

UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
Washington, D.C. 20549

FORM 8-K

CURRENT REPORT  
Pursuant to Section 13 or 15(d)  
of the Securities Exchange Act of 1934

Date of Report (Date of earliest event reported): March 18, 2025

**Alumis Inc.**

(Exact name of registrant as specified in its charter)

Delaware  
(State or other jurisdiction  
of incorporation)

001-42143  
(Commission  
File Number)

86-1771129  
(IRS Employer  
Identification No.)

280 East Grand Avenue  
South San Francisco, California 94080  
(Address of principal executive offices)

Registrant's telephone number, including area code: (650) 231-6625

N/A  
(Former name or former address, if changed since last report.)

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:

- Written communications pursuant to Rule 425 under the Securities Act (17 CFR 230.425)
- 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))
- Pre-commencement communications pursuant to Rule 13e-4(c) under the Exchange Act (17 CFR 240.13e-4(c))

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, \$0.0001 par value per share	ALMS	The Nasdaq Global Select Market

Indicate by check mark whether the registrant is an emerging growth company as defined in 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

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.

**Item 8.01. Other Events.**

On March 18, 2025, Alumis Inc. and ACELYRIN, INC. released an updated presentation regarding their proposed merger as announced previously on February 6, 2025.

A copy of the presentation is furnished as Exhibit 99.1 to this Current Report on Form 8-K (the "Report"). The information set forth in this Report, including without limitation the presentation, is not deemed "filed" for purposes of Section 18 of the Securities Exchange Act of 1934, as amended (the "Exchange Act"), or incorporated by reference in any filing under the Securities Act of 1933, as amended, or the Exchange Act, except as may be expressly set forth by specific reference in such a filing.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits.

<b>Exhibit No.</b>	<b>Description</b>
<a href="#">99.1</a>	<a href="#">Presentation dated March 2025.</a>
104	Cover Page Interactive Data File (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 hereunto duly authorized.

**Alumis Inc.**

By: /s/ Sara Klein  
Sara Klein  
Chief Legal Officer

Dated: March 18, 2025

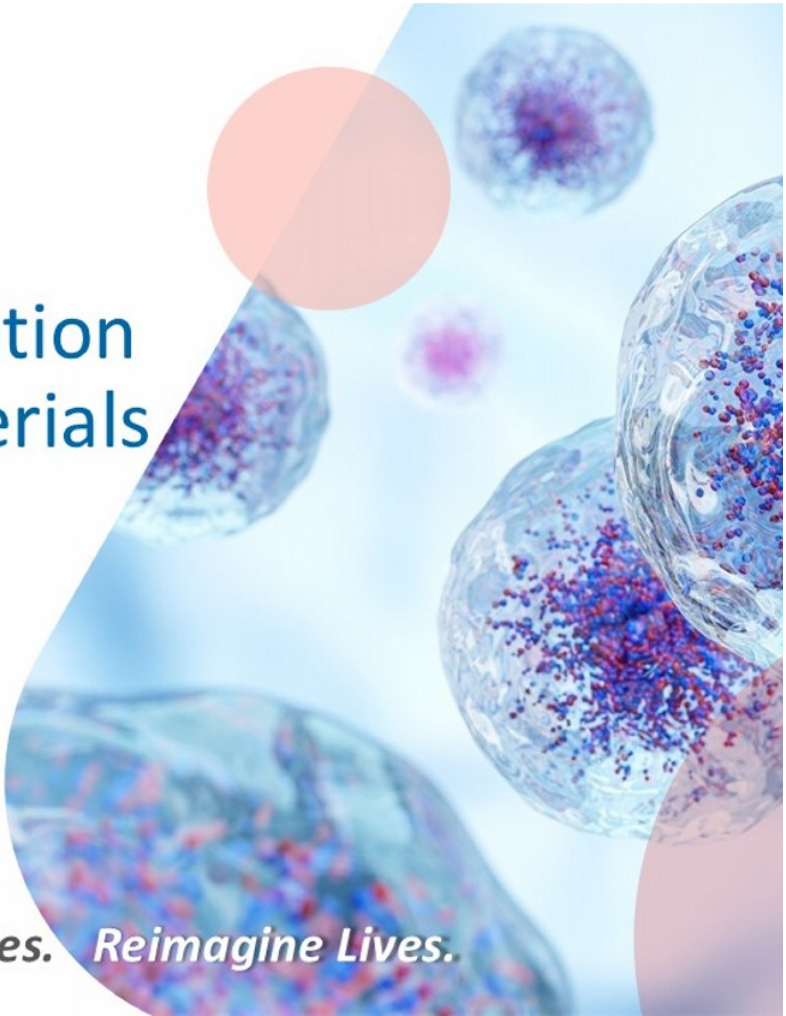
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# Corporate Presentation Supplemental Materials

March 2025

*Transform Therapies. Reimagine Lives.*



#### Forward-Looking Statements

This presentation contains forward looking statements within the meaning of federal securities laws, including the "safe harbor" provisions of the Private Securities Litigation Reform Act of 1995. Such statements are based upon current plans, estimates and expectations of Alumis and ACELYRIN, Inc. ("ACELYRIN") in light of historical results and trends, current conditions and potential future developments, and are subject to various risks and uncertainties that could cause actual results to differ materially from such statements. The inclusion of forward looking statements in this presentation should not be regarded as a representation that such plans, estimates and expectations will be achieved. Words such as "anticipate," "expect," "project," "intend," "believe," "may," "will," "should," "plan," "could," "continue," "target," "contemplate," "estimate," "forecast," "guidance," "pursue," "likely," and words of similar substance used in connection with any discussion of future plans, actions or events identify forward-looking statements. All statements, other than statements of historical facts, including express or implied statements regarding conversion of equity interests contemplated by the agreement and plan of merger, dated as of February 6, 2025, by and among the parties (the "merger agreement"); the issuance of common stock of Alumis contemplated by the merger agreement; the expected filing by Commission (the "SEC") of a registration statement on Form S-4 (the "registration statement") and a joint proxy statement/prospectus of Alumis and ACELYRIN to be included therein (the "joint proxy statement/prospectus"); the expected timing of the closing of the proposed transaction considering the various closing conditions; the expected benefits of the proposed transaction; the sufficiency of the combined company's capital resources; the combined company's cash runway; the competitive ability and position of a pipeline of the combined company; and any assumptions underlying any of the foregoing, are forward-looking statements.

Risks and uncertainties include, among other things, (i) the risk that the proposed transaction may not be completed in a timely basis or at all, which may adversely affect Alumis' and ACELYRIN's businesses and the price of their respective securities; (ii) the potential failure otherwise, the required approvals of the proposed transaction, including stockholder approvals by both Alumis' stockholders and ACELYRIN's stockholders, and the potential failure to satisfy the other conditions to the consummation of the transaction; (iii) the effect of the completion of the proposed transaction on each of Alumis' or ACELYRIN's ability to attract, motivate, retain and hire key personnel and maintain relationships with partners, suppliers and others with whom Alumis or ACELYRIN does business, or on Alumis' or ACELYRIN's or (iv) that the proposed transaction may divert management's attention from each of Alumis' and ACELYRIN's ongoing business operations; (v) the risk of any legal proceedings related to the proposed transaction or otherwise, or the impact of the proposed transaction thereon; (vi) that Alumis or ACELYRIN may be adversely affected by other economic, business and/or competitive factors; (vii) the occurrence of any event, change or other circumstance that could give rise to the termination of the merger agreement, including in circumstances where ACELYRIN is required to pay a termination fee; (viii) the risk that restrictions during the pendency of the proposed transaction may impact Alumis' or ACELYRIN's ability to pursue certain business opportunities or strategic transactions; (ix) the risk that the anticipated benefits and synergies may not be fully realized or may take longer to realize than expected; (x) the impact of legislative, regulatory, economic, competitive and technological changes; (xi) risks relating to the value of Alumis securities to be issued in the proposed transaction; (xii) the risk that integral closing may not occur as anticipated or the combined company may not be able to achieve the growth prospects expected from the transaction; (xiii) the effect of the announcement, pendency or completion of the proposed transaction on the market price of the common stock of Alumis and ACELYRIN; (xiv) the implementation of each of Alumis' and ACELYRIN's business model and strategic plans for product candidates and pipeline, and challenges inherent in developing, commercializing, manufacturing, launching, marketing and selling potential existing and new product candidates, progress, results and costs of developing Alumis' and ACELYRIN's product candidates and any future product candidates, including conducting preclinical studies and clinical trials, and otherwise related to the research and development of Alumis' and ACELYRIN's pipeline of product candidates, including obtaining and maintaining regulatory approval for Alumis' and ACELYRIN's current or future product candidates, and any related restrictions, limitations and/or warnings in the label of any approved product; (xv) the market for, adoption (including rate and degree of reimbursement of Alumis' and ACELYRIN's product candidates, if approved, and their respective abilities to compete with therapies and procedures that are rapidly growing and evolving; (xvi) uncertainties in contractual relationships, including collaborations, partnerships and other arrangements with third party suppliers and manufacturers; (xvii) the ability of each of Alumis and ACELYRIN to establish and maintain intellectual property protection for products or avoid or defend claims of infringement; (xviii) Alumis' ability to successfully integrate ACELYRIN's operations; (xix) potential delays in initiating, enrolling or completing preclinical studies and clinical trials.

These risks, as well as other risks related to the proposed transaction, will be described in the registration statement and the joint proxy statement/prospectus that will be filed with the SEC in connection with the proposed transaction. While the list of factors presented here is not intended to be a complete statement of all potential risks and uncertainties, for additional information about other factors that could cause actual results to differ materially from the forward looking statements, please refer to Alumis' and ACELYRIN's respective periodic reports and other filings with the SEC, including the risk factors identified in Alumis' and ACELYRIN's most recent Quarterly Reports on Form 10-Q and/or Annual Reports on Form 10-K. The risks in the SEC filings cited above are not exclusive and further information concerning Alumis and ACELYRIN and their respective businesses, including factors that potentially could materially affect their respective businesses, financial conditions or operating results, may emerge in the future. Readers should carefully review these forward looking statements, and not to place undue reliance on any forward-looking statements, which speak only as of the date hereof. Readers should also carefully review the risk factors described in other documents filed with the SEC.

The forward-looking statements included in this presentation are made only as of the date hereof. Alumis assumes no obligation and does not intend to update these forward-looking statements, even if new information becomes available in the future, except as required by law.

This presentation contains trademarks, service marks, trade names and copyrights of Alumis and other companies which are the property of their respective owners. This presentation discusses product candidates that are under clinical study and which have not yet been approved by the U.S. Food and Drug Administration. No representation is made as to the safety or effectiveness of these product candidates for the uses for which they are being studied. This presentation also contains estimates and other statistical data made by independent parties and by us, based on data about our industry. These data involve a number of assumptions and limitations, and you are cautioned not to give undue weight to such estimates. We have not independently verified the data generated by independent parties and cannot guarantee their accuracy or completeness. Assumptions, and estimates of our future performance and the future performance of the markets in which we operate are necessarily subject to a high degree of uncertainty and risk.

#### Additional Information and Where to Find It

In connection with the proposed merger, Alumis intends to file with the SEC the registration statement, which will include the joint proxy statement/prospectus. After the registration statement has been declared effective by the SEC, the joint proxy statement/prospectus of Alumis and ACELYRIN, BEFORE MAKING ANY VOTING OR INVESTMENT DECISION, SECURITY HOLDERS OF ALUMIS AND ACELYRIN ARE URGED TO READ THE JOINT PROXY STATEMENT/PROSPECTUS (INCLUDING ALL AMENDMENTS AND SUPPLEMENTS THERETO) AND OTHER DOCUMENTS THAT WILL BE FILED WITH THE SEC WHEN THEY BECOME AVAILABLE BECAUSE THEY WILL CONTAIN IMPORTANT INFORMATION ABOUT THE PROPOSED MERGER. Investors and security holders will be able to obtain copies of the joint proxy statement/prospectus (when available) from Alumis and ACELYRIN with the SEC, without charge, through the website maintained by the SEC at [www.sec.gov](http://www.sec.gov). Copies of the documents filed with the SEC by Alumis will be available free of charge under the SEC Filings heading of the Investor Relations section of Alumis' website at <https://investors.alumis.com>. Copies of the documents filed with the SEC by ACELYRIN will be available free of charge under the Financials & Filings heading of the Investor Relations section of ACELYRIN's website at <https://investors.aceelryn.com>.

#### Participants in the Solicitation

Alumis and ACELYRIN and their respective directors and executive officers may be deemed to be participants in the solicitation of proxies in respect of the proposed transaction. Information about Alumis' directors and executive officers is set forth in Alumis' registration statement on Form S-4, which was filed with the SEC on June 24, 2024. Information about ACELYRIN's directors and executive officers is set forth in the proxy statement for ACELYRIN's 2024 Annual Meeting of Stockholders, which was filed with the SEC on April 22, 2024, and ACELYRIN's registration statement on Form S-4, which was filed with the SEC on May 28, 2024, August 13, 2024 and December 10, 2024. Stockholders may obtain additional information regarding the interests of such participants by reading the registration statement and the joint proxy statement/prospectus and other relevant materials to the proposed merger when they become available. Investors should read the joint proxy statement/prospectus carefully when it becomes available before making any voting or investment decisions.

#### No Offer or Solicitation

This presentation shall not constitute an offer to sell or the solicitation of an offer to buy any securities or a solicitation of any vote or approval, nor shall there be any sale of securities in any jurisdiction in which such offer, solicitation or sale would be unlawful prior to registration under applicable securities laws of any such jurisdiction. No offering of securities shall be made except by means of a prospectus meeting the requirements of Section 10 of the Securities Act of 1933, as amended.



# Creating a Leading I&I Company with Multiple Upcoming Expected Development Milestones and Extended Runway into 2027



- › **Creates a late-stage clinical biopharma company** dedicated to innovating and commercializing transformative therapies for immune-mediated diseases
- › Differentiated pipeline with multiple upcoming milestones expected, including:
  - › Topline data from **Phase 3 ONWARD** trials for Alumis' ESK-001 in plaque psoriasis **now expected for readout in 1Q 2026**
  - › Topline data from **Phase 2b LUMUS** trial in systemic lupus erythematosus for **readout in 2026**
  - › **Phase 2** clinical trial **initiation** for Alumis' A-005 in MS
  - › Addition of **lonigutamab**, a subcutaneously delivered anti-IGF-1R inhibitor for disease (TED), with committed capital for a clinical development program with potential differentiation
- › **Pro forma cash of ~\$737 million** as of December 31, 2024, provides runway **beyond multiple expected clinical readouts**<sup>1</sup>
- › Combined company to benefit from **world-class leadership team with strong operating discipline and capital efficiency**



1. Preliminary, unaudited and subject to change

# Late-Stage Pipeline with Multiple Near-Term Catalysts

TARGET		DEVELOPMENT				
		PRECLINICAL	PHASE 1	PHASE 2	PHASE 3	
ESK-001 (TYK2)	Moderate-to-Severe Plaque Psoriasis (PsO)					14
	Systemic Lupus Erythematosus (SLE)					202
	Add'l indications TBD PsA, IBD, etc.	<i>Other suitable development opportunities for ESK-001 include expanded Psoriasis indications and immunological indications outside the CNS</i>				
Lonigutamab (anti-IGF-1R)	Thyroid Eye Disease (TED)					d
A-005 (TYK2)	Multiple Sclerosis (MS)					202 20.
	Add'l indications TBD PAR, AD, etc.	<i>Expanded opportunities for A-005 include other neuroinflammatory or neurodegenerative conditions</i>				
IRF5, Additional Targets	Undisclosed					YE2

> Post transaction close, we expect to evaluate lonigutamab's promising potential differentiation profile in TED with approximately \$25-50M of committed capital and the combined expertise of Alumis and the lonigutamab team



# Alumis Combination Optimizes Outcome for Patients and ACELYRIN Stockholders

Lonigutamab is a Differentiated Molecule for the Treatment of TED and Deserves Continued Exploration for the Benefit of Patients

- › Potential for commercially attractive product profile with improved safety and IV-like efficacy
- › Robust and compelling Phase 1/2 clinical data that may enable clinical development plan that optimizes efficacy and safety

Alumis is the Right Partner for ACELYRIN to Continue Clinical Development Plan of Lonigutamab

- › World-class leadership team with proven record of developing innovative therapies, operating discipline and capital efficiency
- › The Alumis team has significant experience in optimizing PK/PD and target product profile
- › Alumis has significant ophthalmology expertise, and its Chief Commercial Officer ran the TEPEZZA franchise at Amgen

Independent of the Proposed Merger With Alumis, ACELYRIN Expects to Continue the Development of Lonigutamab

After Careful Consideration, the ACELYRIN Board Determined that the Indication of Interest From Concentra Was Unlikely to Lead to a Superior Proposal



# Comparing Alternative Options for ACELYRIN Stockholders

	Combined Company	Standalone ACELYRIN
<b>Product Candidates</b>	<ul style="list-style-type: none"> <li>• ESK-001</li> <li>• Lonigutamab</li> <li>• A-005</li> <li>• IRF5</li> <li>• Additional Targets</li> </ul> <p><b>Deep Pipeline of Differentiated Assets</b></p>	<ul style="list-style-type: none"> <li>• Lonigutamab</li> </ul>
<b>Upcoming Milestones</b>	<ul style="list-style-type: none"> <li>• <b>Mid-2025:</b> Finalize Plan for Lonigutamab</li> <li>• <b>2H25:</b> A-005 – MS Phase 2 Initiation</li> <li>• <b>2H25:</b> IND Filing for 3<sup>rd</sup> Clinical Candidate</li> <li>• <b>2025:</b> Once-Daily Formulation Established for ESK-001</li> <li>• <b>1Q26:</b> ESK-001 – PsO Phase 3 Topline Data</li> <li>• <b>2026:</b> ESK-001 – SLE Phase 2b Topline Data</li> <li>• <b>2026:</b> A-005 – MS Phase 2 Topline Data</li> </ul> <p><b>Multiple Anticipated Near-Term Milestones Including Major Catalysts</b></p>	<ul style="list-style-type: none"> <li>• Previously announced start of program in Q1 2025 with top line data in Q2 2026</li> </ul>
<b>Clinical-Stage Indications</b>	<ul style="list-style-type: none"> <li>• PsO</li> <li>• SLE</li> <li>• TED</li> <li>• MS</li> </ul> <p><b>Substantial Market Potential Across Immune-Mediated Diseases, Totaling \$60Bn<sup>1</sup></b></p>	<ul style="list-style-type: none"> <li>• TED</li> </ul>
<b>Potential Additional Indications</b>	<ul style="list-style-type: none"> <li>• Additional immunological indications (e.g. PsA, cutaneous lupus, IBD, etc.)</li> <li>• Other neuroinflammatory or neurodegenerative (e.g. PAR, AD, etc.)</li> </ul> <p><b>Broad Potential for Additional Indications</b></p>	



1. GlobalData, market research reports; Market size estimates for 2030 worldwide



# Comprehensive Process Led to Value-Maximizing Opportunity for A Stockholders

## **The Proposed Merger With Alumis Is the Result of a Comprehensive Process Conducted by ACELYRIN Board**

- › This process began in Q2 2024 and encompassed interactions with ~25 potential counterparties
- › A wide variety of potential transactions were explored with the help of a leading independent financial advisor

## **The Combination With Alumis Is Expected To Be Significantly Value Accretive to ACELYRIN Stockholders**

- › Value accretion is driven by synergies, scale and shared ownership of a more diversified portfolio, and Alumis near-term value inflecting catalysts
- › ACELYRIN Board determined that a transaction that supported the continued development of lonigutamab represented the best opportunity for long-term value creation for ACELYRIN stockholders

## **The Transaction Is the Best Path Forward to Exploit the Full Potential Opportunity for Lonigutamab**

- › The Alumis combination provides the assets, resources, and talent for substantial long term value accretion



# Alumis + ACELYRIN: A Well Characterized Late-Stage Portfolio With Opportunity

## Large Opportunity

Three highly differentiated clinical programs, (two TYK2 inhibitors and IGF provide strategic opportunities across a broad range of immune mediated

## High PTS

Genomic, proteomic and clinical insights validate and drive our Research a strategy and increase the probability of technical success (PTS) of our prog

## Market Maker

High efficacy oral and subcutaneous therapies provide significant market r

## Near Term Pivotal Data

ESK-001: Two major value inflection points expected in large indications

## Differentiated MOA

Lonigutamab: A next-generation subcutaneous anti-IGF-1R with best-in-cl  
TED

## First-In-Class

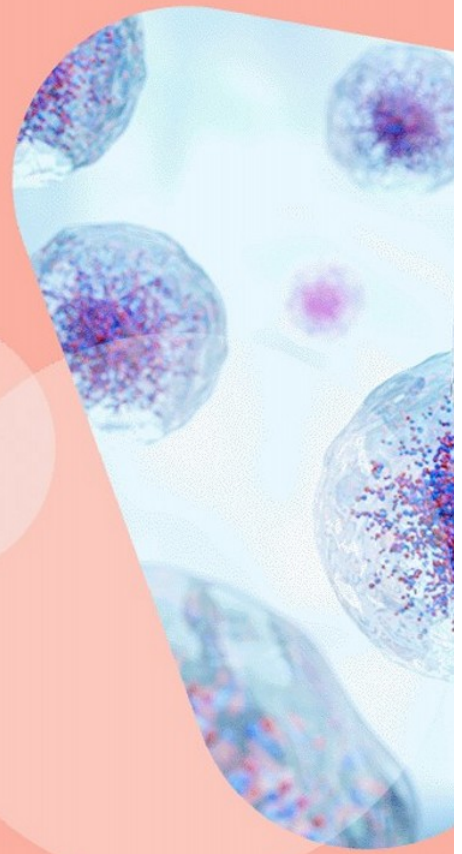
A-005: Potentially first-in-class fully CNS-penetrant TYK2 inhibitor expands addressable indications, including those within the CNS

## Proven Leadership

Experienced team with strong track record in value creation



Thank You!



## Forward-Looking Statements

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These risks, as well as other risks related to the proposed transaction, will be described in the registration statement and the joint proxy statement/prospectus that will be filed with the SEC in connection with the proposed transaction. While the list of factors presented here and the list of factors to be presented in the registration statement are considered representative, no such list should be considered to be a complete statement of all potential risks and uncertainties. For additional information about other factors that could cause actual results to differ materially from those described in the forward-looking statements, please refer to Alumis' and ACELYRIN's respective periodic reports and other filings with the SEC, including the risk factors identified in Alumis' and ACELYRIN's most recent Quarterly Reports on Form 10-Q and/or Annual Reports on Form 10-K. The risks and uncertainties described above and in the SEC filings cited above are not exclusive and further information concerning Alumis and ACELYRIN and their respective businesses, including factors that potentially could materially affect their respective businesses, financial conditions or operating results, may emerge from time to time. Readers are urged to consider these factors carefully in evaluating these forward-looking statements, and not to place undue reliance on any forward-looking statements, which speak only as of the date hereof. Readers should also carefully review the risk factors described in other documents Alumis and ACELYRIN file from time to time with the SEC.

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#### **Participants in the Solicitation**

Alumis and ACELYRIN and their respective directors and executive officers may be deemed to be participants in the solicitation of proxies in respect of the proposed transaction. Information about Alumis' directors and executive officers is set forth in Alumis' registration statement on Form S-1/A (File No. 333-280068), which was filed with the SEC on June 24, 2024. Information about ACELYRIN's directors and executive officers is set forth in the proxy statement for ACELYRIN's 2024 Annual Meeting of Stockholders, which was filed with the SEC on April 22, 2024, and ACELYRIN's Current Reports on Form 8-K filed with the SEC on May 28, 2024, August 13, 2024 and December 10, 2024. Stockholders may obtain additional information regarding the interests of such participants by reading the registration statement and the joint proxy statement/prospectus and other relevant materials to be filed with the SEC regarding the proposed merger when they become available. Investors should read the joint proxy statement/prospectus carefully when it becomes available before making any voting or investment decisions.

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