Appendix 1

National Microbiology Framework Agreement Order Form – C216242 Siemens Healthcare Diagnostics Limited

FROM

Authority:	UK Health Security Agency ("the Authority")
Invoice address:	Post: UKHSA Accounts Payable Team Manor Farm Road Porton Down Salisbury SP4 0JG United Kingdom
Contract Manager:	Name: Phone: E-mail:
Secondary Contact: eg. business operational contact, project manager	Name: Phone: E-mail:
Procurement lead	Name: Phone: E-mail:
Name and address for notices:	Name: Address: UK Health Security Agency 10 South Colonnade Canary Wharf London E14 4PU
Internal reference (if applicable):	To be quoted on all correspondence relating to this Order Form: Contract Code: C216242

то

Supplier:	Siemens Healthcare Diagnostics Ltd (the "Supplier")
	Park View Watchmoor Park Camberley Surrey

	United Kingdom GU15 3YL
Contract Manager:	Name: Phone: E-mail:
Secondary Contact:	Name: E-mail:
Account Manager:	Name: Phone: E-mail:
Name and address for notices:	Siemens Healthcare Diagnostics Ltd. Park View Watchmoor Park Camberley Surrey United Kingdom GU15 3YL

Applicable terms and conditions

The following terms and conditions are applicable to the Contract for this Order:

Appendix A	Call-off Terms and Conditions for the Supply of Goods and the Provision of Services	Applicable to this Contract	
Appendix B	Optional Additional Call-off Terms and Conditions for Installation and Commissioning Services	☐ (only applicable if this box is checked)	
Appendix C	Optional Additional Call-off Terms and Conditions for Maintenance Services	☑ (only applicable if this box is checked)	
Appendix D	Optional Additional Call-off Terms and Conditions for Bespoke Research, Development and Manufacturing Requirements	☐ (only applicable if this box is checked and to the extent the applicable terms are included in Annex A (Order Specific Key Provisions))	
Appendix E	Optional Additional Call-off Terms and Conditions for Reagent Rental	☐ (only applicable if this box is checked)	
Appendix F	Optional Additional Call-off Terms and Conditions for Managed Equipment Services	☐ (only applicable if this box is checked)	

Appendix G	Optional Additional Call-off Terms and Condition for Clinical Laboratory Diagnostic Testing Serv		☐ (only applicable if this box is checked and to the extent the applicable terms are included in Annex A (Order Specific Key Provisions))
Appendix H	Further Optional Additional Call-off Terms and Conditions Each of the following clauses in Appendix H is only		(only applicable if one or more boxes are checked)
	applicable to this Contract if the relevant box is che		checkeu)
	1. TUPE applies at the commencement of the provision of Services		
	2. TUPE on exit		
	3. Different levels and/or types of insurance		
	4. Induction training for Services		
	5. Further Authority obligations		
	6. Assignment of Intellectual Property Rights in deliverables, materials and outputs of the Services		
	7. Inclusion of a Change Control Process		
	8. Authority step-in rights		
	9. Guarantee		
	10. Termination for convenience	\square	
	11. Pre-Acquisition Questionnaire		
	12. Time of the essence (Goods)		
	13. Time of the essence (Services)		
	14. Specific time periods for inspection		
	15. Specific time periods for rights and remedies under Clause 3.6 of Schedule 2 of Appendix A		
	16. Right to terminate following a specified number of material breaches		
	17. Expert Determination	\square	
	18. Consigned Goods		
	19. Improving visibility of Sub-contract opportunities available to Small and Medium		

Order Specific Key Provisions set out at Annex Key Provisions) to this Order Form shall also a	A	☑ (only applicable if this box is checked)
23. Modern slavery		
22. Buffer stock requirements		
21. COVID-19 related enhanced business continuity provisions		
20. Management Charges and Information		
Size Enterprises and Voluntary, Community and Social Enterprises		

1. CONTRACT DETAILS

(1.1) Commencement Date:

The date this Order Form is signed by both Parties (the Authority and the Supplier).

(1.2) Services Commencement Date (if applicable):

The date this Order Form is signed by both Parties (the Authority and the Supplier).

(1.3) Contract Price ((i) breakdown and (ii) payment profile):

- 1.3.1. The value of the Services that can be ordered under this Contract shall be twenty-six thousand, four hundred and thirty-six pounds and sixty-one pence only (£26,436.61) (the "Contract Price").
- 1.3.2. The Contract Price is comprised of the components listed in Table 1 Contract Price Breakdown, along with their corresponding prices. However, please note that this price does not cover any additional parts or labour that might be required, as their necessity may only become apparent after the repair work has started. At such time, the Authority will uplift the Purchase Order ("PO").
- 1.3.3. For the avoidance of doubt, the Authority is not committed to pay the full Contract Price.
- 1.3.4. The Contract Price excludes VAT at the applicable rate but is inclusive of all Supplier visits, labour and spare part fees.
- 1.3.5. Only POs placed directly by the Authority are binding under this Contract.
- 1.3.6. POs issued by the Authority in respect of this Contract do not form part of this Contract.
- 1.3.7. Full detail of the Contract Price is contained within Table 1 Contract Price Breakdown:

Table 1 – Contract Price Breakdown

ESTIMATE SERIAL NUMBER - IRL78040709 SYSTEM DESCRIPTION - ASSY CENTAUR XP SYSTEM ENGRPL=10364455					
QTY	PART DESCRIPTION	PART NUMBER		UNIT PRICE	TOTAL PRICE
1	PWR SUPPLY, PDSI UI MODULE	6035289			
1	MODULE USER INTERFACE XP 5.2.3.1	10484109			
1	PCASY BACK PLANE PULL OUT CAGE	2907605			
2	ASSY & SCH SEQUENCE CONTROLLER LLC	11221918			
2	PCB ASSY LLC STEPPER (SERVICE)	95257 <mark>6</mark> 7			
1	RT SPARC BOARD SERVICE ASSEMBLY	6447307			
1	POWER SUPPLY ASSAY, SPARES	10813520			
1	5 VDC POWER SUPPLY ASSY	5530839			

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	Total parts costs
	Total labour costs
	Total travel costs
	Total charges excluding VAT
(1.4) T	Ferm of Contract:
1.4.1	This Contract shall commence on the date this Order Form is signed by both Parties (the
	"Commencement Date").
1.4.2	It shall, unless terminated earlier in accordance with its terms, expire on 31st January 2024,
	or the date the services have been signed off as complete by both Parties, whichever is
	sooner (the " Term ").
1.4.3	The Authority may terminate the Contract for convenience at any time pursuant to Clause
	10 (Termination for convenience) of Appendix H (Further Optional Additional Call-off Terms
	and Conditions) of this Contract provided the Authority gives the Supplier no less than 3
	(three) months written notice.

(1.5) Term extension options:

Not applicable.

2. GOODS AND/OR SERVICES REQUIREMENTS

(2.1) Description of the Goods / Services:

- 2.1.1 The Supplier shall provide servicing and repair to the Authority's instrument stated in Table 1 contained within clause 1.3.6 of this Order Form (the "Instrument") at the Premises and Locations listed in Section 2.2.
- 2.1.2 The Supplier shall provide the services stated in Table 2 Specification of the Services below (the **"Specification"**) for the Instrument listed in Table 1 of this Order Form (the **"Services"**).

Table 2 – Specification of the Services

The Services: Rebuild of the system, then troubleshoot any further issues that arise SERIAL NUMBER - IRL78040709 SYSTEM DESCRIPTION - ASSY CENTAUR XP SYSTEM ENGRPL=10364455				
QTY	PART DESCRIPTION	PART NUMBER		
1	PWR SUPPLY, PDSI UI MODULE	6035289		
1	MODULE USER INTERFACE XP 5.2.3.1	10484109		
1	PCASY BACK PLANE PULL OUT CAGE	2907605		
2	ASSY & SCH SEQUENCE CONTROLLER LLC	11221918		
2	PCB ASSY LLC STEPPER (SERVICE)	9525767		
1	RT SPARC BOARD SERVICE ASSEMBLY	6447307		
1	POWER SUPPLY ASSAY, SPARES	10813520		
1	5 VDC POWER SUPPLY ASSY 5530839			

(2.2) Premises and Location(s) at which the Services are to be provided:

- 2.2.1 The Supplier shall provide the Services to the site detailed in Appendix 2 Premises and Location ("**Premises and Location**").
- 2.2.2 All planned performance of the Services shall be pre-advised by the Supplier to the Authority's delivery contact stated below at least 2 (two) business days prior to the Services being performed on the Instrument:
 - (the "Delivery Contact").
- 2.2.3. The Supplier shall provide the following information when notifying the Delivery Contact:
 - a. Supplier name;
 - b. Authority's purchase order ("PO") number.

2.2.4. The Delivery Contact will confirm:

- a. Booking reference number;
- b. Date and time of Supplier attending the relevant Premises and Location: and
- c. Premises and Location address where the Services shall be performed.
- 2.2.5 The Authority may refuse unscheduled performance of Services. In such event, the Supplier shall rearrange such performance of Services utilising the service delivery process set out in this Clause 2.2.

2.2.6 The Delivery Contact will report any technical issues to:		
Email:		
Phone:		
(2.3) Key personnel of the Supplier to be involved in the Goods / Services:		
Name:		
Phone:		
E-mail:		
(2.4) Performance standards:		
2.4.1 The Supplier shall deliver the Services in accordance with Good Industry Practice.		
2.4.2 Timely delivery of the Services.		
2.4.3 Quality of Services i.e., Services performed in accordance with the Specification as stated		
in section 2.1 & 2.5.		
2.4.4 Proof of the Services having been performed in accordance with Annex A, clause 1.		
(2.5) Quality standards:		
2.5.1 The Supplier shall repair the equipment to the level of the Supplier's manufactured		
specifications, as stated in Appendix 1 of this Order Form, as sold by the Supplier to the		
Authority.		
(2.6) Contract monitoring arrangements:		
Not Applicable.		
(2.7) Management information and meetings:		
Not Applicable.		
3. CONFIDENTIAL INFORMATION (if applicable)		

(3.1) The following information shall be deemed Confidential Information:

- a. Supplier pricing.
- b. Contact details including, but not limited to, email addresses, landline / mobile phone numbers, etc. of Supplier representatives.

c. Contact details including, but not limited to, email addresses, landline / mobile phone numbers, etc. of Authority's representatives.

(3.2) Duration that the information shall be deemed Confidential Information:

For a period of 3 (three) years after the expiry or earlier termination of this Contract unless otherwise agreed in writing by the Parties.

4. DATA PROCESSING (if applicable)

(4.1) Personal Data to be processed by the Supplier:

Not applicable.

5. LEASE / LICENSE (if applicable)

(5.1) The Authority is granting the following lease or licence to the Supplier:

Not applicable.

For and on behalf of the Authority

DocuSigned by:



Date Signed: 15th November 2023

For and on behalf of the Supplier

DocuSigned by:



Date Signed: 14th November 2023

Annex A

Order Specific Key Provisions

1. Acceptance

- 1.1. The Supplier shall perform the Services at the Premises and Location set out in Appendix2 of this Order Form.
- 1.2. The following criteria for the acceptance of the Services performed by the Supplier by the Authority shall apply ("Acceptance"):
 - a. Upon performance of the Services the Supplier shall produce and submit to the Authority a service report for signature by the Authority's authorised representative in accordance with Appendix 3 Service Report (Sample) ("Service Report").
- 1.3. If Services are deemed not to be Accepted by the Authority, the Supplier shall reperform the Services at their own cost.

2. Invoicing Terms

- 2.1. Payment terms are net 30 (thirty) days from receipt of a valid invoice.
- 2.2. Following receipt of the Supplier's countersigned copy of the Contract, the Authority will send a unique PO number. The Supplier must be in receipt of a valid PO number before submitting an invoice.
- 2.3. All invoices presented by the Supplier to the Authority shall be for Services performed by the Supplier and Accepted by the Authority.
- 2.4. All invoices must be sent for approval and shall include the proof of Acceptance to the Authority's designated finance mailbox e-mail: and their agreed and their agreed representative before being submitted for payment.
- 2.5. All invoices must be sent quoting a valid PO number. The Supplier shall provide a current statement of accounts on a monthly basis; this is a standard commercial process and should show all invoices raised and amounts outstanding.
- 2.6. The Supplier shall provide compliant invoices that include, as a minimum, a valid PO number, PO line item number (if applicable), PO line description, and the details (name and

telephone number) of the Authority's authorised representative. Non-compliant invoices will be sent back to the Supplier, which may lead to a delay in a payment.

2.7. In support of the Services provided, the Supplier shall provide to the Authority a Service Report confirming provision of the Services at the Authority's nominated Premises and Location in accordance with clauses 1.2 a) above.

Appendix 1 – Technical Specifications of Advia Centaur XP Immunoassay System



ADVIA Centaur XP Immunoassay System

Engineered for continuous operation and timely, accurate results, the high-performance ADVIA Centaur® XP Immunoassay System is always ready to stay ahead of increasing workflow demands. Its extensive onboard reagent capacities and dedicated STAT capabilities increase productivity, regardless of volume or types of tests.

siemens-healthineers.com



Technical Specifications

Product specifications	
System description	Random-access immunoassay system with direct chemiluminescence testing methodology using advanced acridinium ester technology
Test throughput	Up to 240 tests per hour in batch or random-access mode
Walkaway time	2.5 hours (at maximum throughput)
Sample handling	
Validated sample types	Serum, plasma, urine (varies by assay)
Sample integrity control	Pressure-based level sensing, short-sample detection and flagging, clot detection and flagging, foamy sample detection, ambient temperature detection and flagging
Auto-repeat	User-defined automatic repeat testing from original sample
Sample dilution	Can be auto-diluted, up to 1:1000 (varies by assay)
Auto-reflex testing	User-defined automatic reflex testing
Sample carryover prevention	Disposable pipette tips eliminate sample carryover
Sample volume per test	10–200 μL of sample (varies by assay)
Sample bar code	Code 39; Codabar; Code 128; Interleaved 2 of 5 (any of the symbologies may be active at one time
Sample tubes	1 mL, 2 mL sample cups; 3 mL, 5 mL, 7 mL, 10 mL tubes; microtainer tubes
Sample capacity	180 sample continuous loading in universal 5-position sample racks
Pipette tips	ADVIA Centaur type; 840 onboard, automatic tracking and notification
STAT handling	Dedicated STAT position accepts samples any time
Reaction area	
Reaction area Reaction cuvettes	Onboard capacity of 1000 ADVIA Centaur cuvettes
	Onboard capacity of 1000 ADVIA Centaur cuvettes 18 minutes; results every 15 seconds thereafter, assay-dependent
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Reaction cuvettes	8000 801 80 11 000 and 50000 01 01 00000 and 50 00 00 00 00 00 00 00 00 00 00 00 00
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Reaction cuvettes Assay times Reagent handling	18 minutes; results every 15 seconds thereafter, assay-dependent
Reaction cuvettes Assay times Reagent handling Assays onboard	18 minutes; results every 15 seconds thereafter, assay-dependent
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F)
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F)
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory,
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control Reagent inventory management	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification, of remaining tests, onboard residency, calibration,
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control Reagent inventory management Reagent preparation	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control Reagent inventory management Reagent preparation Bar code-labeled packs	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control Reagent inventory management Reagent preparation Bar code-labeled packs Calibration/QC	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required Yes Bar-coded labels containing lot-specific data; human- and system-readable through handheld bar-code scanner
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent ncillaries Reagent packs Reagent integrity control Reagent inventory management Reagent preparation Bar code-labeled packs Calibration/QC Calibration identification Calibration status	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required Yes Bar-coded labels containing lot-specific data; human- and system-readable through handheld bar-code scanner Tracking and notification of calibration status, including advance notice of pending expiration
Reaction cuvettes Assay times Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent packs Reagent integrity control Reagent inventory management Reagent preparation Bar code-labeled packs Calibration/QC Calibration identification	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required Yes Bar-coded labels containing lot-specific data; human- and system-readable through handheld bar-code scanner Tracking and notification of calibration status, including advance notice of pending expiration, user-defined processes for response of calibration expiration
Reaction cuvettes Assay times Reagent handling Assays onboard Primary ancillaries Reagent ancillaries Reagent packs Reagent integrity control Reagent inventory management Reagent preparation Bar code-labeled packs Calibration/QC Calibration identification Calibration status QC package	18 minutes; results every 15 seconds thereafter, assay-dependent 30 30-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) 25-position cooled storage with refrigeration at 4–8°C (39.2–46.4°F) ReadyPack® cartridge Reagent pack bar-code identification; automatic tracking and notification of inventory, calibration validity, reagent onboard residency, reagent expired/low flags Automatic tracking and notification of remaining tests, onboard residency, calibration, and reagent expired/low flags None required Yes Bar-coded labels containing lot-specific data; human- and system-readable through handheld bar-code scanner Tracking and notification of calibration status, including advance notice of pending expiration user-defined processes for response of calibration expiration Advanced QC package for long-term monitoring, including L–J plots and Westgard rules



Maintenance	
Daily	Automated: 60 minutes; hands-on: 15 minutes
Weekly	Hands-on: 30 minutes
Monthly	Hands-on: 60 minutes
User interface/data management	
Monitor	19-inch diagonal LCD touchscreen
Operating system	MICROSOFT WINDOWS 7
Remote access and service	Smart Remote Services
General specifications	
Power requirements	208–240 V, 50/60 Hz
Water input requirements	Type 1 reagent water guidelines as specified by the Clinical Laboratory and Standards Institute (CLSI); at minimum, water quality should meet CLSI Type II guidelines
Water quality requirements	Bacterial content: >50 ppb total organic compounds Maximum resistivity: 1 megohm/cm
Drain requirements	7.5 L waste bottle (assembly is included)
Dimensions	130.9 (h) x 183.9 (w) x 104.2 (d) cm 51.5 (h) x 72.4 (w) x 41.0 (d) inches (excluding monitor and accessories)
Weight	545 kg (1200 lb) (including monitor and accessories)
Compliance	Complies with international environmental, health, and safety standards, including CE and RoH
Floor load-bearing requirement	390 kg/m² (80 lb/ft²)
Noise emission	Up to 58.8 dB in ready state, 61.3 dB in operating state
Ambient temperature	18–30°C (64–90°F)
Ambient humidity	20–80% noncondensing
Ventilation	10,000 BTU/hour
Overvoltage classification	Main supply voltage fluctuations must be within ±10% of the nominal voltage
Pollution classification	IEC 1010-1 Pollution Degree 2

Siemens Healthineers Headquarters Siemens Healthcare GmbH Henkestr. 127 91052 Erlangen, Germany Phone: +49 9131 84-0 siemens-healthineers.com

Legal Manufacturer

Siemens Healthcare Diagnostics Inc. Laboratory Diagnostics 511 Benedict Avenue Tarrytown, NY 10591-5005 USA Phone: +1 914-631-8000

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Site	Address	Postcode	Delivery Contact	Email and Contact Number
Cambridge Addenbrookes	Cambridge Addenbrookes Hospital UKHSA Laboratory Cambridge	CB2 0QW		

Appendix 2 – Premises and Locations

Appendix 3 – Service Report (Sample)

Healthin	ENS . eers				Service Report		
Siemens Healthineers 3rd Floor Park View Watchmoor Park Camberley GU15 3YL United Kingdom			Account # Account Name	315502 Public Health England Addenbrooke's Hospital, Level 6 Cambridge Microbiology Lab			
			Street City Country	PO Box 236, H CB2 2QW Cam United Kingdon	bridge		
Contact Cust. Inventory # Report # SR Open Date Severity		Dumo Mururi Virology 1-4011562338 17.01.2023 09:42 A-Srvodown(Emergency)	Contact Email Model Configuration Serviced Component Serial # PO #	ADVIA CENTA ASSY CENTAU IRL90120839	UR XP IR XP SYSTEM ENGLISH E		
Problem Descrip Failing the v		Lots of water in trap.					
Activity Type Employee Name		Field Repair	Arrival Date/T End Date/Tim	Arrival Date/Time			
Quantity	Part#	Manufacturing #	Product		18.01.2023 15:00		
1 1 1 6 2.5	10811323 11222226 11223787 2598122 LB TT	10811323 11222226 11223767 078-K049-02 LB TT	PUMP BELLOWS CENTAUR VACUUM PUMP FILTER WATER MALE TO FEMALE DEGASSER SVC KIT CENTAUR Labor Travel time				
52066	25.20	A.	2.100000-01062				
Bellows pur and ran daily	np has also y clean.	failed causing back up and co	acheive the required vacuum. Intamination of the water lines to advised to perform monthly clea	therefore re-tube	d, replaced bellows pump		
The Instrument w	vas handed o	ver in a proper condition.					
Date	FSR	lame:	Customer Signature				
This electronic do	ocument is al	so valid without a signature.					