

K842111 KOGAN ENDOSPECULUM 9-1/2Jun 19, 1984
21 days to decisionK842111 · Product code: **HTZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k842111/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Cutting, Orthopedic (HTZ)
Date received	May 29, 1984
Decision date	Jun 19, 1984
Days to decision	21 days
Third-party review	No

APPLICANT

Company	J. Sklar Mfg. Co., Inc.
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1977-1984

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k842111/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026