FDA 510(k) Premarket Notification

K810078 Q-PAK THERAPEUTIC DRUG MONIT. CONTROL S

CLEARED

Feb 2, 1981
20 days to decision

K810078 · Product code: **DIF** · Chemistry
Source: https://www.510kdatabase.net/k810078/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drug Mixture Control Materials (DIF)
Date received	Jan 13, 1981
Decision date	Feb 2, 1981
Days to decision	20 days
Third-party review	No

APPLICANT

Company	**Hyland Therapeutic Div., Travenol Laboratories**
Location	Mchenry, IL, US
510(k) history	44 submissions · 44 cleared · 1976-1982