FDA 510(k) Premarket Notification

K830573 PRE VUE

CLEARED

Mar 29, 1983
34 days to decision

K830573 · Product code: **FOZ** · General Hospital
Source: https://www.510kdatabase.net/k830573/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Therapeutic, Short-term Less Than 30 Days (FOZ)
Date received	Feb 23, 1983
Decision date	Mar 29, 1983
Days to decision	34 days
Third-party review	No

APPLICANT

Company	**Taut, Inc.**
Location	Walker, MI, US
510(k) history	16 submissions · 16 cleared · 1983-2002