

Corporate Overview

December 2018



Safe Harbor Statement

This presentation contains "forward-looking statements," which are statements related to future events that by their nature address matters that are uncertain. All forwardlooking information is inherently uncertain and actual results may differ materially from assumptions, estimates or expectations reflected or contained in the forward-looking statements as a result of various factors. For details on the uncertainties that may cause our actual results to be materially different than those expressed in our forwardlooking statements, please review our registration statement on Form S-1 and Quarterly Report on Form 10-Q, which contain risk factors. Forward-looking statements address our expected future business, financial performance, financial condition as well as results of operations, and often contain words such as "intends," "estimates," "anticipates," "hopes," "projects," "plans," "expects," "seek," "believes," "see," "should," "will," "would," "target," and similar expressions and the negative versions thereof. Such statements are based only upon current expectations of SI-BONE. Any forward-looking statement speaks only as of the date made. Reliance should not be placed on forward-looking statements because they involve known and unknown risks, uncertainties and other factors which may cause actual results, performance or achievements to differ materially from those expressed or implied. Forward-looking statements include statements that address activities, events or developments that SI-BONE expects, believes or anticipates will or may occur in the future. Forward-looking statements are based on SI-BONE's experience and perception of current conditions, trends, expected future developments and other factors it believes are appropriate under the circumstances and are subject to numerous risks and uncertainties, many of which are beyond SI-BONE's control. With respect to the forward-looking statements, we claim the protection of the safe harbor for forward-looking statements contained in the Private Securities Litigation Reform Act of 1995. We undertake no obligation to publicly update or revise any forward-looking statements to reflect new information or future events or otherwise unless required by law. This presentation includes the use of non-GAAP measures. Please review our Form 8-K filing which includes the furnishing of our earnings release for a reconciliation to the related GAAP measures.



35,000+ Procedures 1,700+ Surgeons 250M+
Covered lives

Market Leadership



ROBUST CLINICAL DATA >60 published papers

2 randomized trials



POSITIVE FINANCIAL PROFILE

\$53.5M revenue¹

16% growth²

91% gross margins²



REIMBURSEMENT ADVANTAGE

254M covered lives

24 payor exclusives



MARKET LEADER

4,300 U.S. procedures in 2017

279,000 U.S. potential patients per year

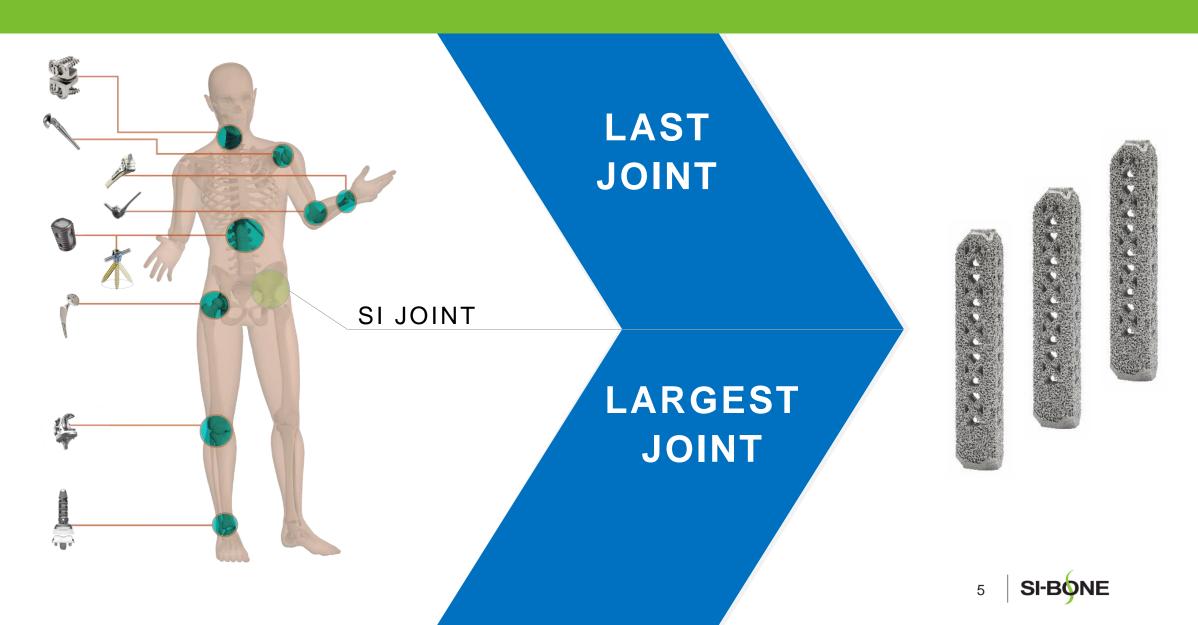
Notes:

1. LTM revenue as of September 30, 2018

2. YTD



Major Joints Market



SI Joint Pain: Prevalent and Highly Burdensome

15%
OF LOWER BACK
PAIN

32 - 43%

SYMPTOMATIC POST-LUMBAR FUSION



30M+ in the U.S. Suffer From Lower Back Pain...

4.7 M SI joint pain sufferers 1.4M

Eligible for surgery

5 years in pain





A Major Gap in Sacroiliac Joint Therapy

NON-SURGICAL MANAGEMENT SURGERY **MIS SI JOINT** MEDICATIONS, **THERAPEUTIC** RADIO-FREQUENCY **OPEN SI JOINT** PHYSICAL THERAPY **INJECTIONS ABLATION FUSION FUSION**

Clinically Proven Minimally Invasive Solution

- Patented unique triangular implants
- 3x the strength of a screw
- 6X the rotational resistance

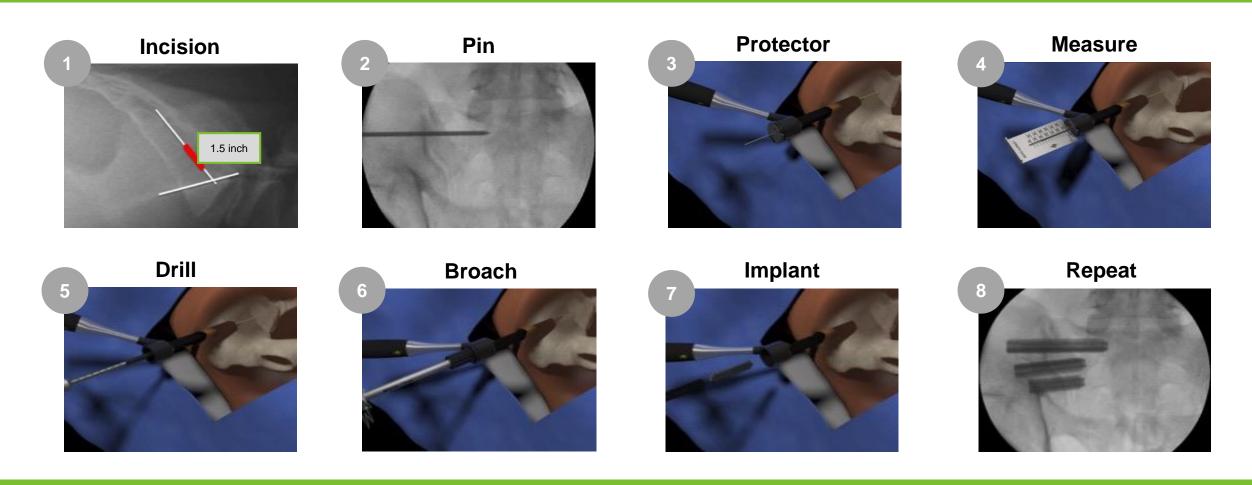


2008 FDA 510(k)

2010 CE MARK

34 COUNTRIES

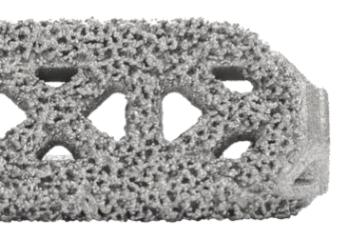
The iFuse Procedure™



Performed inpatient, outpatient or ASC

Platform Technology: iFuse-3DTM

Enhanced Implant Benefits

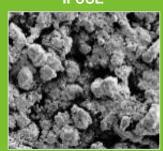


- Proven triangular design and procedure
- Porous, 3D-printed titanium implant
- Bony on-growth, in-growth, through-growth
- FDA/CE clearance in March/May 2017

REPRESENTATIVE COMPETITOR



TPS-COATED



CANCELLOUS BONE



3D-PRINTED IFUSE-3D



3 MONTH SHEEP STUDY



Proprietary, Differentiated Technology

	SI Screws	iFuse
	■ THE THE PARTY	
Rotation	1X resistance	>6x resistance
Safety	 Unknown 	 Very low complication rate
Revision	 Unknown 	3.5% (4-year)
Clinical Evidence	12 publications (no RCTs)	 >60 publications
Surface	- Smooth	- Porous

Patient Experience

VAS Pain Reduction¹

55 POINTS

Clinically meaningful at 20 pts

ODI Disability Improvement¹

28
POINTS

Clinically meaningful at 15 pts

91%

Patient satisfaction

3.5%

Low revision rate²



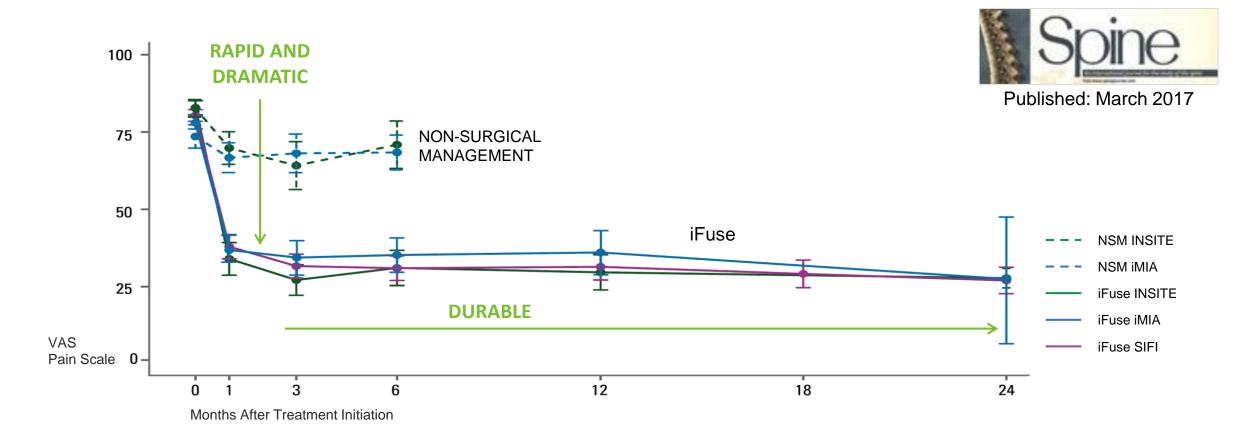
Robust Data and Clinical Evidence

- >60 peer-reviewed published papers
- Long-term 6-year data
- Two level 1 randomized studies



Prospective Outcomes

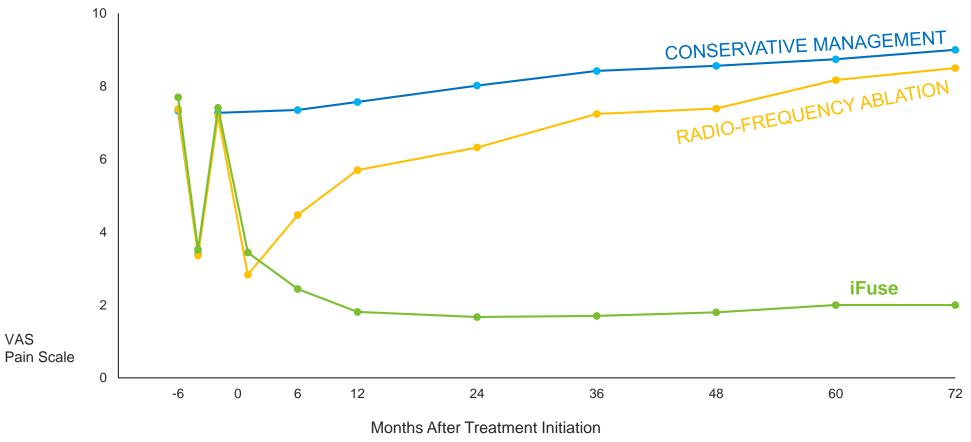
Pooled analysis of two randomized and one large prospective study



6-Year Cohort Study

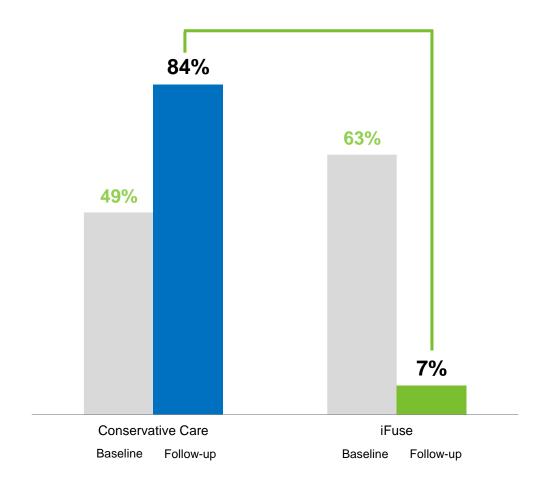


Published: April 2017



Significant Reduction In Opioid Users

6-year study, 2 controls, single center



Economic Payor Benefit

Cost Effective Incremental Cost-Effectiveness Ratio (ICER) **SURGERY** Hip ~\$10K Knee ~\$12K SI Joint ~\$13K Disc Herniation ~\$21K **Spinal Stenosis** ~\$59K Degen Spondy

~\$64K

Payor Savings

\$3,100

Savings per patient when SI joint is included in diagnosis of low back pain



Worker Productivity

Worker productivity improves with iFuse surgery



Rapid Breakeven

2.5 YEARS Outpatient surgery breakeven

\$83 PER MONTH Post-op LBP cost

Optum Study

Accepted for publication August 2018



Health Technology Positive Appraisals







EXCLUSIVE









MAY





iFuse EXCLUSIVE





MAY 2016 **iFuse EXCLUSIVE** **APRIL** 2017

iFuse Only RECOMMENDATION OCTOBER 2018



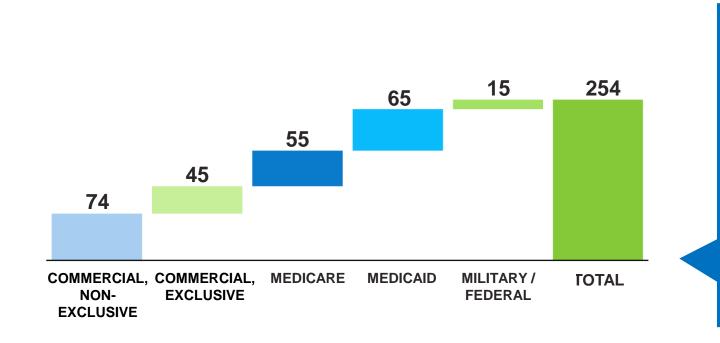




eviCore healthcare

OCTOBER 2018 **iFuse EXCLUSIVE**

Coverage Highlights



119M covered by private payors

All 7 Medicare MACs cover

Growing exclusive iFuse private coverage policies

24 in the U.S.

COVERED LIVES

133 MILLION 162 MILLION

2017

254
MILLION

2016

SEPTEMBER 30, 2018

Commercial Execution

ROBUST CLINICAL DATA

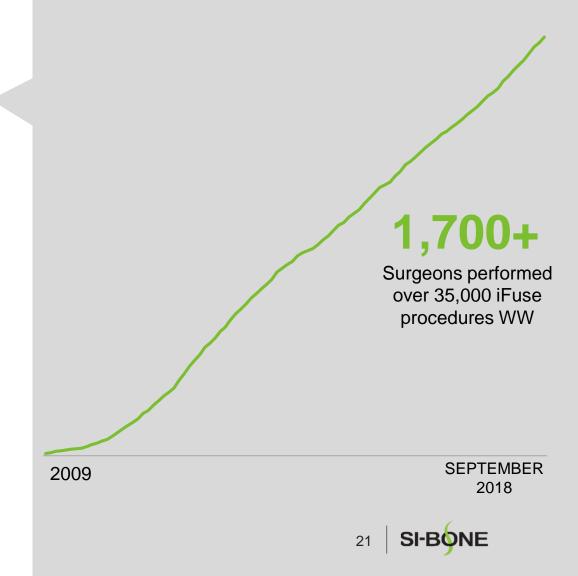
EDUCATION

PHYSICIAN CONFIDENCE

REIMBURSEMENT EXCLUSIVITY

EXPANDING ADOPTION

- New surgeons trained
- Deeper penetration in existing accounts



Diagnostic Algorithm Acceptance and Adoption

Results are as good or better than rules for other lumbar spine conditions

PATIENT HISTORY



PROVOCATIVE TESTS



Series of simple physical exams that lead to an accepted diagnosis

LOCAL ANESTHETIC INJECTION





MEDICARE (MACs)



PRIVATE PAYORS



JAPANESE SACROILIAC JOINT RESEARCH SOCIETY

Surgeon and HCP Education

- Focus on training 7,500 U.S. spine surgeons
- 196 SI-BONE consultants educating HCPs
- Additional Programs
 - Referring community
 - Mid-level/PA
 - Advanced courses

SHIFT

SOCIETY CME DRIVEN EDUCATION















U.S. Commercial Execution

Market and sell through direct sales force and select 3rd party agents

46 DIRECT SALES REPS

- Deep relationships in spine and pain
- Leverage expertise to create referral channels
- Focused technical sales force
- Supported by:
 - Clinical Support Specialists
 - Field Marketing Specialists

>30 SELECT INDEPENDENT AGENTS



Note: As of September 30, 2018

OUS Commercial Execution

29 EUROPEAN EMPLOYEES

INTERNATIONAL SALES FORCE

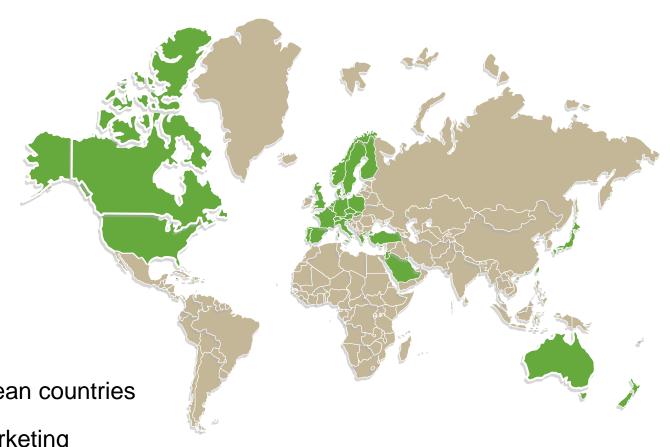
15 DIRECT SALES REPS

21 EXCLUSIVE DISTRIBUTORS

33 COUNTRIES

Focused expansion in major European countries

- UK NICE: recommendation and marketing
- Fédération Hospitalière de France



Expanding and Strengthening SI-BONE

2017

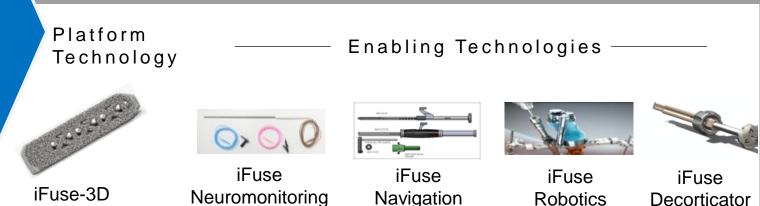
ONE PRODUCT COMPANY

2009 - 2016



SACRO-PELVIC SOLUTIONS COMPANY

2017 - 2018



2017

2018

2018

2018

Intellectual Property Overview

- 40 issued patents: U.S. (34), OUS (6)
- 18 pending patents: U.S. (11), OUS (7)

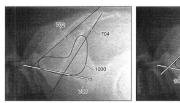
- iFuse patents cover until August 2024
- iFuse-3D patents cover until September 2035

SHAPE



Joint ... fused ... a rectilinear bone fusion implant ... across the joint

APPROACH



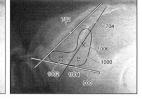
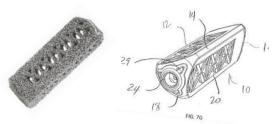


FIG. 10

Lateral insertion path through the ilium and into the sacrum. A postero-lateral insertion path angling through the SI joint.

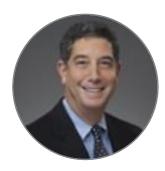
3-D TECHNOLOGY



Fenestration is offset from both the distal end and the proximal end. One repeating internal portion comprising a plurality of apex struts.

Experienced Team

Proven track records



Jeffrey Dunn Chairman, President, CEO & Founder 7 time CEO

TOPEZ, Inc. REILEY ORTHOPEDICS



Chief Financial Officer

PRUKER PWC

McKinsey&Company



Tony Recupero
Chief Commercial Officer

*USSC KYPHONAMEAD OF THE CURVEBAXAND

Mike Pisetsky

General Counsel & Chief Compliance Officer





Scott Yerby

CTO & VP, Engineering





W. Carlton Reckling, MD

Chief Medical Officer

Private practice for 20 years

Joe Powers

VP, Marketing







Nikolas Kerr

VP, Product Marketing





Daniel Cher, MD

VP, Clinical







Andrea Mercanti

VP, EMEA Operations







Troy Wahlenmaier

VP, US Sales







Financial Highlights

Annual Revenue (millions)

GROWTH 15% \$55 - \$55.4 \$48.0 \$42.1 2016 2017 2018E

YTD Revenue (millions)¹



- \$14 million in cash¹
- 91% gross margins¹
 - Revenue
 - Operating Loss

Future Growth Drivers

1. GROW REIMBURSEMENT COVERAGE

- Coverage from remaining major U.S. commercial payors
- More exclusive U.S. policies
- Reimbursement coverage in Japan

2. EXPAND SALES ORGANIZATION

- Increase direct field sales organization
- Add to field support organization
- Increase rep productivity

3. ROLLOUT NEXT GENERATION PRODUCTS

- Build off iFuse-3D introduction
- Launch products in adjacent markets



Investment Highlights

4,300 US PROCEDURES IN 2017

ROBUST **DATA**

>60 PUBLISHED PAPERS

2 RANDOMIZED TRIALS

REIMBURSEMENTADVANTAGE

250+
MILLION
COVERED LIVES

23 PAYOR EXCLUSIVES

POSITIVE FINANCIAL PROFILE

16% GROWTH²

\$53.5M REVENUE¹
91% GROSS
MARGINS²

279,000 US POTENTIAL PATIENTS PER YEAR

LTM revenue as of September 30, 2018
 YTD as if September 30, 2018

SI-BONE

Disclosures

The iFuse Implant System is intended for sacroiliac fusion for conditions including sacroiliac joint dysfunction that is a direct result of sacroiliac joint disruption and degenerative sacroilitis. This includes conditions whose symptoms began during pregnancy or in the peripartum period and have persisted postpartum for more than 6 months.

There are potential risks associated with the iFuse Implant System. It may not be appropriate for all patients and all patients may not benefit. Risk information available upon request.

One or more of the individuals named herein may be past or present SI-BONE employees, consultants, investors, clinical trial investigators, or grant recipients. Research described herein may have been supported in whole or in part by SI-BONE.

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