FDA 510(k) Premarket Notification

K913595 DS-240 AMBULATORY BLOOD PRESSURE MONITORING SYSTEM

CLEARED

Mar 27, 1992
227 days to decision

K913595 · 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