FDA 510(k) Premarket Notification

K961144 MS-2000

CLEARED

Mar 12, 1997
356 days to decision

K961144 · Product code: **DXN** · Cardiovascular
Source: https://www.510kdatabase.net/k961144/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Measurement, Blood-pressure, Non-invasive (DXN)
Date received	Mar 21, 1996
Decision date	Mar 12, 1997
Days to decision	356 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Touritu Engineering Co., Inc.**
Location	San Jaun Capistrano, CA, US
Contact	JAMES R GREENWOOD, PH.D.
510(k) history	1 submissions · 1 cleared · 1997-1997