FDA 510(k) Premarket Notification

K801309 RITTER STERILIZER

CLEARED

Jun 20, 1980
17 days to decision

K801309 · Product code: **FLE** · General Hospital
Source: https://www.510kdatabase.net/k801309/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sterilizer, Steam (FLE)
Date received	Jun 3, 1980
Decision date	Jun 20, 1980
Days to decision	17 days
Third-party review	No

APPLICANT

Company	**Sybron Corp.**
Location	Mchenry, IL, US
510(k) history	37 submissions · 37 cleared · 1977-1986