FDA 510(k) Premarket Notification

K842759 MODEL 503FB

CLEARED

Jan 18, 1985
186 days to decision

K842759 · Product code: **LOS** · Cardiovascular
Source: https://www.510kdatabase.net/k842759/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jul 16, 1984
Decision date	Jan 18, 1985
Days to decision	186 days
Third-party review	No

APPLICANT

Company	**Atrix**
Location	Portland, OR, US
Contact	KIM F DUNCAN
510(k) history	2 submissions · 2 cleared · 1985-1986