FDA 510(k) Premarket Notification

K810323 ACRYTOL, MOUNTING MEDIUM

CLEARED

Mar 6, 1981
28 days to decision

K810323 · Product code: **KEP** · Pathology
Source: https://www.510kdatabase.net/k810323/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Mounting, Oil Soluble (KEP)
Date received	Feb 6, 1981
Decision date	Mar 6, 1981
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**Surgipath**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1981-1982