

K831058 SAF FIXATIVEJun 8, 1983
68 days to decisionK831058 · Product code: **LDW** · Pathology
Source: <https://www.510kdatabase.net/k831058/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fixative, Acid Containing (LDW)
Date received	Apr 1, 1983
Decision date	Jun 8, 1983
Days to decision	68 days
Third-party review	No

APPLICANT

Company	American Micro Scan
Location	Mchenry, IL, US
510(k) history	24 submissions · 24 cleared · 1983-1987

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Device record: <https://www.510kdatabase.net/k831058/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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