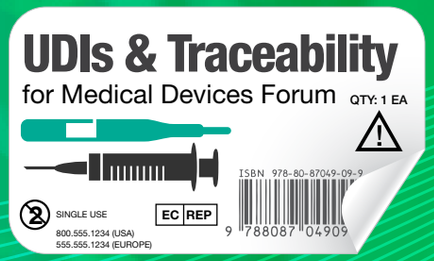
UPDATED FOR 2015



THE UDI FINAL RULE INTERACTIVE COMPLIANCE CALENDAR

Medical Device UDIs & Traceability Forum - 26 - 28 May, 2015 - Munich, Germany

> CLICK HERE <

List of medical devices the FDA classifies as implantable, life-saving, and life-sustaining for purposes of section 614 of FDASIA amending section 519(f) of the FDC Act, September 2013

The FDA Final Rule on unique device identification (UDI) for medical devices is out. Covering both labelling and the GUDID database, manufacturers need to assess their current technical capabilities - both hardware and software - and work out how much they need to do in order to comply. Ahead of the **Medical Device UDIs & Traceability Forum**, taking place 26 - 28 May, 2015 in Munich, Germany, Pharma IQ has put together this one-page compliance calendar to help you understand what impact the September 2013 FDA Final Rule on unique device identification will have on your business.

TO DOWNLOAD THE FDA FINAL RULE
> CLICK HERE <

Hover over boxes below to reveal the requirements

1 Year
after publication of the final rule
(September 24, 2014)

2 Years
after publication of the final rule
(September 24, 2015)

3 Years
after publication of the final rule
(September 24, 2016)

5 Years
after publication of the final rule
(September 24, 2018)

7 Years
after publication of the final rule
(September 24, 2020)

FOR MORE INFORMATION GO TO: **WWW.UDISANDTRACEABILITY.COM**

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Source: www.fda.gov - Compliance dates for all other provisions of the final rule. Except for the provisions listed above, FDA requires full compliance with the final rule as of the effective date that applies to the provision.