## 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): August 09, 2023

## Surrozen, Inc.

(Exact name of Registrant as Specified in Its Charter)

Delaware (State or Other Jurisdiction of Incorporation) 001-39635 (Commission File Number) 98-1556622 (IRS Employer Identification No.)

171 Oyster Point Blvd Suite 400 South San Francisco, California (Address of Principal Executive Offices)

94080 (Zip Code)

Registrant's Telephone Number, Including Area Code: +1 (650) 489-9000

	eck the appropriate box below if the Form 8-K filing is inte owing provisions:	nded to simultaneously s	atisfy the filing obligations of the registrant under any of the				
	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 14	d-2(b) under the Exchan	ge 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 regi	istered pursuant to Sect	ion 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	SRZN	The Nasdaq Capital Market				
Redeemable warrants, each whole warrant exercisable for one share of Common Stock		SRZNW	The Nasdaq Capital Market				
	icate by check mark whether the registrant is an emerging g pter) or Rule 12b-2 of the Securities Exchange Act of 1934		ed in Rule 405 of the Securities Act of 1933 (§ 230.405 of this pter).				
Em	erging 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.  $\Box$ 

#### Item 2.02 Results of Operations and Financial Condition.

On August 9, 2023, Surrozen, Inc. (the "Company") issued a press release announcing its financial results for the quarter ended June 30, 2023. A copy of the press release is attached as Exhibit 99.1 to this Current Report on Form 8-K.

The information set forth under this "Item 2.02. Results of Operations and Financial Condition" (including the exhibit referenced herein) shall not be deemed to be "filed" for the purposes of Section 18 of the Securities Exchange Act of 1934, as amended, nor shall it be incorporated by reference in any filing made by the Company pursuant to the Securities Act of 1933, as amended.

#### Item 9.01 Financial Statements and Exhibits.

#### (d) Exhibits

Exhibit No.	Description
99.1 104	Press Release of Surrozen, Inc. dated August 9, 2023 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.

SURROZEN, INC.

Date: August 9, 2023 By: /s/ Charles Williams

Name: Charles Williams Title: Chief Financial Officer

#### Surrozen Provides Second Quarter 2023 Financial Results

SZN-043 Phase 1a clinical trial in patients with chronic liver disease and in healthy volunteers continues to enroll with safety data expected by the end of 2023

SZN-1326 Phase 1a clinical trial in healthy volunteers continues to enroll with safety data expected by the end of 2023

Corporate prioritization efforts expected to provide cash runway extension into 2025

SOUTH SAN FRANCISCO, Calif., August 9, 2023 (GLOBE NEWSWIRE) -- Surrozen, Inc. ("Surrozen" or the "Company") (Nasdaq: SRZN), a company pioneering targeted therapeutics that selectively activate the Wnt pathway for tissue repair and regeneration, today provided second quarter 2023 financial results and corporate and pipeline updates.

"During the second quarter of 2023, we made important progress in the Phase 1a clinical trials for our lead candidates, SZN-043 and SZN-1326," said Craig Parker, President and Chief Executive Officer of Surrozen. "Continued corporate prioritization activities in 2023 are targeted to deliver proof-of-concept data on our two clinical development candidates in the second half of 2024. We remain focused on clinical validation of our innovative technologies for Wnt pathway modulation to deliver high-value therapies to people with severe diseases."

#### **Research and Development Pipeline Updates**

#### SZN-043

- Enrollment in the Phase 1a clinical trial continues in patients with chronic liver disease and healthy volunteers
- Expect safety data by the end of 2023
- Expect to initiate a Phase 1b clinical trial in patients with alcoholic hepatitis in 2024 and anticipate proof of concept data will potentially be available in the second half of 2024

#### SZN-1326

- Enrollment in the Phase 1a clinical trial continues in healthy volunteers
- Expect safety data by the end of 2023
- Expect to initiate a Phase 1b clinical trial in 2024 in patients with ulcerative colitis and anticipate proof of concept data will potentially be available in the second half of 2024

#### **Corporate Updates**

#### **Corporate Prioritization Activities**

In July 2023, Surrozen implemented a plan to further prioritize and focus resources on the development of the clinical programs for SZN-043 and SZN-1326 which is expected to extend the cash runway into 2025.

#### Corporate Partnerships

Surrozen executed a partnership with Boehringer Ingelheim in the fourth quarter of 2022 to develop a Wnt agonist, SZN-413, for the treatment of people with retinal diseases. We anticipate the potential to nominate the lead Fzd-4 targeted Wnt agonist development candidate in 2024, which would trigger a \$10.0 million milestone payment to the Company.

#### Financial Results for the Second Quarter Ended June 30, 2023

**Cash Position:** Cash, cash equivalents and marketable securities for the second quarter ended June 30, 2023 were \$53.4 million, compared to \$61.7 million as of March 31, 2023.

Research and Development Expenses: Research and development expenses for the second quarter ended June 30, 2023 were \$6.9 million, as compared to \$9.6 million for the same period in 2022. The decrease was primarily as a result of the restructuring plan implemented in the first quarter of 2023 to prioritize and focus our resources on key clinical and discovery programs. Research and development expenses include non-cash stock-based compensation expenses of \$0.3 million for the second quarter ended June 30, 2023, as compared to \$0.4 million for the same period in 2022.

**General and Administrative Expenses:** General and administrative expenses for the second quarter ended June 30, 2023 were \$3.3 million, as compared to \$4.5 million for the same period in 2022. The decrease was primarily related to the employee retention tax credits received in 2023. General and administrative expenses include non-cash stock-based compensation expenses of \$0.8 million for the second quarter ended June 30, 2023, as compared to \$0.6 million for the same period in 2022.

**Interest Income:** Interest income for the second quarter ended June 30, 2023 was \$0.6 million, as compared to \$0.1 million for the same period in 2022. The increase was primarily related to the increase in interest rates on money market funds and marketable securities.

Other Income (Expense), Net: Other income (expense), net for the second quarter ended June 30, 2023 was a net other income of \$0.3 million, as compared to \$0.1 million for the same period in 2022. The increase was primarily due to a decrease in expense related to the commitment shares issued under an equity purchase agreement entered into in 2022.

Net Loss: Net loss for the second quarter ended June 30, 2023 was \$9.4 million, as compared to \$13.9 million for the same period in 2022.

#### **About SZN-043 for Severe Alcoholic Hepatitis**

SZN-043 is the first development candidate using Surrozen's SWEETS™ technology. Surrozen is developing SZN-043 for severe liver diseases, initially focusing on severe alcoholic hepatitis. The company is enrolling patients in a Phase 1a clinical trial in patients with chronic liver disease and healthy volunteers. The Phase 1a clinical trial is ongoing and the company expects safety data by the end of 2023. Surrozen expects to initiate a Phase 1b clinical trial in patients with alcoholic hepatitis in 2024 and anticipates proof of concept data could potentially be available in the second half of 2024.

#### **About SZN-1326 for Ulcerative Colitis**

SZN-1326 is the first development candidate designed using Surrozen's SWAP™ technology and targets the Wnt-signaling pathway in the intestinal epithelium. Surrozen is initially developing SZN-1326 for moderate to severe ulcerative colitis. Enrollment is ongoing in the Phase 1a clinical trial in healthy volunteers. Surrozen expects safety data from the Phase 1a clinical trial in healthy volunteers by the end of 2023. Surrozen also expects to initiate a Phase 1b clinical trial in patients with ulcerative colitis in 2024 and anticipates proof-of-concept data will potentially be available in the second half of 2024.

#### **About SZN-413 for Retinal Diseases**

SZN-413 is a bi-specific antibody targeting Fzd4-mediated Wnt signaling designed using Surrozen's SWAP™ technology. It is currently being developed for the treatment of retinal vascular-associated diseases. Data generated by Surrozen with SZN-413 in preclinical models of retinopathy demonstrated that SZN-413 could potently stimulate Wnt signaling in the eye, induce normal retinal vessel regrowth, suppress pathological vessel growth and reduce vascular leakage. This novel approach could thus potentially allow for regeneration of healthy eye tissue, not only halting retinopathy, but possibly allowing for a full reversal of the patient's disease.

In the fourth quarter of 2022, Surrozen entered into a strategic partnership with Boehringer Ingelheim for the research and development of SZN-413 for the treatment of retinal diseases. Under the terms of the agreement, Boehringer Ingelheim received an exclusive, worldwide license to develop SZN-413 and other Fzd4-specific Wnt-modulating molecules for all purposes, including as a treatment for retinal diseases, in exchange for an upfront payment to Surrozen of \$12.5 million. Surrozen will also be eligible to receive up to \$587.0 million in success-based development, regulatory, and commercial milestone payments, in addition to mid-single digit to low-double digit royalties on sales. After an initial period of joint research, Boehringer Ingelheim will assume all development and commercial responsibilities.

#### **About Wnt Signaling**

Wnt signaling plays key roles in the control of development, homeostasis, and regeneration of many essential organs and tissues, including liver, intestine, lung, kidney, retina, central nervous system, cochlea, bone, and others. Modulation of Wnt signaling pathways has potential for treatment of degenerative diseases and tissue injuries. Surrozen's platform and proprietary technologies have the potential to overcome the limitations in pursuing the Wnt pathway as a therapeutic strategy.

#### **About Surrozen**

Surrozen is a clinical stage biotechnology company discovering and developing drug candidates to selectively modulate the Wnt pathway. Surrozen is developing tissue-specific antibodies designed to engage the body's existing biological repair mechanisms with a current focus on inflammatory bowel, severe liver and eye diseases. For more information, please visit www.surrozen.com.

#### **Forward Looking Statements**

This press release contains certain forward-looking statements within the meaning of the federal securities laws. Forward-looking statements generally are accompanied by words such as "will," "plan," "intend," "potential," "expect," "could," or the negative of these words and similar expressions that predict or indicate future events or trends or that are not statements of historical matters. These forward-looking statements include, but are not limited to, statements regarding Surrozen's discovery, research and development activities, in particular its development plans for its product candidates SZN-1326, SZN-043, and SZN-413 (including anticipated clinical development timelines and the availability of data, the potential for such product candidates to be used to treat human disease), the potential and timeline to nominate the lead development candidate pursuant to its partnership with Boehringer Ingelheim and its expectations with respect to its cash runway. These statements are based on various assumptions, whether or not identified in this press release, and on the current expectations of the management of Surrozen and are not predictions of actual performance. These forward-looking statements are provided for illustrative purposes only and are not intended to serve as, and must not be relied on as, a guarantee, an assurance, a prediction, or a definitive statement of fact or probability. Actual events and circumstances are difficult or impossible to predict and will differ from assumptions. Many actual events and circumstances are beyond the control of Surrozen. These forward-looking statements are subject to a number of risks and uncertainties, including the initiation, cost, timing, progress and results of research and development activities, preclinical or and clinical trials with respect to SZN-1326, SZN-043, SZN-413 and potential future drug candidates; the Company's ability to fund its preclinical and clinical trials and development efforts, whether with existing funds or through additional fundraising; Surrozen's ability to identify, develop and commercialize drug candidates; Surrozen's ability to successfully complete preclinical and clinical studies for SZN-1326, SZN-043, SZN-413, or other future product candidates; the effects that arise from volatility in global economic, political, regulatory and market conditions; and all other factors discussed in Surrozen's Annual Report on Form 10-K for the year ended December 31, 2022 and Surrozen's Quarterly Report on Form 10-Q for the quarter ended June 30, 2023 under the heading "Risk Factors," and other documents Surrozen has filed, or will file, with the Securities and Exchange Commission. If any of these risks materialize or our assumptions prove incorrect, actual results could differ materially from the results implied by these forward-looking statements. There may be additional risks that Surrozen presently does not know, or that Surrozen currently believes are immaterial, that could also cause actual results to differ from those contained in the forward-looking statements. In addition, forward-looking statements reflect Surrozen's expectations, plans, or forecasts of future events and views as of the date of this press release. Surrozen anticipates that subsequent events and developments will cause its assessments to change. However, while Surrozen may elect to update these forward-looking statements at some point in the future, Surrozen specifically disclaims any obligation to do so, except as required by law. These forward-looking statements should not be relied upon as representing Surrozen's assessments of any date after the date of this press release. Accordingly, undue reliance should not be placed upon the forward-looking statements.

#### **Investor and Media Contact:**

Investorinfo@surrozen.com

# SURROZEN, INC. Condensed Consolidated Statements of Operations (In thousands, except per share amounts) (Unaudited)

		Three Months Ended June 30,		Six Months Ended June 30,				
		2023		2022		2023		2022
Operating expenses:								
Research and development	\$	6,937	\$	9,581	\$	15,023	\$	18,952
General and administrative		3,338		4,491		8,637		9,613
Restructuring		_		_		1,207		_
Total operating expenses		10,275		14,072		24,867		28,565
Loss from operations		(10,275)		(14,072)		(24,867)		(28,565)
Interest income		623		60		1,170		109
Other income (expense), net		265		87		13		6,584
Net loss	\$	(9,387)	\$	(13,925)	\$	(23,684)	\$	(21,872)
Net loss per share attributable to common								
stockholders, basic and diluted	\$	(0.31)	\$	(0.40)	\$	(0.79)	\$	(0.63)
Weighted-average shares used in computing net loss per share attributable to common								
stockholders, basic and diluted		30,073		34,945		30,022		34,904

# SURROZEN, INC. Condensed Consolidated Balance Sheets (In thousands)

	June 30, 2023		December 31, 2022 <sup>(1)</sup>		
	(Ui	naudited)			
Assets					
Current assets:					
Cash and cash equivalents	\$	36,496	\$	24,690	
Accounts receivable		1,978		1,978	
Short-term marketable securities		16,899		51,148	
Prepaid expenses and other current assets		2,206		3,489	
Total current assets		57,579		81,305	
Property and equipment, net		2,966		3,630	
Operating lease right-of-use assets		2,655		3,268	
Restricted cash		405		405	
Other assets		394		827	
Total assets	\$	63,999	\$	89,435	
Liabilities and stockholders' equity					
Current liabilities:					
Accounts payable	\$	186	\$	658	
Accrued and other liabilities		3,901		6,848	
Lease liabilities, current portion		2,364		2,226	
Total current liabilities		6,451		9,732	
Lease liabilities, noncurrent portion		2,164		3,376	
Warrant liabilities		326		326	
Total liabilities		8,941		13,434	
Stockholders' equity:					
Preferred stock		_		_	
Common stock		3		3	
Additional paid-in-capital		257,387		254,892	
Accumulated other comprehensive income (loss)		5		(241)	
Accumulated deficit		(202,337)		(178,653)	
Total stockholders' equity		55,058		76,001	
Total liabilities and stockholders' equity	\$	63,999	\$	89,435	

<sup>(1)</sup> Derived from the audited financial statements, included in the Company's Annual Report on Form 10-K for the year ended December 31, 2022.