

# Arrowhead Pharmaceuticals Reports Fiscal 2022 Second Quarter Results

May 10, 2022

- Conference Call and Webcast Today, May 10, 2022 at 4:30 p.m. ET

PASADENA, Calif.--(BUSINESS WIRE)--May 10, 2022-- Arrowhead Pharmaceuticals, Inc. (NASDAQ: ARWR) today announced financial results for its fiscal second quarter ended March 31, 2022. The company is hosting a conference call today, May 10, 2022, 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 <u>http://ir.arrowheadpharma.com/events.cfm</u>. For analysts that wish to participate in the conference call, please dial 855-215-6159 or 315-625-6887 and provide Conference ID 3791265.

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 3791265.

## Selected Recent Events

- Initiated the PALISADE Phase 3 clinical study to evaluate the efficacy and safety of ARO-APOC3, Arrowhead's investigational RNA interference (RNAi) therapeutic designed to inhibit the production of apolipoprotein C-III (APOC3), a key regulator of triglyceride metabolism, in adults with familial chylomicronemia syndrome
- Completed enrollment of 204 patients in the Phase 2b ARCHES-2 clinical study of ARO-ANG3, our investigational medicine designed to reduce production of angiopoietin-like protein 3 ANGPTL3 as a potential treatment for patients with mixed dyslipidemia
  - ARCHES-2 is expected to be complete around the end of 2022 and topline data are anticipated to be available in the first half of 2023
- Initiated the Phase 2 GATEWAY clinical study of ARO-ANG3, Arrowhead's investigational medicine designed to silence the hepatic expression of angiopoietin-like protein 3 (ANGPTL3), in patients with homozygous familial hypercholesterolemia (HoFH)
- Initiated a Phase 1/2 study of ARO-C3, Arrowhead's investigational medicine designed to reduce production of complement component 3 (C3) as a potential therapy for various complement mediated diseases, in up to 24 adult healthy volunteers, up to 24 adult patients with paroxysmal nocturnal hemoglobinuria (PNH), and up to 14 adult patients with complementmediated renal disease
- Filed Clinical Trial Applications (CTA) requesting regulatory clearance to begin clinical studies for two new investigational medicines designed to treat various muco-obstructive and inflammatory pulmonary conditions
  - ARO-MUC5AC, an investigational RNAi therapeutic designed to inhibit the production of mucin 5AC (MUC5AC)
  - ARO-RAGE, an investigational RNAi therapeutic designed to inhibit the production of Receptor for Advanced Glycation End products (RAGE)
- Formed a joint venture, Visirna Therapeutics, with Vivo Capital to expand the reach of innovative medicines in Greater China
- Broke ground on construction of a new drug manufacturing facility and announced awards of up to \$18.5 million in tax incentives from the city of Verona and the Wisconsin Economic Development Corporation

## Selected Fiscal 2022 Second Quarter Financial Results

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

	Three months ended March 31,				Six months ended March 31,			
OPERATING SUMMARY		2022		2021		2022		2021
REVENUE	\$	151,805	\$	32,811	\$	179,244	\$	54,113
OPERATING EXPENSES								
Research and development		75,985		44,697		141,750		81,251
General and administrative expenses		34,267		16,346		59,262		25,147
TOTAL OPERATING EXPENSES		110,252		61,043		201,012		106,398
OPERATING INCOME (LOSS)		41,553		(28,232)		(21,768)		(52,285)

OTHER INCOME/(EXPENSE)		813		1,414	 3,262	<u> </u>	4,735
NET INCOME (LOSS)	\$ 44,	366	\$	(26,818)	\$ (18,506)	\$	(47,550)
NET INCOME (LOSS) PER SHARE (DILUTED)	\$ (	.41	\$	(0.26)	\$ (0.18)	\$	(0.46)
WEIGHTED AVERAGE SHARES OUTSTANDING (DILUTED)	107,	929		103,867	 105,034		103,303
FINANCIAL POSITION SUMMARY	March 31, 2022		September 30, 2021				
CASH AND CASH EQUIVALENTS	\$ 86,	408	\$	184,434			
SHORT-TERM INVESTMENTS AND MARKETABLE SECURITIES	315,	487		183,355			
LONG-TERM INVESTMENTS	201,	590		245,595			
TOTAL CASH RESOURCES (CASH AND INVESTMENTS)	603,	485		613,384			
OTHER ASSETS	100,	090		96,764			
TOTAL ASSETS	703,	575		710,148			
TOTAL CURRENT DEFERRED REVENUE	97,	869		111,055			
TOTAL LONG-TERM DEFERRED REVENUE	89,	754		131,495			
OTHER LIABILITIES	63,	686		58,776			
TOTAL LIABILITIES	251,	309		301,326			
TOTAL STOCKHOLDERS' EQUITY	452,	266		408,822			
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY	\$703,	575	\$	710,148			
SHARES OUTSTANDING	105,	702		104,327			

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

## 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-X, subsequent Quarterly Reports on Form 10-Q and other documents filed with the Securities and Exchange Commission fr

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