FDA 510(k) Premarket Notification

K802359 DISP. FENESTRATED SURG. DRAPE SHEETS

CLEARED

Oct 31, 1980
35 days to decision

K802359 · Product code: **KKX** · General Hospital
Source: https://www.510kdatabase.net/k802359/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drape, Surgical (KKX)
Date received	Sep 26, 1980
Decision date	Oct 31, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**American Convertors Div., American Pharmaseal**
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1980-1984