

OPKO Health Reports Second Quarter 2024 Business Highlights and Financial Results

Conference call begins at 4:30 p.m. Eastern time today

MIAMI, Aug. 07, 2024 (GLOBE NEWSWIRE) -- **OPKO Health, Inc. (NASDAQ: OPK)** reports business highlights and financial results for the three and six months ended June 30, 2024.

Highlights from the second quarter and recent weeks include the following:

- Enrollment underway in the MDX2001 Phase 1 trial for the treatment of solid tumor cancers. MDX2001, a tetraspecific antibody, is designed to optimize T-cell function to stimulate tumor regression while minimizing the likelihood of antigen escape. This Phase 1 open-label trial is expected to enroll 45 cancer patients with a variety of solid tumors at six clinical trial sites, and to evaluate safety, tolerability, pharmacokinetics and early evidence of anti-tumor activity.
- The global commercial launch of NGENLA® is ongoing by OPKO's partner, Pfizer. NGENLA has been launched in all major markets. OPKO is entitled to gross profit sharing based on sales of both NGENLA and Pfizer's daily growth hormone product, Genotropin®. In addition, OPKO is entitled to an additional \$100 million in potential milestone payments associated with approvals for an adult indication for growth hormone deficiency and additional pediatric indications.
- Entered into a \$250 million note purchase agreement with HealthCare Royalty secured by profit share payments related to NGENLA. Under the terms of the agreement, OPKO retains a significant portion of the profit share payments from Pfizer received pursuant to its license agreement relating to NGENLA in the near term with upside over the long term, as well as the full \$100 million of remaining potential milestone payments.
- OPKO's Board of Directors authorized a \$100 million share repurchase program. Under the program, OPKO may repurchase shares of its common stock from time to time through open market purchases, block trades, privately negotiated transactions, accelerated share repurchase transactions and/or pursuant to Rule 10b5-1 plans, in compliance with applicable securities laws and other legal requirements. The Company had approximately 697 million shares outstanding as of June 30, 2024. This new authorization represented approximately 10.1% of shares outstanding at the stock price at the time of the announcement.
- Sale of select assets of BioReference Health expected to close toward the end of the third quarter. OPKO entered into an agreement in March 2024 to sell

BioReference Health's laboratory testing businesses focused on clinical diagnostics and women's health, excluding operations in New York and New Jersey, for \$237.5 million. BioReference Health will continue to offer oncology and urology diagnostic services nationwide, as well as maintain its full operations in New York and New Jersey. Continuing operations accounted for net sales of more than \$400 million in 2023. This transaction is expected to streamline BioReference Health's laboratory services business while retaining its core operations to better position the division for sustained growth and profitability.

Second Quarter Financial Results

- Pharmaceuticals: Revenue from products in the second quarter of 2024 was \$40.5 million compared with \$43.5 million in the second guarter of 2023, reflecting lower sales in OPKO's international operating companies primarily due to foreign currency exchange fluctuations. Revenue from sales of Rayaldee was \$7.2 million compared with \$7.7 million in the same period in 2023. Revenue from the transfer of intellectual property and other was \$12.3 million in the second guarter of 2024, which included \$5.0 million from the BARDA contract, compared with \$94.9 million in the 2023 period, which included milestone revenue of \$90.0 million triggered by the FDA approval of NGENLA. Gross profit share and royalty payments for NGENLA and Pfizer's Genotropin was \$6.3 million in the 2024 guarter compared with \$3.8 million in the same period for 2023. Total costs and expenses increased to \$77.6 million in the second guarter of 2024 from \$74.7 million in the prior-year period primarily due to higher research and development expenses related to increased activity within the ModeX development programs. Operating loss was \$24.8 million in the second guarter of 2024, which included \$17.9 million of depreciation and amortization expense, compared with operating income of \$63.6 million in the second guarter of 2023, after giving effect to the \$90.0 million milestone payment as described above and also included \$17.8 million of depreciation and amortization expense.
- Diagnostics: Revenue from services in the second quarter of 2024 was \$129.4 million compared with \$127.0 million in the prior-year period, with the increase primarily due to higher clinical test reimbursement partially offset by lower clinical test volume. Total costs and expenses were \$156.0 million in the second quarter of 2024 compared with \$171.3 million in the second quarter of 2023, reflecting continued progress with cost-reduction initiatives. Included in second quarter 2024 results were revenue from services of approximately \$25.5 million and total costs and expenses of approximately \$32.5 million related to assets being acquired by Labcorp. Operating loss was \$26.6 million in the second quarter of 2024 compared with \$44.3 million in the 2023 period and included \$6.2 million and \$8.6 million of depreciation and amortization expense, respectively.
- **Consolidated**: Consolidated total revenues for the second quarter of 2024 were \$182.2 million compared with \$265.4 million for the comparable period of 2023. Operating loss for the second quarter of 2024 was \$61.7 million compared with operating income of \$7.0 million for the 2023 quarter, with the 2023 quarter benefiting from the \$90.0 million milestone payment described above. The second quarter of 2024 included non-cash other income of \$60.5 million compared with non-cash other expense of \$19.9 million in the year-ago quarter related to the change in the fair value

of the GeneDx Holdings investment. As a result, net loss for the second quarter of 2024 was \$10.3 million, or \$0.01 per share, compared with net loss of \$19.6 million, or \$0.03 per share, for the 2023 quarter.

• Cash and cash equivalents: Cash and cash equivalents were \$40.6 million as of June 30, 2024 and OPKO's Investments included liquid equity securities which had a market value of \$101.5 million, primarily from the ownership interest in GeneDx. Subsequent to the end of the second quarter, OPKO entered into a \$250 million note purchase agreement secured by OPKO's profit share payments to be received from Pfizer relating to NGENLA. OPKO is also entitled to receive \$237.5 million upon closing of the Labcorp transaction anticipated toward the end of the third quarter of 2024.

Conference Call and Webcast Information

A telephone replay will be available until August 14, 2024, by dialing 877-344-7529 (U.S.) or 412-317-0088 (International) and providing the passcode 6314261. A webcast replay will be available beginning approximately one hour after the completion of the live conference call here.

About OPKO Health

OPKO is a multinational biopharmaceutical and diagnostics company that seeks to establish industry-leading positions in large, rapidly growing markets by leveraging its discovery, development, and commercialization expertise and novel and proprietary technologies. For more information, visit www.opko.com.

Cautionary Statement Regarding Forward Looking Statements

This press release contains "forward-looking statements," as that term is defined under the Private Securities Litigation Reform Act of 1995 (PSLRA), which statements may be identified by words such as "expects," "plans," "projects," "will," "may," "anticipates," "believes," "should," "intends," "estimates," and other words of similar meaning, including statements regarding expected financial performance and expectations regarding the market for and sales of our products, whether our products will launch in all the territories in which they have been approved for sale, the timing of such launches, whether Pfizer will obtain approvals for an adult indication for growth hormone deficiency or additional pediatric indications and accordingly, whether we will be entitled to any additional milestone payments, whether our product development efforts will be successful and whether the expected benefits of our products will be realized, including whether enrollment in a Phase 1 clinical trial for MDX2001will be successful and whether the data will be positive, whether

and how many of our shares we will repurchase under a buyback program, whether NGENLA profits will be sufficient to provide long term upside after satisfying our obligations under the note purchase agreement, whether the relationship with our commercial and strategic partners will be successful, whether our commercial and strategic partners will be able to commercialize our products and successfully utilize our technologies, our ability to market and sell any of our products in development, whether we will continue to successfully advance products in our pipeline and whether they can be commercialized, whether the sale of selected BioReference assets will be completed in the third quarter or at all, and if this transaction is completed, whether BioReference will be able to streamline its laboratory services business and better position the division for sustained growth and profitability, whether BioReference's attempts at returning its core business to profitability will be successful, as well as other non-historical statements about our expectations, beliefs or intentions regarding our business, technologies and products, financial condition, strategies or prospects. Many factors could cause our actual activities or results to differ materially from the activities and results anticipated in forward-looking statements. These factors include those described in our Annual Reports on Form 10-K filed and to be filed with the Securities and Exchange Commission and under the heading "Risk Factors" in our other filings with the Securities and Exchange Commission, as well as the continuation and success of our relationship with our commercial partners, liquidity issues and the risks inherent in funding, developing and obtaining regulatory approvals of new, commerciallyviable and competitive products and treatments. In addition, forward-looking statements may also be adversely affected by general market factors, competitive product development, product availability, federal and state regulations and legislation, the regulatory process for new products and indications, manufacturing issues that may arise, patent positions and litigation, among other factors. The forward-looking statements contained in this press release speak only as of the date the statements were made, and we do not undertake any obligation to update forward-looking statements. We intend that all forward-looking statements be subject to the safe-harbor provisions of the PSLRA.

Contacts:

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—Tables to Follow—

OPKO Health, Inc. and Subsidiaries
Condensed Consolidated Balance Sheets
(in millions)
Unaudited

As of December
June 30, 31, 2024 2023

Assets:		
Cash and cash equivalents	\$ 40.6	\$ 95.9
Assets held for sale	119.7	0.0
Other current assets	 197.7	 213.6
Total current assets	358.0	309.5
In-process research and development and goodwill	725.1	793.3
Other assets	896.8	908.9
Total Assets	\$ 1,979.9	\$ 2,011.7
Liabilities and Equity:		
Accounts payable	\$ 82.2	\$ 69.7
Accrued expenses	94.5	90.1
Liabilities associated with assets held for sale	8.9	0.0
Current portion of convertible notes	0.2	0.0
Other current liabilities	33.7	40.3
Total current liabilities	 219.5	 200.1
Long-term portion of convertible notes	175.9	214.3
Deferred tax liabilities, net	119.1	126.8
Other long-term liabilities, principally leases, and lines of credit	70.1	81.3
Total Liabilities	584.6	 622.5
Equity	1,395.3	1,389.2
Total Liabilities and Equity	\$ 1,979.9	\$ 2,011.7

OPKO Health, Inc. and Subsidiaries Condensed Consolidated Statements of Operations (in millions, except share and per share data) Unaudited

	For the three months ended June 30,			For the six months ended June 30,				
	2024		2023		2024		2023	
Revenues								
Revenue from services	\$	129.4	\$	127.0	\$	256.3	\$	259.4
Revenue from products		40.5		43.5		78.5		83.9
Revenue from transfer of intellectual property and								
other		12.3		94.9		21.1		159.7
Total revenues		182.2		265.4		355.9		503.0
Costs and expenses								
Cost of service revenues		107.1		113.0		216.9		227.1
Cost of product revenues		23.5		25.9		45.2		50.2

Selling, general and administrative	68.8	79.8	139.0	155.4
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Research and	04.4	40.0	40.0	50.0
development	24.1	18.2	46.0	50.8
Contingent consideration	0.0	(0.0)	0.0	0.1
Amortization of intangible				
assets	20.4	21.5	41.9	43.0
Total costs and				
expenses	243.9	258.4	489.0	526.6
Operating loss (income)	(61.7)	7.0	(133.1)	(23.6)
Other income (expense),				
net	51.1	(23.5)	39.4	(9.8)
Loss before income taxes				
and investment losses	(10.6)	(16.5)	(93.7)	(33.4)
Income tax benefit				
(provision)	0.3	(3.1)	1.6	(4.4)
Loss before investment				· · · · · · · · · · · · · · · · · · ·
losses	(10.3)	(19.6)	(92.1)	(37.8)
Loss from investments in	, ,	, ,	, ,	, ,
investees	(0.0)	(0.0)	(0.0)	(0.1)
Net loss	\$ (10.3)	\$ (19.6)	\$ (92.1)	
Loss per share, basic and				
diluted	\$ (0.01)	\$ (0.03)	\$ (0.13)	\$ (0.05)
Weighted average common				
shares outstanding, basic				
and diluted	697,211,592	751,727,383	702,036,148	751,617,431



Source: OPKO Health, Inc.