

K812390 NEUROSPONGE PADDIENov 24, 1981
96 days to decisionK812390 · Product code: **GDY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k812390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Gauze/sponge, Internal, X-ray Detectable (GDY)
Date received	Aug 20, 1981
Decision date	Nov 24, 1981
Days to decision	96 days
Third-party review	No

APPLICANT

Company	Winn Hirsch & Assoc.
Location	Mchenry, IL, US
510(k) history	11 submissions · 10 cleared · 1976-1988

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Device record: <https://www.510kdatabase.net/k812390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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