FDA 510(k) Premarket Notification

K831669 AXCAN SPERM CUP

CLEARED

Jul 12, 1983
50 days to decision

K831669 · Product code: **HDR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k831669/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	May 23, 1983
Decision date	Jul 12, 1983
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Axcan Scientific Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984