

**K111323 SYNTHES SYNPOR HD POROUS POLYETHYLENE
THREER DIMENSIONAL IMPLANTS**Nov 16, 2011
189 days to decisionK111323 · Product code: **KKY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k111323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Material, Polytetrafluoroethylene Vitreous Carbon, For Maxillofacial Reconstruction (KKY)
Date received	May 11, 2011
Decision date	Nov 16, 2011
Days to decision	189 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Synthes, Inc.
Location	19380, PA, US
Contact	ALAN T HALEY
510(k) history	8 submissions · 8 cleared · 2011-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k111323/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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