

K810311 STERI-VERSApr 17, 1981
71 days to decisionK810311 · Product code: **LDS** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k810311/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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