

K010888 SIDEPORT PINCH CLAMP DEVICEJun 27, 2001
96 days to decisionK010888 · 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

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