

K961144 MS-2000Mar 12, 1997
356 days to decisionK961144 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k961144/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026