# Arvinas Announces Changes to its Board of Directors

August 18, 2022

NEW HAVEN, Conn., Aug. 18, 2022 (GLOBE NEWSWIRE) -- Arvinas, Inc. (Nasdaq: ARVN), a clinical-stage biotechnology company creating a new class of drugs based on targeted protein degradation, today announced that Liam Ratcliffe, M.D., Ph.D. has stepped down from the company's Board and John Young has been appointed to join.

"Since joining our board in 2015, Liam has been a valuable member of our team, providing expert guidance, especially as a member of our Compensation Committee," said John Houston, Ph.D., President and Chief Executive Officer at Arvinas. "On behalf of my fellow Board members and Arvinas' leadership team, I would like to thank Liam for his dedication and impactful contributions."

Mr. Young joins Arvinas' Board with nearly 35 years of experience with Pfizer. He most recently served as Group President and Senior Advisor to the Pfizer CEO prior to his retirement from Pfizer in mid-2022 and was formerly Chief Business Officer at Pfizer. He became a member of Pfizer's Executive Leadership Team in 2012 and held a number of senior roles across the organization including Group President, Pfizer Innovative Health, Group President, Pfizer Essential Health, President & General Manager, Primary Care Business Unit, and Regional President, Primary Care, Europe, Canada.

Additionally, Mr. Young currently serves as a board member for Johnson Controls International (JCI), Haleon PLC (HLN), and Imbria Pharmaceuticals Inc., and will serve on the Board of Directors of Haleon, the Consumer Healthcare joint venture between GlaxoSmithKline and Pfizer, when it is separated in 2022. Previously, he served as a board member for the European Federation of Pharmaceutical Industries Associations, The Biotechnology Innovation Organization, and the National Committee on United States-China Relations. Mr. Young received a B.Sc. in Biological Science from Glasgow University and an MBA from Strathclyde Graduate Business School.

"John is an exceptional leader with deep experience and a strong track record in the pharmaceutical industry, and I am pleased to welcome him to Arvinas' Board of Directors," said Dr. Houston. "John is joining us at an exciting time, as Arvinas is on track to become a late-stage development company by year end. I am confident that John will be a tremendous asset as we aim to bring therapeutic options to patients who have few or none."

"I am extremely impressed with the tremendous work Arvinas is doing in the targeted protein degradation space," added Mr. Young. "I am thrilled to join the Arvinas team and contribute to the ongoing success that will hopefully one day impact patient lives."

**About Arvinas**
Arvinas is a clinical-stage biotechnology company dedicated to improving the lives of patients suffering from debilitating and life-threatening diseases through the discovery, development, and commercialization of therapies that degrade disease-causing proteins. Arvinas uses its proprietary PROTAC® Discovery Engine platform to engineer proteolysis targeting chimeras, or PROTAC® targeted protein degraders, that are designed to harness the body's own natural protein disposal system to selectively and efficiently degrade and remove disease-causing proteins. In addition to its robust preclinical pipeline of PROTAC® protein degraders against validated and "undruggable" targets, the company has three investigational clinical-stage programs: bavdegalutamide and ARV-766 for the treatment of men with metastatic castration-resistant prostate cancer; and ARV-471 for the treatment of patients with locally advanced or metastatic ER+/HER2- breast cancer. For more information, visit www.arvinas.com.

**Forward-Looking Statements**
This press release contains forward-looking statements that involve substantial risks and uncertainties, including statements regarding the potential advantages and therapeutic benefits of our product candidates, the future development and potential marketing approval and commercialization of our product candidates, including the initiation of and timing of data from our clinical trials. All statements, other than statements of historical facts, contained in this press release, including statements regarding our strategy, future operations, prospects, plans and objectives of management, are forward-looking statements. The words "anticipate," "believe," "estimate," "expect," "intend," "may," "might," "plan," "predict," "project," "target," "potential," "will," "would," "could," "should," "continue," and similar expressions are intended to identify forward-looking statements, although not all forward-looking statements contain these identifying words.

We may not actually achieve the plans, intentions or expectations disclosed in our forward-looking statements, and you should not place undue reliance on our forward-looking statements. Actual results or events could differ materially from the plans, intentions and expectations disclosed in the forward-looking statements we make as a result of various risks and uncertainties, including but not limited to:, whether we will be able to successfully conduct and complete clinical development of our product candidates, including whether we initiate and receive results from our clinical trials on our expected timelines or at all, obtain marketing approval for and commercialize our product candidates on our current timelines or at all and other important factors discussed in the "Risk Factors" sections contained in our quarterly and annual reports on file with the Securities and Exchange Commission. The forward-looking statements contained in this press release reflect our current views with respect to future events, and we assume no obligation to update any forward-looking statements except as required by applicable law. These forward-looking statements should not be relied upon as representing our views as of any date subsequent to the date of this release.

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