

K801309 RITTER STERILIZERJun 20, 1980
17 days to decisionK801309 · 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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k801309/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026