FDA 510(k) Premarket Notification

K010888 SIDEPORT PINCH CLAMP DEVICE

CLEARED

Jun 27, 2001
96 days to decision

K010888 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k010888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 23, 2001
Decision date	Jun 27, 2001
Days to decision	96 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medivice Systems, Ltd.**
Location	Kfar Saba, IL
Contact	AHAVA STEIN
510(k) history	1 submissions · 1 cleared · 2001-2001