UNITED STATES SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

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CURRENT REPORT PURSUANT TO SECTION 13 OR 15(D) OF THE SECURITIES EXCHANGE ACT OF 1934

Date of Report (Date of earliest event reported): November 22, 2021

Arrowhead Pharmaceuticals, Inc.

(Exact name of registrant as specified in its charter)

Delaware (State or other jurisdiction of incorporation)

001-38042 (Commission File Number)

46-0408024 (IRS Employer Identification No.)

177 E. Colorado Blvd, Suite 700, Pasadena, CA 91105 (Address of principal executive offices) (Zip Code) Registrant's telephone number, including area code (626) 304-3400

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Check the appropriate box below if the Form 8-K filing is intended to simultaneously satisfy the filing obligation of the registrant under any of the following provisions:						
	Soliciting material pursuant to Rule 14a-12 under the Exchange Act (17 CFR 240.14a-12) Pre-commencement communications pursuant to Rule 14d-2(b) under the Exchange Act (17 CFR 240.14d-2(b)					
Indicate by check mark whether the registrant is an emerging growth company as defined in Rule 405 of the Securities Act of 1933 (§230.405 of this chapter) or Rule 12b-2 of the Securities Exchange Act of 1934 (§240.12b-2 of this chapter).						
Emerging growth company \square						
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.						
	evised financial accounting standards provided pursuant to Section 13(a) of the Exchange Act. 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, Par Value \$0.001 per share	ARWR	The Nasdaq Global Select Market			

Item 2.02 Results of Operations and Financial Condition.

On November 22, 2021, Arrowhead Pharmaceuticals, Inc. announced and commented on its fiscal 2021 financial results for the period ended September 30, 2021. A copy of the press release is furnished herewith as Exhibit 99.1.

In accordance with General Instruction B.2 of Form 8-K, the information in items 2.02 of this Current Report on Form 8-K, including Exhibit 99.1, shall not be deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or otherwise subject to the liabilities of that section, nor shall it be deemed incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, whether made before or after the date hereof, except as expressly set forth by specific reference in such filing to this Current Report on Form 8-K.

Item 9.01. Financial Statements and Exhibits.

(d) Exhibits

Exhibit No.	Description
99.1	Press Release, dated November 22, 2021.
104	Cover Page Interactive File (the cover page tags are embedded within the Inline XBRL document).

SIGNATURES

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

Date: November 22, 2021

ARROWHEAD PHARMACEUTICALS, INC.

By: /s/ Kenneth Myszkowski

Kenneth Myszkowski Chief Financial Officer



PRESS RELEASE Nov. 22, 2021

Arrowhead Pharmaceuticals Reports Fiscal 2021 Year End Results

Conference Call and Webcast Today, November 22, 2021 at 4:30 p.m. ET

PASADENA, Calif., Nov. 22, 2021 — Arrowhead Pharmaceuticals, Inc. (NASDAQ: ARWR) today announced financial results for its fiscal year ended September 30, 2021. The company is hosting a conference call today, November 22, 2021, at 4:30 p.m. ET to discuss the results.

Conference Call and Webcast Details

Investors may access a live audio webcast on the Company's website at http://ir.arrowheadpharma.com/events.cfm. For analysts that wish to participate in the conference call, please dial 855-215-6159 or 315-625-6887 and provide Conference ID 8074256.

A replay of the webcast will be available on the company's website approximately two hours after the conclusion of the call and will remain available for 90 days. An audio replay will also be available approximately two hours after the conclusion of the call and will be available for 3 days. To access the audio replay, dial 855-859-2056 or 404-537-3406 and provide Conference ID 8074256.

Selected Recent Events

- Entered into an exclusive license agreement with GlaxoSmithKline (GSK) under which GSK will develop and commercialize ARO-HSD, Arrowhead's investigational RNA interference (RNAi) therapeutic in a Phase 1/2 trial that is currently being developed as a treatment for patients with nonalcoholic steatohepatitis (NASH)
- Presented new clinical data at The Liver Meeting, the Annual Meeting of the American Association for the Study of Liver Disease (AASLD), for the following investigational candidates:

- O JNJ-73763989 (JNJ-3989), formerly called ARO-HBV, being developed by collaborator Janssen Pharmaceuticals, Inc., one of the Janssen Pharmaceutical Companies of Johnson & Johnson (Janssen)
- O ARO-HSD, the investigational RNAi therapeutic being developed as a treatment for patients with NASH and recently licensed to GSK
- O ARO-AAT, also known as TAK-999, the investigational RNAi therapeutic being co-developed with Takeda Pharmaceutical Company Limited as a treatment for the rare genetic liver disease associated with alpha-1 antitrypsin deficiency
- Presented additional Phase 1/2 clinical data on ARO-APOC3, the investigational RNA RNAi therapeutic targeting apolipoprotein C-III (APOC3) being developed as a treatment for patients with hypertriglyceridemia, severe hypertriglyceridemia, and familial chylomicronemia syndrome, at the American Heart Association (AHA) Scientific Sessions 2021
- Initiated and began dosing patients in AROAPOC3-2002, now called MUIR, a Phase 2b clinical study of ARO-APOC3
- Initiated and began dosing patients in AROANG3-2001, now called ARCHES-2, a Phase 2b clinical study of ARO-ANG3, the company's investigational RNAi therapeutic being developed as a treatment for patients with mixed dyslipidemia
- Received Breakthrough Therapy designation from the U.S. Food and Drug Administration for ARO-AAT
- Announced that Janssen disclosed its collaboration with Arrowhead on investigational compound JNJ-75220795, which in a Phase 1 clinical study and is designed to reduce expression in the liver of patatin like phospholipase domain containing 3 (PNPLA3) as a potential treatment for patients with NASH
 - o Earned a \$10 million milestone from Janssen after Janssen dosed the fifth patient in a Phase 1 clinical study

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ARROWHEAD PHARMACEUTICALS, INC. CONSOLIDATED CONDENSED FINANCIAL INFORMATION (in thousands, except per share amounts)

	Y	ear Ended September 30,		
<u>OPERATING SUMMARY</u>		2021		2020
REVENUE	\$	138,287	\$	87,992
OPERATING EXPENSES				
Research and development		206,342		128,875
General and administrative expenses		80,981		52,276
TOTAL OPERATING EXPENSES		287,323		181,151
OPERATING INCOME (LOSS)		(149,036)		(93,159)
OTHER INCOME/(EXPENSE)		8,188		8,606
NET INCOME (LOSS)	\$	(140,848)	\$	(84,553)
NET INCOME (LOSS) PER SHARE (DILUTED)	\$	(1.36)	\$	(0.84)
WEIGHTED AVERAGE SHARES				
OUTSTANDING (DILUTED)		103,745		100,722
FINANCIAL POSITION SUMMARY	Sep	tember 30,	Sep	tember 30,
	-	2021	-	2020
CASH AND CASH EQUIVALENTS	\$	184,434	\$	143,583
SHORT-TERM INVESTMENTS AND				
MARKETABLE SECURITIES		183,355		171,910
LONG-TERM INVESTMENTS		245,595		137,487
TOTAL CASH RESOURCES (CASH AND				
INVESTMENTS)		613,384		452,980
OTHER ASSETS		96,764		69,524
TOTAL ASSETS		710,148		522,504
TOTAL CURRENT DEFERRED REVENUE		111,055		19,291
TOTAL LONG-TERM DEFERRED REVENUE		131,495		-
OTHER LIABILITIES		58,776		41,434
TOTAL LIABILITIES		301,326		60,725
TOTAL STOCKHOLDERS' EQUITY		408,822		461,779
TOTAL LIABILITIES AND STOCKHOLDERS'				
EQUITY	\$	710,148	\$	522,504
SHARES OUTSTANDING		104,327		102,376

About Arrowhead Pharmaceuticals

Arrowhead Pharmaceuticals develops medicines that treat intractable diseases by silencing the genes that cause them. Using a broad portfolio of RNA chemistries and efficient modes of delivery, Arrowhead therapies trigger the RNA interference mechanism to induce rapid, deep, and durable knockdown of target genes. RNA interference, or RNAi, is a mechanism present in

living cells that inhibits the expression of a specific gene, thereby affecting the production of a specific protein. Arrowhead's RNAi-based therapeutics leverage this natural pathway of gene silencing.

For more information, please visit www.arrowheadpharma.com, or follow us on Twitter @ArrowheadPharma. To be added to the Company's email list and receive news directly, please visit http://ir.arrowheadpharma.com/email-alerts.

Safe Harbor Statement under the Private Securities Litigation Reform Act:

This news release contains forward-looking statements within the meaning of the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. Any statements contained in this release except for historical information may be deemed to be forward-looking statements. Without limiting the generality of the foregoing, words such as "may," "will," "expect," "believe," "anticipate," "hope," "intend," "plan," "project," "could," "estimate," or "continue" are intended to identify such forward-looking statements. In addition, any statements that refer to projections of our future financial performance, trends in our business, expectations for our product pipeline or product candidates, including anticipated regulatory submissions and clinical program results, prospects or benefits of our collaborations with other companies, or other characterizations of future events or circumstances are forward-looking statements. These statements are based upon our current expectations and speak only as of the date hereof. Our actual results may differ materially and adversely from those expressed in any forward-looking statements as a result of numerous factors and uncertainties, including the impact of the ongoing COVID-19 pandemic on our business, the safety and efficacy of our product candidates, decisions of regulatory authorities and the timing thereof, the duration and impact of regulatory delays in our clinical programs, our ability to finance our operations, the likelihood and timing of the receipt of future milestone and licensing fees, the future success of our scientific studies, our ability to successfully develop and commercialize drug candidates, the timing for starting and completing clinical trials, rapid technological change in our markets, the enforcement of our intellectual property rights, and the other risks and uncertainties described in our most recent Annual Report on Form 10-K, subsequent Quarterly Reports on Form 10-Q and other documents filed with the Securities and Exchange Commission from time to time. We assume no obligation to update or revise forward-looking statements to reflect new events or circumstances.

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