FDA 510(k) Premarket Notification

K761030 EKOSECTOR I

CLEARED

Dec 13, 1976
31 days to decision

K761030 · Product code: **HGM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k761030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Monitoring, Perinatal (HGM)
Date received	Nov 12, 1976
Decision date	Dec 13, 1976
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Smithkline Diagnostics, Inc.**
Location	Mchenry, IL, US
510(k) history	42 submissions · 42 cleared · 1976-1996

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k761030/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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