry at 2-8 °C.

•	Reconstituted stability of /	days at 2-8°C.

Description	Size	Cat No.	Туре
APO A1/B Calibrator Series	6 x 1 x 1 ml	BXC0413A	LS.

ASO (Single Point) Calibrators

Values are assigned by in house methods and instruments.

- 100% human material designed for matrix conformity. The liquid stable ASO Single point calibrator has a shelf life of 2 years when stored at 2-8 °C. Once opened the calibrator is stable for a period of 30 days when stored without contamination at 2-8 °C.

Description	Size	Cat No.	Туре
ASO (Single Point) Calibrator	1 x 1 ml	BXC0323A	LS
ASO (Single Point) Calibrator	3 x 1 ml	BXC0323B	LS

ASO. CRP. RF Control

This control is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.

- The values are assigned in European Reference Laboratories.
- 100% Human serum designed for matrix conformity.
- Once opened the control is stable for a period of 30 days when stored without contamination at 2-8 °C.
- This material is presented in a liquid stable format and is stable up to expiry at 2-8 °C.

Description	Size	Cat No.	Туре
ASO, CRP, RF Control	3 x 1 ml	BXC0645A	LS

CRP Controls & Calibrators

This control is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.

- The values are assigned in European Reference Laboratories.
- 100% Human serum designed for matrix conformity.
- Once opened the control is stable for a period of 30 days when stored without contamination at 2-8 °C.

Description

CRP (Single Point) Calibrator CRP (Multi Point) Calibrator CRP (ULTRA SENSITIVE) Low Control CRP (ULTRA SENSITIVE) Standard Set CRP Control (Level 1) CRP Control (Level 2)

ESR Controls

CRP Control Set

Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C.

Description

ESR Control (Level 1)

ESR Control (Level 2)

IgE Calibrators & Controls

- Lyophilised product stable to expiry at 2-8 °C.
- Reconstituted stability of 7 days at 2-8°C.
- Reconstituted stability of 30 days at -20°C.

Description

IgE Calibrator

IgE Control Set (Level 1 & Level 2)

Protein Control

- This control is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.
- Values are assigned by reference laboratories based in Europe
- 100% Human material designed for matrix conformity.
- Once opened the control is stable for a period of 30 days when stored without contamination at 2-8 °C.

Albumin	
lpha-1 Acidglycoprotein	
Alpha-1 Antitrypsin	
Alpha-2 Macroglobulin	
ASO	
Antithrombin-III	
Beta-2-Microglobulin	

C1q Inactivator C1 Esterase Inhibitor Ceruloplasmin Complement C3 Complement C4 Ferritin Haptoglobin

CRP

Description

Protein Control (Level 1)

Protein Control (Level 2)

Protein Control Set (Level 3)

Protein Control Set (Level 1, Level 2 & Level 3)

Retinol Binding Protein Control

Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C. Open vial stability of 30 days, at 2-8°C.

Retinol Binding Protein Control

Fortress Diagnostics | Quality Controls Catalogue

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Immunology / Proteins | SECTION 13

Size	Cat No.	Туре
3 x 1 ml	BXC0324A	LS
6 x 1 x 1 ml	BXC0324B	LS
3 x 1 ml	BXC0325A	LS
5 x 1 x 0.5 ml	BXC0327A	LS
5 x 1 ml	BXC0326A	LS
5 x 1 ml	BXC0326B	LS
2 x 1 x1 ml	BXC0326C	LS

Size	Cat No.	Туре
2 x 10 ml	BXC0631A	LS
2 x 10 ml	BXC0632A	LS

Size	Cat No.	Туре
1 x 1 ml	BXC0752A	Lyo.
2 x 1 x 1ml	BXC0753A	Lyo.

Values for Immunoglobulin, Inflammatory proteins, Complements, Carrier and Storage proteins are available in the control.

Analytes

3 x 1 x 1ml

IgA IgG IgM IgE Kappa Light Chains Retinol Binding Protein Free Kappa Light Chain Rheumatoid Factor	Prea Free Lamb Trar	Light Chains albumin da Light Chain nsferrin I Protein
Size	Cat No.	Туре
1 x 1 ml	BXC0638A	LS
1 x 1 ml	BXC0639A	LS
1 x 1 ml	BXC0640A	LS

			0
Size	Cat No.	Туре	
1 x 1 ml	BXC0995A	LS	

BXC0641A

LS

SECTION 13 | Immunology / Proteins

 Lyophilised for long shelf life, stable to expiry at 2-8 °C. Reconstituted stability of 7 days at 2-8°C. Description Size Cat No. Type SAA Control 3 x 1 ml BXC0703A Lyo. 	SAA Control				
SAA Control 3 x 1 ml BXC0703A Lyo.	Description	Size	Cat No.	Туре	
	SAA Control	3 x 1 ml	BXC0703A	Lyo.	

RF (Multi-Point) Calibrators

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- This Calibrator is presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.
 The Calibrator is presented as a single level calibrator which can be diluted as necessary for calibrating RF Turbidimetric Assays.
 Values are assigned by in house methods and instruments.
 100% human material designed for matrix conformity.
 Once opened the control is stable for a period of 30 days. when stored without contamination at 2-8 °C.

Description	Size	Cat No.	Туре
RF (Multi-Point) Calibrator	1 x 2ml	BXC0612A	LS
RF (Multi-Point) Calibrator	3 x 2ml	BXC0612B	LS

Specific Protein Calibrators

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- Fortress Specific Protein Calibrator is a single level multi-analyte calibrator for use in automated systems with auto dilution facility.
 Fortress Specific Protein Calibrator set is a multi analyte multi level calibrator set for calibrating protein assays.
 Both the Calibrators are presented in a liquid stable format and is stable up to expiry when stored at 2-8 °C.
 100% Human material designed for matrix conformity.
 Once opened, the calibrator is stable for a period of 30 days when stored without contamination at 2-8 °C.

	Α	nalytes		
Albumin Alpha-1 Acidglycoprotein Alpha-1 Antitrypsin Alpha-2 Macroglobulin ASO Antithrombin-III Beta-2-Microglobulin	CRP C1q Inactivator C1 Esterase Inhibitor Ceruloplasmin Complement C3 Complement C4 Ferritin Haptoglobin	IgA IgG IgM IgE Kappa Light Chains Retinol Binding Protein Free Kappa Light Chain Rheumatoid Factor	F Free La 1	da Light Chains Yrealbumin mbda Light Chain Iransferrin otal Protein
Description		Size	Cat No.	Туре
Specific Protein Calibrator		1 x 1 ml	BXC0644A	LS
Specific Protein Calibrator Set		6 x 1 x 1 ml	BXC0646A	LS



Lipid Controls

Lipid controls are essential in diagnostics for accurately measuring lipid levels, such as cholesterol and triglycerides, in biological samples. These controls consist of standardized samples with known lipid concentrations, which validate the performance of assays used in lipid testing.

By incorporating lipid controls, laboratories ensure the reliability and precision of test results, which are crucial for diagnosing and managing conditions like hyperlipidemia and cardiovascular diseases. Regular use of these controls helps identify potential issues with reagents or instruments, ultimately supporting effective patient management and enhancing the overall quality of healthcare through accurate lipid profiling. The Fortress Lipid Controls are lyophilised for enhanced stability and are suitable for automated and semi-automated systems. The multi-analyte controls offer test menu consolidation. The controls are supplied with instrument and method specific assigned values.



Glycerol Control

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.

• Open vial stability of 30 days, at 2-8 C.				
Description	Size	Cat No.	Туре	
Glycerol Control	2 x 5 ml	BXC0278A	LS.	

HDL/LDL Calibrators

• Fortress HDL/LDL Cholesterol Calibrator is presented in a Lyophilised format for enhanced stability, stable up to expiry at 2-8 °C.

• Once opened and reconstituted the control is stable for 7 days at 2-8 °C, or 30 days at -20 °C.

• 100% human material designed for matrix conformity.

Analytes				
HDL Cholesterol (Direct) LDL Cholesterol (Direct)				
Description				
Description		Size	Cat No. T	Гуре
HDL/LDL Calibrator		Size 3 x 1 ml		Гуре _yo.

Lipid Controls

The Fortress Lipid Controls are intended for monitoring accuracy and precision.

	Analytes		
Apolipoprotein A1 Apolipoprotein B Cholesterol	HDL Cholesterol (Precipitation) HDL Cholesterol (Direct) LDL Cholesterol (Direct)	LDL Cholesterol (I & Triglyce	
Description	Size	Cat No.	Туре
Lipid Control Level 1 (Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL)	3 x 1 ml	BXC0317A	Lyo.
Lipid Control Level 2 (Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL)	3 x 1 ml	BXC0330A	Lyo.
Lipid Control Level 3 (Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL)	3 x 1 ml	BXC0316A	Lyo.
Lipid Control Set (TRI-LEVELS) (Cholesterol, Triglycerides, APO A1, APO B, HDL, LDL)	3 x 1 x 1 ml	BXC0350A	Lyo.



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Maternal Screening Controls

Maternal screening controls are essential in diagnostics for accurately assessing the health of pregnant individuals and identifying potential risks to the developing fetus. These controls consist of standardised samples with known concentrations of biomarkers, such as hormones and proteins, used to validate the performance of screening assays. By incorporating these controls, laboratories ensure the reliability and accuracy of test results, which are crucial for detecting conditions like gestational diabetes, preeclampsia, and chromosomal abnormalities, thereby informing timely clinical interventions.

The role of maternal screening controls also extends to maintaining quality assurance in laboratory processes, ensuring consistent and reproducible results across different tests. Reliable maternal diagnostics are critical for guiding prenatal care and enabling informed decision-making for expectant parents. Regular use of these controls helps identify any potential issues with reagents or equipment, ultimately supporting effective patient management and improving health outcomes for both mothers and their babies. The Fortress Maternal Screening Controls are available in a lyophilised format offering enhanced stability and are 100% human serum based to negate matrix effects.



Anti-Mullerian Hormone (AMH) Calibrators & Controls

- Lyophilised for long shelf life, stable to expiry at 2-8 °C.
- Reconstituted stability of 7 days at 2-8°C.
- Compatable with all AMH assays.

Description	Size	Cat No.	Туре
Anti-Mullerian Hormone (AMH) Calibrators & Control Set	Calibrator Set: 2 x 1x0.5ml ; Control Set: 2x1x0.5ml	BXC0999A	Lyo.

Maternal Screening Control

For screening of Down's syndrome & Spina Bifida compatible for use on most clinical chemistry systems including immunoassay systems.

• Screening Controls are multi analyte controls intended for use in monitoring accuracy and precision for assays used in first and second trimester screening of Downs Syndrome and Spina Bifida.

- The Controls are provided in a Lyophilised format stable up to expiry at 2-8 $^\circ \! C.$
- Reconstituted stability of 7 days at 2-8 °C.
- 100% human serum.

Analvtes

	-		
AFP Free beta hCG	Inhibin A PAPP-A	Total hCC Unconjugated	-
Description	Size	Cat No.	Туре
Maternal Screening Control (Level 1)	2 x 1 ml	BXC0695A	Lyo.
Maternal Screening Control (Level 2)	2 x 1 ml	BXC0695B	Lyo.
Maternal Screening Control (Level 3)	2 x 1 ml	BXC0695C	Lyo.
Maternal Screening Control Tri-Level	3 x 1 x 1 ml	BXC0699A	Lyo.

PAPP-A & f-beta-hCG Controls

• The Controls are provided in a Lyophilised format stable up to expiry at 2-8 °C.

100% human serum.
Reconstituted stability of 7 days at 2-8 °C.

Description	Size	Cat No.	Туре
PAPP-A & f-beta-hCG Control (Level 1)	1 x 0.5 ml	BXC0337A	Lyo.
PAPP-A & f-beta-hCG Control (Level 2)	1 x 0.5 ml	BXC0338A	Lyo.
PAPP-A & f-beta-hCG Control (Level 3)	1 x 0.5 ml	BXC0339A	Lyo.
PAPP-A Control	1 x 0.5 ml	BXC0240A	Lyo.

Peadiatric Control

• L		
• R		
• C		
Description		
Paediatric Control		

Pre-eclampsia

Pre-eclampsia is a disorder of pregnancy that is associative weeks' gestation and frequently near term. During the and Placental Growth Factor (PLGF) are both predictive increased levels of sFlt-1 and decreased levels of PlGF eclampsia..

Determining serum PIGF concentration, used as a single management and decision making (risk stratification) with

PlGF Testing will be valuable:

- For testing women with signs and symptoms of pre-eclampsia aft
- To help Rule-out and Rule-in suspected pre-eclampsia
 - To monitor women that are at high risk for pre-eclampsia
 - To confirm clinical suspection of pre-eclampsia with symptomatic
 Serum concentrations of sFlt-1 and PIGF can differentiate healthy PIGF levels reflect the severity of the disease.

PIGF as a single marker or sFIt-1/PIGF ratio. PIGF alone compared to sFIt-1, and both options are equally recommended for clinical use. In addition to more simpler and affordable alternative to dual biomarker testing. Using F (decreased serum PIGF level) for identifying pregnancies with placental in

PLGF Controls

For screening of Down's syndrome & Spina Bifida compatible for use of

Description	Details
Placental Growth Factor (PLGF) Control Lev	vel-1 Target: 30pg/ml
Placental Growth Factor (PLGF) Control Lev	vel-2 Target: 100pg/m
Placental Growth Factor (PLGF) Control Lev	vel-3 Target: 600pg/m

Pre-eclampsia Control

Description	Details
Pre-eclampsia Control Level-1	s-FLT-1 (soluble fi
Pre-eclampsia Control Level-2	s-FLT-1 (soluble f

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Size			
Size			
	Cat	No.	Туре
3 x 1ml	BXC	0807A	Lyo.
ated with new-onset hype 2nd and 3rd trimesters so te and diagnostic for pre- in maternal serum can pr narker, or sFlt-1 and PlGF, u n women showing signs and	oluble fms eclampsia edict the s	s-like tyros . It has bee subsequer	sine kinase en shown th at onset of p ve the clinica
women and/or to confirm unclear women from women with pre-ecl /PlGF ratio for pre-eclampsia rule- SFlt-1/PlGF ratio, PLGF alone, with PlGF has additional advantages. St	ampsia. Chan in and rule-ou concentratio udies have sh	ges in sFlt-1 a It has a compa n based cut-o own that PlGI	nd arable performa ffs, could provid ⁼ is a good mark
women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St sufficiency including fetal growth	ampsia. Chan in and rule-ou concentratio udies have sh restriction an	ges in sFlt-1 a It has a compa In based cut-o own that PlGi d/or stillbirth	nd arable performa ffs, could provid F is a good mark
women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St sufficiency including fetal growth	ampsia. Chan in and rule-ou concentratio udies have sh restriction an	ges in sFlt-1 a It has a compa In based cut-o own that PlGi d/or stillbirth	nd arable performa ffs, could provid F is a good mark
women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St Isufficiency including fetal growth	ampsia. Chan in and rule-ou concentratio udies have sh restriction an s including in	ges in sFlt-1 a It has a compa In based cut-o own that PlGf Id/or stillbirth	nd arable performa ffs, could provid is a good mark is a good mark systems.
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women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St isufficiency including fetal growth n most clinical chemistry system	ampsia. Chan in and rule-ou concentratio udies have sh restriction an s including in Size 3 x 2 ml	ges in sFlt-1 a at has a compa n based cut-o own that PlGf d/or stillbirth nmunoassay Cat No. BXC0700A	nd arable performa ffs, could provid is a good mark is a good mark systems. Type LS.
women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St isufficiency including fetal growth n most clinical chemistry system	ampsia. Chan in and rule-ou ic concentratio udies have sh restriction an s including in Size 3 x 2 ml 3 x 2 ml	ges in sFlt-1 a it has a compa in based cut-o own that PlG d/or stillbirth nmunoassay Cat No. BXC0700A BXC0700B	nd arable performa ffs, could provid = is a good mark systems. Type LS. LS.
er week 20 of gestation women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFIL-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St isufficiency including fetal growth n most clinical chemistry system I	ampsia. Chan in and rule-ou ic concentratio udies have sh restriction an s including in Size 3 x 2 ml 3 x 2 ml	ges in sFlt-1 a it has a compa in based cut-o own that PlG d/or stillbirth nmunoassay Cat No. BXC0700A BXC0700B	nd arable performa ffs, could provid is a good mark systems. Type LS. LS. LS.
women and/or to confirm unclear women from women with pre-ecl /PIGF ratio for pre-eclampsia rule- SFlt-1/PIGF ratio, PLGF alone, with PIGF has additional advantages. St isufficiency including fetal growth n most clinical chemistry system	ampsia. Chan in and rule-ou in concentratio udies have sh restriction an s including in Size 3 x 2 ml 3 x 2 ml 3 x 2 ml	ges in sFlt-1 a it has a compa in based cut-o own that PlG d/or stillbirth nmunoassay Cat No. BXC0700A BXC0700B BXC0700C	nd arable performa ffs, could provid = is a good mark systems. Type LS. LS. LS.







Neonatal Screening Controls

Neonatal screening controls are essential in diagnostics for ensuring the accuracy and reliability of tests that detect metabolic and genetic disorders in newborns, enabling timely interventions that can prevent serious health issues and improve long-term outcomes.

The Fortress Neonatal Screening Controls are dried blood spot (DBS) which ensures accurate, reliable and repeatable results.

BCAA (Leucine) Calibrators & Controls

Description	Size	Cat No.	Туре
BCAA (Leucine) Calibrator	2 x 1 Disc	BXC0067A	Disc
BCAA (Leucine) (Level 1)	2 x 1 Disc	BXC0068A	Disc
BCAA (Leucine) (Level 2)	2 x 1 Disc	BXC0068B	Disc
BCAA (Leucine) (Level 3)	2 x 1 Disc	BXC0068C	Disc
BCAA (Leucine) (Level 4)	2 x 1 Disc	BXC0068D	Disc
BCAA (Leucine) (Control Set) (4 Levels)	4 x 1 x 2 Disc	BXC0069A	Disc

Dried Bloodspot Calibrators on Filter Paper for Neonatal Screening

Description	Size	Cat No.	Туре
Dried Bloodspot Calibrators on filter paper for Neonatal Screening	4 x 1 x 1 Disc	BXC0059A	Disc
G-6-PDH Controls			0

G-6-PDH Controls

• Values assigned by reference Laboratories in Europe.

- Stable to expiry at 2-8 °C.
 Reconstituted stability of 5 days at 2-8 °C.
 Typical Levels:

G-6-PDH Deficient Control : 100 - 200 U/l.

G-6-PDH Normal Control : 900 - 1400 U/l.

Description	Size	Cat No.	Туре
G-6-PDH Control (Normal)	1 x 1 ml	BXC0573B	Lyo.
G-6-PDH Control (Deficient)	1 x 1 ml	BXC0572B	Lyo.

PKU Calibrators & Controls

4 Levels of Phenylalanine.			
Description	Size	Cat No.	Туре
PKU Calibrator	2 x 1 Disc	BXC0064A	Disc
PKU (Level 1)	2 x 1 Disc	BXC0065A	Disc
PKU (Level 2)	2 x 1 Disc	BXC0065B	Disc
PKU (Level 3)	2 x 1 Disc	BXC0065C	Disc
PKU (Level 4)	2 x 1 Disc	BXC0065D	Disc
PKU (Control Set) (4 Levels)	4 x 1 x 2 Disc	BXC0066A	Disc

Total Galactose Calibrators & Controls

Description	Size	Cat No.	Туре
Total Galactose Calibrator	1 x 1ml	BXC0060A	LS.
Total Galactose (Level 1)	1 x 1ml	BXC0061A	LS.
Total Galactose (Level 2)	1 x 1ml	BXC0061B	LS.
Total Galactose (Level 3)	1 x 1ml	BXC0061C	LS.
Total Galactose (Level 4)	1 x 1ml	BXC0061D	LS.
Total Galactose (Control Set) (4 Levels)	1 x 1ml	BXC0062A	LS.





Paediatric Screening Control

Paediatric screening controls are crucial in ensuring early detection of health conditions and developmental issues in children, which allows for timely interventions that can significantly improve long-term outcomes. By identifying potential problems before symptoms appear, screenings provide an opportunity to prevent complications and enable treatments that can mitigate the impact of various disorders.

Early detection is particularly important for conditions like metabolic disorders, which, if untreated, can lead to severe consequences, but are manageable when caught early. The importance of paediatric screening also lies in its ability to support families in understanding and addressing their child's health needs. Regular screenings provide parents with reassurance and guidance, helping them monitor their child's development and identify any potential issues early on. This proactive approach reduces the risk of delayed diagnosis, which can lead to more complex and costly treatments in the future. By focusing on preventive care and early intervention, paediatric screening controls contribute to better health outcomes, reduced healthcare burdens, and improved quality of life for both children and their families.

The Fortress Paediatric Control is lyophilised for enhanced stability and ensures accuracy and precision. The controls are supplied with instrument and method specific assigned values.



Description	Size	Cat No.	Туре
Paediatric Control	3 x 1 ml	BXC0807A	Lyo.





Speciality Controls

Serum indices are crucial in diagnostics for providing valuable information about the concentrations of various substances in the blood, such as proteins, enzymes, and electrolytes, which are essential for evaluating overall health and diagnosing specific conditions. These indices, which often include measures like albumin, globulin, and total protein levels, help assess organ function, nutritional status, and the presence of inflammatory or pathological processes. By accurately measuring serum indices, healthcare providers can gain insights into a patient's health status, facilitating timely diagnosis and effective treatment planning.

The role of serum indices in diagnostics extends to maintaining quality assurance in laboratory practices, ensuring that test results are consistent and reliable across different assays and patient samples. Regularly utilising controls and standardised methods in serum index testing helps identify potential discrepancies, such as issues with reagents or instruments, thereby supporting accurate clinical decision-making. Overall, serum indices significantly enhance the quality of healthcare by providing essential diagnostic information that guides treatment strategies and improves patient outcomes.

The Fortress Serum Indices Controls are designed to monitor the ability of an instrument to accurately measure lipaemia, icterus and haemolysis (LIH) in specimens. The controls are 100% human serum based to negate matrix effects and are compatible with a variety of clinical chemistry analysers.



Serum Indices (LIH)

- Available in lypholised and liquid stable formats.
- 100% Human based
- Stable to expiry date at 2-8°C.
- Reconstituted stability of 14 days at 2-8 °C.
- 4 levels available.

Analytes				
Lipemia	Lipemia Icterus		Haemolysis	
Description	Details	Cat No.	Size	
Serum Indices (LIHN)	Lyophilised For Beckman Coulter Series	BXC0599F	3 x 4 x 5 ml	
Serum Indices (LIH)	Lyophilised For Beckman Coulter Series	BXC0599E	4 x 5 ml	
Serum Indices (LIH)	Lyophilised For Beckman Coulter Series	BXC0599D	4 x 5 ml	
Serum Indices (LIH)	Lyophilised For Beckman Coulter Series	BXC0599C	4 x 5 ml	
Serum Indices (LIHN)	Lyophilised For Beckman Coulter Series	BXC0599B	4 x 5 ml	
Serum Indices (LIH)	Lyophilised For Beckman Coulter Series	BXC0599A	4 x 1 x 5 ml	
Serum Indices (LIHN)	Liquid Frozen For Beckman Coulter Series	BXC0600F	3 x 4 x 5 ml	
Serum Indices (LIH)	Liquid Frozen For Beckman Coulter Series	BXC0600E	4 x 5 ml	
Serum Indices (LIH)	Liquid Frozen For Beckman Coulter Series	BXC0600D	4 x 5 ml	
Serum Indices (LIH)	Liquid Frozen For Beckman Coulter Series	BXC0600C	4 x 5 ml	
Serum Indices (LIH)	Liquid Frozen For Beckman Coulter Series	BXC0600B	4 x 5 ml	
Serum Indices (LIHN)	Liquid Frozen For Beckman Coulter Series	BXC0600A	4 x 1 x 5 ml	
Serum Indices (LIHN)	Lyopholised Frozen for Roche Series	BXC0604F	3 x 4 x 5 ml	
Serum Indices (LIH)	Lyopholised Frozen for Roche Series	BXC0604E	4 x 5 ml	
Serum Indices (LIH)	Lyopholised Frozen for Roche Series	BXC0604D	4 x 5 ml	
Serum Indices (LIH)	Lyopholised Frozen for Roche Series	BXC0604C	4 x 5 ml	
Serum Indices (LIH)	Lyopholised Frozen for Roche Series	BXC0604B	4 x 5 ml	
Serum Indices (LIHN)	Lyopholised Frozen for Roche Series	BXC0604A	4 x 1 x 5 ml	
Serum Indices (LIHN)	Liquid Frozen For Roche Series	BXC0605F	3 x 4 x 5 ml	

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		Speciality Controls SECTION	
Description	Details	Cat No.	Size
Serum Indices (LIH)	Liquid Frozen For Roche Series	BXC0605E	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Roche Series	BXC0605D	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Roche Series	BXC0605C	4 x 5 ml
Serum Indices (LIHN)	Liquid Frozen For Roche Series	BXC0605B	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Roche Series	BXC0605A	4 x 1 x 5 ml
Serum Indices (LIHN)	Lyophilised Frozen for Abbott Series	BXC0606F	3 x 4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Abbott Series	BXC0606E	4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Abbott Series	BXC0606D	4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Abbott Series	BXC0606C	4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Abbott Series	BXC0606B	4 x 5 ml
Serum Indices (LIHN)	Lyophilised Frozen for Abbott Series	BXC0606A	4 x 1 x 5 ml
Serum Indices (LIHN)	Liquid Frozen For Abbott Series	BXC0607F	3 x 4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Abbott Series	BXC0607E	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Abbott Series	BXC0607D	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Abbott Series	BXC0607C	4 x 5 ml
Serum Indices (LIHN)	Liquid Frozen For Abbott Series	BXC0607B	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Abbott Series	BXC0607A	4 x 1 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Vitros Series	BXC0608F	3 x 4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Vitros Series	BXC0608E	4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Vitros Series	BXC0608D	4 x 5 ml
Serum Indices (LIH)	Lyophilised Frozen for Vitros Series	BXC0608C	4 x 5 ml
Serum Indices (LIHN)	Lyophilised Frozen for Vitros Series	BXC0608B	4 x 5 ml
Serum Indices (LIHN)	Lyophilised Frozen for Vitros Series	BXC0608A	4 x 1 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Vitros Series	BXC0609F	3 x 4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Vitros Series	BXC0609E	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Vitros Series	BXC0609D	4 x 5 ml
Serum Indices (LIHN)	Liquid Frozen For Vitros Series	BXC0609C	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Vitros Series	BXC0609B	4 x 5 ml
Serum Indices (LIH)	Liquid Frozen For Vitros Series	BXC0609A	4 x 1 x 5 ml







Thalassaemia Controls

Thalassemia controls are essential in diagnostics for accurately detecting and managing thalassemia, a group of inherited blood disorders characterised by abnormal hemoglobin production. These controls consist of standardised samples with known hemoglobin patterns and genetic markers, utilised to validate the performance of diagnostic assays such as hemoglobin electrophoresis and molecular testing. By incorporating these controls, laboratories can ensure the reliability and accuracy of test results, facilitating early diagnosis and appropriate management of thalassemia to prevent complications like anemia and organ damage.

Additionally, thalassemia controls play a crucial role in maintaining quality assurance within laboratory testing processes, ensuring that results are consistent and reproducible across different assays and platforms. Reliable diagnostics are critical for guiding treatment decisions, including the need for blood transfusions or iron chelation therapy, and for providing informed family planning options for affected individuals. Regular use of these controls helps identify any potential issues with reagents or instruments, ultimately supporting effective patient management and enhancing health outcomes for those with thalassemia and their families.

The Fortress Thalassaemia Controls are designed to monitor the accuracy and precision of haemoglobin variant measurements associated with thalassaemia.



Thalassaemia Controls

• Designed for monitoring accuracy and precision of Haemoglobin Variant measurements associated with Thalassaemia.

• The controls are supplied in a Lyophilised format stable up to expiry at 2-8 °C.

100% Human blood.

	Analytes		
Heamoglobin F	Haemoglobin A2	Haemoglobin S	
Description	Size	Cat No. Type	
Thalassaemia (Alpha & Beta) Control (Level 1)	2 x 0.5ml	BXC0665A LS.	
Thalassaemia (Alpha & Beta) Control (Level 2)	2 x 0.5ml	BXC0665B LS.	
HbA2 Control Set	2 x 2 x 0.5ml	BXC0778A LS.	





Therapeutic Drug Controls

Therapeutic drug controls are crucial in diagnostics for ensuring the accuracy and reliability of tests that measure drug levels in the bloodstream, which is essential for optimising pharmacotherapy and ensuring patient safety. These controls consist of standardised samples with known concentrations of specific therapeutic drugs, used to validate the performance of assays such as immunoassays and chromatography. By incorporating these controls, laboratories can monitor assay precision and detect any discrepancies, thereby minimising the risk of subtherapeutic or toxic drug levels that could lead to ineffective treatment or adverse reactions.

Furthermore, the role of therapeutic drug controls extends to maintaining quality assurance in laboratory practices, ensuring consistent and reproducible results across different testing environments. Reliable therapeutic drug monitoring is vital for guiding clinical decisions, adjusting medication dosages, and preventing drug interactions. Regular use of these controls helps identify potential issues with reagents or equipment, ultimately supporting effective patient management and improving health outcomes. Overall, therapeutic drug controls significantly enhance the quality of healthcare by providing accurate diagnostic information that informs treatment strategies and optimises patient care.

The Fortress Therapeutic Drug Controls are manufactured from 100% human serum to negate matric effects and are lyophilised for enhanced stability.



Therapeutic Drug Monitoring

• The Fortress Therapeutic Drug Monitoring controls are intended for checking accuracy and precision.

- The controls are presented as multi analyte, lyophilised controls stable up to expiry at 2-8 °C.
- Reconstituted stability of 14 days when stored at 2-8 °C or 60 days at -20 °C.

100% Human serum designed for matrix conformity.

		Anatyces		
Amikacin Caffeine Carbamazepine Cyclosporine	Digoxin Ethosuxamide Gentamicin Lithium	Methotrexate Paracetamol Phenobarbitone Phenytoin	Primidone Salicylate Theophylline Tobramycin	Valproic Acid Vancomycin
Description		Size	Cat No.	Туре
Therapeutic Drug Monitoring) (Level 1)	5 x 5 ml	BXC0781A	LS.
Therapeutic Drug Monitoring	g (Level 2)	5 x 5 ml	BXC0782A	LS.
Therapeutic Drug Monitoring) (Level 3)	5 x 5 ml	BXC0783A	LS.

Analytes





Urine Controls

Urine controls are vital in diagnostics for ensuring the accuracy and reliability of tests that analyse urine samples, which are commonly used to evaluate kidney function, identify metabolic disorders, and screen for drugs or other substances. These controls include standardised samples with known concentrations of various analytes, such as glucose, protein, and specific gravity, which are utilised to validate the performance of assays like dipstick tests and urinalysis. By incorporating these controls, laboratories can monitor assay precision and identify inconsistencies, thereby reducing the risk of incorrect results that could lead to misdiagnosis or inappropriate treatment.

Moreover, urine controls are crucial for maintaining quality assurance in laboratory procedures, ensuring consistent and reproducible results across different assays and platforms. Reliable urine diagnostics are essential for guiding clinical decision-making, particularly in diagnosing conditions such as urinary tract infections, diabetes, and renal diseases. Regular use of these controls helps detect potential issues with reagents or equipment, ultimately promoting effective patient management and improving health outcomes. Overall, urine controls significantly contribute to healthcare guality by providing reliable diagnostic information that informs treatment strategies and enhances patient care.

The Fortress Urine Controls are manufactured from 100% human serum to negate matrix effects and are available in both lyophilised and liquid stable formats offering flexibility to laboratories.



• The controls are presented in a Liquid format and are stable up to expiry at 2-8 °C.

• Open vial stability of 30 days, at 2-8°C.

Description	Size	Cat No.	Туре
Citrate (Urinary) Calibrator [200mg/L]	2 x 1 ml	BXC0136A	LS
Citrate (Urinary) Control (Level 1) [50mg/L]	2 x 1 ml	BXC0137A	LS
Citrate (Urinary) Control (Level 2) [600mg/L]	2 x 1 ml	BXC0139A	LS

Microalbumin Calibrators & Controls

Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.

Open vial stability of 30 days, at 2-8°C.

100% Human urine

Description	Size	Cat No.	Туре
Microalbumin Calibrator Series	5 x 1 ml	BXC0329A	LS
Microalbumin Control (Level 1)	2 x 1 ml	BXC0328A	LS
Microalbumin Control (Level 2)	2 x 1 ml	BXC0328B	LS

Oxolate Calibrators & Controls

• Liquid Stable (Ready-to-use) stable to expiry at 2-8 °C.

•	Open vial stability of 30 days, at 2-8°C.	

Description	Size	Cat No.	Туре
Oxalate (Urinary) Calibrator [0.5 mmol/L]	1 x 5 ml	BXC0147A	LS
Oxalate (Urinary) Control (Level 1) [0.2 mmol/L]	1 x 5 ml	BXC0148A	LS
Oxalate (Urinary) Control (Level 2) [0.7 mmol/L]	1 x 5 ml	BXC0149A	LS

Urine Assayed Controls

- Fortress Urine assayed controls are intended for monitoring precision and accuracy for Urine chemistry assays.
- Values are assigned by reference laboratories in Europe.
- Reconstituted stability of 5 days 2-8 °C or 15 days at -20 °C.
- 100% human urine designed for matrix conformity.

5HIAA Amylase Calcium Chloride Copper Cortisol	Creatinine Dopamine Epinephrine Glucose Magnesium Metanephrine	Microalbumin Norepinephrine Normetanephrine Osmolality Oxalate Inorganic Phosphorous		Potassium Protein Sodium Urea Uric Acid VMA	
scription		Size	Cat No.	Туре	
ne Assayed Control (Level 1)		10 x 10 ml	BXC0661A	Lyo.	
ne Assayed Control (Level 2)		10 x 10 ml	BXC0661B	Lyo.	

Des

Urin

Urin

Urine Strip Controls

For Urine and hCG Strips Only

- Fortress urine Strip controls are designed for quality control of urine strips.
- Values are assigned by in house methods.
- Open vial stability of 30 days when stored without contamination at 2-8 °C.
- Compatible with all urine strips from a variety of manufacturers.
- Reconstituted stability of 5 days 2-8 °C or 15 days at -20 °C.
- 100% human urine designed for matrix conformity.

Albumin Bilirubin Creatinine Blood	Glucose hCG Ketones Leukocytes	Nitrites pH Protein Specific Gravity	U	robilinogen	
Description		Size	Cat No.	Туре	
Jrine Strip Control (Level 1 & Level 2)		2 x 3 x 12 ml	BXC0663A	LS	

De





• The controls are presented as Lyophilised materials for increased stability, stable up to expiry at 2-8 °C.

Ana	lytes



• The Fortress Urine Strip controls are provided in a liquid stable format with a increased stability, stable up to expiry at 2-8 °C.

Analytes





Xanthochromia Quality Controls

Xanthochromia is derived from the Greek word "xanthos," meaning yellow. The term was first used to describe the pink or yellow pigmentation of cerebrospinal fluid (CSF). This colour change is attributed to varying concentrations of pigmented compounds such as oxyhaemoglobin, bilirubin, and methaemoglobin, which are typically the by-products of red blood cell degradation.

The term is now more widely accepted to represent the yellow colour created by the presence of bilirubin in the CSF. The presence of bilirubin, resulting in yellow discoloration of the CSF, is the contemporary definition of xanthochromia. It can be diagnosed by 2 methods: the traditional visualization or eye test, and the more sensitive and specific spectrophotometry. Detection of Xanthochromia has been facilitated since the 1950s by the spectrophotometric analysis of CSF but is usually possible by simple visual inspection of a sample held in front of a white piece of paper. Heme pigments can be found in CSF within 12 hours of subarachnoid bleeding, reach peak levels after 36–48 hours, and typically disappear over the next 7–10 days.

Bilirubin accumulation in CSF may occur via intrathecal conversion from haemoglobin-heme within macrophages and other leptomeningeal cells, or by passive diffusion from the circulation during periods of severe jaundice. In this latter setting, CSF is not usually stained until the total plasma bilirubin reaches 10–15 mg/dl. The presence of bilirubin in CSF is the main cause of xanthochromia associated with high spinal fluid protein content, usually found above levels of 150 mg/dl.

Fortress Diagnostics has been manufacturing Xanthochromia Controls for a number of years intended for use as a true third party control in monitoring the performance of Bilirubin and Oxyhaemoglobin in Cerebrospinal Fluid. Xanthochromia Controls are supplied as unassayed and are suitable for use as IQC with various UV spectrophotometers. It is designed to deliver a cost-effective control to users.



Xanthochromia Quality Control

Lyophilised format for convenience and longer shelf life.

Human Based Material.

True Third Party control providing unbiased assessment of performance.

Open reconstituted Vial stability of 2 days at 2° to 8°C. Stable for 1 year when stored unopened at 2° to 8°C.

	Analytes			
Billirubin	Oxyhaemoglobin			
Description	Size	Cat No. Type		
Xanthochromia Positive Control	3 x 1 ml	BXC0355A Lyo.		



8 🗠



Seraqual 365: IQC Management Software

The Seraqual 365 IQC Management Software is a peer data QC programme designed for use with Seraqual Controls. It provides a cost effective and comprehensive package for labs who need to manage their daily QC activities on a centralized platform in real time.







Simple, Easy to Use Peer Comparison Tool

Real Time Data on a Centralised Platform



Cost Effective & Comprehensive QC Data Management Software

The system has a number of features to enhance data review and trouble shooting within a lab to include:

- A dashboard interface that presents a condensed panel of the features of the system showing options most frequented. It also flags any exceptions for review
- A simple easy to use interface that allow labs to configure and enter results easily with a need for minimal training
- Interactive charts for quick and easy performance review including Histogram and Levey –Jennings
- Exceptions reports so data from a specified date range can be reviewed quickly
- Statistical Comparison Report can be specified by date range at the instrument, method or all method level.
- **Peer Group Statistics** allowing you to compare your results with other labs using the same QC material
- Advanced Statistical Analysis that includes performance indicators such as Uncertainty of Measurement (UM), Bias, Total Error (TE) and Sigma Scores.



Description	Сус
Seraqual 365 Internal Quality Assesment Programme	1 C

Software Overview



Presents a condensed panel of the features of the system showing options most frequented.

A full list of features that the system offers is available from the drop down at the top left hand corner.

It also flags any exceptions for review to allow rapid identification of any QC problems and allow quick and effective corrective action.

QC	Tests				
instrument ID	Cobas				
Location	Chemistry Lab				
Instrument Name	Roche Cobas c311				
Control Type	Clinical Chemistry				
Lots	H5N029 - H5E029				
Show Current Assays		All Assays		Add	l Assay
Current Assays		U All Assays			
Analyte	Method	Reagent Supplier	Unit	Temperature (°C)	Curren
Acid Phosphatase Prostatic	Naphthyl with Pentandiol	Roche Diagnostics	U/L	37	
	DCA	3P Bioservice S.R.L.	µmol/L	0	~
			µmol/L	0	

This option allows a lab to quickly see the instruments registered, the programmes and lots associated with these instruments and then the associated QC test.

Here a lab can also set the Target mean and SD of a test (cumulative, user defined or Seraqual 365 (targets available for selected assayed lots)).

Fortress Diagnostics | Quality Controls Catalogue









The Levey Jennings and Histograms interactive charts can be generated within a specific date range proving invaluable for identifying data shifts and trends.

The charts can be downloaded, saved, annotated and printed for the users convenience.



5

Exceptions Report

The Exceptions Report presents all results that have violated the multi rules set in a date range that can be specified.

Through colour coding it identifies the violated rules that are 'Alert' and those that are 'Reject'.

Any comments associated with a result are also detailed.

Exceptions Report						
Cobas - Chemistry Lab - Roche Cobas c311						
Acid Phos	phatase Pr	ostatic, Naphthyl	with Pentandiol, R	oche Diagnostics,	U/L,37°C	
			HSN	029	HSE0	29
DATE	TIME	OPERATOR	RESULT	SDI	RESULT	SDI
06/06/2023	11:06:38	FD	15.4000 10 .x+	0.5583	36.4000 10:x+	0.5382
Bilirubin D	irect, DCA,	3P Bioservice S.	R.L., µmol/L			
			HSN	029	HSE0	29
DATE	TIME	OPERATOR	RESULT	SDI	RESULT	SDI
07/06/2023	14:20:48	FD	8.2000 1:25	2.0474	25.5000	1.8610

Peer Group Statistics

A lab can select World or Country Statistics for a chosen Control Type and Lot.

The peer group can be at a number of different levels as seen in the image.

The report includes Mean, SD, CV% and the number of labs.

8

Advanced Statistics Report

This report draws on a number of important statistical metrics that include Mean, SD, CV%, Uncertainty of Measurement (UM), Bias, TE, Sigma (BV) and Sigma (CLIA).

The report can be based at either the Instrument, Method or All Method Level and if required within a specified date range.

6

Statistics Comparison Report

A worldwide and country comparison of statistics at either the Instrument, Method or All Method level is available in this report.

A date range can be specified.

This report includes performance indicators such as Mean, SD, CV%, Bias %, SDI and CVI.

	stics Comparison						C	omparis	on Level:	instrume	nt								
Blood Gas	& Electrolytes, Chloride, Enzyma	tic, Roci	he Diagnos	tics, m	noiL														
LOT	INSTRUMENT	NO	OF RESULTS			MEAN			50			CV%		SIAST		501		CVI	
		YOU	COUNTRY	WORLD	YOU	COUNTRY	world	YOU	COUNTRY	WORLD	YOU	COUNTRY	WORLD	COUNTRY	WORLD	COUNTRY	WORLD	COUNTRY	
105-01-005	Roche Cobas c311 - Chemistry Lab 2	109	109	109	14,105	54,554	14,104	0.63	0.63	0.63	4.42	4.42	4.42	NA	NA	NA	NA	NA	
108-02-008	Roche Cobas c311 - Chemistry Lab 2	109	109	109	72.625	72.620	72.620	1.15	1.15	1.15	1.58	1.58	1.58	NA	NA	NA	NA	NA	
108-03-006	Roche Cobas c311 - Chemistry Lab 2	108	108	108	233.267	233,254	233,254	2.41	3.41	3.41	1.45	1.45	1.45	NA	NA	NA	NA	NA	
Clinical Ch	hemistry, Acid Phosphatase Prost	atic, Naj	phthyl with	Pentar	diol, Ro	che Diagn	ostics, U	L, 37°C											
LOT	INSTRUMENT	NO	OF RESULTS			MEAN			50			CV%		BIAST		501		CVI	
		YOU	COUNTRY	WORLD	YOU	COUNTRY	WORL2	YOU	COUNTRY	WORLD	YOU	COUNTRY	WORLD	COUNTRY	WORLD	COUNTRY	WORLD	COUNTRY	
H5E029	Roche Cobas c311 - Chemistry Lab	192	243	243	35,218	35,230	35,230	2.64	2.54	2.64	7.51	7.45	7.48	-0.04	-0.04	0.00	0.00	1.00	
	Roche Cobas c311 - Chemistry Lab 2	142	243	243	35.106	35,230	35,230	2.33	2.64	2.64	6.63	7.48	7.48	-0.35	-0.35	-0.05	-0.05	0.89	
		142	243	243	35,106	35,230	35,230	2.33	2.64	2.64	6.63	7.48	7.48	-0.35	-0.35	-0.05		-0.05	-0.05 0.89

243 243 14.981 15.171 15.171 1.27 1.82 1.82 192

Additional Features

Programme available in English, Spanish, Chinese, Farsi and Arabic.



For further information please contact admin@seraqual365.com

Fortress Diagnostics | Quality Controls Catalogue

IQC & EQA Programmes - Seraqual 365 | SECTION 23

9

Show Statistics For		Pee	r Group				
O World		0	Analyte				
O United Kingdom		Analyte/Method					
Control Type	Control Type Clinical Chemistry		Analyte/Meth	od/Instrumen	t		
	,,		Analyte/Method/Reagent Supplier				
Lot	HSN055 Q		od/Instrumen	ument/Reagent Supplier			
					SD	CV%	No. Lab
Peer Group		SI Unit	Results	Mean	30		
Peer Group		SI Unit	Results 90	Mean 27.3941	1.079	3.94	
							2

QC Data Review

A lab selects a date range (instrument, lot and assay selection is optional) and the report generated displays any results that have not yet been reviewed.

Results violating a multirule are highlighted according to their 'Alert' or 'Reject' status.

A reviewer has the ability to set all the results to reviewed or individually look at results and comment.

This function allow quick and easy identification of tests that may be problematic.



Instruction for Use available on the Software



Automatic deactivations of peer data to maintain the integrity of the data



SEQAS: EQA Management Software

SEQAS is an EQA scheme that forms part of the Fortress Quality Control Platform. EQA helps to assure customers, such as physicians, patients, and health authorities, that the laboratory can produce reliable results. Individual laboratories can use EQA to identify problems in laboratory practices, allowing for appropriate corrective action. EQA participation will help to evaluate reliability of methods, materials, and equipment, and to evaluate and monitor training impact.



For laboratories performing public health-related testing, EQA can help to assure that results from different laboratories during surveillance activities are comparable. EQA participation is usually required for accreditation. Samples received for EQA testing, as well as the information shared by the EQA provider, are useful for conducting educational and professional development activities.

Participation in an external quality assessment program provides valuable data and information for the laboratory which:

- allows peer comparison of performance and results among different test sites;
- provides early warning for systematic problems associated with kits or operations;
- provides objective evidence of testing quality;
- indicates areas that need improvement;
- identifies training needs.



A fast easy-to-use data capture system



Data analysis that is easy to understand & interpret



reports



Certificates of registration

SEQAS Programmes

Description	Cycle	Cat No.	Size
SEQAS External Quality Assessment Programme		SEQAS00P	
SEQAS Monthly Clinical Chemistry (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS001	1 x 12 x 5ml
SEQAS Monthly Immunoassay Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS002	1 x 12 x 1ml
SEQAS Haematology Programme (Apr-Mar)	2 Cycles = 12 Months (1 Sample/Month)	SEQAS003	2 x 6 x 3ml
SEQAS Monthly HbA1c Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS004	1 x 12 x 0.5ml
SEQAS Monthly Coagulation Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS005	1 x 12 x 1ml
SEQAS Monthly Lipids Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS006	1 x 12 x 1ml
SEQAS Monthly Cardiac Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS007	1 x 12 x 2ml
SEQAS Monthly Blood Gas & Electrolytes Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS008	1 x 12 x 2ml
SEQAS Monthly Specific Protein Panel Programme (Jan-Dec)	1 Cycle = 12 Months (1 Sample/Month)	SEQAS009	1 x 12 x 1ml

Fortress Quality Control Inserts Portal

Download e-inserts for the Fortress Diagnostics Quality Controls & Calibrators range through our online portal.

Fortress Diagnostics are excited to launch the new Fortress QC Inserts Portal. The portal allows customers to easily search our database of inserts by product name, catalogue number or batch number in our easy-to-use filter tool. Inserts can be viewed online or downloaded in PDF format.





Monthly cycles for a range of

programmes

insert.fortressdiagnostics.com



Terms & Conditions

1. General

All quotations made, all orders accepted, all goods sold, all service and advice rendered are subject to these conditions and to the exclusion of any previous conditions of purchase. Any variations to these must be expressly accepted by us in writing.

2. Specifications

We reserve the right to supply goods to the specification and/or design current at the date of dispatch. We cannot accept a request for goods as previously supplied as binding.

3. Admin & Handling Charge

An admin & handling charge of £25.00 will be added to all orders.

4. International Bank Charges

For international orders a £15.00 international bank charge will be added to all BACS payments unless the customer agrees to pay these charges at their end.

5. PayPal/Credit Card Transactions

We accept PayPal, credit card and debit card payments. There is a 3.5% PayPal processing fee added to all customer orders who choose to pay via PayPal this will be detailed on your invoice. This is the same charge that Fortress incur to receive this method of payment.

6. Prices

Although we take great care to ensure price stability, we reserve the right to change prices without notice. Goods will be invoiced at the prices valid on the date of confirmation of the Buyer's order. All prices given are exclusive of VAT unless otherwise stated.

Fortress Diagnostics cannot accept responsibility for typographical errors in pricing within our catalogues, advertising literature or guotations.

Prices guoted are "Ex Works" (Incoterms 2010). Freight, insurance and all other respective costs e.g. Certificate of origin, legalised and notarised documents will be invoiced separately.

7. Delivery

Goods are shipped or dispatched at Buyer's risk. Our standard dispatch dates are 21 days after orders are confirmed and paid in full (unless you have credit terms).

Delivery times quoted are to be treated as an estimate only and Fortress Diagnostics shall not be liable for any failure to deliver within such times whether a time for delivery be quoted or not. We shall not be held responsible for delays caused by anything beyond our reasonable control. Any special delivery requirements must be notified to us in writing at the time of placing the order.

We shall have the right at our discretion to make a reasonable charge for special handling and/or delivery.

The Seller is entitled to make delivery in instalments unless such delivery would be unreasonable or unless otherwise agreed upon and confirmed by the Seller in writing. In the case that a fixed period for delivery is agreed upon and the Seller is in default with the supply, the Buyer shall grant the Seller a reasonable additional period of time, normally of 4 weeks.

8. Order Additions

Any additions will result in a new order being created in line with our standard delivery terms. Where possible we will attempt to dispatch your order additions with your original order but cannot guarantee this.

9. Damage/Loss/Shortage

No claim for breakages or missing goods can be recognised unless notified to Fortress Diagnostics and the carriers within 72 hours of receipt of the goods. If damages or shortages are immediately apparent upon delivery please:

1. Notify the carrier when signing for receipt of the goods.

2. Advise Fortress Diagnostics customer services team as soon as possible.

In the case of non-delivery of a complete consignment notification must be received by us within 10 days of invoice date.

10. Return of Goods

Notification of returns must be made within 1 week of receipt of goods. Goods which have been received damaged or in error must be held in the correct storage conditions

Do not return the goods until you contact our customer services team for a returned goods authorisation number. Clearly mark the authorisation number on the package to ensure your goods can be speedily processed.

You will appreciate that we cannot accept responsibility for loss or damage to goods being returned to us unless we arranged collection. If the mistake was ours there is of course no charge to you. However, if the goods were incorrectly ordered an administration and re-stocking charge of 25% of the invoice value excluding VAT is chargeable. Temperature controlled products incorrectly ordered will not normally be accepted for return.

11. VAT

We are required to charge VAT on all UK orders at the current VAT rate.

For organisations that are VAT exempt each order must be accompanied by an exemption certificate unless special arrangements have been made in writing. Under circumstances where a VAT exemption certificate does not accompany an order we are obliged by law to charge VAT. If subsequently you reclaim a VAT credit it must be completed within 30 days of invoice

12. Payment Terms

Payments are to be made within the agreed payment terms set with our customers. We may at our discretion charge interest at the rate of 5% above the base lending rate of Barclays Bank plc on all overdue accounts to the date of actual payment. Payment in advance or financial guarantees may be requested from customers who do not have an account.

13. Ouotations Quotations are valid for 30 days.

14. Cancellation

Orders may not be cancelled without our prior written agreement and may be subject to a cancellation charge. Standing orders must be cancelled in writing at least 8 weeks prior to what becomes the last shipment date.

15. Liability

Fortress Diagnostics' liability for any loss or damage suffered by you and arising by way of defects in the goods or otherwise, howsoever shall be limited to the invoice price of the goods in respect of or in relation to which loss or damage is claimed. Our liability under this clause shall be in lieu of any warranty or condition implied by law as to the quality or fitness for any particular purpose of the goods and save as provided.

16. Ownership

All goods remain the property of Fortress Diagnostics Limited until payment is received in full. The responsibility for insurance and care of goods, including 3rd party risks lies with the purchaser.

17. Health & Safety

All products sold by Fortress Diagnostics Limited, should be handled by qualified personnel, exercising care during handling, usage and disposal. For further information please contact Fortress Diagnostics Limited.

Control of Substances Hazardous to Health 2002 (COSHH)

All Fortress products are developed and designed for 'IN VITRO' use only.

COSHH safety data sheets are available for all products on request.





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