IRADIMED CORPORATION Reports Second Quarter 2026 Financial Results

Announces Regular Quarterly Cash Dividend of $0.20 Per Share

Reaffirms Full-Year 2026 Revenue and Earnings Guidance

- Reports revenue of $20.5 million for the second quarter of 2026, which increased by $0.1 million, or 0.5% compared to the same period in 2025.
- Reports GAAP diluted EPS of $0.41 and non-GAAP diluted EPS of $0.46 for the second quarter of 2026.
- Declares a regular quarterly cash dividend of $0.20 per share of common stock for the second quarter of 2026, payable on August 28, 2026.

ORLANDO, Fla., July 31, 2026 (GLOBE NEWSWIRE) — IRADIMED CORPORATION (the "Company" or "Iradimed") (NASDAQ: IRMD) announced today its financial results for the three and six months ended June 30, 2026. The Company is a leader in developing innovative magnetic resonance imaging ("MRI") compatible medical devices and products. The Company is a provider of (i) non-magnetic intravenous ("IV") infusion pump systems and (ii) a non-magnetic patient vital signs monitoring system that are each designed for use during MRI procedures.

"The second quarter marked the first step in the actual launch of the new 3870 MR IV pump and transitioning of IV pump revenue from the older 3860 to this next-generation 3870 platform. Second-quarter revenue was in line with our expectations, and the initial 3870 shipments met our plan. This was our first sizable production ramp of the new pump, with expected startup inertia and challenges, which were well managed by our manufacturing and engineering teams. We anticipate that as experience and volumes build through the balance of the year, we will continue to gain efficiencies, which we expect will be reflected in a higher gross margin in the second half of the year.

"We have been very pleased with our bookings for the quarter as domestic monitor unit bookings set a record, and the new 3870 bookings aligned with our expectations as we continue to get the 3870 in front of more customers. Of note, more than 70% of new pump orders were for our 'quad stack' configuration of four pumps on a single IV pole, an indication that customers are standardizing on this configuration for the 3870, and doubling the number of pump channels from the previous norm. We continue to expect the 3870 revenues to ramp significantly through the second half of the year," said Roger Susi, President and Chief Executive Officer of Iradimed.

"For the third quarter of 2026, we expect revenue of $23.0 million to $24.5 million and GAAP

diluted earnings per share of $0.49 to $0.54 and non-GAAP diluted earnings per share of $0.54 to $0.59. We reaffirm our guidance for the full-year 2026, expecting to report revenue of $91.0 million to $96.0 million, and GAAP diluted earnings per share of $1.90 to $2.05 and non-GAAP diluted earnings per share of $2.09 to $2.24," added Mr. Susi.

The Company's board of directors declared a regular quarterly cash dividend of $0.20 per share of our outstanding common stock. The dividend is payable to stockholders of record as of the close of business on August 14, 2026 and will be paid on August 28, 2026.

Three Months Ended June 30, 2026

For the three months ended June 30, 2026, the Company reported revenue of $20.5 million, compared to $20.4 million for the comparable period of 2025, which increased by $0.1 million or 0.5%. The Company reported net income of $5.2 million, compared to $5.8 million for the comparable period of 2025, which decreased by 10%. Earnings per share decreased by $0.04 per diluted share or 9% to $0.41 per diluted share, for the three months ended June 30, 2026, compared to $0.45 per diluted share for the same period of 2025.

For the three months ended June 30, 2026, the Company reported non-GAAP net income of $5.9 million, which excludes $0.6 million of stock compensation expense, net of tax benefit, compared to $6.4 million, which excludes $0.6 million of stock compensation expense, net of tax benefit, which decreased by 8% over the comparable period of 2025. On a non-GAAP basis, net income per diluted share was $0.46 for the three months ended June 30, 2026, compared to $0.49 per diluted share for the comparable period of 2025, which decreased by 7%.

Six Months Ended June 30, 2026

For the six months ended June 30, 2026, the Company reported revenue of $42.4 million, compared to $39.9 million for the comparable period of 2025, which increased by $2.5 million or 6%. The Company reported net income of $11.0 million, compared to $10.5 million for the comparable period of 2025, which increased by 5%. Earnings per share increased by $0.04 per diluted share or 5% to $0.86 per diluted share, for the six months ended June 30, 2026, compared to $0.82 per diluted share for the same period of 2025.

For the six months ended June 30, 2026, the Company reported non-GAAP net income of $12.2 million, which excludes $1.2 million of stock compensation expense, net of tax benefit, compared to $11.7 million, which excludes $1.2 million of stock compensation expense, net of tax benefit, which increased by 5% over the comparable period of 2025. On a non-GAAP basis, net income per diluted share was $0.95 for the six months ended June 30, 2026, compared to $0.91 per diluted share for the comparable period of 2025, which increased by 4% .

Revenue Information:

(in thousands)	Three Months Ended June 30, 2026 (unaudited)	2025	Six Months Ended June 30, 2026 (unaudited)	2025
Devices:				
MRI Compatible Intravenous (“IV”) Infusion Pump Systems	$ 6,428	$ 8,188	$ 14,092	$ 14,187
MRI Compatible Patient Vital Signs Monitoring Systems	6,665	5,944	13,772	12,489
Ferro Magnetic Detection Systems	756	482	1,377	900
Total devices revenue	13,849	14,614	29,241	27,576
Amortization of extended maintenance agreements	758	592	1,417	1,153
Disposables	4,793	4,204	9,678	9,151
Services and other	1,066	999	2,108	2,040
Total revenue	$ 20,466	$ 20,409	$ 42,444	$ 39,920

For the three months ended June 30, 2026 and 2025, domestic sales were 82% and 89% of total revenue, respectively. For the three months ended June 30, 2026, the gross profit margin was 74% compared to 78% for the comparable period of 2025.

For the six months ended June 30, 2026 and 2025, domestic sales were 82% and 86% of total revenue, respectively. For the six months ended June 30, 2026, the gross profit margin was 75% compared to 77% for the comparable period of 2025.

Cash Flow from Operations

For the three months ended June 30, 2026, cash flow from operations was $5.9 million, compared to $7.7 million for the comparable period of 2025, a decrease of 23%. During the three months ended June 30, 2026, cash provided by operations was positively impacted by higher net income, pre-paid expenses, and inventory purchases, and negatively impacted by accounts receivable. See the compilation of non-GAAP free cash flow in the table later in this release.

For the six months ended June 30, 2026, cash flow from operations was $14.2 million, compared to $12.0 million for the comparable period of 2025, an increase of 18%. During the six months ended June 30, 2026, cash provided by operations was positively impacted by higher net income, and lower cash outflows related to accounts payable, pre-paid expenses, and inventory purchases, and negatively impacted by accounts receivable. See the compilation of non-GAAP free cash flow in the table later in this release.

Financial Guidance

For the third quarter of 2026, the Company expects revenue of $23.0 million to $24.5 million, GAAP diluted earnings per share of $0.49 to $0.54, and non-GAAP diluted earnings per share of $0.54 to $0.59. For the full year 2026, the Company expects to report revenue of $91.0 million to $96.0 million, GAAP diluted earnings per share of $1.90 to $2.05, and non-GAAP diluted earnings per share of $2.09 to $2.24.

The Company's non-GAAP diluted earnings per share guidance excludes stock-based compensation expense, net of tax expense. The Company expects stock-based compensation expense, net of tax, to be approximately $2.5 million and $0.6 million for the full year and third quarter of 2026, respectively.

Use of Non-GAAP Financial Measures

The Company believes using non-GAAP net income, non-GAAP net income per share-diluted, and free cash flow is helpful to our investors. These measures, which we refer to as our non-GAAP financial measures, are not prepared in accordance with accounting principles generally accepted in the United States of America ("GAAP"). These non-GAAP measures are intended to provide the reader with additional supplemental perspectives on operating results, performance trends, and financial condition. Non-GAAP financial measures are not a substitute for GAAP measures; they should be read and used in conjunction with the Company's GAAP financial information. Because non-GAAP financial measures presented in this release are not measurements determined in accordance with GAAP and are susceptible to varying calculations, these non-GAAP financial measures, as presented, may not be comparable to other similarly titled measures presented by other companies.

We calculate non-GAAP net income as net income excluding:

(1) Stock-based compensation expense, net of tax. Because of varying available valuation methodologies, subjective assumptions, and the variety of equity instruments that can impact a company's non-cash expenses, we believe that providing non-GAAP financial measures that exclude stock-based compensation expense allows for meaningful comparisons between our operating results from period to period;

(2) Operating expenses, net of tax, that we believe are not indicative of the Company's ongoing core operating performance; and

(3) Infrequent income tax items are considered based on their nature and are excluded from the provision for income taxes as these costs or benefits are not indicative of our normal or future provision for income taxes.

We calculate free cash flow as net cash provided by operating activities, less net cash used in investing activities for the development of internal software and purchases of property and equipment.

We consider free cash flow to be a liquidity measure that provides useful information to management and investors about the amount of cash generated by our business that can be used for strategic opportunities, including investing in our business, making strategic acquisitions, strengthening our balance sheet and returning cash to our stockholders through various means.

Our non-GAAP financial measures are important tools for financial and operational decision-making and for evaluating our ongoing core operating results.

A reconciliation of the non-GAAP financial measures used in this release to the most comparable GAAP measures for the respective periods can be found in the table later in this release immediately following the condensed statements of operations. These non-GAAP financial measures should not be considered in isolation or as a substitute for a measure of the Company's operating performance or liquidity prepared in accordance with GAAP and are not indicative of net income or cash provided by operating activities.

Conference Call

The Company has scheduled a conference call to discuss this release beginning at 11:00 a.m. Eastern Time, July 31, 2026. Individuals interested in listening to the conference call may do so by registering here, https://register-conf.media-server.com/register/BIc3b3caa4c4d94098970951ede5eb5d2e.

Once registered, a dial-in number, a unique PIN, and instructions will be provided to participants.

The conference call will also be available in real-time via the Internet at http://www.iradimed.com/en-us/investors/events/. A recording of the call will be available on the Company's website following the call's completion.

About IRADIMED CORPORATION

IRADIMED CORPORATION is a leader in developing innovative Magnetic Resonance Imaging ("MRI") compatible medical devices. We design, manufacture, market, and distribute MRI-compatible medical devices, accessories, disposables, and related services.

We are the only known provider of a non-magnetic intravenous ("IV") infusion pump system specifically designed to be safe for use during MRI procedures. We were the first to develop an infusion delivery system that largely eliminates many of the dangers and problems present during MRI procedures. Standard infusion pumps contain magnetic and electronic components that can create radio frequency interference and are dangerous to operate in the presence of the powerful magnet that drives an MRI system. Our patented MRidium® MRI compatible IV infusion pump systems (3860 and 3870) have a non-magnetic ultrasonic

motor, uniquely designed non-ferrous parts, and other special features to safely and predictably deliver anesthesia and other IV fluids during various MRI procedures. Our pump solutions provide a seamless approach that enables accurate, safe, and dependable fluid delivery before, during, and after an MRI scan, which is essential to critically ill patients who cannot be removed from their vital medications and children and infants who must generally be sedated to remain immobile during an MRI scan.

Our 3880 MRI compatible patient vital signs monitoring system has been designed with non-magnetic components and other special features to safely and accurately monitor a patient's vital signs during various MRI procedures. The Iradimed 3880 system operates dependably in magnetic fields up to 30,000 gauss, which means it can operate virtually anywhere in the MRI scanner room. The Iradimed 3880 has a compact, lightweight design, allowing it to travel with the patient from the critical care unit to the MRI and back, resulting in increased patient safety through uninterrupted vital signs monitoring and decreasing the amount of time critically ill patients are away from critical care units. The features of the Iradimed 3880 include wireless ECG with dynamic gradient filtering; wireless SpO2 using Masimo® algorithms; non-magnetic respiratory CO2; invasive and non-invasive blood pressure; patient temperature, and optional advanced multi-gas anesthetic agent unit featuring continuous Minimum Alveolar Concentration measurements. The Iradimed 3880 MRI compatible patient vital signs monitoring system has an easy-to-use design and allows for the effective communication of patient vital signs information to clinicians.

For more information, please visit www.iradimed.com.

Forward-Looking Statements

This release and any oral statements made regarding the subject of this release contain forward-looking statements as defined under Section 27A of the Securities Act of 1933, as amended, and Section 21E of the Securities Exchange Act of 1934, as amended. All statements, other than statements of historical facts, that address activities that the Company assumes, plans, expects, believes, intends, projects, indicates, estimates or anticipates (and other similar expressions) will, should or may occur in the future are forward-looking statements, including statements relating to financial guidance, future quarterly cash dividends, operational issues meeting demand, commercialization, timing and expected revenue ramping of our next-generation 3870 MRI compatible IV infusion pump, and the Company's strategic plans, objectives, and intentions. The forward-looking statements are based on management's current belief, based on currently available information, as to the outcome and timing of future events. The forward-looking statements involve risks and uncertainties, including, among others, that our business plans may change as circumstances warrant.

Readers are cautioned not to place undue reliance on forward-looking statements, which

speak only as of the date that they are made, which reflect management's current estimates, projections, expectations, or beliefs, and which involve
risks and uncertainties that could cause actual results and outcomes to be materially different. Risks and uncertainties that may affect the future results of the Company include, but are not limited to: our ability to receive 510(k) clearance for our products and product candidates, complete inspections conducted by the U.S. Food & Drug Administration (the "FDA") or other regulatory bodies resulting in favorable outcomes, additional actions by or requests from the FDA, including a request to cease domestic distribution of products, or other regulatory bodies and unanticipated costs or delays associated with the resolution of these matters; the timing and likelihood of regulatory approvals or clearances from the FDA or other regulatory bodies and regulatory actions on our product candidates and product marketing activities; unexpected costs, expenses and diversion of management attention resulting from actions or requests posed to us by the FDA or other regulatory bodies; failure to obtain and/or maintain regulatory approvals or clearances and comply with applicable regulations; our primary reliance on a limited number of products; our ability to retain the continued service of our key professionals, including key management, marketing and scientific personnel, and to identify, hire and retain such additional qualified professionals; our expectations regarding the sales and marketing of our products, product candidates and services; our expectations regarding the integrity of our supply chain for our products; the potential for adverse application of environmental, health and safety and other laws and regulations of any jurisdiction on our operations; our expectations for market acceptance of our new products; the potential for our marketed products to be withdrawn due to recalls, patient adverse events or deaths; our ability to successfully prepare, file, prosecute, maintain, defend, including in cases of infringement, and enforce patent claims and other intellectual property rights on our products; our ability to identify and pursue development of additional products; the implementation of our business strategies; the potential for exposure to product liability claims; our financial performance expectations and interpretations thereof by securities analysts and investors; our ability to compete in the development and marketing of our products and product candidates with existing companies and new market entrants in our industry; difficulties or delays in the development, production, manufacturing and marketing of new or existing products and services, including difficulties or delays associated with obtaining requisite regulatory approvals or clearances associated with those activities; changes in laws and regulations or in the interpretation or application of laws or regulations, as well as possible failures to comply with applicable laws or regulations as a result of possible misinterpretations or misapplications; cost-containment efforts of our customers, purchasing groups, third-party payers and governmental organizations; costs associated with protecting our trade secrets and enforcing our patent, copyright and trademark rights, and successful challenges to the validity of our patents, copyrights or trademarks; actions of regulatory bodies and other government authorities, including the FDA and foreign counterparts, that could delay, limit or suspend product development, manufacturing or sales or result in recalls, seizures, consent decrees, injunctions and

monetary sanctions; costs or claims resulting from potential errors or defects in our manufacturing that may injure persons or damage property or operations, including costs from remediation efforts or recalls; the results, consequences, effects or timing of any commercial disputes, patent infringement claims or other legal proceedings or any government investigations; changes in our production capacity, including interruptions in our ability to manufacture our products or an inability to obtain key components or raw materials or increased costs in such key components or raw materials; the failure of third parties to uphold their contractual duties or meet expected deadlines; uncertainties in our industry due to the effects of government-driven or mandated healthcare reform; competitive pressures in the markets in which we operate; potential negative impacts resulting from a future pandemic or epidemic, or natural disaster; the impact on our operations and financial results of any public health emergency and any related policies and actions by governments or other third parties; breaches or failures of our or our vendors' or customers' information technology systems or products, including by cyber-attack, data leakage, unauthorized access or theft; the loss of, or default by, one or more key customers or suppliers; unfavorable changes to the terms of key customer or supplier relationships; weakening of economic conditions, or the anticipation thereof, that could adversely affect the level of demand for our products; the conditions in the U.S. and global economies, including the impact of increasing and/or fluctuating tax and interest rates as well as inflationary pressures on such economies; geopolitical risks, including tariffs, trade disputes, international military conflicts and recent or upcoming elections in the United States and other countries, and the impact of insufficient governmental funding and related operational inefficiencies, which could, among other things, lead to increased volatility in the financial market and in the market in which we operate our business; and other risks detailed in our filings with the United States Securities and Exchange Commission (the "SEC").

Such forward-looking statements are subject to known and unknown risks, uncertainties, assumptions and other important factors, many of which are outside of the Company's control that could cause actual results to differ materially from the results discussed in the forward-looking statements. These risks, uncertainties, assumptions and other important factors include, but are not limited to, those included in Part II, Item 1A, "Risk Factors" of the Company's Quarterly Reports on Form 10-Q, and Part I, Item 1A, "Risk Factors" of the Company's Annual Report on Form 10-K for the year ended December 31, 2025, as well as those otherwise described or updated from time to time in our other filings with the SEC. You are cautioned not to place undue reliance upon any forward-looking statements, which speak only as of the date made, and the Company undertakes no commitment to update or revise the forward-looking statements, whether as a result of new information, future events or otherwise.

IRADIMED CORPORATION
CONDENSED BALANCE SHEETS
(in thousands)

	June 30, 2026 (unaudited)	December 31, 2025 (audited)
ASSETS		
Current assets:		
Cash and cash equivalents	$ 59,080	$ 51,159
Other current assets	27,642	30,044
Total current assets	86,722	81,203
Property and equipment, net	24,208	23,856
Other assets	4,045	3,717
Total assets	$ 114,975	$ 108,776
LIABILITIES AND STOCKHOLDERS' EQUITY		
Current liabilities:		
Accounts payable	$ 1,849	$ 1,820
Deferred revenue	3,469	2,871
Other current liabilities	3,846	5,489
Total current liabilities	9,164	10,180
Deferred revenue, non-current	4,063	3,978
Total liabilities	13,227	14,158
Stockholders' equity:		
Total stockholders' equity	101,748	94,618
Total liabilities and stockholders' equity	$ 114,975	$ 108,776

IRADIMED CORPORATION
CONDENSED STATEMENTS OF OPERATIONS
(Unaudited)
(in thousands, except per share data)

	Three Months Ended June 30,		Six Months Ended June 30,	
	2026	2025	2026	2025
Revenue	$ 20,466	$ 20,409	$ 42,444	$ 39,920
Cost of revenue	5,306	4,454	10,471	9,122
Gross profit	15,160	15,955	31,973	30,798
Operating expenses:				
General and administrative	3,866	4,280	8,435	8,891
Sales and marketing	4,246	4,010	8,298	8,186
Research and development	657	877	1,613	1,502
Total operating expenses	8,769	9,167	18,346	18,579
Income from operations	6,391	6,788	13,627	12,219
Other income, net	497	539	1,030	1,053
Income before provision for income taxes	6,888	7,327	14,657	13,272
Provision for income tax expense	1,670	1,553	3,621	2,811
Net income	$ 5,218	$ 5,774	$ 11,036	$ 10,461

Net income per share:				
Basic	$ 0.41	$ 0.45	$ 0.86	$ 0.82
Diluted	$ 0.41	$ 0.45	$ 0.86	$ 0.82
Weighted average shares outstanding:				
Basic	12,784	12,716	12,783	12,715
Diluted	12,872	12,835	12,871	12,830

IRADIMED CORPORATION
RECONCILIATION OF NON-GAAP FINANCIAL MEASURES
(Unaudited)
(in thousands, except per share data)
Non-GAAP Net Income and Diluted EPS

	Three Months Ended June 30,		Six Months Ended June 30,	
	2026	2025	2026	2025
Net income	$ 5,218	$ 5,774	$ 11,036	$ 10,461
Excluding:				
Stock-based compensation expense, net of tax benefit	640	577	1,183	1,230
Non-GAAP net income	$ 5,858	$ 6,351	$ 12,219	$ 11,691
Weighted-average shares outstanding – diluted	12,872	12,835	12,871	12,830
Non-GAAP net income per share – diluted	$ 0.46	$ 0.49	$ 0.95	$ 0.91

Non-GAAP Free Cash Flow

	Three Months Ended		Six Months Ended	
	2026	2025	2026	2025
Net cash provided by operating activities	$ 5,940	$ 7,746	$ 14,230	$ 12,038
Less:				
Capital Expenditures	443	2,824 [1]	912	6,741 [1]
Free cash flow	$ 5,497	$ 4,922	$ 13,318	$ 5,297

[1]For the three and six months ended June 30, 2025, capital expenditures include approximately $2.1 million and $5.5 million respectively related to construction of new corporate office and manufacturing facility completed in 2025.

Media Contact:
IRADIMED CORPORATION
(407) 677-8022
InvestorRelations@iradimed.com

IRADIMED CORPORATION