FDA 510(k) Premarket Notification

K894264 KDF-2.3

CLEARED

Aug 15, 1989
56 days to decision

K894264 · Product code: **HDR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k894264/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	Jun 20, 1989
Decision date	Aug 15, 1989
Days to decision	56 days
Third-party review	No

APPLICANT

Company	**Unimar, Inc.**
Location	Mchenry, IL, US
Contact	ANTHONY HEMMING
510(k) history	10 submissions · 9 cleared · 1981-1994