

**K062342 TRUEPULSE DISPOSABLE INVASIVE BLOOD
PRESSURE MONITORING KIT**Sep 25, 2006
45 days to decisionK062342 · Product code: **DRS** · Cardiovascular
Source: <https://www.510kdatabase.net/k062342/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Aug 11, 2006
Decision date	Sep 25, 2006
Days to decision	45 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Sentra Medical Devices, LLC
Location	Canton, MI, US
Contact	KARTHIK MARAYAN
510(k) history	1 submissions · 1 cleared · 2006-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062342/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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