

K831580 PARAFFIN INFILTRATION MEDIUMJun 17, 1983
31 days to decisionK831580 · Product code: **KEO** · Pathology
Source: <https://www.510kdatabase.net/k831580/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Formulations, Paraffin, All (KEO)
Date received	May 17, 1983
Decision date	Jun 17, 1983
Days to decision	31 days
Third-party review	No

APPLICANT

Company	Surgipath Medical Industries, Inc.
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1977-1988

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Device record: <https://www.510kdatabase.net/k831580/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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