FDA 510(k) Premarket Notification

K902032 O. B. PACK IV

CLEARED

Sep 28, 1990
147 days to decision

K902032 · Product code: **HFC** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k902032/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspirator, Endocervical (HFC)
Date received	May 4, 1990
Decision date	Sep 28, 1990
Days to decision	147 days
Third-party review	No

APPLICANT

Company	**Customed, Inc.**
Location	Carolina Puerto Rico, US
Contact	FELIX B SANTOS
510(k) history	36 submissions · 26 cleared · 1990-1998