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Kci Rotoprone Bed Manual

With Rotoprone, patients can be easily turned and repositioned with the touch of a button. Kci Rotoprone Bed Manual Transfer Switch/Cost Of Rotoprone Bed - 5100X/DeNovo/Registration & Listing/Adverse Events/Recalls/PMA/IDE/Classification/Standards/CFR Title 21/Radiation-Emitting Products/X-Ray Assemblies/Medical Reports/CLIA/TPLC KCI USA, INC. Unit is not designed to allow manual rotation of the rotoprone frame to the prone position. ROTOPRONE THERAPY Back to Search Results/Device Problem Device Operates Differently Than Expected (2013)/Patient Problem No Known Impact Or Consequence To Patient (2005)/Event Date: 01/05/2010/Event Type: Malfunction/Event Description On: 06/01/2011, the nurse reported that the rotoprone would not rotate to prone. Ensure patient surface is at zero degrees supine and lock pin is fully inserted. The exact nature or cause of the malfunction with the dc motor controller was indeterminate. The right hospital bed and therapy surface is key to delivering effective care. Agilis has partnered with leading manufacturers to provide care facilities with access to a diverse range of specialty hospital beds – such as low, lift and bariatric beds – and therapeutic mattresses that include low air loss (LAL), lateral rotation and fluid. Several unsuccessful attempts were made to obtain further information. Details of the incident were not available.

Follow all applicable safety rules and instruction protocols for transfer and manual proning. Search Alerts/Recalls/New Search | Submit an Adverse Event Report/Kci Rotoprone Bed Manual Transfer Switch. Unbuckle and stow proning packs; remove side supports; open head support Place patient in prone position on rotoprone patient surface following facility patient handling and manual proning protocols, or transfer patient to alternate surface and place in prone position. Unlike manual proning which generally requires 6-8 staff members, proning patients with Rotoprone® can be accomplished by 1-2 caregivers. There was no reported injury. Manufacturer Narrative The bed was tested per quality control procedures on 08/06/2010 and met specifications. Device labeling, available in print and online, states if patient is prone-dependent: do not attempt to manually rotate the patient from supine to prone. The cause for the malfunction of the bed was the dc motor controller that controls the rotation motor. The bed was placed with the patient on 09/09/2011 Kci was able to confirm that the bed would not rotate to prone.

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