

TITLE/TOPIC OF GUIDANCE	CONTACT
Guidance for Saline, Silicone Gel, and Alternative Breast Implants (Revised)	Samie Allen, Center for Devices and Radiological Health (HFZ-410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-3090
Class II Special Controls Guidance Document: Resorbable Calcium Salt Bone Void Filler Device (Final)	Nadine Sloan, Center for Devices and Radiological Health (HFZ-410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-1296
Class II Special Controls Guidance Document: Transcutaneous Electrical Stimulator for Cosmetic Use (Draft)	Robert DeLuca, Center for Devices and Radiological Health (HFZ-450), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-1296
Class II Special Controls Guidance Document: Cutaneous Electrode (Draft)	Do
Class II Special Controls Guidance Document: Electroconductive Media (Draft)	Do
Class II Special Controls Guidance Document: Powered Muscle Stimulator for Muscle Conditioning (Draft)	Do
Class II Special Controls Guidance Document: Powered Muscle Stimulator for Rehabilitation (Draft)	Do
Class II Special Controls Guidance Document: Transcutaneous Electrical Nerve Stimulator for Pain Relief (Draft)	Do
Special Control Guidance for Premarket Notifications for Totally Implanted Spinal Cord Stimulators for Pain Relief (Withdrawal)	Kristen Bowsher, Center for Devices and Radiological Health (HFZ-450), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-1296
Guidance for Technical Reporting in the Submission of Research and Marketing Applications for Totally Implanted Spinal Cord Stimulators (Draft)	Do
CATEGORY—PREMARKET REVIEW FOR OPHTHALMIC AND ENT DEVICES	
Class II Special Controls Guidance Document: Rigid Gas Permeable (RGP) by Contact Lens Finishing Laboratories	James F. Saviola, Center for Devices and Radiological Health (HFZ-460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-1744
Premarket Notification (510(k)) Guidance Document for Class II Daily Wear Contact Lenses (Revised)	Do
Class II Special Controls Guidance Document: Artificial Eye Care Products	Do
Class II Special Controls Guidance Document: Intraocular Gases for Retina Tamponade	Do
Retinal Implants: Guidance for Investigational Device Exemptions (IDE) and Premarket Approval (PMA) Applications (Draft)	Do
Guidance for Premarket Approval Applications of Class III Extended Wear Contact Lenses	Do
Guidance for Post Approval Studies of Class III Extended Wear Contact Lenses Worn Beyond Seven Continuous Nights	Do
Labeling Guidance for Ultraviolet Absorbing Contact Lenses	Do
Intraocular Lens Guidance Document	Donna R. Lochner, Center for Devices and Radiological Health (HFZ-460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-2053
Refractive Implants Guidance Document	Do
Guidance Document for Keratomes and Keratome Blades	Everette T. Beers, Chief, Center for Devices and Radiological Health (HFZ-460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-2018