FDA 510(k) Premarket Notification

K874383 MENTOR PLUS (CONDOM)

CLEARED

Mar 7, 1988
133 days to decision

K874383 · Product code: **HIS** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k874383/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Oct 26, 1987
Decision date	Mar 7, 1988
Days to decision	133 days
Third-party review	No

APPLICANT

Company	**Arcon Corp.**
Location	Stewartville, MN, US
Contact	RICHARD FRYAR
510(k) history	2 submissions · 2 cleared · 1984-1988