MEMORANDUM

Device: Arthrex DX Reinforcement Matrix

Kensey Nash Corporation received FDA 510(k) clearance for Medeor™ Matrix on 22 October 2009 as a Class II device (K#091499). Under an agreement between Kensey Nash Corporation and Arthrex Inc, the device will be commercially distributed by Arthrex as Arthrex DX Reinforcement Matrix. Other than the product trade name, no differences exist between Medeor™ Matrix and Arthrex DX Reinforcement Matrix.

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