Publicly Accessible Databases for MDSAP Audits

	Main page	Market Authorizations	Site Licenses/ Registrations	Adverse Events	Advisory Notices	Other topics
TGA	TGA	eBusiness	NA	Adverse Event Notifications (DAEN)	Recalls (SARA)	TGA Act & Regulations Standards Orders and Medical Devices Clinical Evidence Guidelines IVD Guidelines Other guidance
ANVISA	ANVISA	Product Registration	International Manufacturers ID	Adverse Events and Quality Issues Adverse Events and Quality Issues* *computerized system developed by ANVISA for the	Recalls, Counterfeit, Suspended Products	MD IFU ANVISA Website Guidance

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				news of incidents, adverse events (AE) and technical complaints (QT) related to the use of products and services under sanitary surveillance		
Health Canada	Health Canada Medical Devices	Medical Device Active License Listing (MDALL)	Medical Device Establishment Licence Listing (MDEL)	Medical Device Incidents	Recalls and Safety Alerts	NA
MHLW/ PMDA	PMDA	NA	Foreign Manufacturing Sites	NA	Recalls	PMD Act MHLW MO169 Standards for Medical Devices
FDA	FDA	Device Registration and Listing (online search) Establishment Registration & Device Listing	Device Registration and Listing (online search) Establishment Registration & Device Listing	MDR (online search) MAUDE	List of Device Recalls Medical Device Safety (online search) Recalls	Inspection Classification 510(K) Premarket Notification Premarket Approval (PMA)

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						Total Product Life Cycle Product Code Builder Unique Device Identifier CFR Code of Federal Regulations Title 21 eCFR ICH Guidelines Warning Letters