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Urgent Field Safety Notice - Urgent Medical Device Recall ...

AMS 700™ With MS Pump™ Dear «Users_Name», Boston Scientific Is Voluntarily Implementing A Product Removal Of Unused Inventory Of The AMS 700 MS (Momentary Squeeze) Pump Following An Increase In Co 2th, 2024

URGENT Field Safety Notice: RA2020- 2329946 URGENT …

With The LIFEPAK 1000 Defibrillator, LIFEPAK 500 Defibrillator, And LIFEPAK CR Plus/EXPRESS Defibrillator. Description Of Issue Stryker Has Become Aware That Certain Packages Of Infant Child Reduced Energy Electrodes Produced By Cardinal Health, Inc. May Have Compromise 2th, 2024

URGENT Field Safety Notice: RA2020 2329946 URGENT …

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On Demand Urgent Care Rebrands As Vybe Urgent Care, Opens ...

With Newly Constructed Centers At 1420 Chestnut Street In Center City And 1217 South Broad Street In South Philadelphia, With Plans To Bring More Than 100 New Jobs To The Region. Mayor Jim Kenney Will Join Vybe For An Official Ribbon Cutting At The South Philadelphia Location On Thursday, January 26. 2th, 2024

URGENT MEDICAL DEVICE FIELD SAFETY NOTICE REMOVAL …

195245 Vanguard XP Tibial Tray 63mm 195758 XP-XP Tibial Tray - Interlok 79mm ... Zimmer Biomet Is Conducting A Medical Device Field Safety Corrective Action (Removal) For The Vanguard XP ... Burrs Are Not Present On All Trays And The Surgical Technique 1th, 2024

URGENT – Field Safety Notice (FSN) - NHSGGC

The Sara Plus. ArjoHuntleigh Would Like To Remind Users That The Entire Knee Pad And Support Bracket Assembly Must Be Removed When Required To Do So For Exercise. An Additional Warning Statement Shown Below Will Be Added To The Instruction For Use Manual And Included With Future Purchases Of These Units. 2th, 2024

URGENT FIELD SAFETY NOTICE: RA2012-035C

Date URGENT FIELD SAFETY NOTICE: RA2012-035C Dear Customer Description: Neptune Waste Management System. Catalog # 0702-001-000, 0702-002-000, 0700-001-000, 0700-002-000, 0700-003-000, 0700-007-000 Lot # All Devices/serial Numbers In May 2012 Stryker Instruments Initiated A Product Field Action To Inform Users That The IFU For The Neptune 2 Devices Had Been 3th, 2024

URGENT Field Safety Notice: RA2020- 2310673
Organization And Would Like To Assist You In Replacing Your LIFEPAK 500 Device With The FDA-approved LIFEPAK CR2 AED, LIFEPAK 1000 Defibrillator Or The HeartSine Defibrillators Based On Your Needs. Contact Your Local Stryker Sales Representative Or Authorized Distributor To D 1th, 2024

Urgent Field Safety Notice - IGJ.nl
The LIFEPAK 1000 Defibrillator Operating Instructions Instruct Users To Inspect The Battery Well And Battery Contacts Routinely As Part Of The Maintenance And Testing Schedule. You Must Immediately Follo 3th, 2024

AMENDED - GE Healthcare URGENT FIELD SAFETY NOTICE
GE Healthcare 3000 N. Grandview Blvd. W440 GEHC Ref# 32070-2 To: Director Of Biomedical Engineering Director Of Neonatology/ L And D/ Nurse Manager Risk Manager/Hospital Administrator RE: Giraffe Incubator, Giraffe OmniBed, Giraffe Incubator Carestation And 3th, 2024

GE Healthcare URGENT FIELD SAFETY NOTICE Waukesha, WI ...
You Can Continue To Operate Your Device, Consistent With Good Clinical Practices. Until GE Healthcare Performs The Necessary Testing, Users Should: 1) Disconnect Devices That Use The USB (Giraffe OmniBed Carestation And Giraffe Incubator Carestation Only) 1th, 2024

URGENT FIELD SAFETY NOTICE GE Healthcare 3000 N. …
Centricity PACS-IW With Universal Viewer Software Versions 5.0.x With PACS-IW Foundation. These Issues Do Not Impact Customers Using Centricity Universal Viewer With A Centricity PACS Foundation Or Centricity Universal Viewer Cardiology. Intended Use Centricity Univer 1th, 2024

GE Healthcare URGENT FIELD SAFETY NOTICE GE Healthcare …
Centricity PACS RA1000 Workstation. Centricity PACS Versions 3.2 And Forward Are Affected. Customers Using GE Centricity RIS-I Are Not Impacted. Safety Issue #2 Interrupted Workflow Could Break The Synchronization Between Agfa Talk And Centricity PACS RA1000 Workstation. Centricity 1th, 2024

Urgent Field Safety Notice Device Commercial Name
1. 1. Device Type(s)* Brief Description Of The Device(s) In Plain Language, Including Whether Supplied Sterile. Consider Including A Photo (here Or In An Annex) Where This Would Help With Identification 1. Add As Appendix If Necessary. 2. Commercial Name(s) 1. 3. Unique Device Identifier(s) (UDI-DI) Com 3th, 2024

URGENT: FIELD SAFETY NOTICE (Removal)
Required For The Production Of Gowns And Lab Coats. As A Result, Gowns And Lab Coats Produced Rips, Holes, Tears, Incomplete And/or Open Seams, Staining, Foreign Material Or Debris (see Attachm 2th, 2024

URGENT FIELD SAFETY NOTICE - BfArM
The CARESCAPE Patient Data Module (PDM) Is Used With The Following Monitors: CARESCAPE B450/B650/B850, SOLAR 8000M/I And Transport Pro. If Pace Detection Is Turned ON, On The Monitor, And If An Automated External Defibrillator (AED) Is Used To Perform Defibrillation, The Lo 1th, 2024

Urgent Field Safety Notice
The Purpose Of This Notification Is To Inform You That Alere San Diego, Inc. Is Initiating A Voluntary Removal Of The AlereINRatio ® /INRatio ® 2 PT/INR MonitoringSystem From The Market. This Removal Includes Both The Alere INRatio ® /INRa 3th, 2024

Urgent Field Safety Notice SBN-CPS-2019-010
Coagulation Analyzer In Case A Country Will Commence Installations Of The Cobas T 511 And Cobas T 711 Coagulation Analyzer. Corrective Action: Long Term, Adaption Of The Design To Align The Tolerances Between Cuvette And Incubator B 1th, 2024

Urgent Field Safety Notice MY-FSN-RDS-CoreLab-2021-006
2. Wait Until Analyzer Is In “idle” Status, Then Initialize The U 701. The U 701 Only Initialization Can Be Started From Monitoring -> Analyzer -> U 701 Page. * 3. Wait Until Analyzer Is Again In “idle” Status, Then Continue The Measurement Process By Clicking “start” 1th, 2024

Urgent Field Safety Notice FSN-CPS-2019-010
And The Cuvette. This Was Introduced To The Series Production From SN 5000 (cobas T 511 Coagulation Analyzer) And SN 1500 (cobas T 711 Coagulation Analyzer). Instruments Which Were Already Distributed / Installed At Customers Site: Roche Started The Implementation Process Of The Modificat 3th, 2024

Urgent Field Safety Notice SBN-CPS-2019-013
Roche Diagnostics Regrets To Inform You Of Reported Cases Affecting The Cobas T 511 And Cobas T 711 Coagulation Analyzers. Description Of Situation The Cobas T 511 And Cobas T 711 Coagulation Analyze 1th, 2024

Urgent Field Safety Notice - HPRA
As You Know, The CoaguChek® XS Plus/CoaguChek® XS Pro Instrument Measures Your Patients’ Coagulation Status Up To An INR Value Of 8.0. It Is Known That, In Very Rare Situations, INR Values Greater Than 8.0 Can Occur – For Example If The Patient Is Undergoing Antibiotic Trea 1th, 2024

URGENT FIELD SAFETY NOTICE Covidien Parietex™ Composite ...
Mesh Failure Identified Several Years Following Parastomal Hernia Repair Using The Modified Sugarbaker Repair Technique. In These Reports, Parietex™ Composite Parastomal Mesh Failure Led To Hernia Recurrence Requiring Additional Surgical Treatment. Symptoms Of Hernia Recurren 1th, 2024

Urgent Field Safety Notice - Recall Olympic Brainz Monitor
The Olympic Brainz Monitor (OBM) Cerebral Function Monitor (CFM) Is A Three Channel Electroencephalograph (EEG) Acquisition System Intended To Be Used In A Hospital Environment To Record, Collect, 2th, 2024

URGENT: FIELD SAFETY NOTICE CONMED Corporation …
(Valleylab/Covidien) REM™ Force 1B Force 1C Force 2 Force 300 Force 30 Force 40 Force 40S Force 4 Force 4B Force FX Force Triad FT10 ERBE NESSY (Neutral Electrode Safety System) ERBE VIO KLS Martin Group PCS (Patient Control System) Beamer PCS Birtcher Medical PSS® System 6400 : 2th, 2024