FDA 510(k) Premarket Notification

K932505 RI-STAR

CLEARED

Oct 8, 1993
140 days to decision

K932505 · Product code: **ERA** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k932505/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Otoscope (ERA)
Date received	May 21, 1993
Decision date	Oct 8, 1993
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Rudolf Riester GmbH & Co. KG**
Location	D-72417 Jungingen, DE
Contact	KARLHEINZ RIESTER
510(k) history	30 submissions · 26 cleared · 1993-2000