

**K000060 GEL MARK BIOPSY SITE MARKER, MODELS
MK0001, MK0004, MK0007**

Mar 27, 2000
77 days to decision

K000060 · Product code: **GDW** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k000060/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Jan 10, 2000
Decision date	Mar 27, 2000
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Senorx, Inc.
Location	Irvine, CA, US
Contact	AMY BOUCLY
510(k) history	30 submissions · 26 cleared · 2000-2023

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k000060/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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