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2024Vestibular Evoked Myogenic Potential TestingVestibular Analysis Devices Are Currently Regulated By The U.S. Food And Drug Administration (FDA) Through The 510(k) Pathway, Under FDA Product Code LXV. The Term "vestibular Analysis Devices" Includes Both Diagnostic Devices (e.g., Rotary Chairs, Multiaxial Chairs) And Jul 1th, 2024Evoked Potential Studies MP-4 - AAPCTo Aid In The Diagnosis And Monitoring Of Multiple Sclerosis (MS). Recordings Made In The Acute Phase Of MS Disclose The Presence Of Active, Demyelinating Plaques. Recordings Made In The Chronic Phase Of MS Indicate The Presence Of Subclinical Plaques; Or To Localize The Basis For Vi May 3th, 2024.

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