

**K112507 RESTORIS PARTIAL KNEE APPLICATION (PKA), RIO  
STANDARD SYSTEM**Mar 1, 2012  
184 days to decisionK112507 · Product code: **OLO** · Orthopedic  
Source: <https://www.510kdatabase.net/k112507/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthopedic Stereotaxic Instrument (OLO)
Date received	Aug 30, 2011
Decision date	Mar 1, 2012
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Mako Surgical Corporation</b>
Location	Fort Lauderdale, FL, US
Contact	DIONNE SANDERS
510(k) history	8 submissions · 8 cleared · 2008-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k112507/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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