FDA 510(k) Premarket Notification

K913311 SAFETY INJECTION PORT, MODIFICATION

CLEARED

Sep 13, 1991
56 days to decision

K913311 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k913311/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jul 19, 1991
Decision date	Sep 13, 1991
Days to decision	56 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Riverside Technology, Inc.**
Location	Lisle, IL, US
Contact	W. C MILLER
510(k) history	1 submissions · 1 cleared · 1991-1991