

**K903601 ESPI NON-INSULATED DOUBLE-ACTION GRASPER
#8204**Sep 12, 1990
35 days to decisionK903601 · Product code: **HET** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k903601/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, Gynecologic (and Accessories) (HET)
Date received	Aug 8, 1990
Decision date	Sep 12, 1990
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Dixon Medical, Inc.
Location	Atlanta, GA, US
Contact	WILLIAM C DIXON
510(k) history	12 submissions · 12 cleared · 1990-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k903601/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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