FDA 510(k) Premarket Notification

K831335 DISPOSABLE BASE MOLD

CLEARED

May 16, 1983
21 days to decision

K831335 · 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