FDA 510(k) Premarket Notification

K040300 STRYKER CONSOLIDATED OPERATING ROOM EQUIPMENT (CORE)

CLEARED

Mar 3, 2004
23 days to decision

K040300 · Product code: **ERL** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k040300/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Surgical, Ent (electric Or Pneumatic) Including Handpiece (ERL)
Date received	Feb 9, 2004
Decision date	Mar 3, 2004
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stryker Instruments**
Location	Kalamazoo, MI, US
Contact	JEAN W SHEPPARD
510(k) history	73 submissions · 73 cleared · 1994-2026