FDA 510(k) Premarket Notification

K810240 CLINIGUARD STERILE

CLEARED

Feb 19, 1981
22 days to decision

K810240 · Product code: **KMF** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810240/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bandage, Liquid (KMF)
Date received	Jan 28, 1981
Decision date	Feb 19, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Clinipad Corp.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 12 cleared · 1980-1994