This addendum describes a new label on your product. The new label complies with a United States Food and Drug Administration directive for Unique Device Identification (UDI). For more information about UDI, go to *http://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/UniqueDeviceIdentification/UDIBasics/default.htm*.

The new nameplate label includes UDI information, and is placed on the rear of the device. See the following example label.

Place this document with your product manuals for future reference.