(Mark One)

UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

WASHINGTON, D.C. 20549

FORM 10-Q

\boxtimes	Quarterly Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934 For the Quarterly Period Ended September 30, 2022 Or									
	□ Transition Report Pursuant to Section 13 or 15(d) of the Securities Exchange Act of 1934 For the transition period from to									
Commis	sion File No. 0-23047									
	SIG	A Technologies,	Inc.							
	(Exact nam	ne of registrant as specified in	its charter)							
	Delaware 13-3864870									
	(State or other jurisdiction of incorporation or organiza	ation)	(IRS Employer Identification No.)							
	31 East 62nd Street		10065							
	New York, NY		(zip code)							
	(Address of principal executive offices)									
	Registrant's telepho	one number, including area co	de: (212) 672-9100							
Securitie	s 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, \$.0001 par value	SIGA	The Nasdaq Global Market							
	by check mark whether the registrant (1) has filed all reports reas (or for such shorter period that the registrant was required to	-								
	by check mark whether the registrant has submitted electronical 5 of this chapter) during the preceding 12 months (or for such s			n S-T						
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," "smaller reporting company" and "emerging growth company" in Rule 12b-2 of the Exchange Act.										
Large accelerated filer \square Non-accelerated filer \square			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. \Box										
Indicate	by check mark whether the registrant is a shell company (as def	fined in Rule 12b-2 of the Exc	change Act) Yes □ No ⊠.							
As of Oc	tober 21, 2022, the registrant had outstanding 73,024,147 share	es of common stock, par value	s.0001, per share.							

SIGA TECHNOLOGIES, INC. FORM 10-Q

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PART I - FINANCIAL INFORMATION

Item 1 - Condensed Consolidated Financial Statements

SIGA TECHNOLOGIES, INC. CONDENSED CONSOLIDATED BALANCE SHEETS (UNAUDITED)

ASSETS	S	September 30, 2022		December 31, 2021
Current assets				
Cash and cash equivalents	\$	109,749,555	\$	103,138,819
Accounts receivable		54,925,558		83,650,450
Inventory		31,261,346		19,510,379
Prepaid expenses and other current assets		964,065		2,453,444
Total current assets		196,900,524		208,753,092
Property, plant and equipment, net		1,979,517		2,365,957
Deferred income taxes, net		4,183,886		2,422,607
Goodwill		898,334		898,334
Other assets		261,814		286,585
Total assets	\$	204,224,075	\$	214,726,575
LIABILITIES AND STOCKHOLDERS' EQUITY				
Current liabilities				
Accounts payable	\$	2,445,390	\$	2,028,004
Accrued expenses and other current liabilities		15,887,359		9,252,812
Income tax payable		9,123,153		19,207,042
Total current liabilities		27,455,902		30,487,858
Warrant liability		_		6,521,441
Other liabilities		3,630,046		3,402,869
Total liabilities		31,085,948		40,412,168
Commitments and contingencies				
Stockholders' equity				
Common stock (\$.0001 par value, 600,000,000 shares authorized, 73,024,147 and 73,543,602, issued and				
outstanding at September 30, 2022 and December 31, 2021, respectively)		7,302		7,354
Additional paid-in capital		233,271,351		226,070,308
Accumulated deficit		(60,140,526)		(51,763,255)
Total stockholders' equity		173,138,127		174,314,407
Total liabilities and stockholders' equity	\$	204,224,075	\$	214,726,575

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

SIGA TECHNOLOGIES, INC. CONDENSED CONSOLIDATED STATEMENTS OF OPERATIONS AND COMPREHENSIVE INCOME (LOSS) (UNAUDITED)

	Three Months Ended September				-				
		30),			3	U,		
		2022		2021		2022		2021	
Revenues									
Product sales and supportive services	\$	65,621,511	\$	2,346,110	\$	81,558,148	\$	12,793,615	
Research and development		6,589,616		2,500,062		17,859,323		5,519,591	
Total revenues		72,211,127		4,846,172		99,417,471		18,313,206	
Operating expenses									
Cost of sales and supportive services		3,948,974		83,276		9,551,186		1,330,114	
Selling, general and administrative		19,656,138		4,471,640		29,241,565		14,113,384	
Research and development		5,732,982		3,235,145		16,119,858		7,801,901	
Total operating expenses		29,338,094		7,790,061		54,912,609		23,245,399	
Operating income/(loss)		42,873,033		(2,943,889)		44,504,862		(4,932,193)	
Gain/(loss) from change in fair value of warrant liability		_		(1,066,522)		400,663		294,548	
Other income, net		258,975		26,252		354,670		76,055	
Income/(loss) before income taxes		43,132,008		(3,984,159)		45,260,195		(4,561,590)	
(Provision)/benefit for income taxes		(10,091,420)		870,801		(10,543,595)		805,328	
Net and comprehensive income/(loss)	\$	33,040,588	\$	(3,113,358)	\$	34,716,600	\$	(3,756,262)	
Basic income/(loss) per share	\$	0.45	\$	(0.04)	\$	0.48	\$	(0.05)	
Diluted income/(loss) per share	\$	0.45	\$	(0.04)	\$	0.47	\$	(0.05)	
Weighted average shares outstanding: basic		73,024,147		74,840,846		72,924,178		75,822,713	
Weighted average shares outstanding: diluted		73,259,272		74,840,846		73,616,837		76,634,963	

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

${\bf SIGA\ TECHNOLOGIES, INC.}$ CONDENSED CONSOLIDATED STATEMENTS OF CASH FLOWS (UNAUDITED)

	Nine Months Ended September 30,			
	2022			2021
Cash flows from operating activities:				
Net income (loss)	\$	34,716,600	\$	(3,756,262)
Adjustments to reconcile net income (loss) to net cash used in operating activities:				
Depreciation and other amortization		386,440		393,667
Gain on change in fair value of warrant liability		(400,663)		(294,548)
Stock-based compensation		1,092,893		957,320
Write down of inventory, net		168,446		548,551
Deferred income taxes, net		(1,761,279)		(408,233)
Changes in assets and liabilities:				
Accounts receivable		28,724,892		(482,561)
Inventory		(11,919,413)		(9,956,076)
Prepaid expenses and other assets		1,514,150		(139,984)
Accounts payable, accrued expenses and other liabilities		914,139		9,095,055
Income tax payable		(10,083,889)		(871,298)
Deferred revenue		6,364,971		306,443
Net cash provided by/(used in) operating activities		49,717,287		(4,607,926)
Cash flows from investing activities:				
Capital expenditures		_		(24,424)
Cash used in investing activities		_		(24,424)
Cash flows from financing activities:				
Payment of employee tax obligations for common stock tendered		(12,533)		(173,918)
Repurchase of common stock		(10,149,704)		(20,264,547)
Payment of dividend		(32,944,314)		
Cash used in financing activities		(43,106,551)	-	(20,438,465)
Net increase/(decrease) in cash and cash equivalents		6,610,736		(25,070,815)
Cash and cash equivalents at the beginning of period		103,138,819		117,890,240
Cash and cash equivalents at end of period	\$	109,749,555	\$	92,819,425
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Supplemental disclosure of non-cash financing activities:				
Conversion of warrant to common stock	\$	6,120,778	\$	_

The accompanying notes are an integral part of these financial statements

SIGA TECHNOLOGIES, INC. NOTES TO CONDENSED CONSOLIDATED FINANCIAL STATEMENTS (UNAUDITED)

1. Condensed Consolidated Financial Statements

The financial statements of SIGA Technologies, Inc. ("we," "our," "us," "SIGA" or the "Company") are presented in accordance with accounting principles generally accepted in the United States of America ("U.S. GAAP") for interim financial information and the rules and regulations of the Securities and Exchange Commission (the "SEC") for quarterly reports on Form 10-Q and should be read in conjunction with the Company's audited financial statements and notes thereto for the year ended December 31, 2021, included in the Company's 2021 Annual Report on Form 10-K filed on March 3, 2022 (the "2021 Form 10-K"). All terms used but not defined elsewhere herein have the meaning ascribed to them in the 2021 Form 10-K. In the opinion of management, all adjustments (consisting of normal and recurring adjustments) considered necessary for a fair statement of the results of the interim periods have been included. The 2021 year-end condensed consolidated balance sheet data were derived from the audited financial statements but do not include all disclosures required by U.S. GAAP. The results of operations for the three and nine months ended September 30, 2022 are not necessarily indicative of the results expected for the full year.

2. Summary of Significant Accounting Policies

Revenue Recognition

The Company accounts for revenue in accordance with ASC Topic 606, *Revenue from Contracts with Customers* ("ASC 606"). In all transactions, the Company is the principal as it controls the specified good or service before it is transferred to the customer and therefore recognizes revenue on a gross basis. A contract's transaction price is allocated to distinct performance obligations and recognized as revenue when, or as, a performance obligation is satisfied. The Company accounts for shipping and handling activities as fulfillment costs rather than as an additional promised service. As of September 30, 2022, the Company's active contractual performance obligations are referenced in Note 3 and consist of the following: five performance obligations relate to research and development services; and five relate to manufacture and delivery of product. The aggregate amount of the transaction price allocated to remaining performance obligations was \$82.9 million as of September 30, 2022. Remaining performance obligations represent the transaction price for which work has not been performed and excludes unexercised contract options. The Company expects to recognize this amount as revenue within the next three years as the specific timing for satisfying performance obligations is subjective and largely outside the Company's control.

Performance Obligations

A performance obligation is a promise in a contract to transfer a distinct good or service to the customer and is the unit of account in ASC 606. A contract's transaction price is allocated to each distinct performance obligation and recognized as revenue when, or as, the performance obligation is satisfied.

Contract modifications may occur during the course of performance of our contracts. Contracts are often modified to account for changes in contract specifications or requirements. In most instances, contract modifications are for services that are not distinct, and, therefore, are accounted for as part of the existing contract.

The Company's performance obligations are satisfied over time as work progresses or at a point in time. All of the Company's revenue related to current research and development performance obligations is recognized over time, because the customer simultaneously receives and consumes the benefits provided by the services as the Company performs these services. The Company recognizes revenue related to these services based on the progress toward complete satisfaction of the performance obligation and measures this progress under an input method, which is based on the Company's cost incurred relative to total estimated costs. Under this method, progress is measured based on the cost of resources consumed (i.e., cost of third-party services performed, cost of direct labor hours incurred, and cost of materials consumed) compared to the total estimated costs to completely satisfy the performance obligation. Incurred costs represent work performed, which corresponds with, and thereby best depicts, the transfer of control to the customer. The incurred and estimated costs used in the measure of progress include third-party services performed, direct labor hours, and material consumed.

Contract Balances

The timing of revenue recognition, billings and cash collections may result in billed accounts receivable, unbilled receivables (contract assets) and customer advances and deposits (contract liabilities) in the condensed consolidated balance sheets. Generally, amounts are billed as work progresses in accordance with agreed-upon contractual terms either at periodic intervals (monthly) or upon achievement of contractual milestones; as of September 30, 2022, the accounts receivable balance in the condensed balance sheet includes approximately \$46.3 million of unbilled receivables. This amount includes net proceeds, net of Meridian fee (as defined below) from international sales, which will be billed and collected by Meridian and paid to SIGA. Under typical payment terms of fixed price arrangements, the customer pays the Company either performance-based payments or progress payments. For the Company's cost-type arrangements, the customer generally pays the Company for its actual costs incurred, as well as its allocated overhead and general and administrative costs. Such payments occur within a short period of time from billing. When the Company receives consideration, or such consideration is unconditionally due, prior to transferring goods or services to the customer under the terms of a sales contract, the Company records deferred revenue, which represents a contract liability. During the nine months ended September 30, 2022, the Company recognized \$3.5 million of revenue that was included in deferred revenue at the beginning of the period.

Repurchase of shares

When shares recognized as equity are repurchased, the amount of the consideration paid, which includes directly attributable costs, is recognized as a deduction from equity. The excess of the purchase price above par value of repurchased shares that are retired is presented as an increase to accumulated deficit (or a reduction of retained earnings, if any).

3. Procurement Contracts and Research Agreements

19C BARDA Contract

On September 10, 2018, the Company entered into a contract with the U.S. Biomedical Advanced Research and Development Authority ("BARDA") pursuant to which SIGA agreed to deliver up to 1,488,000 courses of oral TPOXX® to the U.S. Strategic National Stockpile ("Strategic Stockpile"), and to manufacture and deliver to the Strategic Stockpile, or store as vendor-managed inventory, up to 212,000 courses of the intravenous (IV) formulation of TPOXX® ("IV TPOXX®"). Additionally, the contract includes funding from BARDA for a range of activities, including: advanced development of IV TPOXX®, post-marketing activities for oral and IV TPOXX®, and procurement activities. As of September 30, 2022, the contract with BARDA (as amended, modified, or supplemented from time to time, the "19C BARDA Contract") contemplates up to approximately \$602.5 million of payments, of which approximately \$51.7 million of payments are included within the base period of performance of five years, approximately \$268.9 million of payments are related to exercised options and up to approximately \$281.9 million of payments are currently specified as unexercised options. BARDA may choose in its sole discretion when, or whether, to exercise any of the unexercised options. The period of performance for options is up to ten years from the date of entry into the 19C BARDA Contract and such options could be exercised at any time during the contract term, including during the base period of performance.

The base period of performance specifies potential payments of approximately \$51.7 million for the following activities: payments of approximately \$11.1 million for the delivery of approximately 35,700 courses of oral TPOXX® to the Strategic Stockpile; payments of \$8.0 million for the manufacture of 20,000 courses of final drug product of IV TPOXX® ("IV FDP"), of which \$3.2 million of payments are related to the manufacture of bulk drug substance ("IV BDS") to be used in the manufacture of IV FDP; payments of approximately \$32.0 million to fund advanced development of IV TPOXX®; and payments of approximately \$0.6 million for supportive procurement activities. As of September 30, 2022, the Company has received \$11.1 million for the delivery of approximately 35,700 courses of oral TPOXX® to the Strategic Stockpile, \$3.2 million for the manufacture of IV BDS, \$4.3 million for the delivery of IV FDP to the Strategic Stockpile and \$17.1 million for other base period activities. IV BDS has been used for the manufacture of courses of IV FDP. The \$3.2 million received for the completed manufacture of IV BDS had been recorded as deferred revenue as of December 31, 2021, but with the delivery of IV FDP to the Strategic Stockpile during the nine months ended September 30, 2022, \$2.9 million was recognized as revenue. The remaining \$0.3 million of deferred revenue will be recognized as IV FDP containing such IV BDS is delivered to the Strategic Stockpile.

The options that have been exercised to date provide for payments up to approximately \$268.9 million. There are exercised options for the following activities: payments up to \$11.2 million for the procurement of raw materials used in the 2020 manufacture of certain courses of oral TPOXX®; payments up to \$213.9 million for the delivery of up to 726,140 courses of oral TPOXX®; payments up to \$25.6 million for the manufacture of courses of IV FDP, of which \$10.2 million of payments relate to the manufacture of IV BDS to be used in the manufacture of IV FDP, payments of up to approximately \$3.6 million to fund post-marketing activities for IV TPOXX®, and payments of up to \$14.6 million for funding of post-marketing activities for oral TPOXX®. As of September 30, 2022, the Company has received \$225.1 million for the delivery (and related procurement of raw materials) of oral TPOXX® to the Strategic Stockpile; received \$10.2 million for the completed manufacture of IV BDS, which has been recorded as deferred revenue as of September 30, 2022; and received or billed \$8.1 million in connection with post-marketing activities for oral TPOXX®.

Unexercised options specify potential payments up to approximately \$281.9 million in total (if all such options are exercised). There are options for the following activities: payments of up to \$225.1 million for the delivery of oral TPOXX® to the Strategic Stockpile; payments of up to \$51.2 million for the manufacture of courses of IV FDP, of which up to \$20.5 million of payments would be paid upon the manufacture of IV BDS to be used in the manufacture of IV FDP; and payments of up to approximately \$5.6 million for supportive procurement activities.

The options related to IV TPOXX® are divided into two primary manufacturing steps. There are options related to the manufacture of bulk drug substance ("IV BDS Options"), and there are corresponding options (for the same number of IV courses) for the manufacture of final drug product ("IV FDP Options"). BARDA may choose to exercise any, all, or none of these options in its sole discretion. The 19C BARDA Contract includes: three separate IV BDS Options, each providing for the bulk drug substance equivalent of 64,000 courses of IV TPOXX®; and three separate IV FDP Options, each providing for 64,000 courses of final drug product of IV TPOXX®. BARDA has the sole discretion as to whether to simultaneously exercise IV BDS Options and IV FDP Options, or whether to exercise options at different points in time (or alternatively, to only exercise the IV BDS Option but not the IV FDP Option). To date, BARDA has exercised one of the three IV BDS options and one of the three IV FDP options, both of which were exercised simultaneously in 2022. If BARDA decides to only exercise the remaining IV BDS Options, then the Company would receive payments up to \$20.5 million; alternatively, if BARDA decides to exercise all the remaining IV BDS Options and IV FDP Options, then the Company would receive payments up to \$51.2 million. For each set of options relating to a specific group of courses (for instance, the IV BDS and IV FDP options that reference the same 64,000 courses), BARDA has the option to independently purchase IV BDS or IV FDP.

Revenues in connection with the 19C BARDA Contract are recognized either over time or at a point in time. Performance obligations related to product delivery generate revenue at a point in time. Revenue from other performance obligations under the 19C BARDA Contract are recognized over time using an input method using costs incurred to date relative to total estimated costs at completion. For the three months ended September 30, 2022 and 2021, the Company recognized revenues of \$2.7 million and \$1.1 million, respectively, on an over time basis. For the nine months ended September 30, 2022 and 2021, the Company recognized revenues of \$4.6 million and \$2.8 million, respectively, on an over time basis. Revenue recognized for product delivery, and therefore at a point in time, for the nine months ended September 30, 2022 was \$7.2 million. In contrast, no revenue was recognized for product delivery, and therefore no revenue was recognized at a point in time, for the three months ended September 30, 2022 or for the three and nine months ended September 30, 2021.

U.S. Department of Defense Procurement Contracts

On May 12, 2022, the Company announced a contract with the U.S. Department of Defense ("DoD") for the procurement of oral TPOXX® ("DoD Contract #1"). The DoD Contract #1 included a firm commitment for the DoD to procure approximately \$3.6 million of oral TPOXX®, and an option, exercisable at the sole discretion of the DoD, for the procurement of approximately \$3.8 million of oral TPOXX®. In the second quarter of 2022, the Company delivered and recognized revenue of \$3.6 million for the delivery of oral TPOXX® to the DoD, fulfilling the firm commitment in DoD Contract #1. In the third quarter of 2022, the DoD exercised the option for \$3.8 million of oral TPOXX® and the Company satisfied its obligation by delivering product and recognized the related revenue in September 2022.

On September 28, 2022, the Company and the DoD signed a new procurement contract ("DoD Contract #2"). The DoD Contract #2 includes a firm commitment for the DoD to procure approximately \$5.2 million of oral TPOXX®, and an option, exercisable at the sole discretion of the DoD for the procurement of approximately \$5.5 million of oral TPOXX®.

International Procurement Contracts

This year, through October 31, 2022, the Company has received firm commitment orders from 13 international customers (including Canada) for the delivery of approximately \$77 million of oral TPOXX®, of which approximately \$39 million is for Canada and approximately \$38 million is for jurisdictions in Europe, Asia-Pacific, and the Middle East. Additionally, the contract with CDND (defined below) has an option until March 31, 2024, exercisable at the sole discretion of CDND, for the purchase of up to an additional \$6 million of oral TPOXX®. With respect to the \$77 million of firm commitment orders that have been received this year, approximately \$66 million of oral TPOXX® was delivered and recorded as revenue in the nine months ended September 30, 2022. Through an International Promotion Agreement (defined below), Meridian Medical Technologies, Inc. ("Meridian") is the counterparty to international contracts under which orders are placed for the purchase of oral TPOXX®. The Public Health Agency of Canada ("PHAC") and the Canadian Department of National Defence ("CDND") are among the contracting parties for the purchase of oral TPOXX® (see below for a summary description of these contracts).

On January 13, 2021, PHAC awarded a contract to Meridian (the "PHAC Contract") for the purchase of up to approximately \$33 million of oral TPOXX® (tecovirimat) within five years. In March 2022 and July 2022, PHAC executed amendments in which total procurement of oral TPOXX® under the PHAC Contract was increased to an amount of approximately \$45 million. Prior to 2022, approximately \$10 million of oral TPOXX® had been ordered and delivered to PHAC. No courses of oral TPOXX® were delivered under this contract for the first six months of 2022. During the three months ended September 30, 2022, approximately \$35 million of oral TPOXX® was delivered to PHAC and recognized as revenue. Including the deliveries in the third quarter of 2022, there are no current remaining amounts specified in the PHAC Contract.

On April 3, 2020, the Company announced that the CDND awarded a contract (the "Canadian Military Contract") to Meridian, pursuant to which the CDND would purchase up to approximately \$14 million of oral TPOXX® over four years. Prior to 2022, approximately \$4 million of oral TPOXX® had been ordered and delivered to CDND. No courses of oral TPOXX® were delivered under this contract for the first nine months of 2022. As of September 30, 2022, an approximate firm commitment order of \$4 million remains to be delivered under this contract. Additionally, there are approximately \$6 million of unexercised options, exercisable at the sole discretion of CDND, remaining under this contract.

The above-listed contract awards were coordinated between SIGA and Meridian under the international promotion agreement (as amended, the "International Promotion Agreement") that was entered into by the parties on June 3, 2019. Under the International Promotion Agreement, Meridian is the counterparty in connection with international contracts for oral TPOXX® and SIGA is responsible for manufacture and delivery of any oral TPOXX® purchased thereunder.

Under the terms of the International Promotion Agreement, Meridian was granted exclusive rights to market, advertise, promote, offer for sale, or sell oral TPOXX® in a field of use specified in the International Promotion Agreement in all geographic regions except for the United States (the "Territory"), and Meridian has agreed not to commercialize any competing product, as defined in the International Promotion Agreement, in the specified field of use in the Territory. SIGA retains ownership, intellectual property, distribution and supply rights and regulatory responsibilities in connection with TPOXX®, and, in the United States market, also retains sales and marketing rights with respect to oral TPOXX®. SIGA's consent is required for the entry into any sales arrangement pursuant to the International Promotion Agreement.

The fee Meridian retains pursuant to the International Promotion Agreement is a specified percentage of the collected proceeds of sales of oral TPOXX® net of certain expenses, for calendar years in which customer collected amounts net of such expenses are less than or equal to a specified threshold, and a higher specified percentage of such collected net proceeds for calendar years in which such net collected amounts exceed the specified threshold. It is probable that we will exceed the specified threshold in 2022 and, as a result, the Company has recorded and will continue to record the higher specified percentage for all International Promotion Agreement sales in 2022.

Revenue in connection with international procurement contracts for the delivery of product are recognized at a point in time on a gross basis, as the Company acts as the principal in the transaction. During the three and nine months ended September 30, 2022, the Company recognized \$61.3 million and \$66.2 million of sales, respectively, in connection with international contracts. During the three and nine months ended September 30, 2021, the Company recognized \$2.3 million and \$12.7 million of sales, respectively, for deliveries to PHAC and CDND.

Research Agreements and Grants

In July 2019, the Company was awarded a multi-year research contract valued at a total of \$19.5 million, with an initial award of \$12.4 million, from the DoD to support work in pursuit of a potential label expansion for oral TPOXX® that would include post-exposure prophylaxis ("PEP") of smallpox (such work known as the "PEP Label Expansion Program" and the contract referred to as the "PEP Label Expansion R&D Contract"). In subsequent modifications, the DoD increased the scope and the available funding under the PEP Label Expansion R&D Contract to approximately \$27 million. The period of performance for this contract, as modified, terminates on January 31, 2025. As of September 30, 2022, remaining revenue to be recognized in the future under the PEP Label Expansion R&D Contract is up to \$11.2 million. Revenue from the performance obligation under the PEP Label Expansion R&D Contract is recognized over time using an input method using costs incurred to date relative to total estimated costs at completion. For the three months ended September 30, 2022 and 2021, the Company, under the PEP Label Expansion R&D Contract, recognized revenue of \$4.2 million and \$1.3 million, respectively, on an over time basis. For the nine months ended September 30, 2022 and 2021, the Company, under the PEP Label Expansion R&D Contract, recognized revenue of \$13.1 million and \$2.0 million, respectively, on an over time basis.

Contracts and grants include, among other things, options that may or may not be exercised at the U.S. Government's discretion. Moreover, contracts and grants contain customary terms and conditions including the U.S. Government's right to terminate or restructure a contract or grant for convenience at any time. As such, the Company may not be eligible to receive all available funds.

4. Inventory

Inventory includes costs related to the manufacture of TPOXX®. Inventory consisted of the following:

		As of				
		September 30, 2022	December 31, 2021			
Raw materials	\$	22,047	\$	22,047		
Work in-process		25,075,163		17,453,358		
Finished goods		6,164,136		2,034,974		
Inventory	\$	31,261,346	\$	19,510,379		

5. Property, Plant and Equipment

Property, plant and equipment consisted of the following:

	As of				
	S	September 30, 2022	Γ	December 31, 2021	
Leasehold improvements	\$	2,420,028	\$	2,420,028	
Computer equipment		471,286		511,062	
Furniture and fixtures		347,045		377,859	
Operating lease right-of-use assets	3,678,647		3,678,647		
		6,917,006		6,987,596	
Less - accumulated depreciation and amortization		(4,937,489)		(4,621,639)	
Property, plant and equipment, net	\$	1,979,517	\$	2,365,957	

Depreciation and amortization expense on property, plant, and equipment was \$0.4 million for each of the nine months ended September 30, 2022 and 2021.

6. Accrued Expenses and Other Current Liabilities

Accrued expenses and other current liabilities consisted of the following:

	As of				
	Se	ptember 30, 2022	D	ecember 31, 2021	
Deferred revenue	\$	10,129,667	\$	3,764,696	
Compensation		1,997,717		2,811,700	
Research and development vendor costs		1,107,791		256,397	
Professional fees		900,205		527,026	
Other		852,370		938,082	
Lease liability, current portion		519,198		466,830	
Inventory		380,411		488,081	
Accrued expenses and other current liabilities	\$	15,887,359	\$	9,252,812	

7. Financial Instruments

2016 Warrant

On September 2, 2016, the Company entered into a loan and security agreement (as amended from time to time, the "Loan Agreement") with OCM Strategic Credit SIGTEC Holdings, LLC ("Lender"). The Company voluntarily prepaid this Loan Agreement in 2020. Upon such prepayment and release, the Loan Agreement was terminated. In connection with the entry into the Loan Agreement, the Company issued a warrant (the "Warrant") to the Lender on September 2, 2016 to purchase a number of shares of the Company's common stock equal to \$4.0 million divided by the lower of (i) \$2.29 per share and (ii) the subscription price paid in connection with the Rights Offering. The Warrant provides for weighted average anti-dilution protection and is exercisable in whole or in part for ten (10) years from the date of issuance. The per share subscription price paid was \$1.50 in connection with the Rights Offering; accordingly, the exercise price of the Warrant was set at \$1.50 per share, and there were 2.7 million shares underlying the Warrant. Taking into account partial exercises of the Warrant, there were approximately 1.0 million shares underlying the outstanding Warrant as of December 31, 2021.

In the second quarter of 2022, the Warrant was fully exercised, and therefore there are no remaining underlying shares as of September 30, 2022. See Note 8. For the nine months ended September 30, 2022, we recorded a gain of approximately \$0.4 million, reflecting a decrease in the fair value of the liabilityclassified warrant primarily due to the decrease in our stock price prior to the exercise of the Warrant.

As of December 31, 2021, there were approximately 1.0 million shares underlying the outstanding Warrant and the fair value of the Warrant was \$6.5 million. The fair value of the liability-classified Warrant was calculated using the following assumptions: risk-free interest rate of 1.21%; no dividend vield; an expected life of 4.7 years; and a volatility factor of 55%.

8. Fair Value of Financial Instruments

The carrying value of cash equivalents, accounts receivable, accounts payable, accrued expenses and other current liabilities, and income tax payable approximates fair value due to the relatively short maturity of these instruments. Prior to being fully exercised, common stock warrants, which were classified as a liability, were recorded at their fair market value as of each reporting period.

The measurement of fair value requires the use of techniques based on observable and unobservable inputs. Observable inputs reflect market data obtained from independent sources, while unobservable inputs reflect our market assumptions. The inputs create the following fair value hierarchy:

- Level 1 Quoted prices for identical instruments in active markets.
- Level 2 Quoted prices for similar instruments in active markets; quoted prices for identical or similar instruments in markets that are not active; and model-derived valuations where inputs are observable or where significant value drivers are observable.
- Level 3 Instruments where significant value drivers are unobservable to third parties.

There were no transfers between levels of the fair value hierarchy for the nine months ended September 30, 2022. As of each of September 30, 2022 and December 31, 2021, the Company had approximately \$0.1 million of cash equivalents classified as Level 1 financial instruments. As of September 30, 2022, the Company had approximately \$35.1 million of cash equivalents classified as Level 2 financial instruments. There were no Level 2 financial instruments as of December 31, 2021.

The following table presents changes in the liability-classified warrant measured at fair value using Level 3 inputs:

	Fair Value Measurements of Level 3 liability- classified warrant
Warrant liability at December 31, 2021	\$ 6,521,441
Decrease in fair value of Warrant liability	(400,663)
Exercise of Warrant	(6,120,778)
Warrant liability at September 30, 2022	<u>\$</u>
10	

9. Per Share Data

The Company computes, presents and discloses earnings per share in accordance with the authoritative guidance, which specifies the computation, presentation and disclosure requirements for earnings per share of entities with publicly held common stock or potential common stock. The objective of basic EPS is to measure the performance of an entity over the reporting period by dividing income (loss) by the weighted average shares outstanding. The objective of diluted EPS is consistent with that of basic EPS, except that it also gives effect to all potentially dilutive common shares outstanding during the period.

The following is a reconciliation of the basic and diluted loss per share computation:

	Tł	nree Months E 30	l September	Nine Months Ended Septembe 30,				
		2022	2021	2022			2021	
Net income/(loss) for basic earnings per share	\$	33,040,588	\$ (3,113,358)	\$	34,716,600	\$	(3,756,262)	
Less: Change in fair value of warrants		_	_		400,663		294,548	
Net income/(loss), adjusted for change in fair value of warrants for diluted			_					
earnings per share	\$	33,040,588	\$ (3,113,358)	\$	34,315,937	\$	(4,050,810)	
Weighted-average shares		73,024,147	74,840,846		72,924,178		75,822,713	
Effect of potential common shares		235,125	_		692,659		812,250	
Weighted-average shares: diluted		73,259,272	74,840,846		73,616,837		76,634,963	
Income/(loss) per share: basic	\$	0.45	\$ (0.04)	\$	0.48	\$	(0.05)	
Income/(loss) per share: diluted	\$	0.45	\$ (0.04)	\$	0.47	\$	(0.05)	

For the nine months ended September 30, 2022, the diluted earnings per share calculation reflects the effect of the exercise of outstanding warrants and any corresponding elimination of the impact included in operating results from the change in fair value of the warrants. Weighted-average diluted shares include the dilutive effect of in-the-money options, stock-settled RSUs and warrants. The dilutive effect of warrants, stock-settled RSUs and options is calculated based on the average share price for each fiscal period using the treasury stock method. Under the treasury stock method, the amount the employee must pay for exercising stock options, the average amount of compensation cost for future service that the Company has not yet recognized, and the amount of tax benefits that would be recorded in additional paid-in capital when the award becomes deductible, are collectively assumed to be used to repurchase shares. Cash-settled RSUs were presumed to be cash-settled and therefore excluded from the diluted earnings per share calculations for the three and nine months ended September 30, 2022 because the net effect of their inclusion, including the elimination of the impact in the operating results of the change in fair value of these RSUs, would have been anti-dilutive. For the three and nine months ended September 30, 2022, the weighted average number of shares under the cash-settled RSUs excluded from the calculation of diluted earnings per share were 32,180 and 12,901, respectively.

For the three and nine months ended September 30, 2021, the Company incurred losses and as a result, the equity instruments listed below were excluded from the calculation of diluted earnings (loss) per share as the effect of the exercise, conversion or vesting of such instruments would have been anti-dilutive. The weighted average number of equity instruments excluded consists of:

	Three Months Ended September 30, 2021	Nine Months Ended September 30, 2021
Warrants	1,047,296	_
Stock options	150,000	153,055
Restricted stock units	204.591	181.730

10. Commitments and Contingencies

From time to time, we may be involved in a variety of claims, suits, investigations and proceedings arising from the ordinary course of our business, collections claims, breach of contract claims, labor and employment claims, tax and other matters. Although such claims, suits, investigations and proceedings are inherently uncertain and their results cannot be predicted with certainty, we believe that the resolution of such current pending matters, if any, will not have a material adverse effect on our business, consolidated financial position, results of operations or cash flow. Regardless of the outcome, litigation can have an adverse impact on us because of legal costs, diversion of management resources and other factors.

Purchase Commitments

In the course of our business, the Company regularly enters into agreements with third party organizations to provide contract manufacturing services and research and development services. Under these agreements, the Company issues purchase orders, which obligate the Company to pay a specified price when agreed-upon services are performed. In connection with many CMO purchase orders, reimbursement by CMOs for inventory losses is limited. Commitments under the purchase orders do not exceed our planned commercial and research and development needs. As of September 30, 2022, the Company had approximately \$21.8 million of purchase commitments associated with manufacturing obligations.

11. Related Party Transactions

Board of Directors and Outside Counsel

A former member of the Company's Board of Directors who did not stand for re-election at the Company's 2021 annual meeting of stockholders is a partner at a law firm used by the Company. The Company did not incur any expenses related to services provided by the outside counsel during the three months ended September 30, 2022 and 2021 or the nine months ended September 30, 2022. During the nine months ended September 30, 2021, the Company incurred \$0.1 million of expenses related to services provided by the outside counsel. The Company had no outstanding payables or accrued expenses related to services performed by the outside counsel as of September 30, 2022.

Real Estate Leases

On May 26, 2017, the Company and MacAndrews & Forbes Incorporated ("M&F") entered into a ten-year Office Lease agreement (the "HQ Lease"), pursuant to which the Company agreed to lease 3,200 square feet at 31 East 62nd Street, New York, New York. The Company is utilizing premises leased under the HQ Lease as its new corporate headquarters. The Company's rental obligations consist of a fixed rent of \$25,333 per month in the first sixty-three months of the term, subject to a rent abatement for the first six months of the term. From the first day of the sixty-fourth month of the term through the expiration or earlier termination of the lease, the Company's rental obligations consist of a fixed rent of \$29,333 per month. In addition to the fixed rent, the Company will pay a facility fee in consideration of the landlord making available certain ancillary services, commencing on the first anniversary of entry into the lease. The facility fee was \$3,333 per month for the second year of the term and increases by five percent each year thereafter, to \$4,925 per month in the final year of the term. During the three and nine months ended September 30, 2022, the Company paid expenses associated with this lease of \$0.1 million and \$0.3 million, respectively.

12. Revenues by Geographic Region

Revenues by geographic region were as follows:

	Three Months Ended September 30,				Nine Months Ended Septem 30,			
		2022		2021		2022		2021
United States	\$	10,925,562	\$	2,475,689	\$	33,180,163	\$	5,387,145
International								
Asia-Pacific		8,946,221		_		13,875,644		_
Canada		35,120,212		2,370,483		35,120,212		12,926,061
Europe, Middle East and Africa (EMEA)		16,255,715		_		16,278,035		_
Other		963,417		_		963,417		_
Total International		61,285,565		2,370,483		66,237,308		12,926,061
Total revenues	\$	72,211,127	\$	4,846,172	\$	99,417,471	\$	18,313,206
	12							

13. Income Taxes

The Company's provision for income taxes consists of federal and state taxes, as applicable, in amounts necessary to align the Company's year-to-date tax provision with the effective rate that it expects to achieve for the full year. Each quarter the Company updates its estimate of the annual effective tax rate and records cumulative adjustments as necessary.

For the three months ended September 30, 2022 and 2021, we recorded pre-tax income of \$43.1 million and losses of (\$4.0) million, respectively, and a corresponding income tax (provision)/benefit of (\$10.1) million and \$0.9 million, respectively.

For the nine months ended September 30, 2022 and 2021, we recorded pre-tax income of \$45.3 million and losses of (\$4.6) million, respectively, and a corresponding income tax (provision)/benefit of (\$10.5) million and \$0.8 million, respectively.

The effective tax rate for the three months ended September 30, 2022 was 23.4% compared to 21.9% for the three months ended September 30, 2021. The effective tax rate for the three months ended September 30, 2022 differs from the U.S. statutory rate of 21% primarily as a result of state taxes, and various non-deductible expenses, including executive compensation under Internal Revenue Code Section 162(m).

The effective tax rate for the nine months ended September 30, 2022 was 23.3% compared to 17.7% for the nine months ended September 30, 2021. The effective tax rate for the nine months ended September 30, 2022 differs from the U.S. statutory rate of 21% primarily as a result of state taxes, and various non-deductible expenses, including executive compensation under Internal Revenue Code Section 162(m).

The Inflation Reduction Act of 2022 (the "Act") was signed into U.S. law on August 16, 2022. The Act includes various tax provisions, including an excise tax on stock repurchases, expanded tax credits for clean energy incentives, and a corporate alternative minimum tax that generally applies to U.S. corporations with average adjusted annual financial statement income over a three-year period in excess of \$1 billion. The Company does not expect the Act to materially impact its consolidated financial statements.

14. Equity

The tables below present changes in stockholders' equity for the three and nine months ended September 30, 2022 and 2021.

	Common	ı Sto	ck	Additional Paid-in	Accumulated	Other Comprehensive	Total Stockholders'
	Shares	Α	mount	Capital	Deficit	Income	Equity
Balances at June 30, 2022	73,024,147	\$	7,302	\$232,942,666	\$ (93,181,114)	\$ —	\$139,768,854
Net income	_		_		33,040,588	_	33,040,588
Stock-based compensation				328,685			328,685
Balances at September 30, 2022	73,024,147	\$	7,302	\$233,271,351	\$(60,140,526)	\$ —	\$173,138,127
•							
				Additional		Other	Total
	Common	Stoc	:k	Paid-in	Accumulated	Comprehensive	Stockholders'
	Shares	Aı	mount	Capital	Deficit	Income	Equity
Balances at December 31, 2021	73,543,602	\$	7,354	\$226,070,308	\$(51,763,255)	\$ —	\$174,314,407
Net income			_	_	34,716,600	_	34,716,600
Repurchase of common stock	(1,474,781)		(147)	_	(10,149,557)	_	(10,149,704)
Payment of common stock tendered for employee stock-							
based compensation tax obligations	(1,973)		_	(12,533)	_	_	(12,533)
Issuance of common stock upon vesting of RSUs	132,396		13	(13)	_	_	_
Issuance of common stock upon exercise of warrants	824,903		82	6,120,696	_	_	6,120,778
Cash dividend (\$0.45 per share)	_		_	_	(32,944,314)	_	(32,944,314)
Stock-based compensation	_		_	1,092,893	_	_	1,092,893
Balances at September 30, 2022	73,024,147	\$	7,302	\$233,271,351	\$(60,140,526)	<u> </u>	\$173,138,127
		13					

	Common	Stoc	ck	Additional Paid-in	Accumulated		Other prehensive	Total Stockholders'
	Shares	An	nount	Capital	Deficit	I	ncome	Equity
Balances at June 30, 2021	75,389,417	\$	7,539	\$225,678,876	\$(108,965,453)	\$	_	\$116,720,962
Net loss	_		_	_	(3,113,358)		_	(3,113,358)
Repurchase of common stock	(1,110,763)		(111)	_	(7,134,577)		_	(7,134,688)
Payment of common stock tendered for employee stock- based compensation tax obligations	(25,203)		(2)	(160,554)	_		_	(160,556)
Issuance of common stock upon vesting of RSUs	53,334		5	(5)	_		_	_
Stock-based compensation	_		_	243,502	_		_	243,502
Balances at September 30, 2021	74,306,785	\$	7,431	\$225,761,819	\$ (119,213,388)	\$	_	\$106,555,862
	Common	Stoc	ck	Additional Paid-in	Accumulated		Other prehensive	Total Stockholders'
	Common Shares		ck mount		Accumulated Deficit	Com	Other prehensive ncome	
Balances at December 31, 2020				Paid-in		Com	prehensive	Stockholders'
Balances at December 31, 2020 Net loss	Shares	Ar	mount	Paid-in Capital	Deficit	Com I	prehensive	Stockholders' Equity
,	Shares	Ar	mount	Paid-in Capital	Deficit \$ (95,192,881)	Com I	prehensive	Stockholders' Equity \$129,793,269
Net loss Repurchase of common stock Payment of common stock tendered for employee stock-	Shares 77,195,704	Ar	7,720	Paid-in Capital	Deficit \$ (95,192,881) (3,756,262)	Com I	prehensive	Stockholders' Equity \$129,793,269 (3,756,262)
Net loss Repurchase of common stock Payment of common stock tendered for employee stock-based compensation tax obligations	Shares 77,195,704	Ar	7,720	Paid-in Capital	Deficit \$ (95,192,881) (3,756,262)	Com I	prehensive	Stockholders' Equity \$129,793,269 (3,756,262)
Net loss Repurchase of common stock Payment of common stock tendered for employee stock-	Shares 77,195,704 — (3,024,690)	Ar	7,720 — (302)	Paid-in Capital \$224,978,430 — —	Deficit \$ (95,192,881) (3,756,262)	Com I	prehensive	Stockholders' Equity \$129,793,269 (3,756,262) (20,264,547)
Net loss Repurchase of common stock Payment of common stock tendered for employee stock-based compensation tax obligations	Shares 77,195,704 — (3,024,690) (27,105)	Ar	7,720 — (302)	Paid-in Capital \$224,978,430 — — (173,915)	Deficit \$ (95,192,881) (3,756,262)	Com I	prehensive	Stockholders' Equity \$129,793,269 (3,756,262) (20,264,547)

On August 2, 2021, the Company's Board of Directors authorized a share repurchase program ("New Repurchase Authorization") under which the Company may repurchase up to \$50 million of the Company's common stock through December 31, 2023. The Company started repurchasing shares under this program in the fourth quarter of 2021. Repurchases under the New Repurchase Authorization may be made from time to time at the Company's discretion in open market transactions, through block trades, in privately negotiated transactions and pursuant to any trading plan that may be adopted by the Company's management in accordance with Rule 10b5-1 under the Securities Exchange Act of 1934, as amended (the "Exchange Act") or otherwise. The timing and actual number of shares repurchased will depend on a variety of factors, including: timing of procurement orders under government contracts; alternative opportunities for strategic uses of cash; the stock price of the Company's common stock; market conditions; alternative capital management uses of cash; and other corporate liquidity requirements and priorities. During the nine months ended September 30, 2022, the Company repurchased approximately 1.5 million shares of common stock under the New Repurchase Authorization for approximately \$10.1 million. No shares were repurchased in the three months ended September 30, 2022.

Prior to the effective date of the New Repurchase Authorization, the Company repurchased shares under a program that was announced in March 2020. Under this program, \$50 million of the Company's common stock was repurchased, including approximately 1.1 million shares of common stock for approximately \$7.1 million that was repurchased during the three months ended September 30, 2021, and 3.0 million shares of common stock for approximately \$20.3 million that was repurchased during the nine months ended September 30, 2021.

On May 5, 2022, the Board of Directors declared a special dividend of \$0.45 per share on the common stock of the Company, which resulted in an overall dividend payment of \$32.9 million. The special dividend was paid on June 2, 2022 to shareholders of record at the close of business on May 17, 2022.

15. Leases

The Company leases its Corvallis, Oregon, facilities and office space under an operating lease, which was signed on November 3, 2017 and commenced on January 1, 2018. The initial term of this lease was to expire on December 31, 2019 after which the Company had two successive renewal options; one for two years and the other for three years. In the second quarter of 2019, the Company exercised the first renewal option, which extended the lease expiration date to December 31, 2021. In the second quarter of 2021, the Company exercised the second renewal option, which extended the lease expiration date to December 31, 2024. In connection with the exercise of the second renewal option, the Company recorded an increase to operating lease right-of-use assets and operating lease liabilities of approximately \$0.7 million in the second quarter of 2021.

On May 26, 2017 the Company and M&F entered into the HQ Lease, a ten-year office lease agreement, pursuant to which the Company agreed to lease 3,200 square feet in New York, New York. The Company is utilizing premises leased under the HQ Lease as its corporate headquarters. The Company has no leases that qualify as finance leases.

Operating lease costs totaled \$0.2 million and \$0.1 million for the three months ended September 30, 2022 and 2021, respectively. Operating lease costs totaled \$0.5 million and \$0.4 million for the nine months ended September 30, 2022 and 2021, respectively. Cash paid for amounts included in the measurement of lease liabilities from operating cash flows was \$0.2 million and \$0.1 million for the three months ended September 30, 2022 and 2021, respectively. Cash paid for amounts included in the measurement of lease liabilities from operating cash flows was \$0.5 million and \$0.4 million for the nine months ended September 30, 2022 and 2021, respectively. As of September 30, 2022, the weighted-average remaining lease term of the Company's operating leases was 3.88 years while the weighted-average discount rate was 4.53%.

Future cash flows under operating leases as of September 30, 2022 are expected to be as follows:

2022	\$ 110,340
2023	669,048
2024	678,627
2025	406,994
2026	409,971
Thereafter	165,916
Total undiscounted cash flows under leases	2,440,896
Less: Imputed interest	(224,813)
Present value of lease liabilities	\$ 2,216,083

As of September 30, 2022, approximately \$1.7 million of the lease liability is included in Other liabilities on the condensed consolidated balance sheet with the current portion included in accrued expenses.

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

The following discussion should be read in conjunction with our condensed consolidated financial statements and notes to those statements and other financial information appearing elsewhere in this Quarterly Report on Form 10-Q and in the Company's Annual Report on Form 10-K filed on March 3, 2022 (the "2021 Form 10-K"). In addition to historical information, the following discussion and other parts of this Quarterly Report contain forward-looking information that involves risks and uncertainties. SIGA's actual results could differ materially from those anticipated by such forward-looking statements due to a number of factors. See the factors set forth under the heading "Safe Harbor Statement" at the end of this Item 2.

Overview

We are a commercial-stage pharmaceutical company. Our lead product, TPOXX® ("oral TPOXX®", also known as "tecovirimat" in certain international markets), is an oral formulation antiviral drug for the treatment of human smallpox disease caused by variola virus. On July 13, 2018, the United States Food & Drug Administration ("FDA") approved oral TPOXX® for the treatment of smallpox. The Company has been delivering oral TPOXX® to the U.S. Strategic National Stockpile ("Strategic Stockpile") since 2013.

In addition to being approved by the FDA, oral TPOXX® (tecovirimat) has regulatory approval with the European Medicines Agency ("EMA"), Health Canada and the Medicines and Healthcare Products Regulatory Agency ("MHRA") of the United Kingdom. The EMA and MHRA approved label indication covers the treatment of smallpox, monkeypox, cowpox, and vaccinia complications following vaccination against smallpox. The Health Canada approved label indication covers the treatment of smallpox.

With respect to the regulatory approvals by the EMA, MHRA and Health Canada, oral tecovirimat represents the same formulation that was approved by the FDA in July 2018 under the brand name TPOXX®.

For the intravenous formulation of TPOXX@ ("IV TPOXX@"), SIGA announced on May 19, 2022 that the FDA approved this formulation for the treatment of smallpox.

Monkeypox Outbreak

Starting in June 2022, procurement orders for oral TPOXX® from new international jurisdictions, as well as orders under existing contracts, have occurred as SIGA has received ongoing inquiries about accessing oral TPOXX® in connection with a global monkeypox outbreak. The Company believes that a portion of the courses of oral TPOXX® delivered under these orders are being used for the treatment of active monkeypox cases as part of a response to this outbreak by the global health community.

Monkeypox is a disease caused by infection with the monkeypox virus. The monkeypox virus is part of the same family of viruses as smallpox. Monkeypox symptoms are similar to smallpox but are not as severe, with historical fatality in Africa of less than 1% to 10% depending on region and clade. The first human case of monkeypox was recorded in 1970. Since then, monkeypox has been reported in several central and western African countries, with case numbers greatly increasing in recent years. Prior to the ongoing 2022 outbreak, nearly all monkeypox cases in people outside of Africa were linked to international travel to countries where the disease commonly occurs, or through imported animals, including two cases in the United States in 2021. These cases are currently occurring on multiple continents. On July 23, 2022, the World Health Organization declared the monkeypox outbreak as a public health emergency of international concern. On August 4, 2022, the U.S. Government declared the monkeypox outbreak as a public health emergency.

Starting in the third quarter of 2022, in connection with the monkeypox outbreak, randomized, placebo-controlled clinical trials have been initiated in the United States, United Kingdom and Democratic Republic of Congo ("DRC") to further assess the safety and efficacy of TPOXX® in participants with monkeypox. These randomized clinical trials are now enrolling patients to collect data on the potential benefits of using TPOXX® as an antiviral treatment for active monkeypox disease.

Study of Tecovirimat for Human Monkeypox Virus (STOMP; A5418) is a U.S.-based clinical trial sponsored by the National Institute of Allergy and Infectious Diseases ("NIAID"), part of the National Institutes of Health. The NIAID-funded AIDS Clinical Trials Group is leading the study, which may later expand to international sites. Study investigators aim to enroll more than 500 participants, including children and those who are pregnant or breastfeeding, from clinical research sites. The trial will also include an open label arm that will include children, pregnant/breastfeeding individuals and those who are immunocompromised or have severe monkeypox disease.

PLATINUM is a U.K.-based clinical trial commissioned and funded by the National Institute for Health Care and Research. The trial is led by researchers at Oxford University and aims to recruit at least 500 participants, including children weighing \geq 13 kg, across the U.K.

PALM 007 is a DRC-based clinical trial sponsored by NIAID and Institute National de Recherche Biomédicale. Study investigators aim to enroll more than 450 participants, including children weighing ≥3 kg and women who are pregnant or breastfeeding, at clinical sites in the DRC.

COVID-19 Pandemic

The COVID-19 pandemic has caused significant societal and economic disruption. The continuing direct and indirect impacts of the pandemic are significant and broad-based, including supply chain disruptions and labor shortages that started during the pandemic and continue to represent business and financial risks. As such, the Company is continually coordinating with service providers and vendors, in particular Contract Manufacturing Organizations ("CMOs") that constitute our supply chain, with respect to risks and mitigating actions.

As of the filing date of this report, the Company has not identified or been notified by government customers of impediments to the continued full performance of their government contracts. With regard to day-to-day operations, the COVID-19 pandemic, and the secondary effects of the pandemic, have at times slowed the pace of execution of government contracts as well as new contract generation. Additionally, the COVID-19 pandemic, and the secondary effects of the pandemic have increased the risk of delays in connection with a broad range of operational activities, including: supply chain procurement of raw materials and manufacturing; and certain research and development activities, such as those that involve clinical trials. Furthermore, the pandemic and related secondary effects could result in a slower pace of future product deliveries if there are shortages or delays in the receipt by the supply chain of raw materials or supplies, or if labor shortages become more acute. While the Company does not currently expect such

delays to have a material adverse impact on the financial condition of the Company or its long-term operating performance, and while the COVID-19 pandemic has not adversely affected the liquidity position of the Company, the Company cannot give assurances as to the full extent of the impact at this time.

Procurement Contracts with the U.S. Government

19C BARDA Contract

On September 10, 2018, the Company entered into a contract with the U.S. Biomedical Advanced Research and Development Authority ("BARDA") pursuant to which SIGA agreed to deliver up to 1,488,000 courses of oral TPOXX® to the U.S. Strategic National Stockpile ("Strategic Stockpile"), and to manufacture and deliver to the Strategic Stockpile, or store as vendor-managed inventory, up to 212,000 courses of the intravenous (IV) formulation of TPOXX® ("IV TPOXX®"). Additionally, the contract includes funding from BARDA for a range of activities, including: advanced development of IV TPOXX®, post-marketing activities for oral and IV TPOXX®, and procurement activities. As of September 30, 2022, the contract with BARDA (as amended, modified, or supplemented from time to time, the "19C BARDA Contract") contemplates up to approximately \$602.5 million of payments, of which approximately \$51.7 million of payments are included within the base period of performance of five years, approximately \$268.9 million of payments are related to exercised options and up to approximately \$281.9 million of payments are currently specified as unexercised options. BARDA may choose in its sole discretion when, or whether, to exercise any of the unexercised options. The period of performance for options is up to ten years from the date of entry into the 19C BARDA Contract and such options could be exercised at any time during the contract term, including during the base period of performance.

The base period of performance specifies potential payments of approximately \$51.7 million for the following activities: payments of approximately \$11.1 million for the delivery of approximately 35,700 courses of oral TPOXX® to the Strategic Stockpile; payments of \$8.0 million for the manufacture of 20,000 courses of final drug product of IV TPOXX® ("IV FDP"), of which \$3.2 million of payments are related to the manufacture of bulk drug substance ("IV BDS") to be used in the manufacture of IV FDP; payments of approximately \$32.0 million to fund advanced development of IV TPOXX®; and payments of approximately \$0.6 million for supportive procurement activities. As of September 30, 2022, the Company has received \$11.1 million for the delivery of approximately 35,700 courses of oral TPOXX® to the Strategic Stockpile, \$3.2 million for the manufacture of IV BDS, \$4.3 million for the delivery of IV FDP to the Strategic Stockpile and \$17.1 million for other base period activities. IV BDS has been used for the manufacture of courses of IV FDP. The \$3.2 million received for the completed manufacture of IV BDS had been recorded as deferred revenue as of December 31, 2021, but with the delivery of IV FDP to the Strategic Stockpile during the nine months ended September 30, 2022, \$2.9 million was recognized as revenue. The remaining \$0.3 million of deferred revenue will be recognized as IV FDP containing such IV BDS is delivered to the Strategic Stockpile.

The options that have been exercised to date provide for payments up to approximately \$268.9 million. There are exercised options for the following activities: payments up to \$11.2 million for the procurement of raw materials used in the 2020 manufacture of certain courses of oral TPOXX®; payments up to \$213.9 million for the delivery of up to 726,140 courses of oral TPOXX®; payments up to \$25.6 million for the manufacture of courses of IV FDP, of which \$10.2 million of payments relate to the manufacture of IV BDS to be used in the manufacture of IV FDP, payments of up to approximately \$3.6 million to fund post-marketing activities for IV TPOXX®, and payments of up to \$14.6 million for funding of post-marketing activities for oral TPOXX®. As of September 30, 2022, the Company has received \$225.1 million for the delivery (and related procurement of raw materials) of oral TPOXX® to the Strategic Stockpile; received \$10.2 million for the completed manufacture of IV BDS, which has been recorded as deferred revenue as of September 30, 2022; and received or billed \$8.1 million in connection with post-marketing activities for oral TPOXX®.

Unexercised options specify potential payments up to approximately \$281.9 million in total (if all such options are exercised). There are options for the following activities: payments of up to \$225.1 million for the delivery of oral TPOXX® to the Strategic Stockpile; payments of up to \$51.2 million for the manufacture of courses of IV FDP, of which up to \$20.5 million for payments would be paid upon the manufacture of IV BDS to be used in the manufacture of IV FDP; and payments of up to approximately \$5.6 million for supportive procurement activities.

The options related to IV TPOXX® are divided into two primary manufacturing steps. There are options related to the manufacture of bulk drug substance ("IV BDS Options"), and there are corresponding options (for the same number of IV courses) for the manufacture of final drug product ("IV FDP Options"). BARDA may choose to exercise any, all, or none of these options in its sole discretion. The 19C BARDA Contract includes: three separate IV BDS Options, each providing for the bulk drug substance equivalent of 64,000 courses of IV TPOXX®; and three separate IV FDP Options, each providing for 64,000 courses of final drug product of IV TPOXX®. BARDA has the sole discretion as to whether to simultaneously exercise IV BDS Options and IV FDP Options, or whether to exercise options at different points in time (or alternatively, to only exercise the IV BDS Option but not the IV FDP Option). To date, BARDA has exercised one of the three IV BDS options and one of the three IV FDP options, both of which were exercised simultaneously in 2022. If BARDA decides to only exercise the remaining IV BDS Options, then the Company would receive payments up to \$20.5 million; alternatively, if BARDA decides to exercise all the remaining IV BDS Options and IV FDP Options, then the Company would receive payments up to \$51.2 million. For each set of options relating to a specific group of courses (for instance, the IV BDS and IV FDP options that reference the same 64,000 courses), BARDA has the option to independently purchase IV BDS or IV FDP. The Company estimates that sales of the IV formulation under this contract (under current terms), assuming the IV FDP Options were exercised, would have a gross margin (sales less cost of sales, as a percentage of sales) that is less than 40%.

Under the terms of this contract, exercise of procurement options is at the sole discretion of BARDA. The request for proposal that preceded the award of the 19C BARDA Contract indicated that the expected purpose of the contract was to maintain the level of smallpox antiviral preparedness in the Strategic Stockpile. Based on prior product delivery activity, and current FDA-approved shelf life of oral TPOXX®, the Company estimates that approximately 940,000 courses of smallpox antiviral treatment would need to be delivered to the U.S. Government between 2022 and 2024 in order to maintain stockpile levels of unexpired smallpox antiviral treatment during this period.

U.S. Department of Defense Procurement Contracts

On May 12, 2022, the Company announced a contract with the U.S. Department of Defense ("DoD") for the procurement of oral TPOXX® ("DoD Contract #1"). The DoD Contract #1 included a firm commitment for the DoD to procure approximately \$3.6 million of oral TPOXX®, and an option, exercisable at the sole discretion of the DoD, for the procurement of approximately \$3.8 million of oral TPOXX®. In the second quarter of 2022, the Company delivered and recognized revenue of \$3.6 million for the delivery of oral TPOXX® to the DoD, fulfilling the firm commitment in DoD Contract #1. In the third quarter of 2022, the DoD exercised the option for \$3.8 million of oral TPOXX® and the Company satisfied its obligation by delivering product and recognized the related revenue in September 2022.

On September 28, 2022, the Company and the DoD signed a new procurement contract ("DoD Contract #2"). The DoD Contract #2 includes a firm commitment for the DoD to procure approximately \$5.2 million of oral TPOXX®, and an option, exercisable at the sole discretion of the DoD for the procurement of approximately \$5.5 million of oral TPOXX®.

International Procurement Contracts

This year, through October 31, 2022, the Company has received firm commitment orders from 13 international customers (including Canada) for the delivery of approximately \$77 million of oral TPOXX®, of which approximately \$39 million is for Canada and approximately \$38 million is for jurisdictions in Europe, Asia-Pacific, and the Middle East. Additionally, the contract with CDND (defined below) has an option until March 31, 2024, exercisable at the sole discretion of CDND, for the purchase of up to an additional \$6 million of oral TPOXX®. With respect to the \$77 million of firm commitment orders that have been received this year, approximately \$66 million of oral TPOXX® was delivered and recorded as revenue in the nine months ended September 30, 2022, and the remaining orders are expected to be fulfilled between November 1, 2022 and July 31, 2023. Through an International Promotion Agreement (defined below), Meridian Medical Technologies, Inc. ("Meridian") is the counterparty to international contracts under which orders are placed for the purchase of oral TPOXX®. The Public Health Agency of Canada ("PHAC") and the Canadian Department of National Defence ("CDND") are among the contracting parties for the purchase of oral TPOXX® (see below for a summary description of these contracts).

On January 13, 2021, PHAC awarded a contract to Meridian (the "PHAC Contract") for the purchase of up to approximately \$33 million of oral TPOXX® (tecovirimat) within five years. In March 2022 and July 2022, PHAC executed amendments in which total procurement of oral TPOXX® under the PHAC Contract was increased to an amount of approximately \$45 million. Prior to 2022, approximately \$10 million of oral TPOXX® had been ordered and delivered to PHAC. No courses of oral TPOXX® were delivered under this contract for the first six months of 2022. During the three months ended September 30, 2022, approximately \$35 million of oral TPOXX® was delivered to PHAC and recognized as revenue. Including the deliveries in the third quarter of 2022, there are no current remaining amounts specified in the PHAC Contract.

On April 3, 2020, the Company announced that the CDND awarded a contract (the "Canadian Military Contract") to Meridian, pursuant to which the CDND would purchase up to approximately \$14 million of oral TPOXX® over four years. Prior to 2022, approximately \$4 million of oral TPOXX® had been ordered and delivered to CDND. No courses of oral TPOXX® were delivered under this contract for the first nine months of 2022. As of September 30, 2022, an approximate firm commitment order of \$4 million remains to be delivered under this contract. Additionally, there are approximately \$6 million of unexercised options, exercisable at the sole discretion of CDND, remaining under this contract.

The above-listed contract awards were coordinated between SIGA and Meridian under the international promotion agreement (as amended, the "International Promotion Agreement") that was entered into by the parties on June 3, 2019. Under the International Promotion Agreement, Meridian is the counterparty in connection with international contracts for oral TPOXX® and SIGA is responsible for manufacture and delivery of any oral TPOXX® purchased thereunder.

International Promotion Agreement

Under the terms of the International Promotion Agreement, Meridian was granted exclusive rights to market, advertise, promote, offer for sale, or sell oral TPOXX® in a field of use specified in the International Promotion Agreement in all geographic regions except for the United States (the "Territory"), and Meridian has agreed not to commercialize any competing product, as defined in the International Promotion Agreement, in the specified field of use in the Territory. SIGA retains ownership, intellectual property, distribution and supply rights and regulatory responsibilities in connection with TPOXX®, and, in the United States market, also retains sales and marketing rights with respect to oral TPOXX®. SIGA's consent is required for the entry into any sales arrangement pursuant to the International Promotion Agreement.

The fee Meridian retains pursuant to the International Promotion Agreement is a specified percentage of the collected proceeds of sales of oral TPOXX® net of certain expenses, for calendar years in which customer collected amounts net of such expenses are less than or equal to a specified threshold, and a higher specified percentage of such collected net proceeds for calendar years in which such net collected amounts exceed the specified threshold. It is probable that we will exceed the specified threshold in 2022 and, as a result, the Company has recorded and will continue to record the higher specified percentage for all International Promotion Agreement sales in 2022. Taking into account Meridian's fee and manufacturing costs of oral TPOXX®, it is currently estimated by the Company that international sales of oral TPOXX® will have a contribution margin (as expressed as a percentage of product sales, and before any consideration of expenses not directly related to manufacturing or Meridian activities) of between approximately 65% and 80%, depending on the international sales levels each year. For purposes of this disclosure, contribution margin (in amount) represents international product sales less applicable cost of sales and the Meridian fee (which is included within selling, general and administrative expenses within the income statement).

Research Agreements and Grants

In July 2019, the Company was awarded a multi-year research contract valued at a total of \$19.5 million, with an initial award of \$12.4 million, from the DoD to support work in pursuit of a potential label expansion for oral TPOXX® that would include post-exposure prophylaxis ("PEP") of smallpox (such work known as the "PEP Label Expansion Program" and the contract referred to as the "PEP Label Expansion R&D Contract"). In subsequent modifications, the DoD increased the scope and the available funding under the PEP Label Expansion R&D Contract to approximately \$27 million. The period of performance for this contract, as modified, terminates on January 31, 2025. As of September 30, 2022, remaining revenue to be recognized in the future under the PEP Label Expansion R&D Contract is up to \$11.2 million. Revenue from the performance obligation under the PEP Label Expansion R&D Contract is recognized over time using an input method using costs incurred to date relative to total estimated costs at completion.

Contracts and grants include, among other things, options that may or may not be exercised at the U.S. Government's discretion. Moreover, contracts and grants contain customary terms and conditions including the U.S. Government's right to terminate or restructure a contract or grant for convenience at any time. As such, the Company may not be eligible to receive all available funds.

Critical Accounting Estimates

The methods, estimates and judgments we use in applying our accounting policies have a significant impact on the results we report in our condensed consolidated financial statements, which we discuss under the heading "Results of Operations" following this section of our Management's Discussion and Analysis of Financial Condition and Results of Operations. Some of our accounting policies require us to make difficult and subjective judgments, often as a result of the need to make estimates of matters that are inherently uncertain. Information regarding our critical accounting policies and estimates appears in Part II, Item 7, Management's Discussion and Analysis of Financial Condition and Results of Operations of our 2021 Form 10-K. Our most critical accounting estimates include revenue recognition over time, the valuation of stock-based awards issued by the Company and income taxes.

Results of Operations

Three Months Ended September 30, 2022 and 2021

For the three months ended September 30, 2022, revenues from product sales and supportive services were \$65.6 million. Such revenues primarily relate to international sales of oral TPOXX® of approximately \$61.3 million and sales of oral TPOXX® to the DoD of approximately \$3.8 million. For the three months ended September 30, 2021, revenues from product sales and supportive services were \$2.3 million, which primarily relate to the acceptance of courses of oral TPOXX® delivered to CDND.

Revenues from research and development activities for the three months ended September 30, 2022 and 2021, were \$6.6 million and \$2.5 million, respectively. The increase of \$4.1 million of revenue is primarily related to an increase in clinical trial activity under the PEP Label Expansion R&D Contract in connection with the PEP development program, as well as an increase in R&D activity under the BARDA Contract.

Cost of sales and supportive services for the three months ended September 30, 2022 and 2021 were \$3.9 million and \$0.1 million, respectively. Such costs in 2022 were associated with the manufacture and delivery of courses of oral TPOXX® for international sales and sales to the DoD. Such costs in 2021 were primarily associated with the manufacture and delivery of courses of oral TPOXX® to CDND.

Selling, general and administrative ("SG&A") expenses for the three months ended September 30, 2022 and 2021 were \$19.7 million and \$4.5 million, respectively. The increase of approximately \$15.2 million mostly reflects promotion fees incurred in connection with international sales that substantially increased from the comparable period in 2021.

Research and development ("R&D") expenses for the three months ended September 30, 2022 and 2021 were \$5.7 million and \$3.2 million, respectively, reflecting an increase of approximately \$2.5 million. The increase is primarily attributable to an increase in direct vendor-related expenses incurred in connection with activities under the PEP Label Expansion R&D Contract and the BARDA Contract.

Changes in the fair value of the liability-classified warrant to acquire common stock were recorded within the statement of operations. The warrant was fully exercised during the three months ended June 30, 2022. For the three months ended September 30, 2022, we recorded no activity. For the three months ended September 30, 2021, we recorded a loss of approximately \$1.1 million, reflecting an increase in the fair value of the liability-classified warrant primarily due to the increase in our stock price.

For the three months ended September 30, 2022 and 2021, we recorded pre-tax income/(loss) of \$43.1 million and (\$4.0) million, respectively, and a corresponding income tax (provision)/benefit of (\$10.1) million and \$0.9 million, respectively. The effective tax rates during the three months ended September 30, 2022 and 2021 were 23.4% and 21.9%, respectively. Our effective tax rate for the periods ended September 30, 2022 and 2021 differ from the statutory rate primarily as a result of state taxes, and non-deductible executive compensation under Internal Revenue Code Section 162(m).

Nine Months Ended September 30, 2022 and 2021

For the nine months ended September 30, 2022, revenues from product sales and supportive services were \$81.6 million. Such revenues primarily relate to approximately \$66.2 million of international sales of oral TPOXX®; approximately \$7.5 million of oral TPOXX® sales to the DoD; and approximately \$7.2 million of sales of IV TPOXX® to the U.S. Government under the 19C BARDA Contract. For the nine months ended September 30, 2021, revenues from product sales and supportive services were \$12.8 million. Such revenues in 2021 primarily relate to the delivery and acceptance of courses of oral TPOXX® delivered to PHAC and CDND.

Revenues from research and development activities for the nine months ended September 30, 2022 and 2021, were \$17.9 million and \$5.5 million, respectively. Most of the increase of \$12.4 million relates to clinical trial activity under the PEP Label Expansion R&D Contract in connection with the PEP development program, with R&D activity under the BARDA Contract also contributing to the increase.

Cost of sales and supportive services for the nine months ended September 30, 2022 and 2021 were \$9.6 million and \$1.3 million, respectively. The increase mostly relates to an increase in international sales of oral TPOXX® and approximately \$4.4 million of cost for the manufacture and sale of IV TPOXX® in 2022; manufacturing costs per unit are higher for IV TPOXX® than oral TPOXX®.

SG&A expenses for the nine months ended September 30, 2022 and 2021 were \$29.2 million and \$14.1 million, respectively. The increase of approximately \$15.1 million mostly reflects promotion fees incurred in connection with international sales that substantially increased from the comparable period in 2021.

R&D expenses for the nine months ended September 30, 2022 and 2021 were \$16.1 million and \$7.8 million, respectively. The increase of \$8.3 million is mostly attributable to an increase in direct vendor-related expenses incurred in connection with activities under the PEP Label Expansion R&D Contract and the BARDA Contract.

Changes in the fair value of the liability-classified warrant to acquire common stock were recorded within the statement of operations. The warrant was fully exercised during the nine months ended September 30, 2022. For the nine months ended September 30, 2022 and 2021, we recorded a gain of approximately \$0.4 million and \$0.3 million, respectively, reflecting a decrease in the fair value of the liability-classified warrant primarily due to the decrease in our stock price.

For the nine months ended September 30, 2022, we recorded pre-tax income of \$45.3 million and a corresponding income tax provision of (\$10.5) million. For the nine months ended September 30, 2021, we recorded a pre-tax loss of (\$4.6) million and a corresponding income tax benefit of \$0.8 million. The effective tax rates during the nine months ended September 30, 2022 and 2021 were 23.3% and 17.7%, respectively. Our effective tax rate for the period ended September 30, 2022 differs from the statutory rate primarily as a result of state taxes and non-deductible executive compensation under Internal Revenue Code Section 162(m). Our effective tax rate for the nine months ended September 30, 2021 differs from the statutory rate primarily as a result of state taxes, non-deductible executive compensation under Internal Revenue Code Section 162(m) and a non-taxable adjustment for the fair market value of the Warrant.

Liquidity and Capital Resources

As of September 30, 2022, we had \$109.7 million in cash and cash equivalents compared with \$103.1 million at December 31, 2021.

Operating Activities

We prepare our condensed consolidated statement of cash flows using the indirect method. Under this method, we reconcile net income/(loss) to cash flows from operating activities by adjusting net income/(loss) for those items that impact net income/(loss) but may not result in actual cash receipts or payments during the period. These reconciling items include but are not limited to stock-based compensation, deferred income taxes, changes in the fair value of our warrant liability, inventory write offs, gains and losses from various transactions and changes in the condensed consolidated balance sheet for working capital from the beginning to the end of the period.

Net cash provided by/(used in) operating activities for the nine months ended September 30, 2022 and 2021 was \$49.7 million and (\$4.6) million, respectively. For the nine months ended September 30, 2022, the receipt of approximately \$80 million for product delivery and acceptance of oral TPOXX® courses delivered to the Strategic Stockpile in December 2021, as well as the receipt of approximately \$23 million in connection with 2022 product deliveries and advance payments were partially offset by the payment of approximately \$19 million of federal income taxes in connection with the 2021 tax year; an increase in inventory investment in connection with a broadening of the customer base for TPOXX® and mitigation of increasing general supply chain risks; and costs in relation to customary operating activities. For the nine months ended September 30, 2021, net cash usage related to support of ordinary working capital (accounts receivable, accounts payable, inventory, and prepaids, among other items) and customary operating activity was partially offset by the receipt of cash from international sales.

Investing Activities

There was no cash-related investing activity for the nine months ended September 30, 2022. For the nine months ended September 30, 2021, we used cash of \$24,424 for capital expenditures.

Financing Activities

Cash used in financing activities for the nine months ended September 30, 2022 was \$43.1 million, which was primarily attributable to our special cash dividend of approximately \$32.9 million. In addition, we repurchased approximately 1.5 million shares of common stock for approximately \$10.1 million. Cash used in financing activities for the nine months ended September 30, 2021 was \$20.4 million, which was substantially all attributable to our repurchase of approximately 3.0 million shares of common stock.

On May 5, 2022, the Board of Directors declared a special dividend of \$0.45 per share on the common stock of the Company, which resulted in an overall dividend payment of \$32.9 million. The special dividend was paid on June 2, 2022 to shareholders of record at the close of business on May 17, 2022.

Future Cash Requirements

As of September 30, 2022, we had outstanding purchase orders associated with manufacturing obligations in the aggregate amount of approximately \$21.8 million.

Off-Balance Sheet Arrangements

The Company does not have any off-balance sheet arrangements.

Recently Issued Accounting Standards

The Company did not adopt any accounting standards this quarter.

Safe Harbor Statement

Certain statements in this Quarterly Report on Form 10-Q, including certain statements contained in the foregoing "Management's Discussion and Analysis of Financial Condition and Results of Operations," constitute "forward-looking statements" within the meaning of Section 27A of the Securities Act of 1933, as amended (the "Securities Act"), and Section 21E of the Exchange Act, including statements relating to the progress of SIGA's development programs and timelines for bringing products to market, delivering products to the Strategic Stockpile, the enforceability of our procurement contracts, such as the 19C BARDA Contract (the "BARDA Contract"), with BARDA, and responding to the global outbreak of monkeypox. The words or phrases "can be," "expects," "may affect," "may depend," "believes," "estimate," "project" and similar words and phrases are intended to identify such forwardlooking statements. Such forward-looking statements are subject to various known and unknown risks and uncertainties, and SIGA cautions you that any forward-looking information provided by or on behalf of SIGA is not a guarantee of future performance. SIGA's actual results could differ materially from those anticipated by such forward-looking statements due to a number of factors, some of which are beyond SIGA's control, including, but not limited to, (i) the risk that BARDA elects, in its sole discretion as permitted under the BARDA Contract, not to exercise all, or any, of the remaining unexercised options under those contracts, (ii) the risk that SIGA may not complete performance under the BARDA Contract on schedule or in accordance with contractual terms, (iii) the risk that the BARDA Contract, DOD Contract #2 or PEP Label Expansion R&D Contract are modified or canceled at the request or requirement of the U.S. Government, (iv) the risk that the nascent international biodefense market does not develop to a degree that allows SIGA to continue to successfully market TPOXX® internationally, (v) the risk that potential products, including potential alternative uses or formulations of TPOXX® that appear promising to SIGA or its collaborators, cannot be shown to be efficacious or safe in subsequent pre-clinical or clinical trials, (vi) the risk that SIGA or its collaborators will not obtain appropriate or necessary governmental approvals to market these or other potential products or uses, (vii) the risk that SIGA may not be able to secure or enforce sufficient legal rights in its products, including intellectual property protection, (viii) the risk that any challenge to SIGA's patent and other property rights, if adversely determined, could affect SIGA's business and, even if determined favorably, could be costly, (ix) the risk that regulatory requirements applicable to SIGA's products may result in the need for further or additional testing or documentation that will delay or prevent SIGA from seeking or obtaining needed approvals to market these products, (x) the risk that the volatile and competitive nature of the biotechnology industry may hamper SIGA's efforts to develop or market its products, (xi) the risk that changes in domestic or foreign economic and market conditions may affect SIGA's ability to advance its research or may affect its products adversely, (xii) the effect of federal, state, and foreign regulation, including drug regulation and international trade regulation, on SIGA's businesses, (xiii) the risk of disruptions to SIGA's supply chain for the manufacture of TPOXX®, causing delays in SIGA's research and development activities, causing delays or the re-allocation of funding in connection with SIGA's government contracts, or diverting the attention of government staff overseeing SIGA's government contracts, (xiv) the risk that the U.S. or foreign governments' responses (including inaction) to national or global economic conditions or infectious diseases, such as COVID-19, are ineffective and may adversely affect SIGA's business, and (xv) risks associated with responding to the current monkeypox outbreak, as well as the risks and uncertainties included in Item 1A "Risk Factors" of our Annual Report on Form 10-K for the year ended December 31, 2021 and SIGA's subsequent filings with the Securities and Exchange Commission. SIGA urges investors and security holders to read those documents free of charge at the SEC's website at http://www.sec.gov. All such forward-looking statements are current only as of the date on which such statements were made. SIGA does not undertake any obligation to update publicly any forward-looking statement to reflect events or circumstances after the date on which any such statement is made or to reflect the occurrence of unanticipated events. The information contained on any website referenced in this Form 10-Q is not incorporated by reference into this filing.

Item 3. Quantitative and Qualitative Disclosures About Market Risk

Our investment portfolio includes cash and cash equivalents. Our main investment objectives are the preservation of investment capital. We believe that our investment policy is conservative, both in the duration of our investments and the credit quality of the investments we hold. We do not utilize derivative financial instruments, derivative commodity instruments or other market risk sensitive instruments, positions or transactions to manage exposure to interest rate changes. As such, we believe that the securities we hold are subject to market risk and changes in the financial standing of the issuers of such securities and our interest income is sensitive to changes in the general level of U.S. interest rates.

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 of September 30, 2022. The term "disclosure controls and procedures" is defined in Rules 13a-15(e) and 15d-15(e) under the Exchange Act. Management recognizes that any disclosure controls and procedures, no matter how well designed and operated, can only provide reasonable assurance of achieving their objectives, and management necessarily applies its judgment in evaluating the cost-benefit relationship of possible controls and procedures.

Based on that evaluation, our Chief Executive Officer and Chief Financial Officer have concluded that our disclosure controls and procedures were effective as of September 30, 2022.

Changes in Internal Control over Financial Reporting

There have been no changes in our internal control over financial reporting during the quarter ended September 30, 2022 that 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 involved in a variety of claims, suits, investigations and proceedings arising from the ordinary course of our business, including collections claims, breach of contract claims, labor and employment claims, tax related matters and other matters. Although such claims, suits, investigations and proceedings are inherently uncertain and their results cannot be predicted with certainty, we believe that the resolution of such current pending matters, if any, will not have a material adverse effect on our business, condensed consolidated financial position, results of operations or cash flow. Regardless of the outcome, litigation can have an adverse impact on us because of legal costs, diversion of management resources and other factors.

Item 1A. Risk Factors

Our results of operations and financial condition are subject to numerous risks and uncertainties described in our 2021 Annual Report on Form 10-K for the fiscal year ended December 31, 2021. There have been no material changes to the risk factors described in Part I, Item 1A "Risk Factors" of our 2021 Form 10-K.

Item 2. Unregistered Sales of Equity Securities and Use of Proceeds

ISSUER PURCHASES OF EQUITY SECURITIES

			Total Number	
			of Shares	Dollar Value
			Purchased as	of Shares That
			Part of	May Yet Be
	Total Number		Publicly	Purchased
	of Shares	Average Price	Announced	Under the
Period	Purchased	Paid per Share	Programs	Programs
July 1, 2022 to July 31, 2022		\$		\$ 35,325,830
August 1, 2022 to August 31, 2022	-			35,325,830
September 1, 2022 to September 30, 2022			- <u>-</u>	35,325,830
	-	\$		

On August 5, 2021, the Company announced that the Board of Directors authorized a share repurchase program under which the Company may repurchase up to \$50 million of the Company's common stock through December 31, 2023. The Company started repurchasing shares under this program in the fourth quarter of 2021. The timing and actual number of shares repurchased will depend on a variety of factors, including: the timing of procurement orders under government contracts; alternative opportunities for strategic uses of cash; the stock price of the Company's common stock; market conditions; alternative capital management uses of cash; and other corporate liquidity requirements and priorities. No shares were repurchased in the three months ended September 30, 2022.

Item 3. Defaults upon Senior Securities

None.

Item 4. Mine Safety Disclosures

No disclosure is required pursuant to this item.

Item 5. Other Information

No disclosure is required pursuant to this item.

Item 6. Exhibits

Description

Exhibit No.

104

Emiloit 110.	Bescription
<u>3.1</u>	Amended and Restated Certificate of Incorporation of SIGA Technologies, Inc. (incorporated by reference to Exhibit 3.1 to the Current
	Report on Form 8-K of the Company filed on June 16, 2022).
<u>3.2</u>	Amended and Restated By-laws of SIGA Technologies, Inc. (incorporated by reference to Exhibit 3.1 to the Current Report on Form 8-K
	of the Company filed on December 15, 2021).
<u>10.1</u> †	Amendment of Solicitation/Modification of Contract 00009, dated January 27, 2022, to Agreement, dated September 10, 2018, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>10.2</u>	Amendment of Solicitation/Modification of Contract 00010, dated March 29, 2022, to Agreement, dated September 10, 2018, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>10.3</u>	Amendment of Solicitation/Modification of Contract 000011, dated July 26, 2022, to Agreement, dated September 10, 2018, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>10.4</u> †	Amendment of Solicitation/Modification of Contract 000012, dated August 5, 2022, to Agreement, dated September 10, 2018, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>10.5</u> †	Amendment of Solicitation/Modification of Contract 000021, dated September 28, 2022, to Agreement, dated May 13, 2011, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>10.6</u>	Amendment of Solicitation/Modification of Contract 000022, dated September 29, 2022, to Agreement, dated May 13, 2011, by and
	between SIGA and the Biomedical Advanced Research and Development Authority of the United States Department of Health and
	Human Services.
<u>31.1</u>	Certification of Chief Executive Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
<u>31.2</u>	Certification of Chief Financial Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
<u>32.1</u>	Certification of Chief Executive Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
<u>32.2</u>	Certification of Chief Financial Officer Pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
	Inline XBRL Instance Document (the instance document does not appear in the Interactive Data File because its XBRL tags are
101.INS	embedded within the Inline XBRL document)
101.SCH	Inline XBRL Taxonomy Extension Schema
101.CAL	Inline XBRL Taxonomy Extension Calculation Linkbase
101.DEF	Inline XBRL Taxonomy Extension Definition Linkbase
101.LAB	Inline XBRL Taxonomy Extension Label Linkbase
101.PRE	Inline XBRL Taxonomy Extension Presentation Linkbase

[†] Certain portions of this exhibit have been omitted pursuant to Item 601(b)(10) of Regulation S-K.

Cover Page Interactive Data File (formatted in Inline XBRL and contained in Exhibit 101).

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.

SIGA TECHNOLOGIES, INC. (Registrant)

ate: November 3, 2022 By: /s/ Daniel J. Luckshire

Daniel J. Luckshire

Executive Vice President and Chief Financial Officer

(Duly Authorized Officer, Principal Financial Officer and Principal

Accounting Officer)

CONFIDENTIAL INFORMATION CONTAINED IN THIS DOCUMENT, MARKED BY "[***]," HAS BEEN OMITTED BECAUSE IT IS BOTH (I) NOT MATERIAL AND (II) THE TYPE THAT THE REGISTRANT TREATS AS PRIVATE OR CONFIDENTIAL.

2. AMENDMENT/MODIFICATION NO. P00009 6. ISSUED BY COI	3. EFFECTIVE DATE See Block 16C DE ASPR-BARDA	NO.	TION/PURCHASE REQ.	1 2 5. PROJECT NO. (If applicable) ASPR-21-02282
6. ISSUED BY COL	DE ASPR-BARDA	7 ADMIN		A3FR-21-02202
		Item 6)	NISTERED BY (If other than	CODE
ASPR-BARDA 200 Independence Ave., S.W. Room 640-G Washington DC 20201 8. NAME AND ADDRESS OF CONTRACTOR (No., street, Code)	county, State and ZIP	(x)	9A. AMENDMENT OF SOI	LICITATION NO.
SIGA TECHNOLOGIES, INC. 1385150 Attn: Daniel Luckshire SIGA TECHNOLOGIES, INC. 31 East 62nd street NEW YORK NY 100658446				
1.2.1 10140111 100000110			9B. DATED (SEE ITEM 11)	
<u>. </u>		X	10A. MODIFICATION OF C NO. HHSO100201800019C	
CODE 1385150	FACILITY CODE		10B. DATED (SEE ITEM 13 09/10/2018	
11. THIS ITEM ONLY API	LIES TO AMENDA	LENTS OF		
amendment on each copy of the offer submitted; or (c) By solicitation and amendment numbers. FAILURE OF YOU FOR THE RECEIPT OF OFFERS PRIOR TO THE HOUR virtue of this amendment you desire to change an offer alre provided each letter or electronic communication makes re hour and date specified. 12. ACCOUNTING AND APPROPRIATION DATA (<i>If requ</i>	R ACKNOWLEDGE R AND DATE SPECII ady submitted, such c ference to the solicitat	MENT TO FIED MAY hange may	BE RECEIVED AT THE PLATESULT IN REJECTION OF be made by letter or electronic	ACE DESIGNATED F YOUR OFFER. If by c communication,
See Schedule 13. THIS ITEM ONLY APPLIES TO MODIFICATION	OF CONTRACTS/0	ORDERS.	IT MODIFIES THE CONTI	RACT/ORDER NO.
AS	DESCRIBED IN ITE	EM 14.		
	S SET FORTH IN IT		URSUANT TO: (Specify auth E MADE IN THE CONTRAC	
ADMINIS SET FOR	STRATIVE CHANGE TH IN ITEM 14, PUF	S (such as SUANT T	CT/ORDER IS MODIFIED TO changes in paying office, appr O THE AUTHORITY OF FAI T IS ENTERED INTO PURSI	opriation data, etc.) R 43.103(b).
AUTHOR		A 14 37		
	43-1 Changes Fixed-P R (<i>Specify type of mod</i>		nd authority)	
			copies to the issuing office.	
14. DESCRIPTION OF AMENDMENT/MODIFICATION (Owhere feasible.) Tax ID Number: 13-3864870 DUNS Number: 932651516				ract subject matter
The purpose of this No-Cost Extension (NCE) modification is contract items No 7 & 8 from 12/31/2021 to 03/31/2022. This clarifies the contract's full performance end date which remai under the base award. In addition, this modification revises Part 1 - The Schedule, A Officer is changed from James Harris to Monica Watson. No other changes apply.	modification adminis ns 09/09/2028 as estal	tratively olished		

Continued ...

Except as provided herein, all terms and conditions of the document referenced in Item 9A or 10A, as heretofore changed, remains unchanged and in full force and effect.

15A. NAME AND TITLE OF SIGNER (Type or print)		16A. NAME AND TITLE OF CONTRAC	CTING OFFICER
DENNIS HRUBY, CSO & EVP		(Type or print)	
		MONICA WATSON	
15B. CONTRACTOR/OFFEROR	15C. DATE	16B. UNITED STATES OF AMERICA	16C. DATE
	SIGNED		SIGNED
/s/ Dennis Hruby		/s/ Monica Watson	
(Signature of person authorized to sign)	01/27/2022	(Signature of Contracting Officer)	01/27/2022
Previous edition unusable	•	1, 2	STANDARD FORM
			30 (REV. 11/2016)
			Prescribed by GSA
		I	FAR (48 CFR) 53.243

CONTINUATION SHEET	REFERENCE NO. OF DOCUMENT BEING CONTINUED HHSO100201800019C/P00009	PA	GE OF
		2	2

NAME OF OFFEROR OR CONTRACTOR SIGA TECHNOLOGIES INC 1385150

	(B)	QUANTITY (C)	UNIT (D)	PRICE	AMOUNT
			(D)	(E)	(F)
	Terms: 1/30, NET 30P				
	Performance: 01/01/2020 to 09/09/2028				
	em 7 to read as follows (amount shown is the obligated amount):				
[***]	ery of [***] courses [***] =				0.00
Obligated	Amount: \$0.00				
Accountin 2021.1998	g Info: SN20.26088 Appr. Yr.: 2021 CAN: 199SN20				
Object Cla Funded: \$	ass: 26088				
Change It	em 8 to read as follows (amount shown is the obligated amount):				
8 Second de	livery of [***] courses [***] =				0.00
	Amount: \$0.00				
Accountin	g Info: SN11.26088 Appr. Yr.: 2021 CAN: 199SN11				
Object Cla Funded: \$	ass: 26088				
i unαcα. φ	0.00				
	ct code: ASPR-21-02282; BARDA project codes: ASPR-20-01588 -20-02051; PSC: AN13 NAICS: 541714 HHS/BARDA COR is Dr.				
	Lu, xi.lu@hhs.gov, (202) 604-5814.				

NSN 7540-01-152-8067 OPTIONAL FORM 336 (4-86) Sponsored by GSA FAR (48 CFR) 53.110

AMENDMENT OF SOLICITATION/MODIFICATION OF CONTRACT			1. CONTRACT ID CODE					
2. AMENDMENT/MODIFICAT P00010	TION NO.	3. EFFECTIVE DATE See Block 16C	4.REC	5. PROJECT NO. (If applicable) ASPR-21-02282	1 2			
6. ISSUED BY	CODE	ASPR- BARDA	7. AD	` '	CODE			
ASPR-BARDA 200 Independence Ave., S.W. Room 640-G Washington DC 20201								
8. NAME AND ADDRESS OF county, State and ZIP Code)	CONTRACTOR (I	No., street,	(x)	9A. AMENDMENT OF SOLICITATI	ON NO.			
SIGA TECHNOLOGIES, INC. Attn: Daniel Luckshire SIGA TECHNOLOGIES, INC. 31 East 62nd street NEW YORK NY 100658446	1385150							
1.2.1. 1.01.01.1.1 1.00.000 1.10				9B. DATED (SEE ITEM 11)				
			X	10A. MODIFICATION OF CONTRA	CT/ORDER NO.			
CODE 1385150		FACILITY	1	HHSO100201800019C 10B. DATED (<i>SEE ITEM 13</i>)				
	THE TEN ON	CODE		09/10/2018				
				ENDMENTS OF SOLICITATIONS 1. The hour and date specified for recei	ot of Offers			
following methods: (a) By con amendment on each copy of the solicitation and amendment of DESIGNATED FOR THE RE YOUR OFFER. If by virtue o	mpleting Items 8 ame offer submitted; umbers. FAILURE CEIPT OF OFFEI f this amendment yovided each letter conour and date speci	nd 15, and return or (c) By separa or (c) By separa or YOUR AC RS PRIOR TO To you desire to chap or electronic conified.	ning ate lette KNOW THE HO ange an	ad date specified in the solicitation or as copies of the amendment; (b) By or or electronic communication which in ILEDGEMENT TO BE RECEIVED A DUR AND DATE SPECIFIED MAY R offer already submitted, such change nation makes reference to the solicitation	acknowledging receipt of this ncludes a reference to the T THE PLACE ESULT IN REJECTION OF nay be made by letter or			
See Schedule		TICATION OF		RACTS/ORDERS. IT MODIFIES T	HE CONTRACT/ORDER			
CHECK ONE	A. THIS CHANG			D IN ITEM 14. PURSUANT TO: (Specify authority) T	THE CHANGES SET			
				IE CONTRACT ORDER NO. IN ITEM				
	ADMINISTRATI	VE CHANGES	(such a	ACT/ORDER IS MODIFIED TO REFI s changes in paying office, appropriati THORITY OF FAR 43.103(b).				
X	C. THIS SUPPLE FAR Part 43.103(a			NT IS ENTERED INTO PURSUANT ons	TO AUTHORITY OF:			
	D. OTHER (Speci	ify type of modifi	ication	and authority)				
E. IMPORTANT: Contractor 14. DESCRIPTION OF AMENI matter where feasible.) Tax ID Number: 13-3864870 DUNS Number: 932651516 The purpose of this no cost bilate 352.232-71: Electronic Submissi 1. The following is hereby incorp Government, into Section I:	DMENT/MODIFIC eral modification is on of Payment Rec	CATION (<i>Organ</i> s to incorporate l quests	ized by HHSAF					
HHSAR Clause 352.232-71 - Ele	ectronic Submissio	on of Payment R	equests					
	oucher, invoice, or terms and conditi	-		ancing payment eferenced in Item 9A or 10A, as here	tofore changed, remains			
unchanged and in full force an 15A. NAME AND TITLE OF S		rint)	16A. N	NAME AND TITLE OF CONTRACTI	NG OFFICER(Type or print))		
DENNIS HRUBY, CSO & EVP				JONATHAN F. GONZALEZ				

15B. CONTRACTOR/OFFEROR /s/ Dennis Hruby (Signature of person authorized to sign) 15C. DATE SIGNED 03/24/2022

16B. UNITED STATES OF

AMERICA /s/ Jonathan F. Gonzalez (Signature of Contracting Officer) 16C. DATE SIGNED

03/29/2022

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STANDARD FORM 30 (REV. 11/2016) Prescribed by GSA FAR (48 CFR) 53.243

CONTINUATION	
SHEET	

REFERENCE NO. OF DOCUMENT BEING	PAGE OF
CONTINUED	
HHSO100201800019C/P00010	
	1 2

NAME OF OFFEROR OR CONTRACTOR
SIGA TECHNOLOGIES, INC. 1385150

ITEM NO	SUPPLIES/SERVICES	QUANTITY	UNIT	UNIT	MOUN
(A)	(B)	(C)	(D)	UNIT PRICE (E)	(F)
	with associated supporting documentation. The payment request must comply with the requirements identified in FAR 32.905(b), ?Content of Invoices? and the applicable Payment clause included in this contract.				
	(b) Except as provided in paragraph (c) of this clause, the Contractor shall submit payment requests electronically using the Department of Treasury Invoice Processing Platform (IPP) or successor system. Information regarding IPP, including IPP Customer Support contact information, is available at www.ipp.gov or any successor site.				
	(c) The Contractor may submit payment requests using other than IPP only when the Contracting Officer authorizes alternate procedures in writing in accordance with HHS procedures.				
	(d) If alternate payment procedures are authorized, the Contractor shall include a copy of the Contracting Officer's written authorization with each payment request.				
	(End of clause)				
	2. The contracting Officer is hereby changed to Jonathan Gonzalez.				
	3. The total amount, scope, period of performance and all other terms and conditions of the contract remain unchanged.				
	4. By signing this modification, SIGA Technologies Inc., hereby releases the Government from any and all liability under this contract for further equitable adjustments attributable to such fact or circumstance giving rise to this modification. Discount Terms: 1/30, NET 30P Period of Performance: 01/01/2020 to 09/09/2028				
	END OF MODIFICATION				

NSN 7540-01-152-8067

OPTIONAL FORM 336 (4-86) Sponsored by GSA FAR (48 CFR) 53.110

AMENDMENT OF SOLICITATION/MODIFICATION OF CONTRACT			1. CONTRACT ID CODE PAC			
2. AMENDMENT/MODIFICATION NO. P00011	3. EFFECTIVE DATE See Block 16C	4. REQUISITION/PURCHASE REQ. NO. 5. PROJECT NO. (If applicable)				
6. ISSUED BY CODE	E ASPR-BARDA	7. ADMINISTERED BY (If other than CODE Item 6)				
ASPR-BARDA 200 Independence Ave., S.W. Room 640-G Washington DC 20201 8. NAME AND ADDRESS OF CONTRACTOR (N	o., street, county,	(x)	9A. AMENDMENT OF SOLICITAT	ION NO.		
State and ZIP Code)	,,,					
SIGA TECHNOLOGIES, INC. 1385150 Attn: Daniel Luckshire SIGA TECHNOLOGIES, INC. 31 East 62nd street NEW YORK NY 100658446			9B. DATED (SEE ITEM 11)			
		X	10A. MODIFICATION OF CONTRA HHSO100201800019C	ACT/ORDER N	0.	
CODE 1385150	FACILITY CODE		10B. DATED (SEE ITEM 13) 09/10/2018		-	
11. THIS ITEM ON		AME	NDMENTS OF SOLICITATIONS			
extended.	d 15, and returning or (c) By separate le OF YOUR ACKNO IE HOUR AND DA an offer already sul	etter or o DWLEI ATE SP omitted	copies of the amendment; (b) By acking copies of the amendment; (b) By acking communication which included a CE RECEIVED AT TO BE RECEIVED AT TO BE RESULT IN REJECT AND AND ASSESSED ASSESSEDAS ASSESSED ASSESSED ASSESSED ASSESSED ASSESSED ASSESSED ASSESSED	nowledging rece des a reference HE PLACE DE ION OF YOUR r electronic com	eipt of this to the SIGNATED & OFFER. If nmunication,	
12. ACCOUNTING AND APPROPRIATION DATA	\ (If required)					
See Schedule 13. THIS ITEM ONLY APPLIES TO MODIFICE 13. THIS ITEM ONLY APPLIES TO MODIFICE 13. THIS ITEM ONLY APPLIES TO MODIFICE 14. THIS ITEM ONLY APPLIES TO MODIFICE 15. THIS ITEM ONLY APPLIES TO MODIFICE 16. THIS ITEM ONLY APPLIES TO MODIFICE 17. THIS ITEM ONLY APPLIES TO MODIFICE 18. THIS ITEM ONLY APPLIES TO MO		TDAC	TS/ODDEDS IT MODIEIES THE	CONTRACT/C	DDDED NO	
	AS DESCRIE	BED IN	ITEM 14.			
	ER IS ISSUED PURSUANT TO: (Specify authority) THE CHANGES SET E MADE IN THE CONTRACT ORDER NO. IN ITEM 10A. ERED CONTRACT/ORDER IS MODIFIED TO REFLECT THE ANGES (such as changes in paying office, appropriation data, etc.) SET RSUANT TO THE AUTHORITY OF FAR 43.103(b).					
ADMIN						
C. THIS OF:	S SUPPLEMENTA	L AGR	EEMENT IS ENTERED INTO PURS	SUANT TO AU	ΓHORITY	
X FAR Part 43.103(a) - Bilateral Modifications						
			ication and authority)			
			ent and return <u>1</u> copies to the issuing o		•	
14. DESCRIPTION OF AMENDMENT/MODIFICA where feasible.) Tax ID Number: 13-3864870 DUNS Number: 932651516 UEI: VJRNRTSL22K4 The purpose of this no cost bilateral modification is a full. The total amount, scope, period of performance and By signing this modification, SIGA Technologies Inc. Continued	to correct the payme	ent tern	ns that were changed via P00008. Veri		•	
Except as provided herein, all terms and condition unchanged and in full force and effect.				re changed, rei	nains	
15A. NAME AND TITLE OF SIGNER (<i>Type or pri</i> DENNIS HRUBY, CSO & EVP	16A. NAME AND TITLE OF CONTRACTING OFFICER (<i>Type or</i> print)					
	l.=0 = :	_	THAN F. GONZALEZ	lia- : :		
15B. CONTRACTOR/OFFEROR	15C. DATE SIGNED		UNITED STATES OF AMERICA	16C. DATE S	IGNED	
<u>/s/ Dennis Hruby</u> 7/26/2022		/s/ Jor	nathan F. Gonzalez	7/26/2022		

(Signature of person authorized to sign) Previous edition unusable

(Signature of Contracting Officer)

STANDARD FORM 30 (REV. 11/2016) Prescribed by GSA FAR (48 CFR) 53.243

	REPERENCE NO. OF DOCUMENT DEING CONTINUED			FAG	L OI
SHEET	HHSO100201800019C/P00011				
				2	2
	ROR OR CONTRACTOR				
	OGIES, INC. 1385150	ľ	Ī		
ITEM NO	SUPPLIES/SERVICES	QUANTITY	UNIT	UNIT PRICE	
(A)	(B)	(C)	(D)	(E)	(F)
	any and all liability under this contract for further equitable adjustments				
	attributable to such fact or circumstance giving rise to this modification.				
	Discount Terms: HHS NET 30P				
	Period of Performance: 01/01/2020 to 09/09/2028				

PAGE OF

OPTIONAL FORM 336

Sponsored by GSA FAR (48 CFR) 53.110

(4-86)

CONTINUATION REFERENCE NO. OF DOCUMENT BEING CONTINUED

NSN 7540-01-152-8067

CONFIDENTIAL INFORMATION CONTAINED IN THIS DOCUMENT, MARKED BY "[***]," HAS BEEN OMITTED BECAUSE IT IS BOTH (I) NOT MATERIAL AND (II) THE TYPE THAT THE REGISTRANT TREATS AS PRIVATE OR CONFIDENTIAL.

AMENDMENT OF SOLICITATION/MODIFICATION OF CONTRACT					1. CONTRACT ID CO		AGE OF AGES 1 5
2. AME P00012	NDMENT/MODIFICATION NO.	3. EFFECTIVE DATE See Block 16C		UISITION/PURCHASE REQ. NO. nedule		5. PROJEC applicable)	T NO. (If
6. ISSU	ED BY CODE	ASPR- BARDA	7. ADM 6)	IINISTERED BY (If other than Item	1	CODE	
Room 6	ependence Ave., S.W.		-			L	
	E AND ADDRESS OF CONTRACTOR State and ZIP Code)	(No., street,	(x)	9A. AMENDMENT OF SOLICITATION NO.			-
	ECHNOLOGIES, INC. 1385150 aniel Luckshire			9B. DATED (SEE ITEM 11)			
SIGA T 31 East	ECHNOLOGIES, INC. 62nd street ORK NY 100658446		X	10A. MODIFICATION OF CONTRACT/ORDER NO. HHSO100201800019C			-
CODE	1385150	FACILITY CODE		10B. DATED (SEE ITEM 13) 09/10/2018			
	11. THIS ITI	EM ONLY API	PLIES T	TO AMENDMENTS OF SOLICIT	TATIONS		<u> </u>
extended Offers methology copy of numb PRIO chang makes	s must acknowledge receipt of this amend ods: (a) By completing Items 8 and 15, and of the offer submitted; or (c) By separate ers. FAILURE OF YOUR ACKNOWLES R TO THE HOUR AND DATE SPECIFICE an offer already submitted, such changes reference to the solicitation and this ame COUNTING AND APPROPRIATION DATE	ment prior to the dreturning letter or electro DGEMENT TO ED MAY RESUE may be made bendment, and is	e hour a _ copies nic com BE REO JLT IN 1	nd date specified in the solicitation of s of the amendment; (b) By acknowle munication which includes a referen CEIVED AT THE PLACE DESIGN. REJECTION OF YOUR OFFER. If or electronic communication, provid	or as amended , by one edging receipt of this a ce to the solicitation at ATED FOR THE REC by virtue of this amended each letter or electr	of the follomendment amendment amendment CEIPT OF (dment you	t on each nent OFFERS desire to
See Sch		DIFICATION (OF CON	TRACTS/ORDERS, IT MODIFU	ES THE CONTRAC	T/ORDER	NO AS
	A. THIS CHANGE ORDER IS ISSUED CONTRACT ORDER NO. IN ITEM 10	D PURSUANT	ESCRII	BED IN ITEM 14.			_
	B. THE ABOVE NUMBERED CONTF in paying office, appropriation data, etc.						ıs changes -
X	C. THIS SUPPLEMENTAL AGREEMI FAR Part 43.103(a) - Bilateral Modifi	ications		O PURSUANT TO AUTHORITY (OF:		
	D. OTHER (Specify type of modified						
E. IMP	ORTANT : Contractor □ is not	⊠ is requi	red to si	gn this document and return	1copies to the	e issuing of	lfice
feasible.	SCRIPTION OF AMENDMENT/MODIF .) Number: 13-3864870	ICATION (Orgo	anized b	y UCF section headings, including s	olicitation/contract su	bject matte	er where

The purpose of this bilateral modification is to exercise and fully fund Option CLINs 0008, 0013, 0015, and 0017.

The total amount, scope, period of performance and all other terms and conditions of the contract remain unchanged.

By signing this modification, SIGA Technologies Inc., hereby releases the Government from Continued...

DUNS Number: 932651516 UEI: VJRNRTSL22K4

15A. NAME AND TITLE OF SIGNER (Type or print) DENNIS HRUBY, CSO & EVP		16A. NAME AND TITLE OF CONTRACTING OFFICER (Type or print) JONATHAN F. GONZALEZ		
15B.	15C. I	DATE	16B. UNITED STATES OF	16C.
CONTRACTOR/OFFEROR	SIGN	ED	AMERICA	DATE SIGNED
/s/ Dennis Hruby		Aug. 5, 2022	/s/ Jonathan F. Gonzalez	Aug. 5, 2022
(Signature of person			(Signature of Contracting Officer)	_
authorized to sign)				
Previous edition unusable				STANDARD FORM 30
				(REV. 11/2016)
				Prescribed by GSA
				FAR (48 CFR) 53.243

CONTI	NUATION SHEET REFERENCE NO. OF DOCUMENT BEING CONTINUED				PAGE OF
	HHSO100201800019C/P00012				2 5
	OF OFFEROR OR CONTRACTOR				
	ECHNOLOGIES, INC. 1385150			, ,	
ITEM	SUPPLIES/SERVICES	QUANTITY			AMOUNT
NO.	(B)	(C)	(D)	PRICE	(F)
(A)				(E)	
	any and all liability under this contract for further equitable adjustments attributable to such fact or				
	circumstance giving rise to this modification.				
	Discount Terms: HHS NET				
	30P				
	Period of Performance:				
	01/01/2020 to 09/09/2028				
	Add Item 9 as follows:				
9	ASPR-22-01725 Siga Technologies to				3,586,806.00
	exercise				
	CLIN0008 post-marketing field study for [***]				
	Accounting Info:				
	2022.1992126.25106 Appr. Yr.: 2022 CAN:				
	1992126 Object Class: 25106				
	Funded: \$3,586,806.00				
	μ απατά, φο,ουο,ουο				
	Add Item 10 as follows:				
10	ASPR-22-01723 Siga Technologies to procure 64				25,625,600.00
	000 TC of the Intravenous (IV) Tecovirimat as defined by CLIN0013 0015 0017 to be delivered to [***]				
	SNS locations Per contract definition 1 TC				
	14 vials under contract				
	HHSO100201800019C				
	Obligated Amount: \$25,625,600.00				
	Requisition No: OS299284				
	Accounting Info:				
	2022.1992126.25103 Appr. Yr.: 2022 CAN:				
	1992126 Object Class: 25103				
	Funded: \$25,625,600.00				
NSN 75	40-01-152-8067	OPTIONAL	FORM	A 336 (4	1-86)
			004	`	-

Sponsored by GSA FAR (48 CFR) 53.110

Base Period Cost Reimbursement CLIN							
Item	Period of Performance	Supplies/Services	Estimated Cost	Fixed Fee	Cost + Fixed Fee (CPFF)		
0001 Base	[***]	Late Stage development activities towards FDA approval for parenteral (IV) antiviral	[***]	[***]	\$32,009,375 (Funded)		
Total			\$30,197,522	\$1,811,853	\$32,009,375		

Base Period I	Fixed Price CLINS				
Item	Period of Performance	Supplies/Services	Units (# of Doses or Dose Equivalents)	Unit Price (\$)	Total (\$)
0002 Base	[***]	Initial purchase and delivery of nonparenteral (oral) formulated antiviral as final drug product (FDP) to SNS	35,718	[***]	\$11,072,580 (Funded)
0003 Base	[***]	Initial procurement of parenteral (IV) formulated antiviral as bulk drug substance (BDS) *course = 14 vials	20,000	[***]	\$3,200,000 (Funded)
0004 Base	[***]	Fill/finish of final drug product (from bulk drug substance procured under CLIN0003)	20,000	[***]	\$4,800,000 (Funded)
0005 Base	[***]	Storage of final drug product in VMI for 5 years (from bulk drug substance procured under CLIN0003) *Monthly rate/TC = [***]	20,000	[***]	[***]
0006 Base	[***]	Delivery of FDP to the SNS (from bulk drug substance procured under CLIN0003)	20,000	[***]	[***]
Total					[***]

Optional Cost	Reimbursement CLINs				
Item	Period of Performance	Supplies/Services	Estimated Cost	Fixed Fee	Cost + Fixed
					Fee (CPFF)
0007 (Option)	[***]	Phase IV post marketing	[***]	[***]	\$40,812,609
		commitments (no parenteral			Funded
		(oral) formulation))			
		including [***]			
8000	[***]	Phase IV post	[***]	[***]	\$3,586,806
(Option)		marketing			Funded
		commitments (parenteral			
		(IV) formulation))			
		including [***]			
Total			\$41,886,242	\$2,513,173	[***]

Optional Fixed CL	1		· · · · · · · · · · · · · · · · · · ·		
Item	Period of Performance	Supplies/Services	Treatment	Unit Price (\$)	Total (\$)
	Periorilance		Courses (# of Product)		
0009A (Option Funded)	1/1/19-12/31/20	Procurement of raw materials used in manufacturing of unmicronized API in sufficient quantity to support the production of 363,070 courses of nonparenteral (oral) formulated antiviral for SNS replenishment. Such raw materials may be forward processed.		[***]	\$11,255,170 Funded
0009B (Option Funded)	1/1/19-12/31/20	Additional procurement of nonparenteral (oral) formulated antiviral as FDP and delivery to the SNS	121,023 (raw material)	[***]	\$33,765,417 Funded
0009C	1/1/19-12/31/20	Additional procurement of	121,023 (raw material)	[***]	\$33,765,417

(Option Funded)		nonparenteral (oral) formulated antiviral as FDP and delivery to the SNS			Funded
0009D (Option Funded)	1/1/19-12/31/20	-	121,024 (raw material)	[***]	\$33,765,696 Funded
0010 (Option Funded)	1/1/20-12/31/21	Additional procurement of nonparenteral (oral) formulated antiviral as FDP and delivery to the SNS	363,070	[***]	\$112,551,700 Funded
0011 (Option)	[***]	Additional procurement of a nonparenteral (oral) formulated antiviral as FDP and delivery to the SNS	363,070	[***]	\$112,551,700
0012 (Option)	[***]	Additional procurement of a nonparenteral (oral) formulated antiviral as FDP and delivery to the SNS	363,072	[***]	\$112,552,320
0013 (Option)	[***]	Surge Capacity – Additional procurement of parenteral (IV) formulated antiviral as bulk drug substance (BDS) *course = 14 vials	64,000	[***]	\$10,240,000 (Funded)
0014 (Option)	[***]	Surge Capacity – Storage of parenteral (IV) formulated antiviral as bulk drug substance (BDS) in VMI for 5 years (from bulk drug substance procured under CLIN0013). *Monthly rate per TC = [***]	64,000	[***]	[***]
0015 (Option)	[***]	Surge Capacity – Fill/finish of final drug product (from bulk drug substance procured under CLIN0013)	64,000	[***]	\$15,360,000 (Funded)
0016 (Option)	[***]	Surge Capacity – Storage of final drug product in VMI for 5 years (from bulk drug substance procured under CLIN0013). *Monthly rate per TC = [***]	64,000	[***]	[***]
0017 (Option)	[***]	Surge Capacity – Delivery of FDP to the SNS (from bulk drug substance procured under CLIN0013) *[***] SNS Shipments @ [***]	64,000	[***]	[***] (Funded)
0018 (Option)	[***]	Surge Capacity – Additional procurement of parenteral (IV) formulated antiviral as bulk drug substance (BDS) *course = 14 vials	64,000	[***]	\$10,240,000
0019 (Option)	[***]	Surge Capacity – Storage of parenteral (IV) formulated antiviral as bulk drug substance (BDS) in VMI for 5 years (from bulk drug substance procured under CLIN0018). *Monthly rate per TC = [***]	64,000	[***]	[***]
0020 (Option)	[***]	Surge Capacity – Fill/finish of final drug product from bulk drug substance procured under CLIN0018).	64,000	[***]	\$15,360,000
0021 (Option)	[***]	Surge Capacity – Storage of final drug product in VMI for 5 years (from bulk drug substance procured under CLIN0018). *Monthly rate per TC: [***]	64,000	[***]	[***]
0022 (Option)	[***]	Surge Capacity – Delivery of FDP to the SNS (from bulk drug substance procured under	64,000	[***]	[***]

		CLIN0018). *[***] SNS Shipments @ [***]			
0023 (Option)	[***]	Surge Capacity – Additional procurement of parenteral (IV) formulated antiviral as bulk drug substance (BDS) *course = 14 vials	64,000	[***]	\$10,240,000
0024 (Option)	[***]	Surge Capacity – Storage of parenteral (IV) formulated antiviral as bulk drug substance (BDS) in VMI for 5 years (from bulk drug substance procured under CLIN0023). *Monthly rate per TC = [***]	64,000	[***]	[***]
0025 (Option)	[***]	Surge Capacity – Fill/finish of final drug product (from bulk drug substance procured under CLIN0023).	64,000	[***]	\$15,360,000
0026 (Option)	[***]	Surge Capacity – Storage of final drug product in VMI for 5 years (from bulk drug substance procured under CLIN0023). *Monthly rate per TC: [***]	64,000	[***]	[***]
0027 (Option)	[***]	Surge Capacity – Delivery of FDP to the SNS (from bulk drug substance procured under CLIN0023). *[***] SNS Shipments @ [***]	64,000	[***]	[***]
Total					[***]

Total Funded	
	\$320,585,711

End of Modification P00012

CONFIDENTIAL INFORMATION CONTAINED IN THIS DOCUMENT, MARKED BY "[***]," HAS BEEN OMITTED BECAUSE IT IS BOTH (I) NOT MATERIAL AND (II) THE TYPE THAT THE REGISTRANT TREATS AS PRIVATE OR CONFIDENTIAL.

AMENDMENT OF SOLICITATION OF CONTRACT	/MODIFICATION		1. CONTRACT ID CODE			E OF GES 3
2. AMENDMENT/MODIFICATION N P00021	O. 3. EFFECTIVE DATE See Block 16C	4. REQUISITION/PURCHASE OS302885	REQ. NO.	5. PROJEC applicable)		
6. ISSUED BY COL	DE ASPR-BARDA	7. ADMINISTERED BY (If other	er than Item 6)	CODE		
ASPR-BARDA 200 Independence Ave., S.W. Room 640-G Washington DC 20201						
8. NAME AND ADDRESS OF CONTI	RACTOR (No., street	, (x) 9A. AMENDMENT OF SO	LICITATION NO.			
SIGA TECHNOLOGIES, INC. 138515 Attn: Daniel Luckshire SIGA TECHNOLOGIES, INC. 31 31 E 62ND ST NEW YORK NY 100658014	0 E 6	9B. DATED (SEE ITEM 11) x 10A. MODIFICATION OF (NO. HHSO100201100001C				
CODE 1385150 FAC	ILITY CODE	10B. DATED (SEE ITEM 13 05/13/2011	?)			
11.	THIS ITEM ONLY	APPLIES TO AMENDMENTS	OF SOLICITATIONS			
☐ The above numbered solicitation is a extended. Offers must acknowledge receipt of methods: (a) By completing Items 8 copy of the offer submitted; or (c) B numbers. FAILURE OF YOUR ACT PRIOR TO THE HOUR AND DAT change an offer already submitted, s makes reference to the solicitation at	this amendment prior and 15, and returning y separate letter or elo KNOWLEDGEMEN E SPECIFIED MAY uch change may be m	to the hour and date specified in to the hour and date specified in to get to the amendment communication which income to be RECEIVED AT THE PRESULT IN REJECTION OF YOu adde by letter or electronic communication.	the solicitation or as ament; (b) By acknowledging reludes a reference to the solution DESIGNATED FOOUR OFFER. If by virtue conication, provided each le	eceipt of this amen olicitation and ame R THE RECEIPT of this amendment	followindment on OF OFF	FERS ire to
12. ACCOUNTING AND APPROPRIA	ATION DATA (If requ	uired)				
See Schedule 13. THIS ITEM ONLY APPLIES	TO MODIFICATIO	ON OF CONTRACTS/ORDERS DESCRIBED IN ITEM 14.	S. IT MODIFIES THE C	ONTRACT/ORE	ER NO	. AS
	GE ORDER IS ISSUE CONTRACT ORDER	ED PURSUANT TO: (Specify auth NO. IN ITEM 10A.	nority) THE CHANGES S	ET FORTH IN IT	EM 14 <i>A</i>	ARE
as changes in pay 43.103(b).	ing office, appropriat	TRACT/ORDER IS MODIFIED T ion data, etc.) SET FORTH IN IT	EM 14, PURSUANT TO	THE AUTHORIT		
	EMENTAL AGREEN (a) - Bilateral Modific	MENT IS ENTERED INTO PURS cations.	UANT TO AUTHORITY	OF:		
	ify type of modification		. ,			
E. IMPORTANT: Contractor is		o sign this document and return 1				
14. DESCRIPTION OF AMENDMENT feasible.) Tax ID Number: 13-3864870	I/MODIFICATION (Organizea by UCF section headin	gs, incluaing solicitation/o	contract subject m	utter wh	ere

The purpose of this modification is to de-obligate excess funds that will cancel at the end of the fiscal year. In addition, replacement funds for those cancelling funds have been added to continue the tasks as outlined in the SOW. The period of performance is also extended until 12/31/2023.

All other terms and conditions remain unchanged.

Continued ...

DUNS Number: 932651516 UEI: VJRNRTSL22K4

Except as provided herein, all terms and conditions of the d	locument referenced	l in Item 9 A or 10A, as heretof	ore changed, remains unchanged		
and in full force and effect.					
15A. NAME AND TITLE OF SIGNER (Type or print)	16A. NAME AND TITLE O	16A. NAME AND TITLE OF CONTRACTING OFFICER (Type			
		or print)			
Dennis Hruby, CSO & EVP					
	JONATHAN F. GONZALEZ	JONATHAN F. GONZALEZ			
15B.	15C.	16B. UNITED STATES OF	16C. DATE SIGNED		
CONTRACTOR/OFFEROR	DATE	AMERICA			
	SIGNED				
/s/ Dennis Hruby	Sept. 26, 2022	/s/ Jonathan F. Gonzalez	Sept. 28, 2022		
		(Signature of Contracting			
(Signature of person authorized to sign)		Officer)			
Previous edition			STANDARD FORM 30		
unusable			(REV. 11/2016)		
			Prescribed by GSA FAR		
			(48 CFR) 53.243		

	REFERENCE NO. OF DOCUMENT BEING]	PAGE	
CONTINUATION SHEET	CONTINUED	OF		
	HHSO100201100001C/P00021			
		2		3

NAME OF OFFEROR OR CONTRACTOR

(B) Discount Terms: HHS NET 14P Period of Performance: 06/28/2016 to 12/31/2023 Change Item 1 to read as follows (amount shown is the obligated amount): Smallpox Antiviral Drug for the Strategic National Stockpile	(C)	(D)	(E)	(F)
Period of Performance: 06/28/2016 to 12/31/2023 Change Item 1 to read as follows (amount shown is the obligated amount): Smallpox Antiviral Drug for the Strategic				
Change Item 1 to read as follows (amount shown is the obligated amount): Smallpox Antiviral Drug for the Strategic				
is the obligated amount): Smallpox Antiviral Drug for the Strategic				
is the obligated amount): Smallpox Antiviral Drug for the Strategic				
Smallpox Antiviral Drug for the Strategic				
National Stockpile				-269,641.
Accounting Info:				
2011.1990001.26402 Appr. Yr.: 2011 CAN:				
1990001 Object Class: 26402				
Funded: \$0.00				
Funded: -\$269,641.07				
A accounting Info.				
is the obligated amount):				
CLIN 0007 Smallpox Antiviral Drug for				-2,336,38
the -				
Strategic National Stockpile [***]				
Accounting Info:				
2013.1992002.25106 Appr. Yr.: 2013 CAN:				
1992002 Object Class: 25106				
Funded: \$0.00				
Funded: -\$1,120,983.54				
Accounting Info				
Accounting Info:				
2016.1992016.25103 Appr. Yr.:2016 CAN				
1992016 Object Class: 25103				
Funded: -\$1,215,399.49				
Continued				
	Strategic National Stockpile [***] Accounting Info: 2013.1992002.25106 Appr. Yr.: 2013 CAN: 1992002 Object Class: 25106 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$1,120,983.54 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN: 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN: 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN: 1992016 Object Class: 25103 Funded: -\$1,215,399.49 FOB: Destination	2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$269,641.07 Accounting Info: 2018.199TWNP.26201 Appr. Yr.: 2018 CAN: 199TWNP Object Class: 26201 Funded: \$0.00 FOB: Destination No-cost time extension for CLIN 0008. [***] Change Item 3 to read as follows (amount shown is the obligated amount): CLIN 0007 Smallpox Antiviral Drug for the - Strategic National Stockpile [***] Accounting Info: 2013.1992002.25106 Appr. Yr.: 2013 CAN: 1992002 Object Class: 25106 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$1,120,983.54 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$1,215,399.49 FOB: Destination	2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$269,641.07 Accounting Info: 2018.199TWNP.26201 Appr. Yr.: 2018 CAN: 199TWNP Object Class: 26201 Funded: \$0.00 FOB: Destination No-cost time extension for CLIN 0008. [***] Change Item 3 to read as follows (amount shown is the obligated amount): CLIN 0007 Smallpox Antiviral Drug for the - Strategic National Stockpile [***] Accounting Info: 2013.1992002.25106 Appr. Yr.: 2013 CAN: 1992002 Object Class: 25106 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$1,120,983.54 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: -\$1,215,399.49 FOB: Destination	2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$269,641.07 Accounting Info: 2018.199TWNP.26201 Appr. Yr.: 2018 CAN: 199TWNP Object Class: 26201 Funded: \$0.00 FOB: Destination No-cost time extension for CLIN 0008. [***] Change Item 3 to read as follows (amount shown is the obligated amount): CLIN 0007 Smallpox Antiviral Drug for the - Strategic National Stockpile [***] Accounting Info: 2013.1992002.25106 Appr. Yr.: 2013 CAN: 1992002 Object Class: 25106 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$1,120,983.54 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00 Accounting Info: 2016.1992016.25103 Appr. Yr.:2016 CAN 1992016 Object Class: 25103 Funded: \$0.00

OPTIONAL FORM 336 (4-86) Sponsored by GSA FAR (48 CFR) 53.110

CONTINUATION	REFERENCE NO. OF DOCUMENT BEING CONTINUED	P	AGE
SHEET	HHSO100201100001C/P00021		OF
		3	3

NAME OF OFFEROR OR CONTRACTOR SIGA TECHNOLOGIES, INC. 1385150

ITEM NO	SUPPLIES/SERVICES	QUANTITY	UNIT	UNIT PRICE	AMOUNT
(A)	(B)	(C)	(D)	(E)	(F)
	No-cost time extension to comply with [***] CLIN 0007.12 Specific PoP extended to 04/22/2023 CLIN 0007.14 Specific PoP extended to 09/21/2023 Change Item 5 to read as follows (amount shown is the obligated amount):				
5	No-cost extension of [***] PoP changed to 06/28/2016 through 12/09/2022				-6,189.84
	Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$6,189.84				
	No-cost time extension for [***]				
	Change Item 6 to read as follows (amount shown is the obligated amount):				
6	No-cost extension of CLIN 00021, [***]				-6,079.60
	Accounting Info: 2016.1992016.25103 Appr. Yr.: 2016 CAN: 1992016 Object Class: 25103 Funded: -\$6,079.60				
	No-cost time extension for [***]				
	Add Item 8 as follows:				
8	ASPR-22-02268- Additional funding to SIGA Technologies to cover the cost for completing tasks remaining under Contract HHSO10020110001C Obligated Amount: \$2,618,293.54				2,618,293.54
	Accounting Info: 2022.1992022.25106 Appr. Yr.: 2022 CAN: 1992022 Object Class: 25106 Funded: \$2,618,293.54				
	PSC: 6505 NAICS: 541714 COR is Dr. Annie Xi Lu, (202) 604-5814, Xi.Lu@hhs.gov				

NSN 7540-01-152-8067

OPTIONAL FORM 336 (4-86) Sponsored by GSA FAR (48 CFR) 53.110

AMENDMENT OF SOLICITATION/MODE CONTRACT	IFIC/	ATION OF	1. 0	CONTRACT ID CODE		PAGE OF PAGES
CONTRACT						1 2
2. AMENDMENT/MODIFICATION NO. P00022		EFFECTIVE DATE		REQUISITION/PURCHASE REQ. NO. 302885	5. 1	PROJECT NO. (If applicable)
		See Block 16C				
6. ISSUED BY	CODE	ASPR- BARDA		ADMINISTERED BY (If CODE er than Item 6)	E	
ASPR-BARDA 200 Independence Ave., S.W. Room 640-G Washington DC 20201						
8. NAME AND ADDRESS OF CONTRACTO county, State and ZIP Code)	R (N	o., street,	(x)	9A. AMENDMENT OF SOLICITATION NO.	•	
SIGA TECHNOLOGIES, INC. 1385150 Attn: Daniel Luckshire SIGA TECHNOLOGIES, INC. 31 E 6 31 E 62ND ST NEW YORK NY 100658014						
				9B. DATED (SEE ITEM 11)		
			Х	10A. MODIFICATION OF CONTRACT/ORD HHSO100201100001C	DER	NO.
CODE 1385150		FACILITY CODE		10B. DATED (<i>SEE ITEM 13</i>) 05/13/2011		
				S TO AMENDMENTS OF SOLICITATION m 14. The hour and date specified for receipt of		ers \square is extended.
methods: (a) By completing Items 8 and 15, copy of the offer submitted; or (c) By separa numbers. FAILURE OF YOUR ACKNOW	and ro te lett LEDC	eturning er or electron GEMENT TO	ic co BE	ur and date specified in the solicitation or as amoropies of the amendment; (b) By acknowledging communication which includes a reference to the RECEIVED AT THE PLACE DESIGNATED IN REJECTION OF YOUR OFFER. If by virtu	g rec soli FOR	eipt of this amendment on each citation and amendment THE RECEIPT OF OFFERS
				ter or electronic communication, provided each		er or electronic communication
12. ACCOUNTING AND APPROPRIATION	DATA	(If required	ecei	ved prior to the opening hour and date specified	1.	
2022.1992022.25100		ICATION O	F C	ONTRACTS/ORDERS. IT MODIFIES THE	E CO	NTRACT/ORDER NO. AS
CHECK ONE A. THI	S CH.			RIBED IN ITEM 14. S ISSUED PURSUANT TO: (Specify authority)) TH	E CHANGES SET FORTH IN
ITEM 1	14 AR	E MADE IN	TH	E CONTRACT ORDER NO. IN ITEM 10A.	•	
X CHAN TO TH	B. THE ABOVE NUMBERED CONTRACT/ORDER IS MODIFIED TO REFLECT THE ADMINISTRATIVE X CHANGES (such as changes in paying office, appropriation data, etc.) SET FORTH IN ITEM 14, PURSUANT TO THE AUTHORITY OF FAR 43.103(b).					
				GREEMENT IS ENTERED INTO PURSUAN	T T	O AUTHORITY OF:
				odification and authority) s document and return _copies to the issuing offi	ice.	
14. DESCRIPTION OF AMENDMENT/MOD feasible.) Tax ID Number: 13-3864870 DUNS Number: 932651516 UEI: VJRNRTSL22K4	IFIC	ATION (Orga	nize	d by UCF section headings, including solicitation		ontract subject matter where
The purpose of this modification is to correct M Mod 21 did not add funds it only de-obligated All other terms and conditions remain unchang Discount Terms: HHS NET 14P Continued Except as provided herein, all terms and con and in full force and effect.	cancel ed.	lling funds.		ystem and add funds. nt referenced in Item 9A or 10A, as heretofor	re ch	anged, remains unchanged
15A. NAME AND TITLE OF SIGNER (Type	or pri	nt)	16 <i>F</i>	A. NAME AND TITLE OF CONTRACTING O	FFI	CER (Type or print)
Dennis E. Hruby, CSO & EVP				NATHAN F. GONZALEZ		
15B. CONTRACTOR/OFFEROR		15C. DATE SIGNED		3. UNITED STATES OF IERICA	160	C. DATE SIGNED
/s/ Dennis E. Hruby (Signature of person authorized to sign)			/s/ J	Jonathan F. Gonzalez gnature of Contracting	29 9	Sep 2022
		1	1 ~		1	

Previous edition unusable

STANDARD FORM 30 (REV. 11/2016)
Prescribed by GSA FAR (48 CFR) 53.243

NAME OF OFFEI	ROR OR CONTRACTOR				
SIGA TECHNOL	OGIES, INC. 1385150				
ITEM NO	SUPPLIES/SERVICES	QUANTITY	UNIT	UNIT PRICE	AMOUNT
(A)	(B)	(C)	(D)	(E)	(F)
	Appr. Yr.: 2022 CAN: 1992022 Object Class: 25106 Period of Performance: 06/28/2016 to 12/31/2023 Add Item 8 as follows:				
8	Add field 6 as follows. ASPR-22-02268 – Additional funding to SIGA Technologies to cover the cost for completing tasks remaining under Contract HHSO10020110001C Obligated Amount: \$2,618,293.54 PSC: 6505 NAICS: 541714 COR is Dr. Annie Xi Lu, (202) 604-5814, Xi.Lu@hhs.gov				2,618,293.54
	00 1 00 1 i, 14.24@.iii.s.gov				

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OPTIONAL FORM 336 (4-

Sponsored by GSA FAR (48 CFR)

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53.110

CONTINUATION REFERENCE NO. OF DOCUMENT BEING CONTINUED

HHSO100201100001C/P00022

SHEET

NSN 7540-01-152-8067

Certification by Chief Executive Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002

I, Phillip L. Gomez, Ph.D., certify that:

- 1. I have reviewed this quarterly report on Form 10-Q of SIGA Technologies, 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.

Date: November 3, 2022

/s/ Phillip L. Gomez, Ph.D.

Phillip L. Gomez, Ph.D. Chief Executive Officer

Certification by Chief Financial Officer Pursuant to Section 302 of the Sarbanes-Oxley Act of 2002

I, Daniel J. Luckshire, certify that:

- 1. I have reviewed this quarterly report on Form 10-Q of SIGA Technologies, 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.

Date: November 3, 2022

/s/ Daniel J. Luckshire

Daniel J. Luckshire Executive Vice President and Chief Financial Officer

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 of SIGA Technologies, Inc. (the "Company") on Form 10-Q for the quarterly period ended September 30, 2022 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Phillip L. Gomez, Ph. D., Chief Executive Officer of the Company, certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that to the best of my knowledge:

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

/s/ Phillip L. Gomez, Ph.D.

Phillip L. Gomez, Ph.D. Chief Executive Officer November 3, 2022

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 of SIGA Technologies, Inc. (the "Company") on Form 10-Q for the quarterly period ended September 30, 2022 as filed with the Securities and Exchange Commission on the date hereof (the "Report"), I, Daniel J. Luckshire, Executive Vice President and Chief Financial Officer of the Company, certify, pursuant to 18 U.S.C. § 1350, as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002, that to the best of my knowledge:

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

/s/ Daniel J. Luckshire

Daniel J. Luckshire Executive Vice President and Chief Financial Officer November 3, 2022