

**K251479 Mobarn 80:20 Lumbar Interbody Fusion System  
(Posterior Lumbar Interbody Fusion Cage (PLIF))**Sep 18, 2025  
128 days to decisionK251479 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k251479/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 13, 2025
Decision date	Sep 18, 2025
Days to decision	128 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Mobarn 80:20 Lumbar Interbody Fusion System (Transforaminal Lumbar Interbody Fusion Cage (TLIF)); Mobarn 80:20 Lumbar Interbody Fusion System (Direct Lateral Lumbar Interbody Fusion Cage (DLIF)); Mobarn 80:20 Lumbar Interbody Fusion System (Anterior Lumbar Interbody Fusion Cage (ALIF))

**APPLICANT**

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Company	<b>Mobarn Medical Devices, LLC</b>
Location	New Orleans, LA, US
Contact	Richard Ehret
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Hypoid Ventures, LLC</b>
Contact	Lisa Ferrara

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