FDA 510(k) Premarket Notification

K220792 SMR Reverse Liner

CLEARED

May 19, 2022
62 days to decision

K220792 · Product code: **MBF** · Orthopedic
Source: https://www.510kdatabase.net/k220792/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Shoulder, Semi-constrained, Metal/polymer, Uncemented (MBF)
Date received	Mar 18, 2022
Decision date	May 19, 2022
Days to decision	62 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Lima Corporate S.P.A.**
Location	Winona Lake, IN, US
Contact	Michela Zanotto
510(k) history	64 submissions · 64 cleared · 2011-2026

REGULATORY CONSULTANT

Consulting firm	**Lima U.S.A., Inc.**
Contact	Lacey Harbour

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov