UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, DC 20549

FORM 10-Q

(Mark one)

☒ **QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the quarterly period ended September 30, 2026

OR

☐ **TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934**

For the transition period from _____ to _____.

Commission File Number: 0-19961



ORTHOFIX MEDICAL INC.
(Exact name of registrant as specified in its charter)

Delaware	**98-1340767**
(State or other jurisdiction of incorporation or organization)	**(I.R.S. Employer Identification No.)**
3451 Plano Parkway, Lewisville, Texas	**75056**
(Address of principal executive offices)	**(Zip Code)**

(214) 937-2000
(Registrant's telephone number, including area code)

Not applicable
(Former name, former address and former fiscal year, if changed since last report)

Indicate by check mark whether the registrant: (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. ☒ Yes ☐ No

Indicate by check mark whether the registrant has submitted electronically every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit such files). ☒ Yes ☐ No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, smaller reporting company, or an emerging growth company. See the definitions of "large accelerated filer," "accelerated filer," "smaller reporting company," and "emerging growth company" in Rule 12b-2 of the Exchange Act.

Large Accelerated filer	☐	Accelerated filer	☒
Non-Accelerated filer	☐	Smaller Reporting Company	☐
	.	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. ☐

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). ☐ Yes ☒ No

As of November 5, 2026, 40,759,801 shares of common stock were issued and outstanding.

Securities registered pursuant to Section 12(b) of the Act:

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common stock, $0.10 par value per share	OFIX	Nasdaq Global Select Market

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Forward-Looking Statements

This quarterly report contains forward-looking statements within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), and Section 27A of the Securities Act of 1933, as amended (the "Securities Act"), relating to our business and financial outlook, which are based on our current beliefs, assumptions, expectations, estimates, forecasts, and projections. All statements, other than statements of historical fact, contained in this report, are forward-looking statements. In some cases, you can identify forward-looking statements by terminology such as "may," "will," "should," "expects," "plans," "anticipates," "believes," "estimates," "projects," "intends," "predicts," "potential," "positioned," "deliver," or "continue" or other comparable terminology. Forward-looking statements include, but are not limited to, statements about:

- our intentions, beliefs, and expectations regarding our operations, sales, expenses, and future financial performance;
- our operating results;
- our plans for future products and enhancements of existing products;
- anticipated growth and trends in our business;
- the timing of, and our ability to maintain and obtain, regulatory clearances or approvals;
- our belief that our cash and cash equivalents, investments, and access to our credit facilities will be sufficient to satisfy our anticipated cash requirements;
- our expectations regarding our revenues, customers, and distributors;
- our ability to persuade surgeons to use our products;
- our ability to attract and retain distributors and/or sales and other personnel;
- our expectations regarding our costs, suppliers, and manufacturing abilities;
- our beliefs and expectations regarding our market penetration and expansion efforts;
- our anticipated trends and challenges in the markets in which we operate; and
- our expectations and beliefs regarding, and the impact of, investigations, claims, and litigation.

Forward-looking statements are not guarantees of future performance and involve risks, uncertainties, estimates, and assumptions that are difficult to predict. Any or all forward-looking statements that we make may turn out to be wrong (due to inaccurate assumptions that we make or otherwise), and our actual outcomes and results may differ materially from those expressed in forward-looking statements. Potential risks and uncertainties that could cause actual results to differ materially include, but are not limited to, those set forth in Part I, Item 1A under the heading *Risk Factors* of our Annual Report on Form 10-K for the year ended December 31, 2025 ("2025 10-K"); Part II, Item 7 *Management's Discussion and Analysis of Financial Condition and Results of Operations* of the 2025 10-K*;* and elsewhere throughout the 2025 10-K, and in our reports filed with the U.S. Securities and Exchange Commission (the "SEC") subsequent to the date we filed the 2025 10-K with the SEC. You should not place undue reliance on any forward-looking statements. Further, any forward-looking statement in this report speaks only as of the date hereof, unless it is specifically otherwise stated to be made as of a different date. Except as required by law, we undertake no obligation to update, and expressly disclaim any duty to update, our forward-looking statements, whether as a result of circumstances or events that arise after the date hereof, new information, or otherwise.

Trademarks

Solely for convenience, our trademarks and trade names in this report are referred to without the ® and ™ symbols, but such references should not be construed as an indicator that we will not assert, to the fullest extent under applicable law, our rights thereto.

PART I. FINANCIAL INFORMATION

Item 1. Financial Statements

ORTHOFIX MEDICAL INC.
Condensed Consolidated Balance Sheets

(U.S. Dollars, in thousands, except par value data)	September 30, 2026 (Unaudited)	December 31, 2025
Assets		
Current assets		
Cash and cash equivalents	$ 86,322	$ 82,025
Restricted cash	598	3,090
Accounts receivable, net of allowances of $11,583 and $8,308, respectively	131,719	135,746
Inventories	190,357	172,319
Prepaid expenses and other current assets	21,791	23,667
Total current assets	430,787	416,847
Property, plant, and equipment, net	133,388	129,399
Intangible assets, net	62,904	72,765
Goodwill	194,934	194,934
Other long-term assets	33,748	36,702
Total assets	**$ 855,761**	**$ 850,647**
Liabilities and shareholders' equity		
Current liabilities		
Accounts payable	$ 58,032	$ 58,392
Current portion of finance lease liability	136	837
Other current liabilities	80,217	111,253
Total current liabilities	138,385	170,482
Long-term debt	221,847	157,391
Long-term portion of finance lease liability	12,868	17,060
Other long-term liabilities	53,094	55,677
Total liabilities	426,194	400,610
Contingencies (Note 7)		
Shareholders' equity		
Common shares $0.10 par value; 100,000 shares authorized; 40,749 and 39,834 issued and outstanding as of September 30, 2026, and December 31, 2025, respectively	4,075	3,983
Additional paid-in capital	837,165	813,769
Accumulated deficit	(411,113)	(368,333)
Accumulated other comprehensive income (loss)	(560)	618
Total shareholders' equity	429,567	450,037
Total liabilities and shareholders' equity	**$ 855,761**	**$ 850,647**

The accompanying notes form an integral part of these condensed consolidated financial statements.

ORTHOFIX MEDICAL INC.

Condensed Consolidated Statements of Operations and Comprehensive Loss

(Unaudited, U.S. Dollars, in thousands, except per share data)	Three Months Ended September 30, 2026	Three Months Ended September 30, 2025	Nine Months Ended September 30, 2026	Nine Months Ended September 30, 2025
Net sales	$ 207,862	$ 205,634	$ 615,503	$ 602,401
Cost of sales	61,316	57,111	179,704	192,726
Gross profit	146,546	148,523	435,799	409,675
Sales, general, and administrative	125,330	148,102	398,271	417,576
Research and development	15,277	14,774	46,541	50,474
Acquisition-related amortization, impairment, and remeasurement (Note 11)	4,394	2,693	12,012	23,547
Operating income (loss)	1,545	(17,046)	(21,025)	(81,922)
Interest expense, net	(5,889)	(4,681)	(17,638)	(13,137)
Other income (expense), net	(904)	(535)	(2,416)	6,441
Loss before income taxes	(5,248)	(22,262)	(41,079)	(88,618)
Income tax expense	(826)	(533)	(1,701)	(1,352)
Net loss	**$ (6,074)**	**$ (22,795)**	**$ (42,780)**	**$ (89,970)**
Net loss per common share:				
Basic	$ (0.15)	$ (0.57)	$ (1.05)	$ (2.28)
Diluted	(0.15)	(0.57)	(1.05)	(2.28)
Weighted average number of common shares:				
Basic	41,178	39,766	40,842	39,468
Diluted	41,178	39,766	40,842	39,468
Other comprehensive income (loss), before tax				
Currency translation adjustment	(283)	(274)	(1,178)	4,936
Other comprehensive income (loss), before tax	(283)	(274)	(1,178)	4,936
Income tax expense related to other comprehensive income (loss)	—	—	—	—
Other comprehensive income (loss), net of tax	(283)	(274)	(1,178)	4,936
Comprehensive loss	**$ (6,357)**	**$ (23,069)**	**$ (43,958)**	**$ (85,034)**

The accompanying notes form an integral part of these condensed consolidated financial statements.

ORTHOFIX MEDICAL INC.

Condensed Consolidated Statements of Changes in Shareholders' Equity

(Unaudited, U.S. Dollars, in thousands)	Number of Common Shares Outstanding	Common Shares	Additional Paid-in Capital	Accumulated Deficit	Accumulated Other Comprehensive Income (Loss)	Total Shareholders' Equity
At December 31, 2025	39,834	$ 3,983	$ 813,769	$ (368,333)	$ 618	$ 450,037
Net loss	—	—	—	(20,908)	—	(20,908)
Other comprehensive loss, net of tax	—	—	—	—	(476)	(476)
Share-based compensation expense	—	—	6,533	—	—	6,533
Common shares issued, net	548	55	(55)	—	—	—
At March 31, 2026	**40,382**	**$ 4,038**	**$ 820,247**	**$ (389,241)**	**$ 142**	**$ 435,186**
Net loss	—	—	—	(15,798)	—	(15,798)
Other comprehensive loss, net of tax	—	—	—	—	(419)	(419)
Share-based compensation expense	—	—	7,735	—	—	7,735
Common shares issued, net	348	35	2,441	—	—	2,476
At June 30, 2026	**40,730**	**$ 4,073**	**$ 830,423**	**$ (405,039)**	**$ (277)**	**$ 429,180**
Net loss	—	—	—	(6,074)	—	(6,074)
Other comprehensive loss, net of tax	—	—	—	—	(283)	(283)
Share-based compensation expense	—	—	6,744	—	—	6,744
Common shares issued, net	19	2	(2)	—	—	—
At September 30, 2026	**40,749**	**$ 4,075**	**$ 837,165**	**$ (411,113)**	**$ (560)**	**$ 429,567**

(Unaudited, U.S. Dollars, in thousands)	Number of Common Shares Outstanding	Common Shares	Additional Paid-in Capital	Accumulated Deficit	Accumulated Other Comprehensive Income (Loss)	Total Shareholders' Equity
At December 31, 2024	38,486	$ 3,849	$ 779,718	$ (276,141)	$ (4,302)	$ 503,124
Net loss	—	—	—	(53,094)	—	(53,094)
Other comprehensive income, net of tax	—	—	—	—	1,746	1,746
Share-based compensation expense	—	—	6,469	—	—	6,469
Common shares issued, net	610	61	(12)	—	—	49
At March 31, 2025	**39,096**	**$ 3,910**	**$ 786,175**	**$ (329,235)**	**$ (2,556)**	**$ 458,294**
Net loss	—	—	—	(14,081)	—	(14,081)
Other comprehensive income, net of tax	—	—	—	—	3,464	3,464
Share-based compensation expense	—	—	7,824	—	—	7,824
Common shares issued, net	387	38	2,808	—	—	2,846
At June 30, 2025	**39,483**	**$ 3,948**	**$ 796,807**	**$ (343,316)**	**$ 908**	**$ 458,347**
Net loss	—	—	—	(22,795)	—	(22,795)
Other comprehensive income, net of tax	—	—	—	—	(274)	(274)
Share-based compensation expense	—	—	7,181	—	—	7,181
Common shares issued, net	36	4	23	—	—	27
At September 30, 2025	**39,519**	**$ 3,952**	**$ 804,011**	**$ (366,111)**	**$ 634**	**$ 442,486**

The accompanying notes form an integral part of these condensed consolidated financial statements.

ORTHOFIX MEDICAL INC.

Condensed Consolidated Statements of Cash Flows

(Unaudited, U.S. Dollars, in thousands)	Nine Months Ended September 30, 2026	2025
Cash flows from operating activities		
Net loss	$ (42,780)	$ (89,970)
Adjustments to reconcile net loss to net cash from operating activities		
Depreciation, amortization, and impairment	39,938	64,243
Inventory reserve expenses	13,941	27,776
Amortization of operating lease assets, debt costs, and other assets	3,332	3,959
Provision for expected credit losses	4,833	2,466
Deferred income taxes	924	333
Share-based compensation expense	21,012	21,474
Loss on disposal of fixed assets	521	861
Change in valuation of investment securities	(16)	(41)
Change in fair value of contingent consideration	3,014	(1,800)
Other	713	1,127
Changes in operating assets and liabilities		
Accounts receivable	(1,292)	3,131
Inventories	(32,908)	(9,469)
Prepaid expenses and other current assets	1,766	797
Accounts payable	(3,316)	(5,656)
Other current liabilities	(31,178)	(9,052)
Contingent consideration milestone payment	(2,864)	(1,340)
Other long-term assets and liabilities	889	(3,189)
Net cash provided by (used in) operating activities	**(23,471)**	**5,650**
Cash flows from investing activities		
Capital expenditures	(37,810)	(23,749)
Other investing activities	146	22
Net cash used in investing activities	**(37,664)**	**(23,727)**
Cash flows from financing activities		
Proceeds from issuance of common shares	2,626	3,090
Payments related to tax withholdings for share-based compensation	(150)	(168)
Payments related to finance lease obligation	(79)	(564)
Proceeds from credit facility	64,025	—
Contingent consideration milestone payment	(3,170)	(4,990)
Payment of debt issuance costs and other financing activities	(24)	(531)
Net cash provided by (used in) financing activities	**63,228**	**(3,163)**
Effect of exchange rate changes on cash	(288)	1,448
Net change in cash, cash equivalents, and restricted cash	1,805	(19,792)
Cash, cash equivalents, and restricted cash at the beginning of period	85,115	85,738
Cash, cash equivalents, and restricted cash at the end of period	**$ 86,920**	**$ 65,946**
Components of cash, cash equivalents, and restricted cash at the end of period		
Cash and cash equivalents	$ 86,322	$ 62,860
Restricted cash	598	3,086
Cash, cash equivalents, and restricted cash at the end of period	**$ 86,920**	**$ 65,946**
Noncash investing activities - Accrued purchases of capital expenditures	$ 16,096	$ 12,923
Noncash investing activities - Purchase of intangible assets	—	40

The accompanying notes form an integral part of these condensed consolidated financial statements.

ORTHOFIX MEDICAL INC.
Notes to the Unaudited Condensed Consolidated Financial Statements

1. Business and basis of presentation

Description of the Business

Orthofix Medical Inc. (the "Company" or "Orthofix") is a global medical technology company dedicated to advancing healing and restoring mobility for patients with complex musculoskeletal conditions. Headquartered in Lewisville, Texas, the Company offers a differentiated portfolio of spinal implants, therapeutic solutions, limb reconstruction systems, biologics, and enabling technologies, including the 7D FLASH Navigation System. The Company's technology-enabled solutions are designed to support surgeons across the continuum of care and improve outcomes for patients.

Basis of Presentation

The accompanying unaudited condensed consolidated financial statements have been prepared in accordance with generally accepted accounting principles in the United States ("U.S. GAAP") for interim financial information and with the instructions to Form 10-Q and Rule 10-01 of Regulation S-X. Pursuant to these rules and regulations, certain information and note disclosures normally included in financial statements prepared in accordance with U.S. GAAP have been condensed or omitted. In the opinion of management, all adjustments (consisting of normal recurring items) considered necessary for a fair statement have been included. These condensed consolidated financial statements should be read in conjunction with the consolidated financial statements and related notes contained in the Company's Annual Report on Form 10-K for the year ended December 31, 2025. Operating results for the three and nine months ended September 30, 2026, are not necessarily indicative of the results that may be expected for other interim periods or the year ending December 31, 2026.

The preparation of financial statements in conformity with U.S. GAAP requires management to make estimates and assumptions that affect the amounts reported in the financial statements and accompanying notes. On an ongoing basis, the Company evaluates its estimates, including those related to revenue recognition; contractual allowances; allowances for expected credit losses; inventories; valuation of intangible assets; goodwill; fair value measurements, including contingent consideration; litigation and contingent liabilities; tax matters; and share-based compensation. Actual results could differ from these estimates.

2. Recently adopted accounting standards and recently issued accounting pronouncements

Recently Issued Accounting Pronouncements

Topic	Description of Guidance	Effective Date	Status of Company's Evaluation
Disclosure Improvements - Codification Amendments in Response to the SEC's Disclosure Update and Simplification Initiative (Accounting Standard Update "ASU" 2023-06)	Adds interim and annual disclosure requirements to a variety of subtopics in the Accounting Standards Codification, including those focusing on accounting changes, earnings per share, debt, and repurchase agreements. The guidance will be applied prospectively. The effective date will be the date when the SEC's removal of the related disclosure requirement becomes effective, with early adoption prohibited.	Various	The Company is currently evaluating the impact this ASU may have on its consolidated financial statements.
Disaggregation of Income Statement Expenses (ASU 2024-03)	Improves financial reporting by requiring that public business entities disclose additional information about specific expense categories in the notes to the financial statements at interim and annual reporting periods. The amendments are to be applied prospectively to financial statements issued for reporting periods after the effective date or retrospectively to all prior periods presented in the financial statements.	January 1, 2027	The Company is currently evaluating the impact this ASU may have on its consolidated financial statements.
Targeted Improvements to the Accounting for Internal-Use Software (ASU 2025-06)	Aligns the accounting for internal-use software with how software is developed to increase the operability of the recognition and capitalization of internal-use software costs in accordance with Subtopic 350-40. Early adoption is permitted as of the beginning of an annual reporting period. The guidance is to be applied prospectively to new software costs incurred as of the beginning of the adoption period for all projects, including in-process projects.	January 1, 2028	The Company is currently evaluating the impact this ASU may have on its consolidated financial statements.
Narrow-Scope Requirements for Interim Reporting (ASU 2025-11)	Clarifies interim disclosure requirements and applicability of Topic 270, *Interim Reporting* for events since the end of the last annual reporting period that have a material impact on the entity. Early adoption is permitted and the amendments are to be applied prospectively or retrospectively to any or all periods presented in the financial statements.	January 1, 2028	The Company is currently evaluating the impact this ASU may have on its consolidated financial statements.

Other recently issued ASUs, excluding those ASUs which have already been disclosed as adopted or described above, were assessed and determined not applicable, or are expected to have minimal impact on the Company's condensed consolidated financial statements.

3. Inventories

Inventories were as follows:

(U.S. Dollars, in thousands)	September 30, 2026	December 31, 2025
	(Unaudited)	
Raw materials	$ 31,981	$ 22,865
Work-in-process	61,576	63,255
Finished products	96,800	86,199
Inventories	$ 190,357	$ 172,319

4. Leases

A summary of the Company's lease portfolio as of September 30, 2026, and December 31, 2025, is presented in the table below:

(U.S. Dollars, in thousands)	Classification	September 30, 2026	December 31, 2025
		(Unaudited)	
Assets			
Operating leases	Other long-term assets	$ 20,926	$ 22,279
Finance leases	Property, plant, and equipment, net	9,124	14,442
Total lease assets		$ 30,050	$ 36,721
Liabilities			
Current			
Operating leases	Other current liabilities	$ 3,296	$ 3,147
Finance leases	Current portion of finance lease liability	136	837
Long-term			
Operating leases	Other long-term liabilities	23,939	25,413
Finance leases	Long-term portion of finance lease liability	12,868	17,060
Total lease liabilities		$ 40,239	$ 46,457

The Company entered into an amendment for its corporate headquarters location in Lewisville, Texas in January 2026, extending the contractual term of its lease to 2040, while also maintaining its existing lease term extension options within the agreement. Accordingly, the Company reassessed the lease classification and remeasured the lease liability based on the terms of the amendment. This resulted in a decrease in the recognized finance lease liability and finance lease right-of-use asset of approximately $4.8 million, respectively, with this decrease largely driven by an increase in the Company's incremental borrowing rate.

Supplemental cash flow information related to leases was as follows:

	Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025
Cash paid for amounts included in the measurement of lease liabilities		
Operating cash flows from operating leases	$ 6,652	$ 6,595
Operating cash flows from finance leases	1,109	602
Financing cash flows from finance leases	79	564
ROU assets obtained in exchange for lease obligations		
Operating leases	1,030	12,192
Finance leases	—	28

5. Long-term debt

The carrying values of the Company's outstanding debt obligations as of September 30, 2026, and December 31, 2025, were as follows:

(U.S. Dollars, in thousands)	September 30, 2026	December 31, 2025
	(Unaudited)	
Outstanding Term Loans		
Principal amount	$ 225,000	$ 160,000
Unamortized original debt discount	(2,263)	(1,839)
Unamortized debt issuance costs and lenders fees	(890)	(770)
Total indebtedness from outstanding term loans	$ 221,847	$ 157,391
Current portion of long-term debt	$ —	$ —
Long-term debt	221,847	157,391
Total indebtedness outstanding	$ 221,847	$ 157,391

On November 7, 2024, the Company, as borrower, and its U.S. subsidiaries entered into a $275.0 million secured credit agreement (the "Credit Agreement") with Oxford Finance LLC, as administrative agent and as collateral agent ("Oxford") and certain lenders party thereto, including Oxford, K2 HealthVentures LLC, and HSBC Ventures USA Inc. The Credit Agreement provides for a $160.0 million senior secured term loan (the "Initial Term Loan") and a $65.0 million senior secured delayed draw term loan facility (the "Term B Loan"). In addition, at Oxford's discretion, an additional $50.0 million of draw capacity is available through January 1, 2029 (the "Term C Loan" and, collectively with the Term B Loan and the Initial Term Loan, the "Credit Facilities").

On January 15, 2026, the Company borrowed $65.0 million via the Term B Loan to provide capital flexibility ahead of the expiration of the tranche. The Credit Facilities, to the extent ultimately drawn, will each mature in November 2029, following an interest-only payment period ending December 2028, and monthly amortization of principal and accrued interest between January 2029 and November 2029. The Credit Agreement contains financial covenants requiring the Company to maintain (i) a minimum level of liquidity at all times and (ii) a maximum total debt-to-EBITDA leverage ratio (measured on a quarterly basis) during the term of the facility. As of September 30, 2026, the Company was in compliance with all required financial covenants.

As of September 30, 2026, the Company had no borrowings on its available lines of credit in Italy, which provide up to an aggregate amount of €5.5 million ($6.2 million).

6. Fair value measurements

The fair value measurements of the Company's financial assets and liabilities measured on a recurring basis were as follows:

	September 30, 2026				December 31, 2025
(U.S. Dollars, in thousands)	Level 1	Level 2	Level 3	Total	Total
				(Unaudited)	
Liabilities					
Lattus Contingent Consideration	$ —	$ —	$ 4,910	$ 4,910	$ 7,930
Deferred compensation plan	—	1,519	—	1,519	1,720
Total	$ —	$ 1,519	$ 4,910	$ 6,429	$ 9,650

Lattus Contingent Consideration

In connection with the merger with SeaSpine Holdings Corporation ("SeaSpine") in 2023 (the "Merger"), the Company assumed a contingent consideration obligation under a purchase agreement between SeaSpine and Lattus Spine LLC ("Lattus") executed in December 2022. Under the terms of this agreement, the Company may be required to make installment payments to Lattus (the "Lattus Contingent Consideration") at certain dates based on future net sales of certain products (the "Lateral Products").

The estimated fair value of the Lattus Contingent Consideration is determined using a Monte Carlo simulation and a discounted cash flow model requiring significant inputs which are not observable in the market. The significant inputs include assumptions related to

the estimated future sales of Lateral Products, revenue risk-adjusted discount rates, revenue volatility, and discount rates matched to the timing of payments. The following table provides a reconciliation of the beginning and ending balances for the Lattus Contingent Consideration measured at estimated fair value using significant unobservable inputs (Level 3):

(Unaudited, U.S. Dollars, in thousands)	2026	2025
Lattus Contingent Consideration estimated fair value at January 1	$ 7,930	$ 15,400
Change in fair value recognized in acquisition-related amortization, impairment, and remeasurement	3,014	(1,800)
Installment payment	(6,034)	(6,330)
Lattus Contingent Consideration estimated fair value at September 30	$ 4,910	$ 7,270

The estimated fair value of the Lattus Contingent Consideration as of September 30, 2026, was $4.9 million; however, the actual amount ultimately paid could be higher or lower. As of September 30, 2026, the Company classified the remaining Lattus Contingent Consideration of $4.9 million within other current liabilities.

The following table provides quantitative information related to certain key assumptions utilized within the valuation as of September 30, 2026:

(Unaudited, U.S. Dollars, in thousands)	Fair Value as of September 30, 2026	Unobservable inputs	Estimate
Lattus Contingent Consideration	$ 4,910	Counterparty discount rates	10.2%
		Revenue risk-adjusted discount rates	6.2%

7. Commitments and Contingencies

Arbitration claims with former executives

In September 2023, the Company's Board of Directors (the "Board") terminated the employment of Keith Valentine, John Bostjancic, and Patrick Keran, who had served respectively as the Company's President and Chief Executive Officer, Chief Financial Officer, and Chief Legal Officer (collectively, the "Former Executives"). The Board's decision followed an investigation conducted by independent outside legal counsel and directed and overseen by a committee of certain of the Company's independent directors. At the time of termination, the Company notified each of the Former Executives that their respective terminations of employment were being made for "Cause," as such term is defined in applicable employment-related agreements (including each executive's respective Change in Control and Severance Agreement, dated June 19, 2023 (the "CIC and Severance Agreements"). The Former Executives subsequently made claims against the Company in arbitration in the State of California, asserting breach of contract because each of them was entitled to the severance payments and other equity-based rights that would be owed to them if their respective termination had been made "without Cause" under the CIC and Severance Agreements, and further asserting damages for purported defamation, false light invasion of privacy, and deceit, as well as indemnification and advancement for attorneys' fees.

On January 26, 2026, the arbitrator in Mr. Valentine's matter issued a decision denying Mr. Valentine's defamation, false light invasion of privacy, and deceit claims, and his indemnification of fees claim. Based on the evidence presented during the arbitration process, the arbitrator found that Mr. Valentine's conduct met the legal definition of "acts of moral turpitude" and that the public statements that the Company made about Mr. Valentine in a press release and filings with the SEC subsequent to the termination of his employment were true. Although Mr. Valentine's conduct was found to meet the legal definition of "acts of moral turpitude" for purposes of his defamation and other tort claims, and although engaging in "material acts of moral turpitude" would constitute "Cause" under the CIC and Severance Agreement, the arbitrator maintained his preliminary decision issued on October 2, 2025, finding that (i) Mr. Valentine's conduct prior to his entry into the CIC and Severance Agreement on June 19, 2023 could not be considered for purposes of determining whether "Cause" existed under such agreement, and (ii) his conduct between that date and his termination of employment on September 11, 2023, did not amount to "Cause". As a result, the arbitrator issued an interim award to Mr. Valentine for breach of contract damages in the amount of $11.8 million, finding such amount to be equivalent to the severance and equity-based rights that Mr. Valentine would have received in a "without Cause" termination. The arbitrator also awarded accrued interest, at the pre-judgment interest rate of 10% under California law. On April 22, 2026, the Company made a payment in satisfaction of the arbitrator's interim award plus accrued interest, in the amount of $14.8 million. On June 24, 2026, the arbitrator issued his final award. The final award (i) confirmed the amounts in the interim award (which were paid by the Company

on April 22, 2026), and (ii) awarded an additional $1.1 million to Mr. Valentine for attorneys' fees, costs, and expenses, which the Company has accrued as of September 30, 2026, and expects to pay once the final award is confirmed in California State Court.

On April 15, 2026, Mr. Keran's arbitration claims were settled for $4.25 million. On April 22, 2026, the Company and Mr. Bostjancic entered into a settlement agreement covering all of Mr. Bostjancic's claims, in the amount of $4.25 million. Payments were made to each of Mr. Keran and Mr. Bostjancic in the amount of $4.25 million, respectively, in the second quarter of 2026.

In addition to the Former Executives' arbitration claims, in September 2024 Messrs. Valentine, Bostjancic and Keran filed an action in California State Court against former director and interim CEO Catherine Burzik and current director Wayne Burris, seeking relief for, among other things, alleged defamation, false light invasion of privacy, intentional misrepresentation, false promise, and tortious interference with contract. Mr. Valentine and Mr. Bostjancic dismissed their claims in this action following the issuance of Mr. Valentine's final arbitration award and execution of Mr. Bostjancic's settlement agreement, respectively. On August 31, 2026, Mr. Keran filed an amended complaint in this action, against current directors Shweta Maniar, Jason Hannon, John Henneman and Michael Paolucci, and former directors Stuart Essig and James Hinrichs, in addition to Mr. Burris and Ms. Burzik, alleging defamation as the sole cause of action. The Company disagrees with the allegations contained in this action against current and former directors and will continue vigorously defending the asserted claims. The Company currently cannot reasonably estimate a possible loss, or range of loss, that may arise from the action.

Securities class action complaints

On August 21, 2024, a securities class action complaint captioned Bernal v. Orthofix Medical Inc., et al., Case No. 24-cv-00690, was filed in the United States District Court for the Eastern District of Texas (the "Bernal Complaint"). The plaintiff, a purported Company shareholder, alleges through the complaint violations of Sections 10(b) and 20(a) of the Exchange Act, and SEC Rule 10b-5 promulgated thereunder, and names as defendants the Company and the following former Company directors and officers: Jon Serbousek (former director and former President and Chief Executive Officer), Keith Valentine (former director and former President and Chief Executive Officer), John Bostjancic (former Chief Financial Officer), and Patrick Keran (former Chief Legal Officer). The complaint alleges that the Company made, and the named former directors and officers caused the Company to make, materially false and misleading statements between October 11, 2022, and September 12, 2023, that, according to the complaint, falsely assured the market of Messrs. Valentine, Bostjancic, and Keran's respective commitments to, among other things, ethical and legal standards and corporate responsibility.

On September 6, 2024, a securities class action complaint captioned O'Hara v. Orthofix Medical Inc., et al., Case No. 24-cv-01593, was filed in the United States District Court for the Southern District of California (the "O'Hara Complaint"). The plaintiff, a purported former shareholder of SeaSpine at the time of the Merger, alleges through the complaint violations of Sections 11, 12 and 15 of the Securities Act, and names most of the same defendants as the Bernal Complaint, as well as certain additional current and/or former Company directors and officers. The complaint makes similar assertions to the Bernal complaint, and alleges that the Company's registration statement on Form S-4 filed in 2022 in connection with the Merger, as well as related written and oral offering materials, contained untrue statements of material fact and material omissions, including, among other things, with respect to the effectiveness of the Company's internal controls. On November 26, 2024, the O'Hara Complaint was transferred to the Eastern District of Texas, and on December 11, 2024, the O'Hara Complaint was consolidated with the Bernal Complaint. On April 17, 2025, the plaintiffs filed an amended complaint in the consolidated action, captioned In re Orthofix Medical Inc. Securities Litigation, with substantially the same allegations contained in the Bernal Complaint and the O'Hara Complaint. The consolidated case is captioned In re Orthofix Medical Inc. Securities Litigation, Case No. 24-cv-00690 and is pending in the Eastern District of Texas. The Company and the individual defendants moved to dismiss the amended complaint on May 15, 2025. On March 9, 2026, the Court granted defendants' motion to dismiss the plaintiffs' claims under the Exchange Act (but not those under the Securities Act), finding that (i) all but one of the Company's statements at issue were immaterial as a matter of law, and (ii) no statements caused loss. The Court provided the plaintiffs with leave to amend their complaint to address pleading deficiencies, and plaintiffs filed an amended complaint on April 8, 2026, which the Company and the individual defendants moved to dismiss on May 8, 2026.

On October 28, 2024, a derivative shareholder complaint was filed against certain of the Company's current and former officers and directors alleging derivative liability for the allegations made in the two complaints noted above. On December 18, 2024, a second derivative shareholder complaint was filed with the same allegations made in the first derivative shareholder complaint. On March 21, 2025, the two derivative shareholder complaints were consolidated into one case. On August 25, 2026, a third derivative shareholder complaint was filed with substantially the same allegations made in the first and second derivative shareholder

complaints. On September 9, 2026, a fourth derivative shareholder complaint was filed against Messrs. Valentine, Bostjancic and Keran, with substantially the same allegations made in the first, second and third derivative shareholder complaints.

The Company disagrees with the legal claims asserted in these complaints and is vigorously defending them. Due in part to the preliminary nature of these three matters, the Company currently cannot reasonably estimate a possible loss, or range of loss, that may arise from the respective complaints.

Commitments

The Company is party to agreements with certain distributor partners that grant the Company the option to purchase, and the distributor the option to require the Company to purchase, the distributor's business on specified future dates. Upon exercise of either option (and, in certain cases, satisfaction of specified conditions), the parties are required to work in good faith to consummate the transaction. The purchase price may be paid in cash, shares of the Company's common stock, or a combination of both, at the Company's election, although in certain cases no more than 50% of the purchase price may be paid in shares.

Italian Medical Device Payback ("IMDP")

In 2015, the Italian Parliament introduced rules for entities that supply goods and services to the Italian National Healthcare System. A key provision of the law is a 'payback' measure, requiring medical device companies in Italy to make payments to the Italian government if medical device expenditures exceed regional maximum ceilings. Companies are required to make payments equal to a percentage of expenditures exceeding maximum regional caps.

In the third quarter of 2022, the Italian Ministry of Health provided guidelines to the Italian regions and provinces on seeking payback of expenditure overruns relating to the 2015 through 2018 calendar years. Since receiving the guidelines, several regions and provinces have requested payment from affected medical device companies, including the Company. The Company has taken legal action to dispute the legality of such measures. In July 2024, the Italian Constitutional Court issued two judgments following public hearings on the matter held in May 2024. These judgments (i) declared the payback system itself as constitutionally legitimate and (ii) extended previously communicated reductions in the payback liability for certain fiscal years to all medical device companies, regardless of whether or not they had waived their legal claims on the matter. In September 2026, the Italian Supreme Administrative Court (Consiglio di Stato) held a hearing in which it verbally indicated that it would issue a written decision finding that the payback law and system are inconsistent with European Union law and would refer the matter to the European Court of Justice for a final decision on the invalidity of the law. While the Company continues to monitor these legal proceedings and evaluate their potential impact on the IMDP, the September 2026 hearing did not result in a final ruling and did not materially change the Company's assessment of its estimated liability as of September 30, 2026.

The Company accounts for the estimated cost of the IMDP as sales, general, and administrative expense and periodically reassesses the liability based upon current facts and circumstances. As a result, the Company recorded expenses of $0.2 million and $0.9 million for the three and nine months ended September 30, 2026, respectively, and expenses of $0.3 million and $0.9 million for the three and nine months ended September 30, 2025. As of September 30, 2026, the Company has accrued $11.2 million related to the IMDP, which it has classified within other long-term liabilities; however, the actual liability could be higher or lower than the amount accrued once all legal proceedings are resolved and upon further clarification of the IMDP by the Italian and European authorities.

8. Accumulated other comprehensive income (loss)

The components of and changes in accumulated other comprehensive income (loss) were as follows:

(Unaudited, U.S. Dollars, in thousands)	Currency Translation Adjustments	Neo Medical Convertible Loan	Accumulated Other Comprehensive Income (Loss)
Balance at December 31, 2025	$ 846	$ (228)	$ 618
Other comprehensive loss	(1,178)	—	(1,178)
Income taxes	—	—	—
Balance at September 30, 2026	$ (332)	$ (228)	$ (560)

9. Revenue recognition and accounts receivable

Revenue recognition

The Company has two reporting segments: Global Spine and Global Limb Reconstruction. Within the Global Spine reporting segment, there are two product categories: (i) Therapeutic Solutions (formerly Bone Growth Therapies), and (ii) Spinal Implants, Biologics, and Enabling Technologies.

The tables below present net sales by product category and reporting segment:

	Three Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	Change
Therapeutic Solutions	$ 63,956	$ 61,211	4.5%
Spinal Implants, Biologics, and Enabling Technologies	107,493	110,852	-3.0%
Global Spine	171,449	172,063	-0.4%
Global Limb Reconstruction	36,413	33,571	8.5%
Net sales	$ 207,862	$ 205,634	1.1%

	Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	Change
Therapeutic Solutions	$ 185,882	$ 178,834	3.9%
Spinal Implants, Biologics, and Enabling Technologies	322,662	326,889	-1.3%
Global Spine	508,544	505,723	0.6%
Global Limb Reconstruction	106,959	96,678	10.6%
Net sales	$ 615,503	$ 602,401	2.2%

Product sales and marketing service fees

The table below presents product sales and marketing service fees, which are both components of net sales:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025	2026	2025
Product sales	$ 196,990	$ 193,813	$ 583,085	$ 566,841
Marketing service fees	10,872	11,821	32,418	35,560
Net sales	$ 207,862	$ 205,634	$ 615,503	$ 602,401

Marketing service fees are received from MTF Biologics ("MTF") based on total sales of biologic tissues and relate solely to the Biologics product category within the Global Spine reporting segment, whereas product sales primarily consist of the sale of Therapeutic Solutions, Spinal Implants, non-MTF sourced Biologics, Enabling Technologies, and Global Limb Reconstruction products. As MTF is the single supplier for certain allografts in the Company's Biologics portfolio, which are derived from deceased donors for their bone grafts and living donors for their amnion grafts, any event or circumstance that would impact MTF's continued access to donors or the Company's ability to market these tissues may adversely impact the Company's financial results.

Accounts receivable and related allowances

The following table provides the detail of changes in the Company's allowance for expected credit losses for the three and nine months ended September 30, 2026, and 2025:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025	2026	2025
Allowance for expected credit losses beginning balance	$ 10,563	$ 8,909	$ 8,308	$ 7,418
Current period provision for expected credit losses	1,242	629	4,833	2,466
Write-offs charged against the allowance and other	(176)	(112)	(1,435)	(870)
Effect of changes in foreign exchange rates	(46)	(13)	(123)	399
Allowance for expected credit losses ending balance	$ 11,583	$ 9,413	$ 11,583	$ 9,413

10. Business segment information

The Company's operations are managed through two reporting segments: Global Spine and Global Limb Reconstruction. These reporting segments represent the operating segments for which the President and Chief Executive Officer, who is also the Chief Operating Decision Maker ("CODM"), reviews financial information and makes resource allocation decisions among businesses. The primary metric used by the CODM in managing the Company is adjusted earnings before interest, tax, depreciation, and amortization ("adjusted EBITDA", a non-GAAP financial measure). Adjusted EBITDA represents earnings before interest income (expense), income taxes, depreciation and amortization, and excludes the impact of share-based compensation and long-term

incentive plan expense; gains and losses related to changes in foreign exchange rates; charges related to the Merger and other strategic investments; restructuring costs and impairments related to the discontinuation of the M6 product lines (as defined in Note 15); acquisition-related fair value adjustments; gains and/or losses on investments; litigation and investigation charges; refunds associated with the Employee Retention Credit established by the Coronavirus Aid, Relief, and Economic Security Act; and certain costs associated with employee transitions.

Corporate activities are comprised of operating expenses not directly identifiable within the two reporting segments, such as human resources, finance, legal, and information technology functions. The Company neither discretely allocates assets, other than goodwill, to its operating segments nor evaluates the operating segments using discrete asset information.

Global Spine

The Global Spine reporting segment offers two primary product categories: (i) Therapeutic Solutions and (ii) Spinal Implants, Biologics, and Enabling Technologies.

The Therapeutic Solutions product category manufactures, distributes, and provides support services for market-leading bone growth stimulation devices that enhance bone fusion. These Class II medical devices with special controls are indicated as an adjunctive, noninvasive treatment to improve fusion success rates in the cervical and lumbar spine as well as a therapeutic treatment for non-spinal, appendicular fractures, treating both fresh or nonunion fractures. These products are sold almost exclusively in the U.S., using distributors and direct sales representatives to provide these devices to healthcare providers and their patients.

Spinal Implants, Biologics, and Enabling Technologies is comprised of (i) a broad portfolio of spine fixation implant products used in surgical procedures of the spine, (ii) one of the most comprehensive biologics portfolios in both the demineralized bone matrix and cellular allograft market segments, and (iii) image-guided surgical solutions to facilitate degenerative, minimally invasive, and complex surgical procedures. Spinal Implants, Biologics, and Enabling Technologies products are sold through a network of distributors and sales representatives to hospitals and healthcare providers on a global basis for Spinal Implants and Enabling Technologies, and primarily within the U.S. for Biologics.

Global Limb Reconstruction

The Global Limb Reconstruction reporting segment offers products and solutions for the underserved limb reconstruction market that encompasses four pillars: deformity correction, limb lengthening, complex fracture management, and limb preservation. This reporting segment specializes in the design, development, and marketing of external and internal fixation limb reconstruction products that are coupled with enabling digital technologies to serve the complete patient treatment pathway. The Company sells these products worldwide through a global network of distributors and sales representatives to hospitals, healthcare organizations, and healthcare providers.

The following table presents adjusted EBITDA, the primary metric used in managing the Company, by reporting segment:

	Three Months Ended September 30, 2026			Nine Months Ended September 30, 2026		
(Unaudited, U.S. Dollars, in thousands)	Global Spine	Global Limb Reconstruction	Total	Global Spine	Global Limb Reconstruction	Total
Segment revenues	$ 171,449	$ 36,413	$ 207,862	$ 508,544	$ 106,959	$ 615,503
Less:						
Non-GAAP Cost of sales	47,037	13,754	60,791	134,440	43,033	177,473
Non-GAAP Sales, general, and administrative	95,150	19,692	114,842	294,163	61,981	356,144
Non-GAAP Research and development	10,840	3,163	14,003	33,765	9,245	43,010
Other segment expenses	82	174	256	404	1,132	1,536
Add:						
Non-GAAP Depreciation, amortization, and share-based compensation expense	8,443	3,356	11,799	27,001	9,895	36,896
Segment Adjusted EBITDA	$ 26,783	$ 2,986	$ 29,769	$ 72,773	$ 1,463	$ 74,236
Reconciling items:						
Corporate operating expenses			$ 3,733			$ 18,368
Interest expense, net			5,889			17,638
Depreciation and amortization			13,172			39,938
Share-based compensation and long-term incentive plan expense			6,969			21,552
Foreign exchange impact			744			2,390
SeaSpine merger-related costs			412			915
Restructuring costs and impairments related to M6 product lines			464			1,802
Strategic investments			509			2,093
Acquisition-related fair value adjustments			1,396			3,014
Interest and loss on investments			—			(16)
Litigation and investigation costs			301			5,402
Employee Retention Credit			—			(951)
Employee transition costs			1,428			3,170
Loss before income taxes			$ (5,248)			$ (41,079)

(Unaudited, U.S. Dollars, in thousands)	Three Months Ended September 30, 2025			Nine Months Ended September 30, 2025		
	Global Spine	Global Limb Reconstruction	Total	Global Spine	Global Limb Reconstruction	Total
Segment Revenues	$ 172,063	$ 33,571	$ 205,634	$ 505,723	$ 96,678	$ 602,401
Less:						
Non-GAAP Cost of sales	44,016	12,611	56,627	131,414	36,748	168,162
Non-GAAP Sales, general, and administrative	93,288	20,379	113,667	282,995	56,290	339,285
Non-GAAP Research and development	11,051	2,764	13,815	34,436	8,290	42,726
Other segment expenses (benefits)	2,223	(37)	2,186	9,130	(206)	8,924
Add:						
Non-GAAP Depreciation, amortization, and share-based compensation expense	9,157	4,037	13,194	28,843	8,793	37,636
Segment Adjusted EBITDA	$ 30,642	$ 1,891	$ 32,533	$ 76,591	$ 4,349	$ 80,940
Reconciling items:						
Corporate operating expenses			$ 7,951			$ 24,283
Interest expense, net			4,681			13,137
Depreciation and amortization			12,941			64,243
Share-based compensation and long-term incentive plan expense			7,181			21,474
Foreign exchange impact			571			(3,224)
SeaSpine merger-related costs			126			6,142
Restructuring costs and impairments related to M6 product lines			6			15,485
Strategic investments			227			4,094
Acquisition-related fair value adjustments			(427)			(1,800)
Interest and loss on investments			(10)			(41)
Litigation and investigation costs			21,548			28,619
Employee Retention Credit			—			(2,854)
Loss before income taxes			$ (22,262)			$ (88,618)

The following table presents depreciation, amortization, and impairment by reporting segment:

(Unaudited, U.S. Dollars, in thousands)	Three Months Ended September 30,		Nine Months Ended September 30,	
	2026	2025	2026	2025
Global Spine	$ 9,842	$ 9,372	$ 30,046	$ 56,125
Global Limb Reconstruction	2,852	2,986	8,437	6,341
Corporate	478	583	1,455	1,777
Total	$ 13,172	$ 12,941	$ 39,938	$ 64,243

Geographical information

The table below presents net sales by geographic destination for each reporting segment and for the consolidated Company:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	**2026**	**2025**	**2026**	**2025**
Global Spine				
U.S.	$ 156,783	$ 160,143	$ 469,190	$ 474,696
International	14,666	11,920	39,354	31,027
Total Global Spine	171,449	172,063	508,544	505,723
Global Limb Reconstruction				
U.S.	10,129	10,198	29,083	29,064
International	26,284	23,373	77,876	67,614
Total Global Limb Reconstruction	36,413	33,571	106,959	96,678
Consolidated				
U.S.	166,912	170,341	498,273	503,760
International	40,950	35,293	117,230	98,641
Net sales	$ 207,862	$ 205,634	$ 615,503	$ 602,401

The following data includes net sales by geographic area:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	**2026**	**2025**	**2026**	**2025**
U.S.	$ 166,912	$ 170,341	$ 498,273	$ 503,760
Italy	6,121	5,658	18,027	16,357
France	2,867	2,691	8,765	8,500
United Kingdom	3,191	3,645	10,758	9,800
Germany	2,591	2,311	7,076	6,784
Brazil	1,358	1,200	4,310	3,323
Others	24,822	19,788	68,294	53,877
Net sales	$ 207,862	$ 205,634	$ 615,503	$ 602,401

The following data includes property, plant, and equipment by geographic area:

(U.S. Dollars, in thousands)	**September 30, 2026**	**December 31, 2025**
	(Unaudited)	
U.S.	$ 119,825	$ 114,483
Italy	9,071	9,893
Germany	855	1,360
Others	3,637	3,663
Total	$ 133,388	$ 129,399

11. Acquisition-related amortization, impairment, and remeasurement

Acquisition-related amortization, impairment, and remeasurement consists of (i) amortization and impairment related to intangible assets acquired through business combinations or asset acquisitions and (ii) remeasurement of any related contingent consideration arrangements. Components of acquisition-related amortization, impairment, and remeasurement are as follows:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025	2026	2025
Amortization and impairment of acquired intangibles	$ 2,998	$ 3,120	$ 8,998	$ 25,347
Changes in fair value of contingent consideration	1,396	(427)	3,014	(1,800)
Total	$ 4,394	$ 2,693	$ 12,012	$ 23,547

12. Share-based compensation

Components of share-based compensation expense are as follows:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025	2026	2025
Cost of sales	$ 234	$ 368	$ 1,027	$ 1,297
Sales, general, and administrative	6,102	6,448	18,824	19,012
Research and development	408	365	1,161	1,165
Total	$ 6,744	$ 7,181	$ 21,012	$ 21,474

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, U.S. Dollars, in thousands)	2026	2025	2026	2025
Stock options	$ 1,424	$ 1,389	$ 4,512	$ 3,852
Market-based stock options	94	447	371	1,685
Time-based restricted stock units	3,128	3,371	9,594	9,893
Market-based / performance-based restricted stock units	1,766	1,585	5,389	4,655
Stock purchase plan	332	389	1,146	1,389
Total	$ 6,744	$ 7,181	$ 21,012	$ 21,474

During each of the three months ended September 30, 2026, and 2025, the Company issued fewer than 0.1 million shares, respectively, of common stock related to stock purchase plan issuances, stock option exercises, and the vesting of restricted stock units. During the nine months ended September 30, 2026, and 2025, the Company issued 0.9 million and 1.0 million shares, respectively, of common stock related to stock purchase plan issuances, stock option exercises, and the vesting of restricted stock units.

13. Income taxes

Generally, income tax provisions for interim periods are based on an estimated annual income tax rate, adjusted for discrete tax items, with any changes affecting the estimated annual effective tax rate recorded in the interim period in which the change occurs. Due to losses in the Company's U.S., Canadian and Italian operations for which no tax benefit is recognized, the Company determined the estimated annual effective tax rate method would not provide a reliable estimate of the Company's overall annual effective tax rate. As such, the Company has calculated the tax provision using the actual effective rate for the nine months ended September 30, 2026. Due to the impact of temporary differences on the Company's current U.S. tax liability for which no deferred tax benefit is recognized, the Company's effective tax rate may vary in future quarters.

For the three months ended September 30, 2026, and 2025, the effective tax rate was (15.7%) and (2.4%), respectively. For the nine months ended September 30, 2026, and 2025, the effective tax rate was (4.1%) and (1.5%), respectively. The primary factors affecting the Company's effective tax rate for the three and nine months ended September 30, 2026, were certain losses for which no tax benefit is recognized and tax amortization on certain acquired intangibles.

In October 2026, the Internal Revenue Service notified the Company of their intention to audit the tax year 2024. The Company does not currently expect the ultimate resolution to have a material impact on the Company's consolidated financial statements.

14. Earnings per share (EPS)

For the three and nine months ended September 30, 2026, no adjustments were made to net income (loss) for purposes of calculating basic and diluted EPS under the treasury stock method. The following is a reconciliation of the weighted average shares used in diluted EPS computations.

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited, In thousands)	2026	2025	2026	2025
Weighted average common shares-basic	41,178	39,766	40,842	39,468
Effect of dilutive securities				
Unexercised stock options and stock purchase plan	—	—	—	—
Unvested restricted stock units	—	—	—	—
Weighted average common shares-diluted	41,178	39,766	40,842	39,468

There were 8.6 million and 9.2 million weighted average outstanding options, time-based restricted stock units, performance-based stock units, and market-based stock units not included in the diluted EPS computation for the three months ended September 30, 2026, and 2025, respectively, and 8.6 million weighted average outstanding options, time-based restricted stock units, performance-based stock units, and market-based stock units not included in the diluted EPS computation for both the nine months ended September 30, 2026, and 2025, respectively, because either (i) inclusion of these awards was anti-dilutive, or (ii) for performance-based stock units and market-based stock units, all necessary conditions had not been satisfied by the end of the respective period.

15. Discontinuation of M6 product lines

In February 2025, the Company announced the discontinuation of its M6-C artificial cervical disc and M6-L artificial lumbar disc product lines (together, the "M6 artificial discs" or "M6 product lines") in order to allocate associated resources and investment to more profitable growth opportunities. Financial results for the Company's M6 product lines continue to be presented within the Company's consolidated statements of operations and comprehensive loss. A summary of impairment charges recognized during the three and nine months ended September 30, 2025, and the associated financial statement lines in which such costs were recognized is shown in the table below. All such changes are included within the Company's Global Spine reporting segment. All related inventory; property, plant, and equipment; and intangible asset balances were fully impaired in the prior year; therefore, there were no further impairments recorded on these assets in the current year.

(Unaudited, U.S. Dollars, in thousands)	Financial Statement Line Item	Three Months Ended September 30, 2025	Nine Months Ended September 30, 2025
Inventory reserve charges	Cost of sales	$ —	$ 11,251
Impairment of property, plant, and equipment	Operating expenses	—	6,834
Impairment of developed technology intangible asset	Acquisition-related amortization, impairment, and remeasurement	—	14,097
Loss on M6 inventories and long-lived assets held for sale		$ —	$ 32,182

16. Subsequent Events

Subsequent to September 30, 2026, the Company and a distributor partner amended a previously exercised purchase option arrangement to defer the acquisition and incorporate the exercised purchase right into a purchase right exercisable in 2029. Accordingly, the Company is not currently required to complete the acquisition as a result of the prior exercise notice.

Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations

The following discussion and analysis of Orthofix Medical Inc.'s (sometimes referred to as the "Company," "we," "us" or "our") financial condition and results of operations should be read in conjunction with the discussion under the heading "Forward-Looking Statements" and our condensed consolidated financial statements and related notes thereto appearing elsewhere in this Form 10-Q.

Executive Summary

We are a global medical technology company dedicated to advancing healing and restoring mobility for patients with complex musculoskeletal conditions. Headquartered in Lewisville, Texas, we offer a differentiated portfolio of spinal implants, therapeutic solutions, limb reconstruction systems, biologics and enabling technologies, including the 7D FLASH Navigation System. Our technology-enabled solutions are designed to support surgeons across the continuum of care and improve outcomes for patients. Learn more at Orthofix.com and follow us on LinkedIn. Information included on our website is not incorporated into, nor does it otherwise create a part of, this report.

Notable financial metrics in the third quarter of 2026 and recent achievements include the following:

- Total reported net sales of $207.9 million, an increase of 1% on a reported basis and 2% on a pro forma constant currency basis compared with the prior year period.
- Reported net loss of $6.1 million and non-GAAP adjusted EBITDA of $26.0 million.
- Global Spine Fixation net sales increased 1% on a reported and constant currency basis compared with the prior year period.
- Global Limb Reconstruction net sales increased 8% on a reported basis and 9% on a constant currency basis compared with the prior year period.
- Therapeutic Solutions net sales increased 4% compared with the prior year period.

Results of Operations

The following table provides certain items in our condensed consolidated statements of operations as a percent of net sales:

	Three Months Ended September 30,		Nine Months Ended September 30,	
(Unaudited)	2026 (%)	2025 (%)	2026 (%)	2025 (%)
Net sales	100.0	100.0	100.0	100.0
Cost of sales	29.5	27.8	29.2	32.0
Gross profit	70.5	72.2	70.8	68.0
Sales, general, and administrative	60.3	72.0	64.7	69.3
Research and development	7.4	7.2	7.6	8.4
Acquisition-related amortization, impairment, and remeasurement	2.1	1.3	1.9	3.9
Operating income (loss)	0.7	(8.3)	(3.4)	(13.6)
Net loss	(2.9)	(11.1)	(7.0)	(14.9)

Net Sales by Product Category and Reporting Segment

Our operations are managed through two reporting segments: Global Spine and Global Limb Reconstruction. The following tables provide net sales by product category and reporting segment:

	Three Months Ended September 30,			
(Unaudited, U.S. Dollars, in millions)	2026	2025	Change	Constant Currency Change
Therapeutic Solutions	$ 64.0	$ 61.2	4.5%	4.5%
Spinal Implants, Biologics and Enabling Technologies*	107.4	108.6	(1.1%)	(1.1%)
Global Spine*	171.4	169.8	0.9%	0.9%
Global Limb Reconstruction	36.4	33.6	8.5%	9.4%
Pro forma net sales*	**207.8**	**203.4**	**2.2%**	**2.3%**
Impact from discontinuation of M6 product lines	0.1	2.2	(96.8%)	(96.8%)
Reported net sales	$ 207.9	$ 205.6	1.1%	1.2%

	Nine Months Ended September 30,			
(Unaudited, U.S. Dollars, in millions)	2026	2025	Change	Constant Currency Change
Therapeutic Solutions	$ 185.9	$ 178.8	3.9%	3.9%
Spinal Implants, Biologics and Enabling Technologies*	322.2	317.8	1.4%	1.4%
Global Spine*	508.1	496.6	2.3%	2.3%
Global Limb Reconstruction	107.0	96.7	10.6%	8.0%
Pro forma net sales*	**615.1**	**593.3**	**3.7%**	**3.2%**
Impact from discontinuation of M6 product lines	0.4	9.1	(95.7%)	(95.9%)
Reported net sales	$ 615.5	$ 602.4	2.2%	1.7%

* Results above for each of Spinal Implants, Biologics, and Enabling Technologies; Global Spine; and pro forma net sales exclude the impact from discontinuation of its M6-C artificial cervical disc and M6-L artificial lumbar disc product lines (together, the "M6 artificial discs" or "M6 product lines"). Since pro forma net sales represent a non-GAAP measure, see the reconciliation above of the Company's pro forma net sales to its reported figures under U.S. GAAP. The Company's reported figures under U.S. GAAP represent each of the pro forma line items discussed above plus the impact from discontinuation of the M6 product lines.

Global Spine

Global Spine offers the following product categories:

- Therapeutic Solutions manufactures, distributes, sells, and provides support services for market-leading devices used adjunctively in high-risk spinal fusion procedures and treats both nonunion and acute fractures in the orthopedic space. Therapeutic Solutions uses distributors and a direct sales channel to sell its devices and provide associated support services to hospitals, healthcare providers, and patients in the U.S.
- Spinal Implants, Biologics, and Enabling Technologies is comprised of a broad portfolio of spine fixation implant products used in surgical procedures of the spine, one of the most comprehensive biologics portfolios in both the demineralized bone matrix and cellular allograft market segments and image-guided surgical solutions to facilitate degenerative, minimally invasive, and complex surgical procedures. Spinal Implants, Biologics, and Enabling Technologies products are sold through a network of distributors and sales representatives to hospitals and healthcare providers on a global basis for Spinal Implants and Enabling Technologies, and primarily within the U.S. for Biologics.

Three months ended September 30, 2026 compared to 2025

Net sales of $171.4 million, a decrease of $0.6 million or 0.4% on a reported basis compared to the prior year period. Net sales from the M6 product lines decreased $2.2 million as a result of the discontinuation of the M6 product lines in 2025 to focus resources and investments on more profitable growth opportunities. Spinal Implants, Biologics, and Enabling Technologies excluding net sales from the M6 product lines decreased $1.2 million, or 1.1% as growth among our Top 30 distributors slowed compared with prior quarters, constrained by supplier related delays, instrument-set availability and the inefficient allocation of existing assets while our smaller distributors continued to underperform. These decreases were partially offset by an increase of $2.7 million in net sales from

Therapeutic Solutions driven by an increase in gross order volumes from our continued investment in our direct sales channels for both the spine and fracture market.

Nine months ended September 30, 2026 compared to 2025

Net sales of $508.5 million, an increase of $2.8 million or 0.6% on a reported basis compared to prior year. Therapeutic Solutions net sales increased $7.0 million driven by an increase in gross order volumes from our continued investment in our direct sales channels for both the spine and fracture market. Spinal Implants, Biologics, and Enabling Technologies excluding sales from the M6 product lines increased $4.5 million primarily due to continued sales growth from our highest-volume distribution partners within Spine Fixation and strong growth in our International markets partially offset by a decline in sales from the M6 product lines as a result of the discontinuation of the M6 product lines in 2025, underperformance with our smaller distributors and our strategy to incentivize purchase commitments of our Spine Fixation and Biologics products over the term of each agreement, over the pursuit of capital sales with our Enabling Technology Voyager Earnout program.

Global Limb Reconstruction

Global Limb Reconstruction offers products and solutions for the underserved limb reconstruction market that encompasses four pillars: deformity correction, limb lengthening, complex fracture management, and limb preservation. Global Limb Reconstruction sells its products through a global network of distributors and sales representatives to hospitals, healthcare organizations, and healthcare providers.

Three months ended September 30, 2026 compared to 2025

Net sales of $36.4 million, an increase of $2.8 million or 8.5% on a reported basis and 9.4% on a constant currency basis compared to the prior year period. International sales increased $3.2 million, or 13.7% on a constant currency basis, driven by favorable timing of stocking distributor orders and commercial execution.

This growth was partially offset by a decline of net sales in the U.S. of $0.1 million, or 0.7%, and $0.3 million due to movement in foreign currency exchange rates, which had an unfavorable impact during the quarter.

Nine months ended September 30, 2026 compared to 2025

Net sales of $107.0 million, an increase of $10.3 million or 10.6% on a reported basis and 8.0% on a constant currency basis compared to prior year. International sales increased $7.7 million, or 11.4% on a constant currency basis, primarily driven by sales of new products launched in the past three years. Net sales increased $2.6 million due to movement in foreign currency exchange rates, which had a favorable impact during the year.

Gross Profit

	Three Months Ended September 30,			Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	**2026**	**2025**	**% Change**	**2026**	**2025**	**% Change**
Net sales	$ 207,862	$ 205,634	1.1%	$ 615,503	$ 602,401	2.2%
Cost of sales	61,316	57,111	7.4%	179,704	192,726	(6.8%)
Gross profit	$ 146,546	$ 148,523	(1.3%)	$ 435,799	$ 409,675	6.4%
Gross margin	70.5%	72.2%	(1.7%)	70.8%	68.0%	2.8%

Three months ended September 30, 2026 compared to 2025

Gross profit decreased $2.0 million, or 1.3%, compared to the prior year period, primarily driven by changes in product and geography mix within our Spine Fixation portfolio.

Nine months ended September 30, 2026 compared to 2025

Gross profit increased $26.1 million, or 6.4%, compared to prior year. The gross margin improvement of 2.8% was attributable to a favorable impact from inventory reserves and manufacturing variances driven by the discontinuation of the M6 product lines and from inventory charges for certain product lines that were rationalized in the prior year as part of our integration activities following the Merger.

Sales, General, and Administrative Expense

(Unaudited, U.S. Dollars, in thousands)	Three Months Ended September 30, 2026	2025	% Change	Nine Months Ended September 30, 2026	2025	% Change
Sales, general, and administrative	$ 125,330	$ 148,102	(15.4%)	$ 398,271	$ 417,576	(4.6%)
As a percentage of net sales	60.3%	72.0%	(11.7%)	64.7%	69.3%	(4.6%)

Three months ended September 30, 2026 compared to 2025

Sales, general, and administrative expense decreased $22.8 million compared to the prior year period primarily due to decreases of (i) $21.2 million associated with certain legal matters, including the arbitration claims with former executives and the related securities class action complaints and (ii) lower commissions.

Nine months ended September 30, 2026 compared to 2025

Sales, general, and administrative expense decreased $19.3 million compared to the prior year primarily due to a decrease of $22.8 million associated with certain legal matters, including the arbitration claims with former executives and the related securities class action complaints partially offset by increases from higher commissions associated with increased sales.

Research and Development Expense

(Unaudited, U.S. Dollars, in thousands)	Three Months Ended September 30, 2026	2025	% Change	Nine Months Ended September 30, 2026	2025	% Change
Research and development	$ 15,277	$ 14,774	3.4%	$ 46,541	$ 50,474	(7.8%)
As a percentage of net sales	7.3%	7.2%	0.1%	7.6%	8.4%	(0.8%)

Three months ended September 30, 2026 compared to 2025

Research and development expense increased $0.5 million compared to the prior year period primarily attributable to increased costs associated with clinical trials and prototypes and from increased legal and professional fees.

Nine months ended September 30, 2026 compared to 2025

Research and development expense decreased $3.9 million compared to prior year primarily attributable to a decrease of $5.4 million related to impairments and costs associated with the discontinuation of the M6 product lines and other organizational restructuring activities in 2025 partially offset by an increase in employee compensation and benefits costs.

Acquisition-related Amortization, Impairment, and Remeasurement

(Unaudited, U.S. Dollars, in thousands)	Three Months Ended September 30, 2026	2025	% Change	Nine Months Ended September 30, 2026	2025	% Change
Acquisition-related amortization, impairment, and remeasurement	$ 4,394	$ 2,693	63.2%	$ 12,012	$ 23,547	(49.0%)
As a percentage of net sales	2.1%	1.3%	0.8%	2.0%	3.9%	(1.9%)

Acquisition-related amortization, impairment, and remeasurement consists of (i) amortization and impairment related to intangible assets acquired through business combinations or asset acquisitions and (ii) remeasurement of related contingent consideration arrangements, which are recognized immediately upon acquisition.

Three months ended September 30, 2026 compared to 2025

Acquisition-related amortization, impairment, and remeasurement increased $1.7 million compared to the prior year period due to an increase of $1.8 million associated with the remeasurement of a contingent consideration obligation with Lattus Spine LLC ("Lattus") assumed in the merger with SeaSpine Holdings Corporation (the "Merger"). This increase was partially offset by a $0.1 million decrease in amortization expense resulting from certain acquired intangibles becoming fully amortized in the prior year.

Nine months ended September 30, 2026 compared to 2025

Acquisition-related amortization, impairment, and remeasurement decreased $11.5 million compared to the prior year primarily driven by a $16.3 million decrease in amortization expense associated with the impairment of certain acquired intangible assets

from the discontinuation of the M6 product lines partially offset by an increase of $4.8 million associated with the remeasurement of a contingent consideration obligation with Lattus assumed in the Merger.

Non-operating Income and Expense

	Three Months Ended September 30,			Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	% Change	2026	2025	% Change
Interest expense, net	$ (5,889)	$ (4,681)	25.8%	$ (17,638)	$ (13,137)	34.3%
Other income (expense), net	(904)	(535)	69.0%	(2,416)	6,441	(137.5%)

Three months ended September 30, 2026 compared to 2025

Interest expense, net increased $1.2 million compared to the prior year period primarily due to the funding of the $65.0 million Term B Loan in January 2026.

Other expense, net increased $0.4 million compared to the prior year period primarily due to the unfavorable change associated with foreign currency exchange rates, as we recorded a non-cash remeasurement loss of $0.7 million in the third quarter 2026 compared to a loss of $0.5 million in the third quarter of 2025.

Nine months ended September 30, 2026 compared to 2025

Interest expense, net increased $4.5 million compared to prior year due to (i) the funding of the $65.0 million Term B Loan in January 2026, (ii) an unfavorable change associated with interest income earned on certain Employee Retention Credit refunds received during 2025 and 2026, and (iii) changes in finance lease obligations and the amortization of debt issuance costs.

Other income (expense), net decreased $8.9 million compared to prior year primarily due to the unfavorable change associated with foreign currency exchange rates, as we recorded a non-cash remeasurement loss of $2.3 million in 2026 compared to a gain of $3.2 million in 2025. In addition, there was an unfavorable change of $3.2 million associated with the receipt of Employee Retention Credit refunds received during 2025 and 2026.

Income Taxes

	Three Months Ended September 30,			Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	% Change	2026	2025	% Change
Income tax expense	$ 826	$ 533	55.0%	$ 1,701	$ 1,352	25.8%
Effective tax rate	(15.7%)	(2.4%)	(13.3%)	(4.1%)	(1.5%)	(2.6%)

Three months ended September 30, 2026 compared to 2025

Income tax expense increased $0.3 million compared to the prior year period primarily due to increased tax on foreign operations and tax expense related to certain long-lived intangible assets. The primary factor affecting our tax expense for the third quarter of 2026 was tax amortization on certain acquired intangibles and financial statement losses for which no benefit is recognized.

Nine months ended September 30, 2026 compared to 2025

Income tax expense increased $0.3 million compared to the prior year primarily due to increased tax on foreign operations and tax expense related to certain long-lived intangible assets. The primary factor affecting our tax expense for 2026 was tax amortization on certain acquired intangibles and financial statement losses for which no benefit is recognized.

Liquidity and Capital Resources

Cash, cash equivalents, and restricted cash at September 30, 2026, totaled $86.9 million compared to $85.1 million at December 31, 2025. The following table presents the net change in cash, cash equivalents, and restricted cash for the nine months ended September 30, 2026, and 2025, respectively:

	Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	Change
Net cash provided by (used in) operating activities	$ (23,471)	$ 5,650	$ (29,121)
Net cash used in investing activities	(37,664)	(23,727)	(13,937)
Net cash provided by (used in) financing activities	63,228	(3,163)	66,391
Effect of exchange rate changes on cash	(288)	1,448	(1,736)
Net change in cash, cash equivalents, and restricted cash	$ 1,805	$ (19,792)	$ 21,597

The following table presents free cash flow, a non-GAAP financial measure, which is calculated by subtracting capital expenditures from net cash from operating activities:

	Nine Months Ended September 30,		
(Unaudited, U.S. Dollars, in thousands)	2026	2025	Change
Net cash provided by (used in) operating activities	$ (23,471)	$ 5,650	$ (29,121)
Capital expenditures	(37,810)	(23,749)	(14,061)
Free cash flow	$ (61,281)	$ (18,099)	$ (43,182)

Operating Activities

Cash flows from operating activities decreased $29.1 million as a result of (i) a favorable change in net loss of $42.8 million, (ii) a decrease of $32.2 million associated with non-cash gains and losses, such as depreciation, amortization, and impairments, and inventory reserve expenses, and (iii) a decrease of $44.1 million relating to changes in working capital accounts, primarily attributable to changes in accounts receivable, inventories, and other current liabilities.

Two of our primary working capital accounts are accounts receivable and inventory. Days sales in receivables were 58 days as of September 30, 2026, compared to 59 days as of September 30, 2025 (calculated using third quarter net sales and ending accounts receivable). Inventory turns decreased to 1.3 times as of September 30, 2026 compared to 1.5 times as of September 30, 2025 (calculated using trailing twelve-month cost of goods sold and ending net inventories).

Investing Activities

Cash flows used in investing activities increased $13.9 million due to an increase in spend of $14.1 million in capital expenditures and partially offset by an increase of $0.1 million in other investing activities.

Financing Activities

Cash flows from financing activities increased $66.4 million primarily attributable to (i) an increase of $64.0 million associated with net borrowing activities related to our credit facilities in the first quarter of 2026 compared to the prior year period, (ii) a favorable change of $1.8 million associated with contingent consideration payments in 2026 compared to the prior year period, and (iii) a favorable change of $0.5 million in debt issuance costs associated with our credit facilities in 2026 compared to the prior year period.

Credit Facilities

On November 7, 2024, we entered into a $275.0 million secured credit agreement (the "Credit Agreement") with Oxford Finance LLC, as administrative agent and as collateral agent ("Oxford") and certain lenders party thereto, including Oxford, K2 HealthVentures LLC, and HSBC Ventures USA Inc. Certain of our foreign subsidiaries joined the Credit Agreement as either a borrower or guarantor shortly after the signing date. The Credit Agreement provides for a $160.0 million senior secured term loan (the "Initial Term Loan") and a $65.0 million senior secured delayed draw term loan facility (the "Term B Loan") which Term B Loan was fully funded on January 15, 2026. In addition, at Oxford's discretion, an additional $50.0 million of draw capacity is available

through January 1, 2029 (the "Term C Loan" and, together with the Term B Loan, the "Delayed Draw Term Loans" and collectively with the Initial Term Loan, the "Credit Facilities").

The Initial Term Loan and Delayed Draw Term Loans, to the extent ultimately drawn, will each mature in November 2029, following an interest-only payment period ending December 2028, and monthly amortization of principal and accrued interest between January 2029 and November 2029.

The Initial Term Loan and Delayed Draw Term Loans, to the extent ultimately drawn, are subject to, among other conditions, our continued compliance with a pro-forma total debt-to-EBITDA leverage ratio of less than 4.0x. EBITDA is a non-GAAP financial measure which represents earnings before interest income (expense), income taxes, depreciation, amortization, and other negotiated addbacks and adjustments.

The Credit Agreement contains financial covenants requiring us to maintain a minimum level of liquidity at all times and to maintain a maximum total debt-to-EBITDA leverage ratio (measured on a quarterly basis) during the term of the facility. As of September 30, 2026, we were in compliance with all required financial covenants.

As of September 30, 2026, we had $225.0 million of outstanding borrowings under the Credit Agreement related to the Initial Term Loan and the Term B Loan. We have not made any borrowings under the Term C Loan as of September 30, 2026.

As of September 30, 2026, we had no borrowings on our available lines of credit in Italy, which provide up to an aggregate amount of €5.5 million ($6.2 million).

Other

For information regarding contingencies, see Note 7 of the Notes to the Unaudited Condensed Consolidated Financial Statements contained herein.

Lattus Contingent Consideration

Under the terms of a contingent consideration obligation in a purchase agreement assumed in the Merger, we may be required to make installment payments to Lattus (the "Lattus Contingent Consideration") at certain dates based on future net sales of certain products (the "Lateral Products"). The estimated fair value of the Lattus Contingent Consideration as of September 30, 2026, was $4.9 million. The actual amount ultimately paid could be higher or lower than the estimated fair value of the Lattus Contingent Consideration. As of September 30, 2026, we classified the remaining Lattus Contingent Consideration liability of $4.9 million within other current liabilities.

In the third quarter of 2026, we made installment payments related to the Lattus Contingent Consideration of $6.0 million. For additional discussion of this matter, see Note 6 of the Notes to the Unaudited Condensed Consolidated Financial Statements.

Off-balance Sheet Arrangements

As of September 30, 2026, we did not have any off-balance sheet arrangements that have or are reasonably likely to have a current or future effect on our financial condition, changes in financial condition, revenues or expenses, results of operations, cash flows, liquidity, capital expenditures or capital resources that are material to investors.

Contractual Obligations

There have been no material changes in any of our material contractual obligations as disclosed in our Annual Report on Form 10-K for the year ended December 31, 2025 ("2025 10-K").

Critical Accounting Estimates

Our discussion of operating results is based upon the condensed consolidated financial statements and accompanying notes. The preparation of these statements requires us to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amount of revenues and expenses during the reporting period. Our critical accounting estimates are described in Item 7 of our 2025 10-K. There have been no significant changes to our critical accounting estimates during the quarter covered by this report.

Recently Issued Accounting Pronouncements

See Note 2 of the Notes to the Unaudited Condensed Consolidated Financial Statements for detailed information regarding the status of recently issued or adopted accounting pronouncements.

Non-GAAP Financial Measures

We believe that providing non-GAAP financial measures that exclude certain items provides investors with greater transparency to the information used by senior management in its financial and operational decision-making. We believe it is important to provide investors with the same non-GAAP financial measures used to supplement information regarding the performance and underlying trends of our business operations to facilitate comparisons to historical operating results and internally evaluate the effectiveness of our operating strategies. Disclosure of these non-GAAP financial measures also facilitates comparisons of our underlying operating performance with other companies in the industry that also supplement their U.S. GAAP results with non-GAAP financial measures.

The non-GAAP financial measures used in this filing may have limitations as analytical tools and should not be considered in isolation or as a replacement for U.S. GAAP financial measures. Some limitations associated with the use of these non-GAAP financial measures are that they exclude items that reflect an economic cost that can have a material effect on cash flows.

Constant Currency

Constant currency is calculated by using foreign currency rates from the comparable, prior year period to present net sales at comparable rates. Constant currency can be presented for numerous U.S. GAAP measures but is most commonly used by management to analyze net sales without the impact of changes in foreign currency rates.

Free Cash Flow

Free cash flow is calculated by subtracting capital expenditures from net cash from operating activities. Management uses free cash flow as an important indicator of how much cash is generated or used by our normal business operations, including capital expenditures. Management uses free cash flow as a measure of progress on its capital efficiency and cash flow initiatives.

Item 3. Quantitative and Qualitative Disclosures About Market Risk

There have been no material changes to our market risks as disclosed in our 2025 10-K.

Item 4. Controls and Procedures

Evaluation of Disclosure Controls and Procedures

We maintain disclosure controls and procedures (as defined in Rule 13a-15(e) of the Exchange Act) designed to provide reasonable assurance that the information required to be disclosed in reports filed or submitted under the Exchange Act are recorded, processed, summarized, and reported within the time periods specified in the SEC's rules and forms. These include controls and procedures designed to ensure that this information is accumulated and communicated to management, including our Chief Executive Officer and our Chief Financial Officer, as appropriate to allow timely decisions regarding required disclosure. Management, with the participation of the Chief Executive Officer and the Chief Financial Officer, evaluated the effectiveness of our disclosure controls and procedures as of September 30, 2026. Based on this evaluation, our Chief Executive Officer and our Chief Financial Officer have concluded that our disclosure controls and procedures were effective as of September 30, 2026.

Changes in Internal Control over Financial Reporting

There was no change in our internal control over financial reporting that occurred during the quarterly period covered by this report that has materially affected, or is reasonably likely to materially affect, our internal control over financial reporting.

PART II. OTHER INFORMATION

Item 1. Legal Proceedings

For information regarding legal proceedings, see Note 7 of the Notes to the Unaudited Condensed Consolidated Financial Statements contained herein, which is incorporated by reference into this Part II, Item 1.

Item 1A. Risk Factors

The U.S. Food and Drug Administration's ("FDA") reclassification of bone growth stimulator devices from Class III to Class II may increase competition and adversely affect our future sales.

We offer the market-leading bone growth stimulation platform and are the only company to provide both pulsed electromagnetic field (PEMF) and low-intensity pulsed ultrasound (LIPUS) bone healing solutions. Historically, our bone growth therapy products were regulated by the FDA as Class III medical devices, subject to the FDA's rigorous premarket approval (PMA) requirements.

The FDA has reclassified bone growth stimulator devices from Class III to Class II, subject to "special controls." These special controls include requirements for clinical data, specific non-clinical performance and biocompatibility data, and labeling, in addition to the Class II requirement that new devices demonstrate substantial equivalence to a legally marketed predicate device. While these controls are intended to provide reasonable assurance of safety and effectiveness, the Class II regulatory pathway is generally less onerous, time-consuming, and costly than the PMA process applicable to Class III devices.

As a result of this reclassification, competitors may be able to enter the market more readily by obtaining FDA clearance for bone growth stimulator devices that are substantially equivalent to existing products. Increased market entry could lead to heightened competition, pricing pressure, and greater marketing and promotional activity by competitors, which may reduce demand for our products or erode our market share.

Although we believe our clinical data, dual-technology platform, and brand recognition differentiate our products, we may be required to increase investments in research and development, clinical studies, sales, marketing, and post-market surveillance to maintain our competitive position. Any such increased costs, or a failure to effectively compete in a more crowded market, could adversely affect our revenues, margins, and results of operations.

Other than as disclosed above, there have been no material changes from the risk factors disclosed in "Part I, Item 1A. Risk Factors" in our Annual Report on Form 10-K for the year ended December 31, 2025.

Item 2. Unregistered Sales of Equity Securities and Use of Proceeds

We have not made any repurchases of our common stock during the third quarter of 2026.

Item 3. Defaults Upon Senior Securities

Not applicable.

Item 4. Mine Safety Disclosures

Not applicable.

Item 5. Other Information

Director and Officer Trading Arrangements

During the last fiscal quarter, none of our directors or officers (as defined in Rule 16a-1(f) of the Exchange Act) adopted, modified or terminated any contract, instruction, or written plan for the purchase or sale of our securities that was intended to satisfy the affirmative defense conditions of Rule 10b5-1(c) of the Exchange Act or any "non-Rule 10b5-1 trading arrangement."

Ratification of Certain Matters

On November 3, 2026, the Company's Board of Directors adopted resolutions ratifying (the "Ratification") under Section 204 of the General Corporation Law of the State of Delaware (the "DGCL"): (a) the grant of certain performance-based restricted stock units and time-based restricted stock units collectively with respect to a maximum aggregate of 441,695 shares (the "Equity Awards") of the Company's common stock (of which Equity Awards with respect to 241,695 shares of common stock may constitute putative stock) granted under the Company's 2012 Long-Term Incentive Plan, as amended (the "2012 LTIP"), due to the failure of the issuance of the Equity Awards to have been authorized in accordance with the 2012 LTIP and Delaware law and (b) an amendment to the 2012 LTIP, effective as of January 1, 2025, to increase the individual sublimit on the number of shares of stock subject to

certain awards that may be granted to individual award recipients in a fiscal year under the 2012 LTIP (the "Applicable Sublimit"), and thereby permit the grant of the Equity Awards. The Company inadvertently issued and delivered, on March 3, 2025, the Equity Awards to the Company's President and Chief Executive Officer in excess of the then-200,000 share Applicable Sublimit. Any claim that any defective corporate act or putative stock ratified pursuant to the Ratification is void or voidable due to the failure of authorization as described above or that the Delaware Court of Chancery should declare in its discretion that the Ratification in accordance with Section 204 of the DGCL not be effective or be effective only on certain conditions must be brought by the later of 120 calendar days from (i) the validation effective time (which was November 3, 2026) and (ii) the giving of this notice (which is deemed given on November 9, 2026, the date that this Quarterly Report on Form 10-Q has been filed with the Securities and Exchange Commission).

Item 6. Exhibits

10.1*	Amendment No. 6 to the Orthofix Medical Inc. Amended and Restated 2012 Long-Term Incentive Plan
31.1*	Rule 13a-14(a)/15d-14(a) Certification of Chief Executive Officer.
31.2*	Rule 13a-14(a)/15d-14(a) Certification of Chief Financial Officer.
32.1#	Section 1350 Certifications of each of the Chief Executive Officer and Chief Financial Officer.
101.INS*	Inline XBRL Instance Document (the instance document does not appear in the Interactive Data File because its XBRL tags are embedded within the Inline XBRL document).
101.SCH*	Inline XBRL Taxonomy Extension Schema Document.
101.CAL*	Inline XBRL Taxonomy Extension Calculation Linkbase Document.
101.DEF*	Inline XBRL Taxonomy Extension Definition Linkbase Document.
101.LAB*	Inline XBRL Taxonomy Extension Label Linkbase Document.
101.PRE*	Inline XBRL Taxonomy Extension Presentation Linkbase Document.
104*	Cover Page Interactive Data File (formatted as inline XBRL and contained in Exhibit 101).

* Filed herewith.
# Furnished herewith.

SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, the Registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

ORTHOFIX MEDICAL INC.

Date: November 9, 2026

By: /s/ MASSIMO CALAFIORE
Name: Massimo Calafiore
Title: President and Chief Executive Officer

Date: November 9, 2026

By: /s/ JULIE ANDREWS
Name: Julie Andrews
Title: Chief Financial Officer

Exhibit 10.1

AMENDMENT NO. 6 TO THE
ORTHOFIX MEDICAL INC.
AMENDED AND RESTATED
2012 LONG-TERM INCENTIVE PLAN

WHEREAS, the Board of Directors (the "***Board***") of Orthofix Medical Inc. (the "***Company***") heretofore established, and the Company's shareholders have approved, the Orthofix Medical Inc. Amended and Restated 2012 Long-Term Incentive Plan (as it has been or may be amended and/or restated from time to time, the "***Plan***"); and

WHEREAS, the Company desires to amend the Plan to increase the annual sublimit amount set forth in Section 6.2(b) of the Plan in order (i) to provide additional flexibility to the Company in making grants of equity compensation to recipients under the Plan and (ii) to permit the Company to continue to make grants of equity compensation at annual grant values determined by the Compensation and Talent Development Committee of the Board that take into account market competitiveness, significance of the individual to the Company's strategy and success, and the level of "total direct compensation" deemed to be market appropriate for such recipient; and

WHEREAS, Section 5.3 of the Plan permits the Board to amend the Plan.

NOW, THEREFORE, BE IT RESOLVED, pursuant to Section 5.3 of the Plan, the Plan is hereby amended as follows, effective as of January 1, 2025:

1. Section 6.2(b) is amended, so that Section 6.2(b), as amended and restated, reads in its entirety as follows:

   "(b) The maximum number of shares of Stock that may be granted under the Plan, other than pursuant to Options or SARs, to a Grantee other than an Outside Director is 1,000,000 shares per fiscal year."

2. Except as specifically set forth herein, the terms of the Plan shall be and remain unchanged.

Exhibit 31.1

CERTIFICATION

I, Massimo Calafiore, certify that:

1. I have reviewed this quarterly report on Form 10-Q for the quarterly period ended September 30, 2026, of Orthofix Medical Inc.;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

   a. designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

   b. designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

   c. evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

   d. disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has material affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

   a. all significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

   b. any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Dated: November 9, 2026

By: /s/ MASSIMO CALAFIORE
Name: Massimo Calafiore
Title: President and Chief Executive Officer, Director

Exhibit 31.2

CERTIFICATION

I, Julie Andrews, certify that:

1. I have reviewed this quarterly report on Form 10-Q for the quarterly period ended September 30, 2026, of Orthofix Medical Inc.;

2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;

3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;

4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:

   a. designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;

   b. designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;

   c. evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and

   d. disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's fourth fiscal quarter in the case of an annual report) that has material affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and

5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):

   a. all significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and

   b. any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Dated: November 9, 2026

By: /s/ JULIE ANDREWS
Name: Julie Andrews
Title: Chief Financial Officer

Exhibit 32.1

CERTIFICATION PURSUANT TO 18 U.S.C. SECTION 1350,
AS ADOPTED PURSUANT TO SECTION 906
OF THE SARBANES-OXLEY ACT OF 2002

In connection with the Quarterly Report of Orthofix Medical Inc. ("Orthofix") on Form 10-Q for the quarterly period ended September 30, 2026, (the "Report"), as filed with the Securities and Exchange Commission on the date hereof, Massimo Calafiore, President and Chief Executive Officer, Director, and Julie Andrews, Chief Financial Officer, each certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that, to his knowledge:

1. The Report fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and

2. The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of Orthofix.

Dated: November 9, 2026

/s/ MASSIMO CALAFIORE
Name: Massimo Calafiore
Title: President and Chief Executive Officer, Director

Dated: November 9, 2026

/s/ JULIE ANDREWS
Name: Julie Andrews
Title: Chief Financial Officer