

**K913595 DS-240 AMBULATORY BLOOD PRESSURE
MONITORING SYSTEM**Mar 27, 1992
227 days to decisionK913595 · Product code: **DXN** · Cardiovascular
Source: <https://www.510kdatabase.net/k913595/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Aug 13, 1991
Decision date	Mar 27, 1992
Days to decision	227 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Koven and Assoc., Inc.
Location	St. Louis, MO, US
Contact	PAUL G KOVEN
510(k) history	8 submissions · 8 cleared · 1989-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k913595/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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