FDA 510(k) Premarket Notification

K831058 SAF FIXATIVE

CLEARED

Jun 8, 1983
68 days to decision

K831058 · 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