FDA 510(k) Premarket Notification

K823790 VENTRICULAR ACCESS SYSTEM

CLEARED

Apr 28, 1983
133 days to decision

K823790 · Product code: **LKG** · General Hospital
Source: https://www.510kdatabase.net/k823790/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Port & Catheter, Implanted, Subcutaneous, Intraventricular (LKG)
Date received	Dec 16, 1982
Decision date	Apr 28, 1983
Days to decision	133 days
Third-party review	No

APPLICANT

Company	**American Heyer Schulte**
Location	Mchenry, IL, US
510(k) history	13 submissions · 13 cleared · 1980-1984