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**UNITED STATES  
SECURITIES AND EXCHANGE COMMISSION  
WASHINGTON, D.C. 20549**

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**FORM 8-K**

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**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 28, 2022**

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**MIND MEDICINE (MINDMED) INC.**

*(Exact name of Registrant as Specified in Its Charter)*

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**British Columbia, Canada**  
(State or Other Jurisdiction  
of Incorporation)

**001-40360**  
(Commission File Number)

**98-1582438**  
(IRS Employer  
Identification No.)

**One World Trade Center, Suite 8500**  
New York, New York  
(Address of Principal Executive Offices)

**10007**  
(Zip Code)

**Registrant's Telephone Number, Including Area Code: (650) 208-2454**

**Not Applicable**

*(Former Name or Former Address, if Changed Since Last Report)*

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

| Title of each class       | Trading Symbol(s) | Name of each exchange on which registered |
|---------------------------|-------------------|---|
| Subordinate Voting Shares | MNMD              | The Nasdaq Stock Market LLC               |

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))

Indicate by check mark whether the registrant is an emerging growth company 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.

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

On March 28, 2022, Mind Medicine (MindMed) Inc. (the “Company”) issued a press release announcing its financial results for its fiscal year ended December 31, 2021. A copy of the press release is furnished as Exhibit 99.1 to this Current Report on Form 8-K.

*The information contained in this Item 2.02 and in the accompanying Exhibit 99.1 shall not be 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 Exchange Act or the Securities Act of 1933, as amended, except as shall be expressly set forth by specific reference in such filing.*

## Item 4.01 Changes in Registrant’s Certifying Accountant

### (a) Dismissal of Independent Registered Public Accounting Firm

On March 28, 2022, the Board of Directors of Mind Medicine (MindMed) Inc. (the “Company”) determined not to reappoint its principal accountant from Ernst & Young (Canada) LLP (“EY”) for the fiscal year ending December 31, 2022.

The principal accountant’s report of EY on the consolidated financial statements of the Company as of December 31, 2021 and 2020 and for each of the three years in the period ended December 31, 2021 did not contain any adverse opinion or disclaimer of opinion and were not qualified or modified as to uncertainty, audit scope or accounting principles.

As at December 31, 2021 and 2020 and for each of the three years in the period ended December 31, 2021 through March 28, 2022, there were no disagreements with EY, whether or not resolved, on any matter of accounting principles or practices, financial statement disclosure, or auditing scope or procedure, which if not resolved to EY’s satisfaction, would have caused it to make a reference to the subject matter of the disagreement in connection with any reports it would have issued. As at December 31, 2021 and 2020 and for each of the three years in the period ended December 31, 2021 through March 28, 2022, except as set forth below, there were no reportable events as that term is defined in Item 304(a)(1)(v) of Regulation S-K. In connection with the preparation of our consolidated financial statements as of and for the fiscal year ended December 31, 2021, EY identified an instance of a material weakness in our internal controls over financial reporting in connection with the Company’s accounting for contracts. This reportable event was discussed among the Audit Committee and EY. EY has been authorized by the Company to respond fully to the inquiries of KPMG LLP, the successor independent registered public accounting firm, concerning this reportable event.

A letter from EY addressed to the Securities and Exchange Commission stating that it concurs with the statements made by the Company with respect to EY in this Current Report on Form 8-K. A copy of such letter is furnished hereto with the filing of this Current Report on Form 8-K.

### (b) Appointment of New Independent Registered Public Accounting Firm

On March 28, 2022, the Board of Directors of the Company approved the engagement of KPMG LLP (“KPMG”) as the Company’s independent registered public accounting firm for the year ending December 31, 2022. The Company’s engagement of KPMG is subject to the approval of the Company’s shareholders at the Company’s 2022 annual meeting.

In connection with the Company’s appointment of KPMG as the Company’s independent registered public accounting firm, the Company has not consulted with KPMG on (i) the application of accounting principles to a specified transaction, either completed or proposed; the type of audit opinion that might be rendered on the Company’s financial statements, and neither a written report nor oral advice was provided that KPMG concluded was an important factor considered by the Company in reaching a decision as to the accounting, auditing or financial reporting issue; or (ii) any matter that was either the subject of a disagreement (as defined in Item 304(a)(1)(iv) of Regulation S-K) or a reportable event (as described in Item 304(a)(1)(v) of Regulation S-K).

## Item 9.01 Financial Statements and Exhibits.

| Exhibit No. | Description   |
|-------------|---|
| 16.1        | <a href="#">Letter of Ernst &amp; Young (Canada) LLP, dated March 28, 2022, regarding change in independent registered public accounting firm</a> |
| 99.1        | <a href="#">Press Release, dated March 28, 2022</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 thereunto duly authorized.

Date: March 28, 2022

MIND MEDICINE (MINDMED) INC.

By: /s/ Cynthia Hu  
Name: Cynthia Hu  
Title: Chief Legal Officer & Secretary





Ernst & Young LLP  
EY Tower  
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Toronto, ON M5H 0B3

Tel: +1 416 864 1234  
Fax: +1 416 864 1174  
ey.com

March 28, 2022

Securities and Exchange Commission 100 F Street, N.E.  
Washington, DC 20549

Ladies and Gentlemen:

We have read Item 4.01 of Form 8-K dated March 28, 2022, of Mind Medicine (MindMed) Inc. and are in agreement with the statements contained in section (a) therein. We have no basis to agree or disagree with other statements of the registrant contained therein.

Regarding the registrant's statement concerning the lack of internal control to prepare consolidated financial statements, included in Item 4.01 (a) therein, we had considered such matter in determining the nature, timing and extent of procedures performed in our audit of the registrant's 2021 consolidated financial statements.

Sincerely,

/s/ Ernst & Young LLP

Chartered Professional Accountants Licensed Public Accountants

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**MindMed Reports Full Year 2021 Financial Results and Business Highlights**

- FDA cleared MindMed's Investigational New Drug (IND) application for Phase 2b dose optimization trial of MM-120 -
- Progressed development programs for all three lead product candidates -
- Cash balance of \$133.5 million at year end 2021 -
- Company to host earnings conference call today at 8:30 AM EDT -

NEW YORK, March 28, 2022 -- **Mind Medicine (MindMed) Inc.** (NASDAQ: MNMD), (NEO: MMED) (the "Company" or "MindMed"), a clinical stage biopharmaceutical company developing novel products to treat brain health disorders, today reported its financial results for the full-year ended December 31, 2021.

"2021 was a year of major advancements across all aspects of MindMed, with significant growth in our organization, development programs and research collaborations. We established a regulatory pathway for MM-120 in the treatment of GAD and for MM-402 launched a first of its kind program to develop a novel treatment for core symptoms of autism spectrum disorder – both of which represent a meaningful leap forward in the field of psychiatry," said Robert Barrow, Chief Executive Officer and Director of MindMed. "We expect 2022 to be a transformational year in which we continue to drive substantial growth across our pharmaceutical and digital medicine pipelines. I am incredibly proud of our team's achievements and I am more confident than ever in our ability to continue advancing our organization and development programs. We are keenly focused on our mission to deliver novel therapies to treat brain health disorders leading to meaningful improvements in patient outcomes in these major areas of unmet medical need."

**Recent Highlights and Anticipated Upcoming Milestones**

*MM-120 (LSD D-tartrate)*: a proprietary, pharmaceutically optimized form of lysergic acid diethylamide (LSD) that is being developed for the treatment of generalized anxiety disorder (GAD). MM-120 is also being studied under various dosing regimens for the treatment of adult attention deficit hyperactivity disorder (ADHD) and for the treatment of chronic pain.

- In January 2022, the U.S. Food and Drug Administration (FDA) cleared the Company's Investigational New Drug (IND) application for the Phase 2b dose-optimization trial of MM-120 for the treatment of GAD.
    - Study MMED008, a Phase 2b dose-optimization study of MM-120 for the treatment of GAD is expected to begin in Q2 2022.
  - In December 2021, Study MMED007, a Phase 2a proof-of-concept study was initiated for the treatment of ADHD. The study is designed to assess the safety and efficacy of repeated low-dose MM-120 administration.
  - A clinical study of MM-120 in a chronic pain condition is expected to be initiated in Q4 2022.
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*MM-110 (zolunicant HCl or 18-MC)*: a derivative of ibogaine that the Company is developing for the treatment of opioid withdrawal. MM-110 is an  $\alpha 3\beta 4$  nicotinic cholinergic receptor antagonist that has been extensively tested in preclinical models of substance use disorders.

- In January 2022, the USAN Council assigned the non-proprietary name “zolunicant” (pronounced: zoe lun’ i kant), to MM-110 or 18-MC.
- In December 2021, the Company completed a Phase 1 study of MM-110 with topline data expected in 2022. These results will inform the design of the planned Phase 2a study in individuals undergoing standardized supervised opioid withdrawal, which is expected to commence in early 2022. The Phase 2a study will evaluate the safety, tolerability and efficacy of MM-110 in mitigating symptoms of opioid withdrawal and facilitating completion of detoxification.

*MM-402 (R(-)-MDMA)*: a synthetic enantiomer of MDMA that exhibits prosocial and empathogenic activity that the Company is developing for the treatment of core symptoms of autism spectrum disorder. Preclinical studies of R(-)-MDMA demonstrate its acute prosocial effects, while its diminished dopaminergic activity suggests that it may exhibit a favorable safety, tolerability and abuse liability profile when compared to racemic MDMA or the S(+)-enantiomer of MDMA.

- IND-enabling studies are currently ongoing, and through the Company’s collaboration with University Hospital Basel, a comparative Phase 1 pharmacokinetic/pharmacodynamic study of R(-), S(+) and  $\pm$  MDMA is expected to commence in mid 2022.

### **Digital Medicine Initiatives**

- In February 2021, the Company completed the acquisition of HealthMode and fully integrated its team to enable rapid progression of digital medicine and business operations functions.
- Having engaged in a productive Pre-Submission meeting with FDA in late 2021, in January 2022, the first subjects were enrolled into the Session Monitoring System (SMS-01) study evaluating the passive collection of sensory data during a consciousness-altering therapeutic session using the MindMed Session Monitoring System (MSMS).
- Anxiety Digital Diagnoses for Precision Psychiatry (ADDAPT, MMED-D001): A Natural History Study and our newly developed mobile application to support the study is expected to launch in private beta in early 2022
- In September 2021 the first participants were enrolled by invitation in the Quantifying the Processes and Events of Psychotherapy at Scale (MM061302) study.

### **Collaborations and Partnerships**

- University Hospital Basel (UHB): The Company continued to support the ongoing collaboration with the Liechti Lab at UHB in Switzerland. MindMed has exclusive worldwide rights to data, compounds and patent rights associated with the Liechti laboratory’s research with LSD and other psychedelic compounds, including data from preclinical studies and over 15 completed and 9 ongoing clinical trials.
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# MindMed

oIn March 2022, the peer-reviewed publication of a double-blind placebo-controlled comparative study of LSD and psilocybin was published in *Neuropsychopharmacology*. The study demonstrated that the key differences between LSD and psilocybin are dose-dependent rather than substance-dependent. These findings have the potential to assist with dose-finding, trial design and inform future studies evaluating the therapeutic utility of psychedelics.

oIn November 2021, the peer-reviewed publication of a randomized, double-blind, placebo-controlled study evaluating the interacting effects of an SSRI and psilocybin in healthy volunteers was published in *Clinical Pharmacology and Therapeutics*. The study suggests psilocybin is safe to take in combination with an SSRI.

▪Nextage Therapeutics Ltd: The Company entered into a collaboration with Nextage Therapeutics in April 2021 to explore the therapeutic potential of noribogine in a proprietary brain targeted liposome drug delivery technology system to mitigate risks of peripheral adverse effects.

▪MindShift Compounds AG: the Company continued to progress its collaborative research and development activities with MindShift Compounds AG in Basel, Switzerland, focusing on the discovery and optimization of next generation compounds, including those with and without acute perceptual effects.

▪The Chopra Foundation: the company ultimately did not reach a definitive agreement for a potential collaboration and discontinued the engagement.

## Director & Officer Appointments

▪In December 2021, [Robert Barrow](#) was appointed as Chief Executive Officer and as a member of the Board of Directors. Mr. Barrow previously served as interim Chief Executive Officer and Chief Development Officer of MindMed and brings strategic expertise and deep industry insight to his role.

▪In December 2021, [Cynthia Hu, JD](#) was appointed as Chief Legal Officer and Corporate Secretary.

▪In November 2021, [Carrie Liao, CPA](#) was appointed as VP, Corporate Controller and Principal Accounting Officer.

▪In September 2021, [Carol Vallone](#) and [Andreas Krebs](#) were appointed to the Board of Directors and were subsequently appointed as chair and vice chair, respectively.

▪In May 2021, [Sarah Vinson, MD](#) was appointed to the Board of Directors.

▪In February 2021, [Daniel Karlin, MD, MA](#) was appointed as Chief Medical Officer.

## Scientific Advisory Board Appointments

•Over the course of 2021, we made significant additions to our Scientific Advisory Board (SAB), including:

•[Robert C. Malenka, MD, PhD](#) (Professor, Stanford University) – SAB chair

•[Maurizio Fava, MD](#) (Psychiatrist-In-Chief, Mass General Hospital; Associate Dean and Professor of Psychiatry, Harvard Medical School)

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# MindMed

- Peter Bergethon, MD (President, Symmetry Research; formerly Vice President, Biogen)
- Robert Dworkin, PhD (Professor, Rochester University; Director, ACTION public-private partnership)
- Bryan Roth, MD, PhD (Professor, University of North Carolina; Director, NIMH Psychoactive Drug Screening Program)
- Maria Oquendo, MD, PhD (Chair and Professor of Psychiatry, University of Pennsylvania; Past President, American Psychiatric Association)

## 2021 Financial Results

*Cash Balance.* As of December 31, 2021, MindMed had cash totaling \$133.5 million compared to \$80.1 million as of December 31, 2020. MindMed believes its available cash and cash equivalents will be sufficient to meet its operating requirements beyond its key development milestones in 2023 and into 2024.

*Net Cash in Operating Activities.* The net cash used in operating activities was \$45.8 million for the year ended December 31, 2021, compared to \$23.6 million for the same period in 2020.

*Research and Development (R&D).* R&D expenses were \$34.8 million for the year ended December 31, 2021, compared to \$18.6 million for the year ended 2020. The increase was primarily due to an increase of \$2.7 million in expenses related to our MM-120 clinical research, \$2.3 million of expense related to our MM-110 clinical research, \$3.5 million in expenses related to preclinical and other research programs, offset by a \$3.5 million decrease of expense in connection with various external R&D collaborations. Internal costs increased \$11.1 million primarily related to an increase non-cash expenses of \$6.6 million of stock-based compensation expenses and \$2.6 million of amortization of our developed technology.

*General and Administrative (G&A).* G&A expenses were \$59.1 million for the year ended December 31, 2021, compared to \$14.4 million for the year ended 2020. The increase was primarily due to an increase of \$28.9 million in non-cash stock-based compensation expenses of which \$21.9 million related to the modification of stock options and RSUs. Other contributors to the increase included higher professional services including accounting, audit, legal, compliance, director and officer insurance, and investor and public relations and personnel costs to support the growth of the company.

*Net Loss.* The net and comprehensive loss for the year ended December 31, 2021 was \$92.3 million, compared to \$33.7 million for the year ended 2020.

## Conference Call and Webcast Reminder

MindMed management will host a conference call at 8:30 AM EDT today to provide a corporate update and review the Company's fiscal year 2021 financial results. Individuals may participate in the call via telephone by dialing (877) 407-0789 (domestic) or (201) 689-8562 (international) and using conference

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ID 13728028. The webcast can be accessed live [here](#) or on MindMed's [Investor Resources](#) webpage. The webcast will be archived on the Company's website for at least 30 days after the conference call.

### **About MindMed**

MindMed is a clinical-stage biopharmaceutical company developing novel products to treat brain health disorders, with a particular focus on psychiatry, addiction, pain and neurology. Our mission is to be the global leader in the development and delivery of treatments that unlock new opportunities to improve patient outcomes. We are developing a pipeline of innovative drug candidates, with and without acute perceptual effects, targeting the serotonin, dopamine and acetylcholine systems.

MindMed trades on NASDAQ under the symbol MNMD and on the Canadian NEO Exchange under the symbol MMED.

### **Forward-Looking Statements**

Certain statements in this news release related to the Company constitute "forward-looking information" within the meaning of applicable securities laws and are prospective in nature. Forward-looking information is not based on historical facts, but rather on current expectations and projections about future events and are therefore subject to risks and uncertainties which could cause actual results to differ materially from the future results expressed or implied by the forward-looking statements. These statements generally can be identified by the use of forward-looking words such as "will", "may", "should", "could", "intend", "estimate", "plan", "anticipate", "expect", "believe", "potential" or "continue", or the negative thereof or similar variations. Forward-looking information in this news release include, but are not limited to, statements regarding anticipated upcoming milestones and studies, results and timing of clinical studies, expected growth and developments of drugs and technologies, continuing collaborations and partnerships, and the availability of cash and cash equivalents. There are numerous risks and uncertainties that could cause actual results and the Company's plans and objectives to differ materially from those expressed in the forward-looking information, including history of negative cash flows; limited operating history; incurrence of future losses; availability of additional capital; lack of product revenue; compliance with laws and regulations; difficulty associated with research and development; risks associated with clinical trials or studies; heightened regulatory scrutiny; early stage product development; clinical trial risks; regulatory approval processes; novelty of the psychedelic inspired medicines industry; as well as those risk factors discussed or referred to herein and the risks described under the headings "Risk Factors" in the Company's filings with the securities regulatory authorities in all provinces and territories of Canada which are available under the Company's profile on SEDAR at [www.sedar.com](http://www.sedar.com) and with the U.S. Securities and Exchange Commission on EDGAR at [www.sec.gov](http://www.sec.gov).

For Media: [media@mindmed.co](mailto:media@mindmed.co)

For Investors: [ir@mindmed.co](mailto:ir@mindmed.co)

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**Mind Medicine (MindMed) Inc.**  
**Consolidated Statements of Operations and Comprehensive Loss**  
(In thousands, except share and per share amounts)

|  | For the Year<br>Ended<br>December 31, 2021 | For the Year<br>Ended<br>December 31, 2020 | For the Period from<br>May 30, 2019 (Date of<br>Incorporation) to<br>December 31, 2019 |
|--|--|--|--|
| Operating expenses:  |  |  |  |
| Research and development                                   | \$ 34,789                                  | \$ 18,631                                  | \$ 7,549   |
| General and administrative                                 | 59,065                                     | 14,399                                     | 3,178  |
| Total operating expenses                                   | 93,854                                     | 33,030                                     | 10,727   |
| Loss from operations                                       | (93,854)                                   | (33,030)                                   | (10,727)   |
| Other income (expense):                                    |  |  |  |
| Interest expense, net                                      | (359)                                      | (164)                                      | 10   |
| Foreign exchange (loss) gain, net                          | (86)                                       | 130  | 18   |
| Other income   | 106  | —  | —  |
| Loss on revaluation of derivative liability                | —  | (873)                                      | —  |
| Total other expense, net                                   | (339)                                      | (907)                                      | 28   |
| Loss before income taxes                                   | (94,193)                                   | (33,937)                                   | (10,699)   |
| Income tax benefit   | (1,157)                                    | —  | —  |
| Net loss   | (93,036)                                   | (33,937)                                   | (10,699)   |
| Other comprehensive gain:                                  |  |  |  |
| Gain on foreign currency translation                       | 762  | 284  | —  |
| Comprehensive loss   | \$ (92,274)                                | \$ (33,653)                                | \$ (10,699)  |
| Net loss per common share, basic and diluted               | \$ (0.23)                                  | \$ (0.13)                                  | \$ (0.10)  |
| Weighted-average common shares, basic and diluted (Note 2) | <u>410,656,231</u>                         | <u>266,220,592</u>                         | <u>102,763,621</u>   |

**Mind Medicine (MindMed) Inc.**  
**Consolidated Balance Sheets**  
(In thousands, except share amounts)

|   | 2021              | December 31, |  | 2020          |
|---|-------------------|--------------|--|---------------|
| <b>Assets</b>   |                   |              |  |               |
| Current assets:   |                   |              |  |               |
| Cash  | \$ 133,539        | \$           |  | 80,094        |
| Prepaid and other current assets  | 3,676             |              |  | 1,425         |
| Total current assets  | 137,215           |              |  | 81,519        |
| Goodwill  | 19,918            |              |  | —             |
| Intangible assets, net  | 6,869             |              |  | —             |
| Total assets  | <u>\$ 164,002</u> | <u>\$</u>    |  | <u>81,519</u> |
| <b>Liabilities and Shareholders' Equity</b>   |                   |              |  |               |
| Current liabilities:  |                   |              |  |               |
| Accounts payable  | \$ 4,178          | \$           |  | 2,022         |
| Accrued expenses  | 6,230             |              |  | 986           |
| Total current liabilities   | 10,408            |              |  | 3,008         |
| Contribution payable  | 1,930             |              |  | 2,643         |
| Total liabilities   | 12,338            |              |  | 5,651         |
| Commitments and contingencies (Note 11)   |                   |              |  |               |
| Shareholders' Equity:   |                   |              |  |               |
| Subordinate voting shares, no par value, unlimited authorized as of December 31, 2021 and 2020; 421,444,157 and 306,135,160 issued and outstanding as of December 31, 2021 and 2020, respectively | —                 |              |  | —             |
| Multiple voting shares, no par value, unlimited authorized as of December 31, 2021 and 2020; 4,521 and 550,000 issued and outstanding as of December 31, 2021 and 2020, respectively              | —                 |              |  | —             |
| Additional paid-in capital  | 288,290           |              |  | 120,220       |
| Accumulated other comprehensive income  | 1,046             |              |  | 284           |
| Accumulated deficit   | (137,672 )        |              |  | (44,636 )     |
| Total shareholders' equity  | 151,664           |              |  | 75,868        |
| Total liabilities and shareholders' equity  | <u>\$ 164,002</u> | <u>\$</u>    |  | <u>81,519</u> |

