

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

FORM 10-Q

(Mark One)

QUARTERLY REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the quarterly period ended **February 28, 2022**

or

TRANSITION REPORT UNDER SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the transition period from [] to []

Commission file number

LEXARIA BIOSCIENCE CORP.

(Exact name of registrant as specified in its charter)

Nevada

State or other jurisdiction of
incorporation or organization

20-2000871

(I.R.S. Employer
Identification No.)

#100 – 740 McCurdy Road, Kelowna BC Canada

(Address of principal executive offices)

V1X 2P7

(Zip Code)

Registrant's Telephone number, including area code: 1.250.765.6424

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

Title of Class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, Par Value \$0.001	LEXX	NASDAQ
Warrants	LEXXW	NASDAQ

Indicate by check mark whether the registrant: (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports) and (2) has been subject to such filing requirements for the last 90 days. Yes No

Indicate by check mark whether the registrant has submitted electronically, every Interactive Data File required to be submitted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit files). Yes No

Indicate by check mark whether the registrant is a large accelerated filer, an accelerated filer, a non-accelerated filer, or a smaller reporting company. See definition of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act.

Large accelerated filer
Non-accelerated Filer

Accelerated filer
Smaller reporting company
Emerging growth company

If an emerging growth company, indicate by a 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

Indicate by check mark whether the registrant is a shell company (as defined in Rule 12b-2 of the Exchange Act). Yes No

Indicate the number of shares outstanding of each of the registrant's classes of common stock as of the latest practicable date.

5,950,998 common shares as of April 11, 2022

DOCUMENTS INCORPORATED BY REFERENCE
None.

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PART I—FINANCIAL INFORMATION

Item 1. Financial Statements

LEXARIA BIOSCIENCE CORP.
CONSOLIDATED BALANCE SHEET
(Expressed in U.S. Dollars)

	<u>February 28,</u> <u>2022</u>	<u>August 31,</u> <u>2021</u>
	(Unaudited)	(Audited)
ASSETS		
Current		
Cash	\$ 8,359,374	\$ 10,917,797
Marketable securities	552,317	833,841
Accounts receivable	506,622	342,401
Inventory	31,927	29,648
Prepaid expenses and deposit	1,121,961	319,253
Total Current Assets	<u>10,572,202</u>	<u>12,442,940</u>
Non-current assets, net		
Lease right of use	72,099	91,041
Intellectual property	411,452	364,623
Property & equipment	358,811	368,213
Total Non-current Assets	<u>842,362</u>	<u>823,877</u>
TOTAL ASSETS	<u>\$ 11,414,564</u>	<u>\$ 13,266,817</u>
LIABILITIES		
Current		
Accounts payable and accrued liabilities	\$ 116,592	\$ 100,723
Due to a related party	-	5,223
Loan payable	7,875	7,926
Lease payable	41,076	39,404
Total Current Liabilities	<u>165,543</u>	<u>153,276</u>
Long Term		
Lease payable	29,080	49,989
Total Long Term Liabilities	<u>29,080</u>	<u>49,989</u>
TOTAL LIABILITIES	<u>194,623</u>	<u>203,265</u>
STOCKHOLDERS' EQUITY		
Share Capital		
Authorized:		
220,000,000 common voting shares with a par value of \$0.001 per share Issued and outstanding: 5,950,998 common shares at February 28, 2022 and 5,726,699 common shares at August 31, 2021		
	5,951	5,727
Additional paid-in capital	46,697,434	45,089,114
Deficit	<u>(35,248,137)</u>	<u>(31,829,204)</u>
Equity attributable to shareholders of the Company	11,455,248	13,265,637
Non-controlling Interest	<u>(235,307)</u>	<u>(202,085)</u>
Total Stockholders' Equity	<u>11,219,941</u>	<u>13,063,552</u>
TOTAL LIABILITIES AND STOCKHOLDERS' EQUITY	<u>\$ 11,414,564</u>	<u>\$ 13,266,817</u>

The accompanying notes are an integral part of these condensed consolidated interim financial statements.



LEXARIA BIOSCIENCE CORP.
CONSOLIDATED STATEMENTS OF OPERATIONS AND COMPREHENSIVE LOSS
(Expressed in U.S. Dollars, except number of shares)

	THREE MONTHS ENDED		SIX MONTHS ENDED	
	February 28		February 28	
	2022	2021	2022	2021
	(Unaudited)		(Unaudited)	
Revenue	\$ 30,650	\$ 192,006	\$ 44,530	\$ 487,662
Cost of goods sold	6,387	30,570	11,957	95,048
Gross profit	24,263	161,436	32,573	392,614
Expenses				
Research and development	275,686	176,398	734,395	368,659
General and administrative	1,197,250	1,089,552	2,750,333	1,841,590
Total operating expenses	1,472,936	1,265,950	3,484,728	2,210,249
Loss from operations	(1,448,673)	(1,104,514)	(3,452,155)	(1,817,635)
Gain on disposal of assets		1,522,704	-	1,522,704
Discontinued operations	-	(25,000)	-	(22,000)
Net and comprehensive income (loss) for the period	\$ (1,448,673)	\$ 393,190	\$ (3,452,155)	\$ (316,931)
Net and comprehensive income (loss) attributable to:				
Common shareholders	\$ (1,425,776)	404,111	\$ (3,418,933)	(291,917)
Non-controlling interest	\$ (22,897)	(10,921)	\$ (33,222)	(25,014)
Basic and diluted income (loss) per share				
Continuing operations	\$ (0.25)	\$ 0.10	\$ (0.59)	\$ (0.08)
Discontinued operations	-	(0.01)	-	(0.01)
	\$ (0.25)	\$ 0.09	\$ (0.59)	\$ (0.09)
Weighted average number of common shares outstanding				
- Basic and diluted	5,911,123	4,052,904	5,818,401	3,524,286

The accompanying notes are an integral part of these consolidated interim financial statements.



LEXARIA BIOSCIENCE CORP.
CONSOLIDATED STATEMENT OF CASH FLOWS
(Expressed in U.S. Dollars)

	SIX MONTHS ENDED	
	February 28,	
	2022	2021
	(Unaudited)	
Cash flows used in operating activities		
Net loss and comprehensive loss	\$ (3,452,154)	\$ (294,931)
Adjustments to reconcile net loss to net cash used in operating activities:		
Stock based compensation	408,544	66,041
Depreciation and amortization	54,093	55,859
Inventory write-off	-	2,482
Bad debt expense	-	37,000
Non-cash right of use lease expense	18,942	17,614
Gain on disposal of assets	-	(1,522,704)
Unrealized loss on marketable securities	281,473	16,952
Shares issued for services	300,000	-
Lease accretion	2,954	4,282
Change in working capital		
Accounts receivable	(164,221)	(69,264)
Inventory	(1,161)	(16,883)
Prepaid expenses and deposits	97,292	(181,651)
Accounts payable and accrued liabilities	15,868	59,574
Due to related parties	(5,223)	148,102
Deferred revenue	-	(43,255)
Net cash used in by operating activities	\$ (2,443,593)	\$ (1,720,782)
Cash flows used in investing activities		
Disposal (acquisition) of assets	(42,375)	273,375
Intellectual property	(50,263)	(8,766)
Net cash (used in) provided by investing activities	\$ (92,638)	\$ 264,609
Cash flows from financing activities		
Repayment of loan payable	-	(23,163)
Lease payments	(22,191)	(21,975)
Proceeds from issuance of equity	-	9,471,495
Net cash provided by (used in) financing Activities	\$ (22,191)	\$ 9,426,357
Net cash provided by discontinued operations	\$ -	\$ 83,000
Net change in cash for the period	(2,558,422)	8,053,184
Cash at beginning of period	10,917,797	1,293,749
Cash at end of period	\$ 8,359,374	\$ 9,346,933
Supplemental information of cash flows:		
Income taxes paid in cash	\$ -	\$ 3,450
Non-cash consideration on asset disposal	\$ -	\$ 1,171,599
Non-cash shares for services included in prepaid expenses	\$ 900,000	\$ -

The accompanying notes are an integral part of these condensed consolidated interim financial statements.



LEXARIA BIOSCIENCE CORP.
CONSOLIDATED STATEMENTS OF STOCKHOLDERS' EQUITY
(Expressed in U.S. Dollars)

	SHARE CAPITAL		ADDITIONAL PAID-IN CAPITAL	DEFICIT	NCI	TOTAL STOCKHOLDERS' EQUITY
	SHARES	AMOUNT				
		\$	\$	\$	\$	\$
Balance August 2020	3,001,476	3,001	30,324,398	(27,802,198)	(42,943)	2,482,258
Stock based compensation	-	-	48,887	-	-	48,887
Net loss	-	-	-	(696,028)	-	(696,028)
Non-controlling interest	-	-	-	-	(14,093)	(14,093)
Balance November 30, 2020	3,001,476	3,001	30,373,285	(28,498,226)	(57,036)	1,821,024
Stock based compensation	-	-	17,154	-	-	17,154
Brokered placement	2,102,856	2,104	9,469,393	-	-	9,471,497
Net Income	-	-	-	404,111	-	404,111
Non-controlling interest	-	-	-	-	(10,921)	(10,921)
Balance February 28, 2021	5,104,332	5,105	39,859,832	(28,094,115)	(67,957)	11,702,865
Stock based compensation	-	-	343,966	-	-	343,966
Warrants issued for services	-	-	785,895	-	-	785,895
Net loss	-	-	-	(2,556,997)	-	(2,556,997)
Non-controlling interest	-	-	-	-	(9,555)	(9,555)
Balance May 31, 2021	5,104,332	5,105	40,989,693	(30,651,112)	(77,512)	10,266,174
Exercise of warrants	610,189	610	4,014,433	-	-	4,015,043
Shares issued for services	12,178	12	84,988	-	-	85,000
Net loss	-	-	-	(1,178,092)	-	(1,178,092)
Non-controlling interest	-	-	-	-	(124,573)	(124,573)
Balance August 31, 2021	5,726,699	5,727	45,089,114	(31,829,204)	(202,085)	13,063,552
Stock based compensation	-	-	408,544	-	-	408,544
Net loss	-	-	-	(1,993,157)	-	(1,993,157)
Non-controlling interest	-	-	-	-	(10,325)	(10,325)
Balance November 30, 2021	5,726,699	5,727	45,497,658	(33,822,361)	(212,410)	11,468,614
Shares issued for services	224,299	224	1,199,776	-	-	1,200,000
Net loss	-	-	-	(1,425,777)	-	(1,425,777)
Non-controlling interest	-	-	-	-	(22,895)	(22,895)
Balance February 28, 2022	5,950,998	5,951	46,697,434	(35,248,138)	(235,305)	11,219,942

The accompanying notes are an integral part of these condensed consolidated interim financial statements.



LEXARIA BIOSCIENCE CORP.
NOTES TO THE CONDENSED INTERIM CONSOLIDATED FINANCIAL STATEMENTS
February 28, 2022
(Expressed in U.S. Dollars)

1. Nature of Business

Lexaria Bioscience Corp. (“Lexaria”, “we”, “our” or the “Company”) is a biotechnology company pursuing the enhancement of the bioavailability of a diverse and broad range of active pharmaceutical ingredients (“APIs”) using our patented drug delivery technology DehydraTECH™. Through continued validation of our research and development our focus is on national and international application for DehydraTECH.

Revenues are primarily derived from licensing fees for the use of the Company’s patented technology to partners who pay either a fee to use DehydraTECH in the manufacturing of their own products or through the purchase of DehydraTECH manufactured products made to their specifications by Lexaria. The Company has relationships with several consumer products companies in the CBD and nutraceuticals spaces that use Lexaria’s technology in consumer goods being sold online and at retailers in the US and Canada.

The Company is headquartered in Kelowna, British Columbia, Canada. The corporate website is www.lexariabioscience.com

Going Concern Analysis

The Company’s consolidated financial statements included herein have been prepared pursuant to the rules and regulations of the Securities and Exchange Commission (“SEC”) and in accordance with accounting principles generally accepted in the United States (“US GAAP”) applicable to a going concern which assumes the Company will have sufficient funds to pay its operational, research and development and capital expenditures for a period of at least 12 months from the date this financial report.

Since inception, the Company has incurred significant operating and net losses. The losses attributable to common shareholders were \$1.2m, \$4.1m and \$4.2m for the years ended August 31, 2021, 2020 and 2019, respectively. As of February 28, 2022, we had an accumulated deficit of \$ 35.25m. We expect to continue to incur significant operational expenses and net losses in the upcoming 12 months. Our net losses may fluctuate significantly from quarter to quarter and year to year, depending on the stage and complexity of our R&D studies and related expenditures, the receipt of additional payments on the licencing of our technology, if any, and the receipt of payments under any current or future collaborations we may enter into.

On January 12, 2021, the Company closed an underwritten public offering for net proceeds of \$9,471,497. In the fourth quarter of the year ended August 31, 2021, the Company received \$4,015,043 from the exercise of warrants. We may offer additional securities for sale during fiscal year 2022 or thereafter in response to market conditions or other circumstances if we believe such a plan of financing is required to advance the Company’s business plans and is in the best interests of our stockholders.

The Company has evaluated whether there are conditions or events, considered in the aggregate, that raise substantial doubt about the Company’s ability to continue as a going concern. As of February 28, 2022, the Company had cash of approximately \$8.4m. We believe this is sufficient to enable the Company to fund its operating and R&D expenses and any capital expenditure requirements through one year from the issuance date of these unaudited consolidated financial statements.



To date, we have obtained cash and funded our operations primarily through equity financings and license agreements. In order to continue the development of our drug candidates, at some point in the future we expect to pursue one or more capital transactions, whether through the sale of equity securities, debt financing, license agreements or entry into strategic partnerships. There can be no assurance that we will be able to continue to raise additional capital in the future.

COVID-19

Impacts of COVID-19 Pandemic

The emergence of the COVID-19 pandemic in 2020 continues to present uncertainty and unforecastable new risks to the Company and its business plans. As of February 28, 2022, there has been no material impact on the Company's financial position as a direct result of the pandemic. However, the Company has experienced some supply chain disruptions and shortages in the timely procurement of ingredients and supplies used in both our R&D activities and production. Management views this situation as transitory but cannot predict the length of time it may take for these disruptions to dissipate or if there will be a significant economic effect on the Company's operations. In the interim, it may cause delays in carrying out our research studies and in our production schedules.

Restrictions on international travel presents a challenge in carrying out normal business activities related to corporate finance efforts and the pursuit of new customers throughout North America who might otherwise access to our licensees' retail products. As a result, the pandemic has increased the risk of lower revenues and higher losses.

During the year ended August 31, 2020, we were in receipt of C\$30,732 in COVID relief under the Canada Emergency Wage Subsidy programs for employees which reduced our employment costs in that year. During fiscal 2020 we also received C\$40,000 from the Canadian Government sponsored Emergency Business Account loan program. As specified by the terms of this program, we have repaid C\$30,000 of the loan in fiscal 2021. The remaining 7,875 (C\$10,000) of the loan payable is anticipated to be forgiven as directed under this program in the year ended August 31, 2023.

We continue to actively monitor the evolving effects of COVID-19 and may take further actions that alter our operations, including those that may be required by federal, state, provincial, or local authorities, or that we determine are in the best interests of our employees and third parties with which we do business. We do not know when it will become practical to relax or eliminate some or all these measures entirely. The economic effect of a prolonged pandemic is difficult to predict and could result in material financial impact in the Company's future reporting periods.

2. Significant Accounting Policies

The significant accounting policies of the Company are consistent with those of our audited financial statements on Form 10-K for the year ended August 31, 2021.



3. Basis of Consolidation

These interim consolidated financial statements include the financial statements of the Company and its wholly owned subsidiaries; Lexaria CanPharm ULC, Lexaria CanPharm Holdings Corp., PoViva Corp., Lexaria Hemp Corp., Kelowna Management Services Corp., and Lexaria Pharmaceutical Corp., and our 83.333% owned subsidiary Lexaria Nicotine LLC (16.667% Altria Ventures Inc., an indirect wholly owned subsidiary of Altria Group, Inc.). All significant intercompany balances and transactions have been eliminated upon consolidation.

4. Basis of Presentation

The Company's unaudited interim consolidated financial statements have been prepared pursuant to the rules and regulations of the Securities and Exchange Commission. Certain information and footnote disclosures normally included in annual financial statements prepared in accordance with United States generally accepted accounting principles (US GAAP) have been condensed or omitted pursuant to such rules and regulations. In the opinion of management, all adjustments considered necessary for a fair presentation have been included. Interim results are not necessarily indicative of results for a full year or any subsequent period.

These unaudited interim consolidated financial statements should be read in conjunction with the audited consolidated annual financial statements and notes thereto included in our annual report filed on Form 10-K for the year ended August 31, 2021.

5. Estimates and Judgements

The preparation of financial statements in conformity with U.S. GAAP requires us to make certain estimates, judgments and assumptions that affect the reported amounts of assets and liabilities, the disclosure of contingent liabilities at the date of the financial statements and the reported amounts of revenue and expenses during the reporting period. Some of the Company's accounting policies require us to make subjective judgments, often as a result of the need to make estimates of matters that are inherently uncertain. These accounting policies involve critical accounting estimates because they are particularly dependent on estimates and assumptions made by management about matters that are highly uncertain at the time the accounting estimates are made. Although we have used our best estimates based on facts and circumstances available to us at the time, different estimates reasonably could have been used. Changes in the accounting estimates used by the Company are reasonably likely to occur from time to time, which may have a material effect on the presentation of financial condition and results of operations.

The Company reviews these estimates, judgments, and assumptions periodically and reflect the effects of revisions in the period in which they are deemed to be necessary. Although we believe that these estimates are reasonable actual results could differ.

In preparing these unaudited interim consolidated financial statements, the significant judgments made by management in applying the Company's accounting policies and the key sources of estimation uncertainty were the same as those applied to the audited consolidated financial statements for the year ended August 31, 2021.



6. Recent Accounting Guidance

Pronouncements Issued but Not Yet Adopted

In October of 2021, the FASB issued an update to Government Assistance (topic 832) to increase the transparency of government assistance and its disclosure in the notes to the financial statements. Amendments in this update take effect for annual periods beginning after December 31, 2021. Early application of the amendments is permitted. The Company does not expect the adoption of these standards to have a material impact on its consolidated financial statements

In June 2016, the FASB issued ASU No. 2016-13, Financial Instruments—Credit Losses (Topic 326): Measurement of Credit Losses on Financial Instruments. The FASB subsequently issued amendments to ASU 2016-13, which have the same effective date and transition date of January 1, 2023. These standards require that credit losses be reported using an expected losses model rather than the incurred losses model that is currently used, and establishes additional disclosures related to credit risks. For available-for-sale debt securities with unrealized losses, these standards now require allowances to be recorded instead of reducing the amortized cost of the investment. These standards limit the amount of credit losses to be recognized for available-for-sale debt securities to the amount by which carrying value exceeds fair value and requires the reversal of previously recognized credit losses if fair value increases. The Company does not currently expect the adoption of these standards to have a material impact on its consolidated financial statements.

7. Accounts and Other Receivables

	February 28, 2022	August 31, 2021
	\$	\$
Trade and deposits receivable	19,016	16,553
Sale of assets - shares receivable	278,107	287,107
Sales tax receivable	209,499	47,741
	506,622	342,401

8. Inventory

	February 28, 2022	August 31, 2021
	\$	\$
Raw materials	31,927	29,648
	31,927	29,648

During the period ended February 28, 2022, the Company wrote down \$Nil (February 28, 2021-\$2,482) in finished goods.



9. Intellectual Property

The following is a list of US capitalized patents held by the Company:

Issued Patent #	Patent Certificate Grant Date	Patent Family
US 9,474,725 B1	10/25/2016	Food and Beverage Compositions Infused With Lipophilic Active Agents and Methods of Use Thereof
US 9,839,612 B2	12/12/2017	
US 9,972,680 B2	05/15/2018	
US 9,974,739 B2	05/22/2018	
US 10,084,044 B2	09/25/2018	
US 10,103,225 B2	10/16/2018	
US 10,381,440	08/13/2019	
US 10,374,036	08/06/2019	
US 10,756,180	08/25/2020	

A continuity schedule for capitalized patents is presented below:

	February 28, 2022	August 31, 2021
	\$	\$
Balance – beginning	364,623	292,000
Addition	50,263	79,493
Amortization	(3,434)	(6,870)
Balance – ending	411,452	364,623

Patents are amortized over their 20 year legal life.

10. Property & Equipment

Six Months Ended February 28, 2022	Cost	Period Amortization	Additions	Accumulated Amortization	Net Balance February 28, 2022
	\$	\$	\$	\$	\$
Leasehold improvements	259,981	(27,019)	-	(167,667)	92,314
Computers	63,964	(8,075)	-	(59,625)	4,339
Furniture fixtures & equipment	31,126	(3,209)	-	(19,629)	11,497
Lab equipment	291,235	13,474	42,375	(82,949)	250,661
	646,306	(51,777)	42,375	(329,870)	358,811

For the six months ended February 28, 2022, amortization of \$1,118 (February 28, 2021 - \$6,010) was included in the cost of goods sold.

Year Ended August 31, 2021	Cost	Period Amortization	Disposal	Accumulated Amortization	Net Balance August 31, 2021
	\$	\$	\$	\$	\$
Leasehold improvements	259,981	(54,038)	-	(140,648)	119,333
Computers	63,964	(19,681)	-	(51,550)	12,414
Furniture fixtures & equipment	34,220	(6,417)	(3,094)	(16,420)	14,706
Lab equipment	291,235	(35,008)	-	(69,475)	221,760
	649,400	(115,144)	(3,094)	(279,093)	368,213



11. Accounts Payable and Accrued Liabilities

	February 28, 2022	August 31, 2021
	\$	\$
Accounts Payable		
Trades payable	60,873	54,668
Sales tax payable	33,244	-
Accrued Liabilities		
Corporate tax payable	-	1,055
Trades payable	22,474	45,000
Balance	116,591	100,723

12. Common Shares and Warrants

In December of 2021, the Company entered a one-year media outreach agreement to SRAX Inc. and issued 224,299 shares as consideration for an aggregate value of \$1.2m.

During the quarter ended February 28, 2022, the Company issued no warrants.

A continuity schedule for warrants is presented below:

	Number of Warrants	Weighted Average Exercise Price \$
Balance August 31, 2020	471,608	16.77
Cancelled/expired	(44,161)	67.50
Exercised	(610,189)	6.58
Issued	2,630,017	6.58
Balance August 31, 2021	2,447,275	8.00
Cancelled/expired	(25,292)	4.57
Balance February 28, 2022	2,421,983	8.04

A summary of warrants outstanding as of February 28, 2022, is presented below:

	# of Warrants	Weighted Average Remaining Contractual Life	Weighted Average Exercise Price \$
	7,500	0.68 years	24.00
	100,000	2.13 years	9.00
	200,000	2.13 years	7.00
	51,814	2.71 years	36.00
	8,984	2.75 years	36.00
	16,667	3.05 years	9.00
	317,190	3.20 years	10.50
	1,719,828	3.88 years	6.58
	2,421,983	3.53 years	8.04

13. Stock Options

The Company has established the Equity Incentive Plan whereby the board of directors may, from time to time, grant stock options up to the equivalent of 10% of the number of common shares issued and outstanding to directors, officers, employees, and consultants. Stock options granted must be exercised within five years from the date of grant or such lesser period as determined by the Company's board of directors. The exercise price of an option is equal to or greater than the closing market price of the Company's common shares on the day preceding the date of grant. The vesting terms of each grant are set by the board of directors. The Company did not grant any options during the quarter ended February 28, 2022.

A continuity schedule for stock options is presented below:

	Options	Weighted Average Exercise Price \$	Weighted Average Remaining Contractual Term (Years)	Aggregate Intrinsic Value \$
Balance August 31, 2020	171,604	11.17		
Cancelled/expired	(50,334)	10.76		
Granted	84,900	5.41		
Balance August 31, 2021	206,170	8.90		
Cancelled	(3,334)	9.60		
Granted	81,800	6.23		
Balance February 28, 2022 (Outstanding)	284,636	8.12	3.61	-
Balance February 28, 2022 (Exercisable)	260,386	8.35	3.59	-

The fair value of share purchase options granted were estimated as of the date of the grant by using the Black-Scholes option pricing model with the following assumptions:

	February 28, 2022
Expected volatility	119%
Risk-free interest rate	0.85%
Expected life	5 years
Dividend yield	0%
Estimated fair value per option	\$ 5.10

14. Revenues

	February 28, 2022 \$	February 28, 2021 \$
Product sales	17,512	231,718
Licensing revenue	16,160	255,844
Other revenue	10,858	100
Income from operations	44,530	487,662

Product revenues of \$17.5k and licensing usage fees of \$16k represent a significant decrease in intermediate product sales and related licensing usage fees during the six months ended February 28, 2022.

15. Related Party Transactions

Due to related parties:

Related party transactions are recorded at the exchange amount established and agreed to between the related parties. As at February 28, 2022, \$884, included in accrued liabilities. At August 31, 2021 - \$5,223 was payable to and included in due to related parties.

16. Segment Information

The Company's operations involve the development and usage, including licensing, of its proprietary DehydrateTECH Technology. Lexaria is centrally managed and its chief operating decision makers, being the President and the CEO, use the consolidated and other financial information supplemented by revenue information by category of alternative health consumer products and technology licensing to make operational decisions and to assess the performance of the Company. The Company has identified two reportable segments: Intellectual Property and Products. Licensing revenues are significantly concentrated on one licensee.

	IP Licensing	Products	Corporate	Consolidated
	\$	\$	\$	Total
Six Months Ended February 28, 2022	\$	\$	\$	\$
Revenue	16,160	17,512	10,858	44,530
Cost of goods sold	-	(11,957)	-	(11,957)
Operating expenses	(1,367,712)	(190,611)	(1,926,406)	(3,484,729)
Segment loss	(1,351,552)	(185,056)	(1,915,548)	3,452,156
Total assets	939,790	87,291	10,387,483	11,414,564
	IP Licensing	Products	Corporate	Consolidated
	\$	\$	\$	Total
Six Months Ended February 28, 2021	\$	\$	\$	\$
External revenue	255,844	231,818	-	487,662
Cost of goods sold	-	(95,048)	-	(95,048)
Operating expenses	1,142,518	(278,374)	(1,551,689)	(687,545)
Discontinued operations	(22,000)	-	-	(22,000)
Segment income (loss)	1,376,362	(141,604)	(1,551,689)	(316,931)
Total assets	696,493	137,282	11,338,633	12,172,408



	Cost US \$	Addition US \$	Net Balance US \$	Cost Canada \$	Net Balance Canada \$	Total Net Balance \$
Capital Asset by Region						
Six Months Ended February 28, 2022						
Leasehold Improvements	-	-	-	259,981	92,314	92,314
Computers	-	-	-	63,964	4,339	4,339
Furniture Fixtures Equipment	-	-	-	31,126	11,497	11,497
Lab Equipment	98,050	42,375	108,144	193,185	142,517	250,661
	98,050	42,375	108,144	548,256	250,667	358,811
Capital Asset by Region						
Year Ended August 31, 2021						
Leasehold Improvements	-	-	-	259,981	119,333	119,333
Computers	-	-	-	63,964	12,414	12,414
Furniture Fixtures Equipment	3,094	(3,094)	-	31,126	14,706	14,706
Lab Equipment	98,050	-	69,580	193,185	152,180	221,760
	101,144	(3,094)	69,580	548,256	298,633	368,213

17. **Commitments, Significant Contracts and Contingencies**

Right of Use Assets - Operating Lease

The Corporate office and R&D laboratory located in Kelowna, British Columbia, Canada is leased until November 15, 2023, with a five-year renewal option. In addition to minimum lease payments, the lease requires us to pay, subject to annual adjustments, property taxes and operating costs.

	February 28, 2022 \$	August 31, 2021 \$
Right of use assets - operating leases	126,920	126,920
Amortization	(54,821)	(35,879)
Total lease assets	<u>72,099</u>	<u>91,041</u>
Liabilities:	89,393	125,431
Lease payments	(22,191)	(43,950)
Interest accretion	2,954	7,912
Total lease liabilities	<u>70,156</u>	<u>89,393</u>
Operating lease cost	72,099	91,041
Operating cash flows for lease	22,191	43,950
Remaining lease term	1.7 Years	2.1 Years
Discount rate	7.50%	7.50%

Pursuant to the terms of the Company's lease agreements in effect, the following table summarizes the Company's maturities of operating lease liabilities as of February 28:

2022	\$ 22,407
2023	44,816
2024	7,469
Thereafter	-
Total lease payments	\$ 74,692
Less: imputed interest	(4,536)
Present value of operating lease liabilities	\$ 70,156
Less: current obligations under leases	(41,076)
Total	\$ 29,080

18. Prepaid Expenses and Deposits

Prepaid expenses consist of the following at February 28, 2022, and August 31, 2021:

	February 28, 2022	August 31, 2021
	\$	\$
Advertising & conferences	1,015,367	168,760
Consulting	-	18,750
Legal fees	25,000	31,380
Licence, filing fees, dues	37,500	19,500
Office & insurance	44,094	80,863
	1,121,961	319,253

19. Marketable Securities

The components of Marketable Securities were as follows:

	Cost Basis \$	Unrealized Gains \$	Unrealized Losses \$	Total \$
August 31, 2021				
Common stock	1,037,025	16,243	(219,427)	
Total	1,037,025	16,243	(219,427)	833,841
February 28, 2022				
Common stock	-	58,893	340,417	(281,524)
Total	1,037,025	73,156	(559,844)	552,317

Unrealized gains and losses from common stock are due to market price movements. Management does not believe any remaining unrealized losses represent other-than-temporary impairments based on our evaluation of available evidence.

20. Discontinued Operations

On November 19, 2020, the Company entered a definitive asset sale agreement through its wholly owned subsidiary Lexaria CanPharm ULC to sell certain non-core business assets to Hill Street Beverage Company (“Hill Street”) (TSX-V: BEER) for gross proceeds of C\$3,850,000.

With the closing of the sale on December 10, 2020, the Company received C\$350,000 in cash, 6,031,363 restricted common shares at a fair value at C\$00,000 as the first required equity-based payment, and a C\$2,000,000 promissory note bearing interest at 10% per annum. The promissory note was included at its nominal value of \$NIL. Pursuant to the terms of the transaction, the Company will receive an additional C\$1,000,000 worth of common shares of Hill Street of which C\$43,939 worth of Hill Street shares were issued to the Company on August 9, 2021, and the remaining C\$356,061 worth of Hill Street shares are to be issued on April 9, 2022.

Gain on asset disposal

Book value of assets sold	\$	-
Cash consideration		273,373
Shares received		468,264
Shares receivable		781,067
Promissory note		-
	\$	<u>1,522,704</u>

The financial results of the group of assets sold are presented as income (loss) from discontinued operations, net of income taxes in our consolidated statement of income. The following table presents financial results of the assets:

	SIX MONTHS ENDED	
	February 28,	
	2022	2021
Revenue	\$ -	\$ 3,000
Operating Expenses	-	25,000
Net Income (loss)	\$ -	\$ (22,000)

The following table presents cash flows of discontinued operations:

	SIX MONTHS ENDED	
	February 28,	
	2022	2021
Cash flows used in discontinued operating activities		
Net income	\$ -	\$ (22,000)
Change in working capital	-	105,000
Net cash used in discontinued operating activities	\$ -	\$ 83,000
Net cash provided by discontinued operations	-	\$ 83,000



21. Subsequent Events

On March 8, 2022 Lexaria granted 36,700 stock options bearing an exercise price of \$3.39 for a period of five years ending March 8, 2027. The Options were issued pursuant to the Company's registered Incentive Equity Plan and any common shares issued upon the exercise of the Options will be unrestricted securities.



Item 2. Management's Discussion and Analysis of Financial Condition and Results of Operations

Cautionary Note Regarding Forward-Looking Statements

This quarterly report contains forward-looking statements as that term is defined in the Private Securities Litigation Reform Act of 1995. Any statements contained herein that are not statements of historical fact may be forward-looking statements. These statements relate to future events or our future financial performance. Any forward-looking statements are based on our present beliefs and assumptions as well as the information currently available to us. In some cases, forward-looking statements are identified by terminology such as “may”, “will”, “should”, “could”, “targets”, “goal”, “expects”, “plans”, “anticipates”, “believes”, “estimates”, “predicts”, “potential” or “continue” or the negative of these terms or other comparable terminology. These statements are only predictions and involve known and unknown risks, uncertainties and other factors, including the risks in the section entitled “Risk Factors” set forth in Item 1(A) in our annual report on Form 10-K, as filed with the Securities and Exchange Commission on November 29, 2021, that may cause our or our industry’s actual results, levels of activity, performance or achievements to be materially different from any future results, levels of activity, performance or achievements expressed or implied by these forward-looking statements.

Although we believe that the expectations reflected in the forward-looking statements are reasonable, we cannot guarantee future results, levels of activity, performance or achievements. We caution you not to place undue reliance on any forward-looking statements as they speak only as of the date on which such statements were made, and we undertake no obligation to update any forward-looking statement or to reflect the occurrence of an unanticipated event. New factors may emerge and it is not possible to predict all factors that may affect our business and prospects. Further, management cannot assess the impact of each factor on the business or the extent to which any factor, or combination of factors, may cause actual results to differ materially from those contained in any forward-looking statements.

Our unaudited interim consolidated financial statements are stated in United States Dollars (“US\$”) and are prepared in accordance with United States Generally Accepted Accounting Principles (“US GAAP”). The following discussion should be read in conjunction with our financial statements and the related notes that appear elsewhere in this quarterly report.

In this quarterly report, unless otherwise specified, all dollar amounts are expressed in United States dollars. All references to “CS” refer to Canadian dollars and all references to “common shares” and “shares” refer to the common shares in our capital stock, unless otherwise indicated. The terms “Lexaria” “we”, “us”, “our” and “Company” mean the Company and/or our subsidiaries, unless otherwise indicated.

The following discussion should be read in conjunction with our condensed financial statements and accompanying notes in this quarterly report on Form 10-Q, and our audited financial statements with notes in our annual report on Form 10-K for the year ended August 31, 2021.

Overview

Lexaria’s patented technology DehydraTECH improves the delivery of bioactive compounds while promoting healthy ingestion methods, lowers overall dosing, and is highly effective in active molecule delivery available in a range of formats from oral ingestible to oral buccal/sublingual to topical products. DehydraTECH substantially improves the rapidity and quantity of Active Pharmaceutical Ingredients (“API”) transport to the blood plasma and brain using the body’s natural process for distributing fatty acids via the oral route. This technology extends across many categories beyond the primary pharmaceutical focus of the Company from foods and beverages to cosmetic products and nutraceuticals.



Research & Development

Lexaria is advancing several R&D activities in both preclinical and clinical programs. Currently, our primary research program is the investigation of cannabidiol (CBD) for the reduction of hypertension with a human clinical trial initiated in Q3, 2022 and three human clinical trials concluded in calendar 2021. Other programs include nicotine for oral pouches and nicotine replacement therapy, antivirals and related compounds for COVID-19 and other viral diseases, PDE5 inhibitors, NSAIDS, hormones, and others. From time to time the Company will engage in contract R&D for third parties who are interested in evaluating DehydraTECH in their products.

During the quarter ended February 28, 2022, Lexaria incurred \$275,213 (February 2021- \$176,398) in R&D expenditures. Specific R&D programs are in ongoing development and align to our financial ability to undertake each research phase for each API. Due to our expanding portfolio coverage, we continually examine accelerated timetable options for testing, research, and development of each API. The first half of fiscal 2022 continued to highlight the direction of our research and development programs with confirmatory results from our ongoing programs. We are devoting an increasing proportion of our resources and focus towards pharmaceutical applications as launched in the prior fiscal year.

In September 2021 the Company reported successful results from its HYPER-H21-2 human study of DehydraTECH-CBD in arterial stiffness. On December 29, 2021, we announced the Company received approval of its study protocols submitted for our study HYPER-H21-4 from the Independent Review Board. This study is expected to consist of 60 volunteers between the ages of 40-70 using three daily doses of DehydraTECH-CBD, every day for the multi-week dosing duration of the study. The Study was initiated during the first week of April, 2022.

HYPER-H21-4 is a double blinded, randomized cross-over design study with a placebo control. Some volunteers will already be using leading standard of care hypertension drugs such as ACE inhibitors with or without diuretics which will help evaluate the efficacy of DehydraTECH-CBD with and without other hypertension treatments. The extended duration of the study will allow Lexaria to gather critical data monitoring of DehydraTECH-CBD over time and will evaluate the potential for longer term health benefits. This study should “de-risk” outcomes prior to Lexaria’s planned entry into regulatory pathways for the use of DehydraTECH-CBD to treat hypertension and possibly other forms of cardiovascular disease.

In October 2021 the Company reported results from its THC-A-21-1 in vivo study of DehydraTECH-THC showing it successfully elevated THC levels in blood plasma, requiring only 15 minutes at levels comparable to those achieved in 45 minutes with concentration-matched controls.

Lexaria also published results from its NIC-A21-1 in vivo trial of DehydraTECH-NICZ (nicotine benzoate) which was delivered via oral pouches. Favorable results were all statistically significant, supporting further evaluation of the candidate. Lexaria plans to progress to a larger investigation in human volunteers of DehydraTECH-nicotine versus leading brands. Lexaria is currently in the design phase of this proposed human clinical study, which will be independently funded with existing capital.

On February 2, 2022, results of our PDE5-A21-1 animal study were published by the Company which illustrated that in as little as four minutes after dosing, the DehydraTECH formulation delivered 74% more sildenafil into the bloodstream on average than the concentration-matched, generic control formulation. Seven minutes after dosing, the DehydraTECH-sildenafil formulation achieved an average blood level higher than the generic sildenafil control formulation reached at any point during the study.



On March 15, 2022, the Company announced that the first phase of its epilepsy research program EPIL-A21-1 has commenced. The EPIL-A21-1 research program consists of two main studies to be performed in rodents following the first phase pilot animal model. The two main studies within the program are expected to begin in May/June 2022, and will involve both an acute seizure model induced by electrical stimulation (“MES”) as well as a chronic chemically induced seizure model (“RISE-SRS”). Lexaria has selected these models because they have been previously employed by other researchers studying the antiepileptic effects of CBD including select study work funded by GW with its Epidiolex® formulation (PubMed Reference Number 30588604).

The Company continues to report progress on its R&D programs through its filing of Form 8-Ks and other public releases. The results of these programs can also be found on the Company’s website: www.LexariaBioscience.com.

Patents

Our current patent portfolio includes patent family applications or grants pertaining to our method of improving bioavailability and taste, and the use of DehydraTECH as a delivery platform for a wide variety of Active Pharmaceutical Ingredients (“APIs”) including, but not limited to, fat soluble vitamins; nonsteroidal anti-inflammatory drugs (“NSAIDs”); anti-viral drugs; phosphodiesterase inhibitors; human hormones; regulated cannabinoids, and nicotine and its analogs.

We will continue to pursue patent protection in more than 40 countries around the world as vigorously as we are able, since the successful granting of more of those applications could lead to material increases in shareholder value. The Company currently has over 50 patent applications pending worldwide.

The Company’s issued patents in the United States, Australia, Europe, India, Mexico, and Japan are as follows:

Issued/Allowed Patent #	Patent Family
US 9,474,725 B1	<p align="center">Food and Beverage Compositions Infused with Lipophilic Active Agents and Methods of Use Thereof</p>
US 9,839,612 B2	
US 9,972,680 B2	
US 9,974,739 B2	
US 10,084,044 B2	
US 10,103,225 B2	
US 10,381,440	
US 10,374,036	
US 10,756,180	
AU 2015274698	
AU 2017203054	
AU 2018202562	
AU 2018202583	
AU 2018202584	
AU 2018220067	
EP 3164141	
JP 6920197	
AU 2016367036	<p align="center">Methods for Formulating Orally Ingestible Compositions Comprising Lipophilic Active Agents</p>
JP 6963507	
MX 388203 B	
AU 2016367037	<p align="center">Stable Ready-to-Drink Beverage Compositions Comprising Lipophilic Active Agents</p>
IN 365864	
JP 6917310	
AU 2019256805	<p align="center">Compositions Infused with Nicotine Compounds and Methods of Use Thereof</p>



In November of 2021, we were advised that our first patent in Mexico, in the Company's second patent family, had been allowed.

Subsequent to the quarter ended February 28, 2022, on March 8, 2022, the Company announced it has been granted a new patent entitled "Compositions Infused with Nicotine Compounds and Methods of Use Thereof". The new Australian patent expands upon Lexaria's international intellectual property rights to apply DehydraTECH enhancement technology to most oral forms of nicotine, including pills, tablets, lozenges, capsules, pouches, gums and sprays. The patent covers many different forms of nicotine including free base nicotine, nicotine salts, polymer resins of nicotine and other forms of nicotine complexes.

Due to the complexity of pursuing patent protection, the quantity of patent applications will vary continuously as each application advances or stalls. The Company is also filing new patent applications for new discoveries that arise from the Company's R&D programs and, due to the inherent unpredictability of scientific discovery, it is not possible to predict if or how often such new applications might be filed.

Reverse Stock Split

On January 11, 2021, the Company filed an amendment and restatement of its articles of incorporation to effectuate a 1-for-30 reverse stock split of the issued and outstanding shares of common stock of the Company. The purpose of the reverse stock split was to meet Nasdaq's minimum stock price requirement. The reverse stock split did not change the number of authorized shares of common stock, which remains at 220,000,000 shares. All warrants, options, share and per share information in this Report gives retroactive effect to the 1-for-30 reverse stock split.

Public Offering

On January 14, 2021, the Company closed an underwritten public offering with the issuance of 2,102,856 shares of the Company's common stock priced at \$5.25 per share with an equivalent number of five-year warrants at an exercise price of \$6.58. Additionally, 227,161 Representative Warrants were issued as partial consideration to the underwriters of the offering that have a five-year term at an exercise price of \$6.58. Net of fees and disbursements, the Company received net proceeds of \$9,471,497.

LEXX Market Listing

The Company's common stock was uplisted from trading on the OTCQX under "LXRP" to the Nasdaq Capital Market where our common stock and certain of our warrants began trading under the symbols "LEXX" and "LEXXW", respectively, effective as of the opening of market trading on January 12, 2021.

The Company, trading under the symbol "LXX", voluntarily delisted from the Canadian Securities Exchanges ("CSE") effective after the closing of trading on Wednesday, July 7, 2021. The overwhelming majority of trading had moved to Nasdaq and by delisting from the CSE the Company expected to realize savings in fees and managerial time and effort that had been required to maintain a dual listing.



Asset Sale

On December 9, 2020, Lexaria CanPharm ULC (“CanPharm”) completed a disposition (the “Disposition”) of its use and licensing rights to use its DehydraTECH technology (the “Assets”) specifically in association with non-pharmaceutical products containing cannabis molecules that contain 0.3% or greater THC. The purpose of the Disposition was to remove the Company’s association with cannabis as it remains a Schedule 1 Drug and thereby eliminating any such regulatory restrictions cannabis products may create. The Disposition assisted the Company in obtaining a listing on the Nasdaq Capital Market (“Nasdaq”) on January 12, 2021. As a result of the Disposition, CanPharm assigned to the purchaser, Hill Street, license agreements with three existing non-related party licensees.

In consideration for the Assets, Hill Street provided CanPharm with C\$350,000 cash, a promissory note bearing a principal amount of C\$2,000,000 and bearing an interest rate of 10% (the “Note”) and C\$1,500,000 in shares of Hill Street, issuable in three tranches by April 9, 2022, of which C\$1,149,939.43 worth of Hill Street stock has been issued to CanPharm to date. The repayment of the Note does not have a fixed maturity date and is based on quarterly installments equal to 5% of the gross sales realized by Hill Street of DehydraTECH enabled products. Due to the uncertainty pertaining to the settlement of the Note, management concluded that the note had \$Nil value at the time of the sale and was recorded as such. Some of the factors considered in the \$Nil valuation of the Note were that the legal sales of THC products in the US and Canada have little or no history which made the expectant quarterly payments very difficult to forecast. Further, Hill Street had no experience selling THC products and at the time of the sale was not licensed to produce and sell such products. Therefore, the Company considered risk of default high and the collectability of the Note as highly doubtful. Since the date of sale Hill Street has repaid \$4,585 in the year-ended August 31, 2021. Subsequent to fiscal 2021, the Company has received a further \$10,858, included in other revenues, as payment toward the balance of the Note and accumulated interest on the Note.

Impact of COVID-19

The COVID-19 pandemic continues to present uncertainty and unforecastable new risks to the Company and its business plan. Restrictions on national and international travel and required business closures present challenges in carrying out normal business activities related to corporate finance efforts and the pursuit of new customers throughout North America who might otherwise access the retail products of our licensees. As a result, the pandemic has increased risk of lower revenues and higher losses. To date, we have not experienced a material impact on our financial statements, impairments of any of our assets or any major business disruptions, including with our vendors.

The extent to which the COVID-19 pandemic will impact the progress of our research and development programs is subject to future developments which are highly uncertain and cannot be predicted with confidence. We continue our efforts to be proactive in managing the impact from the pandemic, including various actions to communicate with suppliers, customers, and project participants as we may deem appropriate.

We have made modifications to our normal operations including requiring team members to work remotely on a staggered basis. To the extent possible, we are conducting business as usual, with necessary or advisable modifications to employee travel. At this time, these measures will continue in force for the near term.

We will continue to actively monitor the rapidly evolving situation related to COVID-19 and may take further actions that alter our operations, including those that may be required by federal, state, provincial, or local authorities, or that we determine are in the best interests of our employees and other third parties with whom we do business. We do not know when, or if, it will become practical to relax or eliminate some or all these measures entirely.



Off-Balance Sheet Arrangements

We have no off-balance sheet arrangements that have or are reasonably likely to have a current or future effect on our financial condition, changes in financial condition, revenues or expenses, results of operations, liquidity, capital expenditures or capital resources that are material to stockholders.

Critical Accounting Estimates

Our consolidated financial statements and accompanying notes are prepared in accordance with US GAAP. These accounting principles require management to make certain estimates, judgments and assumptions that affect the reported amounts of assets, liabilities, revenue, and expenses during the periods reported. These estimates, judgments and assumptions are reasonable based on information available to management at the time that such estimates, judgments and assumptions are made. We believe that understanding the basis and nature of the estimates, judgments and assumptions involved with the following aspects of our financial statements is critical to an understanding of our financials. For a discussion of our critical accounting estimates, please read Note 2 to our financial statements in our Annual Report on Form 10-K for the year ended August 31, 2021. There have been no material changes to the critical accounting estimates previously disclosed in our Annual Report on Form 10-K for the year ended August 31, 2021.

Capital Assets

Capital assets are stated at cost less accumulated depreciation and depreciated using the straight-line method over their useful lives or otherwise by units of production.

Patents

US patent costs, which represent legal costs incurred to establish US granted patents, are capitalized. Capitalized patent costs are amortized on a straight-line basis over the remaining life of the patent. When US patents reach a mature stage, any associated legal costs, including typical maintenance fees, are expensed as incurred. In the period ended February 28, 2022, the Company recognized \$15,840 attributable to capitalized patents. All other patent costs are expensed when incurred.

Revenue Recognition

Product Revenue

Revenue from the sale of products is recognized when persuasive evidence of an arrangement exists, delivery has occurred, the sales price is fixed or determinable, and collectability is reasonably assured, which typically occurs upon shipment. The Company reports its sales net of actual sales returns.

Licensing Revenue from Intellectual Property

We recognize revenue for license fees at a point in time following the transfer of our intellectual property, our patented lipid nutrient infusion technology DehydraTECH for infusing APIs, to the licensee, which typically occurs on delivery of documentation.



Usage Fees from Intellectual Property

We recognize revenue for usage fees when usage of our DehydraTECH intellectual property occurs by licensees infusing an API into one or more of their product lines for sale.

Funding Requirements

We anticipate that our expenditures will increase in connection with our ongoing R&D program, specifically with respect to our animal and human clinical trials of our DehydraTECH formulations for the purposes of treating hypertension and infectious diseases. As we move forward with our Investigational New Drug application with the FDA, we anticipate that our expenditures will further increase and accordingly, we expect to incur increased operating losses and negative cash flows for the foreseeable future.

Through February 28, 2022, we have funded our operations primarily with proceeds from the sale of our common stock. The Company has consistently incurred recurring losses and negative cash flows from operations, including net losses of \$3,452,156 and \$316,931 for the six months ended February 28, 2022, and 2021, respectively.

The continuation of our Company as a going concern is dependent upon our Company raising additional capital and/or attaining and maintaining profitable operations. The accompanying financial statements do not include any adjustment relating to the recovery and classification of recorded asset amounts or the amount and classification of liabilities that might be necessary should our Company discontinue operations. The recurring losses from operations and net capital deficiency do raise doubt about the Company's ability to continue as a going concern within one year following the date that these consolidated financial statements are issued. As of the issuance date of these consolidated interim financial statements, we expect our cash and cash equivalents of approximately \$8.4m as at February 28, 2022, will be sufficient to fund our operating expenses and capital expenditure requirements through the forthcoming 12 months from the issuance date of this report.

Results of Operations for our Period Ended February 28, 2022, and 2021

Our net loss and comprehensive loss for the six months ended and the changes between those periods for the respective items are summarized as follows:

	SIX MONTHS ENDED		
	February 28,		
	2022	2021	Change
	\$	\$	\$
Revenue	44,530	487,662	(443,132)
Research and development	734,395	368,659	365,736
Consulting fees & salaries	1,174,863	839,459	335,404
Legal and professional	365,657	467,252	(101,595)
Other general and administrative	1,209,814	(987,825)	2,197,639
Discontinued operations	-	(22,000)	22,000
Net Loss	<u>(3,452,156)</u>	<u>(316,931)</u>	<u>(3,135,225)</u>



Revenue

Product revenues of \$17k and licensing usage fees of \$16k during period ended February 28, 2022, constitute a significant decline in intermediate product sales and related licensing usage fees. Our primary customer in the B2B product revenue stream has been delayed in chain-store rollouts resulting in overstocked inventory and as such there was no manufacturing & sales of new inventory for this licensee for the quarter ended February 28, 2022. A number of our other licensees are experiencing suspended or curtailed business activities due to the impact of COVID-19 on markets and consumer spending. The abilities of other licensees to generate ongoing sales, thereby increasing usage fees are expected to increase as the effects of the pandemic are eventually diminished and the market acceptance of the products continue to develop. We have continued strong interest in our intermediate products but cannot predict how long the pandemic will affect purchasing decisions of retail customers that ultimately affect the consumer product manufacturers that utilize our intermediate products. Nor can we predict when recovery of the general economy will translate into increasing licensing or usage revenues.

Our licensing revenues consist of IP licensing fees for the transfer of the Technology and usage fees that occur over time. IP licensing fees are due at the signing of definitive agreements for the Technology and can include payments due upon transfer of the Technology and installment payments that are receivable within 12 months.

Research and Development

Expenditures on R&D increased by \$366k for the period ended February 28, 2022, as the company undertook several studies within its 2022 applied research and development program focusing on DehydraTECH-CBD to treat hypertension.

Consulting Fees and Salaries

Our consulting fees increased by \$335k primarily due to non-cash stock-based compensation on options granted and vested (\$287k) in the six months ended February 28, 2022, and increased salaries due to hiring within our R&D, investor relations and accounting departments.

Legal and Professional Fees

Our professional fees decreased by \$102k during the period compared to the same period in the prior year. Prior year expenditures were higher due to increased patent and trademark filings, the up list to the Nasdaq Capital Markets, and additional advisory services utilized. We recognize certain legal fees, tax advice fees, and accounting services all as "Professional Fees."

General and Administrative

Included in general and administrative expenses during the six months ended February 28, 2021 was the gain on the sale of assets (\$1.5m) which accounts for approximately half of the year-over-year change. Other general and administrative expenses increased by \$675k during the period ended February 28, 2022, over the same period last year. The increase is primarily comprised of expenditures on investor relations and advertising (\$505k) and the unrealized losses on marketable securities (\$264k) offset by lower general expenses (\$74k) and bad debts (\$25k).



Liquidity and Financial Condition

Working Capital	February 28, 2022	August 31, 2021
	\$	\$
Current assets	10,572,202	12,442,940
Current liabilities	(165,542)	(153,276)
Net Working Capital	10,406,660	12,289,664

Cash Flows	February 28, 2022	February 28, 2021
	\$	\$
Cash flows (used in) provided by operating activities	(2,443,593)	(1,742,757)
Cash flows (used in) provided by investing activities	(92,638)	264,609
Cash flows (used in) provided by financing activities	(21,191)	9,448,332
Net cash flows (used in) discontinued operations	-	83,000
Increase (decrease) in cash	(2,557,422)	8,053,184

Operating Activities

Net cash used in operating activities for the six months ended February 28, 2022, increased by \$723k for the period compared with cash used in operating during the same period in 2021. This difference was largely due to the increased expenditures pertaining to R&D, professional fees, and our stakeholders outreach programs.

Investing Activities

Net cash used in investing activities increased by \$93k due to increased spending on capitalized US patents filings and the purchase of equipment in the six months ended February 28, 2022, compared to 2021 which saw an inflow of cash from the sale of assets to Hill Street.

Financing Activities

Net cash provided from financing activities during the period ended February 28, 2022, is due to the amortized right-of-use lease payments. Cash provided by financing activities in the six months ended February 28, 2021, of \$9.4m primarily related to the underwritten public offering that closed January 21, 2021.

Liquidity and Capital Resources

We have accumulated a large deficit since inception that has primarily resulted from executing our business plan, including R&D expenditures, in seeking to identify and develop our intellectual property patents for licensing and product creation. We expect to continue to incur losses for at least the short term.

To date, we have obtained cash and funded our operations primarily through equity financings and limited amounts from revenue generation while our licensees ramp up production and market expansions. We expect to continue to evaluate various funding alternatives on an ongoing basis as needed to maintain operations, to continue our research programs and to expand our patent portfolio. If we determine it is advisable to raise additional funds, there is no assurance that adequate funding will be available to us or, if available, that such funding will be available on terms that we or our stockholders view as favorable. Market volatility and global economics may have a significant impact on the availability of funding sources and the terms at which any funding may be available.

Short Term Liquidity

On February 28, 2022, we had \$8.4m cash and \$10.4m of working capital. Based on our current and upcoming research and development programs and our projected general and administration expenditures, we have determined that our cash resources are sufficient to allow us to continue operations through at least the next twelve months from the issuance date of this Quarterly Report.

Long Term Liquidity

It will require substantial cash to achieve our objectives for developing and patenting our intellectual property across all applicable market and industry segments. This process typically takes many years and potentially millions of dollars for each segment. If we pursue full commercial exploitation of all applicable market and industry segment opportunities, we will need to obtain significant funding from existing or new relationships, increasing revenue streams or from other sources of liquidity such as the sale of equity, issuance of debt or other transactions.

Cash requirements will vary depending on the results of our research programs and the requirements of each industry segment pursued. Pursuit of each segment will progress or be curtailed based on available sources of cash with which to execute individual segment business plans. The requirements will also be affected by transactions with existing or new relationships and the depth of regulatory requirements in each segment for compliance required to approve our IP and to market and license it. These changes to requirements and transactions may impact our liquidity as well as affect our expenditures.

Item 3. Controls and Procedures

Management's Report on Disclosure Controls and Procedures

We maintain disclosure controls and procedures that are designed to ensure that information required to be disclosed in our reports filed under the Securities Exchange Act of 1934, as amended, is recorded, processed, summarized and reported within the time periods specified in the Securities and Exchange Commission's rules and forms, and that such information is accumulated and communicated to our management, including our President and Chief Executive Officer (Principal Executive Officer) and our Chief Financial Officer (Principal Financial and Accounting Officer) to allow for timely decisions regarding required disclosure.

As of February 28, 2022, the quarter covered by this report, we carried out an evaluation, under the supervision and with the participation of our CEO, President and CFO, of the effectiveness of the design and operation of our disclosure controls and procedures. Based on the foregoing, our CEO, President, and CFO concluded that our disclosure controls and procedures were effective at a reasonable assurance level as of February 28, 2022.



Management's Report on Internal Control over Financial Reporting

Management is responsible for establishing and maintaining adequate internal control over financial reporting. Responsibility estimates and judgments by management are required to assess the expected benefits and related costs of control procedures. The objectives of internal control include providing management with reasonable, but not absolute, assurance that assets are safeguarded against loss from unauthorized use or disposition, and that transactions are executed in accordance with management's authorization and recorded properly to permit the preparation of consolidated financial statements in conformity with US GAAP. Our management assessed the effectiveness of our internal control over financial reporting as of February 28, 2022. In making this assessment, our management used the criteria set forth by the Committee of Sponsoring Organizations of the Treadway Commission ("COSO") in Internal Control-Integrated Framework. Our management has concluded that, as of February 28, 2022, our internal control over financial reporting is effective in providing reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with US GAAP. Our management reviewed the results of their assessment with our Board.

Inherent limitations on Effectiveness of Controls

Internal control over financial reporting has inherent limitations which include but is not limited to the use of independent professionals for advice and guidance, interpretation of existing and/or changing rules and principles, regulations, segregation of management duties, scale of organization, and personnel factors. Internal control over financial reporting is a process which involves human diligence and compliance and is subject to lapses in judgment and breakdowns resulting from human error. Internal control over financial reporting also can be circumvented by collusion or improper management override. Because of its inherent limitations, internal control over financial reporting may not prevent or detect misstatements on a timely basis, however these inherent limitations are known features of the financial reporting process and it is possible to design into the process safeguards to reduce, though not eliminate, this risk. Therefore, even those systems determined to be effective can provide only reasonable assurance with respect to financial statement preparation and presentation. Projections of any evaluation of effectiveness to future periods are subject to the risk that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies or procedures may deteriorate.

Changes in Internal Control over Financial Reporting

During the quarter ended February 28, 2022, our controls and controls processes remained consistent with our fiscal year ended August 31, 2021. There have been no changes in our internal controls over financial reporting that occurred during the quarter ended February 28, 2022, that have materially or are reasonably likely to materially affect our internal controls over financial reporting.

Our control processes are designed to include remote workers, which we have utilized for many years. The advent of the COVID-19 pandemic has not materially impacted our internal controls over financial reporting other than increasing requirements for social distancing and some additional remote working requirements for staff.



PART II—OTHER INFORMATION

Item 1. Legal Proceedings

We know of no material, existing or pending legal proceedings against our Company, nor are we involved as a plaintiff in any other material proceeding or pending litigation. There are no proceedings in which any of our directors, executive officers or affiliates, or any registered or beneficial stockholder, is an adverse party or has a material interest adverse to our interest.

Item 1A. Risk Factors

Much of the information included in this quarterly report includes or is based upon estimates, projections or other “forward looking statements”. Such forward looking statements include any projections or estimates made by us and our management in connection with our business operations. While these forward-looking statements, and any assumptions upon which they are based, are made in good faith and reflect our current judgment regarding the direction of our business, actual results will almost always vary, sometimes materially, from any estimates, predictions, projections, assumptions or other future performance suggested herein.

The risks associated with our business, common stock and other factors were with those described in the consolidated financial statements for the year ended August 31, 2021.



Item 2. Exhibits, Financial Statement Schedules

a) Financial Statements

- 1) Financial statements for our Company are listed in the index under Item 1 of this document
- 2) All financial statement schedules are omitted because they are not applicable, not material or the required information is shown in the financial statements or notes thereto.

b) Exhibits

c) Exhibit Number	Description
(3)	Articles of Incorporation and Bylaws
3.1	Articles of Incorporation (incorporated by reference as Exhibit 3.1 to our Registration Statement on Form S-1 filed June 3, 2020)
3.2	Bylaws (incorporated by reference as Exhibit 3.2 to our Registration Statement on Form S-1 filed June 3, 2020)
3.3	Amended and Restated Articles of Incorporation (Filed on Form 8-K January 14, 2021 Exh. 3.1)
3.4	Second Amended and Restated Bylaws (incorporated by reference as Exhibit 3.2 to our Current Report on Form 8-K filed January 14, 2021)
3.5	Amended and Restated Bylaws (Filed on Form S-1 June 3, 2020 Exh 3.4)
3.6	Amendment to Articles of Incorporation – Share Consolidation (Filed on Form 8-K June 23, 2009 Exh 3.1)
3.7	Amendment to Articles of Incorporation – Share Expansion (incorporated by reference as Exhibit 3.5 to our Registration Statement on Form S-1 filed June 3, 2020)
3.8	Amendment to Articles of Incorporation – Share Forward Split (Filed on Form 8-K December 16th, 2015 Exh 3.1)
3.9	Amendment to Articles of Incorporation – Name Change (Filed on Form 8-K May 11th, 2016 Exh 99.1)
(10)	Material Contracts
10.1	Executive Employment Agreement dated Dec. 31, 2021 with John Docherty (Filed on Form 10-Q January 14, 2022 Exh 10.1)
10.2	Management Services Agreement dated Dec. 31, 2021 with C.A.B. Financial Services Ltd. (Chris Bunka) (Filed on Form 10-Q January 14, 2022 Exh. 10.2)
(21)	Subsidiaries
21.1	List of Subsidiaries of the Registrant (Filed on Form 10-K November 29, 2021 Exh 21.1)
(31)	Rule 13(a) - 14 (a)/15(d) - 14(a)
31.1	Section 302 Certifications under Sarbanes-Oxley Act of 2002 of Principal Executive Officer
31.2	Section 302 Certifications under Sarbanes-Oxley Act of 2002 of Principal Financial Officer and Principal Accounting Officer
(32)	Section 1350 Certifications
32.1	Section 906 Certification under Sarbanes Oxley Act of 2002 of Principal Executive Officer
32.2	Section 906 Certification under Sarbanes Oxley Act of 2002 of Principal Financial Officer and Principal Accounting Officer
(101)**	Interactive Data Files
101.INS	XBRL Instance Document
101.SCH	XBRL Taxonomy Extension Schema Document
101.CAL	XBRL Taxonomy Extension Calculation Linkbase Document
101.DEF	XBRL Taxonomy Extension Definition Linkbase Document
101.LAB	XBRL Taxonomy Extension Label Linkbase Document
101.PRE	XBRL Taxonomy Extension Presentation Linkbase Document

** Furnished herewith. Pursuant to Rule 406T of Regulation S-T, the Interactive Data Files on Exhibit 101 hereto are deemed not filed or part of any registration statement or prospectus for purposes of Sections 11 or 12 of the Securities Act of 1933, are deemed not filed for purposes of Section 18 of the Securities and Exchange Act of 1934, and otherwise are not subject to liability under those sections.



SIGNATURES

In accordance with Section 13 or 15(d) of the Exchange Act, the registrant caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

LEXARIA BIOSCIENCE CORP.

By: /s/ Christopher Bunka
Christopher Bunka
Chief Executive Officer, Chairman and Director
(Principal Executive Officer)
Date: April 11, 2022

In accordance with the Exchange Act, this Report has been signed below by the following persons on behalf of the registrant and in the capacities and on the dates indicated.

By: /s/ Christopher Bunka
Christopher Bunka
Chief Executive Officer, Chairman and Director
(Principal Executive Officer)
Date: April 11, 2022

By: /s/ John Docherty
John Docherty
President and Director
Date: April 11, 2022

By: /s/ Greg Downey
Greg Downey CPA, CMA
Chief Financial Officer
(Principal Financial and Accounting Officer)
Date: April 11, 2022



**CERTIFICATION PURSUANT TO
18 U.S.C. ss 1350, AS ADOPTED PURSUANT TO
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002**

I, Chris Bunka, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Lexaria Bioscience Corp.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's first fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: April 11, 2022

/s/ " Chris Bunka "

Chris Bunka
CEO and Director
(Principal Executive Officer)

**CERTIFICATION PURSUANT TO
18 U.S.C. ss 1350, AS ADOPTED PURSUANT TO
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002**

I, Gregory Downey, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of Lexaria Bioscience Corp.;
2. Based on my knowledge, this report does not contain any untrue statement of a material fact or omit to state a material fact necessary to make the statements made, in light of the circumstances under which such statements were made, not misleading with respect to the period covered by this report;
3. Based on my knowledge, the financial statements, and other financial information included in this report, fairly present in all material respects the financial condition, results of operations and cash flows of the registrant as of, and for, the periods presented in this report;
4. The registrant's other certifying officer(s) and I are responsible for establishing and maintaining disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) and internal control over financial reporting (as defined in Exchange Act Rules 13a-15(f) and 15d-15(f)) for the registrant and have:
 - a. Designed such disclosure controls and procedures, or caused such disclosure controls and procedures to be designed under our supervision, to ensure that material information relating to the registrant, including its consolidated subsidiaries, is made known to us by others within those entities, particularly during the period in which this report is being prepared;
 - b. Designed such internal control over financial reporting, or caused such internal control over financial reporting to be designed under our supervision, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles;
 - c. Evaluated the effectiveness of the registrant's disclosure controls and procedures and presented in this report our conclusions about the effectiveness of the disclosure controls and procedures, as of the end of the period covered by this report based on such evaluation; and
 - d. Disclosed in this report any change in the registrant's internal control over financial reporting that occurred during the registrant's most recent fiscal quarter (the registrant's first fiscal quarter in the case of an annual report) that has materially affected, or is reasonably likely to materially affect, the registrant's internal control over financial reporting; and
5. The registrant's other certifying officer(s) and I have disclosed, based on our most recent evaluation of internal control over financial reporting, to the registrant's auditors and the audit committee of the registrant's board of directors (or persons performing the equivalent functions):
 - a. All significant deficiencies and material weaknesses in the design or operation of internal control over financial reporting which are reasonably likely to adversely affect the registrant's ability to record, process, summarize and report financial information; and
 - b. Any fraud, whether or not material, that involves management or other employees who have a significant role in the registrant's internal control over financial reporting.

Date: April 11, 2022

/s/ "Gregory Downey"

Gregory Downey CPA, CMA
Chief Financial Officer
(Principal Financial Officer and Principal Accounting
Officer)

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

I, Chris Bunka, hereby certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

- (1) the Quarterly Report on Form 10-Q of Lexaria Bioscience Corp. for the quarter ended February 28, 2022 (the "Report") fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
- (2) the information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of Lexaria Bioscience Corp.

Dated: April 11, 2022

/s/ "Chris Bunka "

Chris Bunka
CEO and Director
(Principal Executive Officer)
Lexaria Bioscience Corp.

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signature that appears in typed form within the electronic version of this written statement required by Section 906, has been provided to Lexaria Bioscience Corp. and will be retained by Lexaria Bioscience Corp. and furnished to the Securities and Exchange Commission or its staff upon request.

CERTIFICATION PURSUANT TO
18 U.S.C. SECTION 1350, AS ADOPTED PURSUANT TO
SECTION 906 OF THE SARBANES-OXLEY ACT OF 2002

I, Gregory Downey, hereby certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

- (1) the Quarterly Report on Form 10-Q of Lexaria Bioscience Corp. for the quarter ended February 28, 2022 (the "Report") fully complies with the requirements of Section 13(a) or 15(d) of the Securities Exchange Act of 1934; and
- (2) the information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of Lexaria Bioscience Corp.

Dated: April 11, 2022

/s/ "Gregory Downey"
Gregory Downey CPA, CMA
Chief Financial Officer
(Principal Financial Officer and Principal Accounting Officer)
Lexaria Bioscience Corp.

A signed original of this written statement required by Section 906, or other document authenticating, acknowledging, or otherwise adopting the signature that appears in typed form within the electronic version of this written statement required by Section 906, has been provided to Lexaria Bioscience Corp. and will be retained by Lexaria Bioscience Corp. and furnished to the Securities and Exchange Commission or its staff upon request.