

**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): **May 17, 2021**

**VYANT BIO, INC.**

(Exact Name of Company as Specified in its Charter)

**Delaware**  
(State or Other Jurisdiction  
of Incorporation)

**001-35817**  
(Commission  
File Number)

**04-3462475**  
(IRS Employer  
Identification No.)

**2 Executive Campus  
2370 State Route 70, Suite 310  
Cherry Hill, NJ 08002**

(Address of Principal Executive Offices) (Zip Code)

Company's telephone number, including area code **(201) 479-8126**

(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 Company under any of the following provisions (see General Instruction A.2. below):

- 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))

Indicate by check mark whether the Company is an emerging growth company as defined by Rule 405 of the Securities Act of 1933 (17 §230.405) or Rule 12b-2 of the Securities Exchange Act of 1934 (17 CFR §240.12b-2).

Emerging growth company

If an emerging growth company, indicate by check mark if the Company 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.

Securities registered pursuant to Section 12(b) of the Act:

<u>Title of each class</u>	<u>Trading Symbol(s)</u>	<u>Name of each exchange on which registered</u>
Common Stock	VYNT	The Nasdaq Capital Market

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

On May 17, 2021, Vyant Bio, Inc. (the "Registrant") issued a press release regarding financial results for the fiscal quarter ended March 31, 2021. A copy of the press release is furnished as Exhibit 99.1 hereto and is incorporated by reference herein.

**Forward-Looking Statements**

This report, including Exhibit 99.1 furnished herewith, contains forward-looking statements within the meaning of the federal securities laws. Forward-looking statements typically are identified by use of terms such as "may," "will," "should," "plan," "expect," "anticipate," "estimate" and similar words, and the opposites of such words, although some forward-looking statements are expressed differently. Forward-looking statements involve known and unknown risks and uncertainties that exist in the Registrant's operations and business environment, which may be beyond the Registrant's control, and which may cause actual results, performance or achievements to be materially different from future results, performance or achievements expressed or implied by such forward-looking statements. All statements other than statements of historical fact are statements that could be forward-looking statements. For example, forward-looking statements include, without limitation: statements regarding prospects for additional customers; market forecasts; projections of earnings, revenues, synergies, accretion or other financial information; and plans, strategies and objectives of management for future operations or transactions. The risks and uncertainties referred to above include, but are not limited to, risks detailed from time to time in the Registrant's filings with the Securities and Exchange Commission, including its Annual Report on Form 10-K for the year ended December 31, 2020 and any subsequent filings. These risks could cause actual results to differ materially from those expressed in any forward-looking statements made by, or on behalf of, the Registrant. Forward-looking statements represent the judgment of management of the Registrant regarding future events. Although the Registrant believes that the expectations reflected in such forward-looking statements are reasonable at the time that they are made, the Registrant can give no assurance that such expectations will prove to be correct. Unless otherwise required by applicable law, the Registrant assumes no obligation to update any forward-looking statements, and expressly disclaims any obligation to do so, whether as a result of new

information, future events or otherwise.

**Item 9.01 Financial Statements and Exhibits.**

(d) Exhibits

As described above, the following exhibits are furnished as part of this report:

[Exhibit 99.1 — Press release, dated May 17, 2021.](#)

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**SIGNATURE**

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.

**VYANT BIO, INC.**

By: /s/ John A. Roberts

Name: John A. Roberts

Title: President and Chief Executive Officer

Date: May 17, 2021

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## Vyant Bio Reports First Quarter 2021 Results and Provides Strategic Business Update

Cherry Hill, N.J., May 17, 2021 – Vyant Bio, Inc. (the “Company”) (Nasdaq: VYNT), is an innovative biotechnology company focused on partnering with pharmaceutical and other biotechnology companies to identify novel and repurposed therapeutics through the integration of human-derived biology with data science technologies and IND-enabling expertise.

### RECENT STRATEGIC AND OPERATIONAL HIGHLIGHTS IN Q-1 2021

- Rebranded the Company from Cancer Genetics to Vyant Bio (Nasdaq: VYNT)
- Completed merger with StemoniX, Inc.
- Completed two equity financing rounds to raise \$27.5 million in cash, entered into Q2 with \$33.1 million of cash on the balance sheet
- Investing in research and development to optimize drug discovery capabilities
- Launched commercial stage, novel disease models in Rett Syndrome and CDKL5
- Initially focusing on novel and repurposed drugs to treat Rett Syndrome and CDKL5

Jay Roberts, Chief Executive Officer of Vyant Bio stated, “in Q1 2021, we reached an important milestone with the closing of the merger with StemoniX, uniquely positioning the combined company to focus our business on discovering applications for novel and repurposed therapeutics. We believe that drug discovery needs to progressively evolve as we know the traditional methods and models for predicting safe and effective drugs have under-performed, as evidenced by the billions of dollars and years of time it takes to bring novel drugs to market. With this as a backdrop, we are focusing our business on converging an impactful approach to drug discovery with data science and biology-driven technologies at the core with engineering disciplines and regulatory expertise.”

“Vyant Bio has commercialized the development, engineering and manufacturing of disease models, built on its induced pluripotent stem cell (“iPSC”) technology, and has developed neural and cardiac screening platforms, which are used to screen novel and repurposed compound targets”, stated Ping Yeh, Vyant Bio’s Chief Innovation Officer. “The most mature disease models are being used to find therapeutic candidates in the central nervous system with its microBrain<sup>®</sup>, driven by a focus on Rett Syndrome and CDKL5 neurological disorders. With the addition of the *vivo*Pharm cancer cell-line assets and scientific expertise in oncology, the Company believes it can also advance models targeting Glioblastoma and Parkinson’s disease. The team has also made progress with our microHeart<sup>®</sup> platform, so we believe there will be continued interest from partners with an interest in Cardiac Fibrosis and Rett Syndrome”, Mr. Yeh continued.

“Our human-derived models, combined with the latest data science and software techniques, can identify and rank order repurposed and novel compounds by target. In our current drug discovery efforts, we aim to leverage our iPSC technology to identify drug candidates for licensure or clinical development. We are in active discussions with prospective pharma partners to offer exclusive licenses to certain disease models, and expect to enter into similar license agreements for access to both novel and repurposed therapies. The Company is striving to receive a mix of upfront payments, licensing fees, milestone-based fees and ongoing royalty payments”, Mr. Yeh concluded.

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The Company filed its quarterly report for Q1 2021 on Form 10-Q today with the Securities and Exchange Commission. The Company formerly known as Cancer Genetics, Inc., StemoniX and CGI Acquisition, Inc. (“Merger Sub”) entered into a merger agreement on August 21, 2020, which was amended on February 8, 2021 and February 26, 2021 (as amended, the “Merger Agreement”). Pursuant to the terms of the Merger Agreement, Merger Sub was merged (“the Merger”) with and into StemoniX on March 30, 2021, with StemoniX surviving the Merger as a wholly owned subsidiary of the Company. The Merger was accounted for as a reverse acquisition with StemoniX being the accounting acquirer of CGI using the acquisition method of accounting.

### FIRST QUARTER 2021 FINANCIAL RESULTS

As StemoniX was deemed to have acquired CGI for accounting purposes and the Merger closed on March 30, 2021, the Company’s first quarter financial results are primarily the StemoniX operations.

Cash and cash equivalents totaled approximately \$33.1 million as of March 31, 2021.

Total revenues increased 32%, or \$54 thousand, to \$222 thousand for the three months ended March 31, 2021, as compared with \$168 thousand for the three months ended March 31, 2020.

Cost of goods sold – service aggregated \$89 thousand and \$132 thousand, respectively, for the three months ended March 31, 2021 and 2020, resulting in a cost of goods sold of 77% and 97%, respectively, of service revenues.

Cost of goods sold – product aggregated \$396 thousand and \$166 thousand for the three months ended March 31, 2021 and 2020, respectively, resulting in a cost of goods sold gross margin deficit of \$290 thousand and \$134 thousand. Our product manufacturing capabilities currently have excess capacity to support future growth.

Research and development expenses decreased by 19%, or \$189 thousand to \$820 thousand for the three months ended March 31, 2021 from \$1.0 million for the three months ended March 31, 2020.

Selling, general and administrative expenses increased by 46%, or \$383 thousand, to \$1.2 million for the three months ended March 31, 2021, as compared with \$833 thousand for the three months ended March 31, 2020.

Merger related costs for the three-month period ended March 31, 2021 were \$2.1 million.

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Total other expense for the three months ended March 31, 2021 was \$2.9 million, which consisted of a number of non-recurring non-cash items related to the conversion of the StemoniX’s capital structure to StemoniX common stock and exchange for Company common stock.

### ABOUT VYANT BIO, INC.

Vyant Bio, Inc. (“Vyant Bio”) (Nasdaq: VYNT) is emerging as an advanced biotechnology drug discovery company. With capabilities in data, science (both biology and chemistry), engineering, and regulatory, we are rapidly identifying small and large molecule therapeutics and derisking decision making through multiple in silico, in vitro, and in vivo modalities. Leveraging these modalities, Vyant Bio is able to capitalize on repurposed and novel compounds and then partner with others to further develop and commercialize valuable therapeutics and new treatments for patients. Vyant Bio operates two wholly-owned subsidiaries, StemoniX and *vivo*Pharm. Formerly known as Cancer Genetics, Inc., the company’s name was changed to Vyant Bio, Inc. in March 2021. Vyant Bio is headquartered in the US, with offices in Europe, and Australia.

StemoniX is empowering the discovery of new medicines through the convergence of novel human biology and software technologies. StemoniX develops and manufactures high-density, at-scale human induced pluripotent stem (iPS) cell-derived neural and cardiac screening platforms for drug discovery and development. Predictive, accurate, and

consistent, these human models enable scientists to quickly and economically conduct research with improved outcomes in a simplified workflow. Through collaborations with drug discovery organizations, StemoniX tests compounds in-house, creates new cell-based disease models, and operationalizes custom human iPSC-derived disease models at large scale for high-throughput screening. With leading-edge iPSC technologies and data science, StemoniX is helping global institutions bring the most promising medicines to patients.

*vivo*Pharm offers proprietary preclinical test systems supporting clinical diagnostic offerings at early stages valued by the pharmaceutical industry, biotechnology companies, and academic research centers. *vivo*Pharm is focused on precision and translational medicine to drive drug discovery and novel therapies. *vivo*Pharm specializes in conducting studies tailored to guide drug development, starting from compound libraries and ending with a comprehensive set of *in vitro* and *in vivo* data and reports, as needed for Investigational New Drug filings. *vivo*Pharm operates in The Association for Assessment and Accreditation of Laboratory Animal Care International (AAALAC) accredited and GLP compliant audited facilities.

For more information, please visit or follow Vyant Bio at:

Internet: [www.vyantbio.com](http://www.vyantbio.com)

LinkedIn: <https://www.linkedin.com/company/vyant-bio>

Twitter: @VyantBio

#### Forward Looking Statements:

This press release contains forward-looking statements within the meaning of the Private Securities Litigation Reform Act of 1995. All statements pertaining to Vyant Bio, Inc.'s (formerly Cancer Genetics, Inc.) expectations regarding future financial and/or operating results, our ability to discover new and repurposed drug candidates, and potential for our services, future revenues or growth, or the potential for future collaborations or partnership transactions in this press release constitute forward-looking statements.

Any statements that are not historical fact (including, but not limited to, statements that contain words such as “will,” “believes,” “plans,” “anticipates,” “expects,” and “estimates”) should also be considered to be forward-looking statements. Forward-looking statements involve risks and uncertainties, including, without limitation, risks inherent in our attempts to adapt to the global coronavirus pandemic, discover drug candidates, partner with pharmaceutical and other biotechnology companies, achieve profitability and increase sales of our services, maintain our existing customer base and avoid cancellation of customer contracts or discontinuance of trials, raise capital to meet our liquidity needs, realize the anticipated benefits of the merger with StemoniX, Inc., and other risks discussed in the Vyant Bio, Inc. Form 10-K for the year ended December 31, 2020, and Form 10-Q for the quarter ended March 31, 2021, along with other filings with the Securities and Exchange Commission. These forward-looking statements speak only as of the date hereof. Vyant Bio disclaims any obligation to update these forward-looking statements.

#### Investor Contacts:

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Investor Relations

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**Vyant Bio, Inc.**  
**(Formerly Known as Cancer Genetics, Inc.)**  
**Consolidated Statements of Operations**  
**(Shares and USD in Thousands)**

	<b>Three months ended March 31,</b>	
	<b>2021</b>	<b>2020</b>
<b>Revenues:</b>		
Service	\$ 116	\$ 136
Product	106	32
Total revenues	<u>222</u>	<u>168</u>
<b>Operating costs and expenses:</b>		
Cost of goods sold – service	89	132
Cost of goods sold – product	396	166
Research and development	820	1,009
Selling, general and administrative	1,216	833
Merger related costs	2,145	-
Total operating costs and expenses	<u>4,666</u>	<u>2,140</u>
Loss from operations	<u>(4,444)</u>	<u>(1,972)</u>
<b>Other (expense) income:</b>		
Change in fair value of warrant liability	214	-
Change in fair value of share-settlement obligation derivative	(250)	-
Loss on debt conversions	(2,518)	-
Interest expense	(368)	(1)
Total other (expense) income	<u>(2,922)</u>	<u>(1)</u>
<b>Loss before income taxes</b>	<u>(7,366)</u>	<u>(1,973)</u>
Income tax expense (benefit)	-	-
<b>Net loss</b>	<u>\$ (7,366)</u>	<u>\$ (1,973)</u>
<b>Net loss per common share:</b>		
Net loss per share attributable to common stock - Basic and Diluted	<u>\$ (2.31)</u>	<u>\$ (0.80)</u>
<b>Weighted average shares outstanding:</b>		
Weighted average common shares outstanding - Basic and Diluted	<u>3,184</u>	<u>2,460</u>

**Vyant Bio, Inc.**  
**(Formerly Known as Cancer Genetics, Inc.)**  
**Consolidated Balance Sheets**  
**(unaudited)**  
**(Shares and USD in Thousands)**

	<u>March 31, 2021</u>	<u>December 31, 2020</u>
<b>Assets</b>		
Current assets:		
Cash and cash equivalents	\$ 33,074	\$ 792
Trade accounts and other receivables	924	357
Inventory	409	415
Prepaid expenses and other current assets	2,134	223
<b>Total current assets</b>	<b>36,541</b>	<b>1,787</b>
Non-current assets		
Goodwill	22,164	-
Intangible assets, net	9,500	-
Fixed assets, net	1,347	1,031
Right-to-use assets, net	1,170	1,095
Long-term prepaid expenses and other assets	1,633	136
<b>Total non-current assets</b>	<b>35,814</b>	<b>2,262</b>
<b>Total assets</b>	<b>\$ 72,355</b>	<b>\$ 4,049</b>
<b>Liabilities, Temporary Equity and Stockholders' Equity (Deficit)</b>		
Current liabilities:		
Accounts payable	\$ 1,412	\$ 1,300
Accrued expenses	3,400	162
Deferred revenue	1,346	92
Income taxes payable	360	-
Obligations under operating leases, current portion	647	486
Obligations under finance leases, current portion	31	-
Other current liabilities	4	9
Other current liabilities - discontinued operations	588	-
<b>Total current liabilities</b>	<b>7,788</b>	<b>2,049</b>
Obligations under operating leases, less current portion	548	627
Obligations under finance leases, less current portion	74	-
Share-settlement obligation derivative	-	1,690
Accrued interest	-	277
Long-term debt	57	6,839
<b>Total liabilities</b>	<b>\$ 8,467</b>	<b>\$ 11,482</b>
<b>Commitments and contingencies</b>		
<b>Temporary equity:</b>		
Series A Convertible Preferred stock, \$0.0001 par value; 4,700 shares authorized, 0 and 4,612 shares issued and outstanding as of March 31, 2021 and December 31, 2020, respectively (liquidation value of \$0 and \$11,732, respectively, as of March 31, 2021 and December 31, 2020)	-	12,356
Series B Convertible Preferred stock, \$0.0001 par value; 4,700 shares authorized, 0 and 3,489 shares issued and outstanding, as of March 31, 2021 and December 31, 2020, respectively (liquidation value of \$0 and \$15,707, respectively, as of March 31, 2021 and December 31, 2020)	-	16,651
Series C Convertible Preferred stock, \$0.0001 par value; 2,000,000 shares authorized, 0 shares issued and outstanding as of March 31, 2021 and December 31, 2020 (liquidation value of \$0 as of March 31, 2021 and December 31, 2020)	-	-
<b>Total temporary equity</b>	<b>-</b>	<b>29,007</b>
<b>Stockholders' equity (deficit):</b>		
Preferred stock, authorized 9,764 shares \$0.0001 par value, none issued	-	-
Common stock, authorized 100,000 shares, \$0.0001 par value, 28,985 and 2,594 shares issued and outstanding as of March 31, 2021 and December 31, 2020, respectively	3	-
Additional paid-in capital	109,205	1,514
Accumulated deficit	(45,320)	(37,954)
<b>Total Stockholders' equity (deficit)</b>	<b>63,888</b>	<b>(36,440)</b>
<b>Total liabilities and Stockholders' equity</b>	<b>\$ 72,355</b>	<b>\$ 4,049</b>