Acquisition of OrthoSpace by Stryker for \$220 million

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InSpace is the industry's only minimally invasive biodegradable, subacromial balloon spacer for arthroscopic treatment of massive, irreparable rotator cuff tears (MIRCTs)



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stryker®

\$220 million

Upfront payment of \$110 million and milestone payments of \$110 million

InSpace with long successful clinical history of over **10 years** and **29,000 balloons** implanted outside of the US

InSpace balloon with **CE mark** and under clinical study **without approval for use** in the US at the time

More than **20,000**patients across **30**countries treated with
InSpace

orthoSpace raised \$8
million for a U.S.
pivotal study in 2015,
followed by another \$7
million a year later including money from
Johnson & Johnson
Innovation, Smith &
Nephew, TriVentures
and
HealthpointCapital.

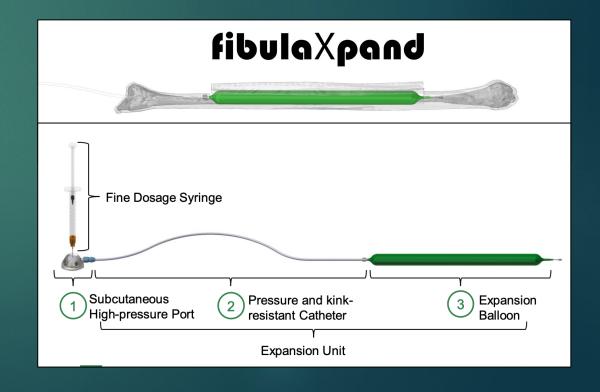


Main Similarities

- Comparable technology (balloon system)
- Expansion via injection of incompressible saline fluid
- Potentially similar regulatory requirements
- Niche applications

Main Differences

- Long-term implantation of InSpace (~1 year)
 vs midterm for fibulaXpand (~30 days)
- InSpace balloon is biodegradable
- fibulaXpand is applied inside the bone
- Gradual expansion (fibulaXpand) vs one-time expansion (InSpace)



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Potential strategic acquisition of fibulaXpand by major medical device companies after successful application in humans























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Sources

Stryker completes the acquisition of OrthoSpace, Ltd., a privately held company founded in 2009 and headquartered in Caesarea, Israel, in an all cash transaction for an upfront payment of \$110 million and future milestone payments of up to an additional \$110 million.

OrthoSpace's product portfolio provides a highly differentiated technology for the treatment of massive irreparable rotator cuff tears. The InSpace product is a biodegradable sub-acromial spacer, which is designed to realign the natural biomechanics of the shoulder. The technology has a long clinical history with over 20,000 patients treated across 30 countries. In the U.S., InSpace is currently under clinical study and not approved for use.

To date, the CE-marked InSpace balloon has been used in outpatient procedures for massive, irreparable tears in more than 20,000 patients in 30 countries, the companies said.

OrthoSpace raised \$8 million in funding for a U.S. pivotal study in 2015, followed by another \$7 million a year later—including money from Johnson & Johnson Innovation, Smith & Nephew, TriVentures and HealthpointCapital.

The InSpace balloon implant has a long successful clinical history of over 10 years and 29,000 balloons implanted outside of the US, as well as the Level I study conducted across North America.

Thank you!

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