ORDIN DE PLATA NR.: 79 TIP.DOC. 1

DATA EMITERII:9 noiembrie 2021

PLATITI: 3500-00 LEI: Trei Mii Cinci Sute lei 00 bani

PLATITOR: (R) S.C. "OXIVIT-MED" S.R.L.
CONTUL DE PLATI/CODUL IBAN
MD44ML000000002251729503
CODUL FISCAL :1007600044280 /

PRESTATORUL PLATITOR
BC"Moldindconbank"S.A. fil."Invest" Chisinau
CODUL BANCII:
:MOLDMD2X329:

BENEFICIAR (R)IMSP Spitalul Clinic Republican "Timofei Mosnea
CONTUL DE PLATI/CODUL IBAN
MD32ML000000002251502448
CODUL FISCAL :1003600150783 /

PRESTATORUL BENEFICIAR
BC"Moldindconbank"S.A.
CODUL BANCII:
:MOLDMD2X :

DESTINATIA PLATII:Pentru garantia pentru: oferta la procedura de achizi?ie public: a nr. ocds-b3wdp1-MD-1634709502119 din 10.11.2021

TIPUL TRANSFERULUI NORMAL/URGENT :N:

L.S.

CODUL TRANZACTIEI:001:
DATA PRIMIRII:09/11/2021
DATA EXECUTARII:09/11/2021 0:00:00:

SEMNATURILE EMITENTULUI

CONDUCATOR:Web Kojevnikov Dmitrii

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(semnatura electronica)

CONTABIL-SEF:Web Kojevnikov Dmitrii

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L.S. (semnatura electronica)

CONDUCATOR: _____
(semnatura manuala)

CONTABIL-SEF: _____
(semnatura manuala)

SEMNATURA PRESTATORUL L.S.

MOTIVUL REFUZULUI L.S.

SEMNATURA BANCII :

Zbul2TC8Vk0YEZb5ldvA4zoyKNGsIdPNcwMszZcn80vf/6PAxb6BmxkNV4/oNXk+3
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I+

Digitally signed by Kojevnikov Dmitrii
Date: 2021.11.09 18:02:18 EET
Reason: MoldSign Signature
Location: Moldova

MOBIAS BANCA

Nr. 12101-304

18.03.2016

CERTIFICAT
PRIVIND EXISTENTA CONTURILOR CURENTE

Prin prezentul, **BC „Mobiasbancă – Groupe Societe Generale" S.A.**, codul băncii (BIC): **MOBBMD22**, confirmă că compania **OXIVIT-MED SRL**, cod fiscal (IDNO) **1007600044280,** deține următoarele conturi curente la BC "Mobiasbancă-Groupe Societe Generale" S.A., Filiala. 1 Stejaur :

1. **MDL - 2224710SV23488147100; IBAN- MD09MO2224ASV23488147100**
2. **EUR - 2224710SV22227957100; IBAN- MD17MO2224ASV22227957100**
3. **USD - 2224710SV22214937100; IBAN- MD86MO2224ASV22214937100**

Certificatul este emis în baza cererii întreprinderii: Oxivit-Med SRL.

L.Ș.

Dumitru Popa
Director filială „Stejaur"

Executor : Mariana Guzun
Tel: 022 812 614

Filiala Nr. 1 „Stejaur"
Bd. Ştefan cel Mare şi Sfânt 196
MD-2004, Chişinău, Moldova
Cod MOBBMD22
Cont de corespondenţă 35213892
la Centrul de Decontări al BNM

Tel. +373 22 81 26 15
Fax. +373 22 81 26 15
www.mobiasbanca.md

BC „Mobiasbancă – Groupe Société Générale" SA
Capital Social: 100 000 000 MDL
Număr de înregistrare de stat - 1002600006089
Sediul Central:
bd. Ştefan cel Mare şi Sfânt 81a
MD-2012, Chişinău, Moldova

GROUPE SOCIETE GENERALE

REPUBLICA MOLDOVA

CERTIFICAT DE ÎNREGISTRARE

Societatea Comercială "OXIVIT-MED" S.R.L.

ESTE ÎNREGISTRATĂ LA CAMERA ÎNREGISTRĂRII DE STAT

Numărul de identificare de stat - codul fiscal

1007600044280

Data înregistrării **30.07.2007**

Data eliberării **30.07.2007**

Bordeianu Tatiana, registrator de stat

Funcția, numele, prenumele persoanei care a eliberat certificatul

semnătura

CAMERA ÎNREGISTRĂRII DE STAT A REPUBLICII MOLDOVA L.Ș.

MD 0067985

I.P. "AGENŢIA SERVICII PUBLICE"

Departamentul înregistrare şi licenţiere a unităţilor de drept

EXTRAS

din Registrul de stat al persoanelor juridice

nr. **8871** din **05.05.2021**

Denumirea completă: **Societatea Comercială «OXIVIT-MED» S.R.L.**
Denumirea prescurtată: **S.C. «OXIVIT-MED» S.R.L.**
Forma juridică de organizare: **Societate cu Răspundere Limitată.**
Numărul de identificare de stat şi codul fiscal: **1007600044280.**
Data înregistrării de stat: **30.07.2007.**
Sediul: **MD-2032, bd. Decebal, 82, ap.(of.) 90, mun. Chişinău, Republica Moldova.**
Modul de constituire: **nou creată.**
Obiectul principal de activitate:
1 Importul, fabricarea, comercializarea, asistenţa tehnică şi (sau) reparaţia dispozitivelor medicale şi (sau) a opticii;
2 Comerţul cu ridicata al parfumurilor şi produselor cosmetice;
3 Comerţul cu amănuntul al produselor cosmetice şi de parfumerie, articolelor de toaletă;
4 Intermedieri pentru vînzarea unui asortiment larg de mărfuri;
5 Alte tipuri de comerţ cu amănuntul în magazine nespecializate;
6 Alte tipuri de comerţ cu ridicata;
7 Închirierea altor maşini şi echipamente.
Capitalul social: **5400** lei.
Administrator: KOJEVNIKOV DMITRII, IDNP 0972305012362,
Asociaţi:
1. KOJEVNIKOV DMITRII , IDNP 0972305012362
cota 5400.00 lei, ce constituie 100 %.

Prezentul extras este eliberat în temeiul art. 34 al Legii nr. 220-XVI din 19 octombrie 2007 privind înregistrarea de stat a persoanelor juridice şi a întreprinzătorilor individuali şi confirmă datele din Registrul de stat la data de: 05.05.2021.

Specialist coordonator
tel. 022-207-840
Lazari Aliona

c/f: 1007600044280; adresa: str. Decebal 82-90, or. Chișinău, Republica Moldova
telefon: + 373 22 808002; fax: + 373 22 808003
web: www.oxivit-med.com; e-mail:info@oxivit-med.com

Lista fondatorilor companiei SRL „Oxivit-Med”

Nr.	Numele, Prenumele	Codul Personal
1	Kojevnikov Dmitrii	0972305012362

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

Manufacturer:

Medtronic, Inc.
710 Medtronic Parkway
Minneapolis MN 55432
USA

Product Category(ies):

- **Autotransfusion Systems and Associated Disposables**
- **Centrifugal Blood Pumps**
- **Bio-Console Drive Units**
- **Flow Monitoring Systems**
- **Bio-Cal Blood Temperature Controller**
- **Temperature Monitoring Systems and Associated Disposables**
- **Blood Monitoring Systems**
- **Cardioplegia Delivery Systems**
- **Disposable Blood Handling Devices used for Open Heart Surgery**
- **Arterial Filters**
- **Oxygenators including Heat Exchangers, with and without Cardiotomy Reservoirs**
- **Cardiotomy Venous Reservoirs**
- **Venous Reservoir Bags**
- **Perfusion Equipment and Disposable Perfusion Devices**
- **Disposable Medical Devices for Drainage Systems**
- **Disposable Medical Devices for use in Cardiopulmonary Surgery: Cardioplegia, Cannulae, Venting, Suction**
- **Pressure Display System & related accessories of class IIa**
- **Tissue Positioning/Stabilizing Devices**
- **Surgical Site Clearing Devices**
- **Intravascular Shunts**
- **Surgical Retractors**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 72150396

Valid from: 2020-02-12
Valid until: 2024-05-26

Date, 2020-02-12

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

Facility(ies):

Medtronic Mexico S.de R.L.de CV
Av. Paseo Cucapah, 10510 El Lago, C.P. 22210 Tijuana, Baja California, MEXICO

Medtronic Perfusion Systems
7611 Northland Drive, Minneapolis, MN 55428, USA

Medtronic, Inc.
710 Medtronic Parkway, Minneapolis MN 55432, USA

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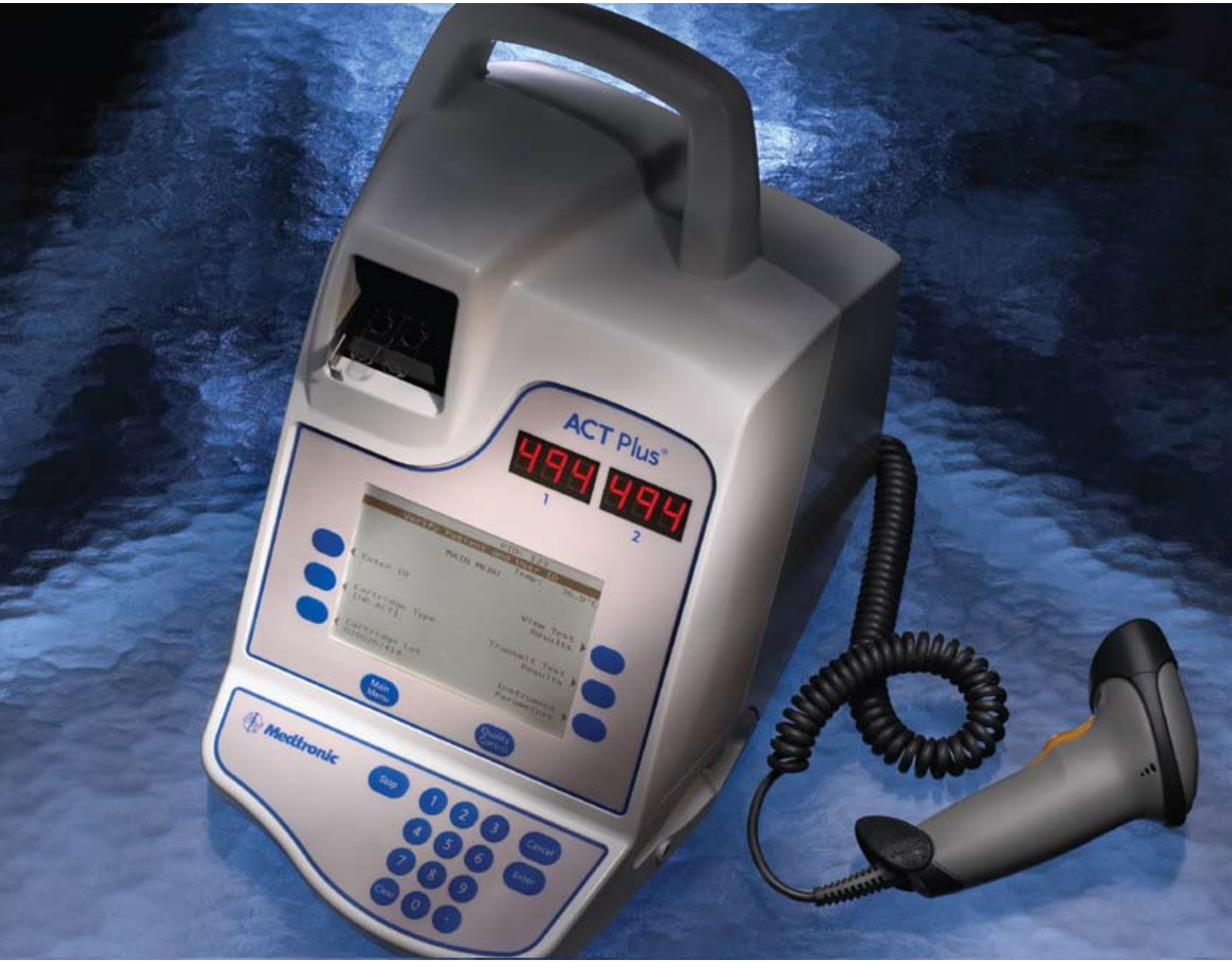
Medtronic

ACT PLUS® SYSTEM

Automated Coagulation Timer



Precise, reliable – the trusted standard

ACT Plus® Sys

The ACT Plus® Automated Coagulation Timer is recognized worldwide as the trusted standard for activated clotting time testing, delivering the accurate, precise and timely results you need.

The ACT Plus® System is used with point-of-care testing during critical procedures, when accurate ACT results are necessary.

- Cardiovascular and vascular surgery
- Cardiac catheterization lab
- Critical care units
- Hemodialysis units
- Pediatric care
- ECMO
- Interventional radiology

Medtronic is the global leader in medical technology — alleviating pain, restoring health, and extending life for millions of people around the world.

As an acknowledged expert in the area of blood management and clot detection technology for more than 20 years, Medtronic leads the way in delivering reliable, cost-effective solutions.

Precise, Reliable Clot Detection Technology

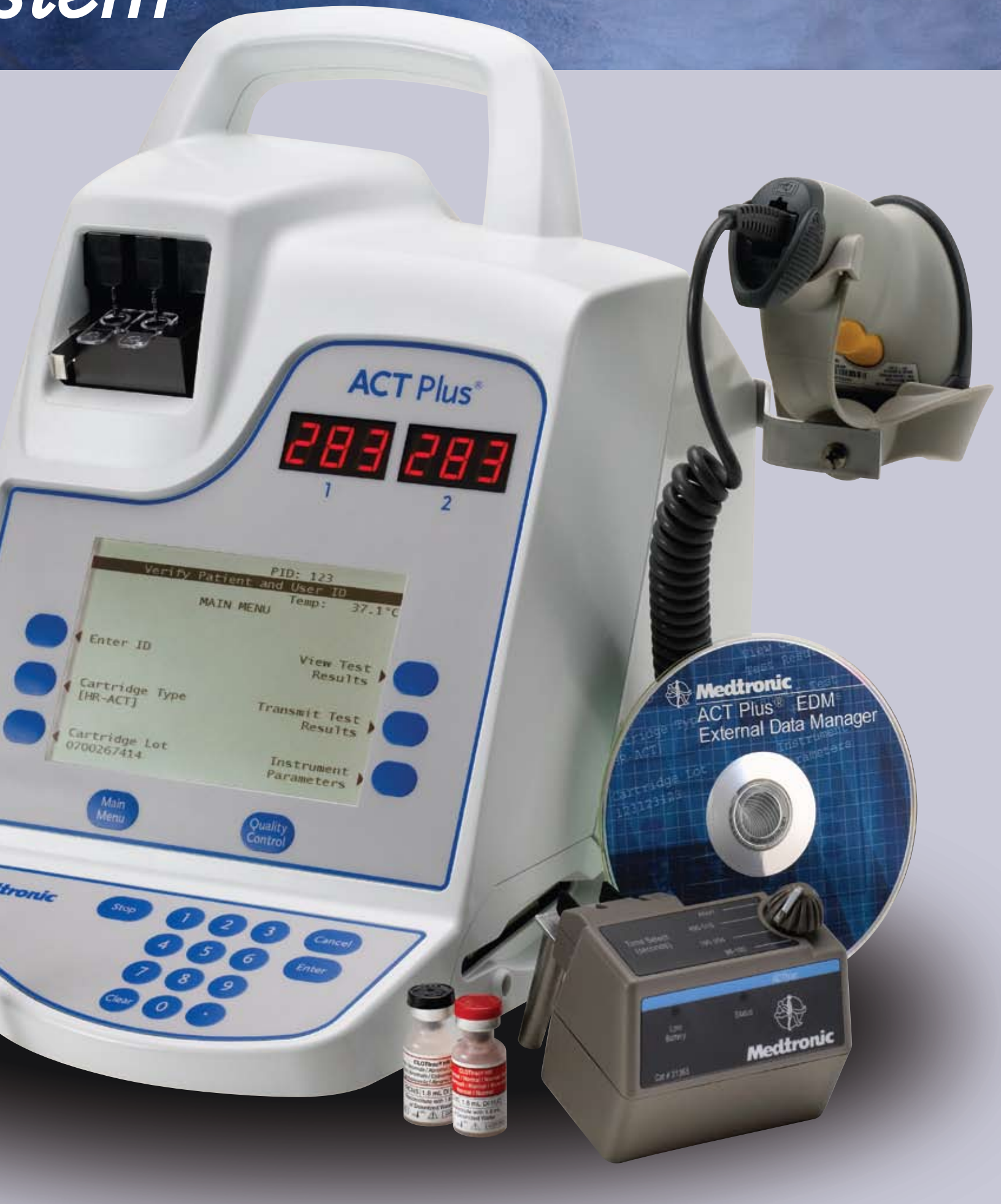
The ACT Plus® System combines the trusted standard in precise clot detection technology with data management and connectivity-ready capabilities. Unlike algorithm-based test measurement, the ACT Plus® System provides real-time clot detection, ensuring accurate and precise clotting time results.

SYSTEM HIGHLIGHTS

- Dual well testing verifies your results are accurate
- Real-time clot detection ensures precise results
- Connects to industry standard point-of-care LIS interfaces, including MAS RALS®-Plus, TELCOR Quick-Linc®, AegisPOC®, and others.*
- Enhanced data management meets all POC testing requirements
  - Stores up to 1000 patient and QC records
  - Holds up to 600 operator IDs
  - Downloads data to a USB or floppy storage device
  - Enables QC and user lockout
  - Bar code scanner available for quick, accurate entry of cartridge and control information, and patient and user IDs

* Check with your local Medtronic representative for country-specific information.

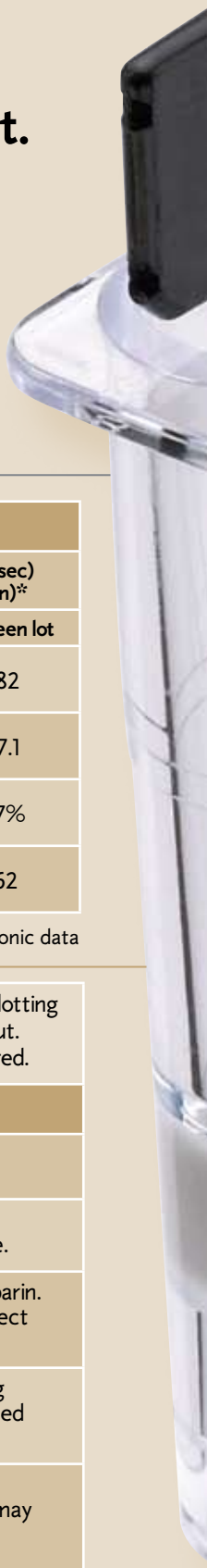
stem


ACT Plus®
283
283
1
2
PID: 123
Verify Patient and User ID
MAIN MENU
Temp: 37.1°C
Enter ID
Cartridge Type
[HR-ACT]
Cartridge Lot
0700267414
View Test Results
Transmit Test Results
Instrument Parameters
Main Menu
Quality Control
Stop
Cancel
Enter
Clear
Medtronic
ACT Plus® EDM
External Data Manager
Medtronic

Cartridges, Controls and Accessor

Our unique dual channel cartridge design allows you to instantly verify your results, eliminating the need to retest.

- **Real-time clot detection** reflects patient's ability to form a clot (rather than mathematically calculated results)
- **Liquid suspended Kaolin activator** for uniform mixing with blood sample
- **Room temperature storage** saves refrigerator space



MULTIPLE TESTING APPLICATIONS

- **High Range Activated Clotting Time (HR-ACT)** for use with fresh whole blood samples in cardiovascular surgery, vascular surgery and percutaneous transluminal coronary angioplasty (PTCA)
- **Low Range Activated Clotting Time (LR-ACT)** for use with fresh whole blood samples in dialysis, ECMO and therapeutic heparin monitoring
- **Recalcified Activated Clotting Time (RACT)** for use with citrated whole blood samples in dialysis and therapeutic heparin monitoring

High Range ACT Precision

	Baseline ACT (sec) (0 units/ml heparin)		Heparinized ACT (sec) (4 units/ml heparin)*	
	Within lot	Between lot	Within lot	Between lot
Mean	130	130	582	582
SD	3.3	6.6	19.0	27.1
%CV	2.6%	5.1%	3.3%	4.7%
N	63	63	162	162

* Internal Medtronic data

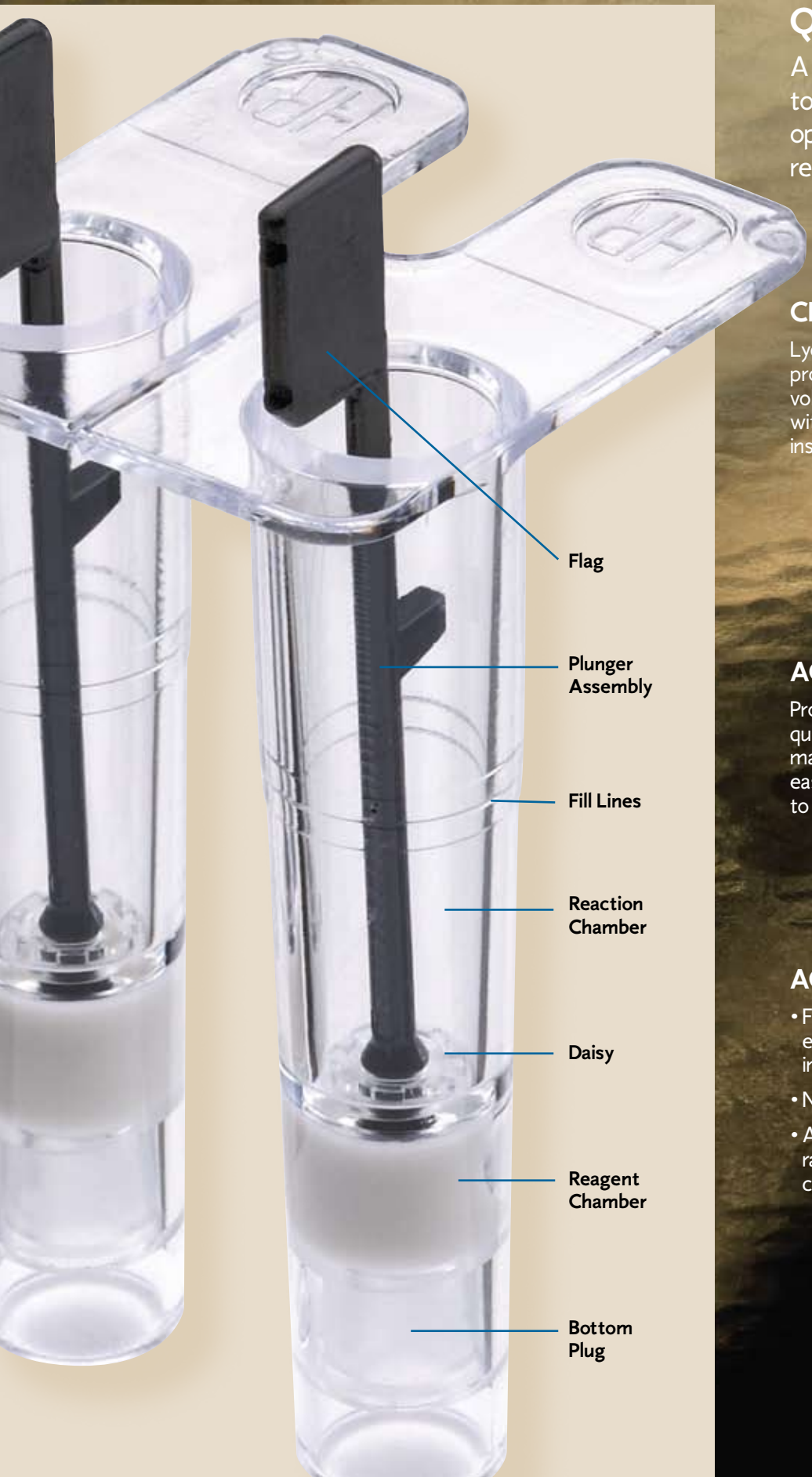
- **Heparinase Test Cartridge (HTC)** for use with fresh whole blood samples to identify the presence of presurgical and postsurgical heparin in various clinical settings, allowing you to:
  - Determine baseline clotting time in the presence of heparin
  - Determine the appropriate time to pull sheath
  - Confirm heparin reversal
  - Identify heparin rebound

The HR Heparinase Cartridge is a two channel activated clotting time test: Channel 1 with heparinase and Channel 2 without. The clotting time results for the two channels are compared.

Example for Interpreting HR Heparinase Results

CHANNEL 1 (with heparin)	CHANNEL 2 (without heparin)	INTERPRETATION
120	120	Patient at baseline. No heparin in sample.
120	260	Sample contains heparin. Extension due to effect of heparin only.
260	260	May be an underlying coagulopathy unrelated to heparin.
260	360	Heparin in the blood sample and patient may have an underlying coagulopathy.

ries



QUALITY CONTROL

A full range of controls are available to verify instrument, cartridge and operator performance and meet regulatory guidelines for testing.

CLOTtrac® Coagulation Controls

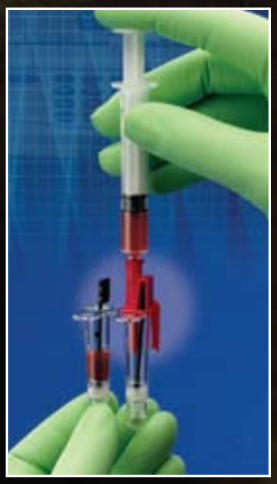
Lyophilized controls provide sufficient volume for testing with multiple instruments

ACTtrac® Electronic Control

Provides a multi-level quality check to make quality control easier and faster to perform

ACT® Easy Fill Accessory

- Fill lines are easily visible, even in low light settings
- Needleless system
- Available for high range and low range cartridges



ACT Plus® EDM External Data Man

Manage your ACT Plus® System data efficiently and effectively.

Get instant access to the charts, reports and spreadsheets you need when you combine the ACT Plus® EDM External Data Manager with the ACT Plus® System. This versatile Microsoft Access-based software solution helps you to analyze test results and make informed decisions more effectively — allowing you to stay focused on improving patient outcomes.

- Manages patient, QC, instrument and user ID data efficiently
- Provides powerful PC-based software to manage data, without an LIS interface
- Delivers 13 preformatted patient and quality control reports

FEATURES AND BENEFITS

Easy-to-Use Interface

- Intuitive features allow users to be up and running with minimal training
- Prompts lead EDM users through the data import function, eliminating all guesswork

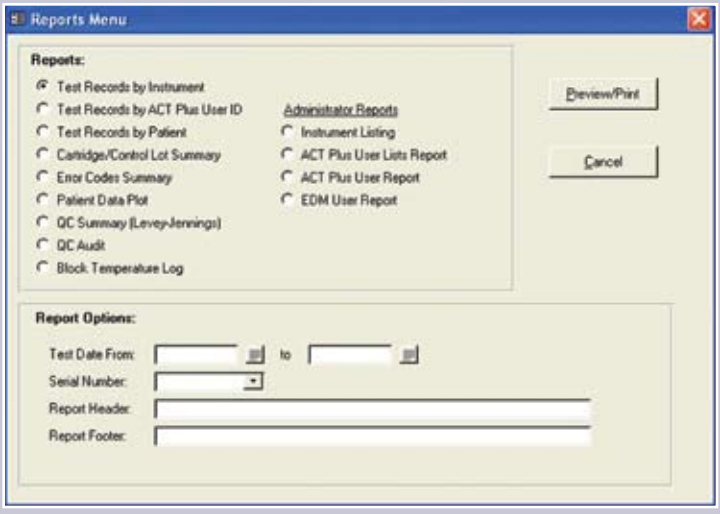
Time-Saving Reports

- Preformatted reports increase efficiency
- Provides visual assessment of QC data and ability to trend QC results
- Ability to track all results for a specific patient

Custom Reporting

- Flexibility in report design and data management
- Data can be exported to Microsoft Excel for graphical and statistical analysis

Reports Menu



Users can select from a list of all preformatted reports to view the desired information and associated options

ager

Simplified Administrative Management

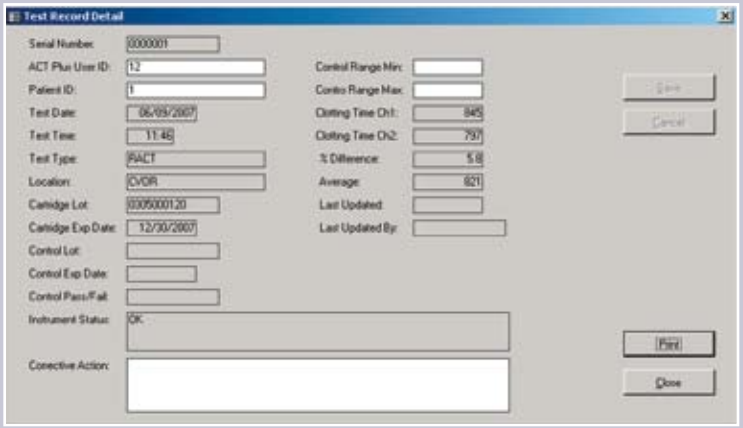
- View the location of all ACT Plus® Systems at your site
- Group ACT Plus® System users into lists to upload into the ACT Plus® System
- Manage your list of all ACT Plus® System users and their certification

Secure System

- Password protection ensures that only authorized EDM users access the information
- Can be backed up onto your network server to ensure safety of and access to information anytime

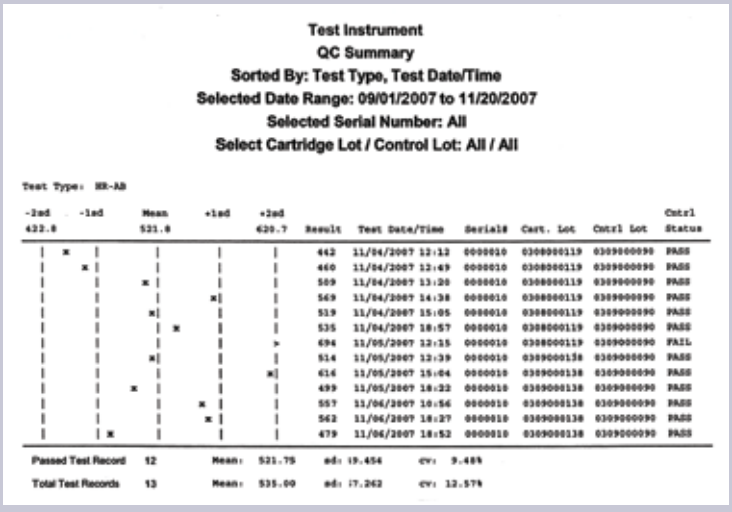
The ACT Plus® EDM External Data Manager is not available in all countries.

Display Detail Option



Users can view a complete listing for a single record and note corrective action taken

QC Summary (Levey-Jennings)



Test Instrument
QC Summary
Sorted By: Test Type, Test Date/Time
Selected Date Range: 09/01/2007 to 11/20/2007
Selected Serial Number: All
Select Cartridge Lot / Control Lot: All / All

Test Type: HR-AB

-2sd 422.8	-1sd	Mean 521.4	+1sd	+2sd 620.7	Result	Test Date/Time	Serial#	Cart. Lot	Cntrl Lot	Cntrl Status
| x |		|	|	|	442	11/04/2007 12:12	0000010	0308000119	0309000090	PASS
| x |		|	|	|	460	11/04/2007 12:49	0000010	0308000119	0309000090	PASS
| | x |		|	|	|	509	11/04/2007 13:20	0000010	0308000119	0309000090	PASS
| | |		x|	|	|	569	11/04/2007 14:38	0000010	0308000119	0309000090	PASS
| | x|		|	|	|	519	11/04/2007 15:05	0000010	0308000119	0309000090	PASS
| | | x		|	|	|	535	11/04/2007 18:57	0000010	0308000119	0309000090	PASS
| | |		|	>	694	11/05/2007 12:15	0000010	0308000119	0309000090	FAIL
| | x|		|	|	|	514	11/05/2007 12:39	0000010	0309000138	0309000090	PASS
| | |		|	|	x|	616	11/05/2007 15:04	0000010	0309000138	0309000090	PASS
| | x |		|	|	|	499	11/05/2007 18:22	0000010	0309000138	0309000090	PASS
| | |	x |	|	|	|	557	11/06/2007 10:56	0000010	0309000138	0309000090	PASS
| | |	x |	|	|	|	562	11/06/2007 18:27	0000010	0309000138	0309000090	PASS
| | x |		|	|	|	479	11/06/2007 18:52	0000010	0309000138	0309000090	PASS

Passed Test Record	12	Mean: 521.75	sd: 19.454	cv: 3.48%	
Total Test Records	13	Mean: 535.00	sd: 37.262	cv: 12.57%	

ORDERING INFORMATION

ACT PLUS® SYSTEM COMPONENTS

Catalog Number	Description
ACT100	ACT Plus® Instrument
ACT200	ACT Plus® Instrument Outside US Version
ACTSC	ACT Plus® Bar Code Scanner
31363	ACTtrac® Electronic Control for ACT Plus®
ACTEDM	ACT Plus® External Data Manager Software

CARTRIDGES AND CONTROLS

Catalog Number	Description	Packaging
ACT® DISPOSABLE TEST CARTRIDGES		
402-03	High Range Activated Clotting Time (HR ACT™)	50 cartridges per box
402-01	Low Range Activated Clotting Time (LR ACT™)	50 cartridges per box
402-07	Heparinase Test Cartridge (HTC)	20 per box
402-02	Recalcified Activated Clotting Time (RACT™)	50 cartridges per box
ACT® CLOTtrac® COAGULATION CONTROLS		
550-07	CLOTtrac® HR Coagulation Control	15 vials of control and deionized water per box
550-08	CLOTtrac® HR Abnormal Coagulation Control	15 vials of control and deionized water per box
550-13	CLOTtrac® HR Control Pack	1 box of HR 550-07 and 1 box of 550-08
550-01	CLOTtrac® CWB Control (normal) – for Low RangeACT® and RACT cartridges	15 vials of control and deionized water per box
550-09	CLOTtrac® LR Abnormal Coagulation Control	15 vials of control and deionized water per box
550-10	CLOTtrac® RACT Abnormal Coagulation Control	15 vials of control and deionized water per box
550-11	Calcium Chloride – for use with the Low Range ACT controls	1 vial per box
550-12	CLOTtrac® HTC Coagulation Control	15 vials of control and deionized water per box
ACT® ACCESSORIES		
313-11	Temperature Verification Cartridge for ACT Plus®	1 per box
313-50	HEPline Kit	Each
303-58	Actuator Cleaning Kit	Each
ACTFILHR	High Range ACT® Easy Fill Accessory	100 per box
ACTFILLR	Low Range ACT® Easy Fill Accessory	100 per box

ACT PLUS® SYSTEM SPECIFICATIONS

DIMENSIONS
Height: 11.9 in (30.2 cm)
Width: 9.0 in (22.9 cm)
Depth: 13.6 in (34.5 cm)
Weight: 11.5 lb (5.22 kg)

DATA PORTS
- Serial Data Port: 19200 baud, 8 data bits, 1 stop bit, no parity
- Bar Code Scanner Port: used with the optional ACT Plus® Bar Code Scanner
- Floppy Drive: PC compatible, 1-44 MB, 3.5 inch floppy disk
- USB: used with Medtronic approved USB storage device

POWER
Voltage: 100-240 V ~ Single Phase
Frequency: 50-60 Hz
Maximum current: 1.0 A (100-240)

OPERATING LIMITS
Temperature: 14°C to 32°C (57°F to 90°F)
Humidity: 10% to 90%, noncondensing

STORAGE LIMITS
Temperature: 0°C to 49°C (32°F to 120°F)
Humidity: 5% to 90%, noncondensing

World Headquarters
Medtronic, Inc.
710 Medtronic Parkway
Minneapolis, MN 55432-5604
USA
Tel: (763) 514-4000
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