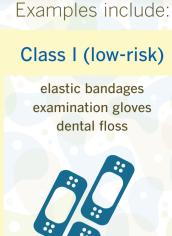
Unique Device Identification: Get to know UDI



UDI is a unique device identification system created and regulated by the U.S. Food and Drug Administration (FDA). It is designed to adequately identify medical devices through their distribution and use. When fully implemented, most medical devices will include a unique device identifier in human and machine-readable forms. When required, these identifiers must not only appear on labels and packaging, but on the devices themselves.

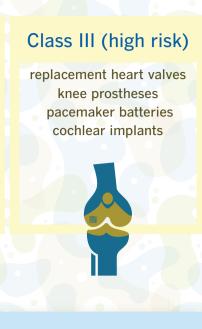
This summary is for informational purposes only and is not intended as legal advice. For a complete description of the Unique Device Identification system, go to: http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/UniqueDeviceIdentification/

What is a medical device?



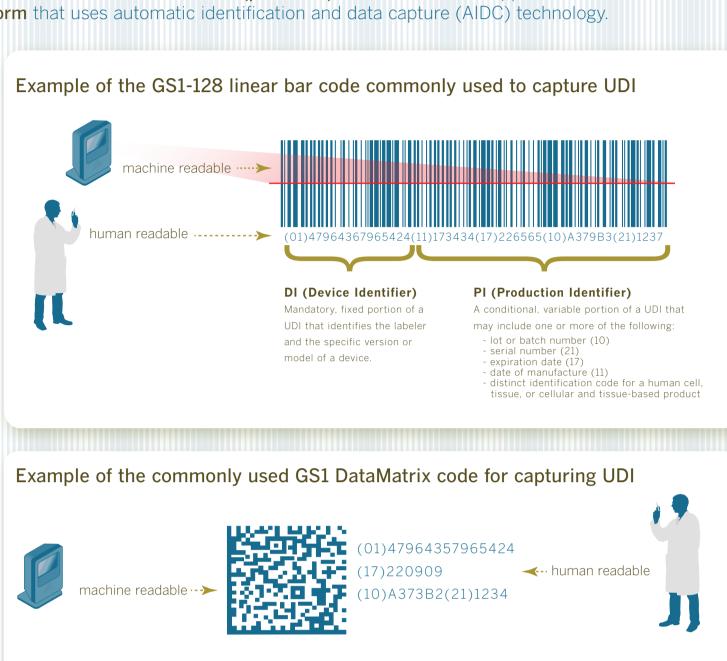






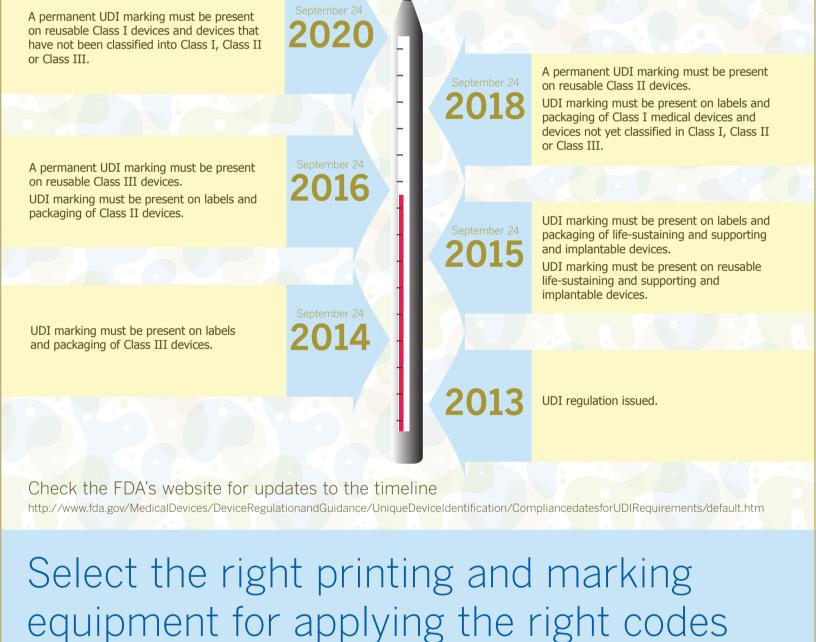
What is a UDI code?

On each medical device, label and package subject to the regulation, a UDI code must be provided in a **human-readable (plain-text) form**. It also must appear in a **machine-readable form** that uses automatic identification and data capture (AIDC) technology.



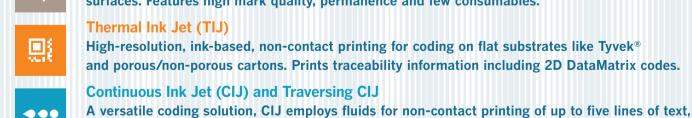
coding requirements

Keep an eye on the UDI timeline for





Meet the demand for accurate codes on paperboard, plastic, glass, labels and



Thermal Transfer Overprinter (TTO)

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Cases

A digitally controlled printhead precisely melts ink from a ribbon directly onto flexible films to provide high-resolution, real time prints.

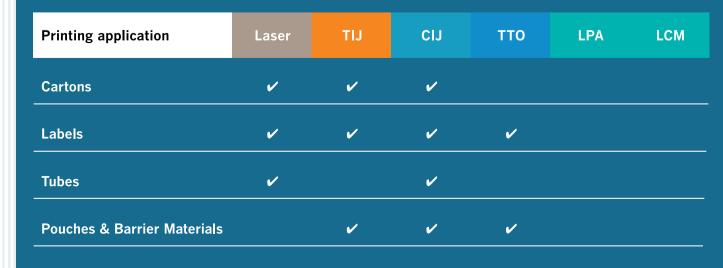
Label Printer Applicator (LPA)

as well as linear and 2D bar codes. Can print on stationary packaging via traversing systems.

Prints and places labels on cases of various sizes for traceability throughout the supply chain.

Large Character Marking (LCM)
Ink-based, non-contact printing of alphanumeric codes, logos and bar codes on cases.

Coding technologies for your packaging type:



Learn more about coding on medical devices
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