

**K152787 ST Internal Fixture System**Jul 22, 2016  
301 days to decisionK152787 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k152787/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Sep 25, 2015
Decision date	Jul 22, 2016
Days to decision	301 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>T-Plus Implant Tech. Co., Ltd.</b>
Location	New Taipei City, TW
Contact	DANA CHENG
510(k) history	4 submissions · 4 cleared · 2013-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k152787/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 28, 2026