

K810078 Q-PAK THERAPEUTIC DRUG MONIT. CONTROL SFeb 2, 1981
20 days to decisionK810078 · 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

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Device record: <https://www.510kdatabase.net/k810078/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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