

**K201660 KMTI Hip Replacement System, Tesera Trabecular Technologies (T3) Acetabular Shell System, Porous Acetabular Cup System**Jul 16, 2020  
28 days to decisionK201660 · Product code: **OQG** · Orthopedic  
Source: <https://www.510kdatabase.net/k201660/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Hip Prosthesis, Semi-constrained, Cemented, Metal/polymer, + Additive, Porous, Uncemented (OQG)
Date received	Jun 18, 2020
Decision date	Jul 16, 2020
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kyocera Medical Technologies, Inc.</b>
Location	Redlands, CA, US
Contact	Anthony DeBenedictis
510(k) history	15 submissions · 15 cleared · 2020-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201660/> 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 May 25, 2026