

K953169 STELKAST ALL POLYETHYLENE ACETABULAR CUPAug 8, 1995
32 days to decisionK953169 · Product code: **LZO** · Orthopedic
Source: <https://www.510kdatabase.net/k953169/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/ceramic/polymer, Cemented Or Non-porous, Uncemented (LZO)
Date received	Jul 7, 1995
Decision date	Aug 8, 1995
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stelkast Company
Location	Pittsburgh, PA, US
Contact	DONALD A STEVENS
510(k) history	40 submissions · 37 cleared · 1994-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k953169/> 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 18, 2026