FDA 510(k) Premarket Notification

K011003 SAFESTING AND SAFESTING HUB

CLEARED

Apr 20, 2001
16 days to decision

K011003 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k011003/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 4, 2001
Decision date	Apr 20, 2001
Days to decision	16 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Diasol, Inc.**
Location	North Hollywood, CA, US
Contact	MONICA ABELES
510(k) history	10 submissions · 10 cleared · 1999-2014

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k011003/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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