FDA 510(k) Premarket Notification

K792349 SUSCEPTIBILITY DISK, CINOXACIN 100 MCG.

CLEARED

Mar 12, 1980
114 days to decision

K792349 · Product code: **JNT** · Microbiology
Source: https://www.510kdatabase.net/k792349/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Chromatographic, Phospholipids (JNT)
Date received	Nov 19, 1979
Decision date	Mar 12, 1980
Days to decision	114 days
Third-party review	No

APPLICANT

Company	**Pfizer, Inc.**
Location	Mchenry, IL, US
510(k) history	30 submissions · 30 cleared · 1977-2018

Pfizer, Inc. is an American multinational pharmaceutical and biotechnology corporation headquartered in Manhattan, New York City. Founded in 1849, Pfizer is one of the oldest pharmaceutical companies in North America. Pfizer's FDA 510(k) regulatory record includes cleared devices from total submissions, spanning 1977 to 2018. The company's device portfolio demonstrates strength in orthopedic devices, including surgical implants and fixation systems. This regulatory activity is now historical, with no clearances recorded in the past five years. The company's cleared device...