FDA 510(k) Premarket Notification

K760602 SORVALL EMBEDDING MEDIUM

CLEARED

Sep 21, 1976
14 days to decision

K760602 · Product code: **KEO** · Pathology
Source: https://www.510kdatabase.net/k760602/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Formulations, Paraffin, All (KEO)
Date received	Sep 7, 1976
Decision date	Sep 21, 1976
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**E.I. Dupont DE Nemours & Co., Inc.**
Location	Mchenry, IL, US
510(k) history	253 submissions · 252 cleared · 1976-1996