FDA 510(k) Premarket Notification

K812080 AEGIS I.V. STRIPS

CLEARED

Aug 3, 1981
12 days to decision

K812080 · Product code: **KMK** · General Hospital
Source: https://www.510kdatabase.net/k812080/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Intravascular Catheter Securement (KMK)
Date received	Jul 22, 1981
Decision date	Aug 3, 1981
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Warner-Lambert Co.**
Location	Mchenry, IL, US
510(k) history	50 submissions · 50 cleared · 1979-2003