

**K082335 SYNTHES MATRIXMANDIBLE PLATE AND SCREW SYSTEM**Nov 10, 2008  
88 days to decisionK082335 · Product code: JEY · Dental  
Source: <https://www.510kdatabase.net/k082335/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Bone (JEY)
Date received	Aug 14, 2008
Decision date	Nov 10, 2008
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	AMNON TALMOR
510(k) history	411 submissions · 394 cleared · 1977-2015

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k082335/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026