

K831335 DISPOSABLE BASE MOLDMay 16, 1983
21 days to decisionK831335 · Product code: **KER** · Pathology
Source: <https://www.510kdatabase.net/k831335/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Container, Embedding (KER)
Date received	Apr 25, 1983
Decision date	May 16, 1983
Days to decision	21 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/k831335/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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