

**K132312 RENOVIS TESERA TRABECULAR TECHNOLOGY (T3)
ACETABULAR SHELL SYSTEM**Apr 11, 2014
260 days to decisionK132312 · Product code: LPH · Orthopedic
Source: <https://www.510kdatabase.net/k132312/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Porous Uncemented (LPH)
Date received	Jul 25, 2013
Decision date	Apr 11, 2014
Days to decision	260 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Renovis Surgical Technologies, LLC
Location	Round Rock, TX, US
Contact	Sharyn Orton, Ph.D.
510(k) history	9 submissions · 9 cleared · 2010-2014

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k132312/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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