

Issuing Org	Reg. Type	MITA ID/Date Code	Title	Expected Release	Actual Release	sent to TRC	Member Feedback Deadline	First Draft Deadline	MITA Finalization Deadline	Final Deadline	MITA Comments Status
FDA	Draft Guidance	2014.PMADData-DG.0722	Balancing PMA Premarket & Postmarket Data Draft Guidance		4/23/2014	4/23/2014			7/22/2014	7/22/2014	no comment
FDA	Draft Guidance	2014.CompModeling.0417	Computational Modeling Draft Guidance			2/4/2014				4/17/2014	submitted
FDA	Agency Report	2014.510kMods.0604	FDA Report to Congress on 510(k) Modifications			2/27/2014				6/4/2014	submitted
FDA	Draft Guidance	2014.DistributingPubs.0502	Distributing Scientific Publications Draft Guidance			4/3/2014				5/2/2014	submitted
FDA	Draft Document	2014.FDAPriorities.0731	Draft FDA Strategic Priorities 2014-2018		June 2014	7/31/2014			7/31/2014	7/31/2014	complete
FDA	Draft Guidance	2014.VolStandards.0811	CDRH Draft Guidance on Voluntary Consensus Standards		5/13/2014	5/12/2014				8/11/2014	no comment
FDA	Draft Guidance	2014.OnlineMisinfo.0916	FDA Draft Guidance on Online Correction of Third Party Misinformation		June 2014	6/18/2014	9/2/2014	9/4/2014	9/10/2014	9/16/2014	
FDA	Draft Guidance	2014.OnlineInfo.0916	FDA Draft Guidance on Character-Limited Online Information		June 2014	6/18/2014	9/2/2014	9/4/2014	9/10/2014	9/16/2014	
FDA	Draft Rule	2014.ClassificationRule.0922	Device Classification Draft Rule		3/25/2014	5/5/2014	9/8/2014	9/10/2014	9/17/2014	9/22/2014	
FDA	Draft Guidance	2014.510kBenefitRisk.1001	510(k) Benefit-Risk Draft Guidance (due October)		7/14/2014	7/28/2014	9/17/2014		9/17/2014	10/1/2014	
FDA	Final Guidance	2014.CDRHAppeals	Center for Devices and Radiological Health Appeals Processes: Questions and Answers About 517A - Guidance for Industry and Food and Drug Administration Staff		7/30/2014	8/1/2014			n/a	n/a	no comment
FDA	FYI		Co-convenership of IEC/SC 62A-ISO/TC 210 JWG 4, Medical devices			8/26/2014	n/a	n/a	n/a	n/a	