

**K954690 STRYKER ORAL MAX SYSTEM**Aug 20, 1996  
314 days to decisionK954690 · Product code: **DZI** · Dental  
Source: <https://www.510kdatabase.net/k954690/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Drill, Bone, Powered (DZI)
Date received	Oct 11, 1995
Decision date	Aug 20, 1996
Days to decision	314 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Corp.</b>
Location	Mchenry, IL, US
Contact	TAMMY LOUNDS
510(k) history	124 submissions · 121 cleared · 1976-2023

Stryker Corp. is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, implants, and patient safety technologies used globally across multiple medical specialties. Stryker has received FDA 510(k) clearances from total submissions since its first clearance in 1976. The company maintains active regulatory engagement, with its latest clearance in 2023. Its product portfolio spans orthopedic devices, neurosurgical implants, surgical instruments, and endoscopy systems, reflecting a broad pr...

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