FDA 510(k) Premarket Notification

K891465 OBSTETRIC CUP (MODIFICATION)

CLEARED

May 11, 1989
57 days to decision

K891465 · Product code: **HDB** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k891465/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Mar 15, 1989
Decision date	May 11, 1989
Days to decision	57 days
Third-party review	No

APPLICANT

Company	**Go Medical Industries Pty. , Ltd.**
Location	Walker, MI, US
Contact	GEORGE O'NEIL
510(k) history	11 submissions · 11 cleared · 1984-2001