

**K810240 CLINIGUARD STERILE**Feb 19, 1981  
22 days to decisionK810240 · Product code: **KMF** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k810240/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k810240/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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