



For Immediate Release

United Therapeutics Corporation Reports Second Quarter 2024 Financial Results

SILVER SPRING, Md. and RESEARCH TRIANGLE PARK, N.C., July 31, 2024: United Therapeutics Corporation (Nasdaq: **UTHR**), a public benefit corporation, today announced its financial results for the quarter ended June 30, 2024. Total revenues in the second quarter of 2024 grew 20 percent year-over-year to \$714.9 million, compared to \$596.5 million in the second quarter of 2023.

“This quarter we drove record revenue from our foundational commercial business. Next year we expect data from our innovative clinical pipeline. All this while we march forward with our revolutionary organ manufacturing programs,” said **Martine Rothblatt, Ph.D.**, Chairperson and Chief Executive Officer of United Therapeutics. “We believe there is no other biotech with our combination of relentless focus, near-term commercial growth, and clinical potential.”

Michael Benkowitz, President and Chief Operating Officer of United Therapeutics, added, “Our fifth straight quarter of record revenue was driven by continued traction for Tyvaso in pulmonary hypertension associated with interstitial lung disease, along with strong fundamentals for our other products in pulmonary arterial hypertension and neuroblastoma.”

Second Quarter 2024 Financial Results

Key financial highlights include (dollars in millions, except per share data):

	Three Months Ended		Dollar Change	Percentage Change
	June 30,			
	2024	2023		
Total revenues	\$ 714.9	\$ 596.5	\$ 118.4	20 %
Net income	\$ 278.1	\$ 259.2	\$ 18.9	7 %
Net income, per basic share	\$ 6.26	\$ 5.53	\$ 0.73	13 %
Net income, per diluted share	\$ 5.85	\$ 5.24	\$ 0.61	12 %

Revenues

The table below presents the components of total revenues (dollars in millions):

	Three Months Ended June 30,		Dollar Change	Percentage Change
	2024	2023		
Net product sales:				
Tyvaso DPI ^{®(1)}	\$ 258.3	\$ 193.6	\$ 64.7	33 %
Nebulized Tyvaso ^{®(1)}	139.9	125.3	14.6	12 %
Total Tyvaso	398.2	318.9	79.3	25 %
Remodulin ^{®(2)}	147.3	127.2	20.1	16 %
Orenitram [®]	107.1	95.1	12.0	13 %
Unituxin [®]	51.7	44.3	7.4	17 %
Adcirca [®]	5.7	7.5	(1.8)	(24)%
Other	4.9	3.5	1.4	40 %
Total revenues	\$ 714.9	\$ 596.5	\$ 118.4	20 %

(1) Net product sales include both the drug product and the respective inhalation device.

(2) Net product sales include sales of infusion devices, including the Remunity[®] Pump.

Total Tyvaso revenues grew by 25 percent to \$398.2 million in the second quarter of 2024, compared to \$318.9 million in the second quarter of 2023. This growth was primarily due to an increase in quantities sold, driven by the commercial launch of Tyvaso DPI in June 2022 and continued growth in commercial utilization by patients with pulmonary hypertension associated with interstitial lung disease and, to a lesser extent, price increases.

The growth in Tyvaso DPI revenues resulted primarily from an increase in quantities sold and, to a lesser extent, price increases. The increase in Tyvaso DPI quantities sold was due to continued growth in the number of patients following the product's launch and, to a lesser extent, increased commercial utilization following the implementation of the Part D redesign under the Inflation Reduction Act (**IRA**).

The growth in Remodulin revenues resulted primarily from an increase in U.S. Remodulin revenues, driven by an increase in quantities sold and, to a lesser extent, lower Medicaid rebates.

The growth in Orenitram revenues resulted primarily from an increase in quantities sold and, to a lesser extent, a price increase. The increase in quantities sold was driven, at least in part, by increased commercial utilization following the implementation of the Part D redesign under the IRA.

The growth in Unituxin revenues resulted from a price increase and an increase in quantities sold.

The table below presents the breakdown of total revenues between the United States and rest-of-world (**ROW**) (in millions):

	Three Months Ended June 30,					
	2024			2023		
	U.S.	ROW	Total	U.S.	ROW	Total
Net product sales:						
Tyvaso DPI ⁽¹⁾	\$ 258.3	\$ –	\$ 258.3	\$ 193.6	\$ –	\$ 193.6
Nebulized Tyvaso ⁽¹⁾	130.2	9.7	139.9	119.6	5.7	125.3
Total Tyvaso	388.5	9.7	398.2	313.2	5.7	318.9
Remodulin ⁽²⁾	122.5	24.8	147.3	103.5	23.7	127.2
Orenitram	107.1	–	107.1	95.1	–	95.1
Unituxin	46.8	4.9	51.7	39.5	4.8	44.3
Adcirca	5.7	–	5.7	7.5	–	7.5
Other	4.6	0.3	4.9	3.2	0.3	3.5
Total revenues	\$ 675.2	\$ 39.7	\$ 714.9	\$ 562.0	\$ 34.5	\$ 596.5

(1) Net product sales include both the drug product and the respective inhalation device.

(2) Net product sales include sales of infusion devices, including the Remunity Pump.

Expenses

Cost of sales. The table below summarizes cost of sales by major category (dollars in millions):

Category:	Three Months Ended		Dollar Change	Percentage Change
	June 30,			
	2024	2023		
Cost of sales	\$ 75.9	\$ 63.2	\$ 12.7	20 %
Share-based compensation expense ⁽¹⁾	1.9	0.9	1.0	111 %
Total cost of sales	\$ 77.8	\$ 64.1	\$ 13.7	21 %

(1) See *Share-based compensation* below.

Cost of sales, excluding share-based compensation. Cost of sales for the three months ended June 30, 2024 increased as compared to the same period in 2023, primarily due to an increase in Tyvaso DPI royalty expense.

Research and development. The table below summarizes the nature of research and development expense by major expense category (dollars in millions):

Category:	Three Months Ended June 30,		Dollar Change	Percentage Change
	2024	2023		
External research and development ⁽¹⁾	\$ 49.4	\$ 49.3	\$ 0.1	– %
Internal research and development ⁽²⁾	44.5	34.7	9.8	28 %
Share-based compensation expense ⁽³⁾	8.6	5.0	3.6	72 %
Impairments ⁽⁴⁾	–	–	–	– %
Other ⁽⁵⁾	37.1	–	37.1	NM ⁽⁶⁾
Total research and development expense	\$ 139.6	\$ 89.0	\$ 50.6	57 %

- (1) *External research and development* primarily includes fees paid to third parties (such as clinical trial sites, contract research organizations, and contract laboratories) for preclinical and clinical studies and payments to third-party contract manufacturers before FDA approval of the relevant product.
- (2) *Internal research and development* primarily includes salary-related expenses for research and development functions, internal costs to manufacture product candidates before FDA approval, and internal facilities-related expenses, including depreciation, related to research and development activities.
- (3) See *Share-based compensation* below.
- (4) *Impairments* primarily includes impairment charges to write down the carrying value of in-process research and development and of certain property, plant, and equipment as a result of research and development activities. There were no impairment charges during the three months ended June 30, 2024 and June 30, 2023.
- (5) *Other* primarily includes upfront fees and milestone payments to third parties under license agreements related to development-stage products and adjustments to the fair value of our contingent consideration obligations.
- (6) Calculation is not meaningful.

Research and development expense, excluding share-based compensation. Research and development expense for the three months ended June 30, 2024 increased as compared to the same period in 2023, primarily due to increased expenditures related to upfront non-refundable licensing payments for drug delivery devices and increased expenditures related to organ manufacturing projects.

Selling, general, and administrative. The table below summarizes selling, general, and administrative expense by major category (dollars in millions):

	Three Months Ended June 30,		Dollar Change	Percentage Change
	2024	2023		
Category:				
General and administrative	\$ 113.0	\$ 102.0	\$ 11.0	11 %
Sales and marketing	25.4	20.1	5.3	26 %
Share-based compensation expense ⁽¹⁾	39.2	7.9	31.3	396 %
Total selling, general, and administrative expense	\$ 177.6	\$ 130.0	\$ 47.6	37 %

(1) See *Share-based compensation* below.

General and administrative, excluding share-based compensation. General and administrative expense for the three months ended June 30, 2024 increased as compared to the same period in 2023, primarily due to an increase in legal expenses related to litigation matters and an increase in personnel expense due to growth in headcount.

Share-based compensation. The table below summarizes share-based compensation expense by major category (dollars in millions):

	Three Months Ended June 30,		Dollar Change	Percentage Change
	2024	2023		
Category:				
Stock options	\$ 8.1	\$ 1.6	\$ 6.5	406 %
Restricted stock units	19.2	13.6	5.6	41 %
Share tracking awards plan (STAP)	21.9	(1.9)	23.8	NM ⁽¹⁾
Employee stock purchase plan	0.5	0.5	–	– %
Total share-based compensation expense	\$ 49.7	\$ 13.8	\$ 35.9	260 %

(1) Calculation is not meaningful.

The increase in share-based compensation expense for the three months ended June 30, 2024, as compared to the same period in 2023, was primarily due to an increase in STAP expense driven by a 39 percent increase in our stock price for the three months ended June 30, 2024, as compared to a one percent decrease in our stock price for the same period in 2023.

Income tax expense. *Income tax expense* for the three months ended June 30, 2024 and 2023 was \$77.2 million and \$76.0 million, respectively. Our effective income tax rate (**ETR**) for the three months ended June 30, 2024 and 2023 was 22 percent and 23 percent, respectively. Our ETR for the three months ended June 30, 2024 decreased compared to our ETR for the three months ended June 30, 2023 primarily due to a lower amount of uncertain tax positions recorded.

Share repurchase. In March 2024, we entered into an accelerated share repurchase agreement (the **ASR agreement**) with Citibank, N.A. (**Citi**). Under the ASR agreement, we made an aggregate upfront payment of \$1.0 billion to Citi and received an aggregate initial delivery of 3,275,199 shares of our common stock on March 27, 2024, representing approximately 80 percent of the total shares that

would be repurchased under the ASR agreement measured based on the closing price of our common stock on March 25, 2024.

The share purchase under the ASR agreement was divided into two tranches, resulting in upfront payments of \$300 million and \$700 million, respectively. The final settlement of the \$300 million tranche occurred in June 2024, and we received an additional 181,772 shares of our common stock upon settlement. At the final settlement of the \$700 million second tranche, which we expect to occur in the third quarter of 2024, we may be entitled to receive additional shares of common stock, or, under certain limited circumstances, be required to make a cash payment to Citi or, if we so elect, deliver shares of common stock to Citi.

The final number of shares that we will ultimately repurchase pursuant to the ASR agreement will be based on the average of the daily volume-weighted average price per share of our common stock during the repurchase period, less a discount and subject to adjustments pursuant to the terms and conditions of the ASR agreement.

Webcast

We will host a webcast to discuss our second quarter 2024 financial results on Wednesday, July 31, 2024, at 9:00 a.m. Eastern Time. The webcast can be accessed live via our website at <https://ir.unither.com/events-and-presentations>. A replay of the webcast will also be available at the same location on our website.

United Therapeutics: Enabling Inspiration

At United Therapeutics, our vision and mission are one. We use our enthusiasm, creativity, and persistence to innovate for the unmet medical needs of our patients and to benefit our other stakeholders. We are bold and unconventional. We have fun; we do good. We are the first publicly-traded biotech or pharmaceutical company to take the form of a public benefit corporation (**PBC**). Our public benefit purpose is to *provide a brighter future for patients through (a) the development of novel pharmaceutical therapies; and (b) technologies that expand the availability of transplantable organs.*

You can learn more about what it means to be a PBC here: unither.com/pbc.

Forward-Looking Statements

Statements included in this press release that are not historical in nature are “forward-looking statements” within the meaning of the Private Securities Litigation Reform Act of 1995. Forward-looking statements include, among others, statements related to our anticipated near-term commercial growth; the anticipated progress of our organ manufacturing programs; our clinical potential, including anticipated clinical trial data next year; our unique position in the biotech industry; and our goals of innovating for the unmet medical needs of our patients and to benefit our other stakeholders, furthering our public benefit purpose of developing novel pharmaceutical therapies and technologies that expand the availability of transplantable organs. These forward-looking statements are subject to certain risks and uncertainties, such as those described in our periodic reports filed with the Securities and Exchange Commission, that could cause actual results to differ materially from anticipated results. Consequently, such forward-looking statements are qualified by the cautionary statements, cautionary language and risk factors set forth in our periodic reports and documents filed with the Securities and Exchange Commission, including our most recent Annual Report on Form 10-K, Quarterly Reports on Form 10-Q, and Current Reports on Form 8-K. We claim the protection of the safe harbor contained in the Private Securities Litigation Reform Act of 1995 for forward-looking statements. We are providing this information as of July 31, 2024, and assume no obligation to update or revise the information contained in this press release whether as a result of new information, future events, or any other reason.

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UNITED THERAPEUTICS CORPORATION
CONSOLIDATED STATEMENTS OF OPERATIONS
(In millions, except per share data)

	Three Months Ended June 30,	
	2024	2023
	(Unaudited)	
Total revenues	\$ 714.9	\$ 596.5
Operating expenses:		
Cost of sales	77.8	64.1
Research and development	139.6	89.0
Selling, general, and administrative	177.6	130.0
Total operating expenses	395.0	283.1
Operating income	319.9	313.4
Interest income	46.2	37.2
Interest expense	(11.6)	(14.8)
Other income (expense), net	0.8	(0.6)
Total other income, net	35.4	21.8
Income before income taxes	355.3	335.2
Income tax expense	(77.2)	(76.0)
Net income	\$ 278.1	\$ 259.2
Net income per common share:		
Basic	\$ 6.26	\$ 5.53
Diluted	\$ 5.85	\$ 5.24
Weighted average number of common shares outstanding:		
Basic	44.4	46.9
Diluted	47.5	49.5

SELECTED CONSOLIDATED BALANCE SHEET DATA
(Unaudited, in millions)

	June 30, 2024
Cash, cash equivalents, and marketable investments	\$ 4,301.9
Total assets	6,723.2
Total liabilities	1,026.0
Total stockholders' equity	5,697.2