

**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 March 31, 2021

OR

**TRANSITION REPORT PURSUANT TO SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934 FOR THE TRANSITION PERIOD FROM TO**

Commission File Number 001-39825

**GBS Inc.**

(Exact name of Registrant as specified in its Charter)

Delaware  
(State or other jurisdiction  
of incorporation or organization)

708 3rd Avenue, 6th Floor, New York  
(Address of principal executive offices)

82-1512711  
(I.R.S Employer  
Identification No.)

10017  
(Zip Code)

Registrant's telephone number, including area code: (646) 828-8258

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

Title of each class	Trading Symbol(s)	Name of each exchange on which registered
Common Stock, par value \$ 0.01 per share	GBS	Nasdaq Global Market

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 past 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 such files).

YES  NO

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

Large accelerated filer   
Non-large accelerated filer

Accelerated filer   
Smaller reporting company   
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.

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

The number of shares of registrant's common stock outstanding as of May 13, 2021 was 12,382,122.

## Table of Contents

<b><u>PART I. FINANCIAL INFORMATION</u></b>	<b>3</b>
<b><u>Item 1. Financial statements (Unaudited)</u></b>	<b>3</b>
<u>Condensed Consolidated Balance Sheets</u>	3
<u>Condensed Consolidated Statements of Operations and Other Comprehensive Loss</u>	4
<u>Condensed Consolidated Statements of Changes in Shareholders' Equity</u>	5
<u>Condensed Consolidated Statements of Cash Flows</u>	7
<u>Notes to Condensed Consolidated Financial Statements</u>	8
<b><u>Item 2. Management's Discussion and Analysis of Financial Condition and Result of Operations</u></b>	<b>18</b>
<b><u>Item 3. Quantitative and Qualitative Disclosures about Market Risk</u></b>	<b>26</b>
<b><u>Item 4. Controls and Procedures</u></b>	<b>26</b>
<b><u>PART II—OTHER INFORMATION</u></b>	<b>27</b>
<b><u>Item 1. Legal Proceedings.</u></b>	27
<u>Item 1a. Risk Factors</u>	27
<b><u>Item 2. Unregistered Sales of Equity Securities and Use of Proceeds.</u></b>	27
<b><u>Item 3. Defaults Upon Senior Securities.</u></b>	28
<b><u>Item 4. Mine Safety Disclosures.</u></b>	28
<b><u>Item 5. Other Information.</u></b>	28
<b><u>Item 6. Exhibits.</u></b>	28
<u>Signatures</u>	29

PART I. FINANCIAL INFORMATION

ITEM 1. FINANCIAL STATEMENTS (UNAUDITED)

**GBS Inc.**  
**CONDENSED CONSOLIDATED BALANCE SHEETS**  
**(Unaudited)**  
**(Amounts in \$)**

	<u>March 31, 2021</u>	<u>June 30, 2020</u>
<b>ASSETS</b>		
Current assets:		
Cash and cash equivalents	\$ 14,261,622	\$ 427,273
Deferred charges	-	1,863,613
Other current assets	<u>2,324,389</u>	<u>49,062</u>
Total current assets	16,586,011	2,339,948
Investment in affiliate	-	135,692
Other non-current assets	<u>866,667</u>	<u>-</u>
<b>TOTAL ASSETS</b>	<b>\$ 17,452,678</b>	<b>\$ 2,475,640</b>
<b>LIABILITIES AND SHAREHOLDERS' EQUITY (DEFICIT)</b>		
Current liabilities:		
Accounts payable and accrued expenses	\$ 1,521,899	\$ 787,469
Related party payables	37,235	1,769,293
Convertible notes payable	-	5,133,706
Total current liabilities	<u>1,559,134</u>	<u>7,690,468</u>
Employee benefit liabilities	<u>18,128</u>	<u>-</u>
Total liabilities	1,577,262	7,690,468
Commitments and contingencies - Note 10		
Shareholders' equity (deficit):		
Preferred stock, \$0.01 par value, 10,000,000 shares authorized, 3,000,000 and 2,370,891 shares issued and outstanding at March 31, 2021 and June 30, 2020, respectively	30,000	23,709
Common stock, \$0.01 par value, 100,000,000 shares authorized, 11,881,322 and 8,630,000 shares issued and outstanding at March 31, 2021 and June 30, 2020, respectively	118,813	86,300
Additional paid-in capital	38,440,097	10,899,942
Accumulated deficit	(22,016,804)	(15,832,517)
Accumulated other comprehensive loss	<u>(642,695)</u>	<u>(363,951)</u>
Total consolidated group equity (deficit)	15,929,411	(5,186,517)
Non-controlling interests	<u>(53,995)</u>	<u>(28,311)</u>
Total shareholders' equity (deficit)	<u>15,875,416</u>	<u>(5,214,828)</u>
<b>TOTAL LIABILITIES AND SHAREHOLDERS' EQUITY</b>	<b>\$ 17,452,678</b>	<b>\$ 2,475,640</b>

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

**GBS Inc.**  
**CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS AND OTHER COMPREHENSIVE LOSS**  
**(Unaudited)**  
**(Amounts in \$)**

	<b>Three Months Ended March 31,</b>		<b>Nine Months Ended March 31,</b>	
	<b>2021</b>	<b>2020</b>	<b>2021</b>	<b>2020</b>
<b>Revenues</b>				
<b>Other income:</b>				
Government support income	\$ 34,290	-	\$ 372,754	\$ -
Shared services	-	(1,533)	-	119,744
<b>Total revenues and other income</b>	<b>34,290</b>	<b>(1,533)</b>	<b>372,754</b>	<b>119,744</b>
<b>Operating expenses:</b>				
General and administrative expenses	1,013,389	53,842	2,205,842	1,752,182
Development and regulatory approval expenses	2,156,316	(58,825)	2,529,074	541,023
Prospectus and capital raising expenses	5,100	32,274	358,674	174,639
<b>Total operating expenses</b>	<b>3,174,805</b>	<b>27,291</b>	<b>5,093,590</b>	<b>2,467,844</b>
<b>Loss from operations</b>	<b>(3,140,515)</b>	<b>(28,824)</b>	<b>(4,720,836)</b>	<b>(2,348,100)</b>
<b>Other (expense) income:</b>				
Interest expense	(18,561)	(82,305)	(1,091,249)	(380,961)
Loss from unconsolidated equity method investment	-	-	(135,692)	-
Realized foreign exchange gain/(loss)	8,774	-	(270,333)	-
Interest income	7,635	22	8,139	91
<b>Total other expense</b>	<b>(2,152)</b>	<b>(82,283)</b>	<b>(1,489,135)</b>	<b>(380,870)</b>
<b>Loss before income taxes</b>	<b>(3,142,667)</b>	<b>(111,107)</b>	<b>(6,209,971)</b>	<b>(2,728,970)</b>
<b>Income tax (expense)/benefit</b>				
Current	-	-	-	-
Deferred	-	-	-	-
<b>Total income tax (expense)/benefit</b>	<b>-</b>	<b>-</b>	<b>-</b>	<b>-</b>
<b>Net loss</b>	<b>(3,142,667)</b>	<b>(111,107)</b>	<b>(6,209,971)</b>	<b>(2,728,970)</b>
Net (loss) income attributable to non-controlling interest	(14,854)	1,485	(25,684)	(22,210)
<b>Net loss attributable to GBS, Inc.</b>	<b>\$ (3,127,813)</b>	<b>\$ (112,592)</b>	<b>\$ (6,184,287)</b>	<b>\$ (2,706,760)</b>
<b>Other comprehensive income</b>				
Foreign currency translation gain/(loss) attributable to non-controlling interest	-	-	-	-
Foreign currency translation gain/(loss) attributable to GBS, Inc.	(262,032)	100,921	(278,744)	(28,129)
<b>Total other comprehensive income</b>	<b>(262,032)</b>	<b>100,921</b>	<b>(278,744)</b>	<b>(28,129)</b>
<b>Comprehensive net loss attributable to GBS, Inc</b>	<b>\$ (3,389,845)</b>	<b>\$ (11,671)</b>	<b>\$ (6,463,031)</b>	<b>\$ (2,734,889)</b>
<b>Net loss per share, basic and diluted</b>	<b>\$ (0.27)</b>	<b>\$ (0.01)</b>	<b>\$ (0.64)</b>	<b>\$ (0.32)</b>
<b>Weighted average number of shares outstanding, basic and diluted</b>	<b>11,795,741</b>	<b>8,510,000</b>	<b>9,667,399</b>	<b>8,510,000</b>

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

**GBS Inc.**  
**CONDENSED CONSOLIDATED STATEMENTS OF CHANGES IN SHAREHOLDERS' EQUITY**  
**(Unaudited)**  
**(Amounts in \$)**

	<u>Preferred stock</u>		<u>Common stock</u>		<u>Additional paid in capital</u>	<u>Accumulated deficit</u>	<u>Other comprehensive (loss) income</u>	<u>Non-controlling interest</u>	<u>Total stockholders' equity (deficit)</u>
	<u>Shares</u>	<u>Amount</u>	<u>Shares</u>	<u>Amount</u>					
Balance, June 30, 2020	2,370,891	\$ 23,709	8,630,000	\$ 86,300	\$ 10,899,942	\$ (15,832,517)	\$ (363,951)	\$ (28,311)	\$ (5,214,828)
Issuance of convertible preferred shares	439,299	4,393	-	-	3,290,352	-	-	-	3,294,745
Foreign currency translation loss	-	-	-	-	-	-	(50,568)	-	(50,568)
Net loss	-	-	-	-	-	(1,072,510)	-	(4,405)	(1,076,915)
Balance, September 30, 2020	<u>2,810,190</u>	<u>28,102</u>	<u>8,630,000</u>	<u>86,300</u>	<u>14,190,294</u>	<u>(16,905,027)</u>	<u>(414,519)</u>	<u>(32,716)</u>	<u>(3,047,566)</u>
Issuance of common stock at initial public offering	-	-	1,270,589	12,706	21,587,307	-	-	-	21,600,013
Issuance cost of common stock at initial public offering	-	-	-	-	(3,867,565)	-	-	-	(3,867,565)
Cancellation of common stock in exchange for preferred shares	3,000,000	30,000	(3,000,000)	(30,000)	-	-	-	-	-
Conversion of convertible notes into common stock at initial public offering	-	-	710,548	7,105	5,126,601	-	-	-	5,133,706
Conversion of convertible preferred shares into common stock at initial public offering	(2,810,190)	(28,102)	2,810,190	28,102	-	-	-	-	-
Beneficial conversion feature	-	-	-	-	905,948	-	-	-	905,948
Series A warrants exercised to purchase common shares	-	-	1,200	12	10,188	-	-	-	10,200
Series A and B warrants acquired	-	-	-	-	3,812	-	-	-	3,812
Foreign currency translation loss	-	-	-	-	-	-	33,856	-	33,856
Net loss	-	-	-	-	-	(1,983,964)	-	(6,425)	(1,990,389)
Balance, December 31, 2020	<u>3,000,000</u>	<u>\$ 30,000</u>	<u>10,422,527</u>	<u>\$ 104,225</u>	<u>\$ 37,956,585</u>	<u>\$ (18,888,991)</u>	<u>\$ (380,663)</u>	<u>\$ (39,141)</u>	<u>\$ 18,782,015</u>
Series A warrants exercised to purchase common shares	-	-	58,600	586	497,514	-	-	-	498,100
Series B warrants exercised to purchase common shares	-	-	1,400,195	14,002	(14,002)	-	-	-	-
Foreign currency translation loss	-	-	-	-	-	-	(262,032)	-	(262,032)
Net loss	-	-	-	-	-	(3,127,813)	-	(14,854)	(3,142,667)
Balance, March 31, 2021	<u>3,000,000</u>	<u>\$ 30,000</u>	<u>11,881,322</u>	<u>\$ 118,813</u>	<u>\$ 38,440,097</u>	<u>\$ (22,016,804)</u>	<u>\$ (642,695)</u>	<u>\$ (53,995)</u>	<u>\$ 15,875,416</u>

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

	Preferred stock		Common stock		Additional paid in capital	Accumulated deficit	Other comprehensive (loss) income	Non-controlling interest	Total stockholders' equity (deficit)
	Shares	Amount	Shares	Amount					
Balance, June 30, 2019	2,064,884	\$ 20,649	8,510,000	\$ 85,100	\$ 8,164,804	(12,668,741)	(216,870)	637,919	\$ (3,977,139)
Reclassification of noncontrolling interest	-	-	-	-	637,056	-	-	(637,056)	-
Balance, June 30, 2019	2,064,884	20,649	8,510,000	85,100	8,801,860	(12,668,741)	(216,870)	863	(3,977,139)
Deemed dividend	-	-	-	-	(976,308)	-	-	-	(976,308)
Issuance of convertible preferred shares	259,007	2,590	-	-	1,939,964	-	-	-	1,942,554
Issuance costs for common and preferred shares	-	-	-	-	(116,402)	-	-	-	(116,402)
Foreign currency translation loss	-	-	-	-	-	-	4,234	-	4,234
Net loss	-	-	-	-	-	(757,850)	-	(6,980)	(764,830)
Balance, September 30, 2019	2,323,891	23,239	8,510,000	85,100	9,649,114	(13,426,591)	(212,636)	(6,117)	(3,887,891)
Foreign currency translation loss	-	-	-	-	-	-	(133,284)	-	(133,284)
Net loss	-	-	-	-	-	(1,836,318)	-	(16,715)	(1,853,033)
Balance, December 31, 2019	2,323,891	\$ 23,239	8,510,000	\$ 85,100	\$ 9,649,114	\$ (15,262,909)	\$ (345,920)	\$ (22,832)	\$ (5,874,208)
Foreign currency translation loss	-	-	-	-	-	-	100,921	-	100,921
Net loss	-	-	-	-	-	(112,592)	-	1,485	(111,107)
Balance, March 31, 2020	2,323,891	\$ 23,239	8,510,000	\$ 85,100	\$ 9,649,114	\$ (15,375,501)	\$ (244,999)	\$ (21,347)	\$ (5,884,394)

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

**GBS Inc.**  
**CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS**  
**(Unaudited)**  
**(Amount in \$)**

	<b>Nine Months Ended March 31,</b>	
	<b>2021</b>	<b>2020</b>
<b>Cash flows from operating activities:</b>		
Net loss	\$ (6,209,971)	\$ (2,728,970)
Adjustments to reconcile net loss to net cash used in operating activities:		
Loss on foreign currency translations (net)	(270,333)	-
Loss on investment in affiliate	135,692	-
Amortization of debt discount and issuance costs	-	2,359
Contingent beneficial conversion feature on convertible notes	905,948	-
Changes in operating assets and liabilities:		
Other receivables	-	118,056
Other current assets	(2,275,327)	94,274
Other non-current assets	(866,667)	-
Accounts payable	680,922	(403,305)
Accounts payable - related party	(1,732,058)	2,223,993
Other long-term liabilities	18,128	-
<b>Net cash used in operating activities</b>	<b>(9,613,666)</b>	<b>(693,593)</b>
<b>Cash flows from investing activities:</b>		
<b>Net cash used in investing activities</b>	<b>-</b>	<b>-</b>
<b>Cash flows from financing activities:</b>		
Proceeds from issuance of warrants	3,812	-
Proceeds from warrant holders for common shares	508,300	-
Proceeds from issuance of preferred stock	3,294,745	648,750
Proceeds from initial public offering	21,600,013	-
Payment of equity issuance costs	(2,003,952)	(116,402)
<b>Net cash provided by financing activities</b>	<b>23,402,918</b>	<b>532,348</b>
Effect of foreign exchange rates on cash and cash equivalents	45,097	(18,624)
Increase in cash and cash equivalents	13,834,349	(179,869)
Cash and cash equivalents, beginning of period	427,273	197,940
Cash and cash equivalents, end of period	<b>\$ 14,261,622</b>	<b>\$ 18,071</b>
<b>Non-cash investing and financing activities</b>		
Reclassification of deferred charges to additional paid in capital upon completion of initial public offering	\$ 1,863,613	\$ -
Conversion of notes to common shares at initial public offering	5,133,706	-
Conversion of preferred shares into common shares	28,102	-
<b>Supplemental disclosure of cash flow information:</b>		
Cash paid for income taxes	\$ -	\$ -
Cash paid for interest	\$ 185,301	\$ 249,627

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

**GBS Inc.**  
**NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS**  
**(Unaudited)**

**NOTE 1. ORGANIZATION AND DESCRIPTION OF THE BUSINESS**

GBS Inc. and its wholly owned subsidiary, GBS Operations Inc. were formed on December 5, 2016 under the laws of the state of Delaware. Glucose Biosensor Systems (Greater China) Pty Ltd (“GBSPL”) was formed on August 4, 2016 under the laws of New South Wales, Australia and was renamed to GBS (APAC) Pty Ltd on October 14, 2020. Glucose Biosensor Systems (Japan) Pty Ltd and GBS (APAC) Pty Ltd were formed under the laws of New South Wales, Australia on February 22, 2017 and February 23, 2017 respectively. These companies (collectively, the “Company” or “Group”) were formed to provide a non-invasive, pain free innovation to make it easier for people to manage diabetes using the Company’s Saliva Glucose Biosensor (“SGB” and, together with the software app that interfaces the SGB with the Company’s digital information system, the “SGT”).

GBS Inc. is a 48.7% owned (by voting rights) affiliate of Life Science Biosensor Diagnostics Pty Ltd (“LSBD”), an Australian company that owns the worldwide intellectual property rights to the biosensor platform from University of Newcastle, Australia. LSBD has licensed to the Company that technology to introduce and launch the platform in the Asia-Pacific Region (“APAC”). The Company will commence this process with the SGT.

On May 29, 2020 a research agreement was executed between LSBD and the Wyss Institute for Biologically Inspired Engineering at Harvard University (Wyss). The Company is not a legal party to the agreement but is expecting to derive a benefit through the Technology Transfer Agreement executed with LSBD and the Company on June 23, 2020, further details which are provided below. The Company has transferred biosensors (research materials) to the Wyss Institute where its research and development scientists have commenced a pilot research program. Since the biosensor architecture is complete and given the pre-existing plans to develop immunology diagnostic tests, it is therefore relatively straightforward and expeditious to develop the SARS-CoV-2 test.

SARS-CoV-2 antibody testing in saliva can play a critically important role in large-scale ‘sero’-surveillance to address key public health priorities and guide policy and decision-making for COVID-19. It is anticipated that FDA review will be under the Emergency Use Authorization program, which means expedited time to market.

On June 23, 2020, the Company entered into a Technology Transfer Agreement global license with LSBD. The significant terms of the license agreement are:

- The Company has the exclusive worldwide rights to a biosensor strip for antibodies against SARS-CoV-2 and associated application for reading devices to:
  - act as the authorized party for the purpose of processing the application of, and obtaining any, regulatory approval for the Licensed Product, including being authorized to process the approval for an investigational device required for the purpose of carrying out clinical studies;
  - manufacture, promote, market, import, offer, sell, and distribute the Licensed Products;
  - provide reasonable customer support services on the use of the Licensed Products to end users of, and health care practitioners referring end users to, the Licensed Products;
  - use the Licensed Products only for the purposes identified and permitted pursuant to regulatory approval; and
  - collect data acquired from the Licensed Products
- The royalty rate is 13%, based upon mutually agreed sales projections on the net sales of the commercial units and dedicated reading devices. This serves as the minimum royalty and falls to 3% at the expiry of the relevant patent(s)



- Each additional year, the sales upon which the minimum royalty is calculated on is increased by the mutually agreed Expected Market Growth rate plus an Additional Growth Percentage rate up to 7% annually. The Additional Growth Percentage Rate is calculated and applied for 10 years
- In the event of a dispute, in relation to the expected market growth or additional percentage, the agreement provides for a dispute resolution by an independent third party.

There are no milestone payments.

### **Initial public offering**

On December 28, 2020, the Company closed its initial public offering (“IPO”) and sold 1,270,589 units, consisting of (a) one share of the Company’s common stock (or, at the purchaser’s election, one share of Series B Convertible Preferred Stock), (b) one Series A warrant (the “Series A Warrants”) to purchase one share of the Company’s common stock at an exercise price equal to \$8.50 per share, exercisable until the fifth anniversary of the issuance date, and (c) one Series B warrant (the “Series B Warrants”) to purchase one share of the Company’s common stock at an exercise price equal to \$17.00 per share, exercisable until the fifth anniversary of the issuance date and subject to certain adjustment and cashless exercise provisions. The public offering price of the shares sold in the IPO was \$17.00 per unit. In aggregate, the units issued in the offering generated \$17,732,448 in net proceeds, which amount is net of \$1,714,001 in underwriters’ discount and commissions, and \$2,153,564 in offering costs. Offering costs include underwriters’ warrants to acquire up to 63,529 shares with an exercise price of \$18.70 per share, exercisable until the fifth anniversary of the issuance date. The Company also issued to the underwriter an option, exercisable one or more times in whole or in part. If, and to the extent, the Over Allotment Option was exercised, the underwriter may purchase up to 190,588 additional shares of common stock and/or Series A Warrants to purchase up to an aggregate of 190,588 shares of common stock and/or Series B Warrants to purchase up to an aggregate of 190,588 shares of common stock, in any combinations thereof, from us at the public offering price per security, less the underwriting discounts and commissions, for 45 days after the date of the IPO to cover over-allotments, if any (the “Over-Allotment Option”).

Upon the closing of the IPO, all shares of preferred stock then outstanding were automatically converted into 2,810,190 shares of common stock, and all convertible notes then outstanding were automatically converted into 710,548 shares of common stock.

Pre-IPO preferred shareholders were issued warrants following the Company’s completed IPO, that allow the holder to acquire 2,736,675 shares of common stock at the IPO price during year two through to year three following the completion of the IPO. At exercise date, the shareholder must hold, for each warrant to be exercised, the underlying common share to exercise the warrant. The warrants are not transferable and apply to the number of shares that were subscribed for.

### **NOTE 2. LIQUIDITY**

The Financial Accounting Standards Board (“FASB”) Accounting Standards Codification (“ASC”) Topic 205-40, *Presentation of Financial Statements - Going Concern* (ASC 205-40) requires management to assess an entity’s ability to continue as a going concern within one year of the date of the financial statements are issued. In each reporting period, including interim periods, an entity is required to assess conditions known and reasonably knowable as of the financial statement issuance date to determine whether it is probable an entity will not meet its financial obligations within one year from the financial statement issuance date. Substantial doubt about an entity’s ability to continue as a going concern exists when conditions and events, considered in the aggregate, indicate it is probable the entity will be unable to meet its financial obligations as they become due within one year after the date the financial statements are issued.

The Company is an emerging growth company and has not generated any revenues to date. As such, the Company is subject to all of the risks associated with emerging growth companies. Since inception, the Company has incurred losses and negative cash flows from operating activities. The Company does not expect to generate positive cash flows from operating activities in the near future until such time, if at all, the Company completes the development process of its products, including regulatory approvals, and thereafter, begins to commercialize and achieve substantial acceptance in the marketplace for the first of a series of products in its medical device portfolio.

The Company incurred a net loss of \$6,209,971 for the nine months ended March 31, 2021 (Net loss \$2,728,970 for the nine months ended March 31, 2020). As at March 31, 2021, the Company has shareholders' equity of \$15,875,416, working capital of \$15,026,877, and an accumulated deficit of \$(22,016,804).

On January 30, 2020, the International Health Regulations Emergency Committee of the World Health Organization (the "WHO") declared the novel coronavirus disease 2019 ("COVID-19") outbreak a public health emergency of international concern and on March 12, 2020 the WHO announced the outbreak was a pandemic. The COVID-19 pandemic is having a negative impact on global markets and business activity, which has had a limited impact on our core business operations. However, due to the nature of our platform technology we are able to quickly adapt to this rapidly evolving environment. As part of the immunology modality of the biosensor platform, LSB D executed an agreement on May 29, 2020 with the Wyss Institute for Biologically Inspired Engineering at Harvard University to use the biosensor platform to develop a COVID-19 rapid diagnostic test. The Company has the rights to the technology from this agreement under a Technology Transfer Agreement global license with LSB D entered into on June 23, 2020.

GBS Inc. is the global licensee and intends to commercialize COVID-19 diagnostic tests across the US, Europe, APAC and the rest of the world through appropriately qualified distributors.

In the near future, the Company anticipates incurring operating losses and does not expect to experience positive cash flows from operating activities and may continue to incur operating losses until it completes the development of its products and seeks regulatory approvals to market such products.

The Company's consolidated financial statements have been prepared on a going concern basis which contemplates the realization of assets and satisfaction of liabilities and commitments in the normal course of business. The consolidated financial statements do not include any adjustments relating to the recoverability and classification of recorded asset amounts or the amounts and classification of liabilities should the Company be unable to continue as a going concern.

As a result of the Company's initial public offering (see Note 1), the Company believes it has sufficient working capital to finance its operations for the next twelve months as such these consolidated financial statements are prepared on the going concern basis.

### **NOTE 3. SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES**

#### ***Basis of presentation***

The unaudited condensed consolidated financial statements of the Company have been prepared in accordance with accounting principles generally accepted in the United States ("U.S. GAAP") and pursuant to the requirements for reporting on Form 10-Q and Article 10 of Regulation S-X and, therefore, omit or condense certain footnotes and other information normally included in financial statements prepared in accordance with U.S. GAAP. In the opinion of management, the condensed consolidated financial statements reflect all adjustments and reclassifications that are necessary for the fair presentation of financial results as of and for the periods presented. The results of operations for an interim period may not give a true indication of the results for the entire year. The June 30, 2020 consolidated balance sheet has been derived from the audited financial statements as of that date.

These condensed consolidated financial statements have been derived from, and should be read in conjunction with, the Company's audited consolidated financial statements and notes thereto as of and for the year ended June 30, 2020 included in the Company's Registration Statement on Form S-1, File No. 333-252277 on file with the U.S. Securities and Exchange Commission (the "SEC"). There have not been any significant changes to the Company's significant accounting policies during the nine months ended March 31, 2021.

### ***Use of estimates***

The preparation of consolidated financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the consolidated financial statements, and the reported amounts of revenue and expenses during the reporting period. Actual results could materially differ from those estimates.

### ***Reclassifications***

Certain reclassifications have been made to prior periods to conform to current period presentation as described below.

In the comparative period (FY 2020), management determined that certain transactions involving the issuance of shares of its subsidiary that occurred during the prior year should have resulted in an adjustment to non-controlling interest (“NCI”) and Additional Paid-in-Capital (“APIC”) to reflect the difference between the fair value of the consideration received and the book value of NCI involving these changes in ownership. As a result, the Company increased its prior year APIC with an offsetting reduction to NCI of \$637,056. Management concluded that this reclassification was not meaningful to the Company’s financial position for the prior year, and as such, this change was recorded in the consolidated balance sheets and statements of shareholder’s equity in the first quarter of the comparative period (FY 2020) as an out-of-period adjustment.

For the three months ended March 31, 2020, amounts presented in the income statement reflect the difference between the nine months ended March 31, 2020 and the previously reported six months ended December 31, 2019 (Form 10Q for the quarter ended December 31, 2019). These quarterly balances was mainly impacted by a reclassification of \$268,457 in overhead reimbursements that has been reclassified from other income to general and administrative expenses for comparative reasons. The Company currently does not generate any revenue. The foreign currency translation gain was also adjusted by \$165,230 to total operating expenses for the same reason.

### ***Revenue recognition***

Revenue from contracts with customers is recognized when, or as, the Company satisfies its performance obligations by delivering the promised goods or service deliverables to the customers. A good or service deliverable is transferred to a customer when, or as, the customer obtains control of that good or service deliverable.

### ***Foreign currency translation***

Assets and liabilities of foreign subsidiaries are translated from local (functional) currency to reporting currency (U.S. dollar) at the rate of exchange in effect on the consolidated balance sheets date; income and expenses are translated at the average rate of exchange prevailing during the year. The functional currency of GBS Inc. is the United States dollar. Foreign currency movements resulted in a gain/(loss) of (\$262,032) and (\$278,744) for the three and nine months ended March 31, 2021, respectively and \$100,921 and (\$28,129) for the three and nine months ended March 31, 2020, respectively.

### ***Income taxes***

In accordance with the provisions of Financial Accounting Standards Board Accounting Standards Codification (“FASB ASC”) 740, *Income Taxes*, tax positions initially need to be recognized in the consolidated financial statements when it is more likely than not that the positions will be sustained upon examination by taxing authorities. It also provides guidance for de-recognition, classification, interest and penalties, accounting in interim periods, disclosure, and transition.

As of March 31, 2021, the Company had no uncertain tax positions that qualified for either recognition or disclosure in the consolidated financial statements. Additionally, the Company had no interest and penalties related to income taxes.

The Company accounts for current and deferred income taxes and, when appropriate, deferred tax assets and liabilities are recorded with respect to temporary differences in the accounting treatment of items for financial reporting purposes and for income tax purposes. Where, based on the weight of all available evidence, it is more likely than not that some amount of the recorded deferred tax assets will not be realized, a valuation allowance is established for that amount that, in management's judgment, is sufficient to reduce the deferred tax asset to an amount that is more likely than not to be realized.

#### ***Debt issuance cost***

Debt issuance costs are amortized using the effective interest rate method over the term of the loan and the amortization expense is recorded as part of interest expense of the consolidated statements of operations.

#### ***Licensing rights***

During the first quarter of the fiscal year ended June 30 2020, the Company purchased the license right procurement assets from LSB D for an amount of \$976,308 (June 30, 2019: \$ nil) in relation to the development and approval process for the Glucose Biosensor Technology. The Company recorded the license at the historical carrying value in the books of LSB D which was \$ nil and recorded the amount paid as a deemed dividend. The Company has agreed to pay royalties of sales & milestones payments as defined.

On July 3, 2019, the Company entered into an amended and restated license agreement. There is no set expiration date for the license. However, the exclusivity of the license granted under the license agreement runs until the expiration of the patent portfolio covered by the agreement which is currently until 2033. No royalties have been incurred through to March 31, 2021 (March 31, 2020: \$ nil).

On March 31, 2021, GBS entered into an agreement with LSB D to provide GBS an option to acquire an exclusive license to use LSB D's intellectual property in the Saliva Glucose Biosensor in North America (the "Option Agreement"). The Option Agreement has a term of two years and the exercise price for the option is \$5 million. The fee of \$0.5 million incurred for the option has been recognized as an expense and included within 'Development and regulatory approval expenses in the consolidated statements of operations.

#### ***Research and development costs***

During the quarter ended March 31, 2021, the Company contributed a total of \$2,600,000 towards budgeted development and commercialization costs to be incurred by BiosensX (North America) Inc. in which the Company has a 50% interest. This represents the Company's contribution towards budgeted development and commercialization costs included in total costs budgeted in the Form S-1. This funding relates to the development and preparation for submission of the Saliva Glucose Biosensor connected with regulatory approval for the U.S market by the U.S Food & Drug Administration. This amount is recognized as a prepayment and will be expensed as incurred over an estimated 18 month period in which the costs are expected to be incurred.

#### ***Net loss per share attributable to common shareholders ("EPS")***

The Company calculates earnings per share attributable to common shareholders in accordance with ASC Topic 260, *Earning Per Share*. Basic net income (loss) per share attributable to common shareholders is calculated by dividing net income (loss) attributable to common shareholders by the weighted-average number of common shares outstanding during the period. Diluted net income (loss) per common share is calculated by dividing net income (loss) attributable to common shareholders by weighted-average common shares outstanding during the period plus potentially dilutive common shares, such as share warrants.

Potentially dilutive common shares shall be calculated in accordance with the treasury share method, which assumes that proceeds from the exercise of all warrants are used to repurchase common share at market value. The number of shares remaining after the proceeds are exhausted represents the potentially dilutive effect of the securities.

As the Company has incurred net losses in all periods, certain potentially dilutive securities, including convertible preferred stock, warrants to acquire common stock, and convertible notes payable have been excluded in the computation of diluted loss per share as the effects are antidilutive.

### ***Recently issued but not yet effective accounting pronouncements***

As the Company is an emerging growth company, it has elected to defer the adoption of new accounting pronouncements until they would apply to private companies.

In August 2020, the FASB issued ASU 2020-06, which simplifies the guidance on the issuer's accounting for convertible debt instruments by removing the separation models for (1) convertible debt with a cash conversion feature and (2) convertible instruments with a beneficial conversion feature. As a result, entities will not separately present in equity an embedded conversion feature in such debt and will account for a convertible debt instrument wholly as debt, unless certain other conditions are met. The elimination of these models will reduce reported interest expense and increase reported net income for entities that have issued a convertible instrument that is within the scope of ASU 2020-06. Also, ASU 2020-06 requires the application of the if-converted method for calculating diluted earnings per share and treasury stock method will be no longer available. ASU 2020-06 is applicable for fiscal years beginning after December 15, 2021, with early adoption permitted no earlier than fiscal years beginning after December 15, 2020. The Company does not intend to early adopt and continues to evaluate the impact of the provisions of ASU 2020-06 on its consolidated financial statements.

In February 2016, the FASB issued ASU No. 2016-02, Leases ("ASU 2016-02"). This update requires all leases with a term greater than 12 months to be recognized on the balance sheet through a right-of-use asset and a lease liability and the disclosure of key information pertaining to leasing arrangements. This new guidance is effective for fiscal years beginning after December 15, 2021, and interim period within fiscal years beginning after December 15, 2022 as amended by ASU 2020-05 with early adoption permitted. The Company has not early adopted the standard and continues to evaluate the impact.

In December 2019, the FASB issued ASU No. 2019-12, Income Taxes (Topic 740): Simplifying the Accounting for Income Taxes ("ASU 2019-12"), which is intended to simplify various aspects of the accounting for income taxes. ASU 2019-12 removes certain exceptions to the general principles in Topic 740 and also clarifies and amends existing guidance to improve consistent application. This standard is effective for fiscal years and interim periods within those fiscal years, beginning after December 15, 2020. Early adoption is permitted. The Company has not early adopted the standard and continues to evaluate the impact.

### ***Concentration of credit risk***

The Company places its cash and cash equivalents, which may at times be in excess of the Australia Financial Claims Scheme or the United States' Federal Deposit Insurance Corporation insurance limits, with high credit quality financial institutions and attempts to limit the amount of credit exposure with any one institution.

### ***Related parties***

The Company has related party transactions with LSBD. See Notes 7 and 8.

### ***Fair value of financial instruments***

The carrying value of financial instruments classified as current assets and current liabilities approximate fair value due to their liquidity and short-term nature.

**NOTE 4. OTHER ASSETS**

Other current assets consist of the following:

	<u>March 31, 2021</u>	<u>June 30, 2020</u>
Goods and services tax receivable	\$ 89,566	\$ 7,509
Prepayments	2,227,221	29,469
Other receivables	7,602	12,084
Total	<u>\$ 2,324,389</u>	<u>\$ 49,062</u>

During the three months ended March 31, 2021, the Company made \$2,600,000 in prepayments relating to research and development contributions. Of the total prepayments, \$866,667 was recorded as a non-current asset as of March 31, 2021 based on the expected outflow of the budgeted research and development costs.

**NOTE 5. ACCOUNTS PAYABLE AND ACCRUED EXPENSES**

	<u>March 31, 2021</u>	<u>June 30, 2020</u>
Accounts and other payables	\$ 1,355,530	\$ 483,576
Accruals	41,810	56,894
Related party payables	37,235	1,769,293
Employee liabilities (current and non-current)	142,687	246,999
Total	<u>\$ 1,577,262</u>	<u>\$ 2,556,762</u>

**NOTE 6. CONVERTIBLE NOTES PAYABLE**

The Company's previously outstanding notes mandatorily converted, at a conversion price equal to 85% of 50% of the unit offering price of the IPO (or \$7.23), for an aggregate of 710,548 shares based on \$5,133,706 of principal and zero accrued interest outstanding at the date of conversion.

The convertible notes had a contingent Beneficial Conversion Features (BCF), with the contingency being the event of IPO. As such, a financing cost of \$905,948 was recognized as interest expense in the consolidated statements of operations and other comprehensive loss in relation to this contingent BCF during the nine months ended March 31, 2021.

**NOTE 7. SHAREHOLDERS' EQUITY***December 2020 Transactions*

On December 14, 2020, the Company agreed to issue to LSB D, in consideration of LSB D's contribution towards the research and development of applications other than glucose and COVID-19 applications to a maximum of \$2 million over a 5-year period, a 5-year non-transferable warrant to purchase 3,000,000 shares of the Company's common stock at the exercise price of \$17.00 per share. As this was a transaction between entities under common control, the \$2 million receivable due from LSB D has been recognized as contra-equity.

On December 18, 2020, the Company entered into an Exchange Agreement (the "EA") with LSB D to exchange 3,000,000 shares of its common stock held by LSB D for 3,000,000 shares of the Company's Series B Convertible Preferred Stock (the "Exchange"). In addition, the parties to the Exchange Agreement entered into a Registration Rights Agreement (the "RRA") pursuant to which the Company agreed to prepare and file within 30 days following the closing of the IPO with the Securities and Exchange Commission a registration statement to register for resale the shares of Common Stock issuable upon conversion of the Series B Convertible Preferred Stock. If and to the extent the Company fails to, among other things, file such resale registration statement or have it declared effective as required under the terms of the RRA, the Company will be required to pay to the holder of such registration rights partial liquidated damages payable in cash in the amount equal to the product of 1.0% multiplied by the aggregate purchase price paid by such holder pursuant to the EA. The EA and the RRA contain customary representations, warranties, agreements and, indemnification rights and obligations of the parties. The common stock acquired in the Exchange was immediately retired. Each share of Series B Convertible Preferred Stock is convertible into 1 share of the Company's common stock, subject to proportional adjustment and beneficial ownership limitations. In the event of the Company's liquidation, dissolution or winding up, holders of Series B Convertible Preferred Stock will participate pari passu with any distribution of proceeds to holders of the Company's common stock. Holders of Series B Convertible Preferred Stock are entitled to receive dividends on shares of Series B Preferred equal (on an as converted to common stock basis) to and in the same form as dividends actually paid on the Company's common stock. Shares of Series B Convertible Preferred Stock generally have no voting rights, except as required by law.

### ***Initial public offering***

In December 2020, the Company completed its initial public offering. For further details refer to Note 1.

### ***March 2021 Transactions***

During the quarter ended March 31, 2021, Series A and Series B warrants held by certain shareholders were exercised. Each warrant is convertible into 1 share of the Company's common stock. A total of 58,600 Series A warrants and 1,400,195 Series B warrants were exercised and converted into common stock.

### **NOTE 8. RELATED-PARTY TRANSACTIONS**

The Company completed certain financing transactions with LSB D as described in Note 7.

Sales to and purchases from related parties are made in arm's length transactions both at normal market prices and on normal commercial terms. The following transactions also occurred with LSB D during the period July 1, 2020 to March 31, 2021 (FY2020: July 1, 2019 to March 31, 2020):

The Company incurred a total of \$23,523 (FY2020: \$541,023) towards the services in connection with development and regulatory approval pathway for the technology, including payments made or expenses incurred on behalf of the Company.

The Company incurred a total of \$nil (FY2020: \$447,440) towards overhead cost reimbursement which includes salaries, rents and other related overheads directly attributable to the Company which are included in general and administration expenses.

The Company recognized income of \$nil (FY2020: \$119,744) in relation to shared labour reimbursement which includes salaries directly attributable to the Company which are included in shared-services revenue.

On March 31, 2021, GBS entered into an Option Agreement with LSB D to provide GBS the option to acquire an exclusive license for LSB D's intellectual property. A fee of \$500,000 was paid to acquire this option. For further details refer to Note 3.

During the quarter ended March 31, 2021, the Company contributed a total of \$2,600,000 towards budgeted development and commercialization costs to be incurred by BiosensX (North America) Inc. relating to the development and preparation for submission of the Saliva Glucose Biosensor connected with regulatory approval for the U.S market by the U.S Food & Drug Administration. For further details refer to Note 3.

### **NOTE 9. INVESTMENT IN AFFILIATE**

On May 29, 2020 LSB D, issued 14,000,000 common shares of BiosensX (North America) Inc. to the Company at par value of \$0.001 per share. This transaction provided the Company with a 50% interest in BiosensX (North America) Inc., the holder of the technology license for the North America region.

The investment in BiosensX (North America) Inc. is accounted for by use of the equity method in accordance with ASC 323 *Investments - Equity Method and Joint Ventures*.

At the date of this transaction, LSB D was the parent of both the Company and BiosensX (North America) Inc., the transfer of BiosensX shares to the Company was deemed to be a common control transaction. As a result of the share transfer, the Company has significant influence over BiosensX (North America) Inc. but in accordance with ASC 810 *Consolidation* LSB D is deemed to have control over BiosensX (North America) Inc. due to its direct ownership of 50% in BiosensX (North America) Inc. and indirect ownership of 50% in BiosensX (North America) Inc. through GBS Inc.

As of March 31, 2021, LSB D holds 48.7% of common Stock of GBS Inc. and therefore still has control over BiosensX (North America) Inc.

The following table summarizes the amount recorded in the consolidated financial statements:

	<b>March 31, 2021</b>	<b>June 30, 2020</b>
Investment value	\$ 135,692	\$ 14,000
(Loss) income from the affiliate	(135,692)	121,692
Carrying amount	<u>\$ -</u>	<u>\$ 135,692</u>

#### **NOTE 10. COMMITMENTS AND CONTINGENCIES**

On January 5, 2021, the Company entered into a certain Research Collaboration Agreement with Harvard College for the purposes of facilitating mutual collaboration in scientific research in connection with the Company's non-exclusive royalty free license to combat COVID-19 coronavirus. The contemplated collaboration includes research teams from the Company and Harvard and will include, among others, exchange of materials and research data, to now progress with the milestone of integrating the Harvard technology with the Company's biosensor with applications for SARS-Cov-2 antibody test for COVID-19. The Company agreed to pay Harvard a total amount of \$609,375 payable in 3 instalments of which \$304,686 remains payable as of March 31, 2021.

On January 21, 2021, the Company entered into a sponsored research agreement with Johns Hopkins Bloomberg School of Public Health to accelerate the development of next-generation saliva-based diagnostic tests. The Company is collaborating with the Bloomberg School of Public Health to optimise the collection of saliva and monitoring of diverse biomarkers across a number of modalities including clinical chemistry and infectious diseases. Johns Hopkins intend to utilise biosensor products to conduct in-field epidemiological studies. The Company agreed to pay Johns Hopkins a total amount of \$423,589 as a part of this sponsored research agreement of which all remains payable as of March 31, 2021.

The Company has no other material future minimum lease commitments or purchase commitments.

From time to time, the Company may become a party to various legal proceedings arising in the ordinary course of business. Based on information currently available, the Company is not involved in any pending or threatened legal proceedings that it believes could reasonably be expected to have a material adverse effect on its financial condition, results of operations or liquidity. However, legal matters are inherently uncertain, and the Company cannot guarantee that the outcome of any potential legal matter will be favorable to the Company.



**NOTE 11. LOSS PER SHARE**

Basic loss per common share is computed by dividing net loss allocable to common stockholders by the weighted average number of shares of common stock or common stock equivalents outstanding. Diluted loss per common share is computed similar to basic loss per common share except that it reflects the potential dilution that could occur if dilutive securities or other obligations to issue common stock were exercised or converted into common stock.

	Three Months Ended		Nine Months Ended	
	March 31, 2021	March 31, 2020	March 31, 2021	March 31, 2020
Net loss attributable to GBS, Inc.	\$ (3,127,813)	\$ (112,592)	\$ (6,184,287)	\$ (2,706,760)
Basic and diluted net loss per share attributed to common shareholders	\$ (0.27)	\$ (0.01)	\$ (0.64)	\$ (0.32)
Weighted-average number of ordinary shares	11,795,741	8,510,000	9,667,399	8,510,000

The following outstanding warrants, options and preferred shares were excluded from the computation of diluted net loss per share for the periods presented because their effect would have been anti-dilutive:

	Three Months Ended		Nine Months Ended	
	March 31, 2021	March 31, 2020	March 31, 2021	March 31, 2020
Warrants - Series A	1,401,377	-	1,401,377	-
Warrants - Series B	60,982	-	60,982	-
Warrants issued to underwriters	63,529	-	63,529	-
Pre IPO warrants	2,736,675	2,250,376	2,736,675	2,250,376
Warrants issued to parent entity	3,000,000	-	3,000,000	-
Preferred stock - Series A	-	2,323,891	-	2,323,891
Preferred stock - Series B	3,000,000	-	3,000,000	-

**NOTE 12. SUBSEQUENT EVENTS**

Subsequent to March 31, 2021 and through to the date of this filing, a total of 500,000 Series B Convertible Preferred Stock was converted into common stock. Each share of Series B Convertible Preferred Stock is convertible into 1 share of the Company's common stock.

Subsequent to March 31, 2021 and through to the date of this filing, a total of 800 Series B Warrants were exercised to purchase one Common Stock per Warrant in a cashless exercise provision as described in Company's Registration Statement on Form S-1, File No. 333-252277 on file with the U.S. Securities and Exchange Commission (the "SEC").

## ITEM 2. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULT OF OPERATIONS

*You should read the following discussion and analysis of our financial condition and results of operations together with our condensed consolidated financial statements and the related notes and other financial information included elsewhere in this Quarterly Report on Form 10-Q. This discussion and other parts of this report contain forward-looking statements. These forward-looking statements involve a number of risks, uncertainties (some of which are beyond our control) or other assumptions that may cause actual results or performance to be materially different from those expressed or implied by these forward-looking statements. These forward-looking statements involve a number of risks, uncertainties (some of which are beyond our control) or other assumptions that may cause actual results or performance to be materially different from those expressed or implied by these forward-looking statements. Our forward-looking statements include, but are not limited to, statements regarding our or our management team's expectations, hopes, beliefs, intentions or strategies regarding the future. In addition, any statements that refer to projections, forecasts or other characterizations of future events or circumstances, including any underlying assumptions, are forward-looking statements. The words "anticipate," "believe," "continue," "could," "estimate," "expect," "intends," "may," "might," "plan," "possible," "potential," "predict," "project," "should," "would" and similar expressions may identify forward-looking statements, but the absence of these words does not mean that a statement is not forward-looking. Factors that might cause or contribute to such forward-looking statements include, but are not limited to, those set forth in the Risk Factors section of the Company's registration statement and prospectus for the Company's initial public offering filed with the SEC. The following discussion should be read in conjunction with our financial statements and related notes thereto included elsewhere in this report.*

### Overview

We are a biosensor diagnostic technology company developing our SARS COV2 antibody ("COV2") test for the world market, the Saliva Glucose Biosensor ("SGB") for the Asia-Pacific region ("APAC") and have a 50% interest for the North America region. This the prelude to a biosensor platform comprising of biochemistry, immunology, tumor markers, hormones, and nucleic acid diagnostic modalities. We were incorporated under the laws of Delaware on December 5, 2016. Our headquarters are in New York. We were formed to provide a non-invasive, pain free innovation to make it easier for people to manage diabetes using the Company's SGB and, together with the software app that interfaces the SGB with the Company's digital information system, the "SGT").

We currently are a 48.7%-owned (by voting rights) affiliate of Life Science Biosensor Diagnostics Pty Ltd ("LSBD"), an Australian company that owns the worldwide intellectual property rights to the biosensor platform from University of Newcastle, Australia. LSBD has licensed to us that technology for us to introduce and launch the platform in the APAC Region. We will commence this process with the SGT.

Our objective is to introduce and launch a COV2 test globally and then the SGB, the second of our diagnostic tests that stem from the Biosensor Platform that we license, in the APAC Region. In the next four years we intend on developing the platform to its full capacity testing across the following diagnostic modalities: immunology, hormones, chemistry, tumour markers and nucleic acid tests.

We believe that the COVID-19 pandemic is likely to remain with us for many years. Development of an improved antibody assays to detect prior infection with SARS-CoV-2 has been identified as one of the top unmet needs in the ongoing COVID-19 pandemic response. Precise knowledge of SARS-CoV-2 infection at the individual level can potentially inform clinical decision-making, whereas at the population level, precise knowledge of prior infection, immunity, and attack rates (particularly asymptomatic infection) is needed to prioritize risk management decision-making about social distancing, treatments, and vaccination. If saliva can support measurements of both the presence of SARS-CoV-2 RNA26-28 as well as antibodies against SARS-CoV-2, this sample type could provide an important opportunity to monitor individual and population-level SARS-CoV-2 transmission, infection, and immunity dynamics over place and time.

We anticipate there to be 3 different applications for the foreseeable future:

1. Population Screening - SARS-CoV-2 antibody testing is urgently needed to estimate the incidence and prevalence of SARS-CoV-2 infection at the general population level. Precise knowledge of population immunity could allow government bodies to make informed decisions about how and when to relax stay-at-home directives and to reopen the economy.
2. Diagnosis – The COV2 Biosensor test can be used as a complement to the (RNA) virus detection tests for patients presenting late after symptoms onset to healthcare facilities and where virus detection tests are negative despite strong indications of infection. In addition, they can potentially be used for informing the decision on discharge of patients who recovered from SARS-CoV-2 infection but remain RNA-positive by RT-PCR for a long time after symptoms have subsided. The degree of protective immunity conferred by or correlated with the antibodies detected in subjects with past SARS-CoV-2 infection is still under investigation. Once this is clarified, the COV 2 antibody tests could be, together with the (RNA) direct virus detection, an essential tool in de-escalation strategies. Currently antibody tests are used for sero-epidemiological surveys and studies.
3. Post vaccination screening - To assess the degree of the elicited potent antigen-specific antibody responses, to COV2 vaccines when developed and administered to humans.

Based on a recent paper publicly available and authored by the team at Johns Hopkins Department of Environmental Health and Engineering, Bloomberg School of Public Health, results indicate it is feasible to accurately measure the salivary IgG response to identify individuals with a prior SARS-CoV-2 infection. A saliva-based approach could serve as a non-invasive approach for accurate and large-scale SARS-CoV-2 “sero”-surveillance.

A saliva antibody test can greatly increase the scale of testing—particularly among susceptible populations—compared to blood and could clarify population immunity and susceptibility to SARS-CoV-2. The team at John Hopkins further demonstrated in the laboratory that when saliva was collected  $\geq 10$  days post symptom onset, the anti-SARS-CoV-2 IgG assay detects SARS-CoV-2 infection with 100% sensitivity and 99% specificity. In addition, the team demonstrated that the temporal kinetics of SARS CoV-2-specific IgG responses in saliva are consistent with those observed in serum and indicate that most individuals seroconvert approximately 10 days after COVID-19 symptom onset or approximately two weeks post-presumed infection.

By utilizing the biosensor platform for detecting COV2 we expect to have lower detection limits, improve on sensitivity and specificity characteristics of current diagnostic methods, be able to provide real time results at the point of care and provide quantitative results as opposed to negative or positive which is how other POCT report the results.

Accurate and scalable point-of-care (POC) tests for the diagnosis of COVID-19 would increase the scope for diagnosis to be made in the community and outside the laboratory setting. They would have the potential to reduce the time to obtaining an actionable result, could support early identification of those with COVID-19 and could also support appropriate use of isolation resources, infection control measures, and recruitment into clinical trials of treatments.

We are progressing with the milestone of integrating Harvard University’s technology with our biosensor applications for SARS-Cov-2 antibody test for COVID-19 by entering on January 5, into a Research Collaboration Agreement with Harvard College for the purposes of facilitating mutual collaboration in scientific research in connection with the Company’s non-exclusive royalty free license to combat COVID-19 coronavirus. The contemplated collaboration includes research teams from the Company and Harvard and will include, among others, exchange of materials and research data.

Our Company has not generated any revenues to date. As such, the Company is subject to all of the risks associated with emerging growth companies. Since inception, the Company has incurred losses and negative cash flows from operating activities. The Company does not expect to generate positive cash flows from operating activities in the near future until such time, if at all, the Company completes the development process of its products, including regulatory approvals, and thereafter, begins to commercialize and achieve substantial acceptance in the marketplace for the first of a series of products in its medical device portfolio.

## ***Recent Developments***

### *December Quarter Developments*

On December 14, 2020, the Company agreed to issue to LSBDD, in consideration of LSBDD’s contribution towards the research and development of applications other than glucose and COVID-19 applications to a maximum of \$2 million over a 5-year period, a 5-year non-transferable warrant to purchase 3,000,000 shares of the Company’s common stock at the exercise price of \$17.00 per share.

On December 18, 2020, the Company entered into an Exchange Agreement (the “EA”) with LSB D to exchange 3,000,000 shares of its common stock held by LSB D for 3,000,000 shares of the Company’s Series B Convertible Preferred Stock. In addition, the parties to the Exchange Agreement entered into a Registration Rights Agreement (the “RRA”) pursuant to which the Company agreed to prepare and file within 30 days following the closing of the IPO with the Securities and Exchange Commission a registration statement to register for resale the shares of Common Stock issuable upon conversion of the Series B Convertible Preferred Stock. If and to the extent the Company fails to, among other things, file such resale registration statement or have it declared as required under the terms of the RRA, the Company will be required to pay to the holder of such registration rights partial liquidated damages payable in cash in the amount equal to the product of 1.0% multiplied by the aggregate purchase price paid by such holder pursuant to the EA. The EA and the RRA contain customary representations, warranties, agreements and, indemnification rights and obligations of the parties.

On December 18, 2020, LSB D entered into a certain Purchase and Assignment Agreement (the “PAA”) with an institutional accredited investor (the “Purchaser”) pursuant to which LSB D sold and assigned to the Purchaser 3,000,000 shares of the Series B Convertible Preferred Stock and assigned to the Purchaser its rights under the EA and the RRA with respect to such preferred shares for a total purchase price of \$2,000,000. The investor’s Series B Convertible Preferred Stock is convertible into 3,000,000 shares of the Company’s common stock, subject to beneficial ownership limitation.

#### *March Quarter Developments*

##### *Point-of-Care Test Commercialization Ecosystem Established*

- Received approval from the Harvard Longwood campus Institutional Review Board (IRB) to commence a validation study to test clinical samples from a COVID-19 repository and to commence clinical studies on the COVID-19 Antibody Biosensor;
- Onboarded and aligned with world-class institutions, Johns Hopkins University, The Wyss Institute for Biologically Inspired Engineering, and the University of Newcastle for the development of saliva-based POCTs for both glucose monitoring and COVID-19 antibody detection;
- Onboarded new top-tier members to GBS’s scientific team to formulate and execute its commercialization plan.

#### *COVID-19 Key Developments*

- Commenced research protocols with The Wyss Institute for Biologically Inspired Engineering to progress with the milestone of integrating this technology with the Company’s Biosensor for SARS-Cov-2 antibody tests;
- Initiated study for the salivary collection protocol with Johns Hopkins University, Bloomberg School of Public Health;
- Completed technical optimization of the Wyss’s eRapid assay performance in relation to SARS-Cov-2 antibody detection at The Wyss Institute to align with the fastest antibody tests currently on market using clinical samples.

#### *Glucose Key Developments*

- Developing a clinical plan for regulatory submission and subsequent approval with Precision Medicine Architects, LLC;
- Commenced global voice of customer survey with Precision Medicine Architects, LLC as part of the process to finalize product development of the device and usability;
- Further development of prototyping for middleware and smart phone application;
- Executed option agreement to acquire the rights to use, make, market, sell and offer to sell Products under the Intellectual Property Rights in the Glucose Field in the North American market for the Saliva Glucose Biosensor .

During the quarter ended March 31, 2021, Series A and Series B warrants held by certain shareholders were exercised. Each warrant is convertible into 1 share of the Company's common stock. A total of 58,600 Series A warrants and 1,400,195 Series B warrants were exercised and converted into common stock.

On March 31, 2021, GBS entered into an agreement with LSB D to provide GBS an option to acquire an exclusive license to use LSB D's intellectual property in the treatment or management of diabetes field in North America (the "Option Agreement"). The Option Agreement has a term of two years and the exercise price for the option is \$5 million.

Subsequent to March 31, 2021, a total of 500,000 Series B Convertible Preferred Stock was converted into common stock. Each share of Series B Convertible Preferred Stock is convertible into 1 share of the Company's common stock as described in the Company's Registration Statement on Form S-1, File No. 333-242277 with the U.S. Securities and Exchange Commission.

### ***Initial public offering & share structure***

On December 28, 2020, the Company closed its initial public offering ("IPO") and sold 1,270,589 units, consisting of (a) one share of the Company's common stock (or, at the purchaser's election, one share of Series B Convertible Preferred Stock), (b) one Series A warrant (the "Series A Warrants") to purchase one share of the Company's common stock at an exercise price equal to \$8.50 per share, exercisable until the fifth anniversary of the issuance date, and (c) one Series B warrant (the "Series B Warrants") to purchase one share of the Company's common stock at an exercise price equal to \$17.00 per share, exercisable until the fifth anniversary of the issuance date and subject to certain adjustment and cashless exercise provisions. The public offering price of the shares sold in the IPO was \$17.00 per unit. In aggregate, the units issued in the offering generated \$17,732,448 in net proceeds, which amount is net of \$1,714,001 in underwriters' discount and commissions, and \$2,153,564 in offering costs. The Company also issued to the underwriter an option, exercisable one or more times in whole or in part, to purchase up to 190,588 additional shares of common stock and/or Series A Warrants to purchase up to an aggregate of 190,588 shares of common stock and/or Series B Warrants to purchase up to an aggregate of 190,588 shares of common stock, in any combinations thereof, from us at the public offering price per security, less the underwriting discounts and commissions, for 45 days after the date of the IPO to cover over-allotments, if any (the "Over-Allotment Option").

Upon the closing of the IPO, all shares of preferred stock then outstanding were automatically converted into 2,810,190 shares of common stock, and all convertible notes then outstanding were automatically converted into 710,548 shares of common stock.

Certain preferred shareholders were issued warrants that, following the Company's completed IPO, allow the holder to acquire 2,736,675 shares of common stock at the IPO price during years two through three following the IPO. At exercise date, the shareholder must hold for each warrant to be exercised, one underlying common share to exercise the option. The warrants are not transferable and apply to the number of shares that were subscribed for.

Accordingly, the share structure as of May 12, 2021 are as follows:

- 12,382,122 of Issued Common Stock
- 1,401,377 of Series A warrant exercisable at \$8.50
- 60,182 of Series B warrants exercisable at \$17 (subject to a cashless exercise provision)
- 63,529 of Warrants issued to the underwriter exercisable at \$18.70
- 2,736,675 of the Pre-IPO Warrants exercisable at \$8.50 (during year two through to year three after the IPO)
- 3,000,000 Warrants issued to LSB D exercisable at \$17
- 2,500,000 Preferred Stock-Series B

## Results of Operations:

### Comparison of the Three and Nine Months Ended March 31, 2021 and 2020

#### Revenue

##### *Government support income*

Government support income increased by \$34,290 to \$34,290 from \$0 for the three months ended March 31, 2021 compared to same period in 2020. This increase was primarily attributable to GBS Inc.'s subsidiary companies receiving COVID-19 related government support in the current period where the companies are located (The purpose of the grant is to support companies in managing its business and payroll costs during the COVID-19 pandemic).

Government support income increased by \$372,754 to \$372,754 from \$0 for the nine months ended March 31, 2021 compared to same period in 2020. This increase was primarily attributable to GBS Inc.'s subsidiary companies receiving Research and Development tax incentives. The purpose of the grant is to incentivize companies with their research and development related activities and other COVID-19 related government support in the current period where the companies are located.

##### *Shared service*

Shared service revenue was \$0 and \$(1,533) for the three months ended March 31, 2021 and 2020, respectively, and \$0 and \$119,744 for the nine months ended March 31, 2021 and 2020, respectively. Shared service revenue is mainly attributable to the recovery of costs from related parties. There were no shared services in the current period.

#### Operating expenses

##### *General and administrative expenses*

General and administrative expenses increased by \$959,547 to \$1,013,389 from \$53,842 for the three months ended March 31, 2021 compared to the same period in 2020. This increase was primarily driven by an increase in operational activities following completion of the IPO in the current period (December 2020).

General and administrative expenses increased by \$453,660 to \$2,205,842 from \$1,752,182 for the nine months ended March 31, 2021 compared to the same period in 2020. This increase was attributable to an increase in operational activities following completion of the IPO in the current period (December 2020).

As the Company's operating activities increase, we expect its general and administrative costs will include additional costs in overhead contribution, consultancy, and travel expenses.

##### *Development and regulatory expenses*

Development and regulatory expenses increased by \$2,215,141 to \$2,156,316 from \$(58,825) for the three months ended March 31, 2021 compared to the same period in 2020. This increase is primarily driven by funding availability since completion of the IPO in December 2020 that has allowed the Company to progress on its milestones, as well as a \$500,000 option that the Company expensed on the basis that there is no FDA approval of the intellectual property held by LSBDD.

Development and regulatory expenses increased by \$1,988,051 to \$2,529,074 from \$541,023 for the nine months ended March 31, 2021 compared to the same period in 2020. This increase is primarily driven by funding availability since completion of the IPO in December 2020 that has allowed the Company to progress on its milestones, as well as a \$500,000 option that the Company expensed on the basis that there is no FDA approval of the intellectual property held by LSBDD.

As the Company's operating activities increase, we expect its development and regulatory expenses to increase in future periods.

*Prospectus and capital raising expenses*

Prospectus and capital raising expenses decreased by \$27,174 to \$5,100 from \$32,274 for the three months ended March 31, 2021 as compared to the same period in 2020. This decrease was attributable to minimal expenditures required by us in the current period having completed our IPO in December 2020.

Prospectus and capital raising expenses increased by \$184,035 to \$358,674 from \$174,639 for the nine months ended March 31, 2021 as compared to the same period in 2020, respectively. This increase was attributable to majority of final expenditures required by us in the current period to successfully complete the IPO in December 2020.

***Other income and expenses***

*Interest expense*

Interest expense decreased \$63,744 to \$18,561 from \$82,305 for the three months ended March 31, 2021 as compared to the same period in 2020. This decrease was attributable to the conversion of convertible notes into common stock at completion of the IPO.

Interest expense increased \$710,288 to \$1,091,249 from \$380,961 for the nine months ended March 31, 2021 as compared to the same period in 2020. This increase was attributable to the non-cash recognition of a beneficial conversion feature associated with convertible notes, offset by conversion of convertible notes into common stock that occurred at IPO.

*Loss (income) from unconsolidated equity method investment*

Loss (income) from unconsolidated equity method investment was \$0 for the three months ended March 31, 2021 and 2020, respectively.

Loss from unconsolidated equity method investment increased \$135,692 to \$135,692 from \$0 for the nine months ended March 31, 2021 compared to the same period in 2020. This increase was attributable to the reduction in the carrying amount of its investment in BiosensX (North America) Inc.

*Realized foreign exchange gain (loss)*

Realized foreign exchange gain was \$8,774 for the three months ended March 31, 2021 compared to \$0 for the same period in 2020. This increase was attributable to the favorable foreign exchange translations upon settling payments in foreign currency.

Realized foreign exchange loss was \$270,333 for the nine months ended March 31, 2021 compared to \$0 the same period in 2020. This increase was largely attributable to the unfavorable foreign exchange translations on capital raisings from AUD to USD.

***Income tax (expense) benefit***

There was no income tax expense for the three and nine months ended March 31, 2021 and 2020 as the Company has established a full valuation allowance for all of its deferred tax assets.

## ***Other comprehensive income***

### ***Foreign currency translation gain/(loss)***

Unrealized foreign currency translation gain/(loss) increased by (\$362,953) to (\$262,032) from \$100,921 for the three months ended March 31, 2021 as compared to the same period in 2020. It is calculated based on the Company's unsettled transactions and balances in currencies other than its functional currency.

Unrealized foreign currency translation gain/(loss) increased by (\$250,615) to (\$278,744) from (\$28,129) for the nine months ended March 31, 2021 and 2020, respectively. It is calculated based on the Company's unsettled transactions and balances in currencies other than its functional currency.

### ***Net loss***

Net loss increased by \$3,031,560 to \$3,142,667 from \$111,107 for the three months ended March 31, 2021 compared to the same period in 2020. This overall increase was largely attributable to further progression on regulatory and development milestones and increased expenditure on general and administrative expenses with funding secured by the IPO. Further contributing to this movement was an option fee of \$500,000 to acquire an exclusive license for LSBD's intellectual property, \$268,457 in overhead reimbursements reversed, as well as changes in the foreign currency exchange rate between AUD and USA due to COVID-19, both within the same period in 2020.

Net loss increased by \$3,481,001 to \$6,209,971 from \$2,728,970 for the nine months ended March 31, 2021 compared to the same period in 2020. This overall increase was largely attributable to the non-cash recognition of a beneficial conversion feature, an option fee to acquire an exclusive license for LSBD's intellectual property and increased expenditure on general and administrative expenses and further progression on regulatory and development milestones with funding secured by the IPO.

## **Liquidity and Capital Resources**

Since our inception, our operations have primarily been financed through the issuance of our common stock, redeemable convertible preferred stock and the incurrence of debt. As of March 31, 2021, we had \$14,261,622 in cash and cash equivalents and \$15,026,877 in working capital.

See "Initial public offering" herein for details about our IPO.

According to our management's estimates, based on our budget and proposed schedules of development, approvals and organization, we believe, although there can be no assurances, that after our IPO we will have sufficient capital resources to enable us to continue to implement our business plan and remain in operation for at least 30 months. During this time, we expect to use the net proceeds available to us for the following purposes:

- to obtain regulatory approvals and establish manufacturing capacities necessary for marketing of the SGT;
- to market the SGT and establish a distribution network in the APAC Region; and
- for working capital and general corporate purposes.

We do not anticipate generating any revenue commencing in the vicinity of 6-10 months from the date of this report, if at all, and our revenues will not immediately be sufficient to finance our ongoing operations. In addition, available resources may be consumed more rapidly than currently anticipated, and there can be no assurance that we will be successful in developing the SGT and generating sufficient revenue in the timeframe set forth above, or at all. We may be unable to meet our targets for regulatory approval and market launch, or we may be unable to generate anticipated amounts of revenue from sales of the system. We may also need additional funding for developing new products and services and for additional sales, marketing and promotional activities. Should this occur, we may need to seek additional capital earlier than anticipated.



In the event we require additional capital, there can be no assurances that we will be able to raise such capital on acceptable terms, or at all. Failure to generate sufficient revenues or raise additional capital through debt or equity financings, or through collaboration agreements, strategic alliances or marketing and distribution arrangements, could have a material adverse effect on our ability to meet our long-term liquidity needs and achieve our intended long-term business plan. Our failure to obtain such funding when needed could create a negative impact on our stock price or could potentially lead to a reduction in our operations or the failure of our company.

### **Contractual Obligations**

We are a smaller reporting company as defined by Rule 12b-2 of the Exchange Act and are not required to provide the information otherwise required under this item.

### **Extended Transition Period for “Emerging Growth Companies”**

We have elected to use the extended transition period for complying with new or revised accounting standards under Section 102(b)(1) of the JOBS Act. This election allows us to delay the adoption of new or revised accounting standards that have different effective dates for public and private companies until those standards apply to private companies. As a result of this election, our financial statements may not be comparable to companies that comply with public company effective dates. Because our financial statements may not be comparable to companies that comply with public company effective dates, investors may have difficulty evaluating or comparing our business, performance or prospects in comparison to other public companies, which may have a negative impact on the value and liquidity of our common stock.

### **Off-Balance Sheet Arrangements**

Through March 31, 2021, we have not entered into any off-balance sheet arrangements as defined by applicable SEC regulations.

### **Critical Accounting Policies, Significant Judgments and Use of Estimates**

Our financial statements have been prepared in accordance with U.S. generally accepted accounting principles, or GAAP. The preparation of these financial statements requires us to make estimates and assumptions that affect the reported amounts of assets and liabilities, the disclosure of contingent assets and liabilities at the date of the financial statements and the reported expenses incurred during the reporting periods. Our estimates are based on our historical experience and on various other factors that we believe are reasonable under the circumstances, the results of which form the basis for making judgments about the carrying value of assets and liabilities that are not readily apparent from other sources. Actual results may differ from these estimates under different assumptions or conditions. We believe that the accounting policies discussed below are critical to understanding our historical and future performance, as these policies relate to the more significant areas involving management’s judgments and estimates.

Our critical accounting policies are described under the notes to the unaudited condensed consolidated financial statements included in “Part I, Item 1 — Financial Statements” of this Quarterly Report on Form 10-Q. During the nine months ended March 31, 2021, except as described in Note 3 to the unaudited interim condensed financial statements appearing elsewhere in this Quarterly Report on Form 10-Q, there were no material changes to our critical accounting policies from those discussed in our final prospectus filed on December 18, 2020.

## Recent Accounting Pronouncements

See “Recent Accounting Pronouncements” in Note 3 to our consolidated financial statements included elsewhere in this Quarterly Report on Form 10-Q for additional information.

## ITEM 3. QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK

We are a smaller reporting company as defined by Rule 12b-2 of the Exchange Act and are not required to provide the information otherwise required under this item.

## ITEM 4. CONTROLS AND PROCEDURES

### *Evaluation of Disclosure Controls and Procedures*

Our management, with the participation of our Chief Executive Officer and Chief Financial Officer, evaluated the effectiveness of our disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) as of the end of the period covered by this Quarterly Report on Form 10-Q, and have concluded that, based on such evaluation, our disclosure controls and procedures were not effective due to the material weakness in our internal control over financial reporting as of March 31, 2021 as described below.

Notwithstanding the conclusion that our disclosure controls and procedures were not effective as of the end of the period covered by this report, we believe that our consolidated financial statements and other information contained in this quarterly report present fairly, in all material respects, our business, financial condition and results of operations for the interim periods presented.

### **Material Weakness**

The Company completed the IPO in December 2020. Prior to the IPO, the Company was a private corporation with limited accounting personnel and other supervisory resources necessary to adequately execute its accounting processes and address its internal controls over financial reporting requirements. As a result, previously existing internal controls are no longer sufficient, and the Company is in the process of updating these controls. The design and implementation of internal control over financial reporting for the Company’s post-IPO has required and will continue to require significant time and resources from management and other personnel.

As part of this updating process, our management identified a material weakness in its internal control over financial reporting. A material weakness is a deficiency, or combination of deficiencies, in internal control over financial reporting, such that there is a reasonable possibility that a material misstatement of our annual or interim financial statements will not be prevented or detected on a timely basis. The material weakness identified relates to the fact that the Company has not yet designed and maintained an effective control environment commensurate with its financial reporting requirements, including a) has not yet completed the formally documented policies and procedures with respect to the review, supervision and monitoring of the Company’s accounting and reporting functions and b) lack of evidence to support the performance of controls and the adequacy of review procedures, including the completeness and accuracy of information used in the performance of controls.

### **Remediation Plan**

Management is committed to continuing with the steps necessary to remediate the control deficiencies that constituted the above material weakness. During 2021, we made the following enhancements to our control environment:

- a. We added accounting and finance personnel to provide additional individuals to allow for segregation of duties in the preparation and review of schedules, calculations, and journal entries that support financial reporting, to provide oversight, structure and reporting lines, and to provide additional review over our disclosures;
- b. We enhanced our controls to improve the preparation and review over complex accounting measurements, and the application of GAAP to significant accounts and transactions, and our financial statement disclosures; and,
- c. We are in the process of engaging outside consultants to assist us in our evaluation of the design, implementation, and documentation of internal controls that address the relevant risks, and that provide for appropriate evidence of performance of our internal controls (including completeness and accuracy procedures).

Under the direction of the audit committee of the board of directors, management will continue to take measures to remediate the material weakness in 2021. As such, we will continue to enhance corporate oversight over process-level controls and structures to ensure that there is appropriate assignment of authority, responsibility, and accountability to enable remediation of our material weakness. We believe that our remediation plan will be sufficient to remediate the identified material weakness and strengthen our internal control over financial reporting.

As we continue to evaluate, and work to improve, our internal control over financial reporting, management may determine that additional measures to address control deficiencies or modifications to the remediation plan are necessary.

### ***Inherent Limitation on the Effectiveness of Internal Controls***

The effectiveness of any system of internal control over financial reporting, including ours, is subject to inherent limitations, including the exercise of judgment in designing, implementing, operating, and evaluating the controls and procedures, and the inability to eliminate misconduct completely. Accordingly, in designing and evaluating the disclosure controls and procedures, management recognizes that any system of internal control over financial reporting, including ours, no matter how well designed and operated, can only provide reasonable, not absolute assurance of achieving the desired control objectives. In addition, the design of disclosure controls and procedures must reflect the fact that there are resource constraints and that management is required to apply its judgment in evaluating the benefits of possible controls and procedures relative to their costs. Moreover, 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. We intend to continue to monitor and upgrade our internal controls as necessary or appropriate for our business, but cannot assure you that such improvements will be sufficient to provide us with effective internal control over financial reporting.

***Changes in Internal Controls over Financial Reporting***

Except as noted above, there were no changes in our internal control over financial reporting that occurred during the three months ended March 31, 2021 that have materially affected, or are reasonably likely to materially affect, our internal control over financial reporting.

## PART II—OTHER INFORMATION

### ITEM 1. LEGAL PROCEEDINGS.

From time to time, we may be subject to legal proceedings and claims arising in the ordinary course of business. We are not currently engaged in any material legal proceedings.

### ITEM 1A. RISK FACTORS

We are a smaller reporting company as defined by Rule 12b-2 of the Exchange Act and are not required to provide the information otherwise required under this item.

### ITEM 2. UNREGISTERED SALES OF EQUITY SECURITIES AND USE OF PROCEEDS.

There were no unregistered sales of equity securities during the period.

On December 28, 2020, the Company closed its initial public offering (“IPO”) and sold 1,270,589 units, consisting of (a) one share of the Company’s common stock (or, at the purchaser’s election, one share of Series B Convertible Preferred Stock), (b) one Series A warrant (the “Series A Warrants”) to purchase one share of the Company’s common stock at an exercise price equal to \$8.50 per share, exercisable until the fifth anniversary of the issuance date, and (c) one Series B warrant (the “Series B Warrants”) to purchase one share of the Company’s common stock at an exercise price equal to \$17.00 per share, exercisable until the fifth anniversary of the issuance date and subject to certain adjustment and cashless exercise provisions. The public offering price of the shares sold in the IPO was \$17.00 per unit. In aggregate, the units issued in the offering generated \$17,732,448 in net proceeds, which amount is net of \$1,714,001 in underwriters’ discount and commissions, and \$2,153,564 in offering costs (including deferred equity offering cost of \$1,863,612). The Company also issued to the underwriter an option, exercisable one or more times in whole or in part, to purchase up to 190,588 additional shares of common stock and/or Series A Warrants to purchase up to an aggregate of 190,588 shares of common stock and/or Series B Warrants to purchase up to an aggregate of 190,588 shares of common stock, in any combinations thereof, from us at the public offering price per security, less the underwriting discounts and commissions, for 45 days after the date of the IPO to cover over-allotments, if any (the “Over-Allotment Option”).

Upon the closing of the IPO, all shares of preferred stock then outstanding were automatically converted into 2,810,190 shares of common stock, and all convertible notes then outstanding were automatically converted into 710,548 shares of common stock. Certain of the preferred shareholders were issued warrants that, following the Company’s completed IPO, allow the holder to acquire 2,736,675 shares of common stock at the IPO price during years two through three following the IPO. At exercise date, the shareholder must hold for each warrant to be exercised, one underlying common share to exercise the option. The warrants are not transferable and apply to the number of shares that were subscribed for.

There has been no material change in the planned use of proceeds from our IPO as described in our final prospectus filed with the SEC on December 28, 2020 pursuant to Rule 424(b). No direct or indirect payments were made by us to any of our directors or officers or their associates, to persons owning ten percent or more of our common stock or to their associates, or to our affiliates, other than payments in the ordinary course of business to officers for salaries. Pending the uses described, we intend to invest the net proceeds in short-term, interest-bearing obligations, investment-grade instruments, certificates of deposit or direct or guaranteed obligations of the U.S. government.

**ITEM 3. DEFAULTS UPON SENIOR SECURITIES.**

Not applicable.

**ITEM 4. MINE SAFETY DISCLOSURES.**

Not applicable.

**ITEM 5. OTHER INFORMATION.**

Not applicable.

**ITEM 6. EXHIBITS.**

<b>Exhibit No.</b>	<b>Description</b>
<b>31.1</b>	<a href="#"><u>Certification of Chief Executive Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.*</u></a>
<b>31.2</b>	<a href="#"><u>Certification of Principal Executive and Principal Financial Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.*</u></a>
<b>32.1</b>	<a href="#"><u>Certification of the Principal Executive Officer pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.</u></a>
<b>32.2</b>	<a href="#"><u>Certification of the Principal Financial Officer pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.</u></a>
<b>101.INS</b>	XBRL Instance Document.
<b>101.SCH</b>	XBRL Taxonomy Extension Schema Document.
<b>101.CAL</b>	XBRL Taxonomy Extension Calculation Linkbase Document.
<b>101.DEF</b>	XBRL Taxonomy Extension Definition Linkbase Document.
<b>101.LAB</b>	XBRL Taxonomy Extension Label Linkbase Document.
<b>101.PRE</b>	XBRL Taxonomy Extension Presentation Linkbase 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.

**GBS Inc.**

Date: May 13, 2021

By: /s/ Harry Simeonidis  
HARRY SIMEONIDIS  
CHIEF EXECUTIVE OFFICER AND PRESIDENT  
(Principal Executive Officer)

Date: May 13, 2021

By: /s/ Spiro Sakiris  
SPIRO SAKIRIS  
CHIEF FINANCIAL OFFICER  
(Principal Financial Officer)

OFFICER'S CERTIFICATION PURSUANT TO  
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Harry Simeonidis, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of GBS, Inc.;
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 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 fourth 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 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.

May 13, 2021

By: /s/ Harry Simeonidis

Harry Simeonidis, Chief Executive Officer  
(Principal Executive Officer)

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OFFICER'S CERTIFICATION PURSUANT TO  
SECTION 302 OF THE SARBANES-OXLEY ACT OF 2002

I, Spiro Sakiris, certify that:

1. I have reviewed this Quarterly Report on Form 10-Q of GBS, Inc.;
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 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:
  - e. 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;
  - f. 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;
  - g. 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
  - h. 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 fourth 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 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.

May 13, 2021

By: /s/ Spiro Sakiris

Spiro Sakiris, Chief Financial Officer  
(Principal Financial and Accounting Officer)

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

In connection with the Quarterly Report on Form 10-Q for the quarter ended March 31, 2021 of GBS, Inc. (the "Company") as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Harry Simeonidis, Chief Executive Officer of the Company, hereby certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

- The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934 (15 U.S.C 78m or 78o(d)); and
- The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: May 13, 2021

By: */s/ Harry Simeonidis*

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Harry Simeonidis  
Chief Executive Officer  
(Principal Executive Officer)

A signed original of this written statement required by Section 906 has been provided to GBS, Inc. and will be retained by GBS, Inc. and furnished to the Securities and Exchange Commission or its staff upon request.

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

In connection with the Quarterly Report on Form 10-Q for the quarter ended March 31, 2021 of GBS, Inc. (the "Company") as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Spiro Sakiris, Chief Executive Officer of the Company, hereby certify, pursuant to 18 U.S.C. Section 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that:

- The Report fully complies with the requirements of section 13(a) or 15(d) of the Securities Exchange Act of 1934 (15 U.S.C 78m or 78o(d)); and
- The information contained in the Report fairly presents, in all material respects, the financial condition and results of operations of the Company.

Date: May 13, 2021

By: /s/ Spiro Sakiris

Spiro Sakiris  
Chief Financial and Accounting Officer  
(Principal Executive Officer)

A signed original of this written statement required by Section 906 has been provided to GBS, Inc. and will be retained by GBS, Inc. and furnished to the Securities and Exchange Commission or its staff upon request.

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