

UNITED STATES
SECURITIES AND EXCHANGE COMMISSION

Washington, D.C. 20549

FORM 8-K

CURRENT REPORT

Pursuant to Section 13 or 15(d) of The Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): September 8, 2026

INTEGRA LIFESCIENCES HOLDINGS CORPORATION

(Exact Name of Registrant as Specified in its Charter)

Delaware	0-26224	51-0317849
(State or Other Jurisdiction of Incorporation or Organization)	(Commission File Number)	(IRS Employer Identification No.)

1100 Campus Road
Princeton, NJ 08540
(Address of principal executive offices) (Zip Code)
Registrant's telephone number, including area code: (609) 275-0500

Not Applicable
(Former name or former address, if changed since last report)

Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions (see General Instruction A.2. below):

- ☐ Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425).
- ☐ Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12).
- ☐ Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b)).
- ☐ Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c)).

Securities Registered Pursuant to Section 12(b) of the Act:

Title of Each Class	Trading Symbol	Name of Exchange on Which Registered
Common Stock, Par Value \$.01 Per Share	IART	Nasdaq Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).

Emerging growth company ☐

If an emerging growth company, indicate by check mark if the registrant has elected not to use the extended transition period for complying with any new or revised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. ☐

Item 7.01 Regulation FD Disclosure.

Integra LifeSciences Holdings Corporation (the “Company”) hereby furnishes the updated investor presentation attached as Exhibit 99.1 to this Current Report on Form 8-K, which the Company may use, from time to time, in presentations to existing or prospective lenders in connection with the Company’s previously-disclosed plans to refinance its outstanding senior credit facility in 2026.

The information in this Item 7.01, including Exhibit 99.1, shall not be deemed “filed” for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the “Exchange Act”) or otherwise subject to the liability of such section, nor shall it be deemed incorporated by reference in any filing of the Company under the Securities Act of 1933, as amended, or the Exchange Act, regardless of any general incorporation language in such filing, unless expressly incorporated by specific reference in such filing.

Item 9.01 Financial Statements and Exhibits.

(d) Exhibits

99.1 [Investor Presentation dated September 8, 2026](#)

104 Cover Page Interactive Data File (embedded within the inline XBRL document)

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned hereunto duly authorized.

INTEGRA LIFESCIENCES HOLDINGS CORPORATION

Date: September 8, 2026

By: /s/ Lea Knight

Lea Knight

Title: Executive Vice President and Chief Financial Officer

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These materials are only available to QIBs and non-US persons.

Investor Presentation

September 2026

This presentation includes information that (i) is publicly available (or could be derived from publicly available information) and (ii) does not constitute material nonpublic information with respect to the company, its subsidiaries or its or their respective securities.



Safe Harbor Statement

This presentation contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995 that involve risks and uncertainties and reflect the Company's judgment as of the date of this presentation. All statements, other than statements of historical fact, are statements that could be deemed forward-looking statements. Some of these forward-looking statements may contain words like "will," "believe," "may," "could," "would," "might," "possible," "should," "expect," "intend," "forecast," "guidance," "plan," "anticipate," "target," or "continue," the negative of these words, other terms of similar meaning or they may use future dates. Forward-looking statements contained in this presentation include, but are not limited to, statements concerning: the future business, operational and financial performance of the Company and the Company's expectations and plans with respect to market opportunity, business and operational performance, strategic initiatives, capabilities, resources, manufacturing capabilities, product development, product availability and regulatory approvals, including expectations regarding the Company's compliance master plan to improve the Company's quality systems. It is important to note that the Company's goals and expectations are not predictions of actual performance. Such forward-looking statements involve risks and uncertainties that could cause actual results to differ materially from predicted or expected results. Such risks and uncertainties include, but are not limited to, the following: increased geopolitical tension, instability and other macroeconomic factors, including trade barriers and related restrictions (including tariffs and related countermeasures), armed conflict and acts of terrorism, supply chain disruptions, and interest rate and foreign currency rate fluctuations on the Company's suppliers, vendors and customers and on the Company's business and financial condition, results of operations and cash flows; the Company's ability to execute its financial, strategic and operating plans effectively; the Company's ability to remediate quality systems violations; difficulties in implementing the Company's compliance master plan; difficulties or delays in obtaining and maintaining required regulatory approvals, including the costs there of potential difficulties, delays and disruptions in manufacturing, distribution or sale of products; the Company's ongoing assessment of the flooding event at its Cincinnati, Ohio manufacturing facility, related financial impacts, recovery efforts, potential assets impairments, unforeseen costs and insurance recoveries; the failure of the company's suppliers, vendors, and other third parties to meet contractual, regulatory and other obligations; the anticipated development of markets the Company sells its products into and the success of the Company's products in these markets; the Company's ability to predict accurately the demand for its products, and products under development; increasing industry competition; the coverage and reimbursement decisions of third-party payors; trends toward healthcare cost containment; difficulties in controlling expenses, including costs to procure and manufacture the Company's products; the ability of the Company to successfully manage leadership and organizational changes and the impact of changes in management or staff levels; the impact of goodwill and intangible asset impairment charges if future operating results of acquired businesses are significantly less than the results anticipated at the time of the acquisitions, the geographic distribution of where the Company generates its taxable income; changes to applicable laws, regulations and enforcement guidance, including tax laws and global healthcare reforms; fluctuations in foreign currency exchange rates; the amount of our bank borrowings outstanding and other factors influencing liquidity; breaches, failures or other disruptions of our or our vendors' or customers' information technology systems or products; the potential impact of our compliance with governmental regulations and accounting guidance; and the economic, competitive, governmental, technological, and other risk factors and uncertainties identified under the heading "Risk Factors" included in Item 1A of Integra's Annual Report on Form 10-K for the year ended December 31, 2025 and information contained in subsequent filings with the Securities and Exchange Commission.

These forward-looking statements are made only as of the date hereof, and the Company undertakes no obligation to update or revise the forward-looking statements, whether as a result of new information, future events, or otherwise, except as otherwise required by law.

Non-GAAP Financial Measures

In addition to our GAAP results, we provide certain non-GAAP measures, including organic revenues, adjusted earnings before interest, taxes, depreciation and amortization ("EBITDA"), adjusted EBITDA margin, adjusted gross profit, adjusted gross margin, free cash flow and net debt. Organic revenues consist of total revenues excluding the effects of currency exchange rates, revenues from current-period acquisitions and product divestitures. Adjusted EBITDA consists of GAAP net income excluding: (i) depreciation and amortization; (ii) other income (expense); (iii) interest income and expense; (iv) income tax expense (benefit); (v) impairment charges; and (vi) those operating expenses also excluded from adjusted net income. The measure of adjusted EBITDA margin is calculated by dividing adjusted EBITDA by GAAP revenues. Adjusted gross profit consists of GAAP gross profit adjusted for: (i) structural optimization charges; (ii) divestiture, acquisition and integration-related charges; (iii) charges related to the transition of Boston-related manufacturing operations to the Company's Braintree, Massachusetts facility; (iv) EU Medical Device Regulation-related charges; and (v) intangible asset amortization expense. The measure of free cash flow consists of GAAP net cash provided by operating activities less purchases of property and equipment. The measure of net debt consists of GAAP total debt (excluding deferred financing costs) less short-term investments, cash and cash equivalents.

Reconciliations of (i) GAAP net income to adjusted EBITDA for the years ended December 31, 2025, 2024, 2023, 2022 and 2021, and for the four quarters and last twelve months ended June 30 2026, (ii) GAAP gross profit to adjusted gross profit, GAAP gross margin to adjusted gross margin, and GAAP operating cash flow to free cash flow for the years ended December 31, 2025, 2024, 2023, 2022 and 2021, and for the last twelve months ended June 30 2026, appear in the Appendix in this presentation.

The Company believes that the presentation of non-GAAP measures provide important supplemental information to management and investors regarding financial and business trends relating to the Company's financial condition and results of operations.

No Offer

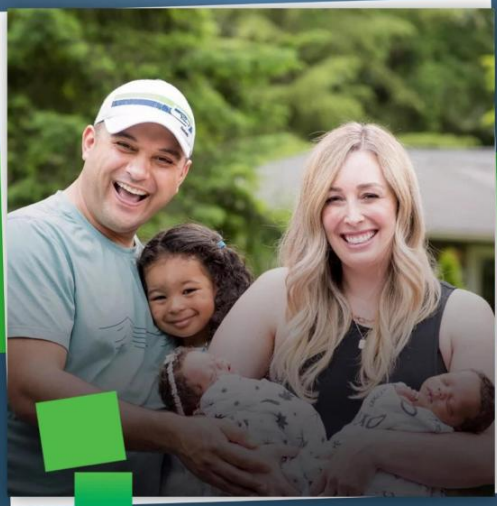
This presentation does not constitute an offer to sell, or the solicitation of an offer to buy, any security, and shall not form the basis of any contract. Any offering of securities will be made solely by means of a separate offering memorandum or other definitive offering document and in accordance with applicable securities laws.

Executive summary

- **Integra Lifesciences Holdings Corporation ("Integra" or the "Company") is a global MedTech leader that engages in the manufacture and sale of medical devices specializing in neurosurgery and regenerative tissue repair through its two segments:**
 - **Specialty Surgery:** provides neurosurgical and neurocritical care devices used across the continuum of care—from pre-operative planning and intraoperative neurosurgery to post-operative critical care—for adult and pediatric patients with traumatic brain injury, hydrocephalus, cerebrovascular conditions, brain tumors, and select ENT related disorders
 - **Tissue Reconstruction:** provides collagen-based and regenerative tissue technologies used in the treatment of acute and chronic wounds, burns, and surgical tendon and ligament repair
- **Integra views 2026 as a key inflection period as it progresses through FDA-mandated manufacturing / quality system remediations, with several operational tailwinds expected to support future growth**
 - All warning letter commitments on track to be completed in 2026
 - SurgiMend expected to relaunch in Q4 2026
 - New state-of-the-art manufacturing site in Braintree, MA ("Braintree") became operational in Q2 2026; on track for commercial revenue being generated in Q4 2026
 - Remaining remediation / compliance spend expected to decrease
- **Key financials highlights**
 - LTM Q2 2026 revenue of \$1,648M, up 1.6% YoY, and Adjusted EBITDA of \$337M, up 11.8% YoY, representing a 20.5% margin
 - YoY FCF growth of \$57M

01

Business overview



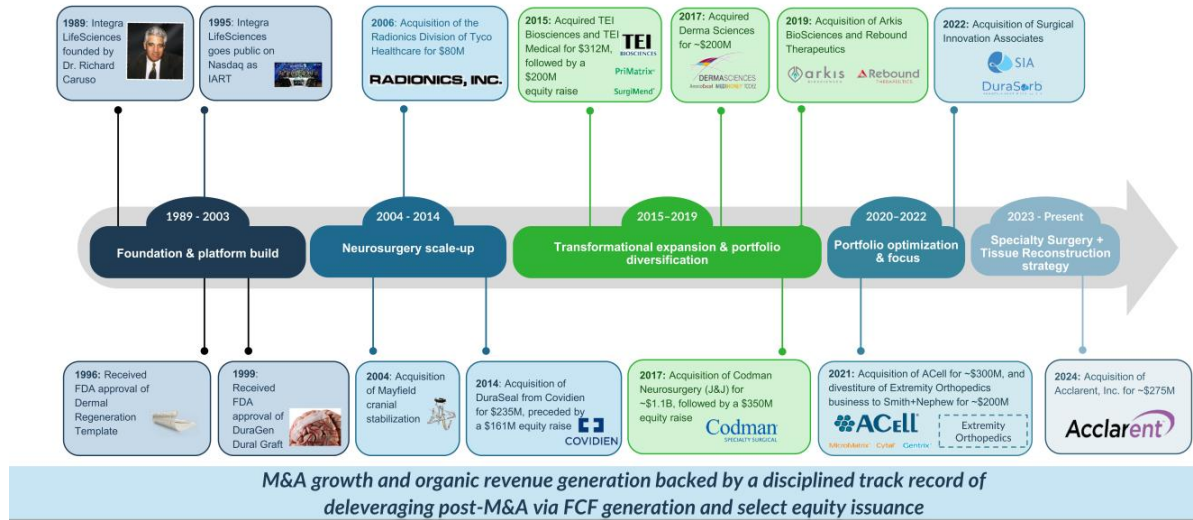
INTEGRA 

A global MedTech leader dedicated to restoring lives



Financial metrics shown for the trailing 12-months ended Jun. 30, 2026
Both adj. gross margin and adj. EBITDA margin are non-GAAP financial measures; See appendix for non-GAAP reconciliations

Evolution of Integra: Building a scaled specialty platform



Where we play

Leading player in attractive, niche markets through focus



¹ TAM excludes private label markets; ² LTM revenue as of 06/30/26

How we play

Leadership in attractive, niche markets through differentiated technologies and commercial excellence

OUR ADVANTAGED POSITION



Unique focus and #1-2 player in attractive markets



Depth and breadth of portfolio and technologies



Tenured, trusted salesforce and channels



Global footprint for technology access and commercialization



Flexible balance sheet with disciplined capital management



Organic and inorganic innovation aimed at accretive growth segments

Growing patient population in chronic disease and emerging markets

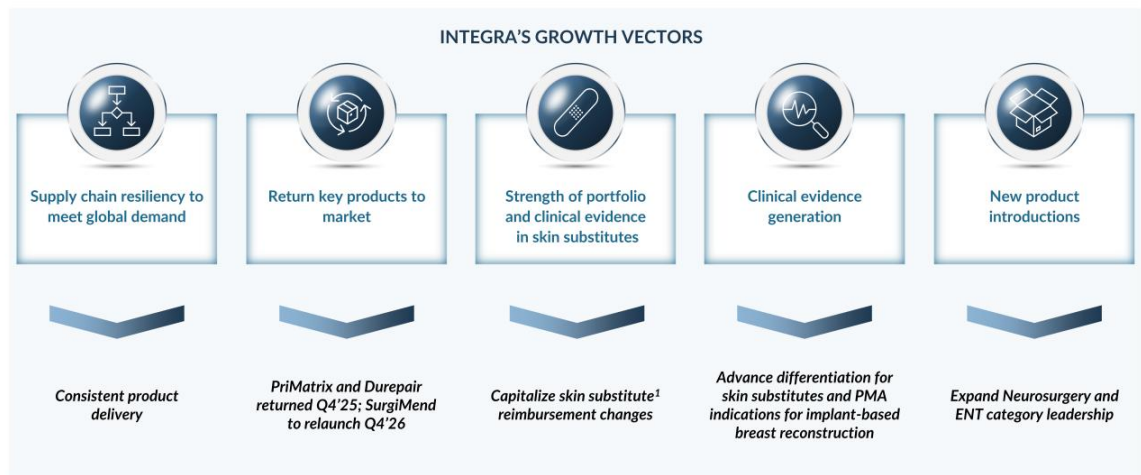
Differentiated technologies and innovation matter

Attractive high growth niche opportunities in core markets

Surgeons have higher influence in product decisions

Building the foundation for sustainable growth

Focused execution enabling innovation and growth



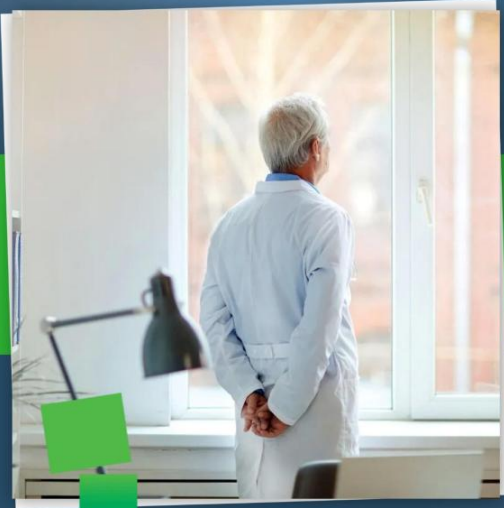
¹ CMS refers to skin substitute as CTP = Cell & Tissue-Based Products

Delivering our vision in 2026 and beyond

Build a culture of quality and execution excellence, solidify market leadership, drive company performance



02

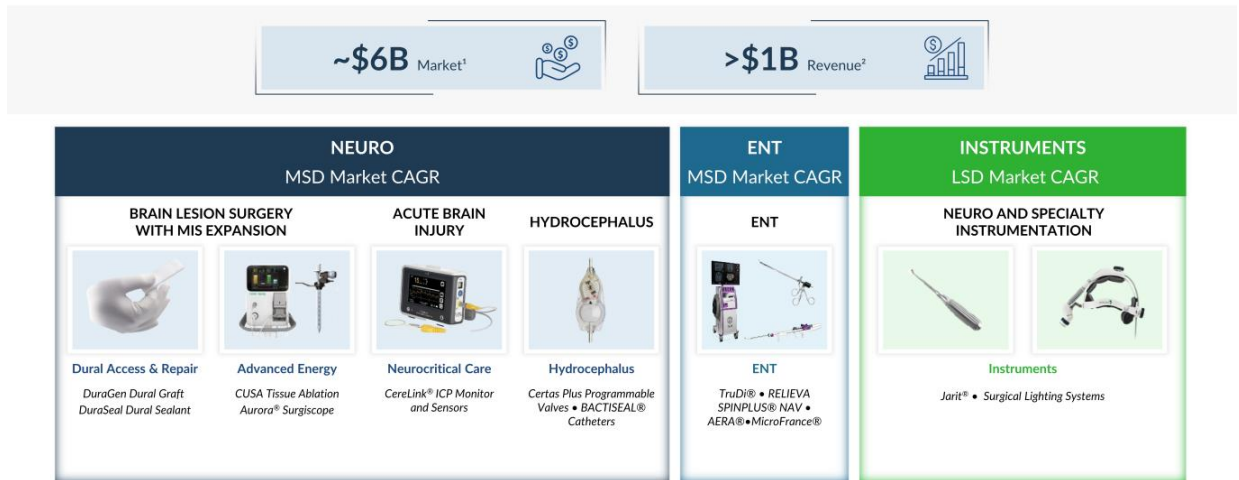


Specialty Surgery

INTEGRA 

Specialty Surgery overview

Established leadership in served markets, portfolio positioned to maintain stable market growth



Note: Market size and growth from third party market reports and internal estimates; ¹ TAM for ENT is US only; ² LTM revenue as of 06/30/26; MIS = Minimally Invasive Surgery; ICP = Increased Intracranial Pressure

Neuro

Comprehensive portfolio of high-utilization products that underpin neurosurgery procedures

DuraGen® Dural Graft Matrix	DuraSeal® Dural Sealant System	BACTISEAL® Catheters	CUSA® Clarity	CereLink® ICP Monitor	Certas® Plus Programmable Valve	Aurora® Surgiscope System & MIS	Mayfield® Cranial Stabilization
							
Absorbable dural graft matrix for sutureless repair of dural defects	Hydrogel sealant to provide watertight closure around suture lines	Antibiotic-impregnated catheter to reduce CSF infection risk	Platform used for fragmentation and aspiration of soft and hard tissues	Advanced digital ICP monitoring platform for neuro-intensive care	Implantable programmable valve for managing hydrocephalus	Tubular retractor system to manage intracerebral hemorrhage	Surgical device to stabilize the patient's head during procedures



1

Addresses complex pathologies and disease states in neurosurgery and surgical adjacencies

2

Portfolio spans the entire continuum of care from pre-operative stabilization to post-operative neuro-critical care

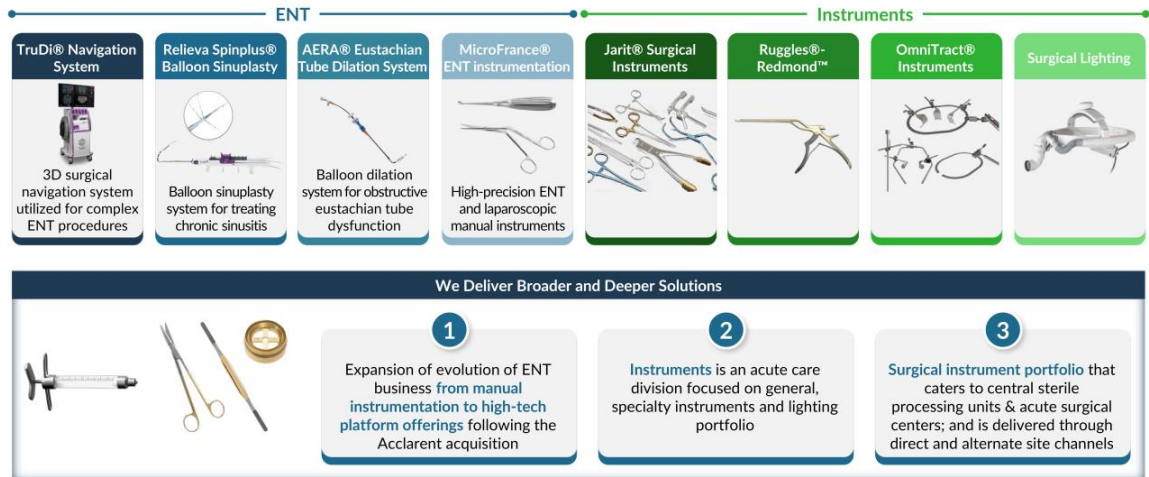
3

Industry-leading sales channel that offers a full portfolio of solutions to specialized customers and surgeons at multiple care sites

CSF = Cerebrospinal Fluid; ICP = Intracranial Pressure; MIS = Minimally Invasive Surgery

ENT and Instruments

High-precision ENT and laparoscopic manual instruments



Integra is #1 or #2 in most categories in which we compete

Global market position and trend by platform

	Dural Access & Repair	Advanced Energy	Neurocritical Care	Hydrocephalus	ENT ¹	Instruments
INTEGRA Market Position	#1	#2	#1	#1	#3	#1
LTM Q2'26A Revenue ²	\$0.3B	\$0.2B	\$0.1B	\$0.2B	\$0.1B	\$0.2B
Other Players						

Durable hospital and neurosurgeon relationships that sustain market leadership

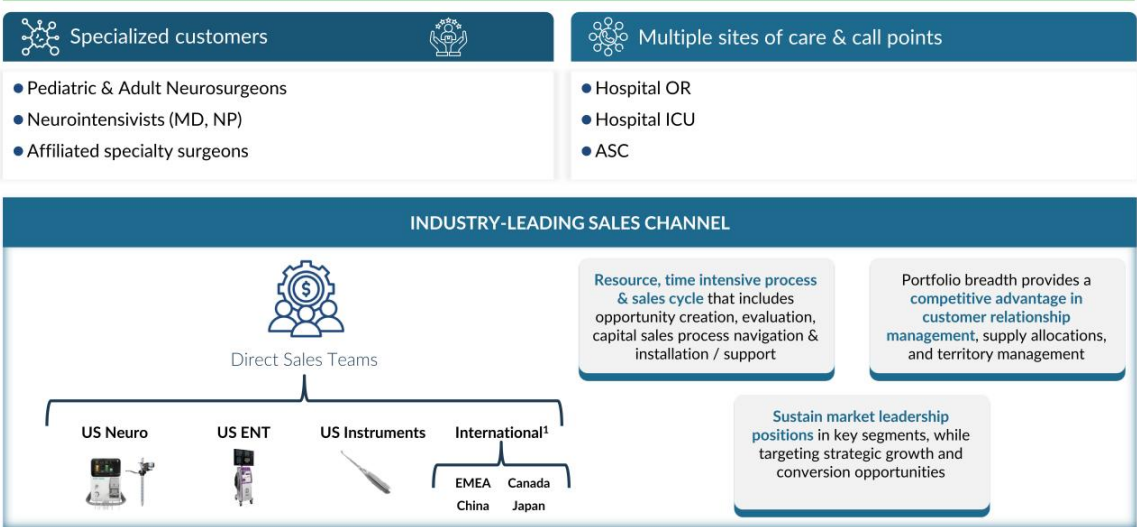
¹ US position for core ENT market; ² LTM revenue as of 06/30/26

Accelerate growth

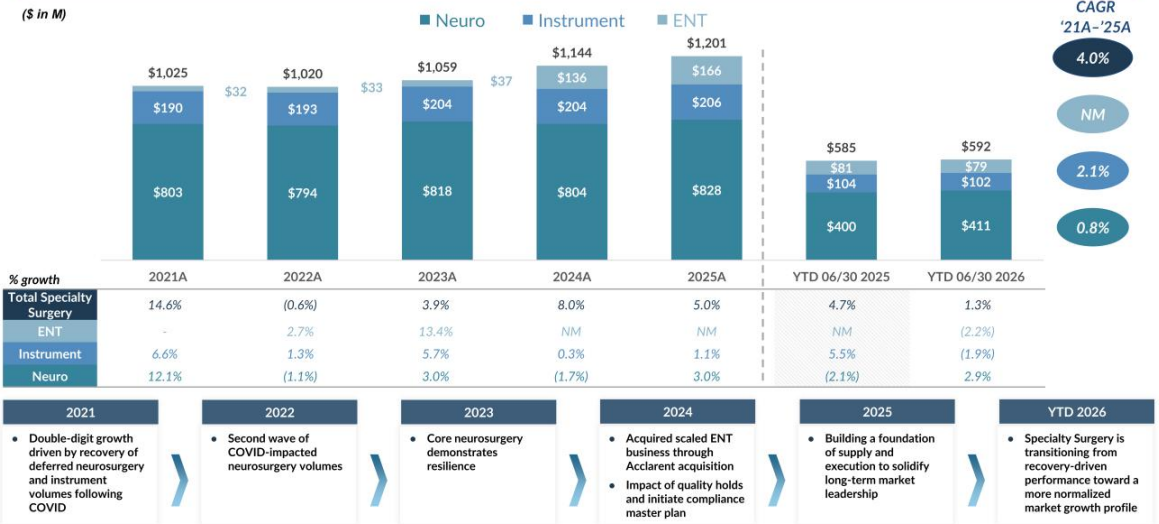
Prioritizing longer-term organic growth



Specialty Surgery: Addressing complex pathologies in neurosurgery & surgical adjacencies



Category-leading technologies fueled by scale and resilience



03

Tissue Reconstruction



INTEGRA 

Tissue Reconstruction Overview

Poised for Growth in Attractive Global Markets

~\$2.5B Market 

~\$0.4B Revenue¹ 

~90% Of sales are to the hospital and surgery center setting 

WOUND RECONSTRUCTION ("WR") MSD Market CAGR

COMPLEX WOUNDS, BURNS, TRAUMATIC, CHRONIC WOUNDS, NERVE AND TENDON REPAIR

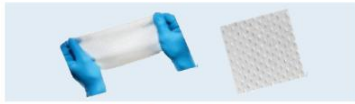


Wound Reconstruction Solutions

Integra® Dermal Matrices • PriMatrix • MicroMatrix • Cytal®
AmnioExcel® • MediHoney® • TCC-EZ • NeuroGen

SOFT TISSUE SOLUTIONS MSD to HSD Market CAGR

SOFT TISSUE AND MUSCLE FLAP REINFORCEMENT, BREAST RECONSTRUCTION, HERNIA REPAIR

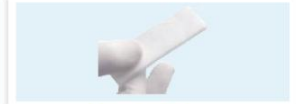


Soft Tissue Solutions

SurgiMend • DuraSorb® • Gentrix®

PRIVATE LABEL MSD Growth

B2B PARTNERSHIPS WITH SELECT COLLAGEN TECHNOLOGIES



Private Label Partners

Note: Market size and growth from third party market reports and internal estimates; ¹ LTM revenue as of 06/30/26

Wound Reconstruction Solutions

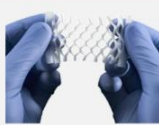
Comprehensive portfolio to treat complex wounds

Engineered Collagen
(Integra®)



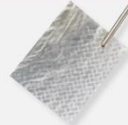
Helps Restore Lost *Function* and Joint Mobility

Fetal Bovine Collagen
(PriMatrix®)



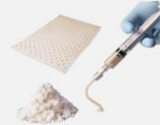
An *Adaptable* Solution for Challenging Wounds

Placental Allograft
(AmnioExcel®)



Strong Enough to go the Distance

UBM
(MicroMatrix® & Cytal®)



Addressing Challenging Wound Microenvironments

We Deliver Broader and Deeper Solutions



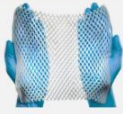
- 1
Complex wound reconstruction can be difficult, *surgeons need choices* to address how wounds may be uniquely presenting
- 2
Integra LifeSciences offers a *broad portfolio* addressing a full spectrum of needs when managing complex wounds
- 3
Integra's specialized wound care *sales team* delivers a full portfolio to meet surgeon and patient needs

UBM = Urinary Bladder Matrix

Soft Tissue Solutions

Comprehensive portfolio for soft tissue reinforcement and repair

**SurgiMend
Biological Matrix**



Source Material: Fetal Bovine Dermis
Category: Xenograft Biologic
Meshed/Fenestrated and Surgical/Hiatal configurations

**DuraSorb®
Monofilament Mesh**



Source Material: Polydioxanone
Category: Bioresorbable Synthetic
Macroporous/Monofilament Mesh

**Gentrix®
Surgical Matrix**



Source Material: UBM
(Urinary Bladder Matrix)
Category: Xenograft Tissue Scaffold
Surgical/Hiatal configurations

We Deliver Broader and Deeper Solutions



- 1
Flagship collagen-based products exhibit biological superiority and are associated with fetal-like healing, rapid tissue remodeling, and superior elasticity
- 2
Broad portfolio that covers high-incidence surgical needs from complex ventral hernia reconstruction to specialized breast procedures
- 3
Pursuing "first-mover" advantage in the IBBR market by seeking formal PMA approval for both its biologic and synthetic platforms

IBBR = Implant-Based Breast Reconstruction; PMA = Pre-market approval; UBM = Urinary Bladder Matrix

Integra is #1 in Wound Reconstruction and an emerging competitor in Soft Tissue Solutions

Global market position and trend by platform (excl. private label¹)

	Wound Reconstruction Solutions	Soft Tissue Solutions
INTEGRA Market Position	#1	#7
LTM Q1'26 Revenue	\$0.3B	<\$0.1B²
Other Players	SmithNephew LifeNet Health ³ ORGANOGENESIS kerecis ⁴ MIMEDx	abbvie / LifeCell BD mtfbiologics stryker

We continue to be a top key market player in the tissue reconstruction subsegments

Note: Tissue Reconstruction market positions are hospital-based and do not include office-based procedures
¹ Q1'26 LTM revenue of \$113M; ² Excludes SurgiMend®; anticipated relaunch in 4Q'26

Accelerate growth

Prioritizing four strategies to deliver innovation and drive organic growth

Tissue Reconstruction

Implant Based Breast Reconstruction

Drive adoption via dual-PMAs of
SurgiMend and DuraSorb

SurgiMend and DuraSorb PMAs
expected to be approved in 2027

Wound Reconstruction Outpatient (OP)

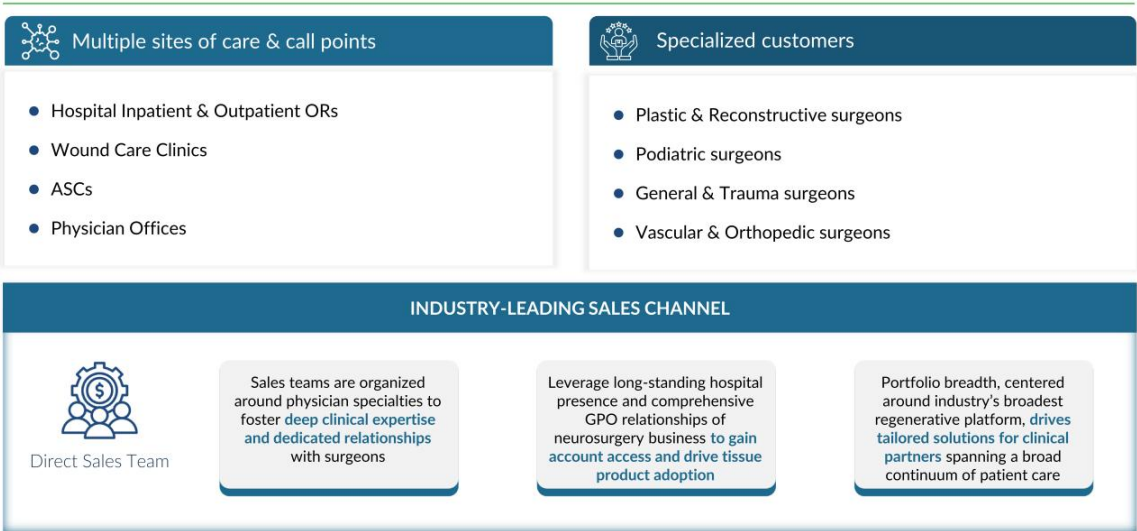
Capitalize on CMS reimbursement
changes

PriMatrix & SurgiMend

Rapidly recover pre-recall revenue
and grow
market share

UBM = Urinary Bladder Matrix

Tissue Reconstruction: Leading the way in regenerative and tissue technologies



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CMS Reimbursement Reform

2026 CMS Wound Care Reimbursement Reform is expected to be a Net Positive for Integra

What CMS Changed (Context)

CMS significantly reduced reimbursement rates for skin substitutes beginning January 1, 2026, from an **average ASP of ~\$1,300/cm²** (across 200+ skin substitutes) to **~\$127/cm²**

Reforms are intended to **address overuse and excessive reimbursement** of skin substitutes in physician office settings

New economics and oversight should result in procedures shifting **out of physician offices and into inpatient (hospital) settings**

Manufacturers with significant physician office volume face meaningful margin pressure under the new reimbursement economics

Why This Benefits Integra (Impact)

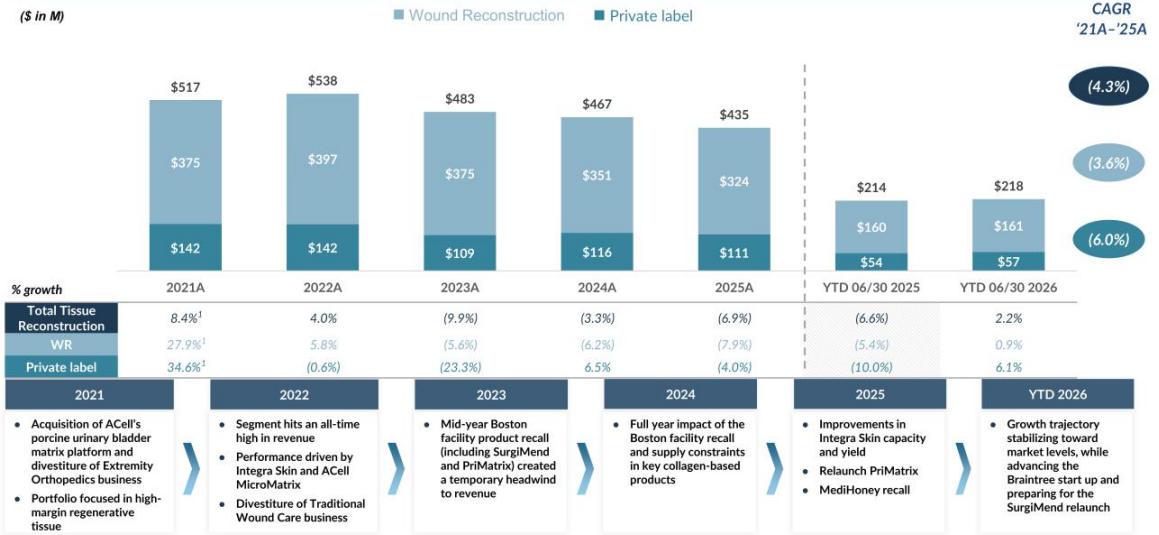
Integra's portfolio is **already priced under ~\$127/cm²** – no price or margin risk on existing or incremental volume

Integra is well positioned commercially to **capture volume shifting away from physician offices**

~90% of Integra's revenue is hospital based, reimbursed under a diagnosis-related group (DRG) and **not exposed to the January 1 CMS reimbursement changes**

Broad portfolio, strong clinical evidence, and appropriate pricing **uniquely positions Integra across inpatient and outpatient settings**

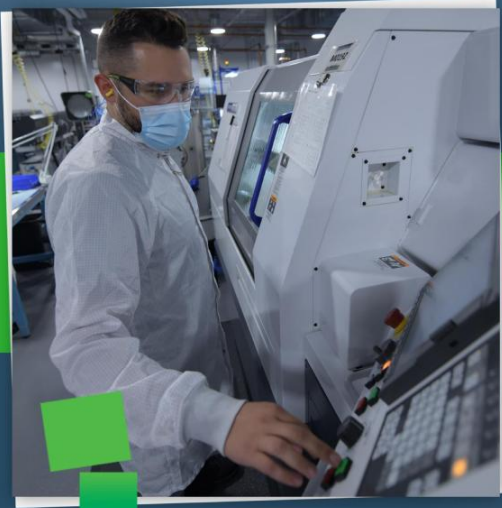
2026 recovering from previous operational headwinds



¹ 2021A growth rates reflect a recovery from pandemic-related contractions in 2020 due to global surgical procedure deferrals

04

Supply Chain / Operations



Global manufacturing and distribution footprint

Global network that supports broad technology portfolio throughout patient journey



Overview of state-of-the-art facilities

ANASCO, PUERTO RICO



- Competencies: Collagen manufacturing & packaging, device assembly & packaging
- ISO7/ISO8 Clean rooms
- Space: ~80k sq. ft.
- Areas of focus: Dural Access & Repair, Regenerative Neurosurgery

PLAINSBORO, NEW JERSEY



- Competencies: Collagen manufacturing & packaging
- ISO7 Clean rooms
- Space: ~84k sq. ft.
- Areas of focus: Advanced Wound Care, Dermal Regeneration

LE LOCLE, SWITZERLAND



- Competencies: Micro molding & assembly, antibiotics impregnation, packaging
- ISO8 Clean rooms
- Space: ~80k sq. ft.
- Areas of focus: Cerebral Spinal Fluid (CSF) Management, Neurosurgical Monitoring

BRAINTREE, MASSACHUSETTS

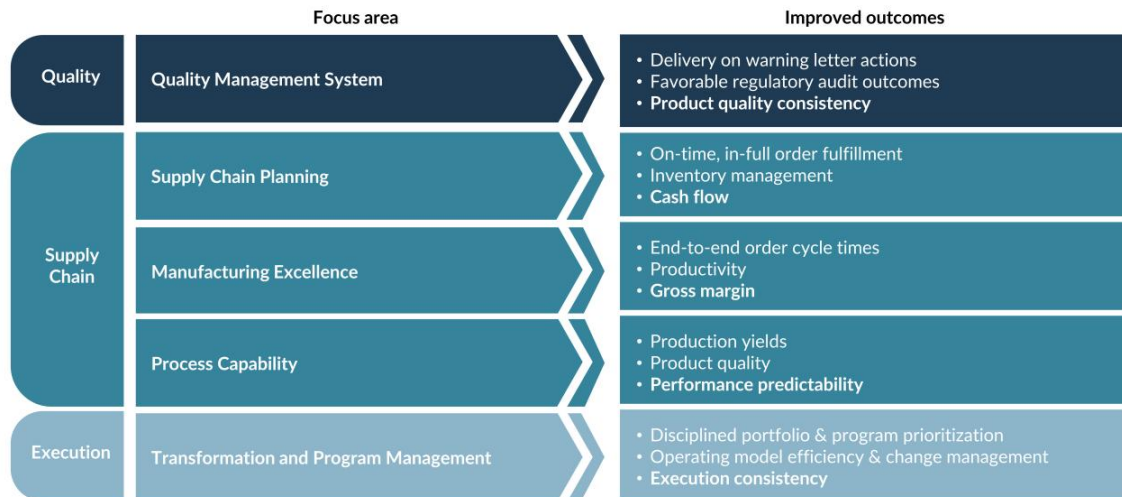


- Competencies: Bovine Matrix processing & packaging
- ISO8 Clean rooms
- Space: ~107k sq. ft.
- Areas of focus: Surgical Reconstruction, Plastic & Reconstructive Surgery

Operational since June 2026

A broad manufacturing footprint with distinct, specialized capabilities creates a durable competitive advantage

Key Capabilities to Drive Operations and Customer Excellence



Summary of global operations opportunity

Milestones

Mid-60%

Return to historical GM

95%+

sustained service levels

5%+

productivity improvement
annually

1

Return to Historical GM
Enhanced gross margins and
improved cashflow

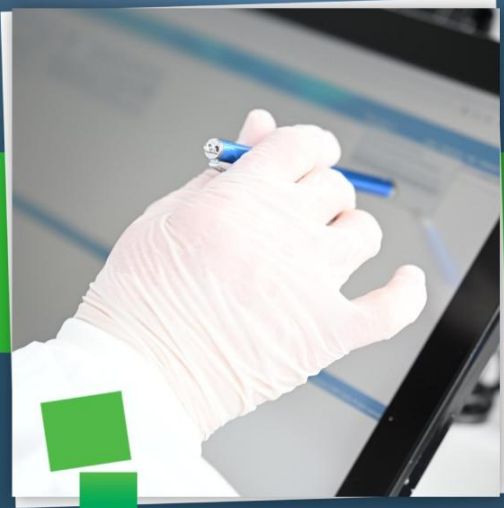
2

Sustained Service Levels
Product quality consistency and
performance predictability

3

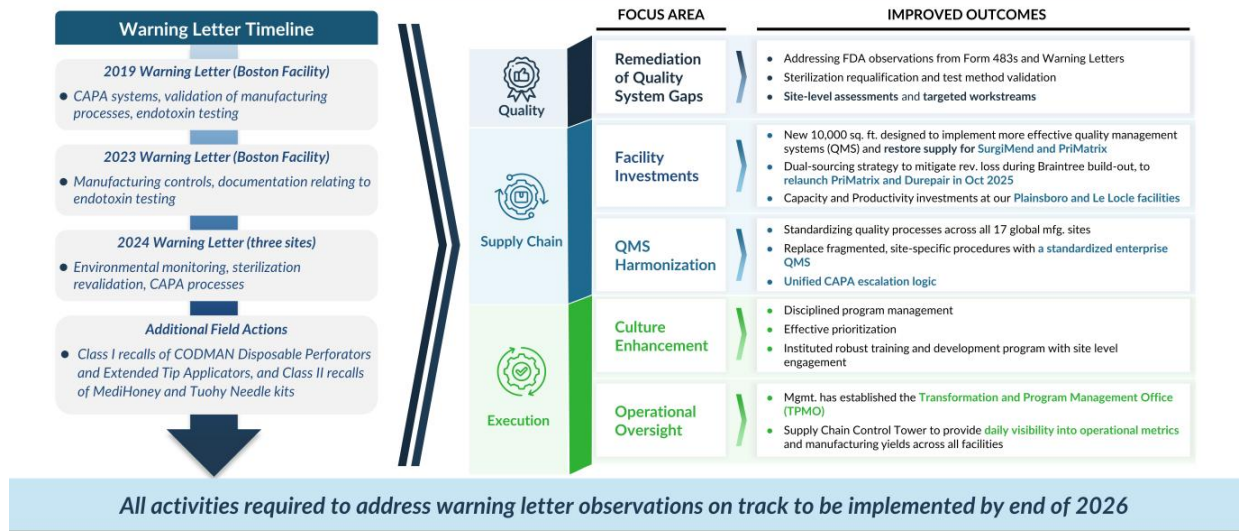
**Productivity Improvement
Annually**
Improved production yields and
order cycle items

05



Remediation update

Beyond Remediation: Building a Stronger Quality and Operating Foundation



Path to full product reintroduction

■ Relaunch completed

Products	Current Status
 Cranial perforators	✓ Relunched
 PriMatrix® Dermal Scaffold	✓ Relunched
 Durepair® Dura Regeneration Matrix	✓ Relunched
 SurgiMend®	Anticipated relaunch in 4Q'26 from the Braintree, MA facility
 CODMAN® Cottonoid® Surgical Strips and Patties	Remediation in process to support release with streamlined product portfolio
 MediHoney® Wound & Burn Dressing	Remediation in process with 3rd party manufacturer to support 2027 release

All products, which historically represented ~\$150mm of annual revenue, are expected to be reintroduced to market by the end of 2027

Braintree: restoring supply and enabling recovery

A new, purpose-built site replacing Integra's legacy Boston manufacturing operation — restoring in-house supply of key products, with start-up delivered on the committed timeline

BRAINTREE, MASSACHUSETTS

Operational & producing since Q2 2026



FROM COMMITMENT TO DELIVERY

Milestone	What management committed to	Status
PriMatrix & Durepair	Restore supply via alternate supplier	✓ Relunched in Q4 2025 — ~12 months ahead of plan
Facility start-up	Braintree online by end of June 2026	✓ Online and producing inventory in Q2 2026 — on time
SurgiMend relaunch	Return SurgiMend to market in Q4 2026	On track: producing inventory to support the Q4 2026 relaunch
Transition-cost wind-down	Reduce transition costs as Braintree ramps	Expect transition costs to decline greater than 50% from 2025 to 2026

Why Braintree matters to the IART story

Restoring supply through disciplined execution

Braintree's on-time start-up marks the completion of a major manufacturing reset, restoring supply and validating the effectiveness of management's recovery plan

Provides the foundation for indication expansion

Braintree will enable the launch of PMA products that will provide a differentiated advantage in implant-based breast reconstruction (IBBR)

Supports the financial recovery

With Braintree operational, remediation activities and associated special charges are expected to decline, supporting earnings, cash flow, and deleveraging

Source: Integra LifeSciences Q2 2026 earnings release and investor presentation (July 29, 2026); Q4 2025 (Feb 2026) and Q1 2026 (May 2026) earnings calls; company filings
Note: ✓ = delivered; remaining milestones on track as noted. Transition cost reflects the Braintree transition adjustment (adjusted net income basis). Dollar figures in USD millions

Key adjustments mapped to Integra's story

Trailing twelve months across meaningful categories

(In \$M)	2024A	2025A	LTM Q2'26A	Q1'26A	Q2'26A	COMMENTARY
EU MDR	44.6	41.9	30.6	7.9	2.4	90% complete, as we continue to reduce headcount of contractors working on EU MDR-related efforts with majority rolled off and Q2 figures reflecting normalized spend
Acquisition Related	33.6	3.6	(3.4)	1.8	2.4	Acclarent integration costs winding down
Structural Optimization	24.2	48.0	48.2	9.3	7.5	Include benefits of \$5.3M, \$20.5M, and \$0.8M in 2024, 2025, and YTD 2026 for contingent consideration adjustments
Braintree Transition	45.0	56.2	45.4	7.7	9.9	Expected to come down in near term, driven by a reduction in idle manufacturing costs incurred during the production remediation period
Operating Non-GAAP Adjustments	\$147.4	\$149.7	\$120.8	\$26.7	\$22.3	Braintree became operational at the end of June 2026, allowing for transition-related expenses to be significantly reduced by end of 2026; on-track to launch SurgiMend and generate commercial revenue in Q4 2026

Substantial reduction between EU MDR and Braintree underpin significant operating cash flow improvement

06



Investment highlights

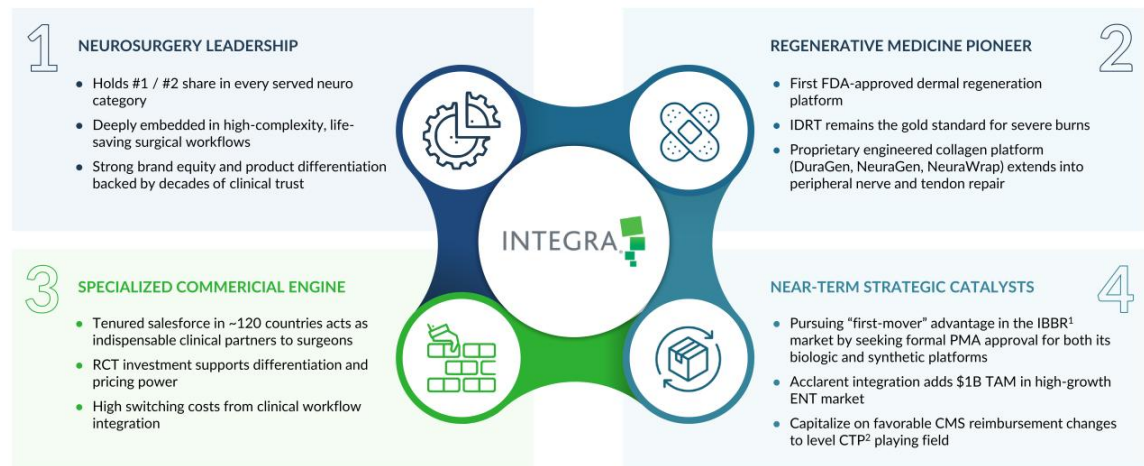
INTEGRA 

Sustained category leadership and operational excellence



1 Global leader in neurosurgery and regenerative medicine

Advancing transformational care, delivering impactful innovation, and enriching life moments



¹ IBBR = Implant-Based Breast Reconstruction; ² CTP = Cell & Tissue-Based Products (i.e. skin substitutes)

2 Large and growing addressable markets



¹ TAM excludes private label markets

3 Diversified portfolio across various specialties

Specialty Surgery		Tissue Reconstruction ¹	
Neuro		Wound Reconstruction Solutions	
 Duragen Dural Graft Matrix		 Integra Dermal Matrices	
 Aurora Surgiscope		 AmnioExcel®	
 CUSA Tissue Ablation		 PriMatrix	
 Duraseal Dural Sealant		 MediHoney	
 CereLink® ICP Monitor and Sensors		 MicroMatrix	
 BACTISEAL Catheters		 TCC-EZ®	
 Certas Plus Programmable Valves		 Cytal®	
 NeuraGen®			
ENT		Instruments	
 TruDi® Navigation System		 Javit	
 RELIEVA SPINPLUS® Balloon Sinuplasty System		 DUO LED Surgical Headlight	
 AERA® Eustachian Tube Dilation System			
 MicroFrance ENT instrumentation			
		Soft Tissue Solutions	
		 SurgiMend	
		 DuraSorb	
		 Gentrix	

¹ Excludes private label

4 Significant brand equity and loyalty

Highly differentiated products trusted by healthcare professionals and systems

Clinically differentiated products

- "First-to-market" innovations, especially in dural repair and ultrasonic tissue ablation
- Outcome reliability establishes brand as a risk-mitigating choice
- High brand recall among neurosurgeons and reconstructive specialists
- Core products function as default selection within key indications



Training-led adoption model

- Incumbency advantage reinforced in time-sensitive surgical settings
- Surgeon muscle memory and technique standardization limit substitution
- Deep adoption across leading academic and tertiary care centers
- On-demand learning resources for HCPs through Integra Institute

Partnership ethos

- Comprehensive onboarding, training, and change management support
- Proactive post-market engagement rather than transactional vendor relationships

5 Clear and methodical remediation plan well underway

Strengthening the foundation to enable a reliable & scalable supply chain

Key Focus Areas

Quality

- Embedding a quality culture to drive results and continuous improvement
- Delivering on key metrics
- Develop supplier management and stabilization program
- Remediating sites
- Driving EU MDR product compliance

Operations

- Implementing full production process ownership & org redesign
- Driving increased alignment with Good Manufacturing Practices
- Deploying Lean & Six Sigma
- Planning and executing Integrated Business Planning
- Increasing visibility and planning via Supply Chain Control Tower
- Enhancing process capabilities

Roadmaps and Targets

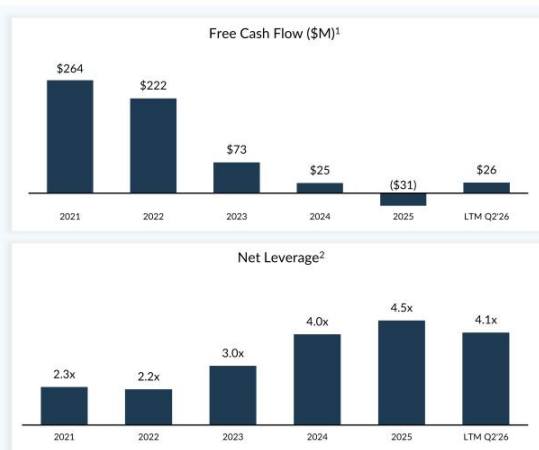
Quality transformation
and site/ production
remediation

Supply chain improvement
and IBP roadmap

Robust governance in place to ensure resourcing and milestone achievement

6 Return to strong FCF generation and deleveraging

Strong Free Cash Flow to support meaningful deleveraging story



Drivers of Margin Expansion & Cash Flow Improvement

- Return Products to Market through completion of remediation activities
- Deliver Cost Savings Initiatives announced and implemented in Q1 2026 as part of a broader margin expansion program
- Simplify Operations through the new operating model, improving efficiency and accountability
- Eliminate Temporary Transformation Costs as EU MDR, Braintree start-up, and structural optimization expenses roll off
- Improve Working Capital Performance through better inventory and receivables management
- Expand Gross Margins through lower remediation costs, reduced scrap, and manufacturing productivity gains
- Normalize Capital Spending as major investments in Braintree and capacity expansion are completed

¹ GAAP net cash provided by operating activities, net of CapEx. Free Cash Flow is a non-GAAP measure. See appendix for non-GAAP reconciliation

² Debt includes undrawn Letters of Credit and non-contingent deferred payment obligations (including certain deferred purchase price obligations and earned but unpaid earn-outs), excluding ordinary-course trade payables

7 Seasoned leadership team with decades of MedTech experience

Executive Leadership					Board of Directors		
 Stuart Essig, Ph.D. President and CEO	 Andrea Caruso Corporate VP, Strategy & Business Development	 Robert T. Davis, Jr. President, Tissue Reconstruction	 Kerri DiPietro Corporate VP, Chief Quality Officer	 Teshar Elavia Corporate VP, CTO	 Stuart Essig, Ph.D. President and CEO	 Keith Bradley, Ph.D.	
 Michael Hutchinson EVP, CLO, Company Secretary	 Topaz Kirlaw Chief Regulatory Officer	 Lea Daniels Knight Chief Financial Officer	 Dimitri Kivares CVP, CIO	 Rick Maveus SVP, Transformation & Program Management	 Shaundra Clay	 Barbara B. Hill	 Jeff Graves, Ph.D.
 Michael McBreen EVP & Chief Commercial Officer	 Harvinder Singh EVP & President, International	 Dr. Raymond Turner Corporate VP & CMO	 Chantal Veillon EVP & CHRO	 Valerie Young Corporate VP, Global Operations & Supply Chain	 Renee Lo	 Christian Schade	
      					     		
Leadership		Organization			Execution		
Deep and specialized MedTech expertise		Structural alignment and enhanced accountability			Delivering on operational and financial Compliance Master Plan		

07



Financial summary

INTEGRA 

Financial policy

Balance Sheet	• Continue to prioritize optimizing the Company's capital structure with a focus on deleveraging, extending maturities, and maintaining adequate liquidity • Decreased net leverage from 4.5x (2025YE) to 4.1x (Q2'2026) – Our target net leverage ratio is 2.5-3.5x • Disciplined portfolio investments with a focus on high-growth, high-margin opportunities that we have a clear right-to-win position in
Strategic Investments	Capital Expenditures • Remediation and project costs expected to be completed in fiscal year 2026 • Braintree facility became operational in Q2 2026; on track for commercial revenue being generated in Q4 2026 • 80% of CapEx focused on improving supply reliability and capacity with remaining 20% focused on growth and innovation
	M&A • Focus on deleveraging • M&A strategy focused on tuck-ins following deleveraging and sustainable cash flow generation
Capital Management	Liquidity • We anticipate increased flexibility to reduce outstanding Revolving Credit Facility borrowings as free cash flow generation improves, enhancing future borrowing capacity
	Share Repurchases • Repurchase shares strategically when it's the best way to return cash to our shareholders and when not in conflict with deleveraging objective • Current program expired in Q4 2025 and has not been renewed

Historical operating performance

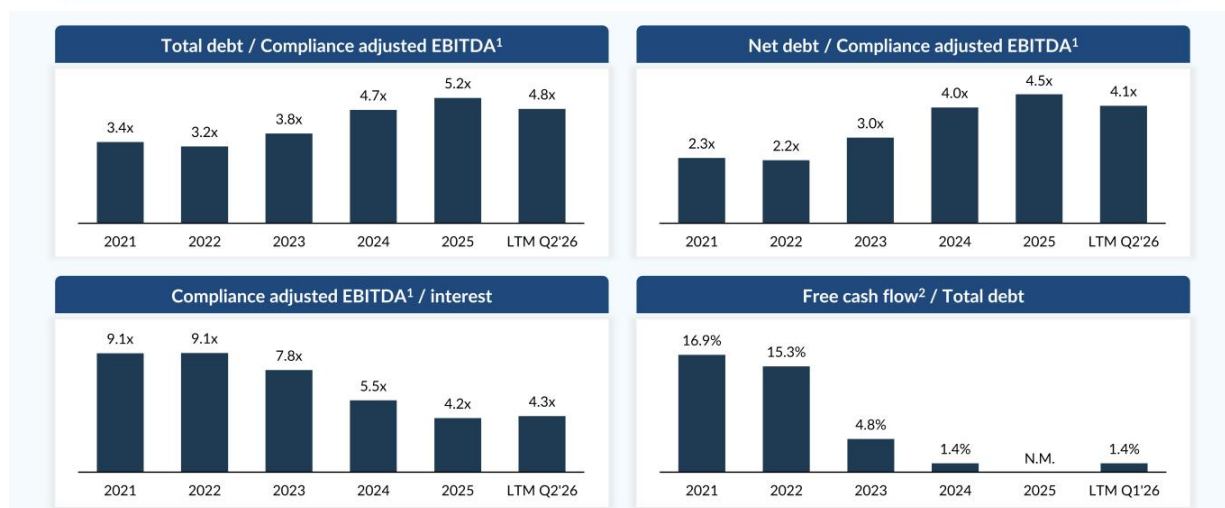


Source: Company Filings

¹ Compliance adjusted EBITDA is a non-GAAP measure. See appendix for non-GAAP reconciliation;

² GAAP net cash provided by operating activities, net of CapEx. Free Cash Flow is a non-GAAP measure. See appendix for non-GAAP reconciliation

Historical credit metrics



Source: Company Filings

Note: Debt includes undrawn Letters of Credit and non-contingent deferred payment obligations (including certain deferred purchase price obligations and earned but unpaid earn-outs), excluding ordinary-course trade payables

¹ Compliance adjusted EBITDA is a non-GAAP measure. See appendix for non-GAAP reconciliation;

² GAAP net cash provided by operating activities, net of CapEx. Free Cash Flow is a non-GAAP measure. See appendix for non-GAAP reconciliation



Appendix

EBITDA reconciliation

Trailing twelve month EBITDA reconciliation

(In \$M)	FY 2021	FY 2022	FY 2023	FY 2024	FY 2025	LTM Q2'26
Total Revenue	\$1,542	\$1,558	\$1,542	\$1,611	\$1,635	\$1,648
GAAP Net Income	\$169	\$181	\$68	\$(7)	\$(516)	\$(7)
Goodwill impairment charges	-	-	-	-	511	-
Depreciation and intangible asset amortization expense	123	118	123	147	151	152
Other (income), net	(14)	(8)	(3)	(4)	3	(8)
Interest expense, net	44	38	34	51	66	73
Income tax expense (benefit)	46	33	13	(11)	(47)	7
Structural optimization charges	20	23	16	24	48	48
EU Medical Device Regulation charges	24	45	47	45	42	31
Boston Recall	-	-	47	45	56	45
COVID-19 related charges	-	-	-	-	-	-
Convertible debt non-cash interest	-	-	-	-	-	-
Expenses related to debt refinancing	-	-	-	-	-	-
Discontinued product lines charges	0	-	-	-	-	-
Acquisition, divestiture and integration-related charges	(12)	(19)	25	34	4	(3)
Total of non-GAAP adjustments	232	231	302	329	834	345
Adjusted EBITDA	\$401	\$411	\$370	\$322	\$317	\$337
Adjusted EBITDA Margin	26.0%	26.4%	24.0%	20.0%	19.4%	20.5%
Stock-based compensation	36	25	20	24	19	25
Interest income	7	12	17	20	18	18
Other income	61	13	4	(1)	(2)	8
Other credit facility adjustments	(47)	(9)	(10)	23	6	8
Total Credit Agreement adjustments	57	41	31	66	41	59
Compliance adjusted EBITDA	\$458	\$452	\$401	\$389	\$359	\$396
Compliance Adjusted EBITDA Margin	29.7%	29.0%	26.0%	24.1%	21.9%	24.1%

FCF & Adjusted Gross Margin reconciliation

Trailing twelve month FCF reconciliation

(In \$M)	FY 2021	FY 2022	FY 2023	FY 2024	FY 2025	LTM Q2'26
Operating Cash Flow	\$312	\$264	\$140	\$129	\$50	\$85
Capex	48	42	67	104	81	60
Free Cash Flow	\$264	\$222	\$73	\$25	\$(31)	\$26

Trailing twelve month Adjusted Gross Margin reconciliation

(In \$M)	FY 2021	FY 2022	FY 2023	FY 2024	FY 2025	LTM Q2'26
Total Revenue	\$1,542	\$1,558	\$1,542	\$1,611	\$1,635	\$1,648
GAAP Gross Profit	\$945	\$970	\$885	\$882	\$832	\$865
Structural optimization charges	9	6	8	16	28	24
Acquisition, divestiture and integration-related charges	20	2	3	9	1	0
Braintree transition	-	-	46	43	55	44
EU MDR	4	5	6	4	4	3
Intangible asset amortization expense	67	64	70	84	92	93
Total of non-GAAP adjustments	99	76	134	156	180	165
Adjusted Gross Profit	\$1,043	\$1,046	\$1,018	\$1,038	\$1,012	\$1,029
Adjusted Gross Margin	67.7%	67.2%	66.1%	64.5%	61.9%	62.5%

