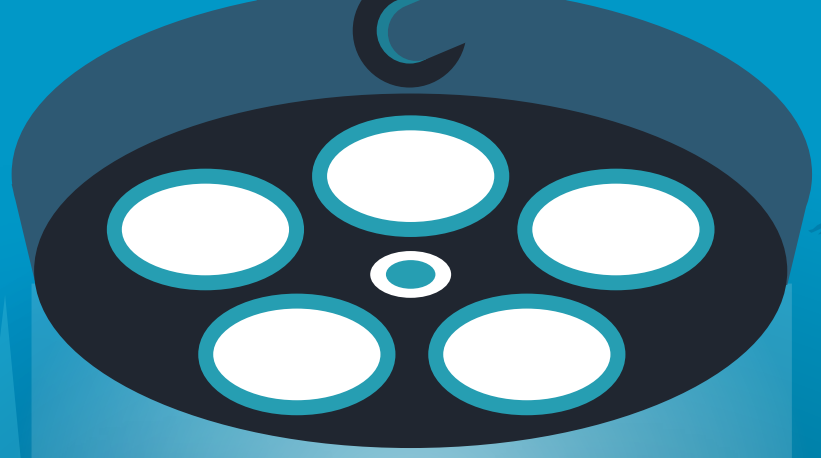
Taking the Pulse of UDI Compliance

We polled approximately 120 professionals in the medical device industry to see where they are with UDI compliance and labeling as of May 2016. According to the FDA, Class II and Class III devices must bear a UDI label by September 24, 2016.



47%
Are not sure their current barcode labeling solution will be able to scale to meet international UDI requirements

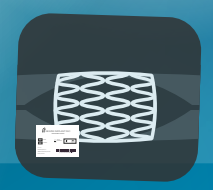
50%
Cite getting all the necessary data on the device label as a top remaining challenge

45%
Cite integrating to enterprise applications for labeling data as a top remaining challenge

15%
Have already achieved UDI compliance for Class II and Class III devices

36%
Expect to meet all Class II and Class III UDI requirements right on the Sept. 24th deadline

56%
Are using a single standardized barcode labeling software solution to meet UDI requirements



75%
Are currently working towards UDI compliance for Class II and Class III devices

47%
Say that internal audits as part of their UDI program have been conducted or are planned

53%
Still need support for ongoing UDI program processes and regulatory requirements

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