

K040300 STRYKER CONSOLIDATED OPERATING ROOM EQUIPMENT (CORE)Mar 3, 2004
23 days to decisionK040300 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k040300/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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