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**labnotes - volume 19, no. 1, 2009 - bd: medical** - and the Safe Medical Devices Act (1990) the potential impact of such environmental factors on the draw volume of evacuated tubes. (1990). Federal Register.

**neil f. o'flaherty | ofw law | washington, d.c** - Neil F. O'Flaherty Principal Attorney plans and strategies for their medical devices and HCT/PS; registered clients Medical Device Information, Vol

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**fda medical devices - official site** - International Medical Device Regulators Forum, (Medical Devices) Medical Device News, Device Registration and Listing; Medical Device Databases;

**medical device register. international volume** - International volume.. [Medical Device Register, Inc.]; Medical device register. International volume datePublished " 1990/1995" ;

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**a lightweight assessment method for medical device** - Federal Register, 2011. vol Development for Medical Devices. In: First International Symposium on assessment method for medical device

**post-marketing surveillance and vigilance for** - Council Directive: 93/42/EEC concerning Medical Devices, OJ L169; Volume 36; 1993 Jul 12

**register as a manufacturer to sell medical** - is the competent authority for the registration of medical devices. MHRA will only register manufacturers or Database for Medical Device Registration.

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**cfr - code of federal regulations title 21** - SUBCHAPTER H--MEDICAL DEVICES: Sec. 820.181 Device master record. Device specifications including appropriate drawings,

**10586 federal register / vol. 80, no. 39/friday**, - Feb 26, 2015 Medical Device Reporting: Manufacturers, International and Federal Register/Vol. 80, No. 39/Friday,

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**sgs - harmonisation of asia's medical device** - of Malaysia s medical devices to international to register with the Medical Device Registered medical device establishments will

**brad a. james, ph.d., p.e., fasm | professionals** | - (ASM International) involving failure analysis, design, and life prediction/validation of medical devices, Registered Professional Engineer,

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