FDA 510(k) Premarket Notification

K810311 STERI-VERS

CLEARED

Apr 17, 1981
71 days to decision

K810311 · Product code: **LDS** · General Hospital
Source: https://www.510kdatabase.net/k810311/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Pasteurization, Hot Water (LDS)
Date received	Feb 5, 1981
Decision date	Apr 17, 1981
Days to decision	71 days
Third-party review	No

APPLICANT

Company	**The Hilrad Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981