ORTHOCON Announces Uninterrupted Availability of HEMA*SORB* Resorbable Hemostatic Bone Putty

IRVINGTON, N.Y., Aug. 23, 2012 /PRNewswire/ -- On August 21, 2012, the U.S. Food and Drug Administration (FDA) announced that a hemostatic bone putty has been subjected to a Class 1 recall because of potential ignition if contacted with electrosurgical cautery systems under certain conditions during surgery. ORTHOCON and its hemostatic bone putty (HEMA*SORB)* are not associated with this recall.

For more information regarding HEMA*SORB*, please visit www.orthocon.com or call 914.357.2612.
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