FDA 510(k) Premarket Notification

K810075 AUTOCLIX

CLEARED

Feb 4, 1981
22 days to decision

K810075 · Product code: **FMK** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k810075/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Single Use Only Blood Lancet With An Integral Sharps Injury Prevention Feature (FMK)
Date received	Jan 13, 1981
Decision date	Feb 4, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	**Biodynamics Corp.**
Location	Walker, MI, US
510(k) history	21 submissions · 21 cleared · 1980-2006

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810075/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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