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

FORM C-AR

UNDER THE SECURITIES ACT OF 1933

(Mark one.)

- ☐ Form C: Offering Statement
- ☐ Form C-U: Progress Update
- ☐ Form C/A: Amendment to Offering Statement
  - ☐ Check box if Amendment is material and investors must reconfirm within five business days.
- ☑ Form C-AR: Annual Report
- ☐ Form C-AR/A: Amendment to Annual Report
- ☐ Form C-TR: Termination of Reporting

Name of issuer
GenoBank.io

Legal status of issuer

Form
Corporation

Jurisdiction of Incorporation/Organization
California

Date of organization
May 29, 2018

Physical address of issuer
55 E Third Ave, San Mateo, CA 94401

Website of issuer
https://GenoBank.io

Current number of employees
2

	Most recent fiscal year-end	Prior fiscal year-end

Total Assets	$6,478.00	$36.00
Cash & Cash Equivalents	$1,027.00	$36.00
Accounts Receivable	-$71.00	-$71.00
Short-term Debt	$0.00	$0.00
Long-term Debt	$0.00	$0.00
Revenues/Sales	$0.00	$0.00
Cost of Goods Sold	$0.00	$0.00
Taxes Paid	$0.00	$0.00
Net Income	-$158,572.00	-$14.00

04-30-2021

FORM C-AR

GenoBank.io

This Form C-AR (including the cover page and all exhibits attached hereto, the "Form C-AR) is being furnished by GenoBank.io, a California Corporation (the "Company," as well as references to "we," "us," or "our") for the sole purpose of providing certain information about the Company as required by the Securities and Exchange Commission ("SEC").

No federal or state securities commission or regulatory authority has passed upon the accuracy or adequacy of this document. The U.S. Securities and Exchange Commission does not pass upon the accuracy or completeness of any disclosure document or literature. The Company is filing this Form C-AR pursuant to Regulation CF (§ 227.100 et seq.) which requires that it must file a report with the Commission annually and post the report on its website at https://GenoBank.io no later than 120 days after the end of each fiscal year covered by the report. The Company may terminate its reporting obligations in the future in accordance with Rule 202(b) of Regulation CF (§ 227.202(b)) by 1) being required to file reports under Section 13(a) or Section 15(d) of the Exchange Act of 1934, as amended, 2) filing at least one annual report pursuant to Regulation CF and having fewer than 300 holders of record, 3) filing annual reports for three years pursuant to Regulation CF and having assets equal to or less than $10,000,000, 4) the repurchase of all the Securities sold pursuant to Regulation CF by the Company or another party, or 5) the liquidation or dissolution of the Company.

The date of this Form C-AR is 04-30-2021.

THIS FORM C-AR DOES NOT CONSTITUTE AN OFFER TO PURCHASE OR SELL SECURITIES.

Forward Looking Statement Disclosure

This Form C-AR and any documents incorporated by reference herein or therein contain forward-looking statements and are subject to risks and uncertainties. All statements other than statements of historical fact or relating to present facts or current conditions included in this Form C-AR are forward-looking statements. Forward-looking statements give the Company's current reasonable expectations and projections relating to its financial condition, results of operations, plans, objectives, future performance and business. You can identify forward-looking statements by the fact that they do not relate strictly to historical or current facts. These statements may include words such as "anticipate," "estimate," "expect," "project," "plan," "intend," "believe," "may," "should," "can have," "likely" and other words and terms of similar meaning in connection with any discussion of the timing or nature of future operating or financial performance or other events.

The forward-looking statements contained in this Form C-AR and any documents incorporated by reference herein or therein are based on reasonable assumptions the Company has made in light of its industry experience, perceptions of historical trends, current conditions, expected future developments and other factors it believes are appropriate under the circumstances. As you read and consider this Form C-AR, you should understand that these statements are not guarantees of performance or results. They involve risks, uncertainties (many of which are beyond the Company's control) and assumptions. Although the Company believes that these forward-looking statements are based on reasonable assumptions, you should be aware that many factors could affect its actual operating and financial performance and cause its performance to differ materially from the performance anticipated in the forward-looking statements. Should one or more of these risks or uncertainties materialize, or should any of these assumptions prove incorrect or change, the Company's actual operating and financial performance may vary in material respects from the performance projected in these forward-looking statements.

Any forward-looking statement made by the Company in this Form C-AR or any documents incorporated by reference herein or therein speaks only as of the date of this Form C-AR. Factors or events that could cause our actual operating and financial performance to differ may emerge from time to time, and it is not possible for the Company to predict all of them. The Company undertakes no obligation to update any forward-looking statement, whether as a result of new information, future developments or otherwise, except as may be required by law.

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About this Form C-AR

You should rely only on the information contained in this Form C-AR. We have not authorized anyone to provide you with information different from that contained in this Form C-AR. You should assume that the information contained in this Form C-AR is accurate only as of the date of this Form C-AR, regardless of the time of delivery of this Form C-AR. Our business, financial condition, results of operations, and prospects may have changed since that date.
Statements contained herein as to the content of any agreements or other document are summaries and, therefore, are necessarily selective and incomplete and are qualified in their entirety by the actual agreements or other documents.

SUMMARY

The following summary is qualified in its entirety by more detailed information that may appear elsewhere in this Form C-AR and the Exhibits hereto.

GenoBank.io (the "Company") is a California Corporation, formed on May 29, 2018.

The Company is located at 55 E Third Ave, San Mateo, CA 94401.

The Company's website is https://GenoBank.io.

The information available on or through our website is not a part of this Form C-AR.

The Business

Genobank allows users to track and establish ownership of their genetic information as private property through the use of blockchain technology. We sell a white-label (patent pending) anonymous DNA/RNA extraction kit that is represented by a non-fungible token based in the ethereum standard ERC721 to allow users/patients establish ownership and control over their biosamples and residual biological datasets.

RISK FACTORS

Risks Related to the Company's Business and Industry

The development and commercialization of our [products/services] is highly competitive.
We face competition with respect to any products that we may seek to develop or commercialize in the future. Our competitors include major companies worldwide. Many of our competitors have significantly greater financial, technical and human resources than we have and superior expertise in research and development and marketing approved [products/services] and thus may be better equipped than us to develop and commercialize [products/services]. These competitors also compete with us in recruiting and retaining qualified personnel and acquiring technologies. Smaller or early stage companies may also prove to be significant competitors, particularly through collaborative arrangements with large and established companies. Accordingly, our competitors may commercialize products more rapidly or effectively than we are able to, which would adversely affect our competitive position, the likelihood that our [products/services] will achieve initial market acceptance and our ability to generate meaningful additional revenues from our products.

We depend on third-party service providers and outsource providers for a variety of services and we outsource a number of our non-core functions and operations.
In certain instances, we rely on single or limited service providers and outsourcing vendors [around the world] because the relationship is advantageous due to quality, price, or lack of alternative sources. If production or service was interrupted and we were not able to find alternate third-party providers, we could experience disruptions in manufacturing and operations including product shortages, higher freight costs and re-engineering costs. If outsourcing services are interrupted or not performed or the performance is poor, this could impact our ability to process, record and report transactions with our customers and other constituents. Such interruptions in the provision of supplies and/or services could result in our inability to meet customer demand, damage our reputation and customer relationships and adversely affect our business.

We depend on third party providers, suppliers and licensors to supply some of the hardware, software and operational support necessary to provide some of our services.
We obtain these materials from a limited number of vendors, some of which do not have a long operating history, or which may not be able to continue to supply the equipment and services we desire. Some of our hardware, software and operational support vendors represent our sole source of supply or have, either through contract or as a result of intellectual property rights, a position of some exclusivity. If demand exceeds these vendors' capacity or if these vendors

experience operating or financial difficulties or are otherwise unable to provide the equipment or services we need in a timely manner, at our specifications and at reasonable prices, our ability to provide some services might be materially adversely affected, or the need to procure or develop alternative sources of the affected materials or services might delay our ability to serve our customers. These events could materially and adversely affect our ability to retain and attract customers, and have a material negative impact on our operations, business, financial results and financial condition.

Quality management plays an essential role in determining and meeting customer requirements, preventing defects, improving the Company's products and services and maintaining the integrity of the data that supports the safety and efficacy of our products.

Our future success depends on our ability to maintain and continuously improve our quality management program. An inability to address a quality or safety issue in an effective and timely manner may also cause negative publicity, a loss of customer confidence in us or our current or future products, which may result in the loss of sales and difficulty in successfully launching new products. In addition, a successful claim brought against us in excess of available insurance or not covered by indemnification agreements, or any claim that results in significant adverse publicity against us, could have an adverse effect on our business and our reputation.

Manufacturing or design defects, unanticipated use of our products, or inadequate disclosure of risks relating to the use of the products can lead to injury or other adverse events.

These events could lead to recalls or safety alerts relating to our products (either voluntary or required by governmental authorities) and could result, in certain cases, in the removal of a product from the market. Any recall could result in significant costs as well as negative publicity that could reduce demand for our products. Personal injuries relating to the use of our products can also result in product liability claims being brought against us. In some circumstances, such adverse events could also cause delays in new product approvals. Similarly, negligence in performing our services can lead to injury or other adverse events.

We [may] [plan to] implement new lines of business or offer new products and services within existing lines of business.

There are substantial risks and uncertainties associated with these efforts, particularly in instances where the markets are not fully developed. In developing and marketing new lines of business and/or new products and services, we may invest significant time and resources. Initial timetables for the introduction and development of new lines of business and/or new products or services may not be achieved and price and profitability targets may not prove feasible. We may not be successful in introducing new products and services in response to industry trends or developments in technology, or those new products may not achieve market acceptance. As a result, we could lose business, be forced to price products and services on less advantageous terms to retain or attract clients, or be subject to cost increases. As a result, our business, financial condition or results of operations may be adversely affected.

In general, demand for our products and services is highly correlated with general economic conditions.

A substantial portion of our revenue is derived from discretionary spending by individuals, which typically falls during times of economic instability. Declines in economic conditions in the U.S. or in other countries in which we operate may adversely impact our consolidated financial results. Because such declines in demand are difficult to predict, we or the industry may have increased excess capacity as a result. An increase in excess capacity may result in declines in prices for our products and services.

The use of individually identifiable data by our business, our business associates and third parties is regulated at the state, federal and international levels.

Costs associated with information security – such as investment in technology, the costs of compliance with consumer protection laws and costs resulting from consumer fraud – could cause our business and results of operations to suffer materially. Additionally, the success of our online operations depends upon the secure transmission of confidential information over public networks, including the use of cashless payments. The intentional or negligent actions of employees, business associates or third parties may undermine our security measures. As a result, unauthorized parties may obtain access to our data systems and misappropriate confidential data. There can be no assurance that advances in computer capabilities, new discoveries in the field of cryptography or other developments will prevent the compromise of our customer transaction processing capabilities and personal data. If any such compromise of our security or the security of information residing with our business associates or third parties were to occur, it could have a material adverse effect on our reputation, operating results and financial condition. Any compromise of our data security may materially increase the costs we incur to protect against such breaches and could subject us to additional legal risk.

Security breaches and other disruptions could compromise our information and expose us to liability, which would cause our business and reputation to suffer.

We [collect and] store sensitive data, including intellectual property, our proprietary business information and that of our customers, [suppliers and business partners,] and personally identifiable information of our [customers and] employees, in our data centers and on our networks. The secure [processing,] maintenance [and transmission] of this information is critical to our operations [and business strategy]. Despite our security measures, our information technology and infrastructure may be vulnerable to attacks by hackers or breached due to employee error, malfeasance or other disruptions. Any such breach could compromise our networks and the information stored there could be accessed, publicly disclosed, lost or stolen. Any such access, disclosure or other loss of information could result in legal claims or proceedings, [liability under laws that protect the privacy of personal information,] [and regulatory penalties,] [disrupt our operations [and the services we provide to customers],] [and] damage our reputation, [and cause a loss of confidence in our products and services], which could adversely affect our [business/operating margins, revenues and competitive position].

[The secure [processing,] maintenance [and transmission] of this information is critical to our operations [and business strategy], and we devote significant resources to protecting our

information [by [DESCRIBE ADDITIONAL SECURITY MEASURES.]] The expenses associated with [protecting our information/ these steps] could reduce our operating margins.]

An intentional or unintentional disruption, failure, misappropriation or corruption of our network and information systems could severely affect our business.

Such an event might be caused by computer hacking, computer viruses, worms and other destructive or disruptive software, "cyber attacks" and other malicious activity, as well as natural disasters, power outages, terrorist attacks and similar events. Such events could have an adverse impact on us and our customers, including degradation of service, service disruption, excessive call volume to call centers and damage to our plant, equipment and data. In addition, our future results could be adversely affected due to the theft, destruction, loss, misappropriation or release of confidential customer data or intellectual property. Operational or business delays may result from the disruption of network or information systems and the subsequent remediation activities. Moreover, these events may create negative publicity resulting in reputation or brand damage with customers.

The Company's success depends on the experience and skill of the board of directors, its executive officers and key employees.

In particular, the Company is dependent on Daniel Uribe and Candy Bellau who are Chief Executive Officer, President, Secretary and CFO, Treasurer of the Company. The Company has or intends to enter into employment agreements with Daniel Uribe and Candy Bellau although there can be no assurance that it will do so or that they will continue to be employed by the Company for a particular period of time. The loss of Daniel Uribe and Candy Bellau or any member of the board of directors or executive officer could harm the Company's business, financial condition, cash flow and results of operations.

We rely on various intellectual property rights, including patents and trademarks in order to operate our business.

Such intellectual property rights, however, may not be sufficiently broad or otherwise may not provide us a significant competitive advantage. In addition, the steps that we have taken to maintain and protect our intellectual property may not prevent it from being challenged, invalidated, circumvented or designed-around, particularly in countries where intellectual property rights are not highly developed or protected. In some circumstances, enforcement may not be available to us because an infringer has a dominant intellectual property position or for other business reasons, or countries may require compulsory licensing of our intellectual property. Our failure to obtain or maintain intellectual property rights that convey competitive advantage, adequately protect our intellectual property or detect or prevent circumvention or unauthorized use of such property, could adversely impact our competitive position and results of operations. We also rely on nondisclosure and noncompetition agreements with employees, consultants and other parties to protect, in part, trade secrets and other proprietary rights. There can be no assurance that these agreements will adequately protect our trade secrets and other proprietary rights and will not be breached, that we will have adequate remedies for any breach, that others will not independently develop substantially equivalent proprietary information or that third parties will not otherwise gain access to our trade secrets or other proprietary rights.

As we expand our business, protecting our intellectual property will become increasingly important. The protective steps we have taken may be inadequate to deter our competitors from using our proprietary information. In order to protect or enforce our patent rights, we may be required to initiate litigation against third parties, such as infringement lawsuits. Also, these third parties may assert claims against us with or without provocation. These lawsuits could be expensive, take significant time and could divert management's attention from other business concerns. The law relating to the scope and validity of claims in the technology field in which we operate is still evolving and, consequently, intellectual property positions in our industry are generally uncertain. We cannot assure you that we will prevail in any of these potential suits or that the damages or other remedies awarded, if any, would be commercially valuable.

From time to time, third parties may claim that one or more of our products or services infringe their intellectual property rights.

Any dispute or litigation regarding patents or other intellectual property could be costly and time-consuming due to [the complexity of our technology and] the uncertainty of intellectual property litigation and could divert our management and key personnel from our business operations. A claim of intellectual property infringement could force us to enter into a costly or restrictive license agreement, which might not be available under acceptable terms or at all, could require us to redesign our products, which would be costly and time-consuming, and/or could subject us to an injunction against development and sale of certain of our products or services. We may have to pay substantial damages, including damages for past infringement if it is ultimately determined that our products infringe on a third party's proprietary rights. Even if these claims are without merit, defending a lawsuit takes significant time, may be expensive and may divert management's attention from other business concerns. Any public announcements related to litigation or interference proceedings initiated or threatened against us could cause our business to be harmed. Our intellectual property portfolio may not be useful in asserting a counterclaim, or negotiating a license, in response to a claim of intellectual property infringement. In certain of our businesses we rely on third party intellectual property licenses and we cannot ensure that these licenses will be available to us in the future on favorable terms or at all.

The amount of capital the Company is attempting to raise in this Offering is not enough to sustain the Company's current business plan.

In order to achieve the Company's near and long-term goals, the Company will need to procure funds in addition to the amount raised in the Offering. There is no guarantee the Company will be able to raise such funds on acceptable terms or at all. If we are not able to raise sufficient capital in the future, we will not be able to execute our business plan, our continued operations will be in jeopardy and we may be forced to cease operations and sell or otherwise transfer all or substantially all of our remaining assets, which could cause an Investor to lose all or a portion of his or her investment.

Although dependent on certain key personnel, the Company does not have any key man life insurance policies on any such people.

The Company is dependent on Daniel Uribe and Candy Bellau in order to conduct its operations and execute its business plan, however, the Company has not purchased any insurance policies with respect to those individuals in the event of their death or disability. Therefore, if any of Daniel Uribe and Candy Bellau die or become disabled, the Company will not receive any compensation to assist with such person's absence. The loss of such person could negatively affect the Company and its operations.

We are subject to income taxes as well as non-income based taxes, such as payroll, sales, use, value-added, net worth, property and goods and services taxes, in both the U.S. [and various foreign jurisdictions].

Significant judgment is required in determining our provision for income taxes and other tax liabilities. In the ordinary course of our business, there are many transactions and calculations where the ultimate tax determination is uncertain. Although we believe that our tax estimates are reasonable: (i) there is no assurance that the final determination of tax audits or tax disputes will not be different from what is reflected in our income tax provisions, expense amounts for non-income based taxes and accruals and (ii) any material differences could have an adverse effect on our financial position and results of operations in the period or periods for which determination is made.

We are not subject to Sarbanes-Oxley regulations and lack the financial controls and safeguards required of public companies.

We do not have the internal infrastructure necessary, and are not required, to complete an attestation about our financial controls that would be required under Section 404 of the Sarbanes-Oxley Act of 2002. There can be no assurance that there are no significant deficiencies or material weaknesses in the quality of our financial controls. We expect to incur additional expenses and diversion of management's time if and when it becomes necessary to perform the system and process evaluation, testing and remediation required in order to comply with the management certification and auditor attestation requirements.

The Company has indicated that it has engaged in certain transactions with related persons.

Please see the section of this Memorandum entitled "Transactions with Related Persons and Conflicts of Interest" for further details.

Changes in employment laws or regulation could harm our performance.

Various federal and state labor laws govern our relationship with our employees and affect operating costs. These laws include minimum wage requirements, overtime pay, healthcare reform and the implementation of the Patient Protection and Affordable Care Act, unemployment tax rates, workers' compensation rates, citizenship requirements, union membership and sales taxes. A number of factors could adversely affect our operating results, including additional government-imposed increases in minimum wages, overtime pay, paid leaves of absence and mandated health benefits, mandated training for employees, increased tax reporting and tax payment [requirements for employees who receive tips, a reduction in the number of states that allow tips to be credited toward minimum wage requirements,] changing regulations from the

National Labor Relations Board and increased employee litigation including claims relating to the Fair Labor Standards Act.

The Company's business operations may be materially adversely affected by a pandemic such as the Coronavirus (COVID-19) outbreak.

In December 2019, a novel strain of coronavirus was reported to have surfaced in Wuhan, China, which spread throughout other parts of the world, including the United States. On January 30, 2020, the World Health Organization declared the outbreak of the coronavirus disease (COVID-19) a "Public Health Emergency of International Concern." On January 31, 2020, U.S. Health and Human Services Secretary Alex M. Azar II declared a public health emergency for the United States to aid the U.S. healthcare community in responding to COVID-19, and on March 11, 2020 the World Health Organization characterized the outbreak as a "pandemic." COVID-19 resulted in a widespread health crisis that adversely affected the economies and financial markets worldwide. The Company's business could be materially and adversely affected. The extent to which COVID-19 impacts the Company's business will depend on future developments, which are highly uncertain and cannot be predicted, including new information which may emerge concerning the severity of COVID-19 and the actions to contain COVID-19 or treat its impact, among others. If the disruptions posed by COVID-19 or other matters of global concern continue for an extended period of time, the Company's operations may be materially adversely affected.

We face risks related to health epidemics and other outbreaks, which could significantly disrupt the Company's operations and could have a material adverse impact on us.

The outbreak of pandemics and epidemics could materially and adversely affect the Company's business, financial condition, and results of operations. If a pandemic occurs in areas in which we have material operations or sales, the Company's business activities originating from affected areas, including sales, materials, and supply chain related activities, could be adversely affected. Disruptive activities could include the temporary closure of facilities used in the Company's supply chain processes, restrictions on the export or shipment of products necessary to run the Company's business, business closures in impacted areas, and restrictions on the Company's employees' or consultants' ability to travel and to meet with customers, vendors or other business relationships. The extent to which a pandemic or other health outbreak impacts the Company's results will depend on future developments, which are highly uncertain and cannot be predicted, including new information which may emerge concerning the severity of a virus and the actions to contain it or treat its impact, among others. Pandemics can also result in social, economic, and labor instability which may adversely impact the Company's business.

If the Company's employees or employees of any of the Company's vendors, suppliers or customers become ill or are quarantined and in either or both events are therefore unable to work, the Company's operations could be subject to disruption. The extent to which a pandemic affects the Company's results will depend on future developments that are highly uncertain and cannot be predicted.

We face risks relating to public health conditions such as the COVID-19 pandemic, which could adversely affect the Company's customers, business, and results of operations.

Our business and prospects could be materially adversely affected by the COVID-19 pandemic or recurrences of that or any other such disease in the future. Material adverse effects from COVID-19 and similar occurrences could result in numerous known and currently unknown ways including from quarantines and lockdowns which impair the Company's business including: [marketing and sales efforts, supply chain, etc.]. [Describe how a quarantine has or may in the future negatively affect your employees and their ability to perform their duties]. [Describe how a quarantine has or may in the future negatively affect your suppliers, their employees, and overall ability to fulfill orders]. If the Company purchases materials from suppliers in affected areas, the Company may not be able to procure such products in a timely manner. The effects of a pandemic can place travel restrictions on key personnel which could have a material impact on the business. In addition, a significant outbreak of contagious diseases in the human population could result in a widespread health crisis that could adversely affect the economies and financial markets of many countries, resulting in an economic downturn that could reduce the demand for the Company's products and impair the Company's business prospects including as a result of being unable to raise additional capital on acceptable terms to us, if at all.

The United States tax rules applicable to an investment in the Securities and the underlying Bitcoins are uncertain and the tax consequences to an investor of an investment in the Securities could differ from the investor's expectations.

The tax rules applicable to the Securities and the underlying Bitcoins held by the Company are complex, and no statutory, judicial, or administrative authority directly addresses the characterization of an investment in Bitcoins. The tax consequences to an Investor of the Securities could differ from the Investor's expectations. Investor's should consult their own tax advisors.

Regulatory changes or actions may alter the nature of an investment in the Securities or restrict the use of Bitcoins or the operation of the Bitcoin network in a manner that adversely affects an investment in the Securities.

Until recently, little or no regulatory attention has been directed toward Bitcoins and the Bitcoin network by U.S. federal and state governments, foreign governments and self-regulatory agencies. As Bitcoins have grown in popularity and in market size, the U.S. Congress and certain U.S. agencies (e.g., FinCEN and the Federal Bureau of Investigation) have begun to examine the operations of the Bitcoin network, Bitcoin users and the Bitcoin exchange market. Local state regulators such as the California Department of Financial Institutions and the New York State Department of Financial Services have also initiated examinations of Bitcoin. Additionally, a U.S. federal magistrate judge in the U.S. District Court for the Eastern District of Texas has ruled that "Bitcoin is a currency or form of money," although there is no indication yet whether other courts or federal or state regulators will follow the federal magistrate's opinion. There is a possibility of future regulatory change altering, perhaps to a material extent, the nature of an investment in the Securities or the ability of the Company to continue to operate. Currently, neither the SEC nor the CFTC has formally asserted regulatory authority over the Bitcoin network or Bitcoin trading and ownership. To the extent that Bitcoins are determined to be a

security, commodity future or other regulated asset, or to the extent that a U.S. or foreign government or quasi-governmental agency exerts regulatory authority over the Bitcoin network or Bitcoin trading and ownership, trading or ownership in Bitcoins or the Securities may be adversely affected.

To the extent that future regulatory actions or policies limit the ability to exchange Bitcoins or utilize them for payments, the demand for Bitcoins will decrease.

New regulations may make it more difficult to acquire and/or use Bitcoins. Furthermore, regulatory actions may limit the ability of end-users to convert Bitcoins into fiat currency (e.g., U.S. Dollars) or use Bitcoins to pay for goods and services. Such regulatory actions or policies would negatively affect our business and decrease the value of the Securities.

Bitcoin currently faces an uncertain regulatory landscape in not only the United States but also in many foreign jurisdictions such as the European Union.

While the German Ministry of Finance has declared Bitcoin to be "Rechnungseinheiten" (a form of private money that is recognized as a unit of account, but not recognized in the same manner as fiat currency), most regulatory bodies have not yet issued official statements regarding intention to regulate or determinations on regulation of Bitcoin, Bitcoin users and the Bitcoin network. Conversely, regulatory bodies in some countries such as Canada and India have declined to exercise regulatory authority when afforded the opportunity. Various foreign jurisdictions may, in the near future, adopt laws, regulations or directives that affect the Bitcoin network and its users, particularly Bitcoin exchanges and service providers that fall within such jurisdictions' regulatory scope. Such laws, regulations or directives may conflict with those of the United States and may negatively impact the acceptance of Bitcoins by users, merchants and service providers outside of the United States and may therefore impede the growth of the Bitcoin economy. We are not able to predict the effect of any future regulatory change on the Company or Bitcoins, but such change could be substantial and adverse to the Company or the value of the Shares.

It may be illegal now, or in the future, to acquire, own, hold, sell or use Bitcoins in one or more countries.

Although currently Bitcoins are not regulated or are lightly regulated in most countries, including the United States, one or more countries may take regulatory actions in the future that severely restricts the right to acquire, own, hold, sell or use Bitcoins or to exchange Bitcoins for fiat currency. Such an action may also result in the restriction of ownership, holding or trading in the Securities. Such a restriction could result in the termination and liquidation of the Company at a time that is disadvantageous to Investor, or may adversely affect an investment in the Company.

The Company may be deemed a "money transmitter."

To the extent that the activities of the Company cause it to be deemed a "money transmitter" under the regulations promulgated by FinCEN under the authority of the U.S. Bank Secrecy Act, the Company may be required to comply with FinCEN regulations, including those that would mandate the Company to implement anti-money laundering programs, make certain reports to

FinCEN and maintain certain records. Such additional regulatory obligations may cause the Company to incur extraordinary expenses, possibly affecting an investment in the Securities in a material and adverse manner. Additionally, certain states including California, Idaho and New York require Bitcoin businesses to register on the state level as money transmitters.

Current and future legislation, CFTC and SEC rulemaking and other regulatory developments may impact the manner in which Bitcoins are treated for classification and clearing purposes.

In particular, Bitcoins may not be excluded from the definition of "commodity future" or "security" by such future CFTC and SEC rulemaking, respectively. As of the date of this Memorandum, the Company is not aware of any rules that have been proposed to regulate Bitcoins as commodity futures or securities. The Company cannot be certain as to how future regulatory developments will impact the treatment of Bitcoins under the law. Such additional registrations may result in extraordinary expenses of the Company thereby materially and adversely impacting the Securities.

We face heavy government regulation, and FDA regulatory approval of our products is uncertain.

The research, testing, manufacturing and marketing of drug products such as those that we are developing are subject to extensive regulation by federal, state and local government authorities, including the FDA. To obtain regulatory approval of a product, we must demonstrate to the satisfaction of the applicable regulatory agency that, among other things, the product is safe and effective for its intended use. In addition, we must show that the manufacturing facilities used to produce the products are in compliance with current Good Manufacturing Practices regulations (cGMP). The process of obtaining FDA and other required regulatory approvals and clearances will require us to expend substantial time and capital. Despite the time and expense expended, regulatory approval is never guaranteed. The number of preclinical and clinical trials that will be required for FDA approval varies depending on the drug candidate, the disease or condition that the drug candidate is in development for, and the requirements applicable to that particular drug candidate. The FDA can delay, limit or deny approval of a drug candidate for many reasons, including that:

* a drug candidate may not be shown to be safe or effective;

* the FDA may not approve our manufacturing process

* the FDA may interpret data from preclinical and clinical trials in different ways than we do; and

* the FDA may not meet, or may extend, the Prescription Drug User Fee Act date with respect to a particular New Drug Application ("NDA").

For example, if certain of our methods for analyzing our trial data are not accepted by the FDA, we may fail to obtain regulatory approval for our product candidates. Moreover, if and when our products do obtain marketing approval, the marketing, distribution and manufacture of such

products would remain subject to extensive ongoing regulatory requirements. Failure to comply with applicable regulatory requirements could result in warning letters, fines, civil penalties, injunctions, recall or seizure of products, total or partial suspension of production, refusal of the government to grant future approvals, withdrawal of approvals, or criminal prosecution.

Any delay or failure by us to obtain regulatory approvals for our product candidates could diminish competitive advantages that we may attain and would adversely affect the marketing of our products. [To date, we have not received regulatory approval to market any of our product candidates in any jurisdiction. Following regulatory approval of any of our drug candidates, we will be subject to ongoing regulatory obligations and restrictions, which may result in significant expense and limit our ability to commercialize our potential products.]

With regard to our drug candidates[, if any,] approved by the FDA or by another regulatory authority, we are held to extensive regulatory requirements over product manufacturing, labeling, packaging, adverse event reporting, storage, advertising, promotion and record keeping. Regulatory approvals may also be subject to significant limitations on the indicated uses or marketing of the drug candidates. Potentially costly follow-up or post-marketing clinical studies may be required as a condition of approval to further substantiate safety or efficacy, or to investigate specific issues of interest to the regulatory authority. Previously unknown problems with the drug candidate, including adverse events of unanticipated severity or frequency, may result in restrictions on the marketing of the drug, and could include withdrawal of the drug from the market.

In addition, the law or regulatory policies governing pharmaceuticals may change. New statutory requirements may be enacted or additional regulations may be enacted that could prevent or delay regulatory approval of our drug candidates. We cannot predict the likelihood, nature or extent of adverse government regulation that may arise from future legislation or administrative action, either in the United States or elsewhere. If we are not able to maintain regulatory compliance, we might not be permitted to market our drugs and our business could suffer.

We [are] [may in the future be] subject to various U.S. federal and state laws pertaining to health care fraud and abuse, including anti-kickback, self-referral, false claims and fraud laws, and any violations by us of such laws could result in fines or other penalties.

[We are] [If one or more of our product candidates is approved, we will likely be] subject to the various U.S. federal and state laws intended to prevent health care fraud and abuse. The federal anti-kickback statute prohibits the offer, receipt, or payment of remuneration in exchange for or to induce the referral of patients or the use of products or services that would be paid for in whole or part by Medicare, Medicaid or other federal health care programs. Remuneration has been broadly defined to include anything of value, including cash, improper discounts, and free or reduced price items and services. Many states have similar laws that apply to their state health care programs as well as private payors. Violations of the anti-kickback laws can result in exclusion from federal health care programs and substantial civil and criminal penalties.

The False Claims Act (FCA) imposes liability on persons who, among other things, present or cause to be presented false or fraudulent claims for payment by a federal health care program. The FCA has been used to prosecute persons submitting claims for payment that are inaccurate

or fraudulent, that are for services not provided as claimed, or for services that are not medically necessary. The FCA includes a whistleblower provision that allows individuals to bring actions on behalf of the federal government and share a portion of the recovery of successful claims. If our marketing or other arrangements were determined to violate the FCA or anti-kickback or related laws, then our revenue could be adversely affected, which would likely harm our business, financial condition, and results of operations.

State and federal authorities have aggressively targeted medical technology companies for alleged violations of these anti-fraud statutes, based on improper research or consulting contracts with doctors, certain marketing arrangements that rely on volume-based pricing, off-label marketing schemes, and other improper promotional practices. Companies targeted in such prosecutions have paid substantial fines in the hundreds of millions of dollars or more, have been forced to implement extensive corrective action plans or Corporate Integrity Agreements, and have often become subject to consent decrees severely restricting the manner in which they conduct their business. If we become the target of such an investigation or prosecution based on our contractual relationships with providers or institutions, or our marketing and promotional practices, we could face similar sanctions, which would materially harm our business.

If we are found to have violated laws protecting the privacy or security of patient health information, we could be subject to civil or criminal penalties, which could increase our liabilities and harm our reputation or our business.

There are a number of U.S. federal and state laws and foreign laws protecting the privacy and security of individually identifiable health information, or "protected health information" including patient records, and restricting the use and disclosure of that protected health information that we are subject to. In the United States, the U.S. Department of Health and Human Services promulgated health information privacy and security rules under the Health Insurance Portability and Accountability Act of 1996 (HIPAA) and then significantly strengthened and broadened the applicability of HIPAA under the Health Information Technology for Economic and Clinical Health Act (HITECH, together HIPAA). HIPAA applies to health care providers engaging in certain standard transactions electronically; health plans and health care clearing houses. These entities are referred to as "covered entities." Certain HIPAA provisions also apply to "business associates" of covered entities, or third party providers of services to covered entities that involve the use or disclosure of protected health information. HIPAA's privacy rules protect medical records and protected health information in all forms by limiting its use and disclosure, giving individuals the right to access, amend and seek accounting of their own health information and limiting, in some circumstances, the use and disclosure of protected health information to the minimum amount reasonably necessary to accomplish the intended purpose of the use or disclosure. HIPAA's security standards require both covered entities and business associates to implement administrative, physical and technical security measures to maintain the security of protected health information in electronic form. Covered entities and business associates must conduct initial and ongoing risk assessments to ensure the ongoing effectiveness of security measures and maintain a written information security plan. We are a [covered entity] [business associate] and as such, we must comply with HIPAA and ensure that all aspects of our operations comply with relevant HIPAA standards. We are subject to random audit by federal authorities, and enforcement by both state and federal regulators. We are also subject to investigation in response to complaints. If we are found to be in violation of the

HIPAA requirements, we could be subject to civil or criminal penalties as well as fines, which could increase our liabilities and harm our reputation or our business.

Beyond HIPAA, most states have adopted data security laws protecting the personal data of state residents. Personal data subject to protection typically includes name coupled with social security number, state-issued identification number, or financial account number. Most states require specific, technical security measures for the protection of all personal data, including employee data, and impose their own breach notification requirements in the event of a loss of personal data. State data security laws generally overlap and apply simultaneously with HIPAA. [Non-U.S. privacy protection requirements such as the European Union's Data Protection Directive governing the processing of personal data, may be stricter than the U.S. law and violation would impose similar or more severe penalties. These laws could create liability for us or increase our cost of doing business, and any failure to comply could result in harm to our reputation, and potentially fines and penalties.

Healthcare legislative reform measures may have a material adverse effect on our business and results of operations.

In the United States, there have been and continue to be a number of legislative initiatives to contain healthcare costs. For example, in March 2010, the Patient Protection and Affordable Care Act, as amended by the Health Care and Education Reconciliation Act, or the Affordable Care Act, was passed, which substantially changed the way health care is financed by both governmental and private insurers, and significantly impacted the U.S. pharmaceutical industry. The Affordable Care Act, among other things, subjected biologic products to potential competition by lower-cost biosimilars, addressed a new methodology by which rebates owed by manufacturers under the Medicaid Drug Rebate Program are calculated for drugs that are inhaled, infused, instilled, implanted or injected, increased the minimum Medicaid rebates owed by manufacturers under the Medicaid Drug Rebate Program and extended the rebate program to individuals enrolled in Medicaid managed care organizations, established annual fees and taxes on manufacturers of certain branded prescription drugs, and a new Medicare Part D coverage gap discount program, in which manufacturers must agree to offer 50% point-of-sale discounts off negotiated prices of applicable brand drugs to eligible beneficiaries during their coverage gap period, as a condition for the manufacturer's outpatient drugs to be covered under Medicare Part D.

We expect that additional state and federal healthcare reform measures will be adopted in the future, any of which could limit the amounts that federal and state governments will pay for healthcare products and services, which could result in reduced demand for our products or additional pricing pressures, which would negatively affect our business.

New product development involves a lengthy, expensive and complex process.

We may be unable to develop or commercialize any of the product candidates we are currently researching. Moreover, even if we develop such candidates, they may be subject to significant regulatory review, approval and other government regulations. We are currently conducting research and development on [name of treatment] for [disease type], such as [example diseases]. There can be no assurance that our technologies will be capable of reliably addressing resistant

infections or that we can develop and commercialize our products at all. New product development involves a lengthy, expensive and complex process and we currently have no fully validated diagnostic candidates. In addition, before we can commercialize any new product candidates, we will need to:

* conduct substantial research and development;

* conduct validation studies;

* expend significant funds;

* develop and scale-up our laboratory processes; and

* obtain regulatory approval and acceptance of our product candidates.

This process involves a high degree of risk and takes several years. Our product development efforts may fail for many reasons, including:

* failure of the product at the research or development stage; and

* lack of clinical validation data to support the effectiveness of the product.

Few research and development projects result in commercial products, and perceived viability in early clinical trials often is not replicated in later studies. At any point, we may abandon development of a product candidate or we may be required to expend considerable resources repeating clinical trials, which would adversely impact the timing for generating potential revenues from those product candidates. In addition, as we develop product candidates, we will have to make significant investments in product development, marketing and sales resources.

We may not be able to conduct clinical trials necessary to commercialize and sell our proposed products and formulations.

In order to conduct clinical trials that are necessary to obtain approval by the FDA to market a formulation or product, it is necessary to receive clearance from the FDA to conduct such clinical trials. The FDA can halt clinical trials at any time for safety reasons or because we or our clinical investigators do not follow the FDA's requirements for conducting clinical trials. If we are unable to receive clearance to conduct clinical trials or the trials are halted by the FDA, we would not be able to achieve any revenue from such product as it is illegal to sell any drug or medical device for human consumption without FDA approval. [Moreover, it is our stated intention to attempt to avail ourselves of the FDA's Fast Track approval procedure, which we believe is less costly and time consuming.] If this approval pathway is not available to us with respect to a particular formulation or product, or at all, the time and cost associated with developing and commercializing such formulations or products may be prohibitive and our business strategy would be materially and adversely affected.

Our long-term viability and growth will depend upon successful clinical trials.

Product development is very expensive and involves a high degree of risk. Only a small number of research and development programs result in the commercialization of a product. Success in preclinical work or early stage clinical trials does not ensure that later stage or larger scale clinical trials will be successful. Conducting clinical trials is a complex, time-consuming and expensive process. Our ability to complete our clinical trials in a timely fashion depends in large part on a number of key factors including protocol design, regulatory and institutional review board approval, the rate of patient enrollment in clinical trials, and compliance with extensive current Good Clinical Practices. [We have opened clinical sites and are enrolling patients in a number of countries where our experience is more limited, and we are in most cases using the services of third party clinical trial providers. If we fail to adequately manage the design, execution and regulatory aspects of our [large, complex and diverse] clinical trials, our studies and ultimately our regulatory approvals may be delayed or we may fail to gain approval for our product candidates. Clinical trials may indicate that our product candidates have harmful side effects or raise other safety concerns that may significantly reduce the likelihood of regulatory approval, result in significant restrictions on use and safety warnings in any approved label, adversely affect placement within the treatment paradigm, or otherwise significantly diminish the commercial potential of the product candidate. Also, positive results in a registrational trial may not be replicated in any subsequent confirmatory trials. Even if later stage clinical trials are successful, regulatory authorities may disagree with our view of the data or require additional studies, and may fail to approve or delay approval of our product candidates or may grant marketing approval that is more restricted than anticipated, including indications for a narrower patient population than expected and the imposition of safety monitoring or educational requirements or risk evaluation and mitigation strategies.] In addition, if another Company is the first to file for marketing approval of a competing orphan drug candidate, that Company may ultimately receive marketing exclusivity for its drug candidate, preventing us from commercializing our orphan drug candidate in the applicable market for several years.

We face significant competition from other biotechnology and pharmaceutical companies.
We are aware of several companies that are working to develop drugs that would compete against our drug candidates. Many of our existing or potential competitors have substantially greater financial, technical and human resources than we do and significantly greater experience in the discovery and development of drug candidates, as well as in obtaining regulatory approvals of those drug candidates in the United States and in foreign countries. Our current and potential future competitors may also have significantly more experience commercializing drugs that have been approved for marketing. Mergers and acquisitions in the pharmaceutical and biotechnology industries could result in even more resources being concentrated among a small number of our competitors.

Competition may increase further as a result of advances in the commercial applicability of technologies and greater availability of capital for investment in these industries. Our competitors may succeed in developing, acquiring or licensing, on an exclusive basis, drug candidates that are more effective or less costly than any drug candidate that we may develop.

Our ability to compete successfully will depend largely on our ability to:

* discover, develop and commercialize drugs that are superior to other products in the market;

* demonstrate through our clinical trials that our drug candidates are differentiated from existing and future therapies;

* attract qualified scientific, product development and commercial personnel;

* obtain patent or other proprietary protection for our drugs and technologies;

* obtain required regulatory approvals; successfully collaborate with pharmaceutical companies in the discovery, development and commercialization of new drugs; and

* negotiate competitive pricing and reimbursement with third party payors

The availability of our competitors' products could limit the demand, and the price we are able to charge, for any drug candidate we develop. The inability to compete with existing or subsequently introduced drug candidates would have a material adverse impact on our business, financial condition and prospects.

Established pharmaceutical companies may invest heavily to accelerate discovery and development of novel compounds or to in license novel compounds that could make our drug candidates less competitive. In addition, any new product that competes with an approved product must demonstrate compelling advantages in efficacy, convenience, tolerability and safety in order to overcome price competition and to be commercially successful. Accordingly, our competitors may succeed in obtaining patent protection, discovering, developing or receiving FDA approval for or commercializing medicines before we do, which would have a material adverse impact on our business.

Our research and development efforts may not succeed in developing commercially successful products and technologies, which may limit our ability to achieve profitability.

We must continue to explore opportunities that may lead to new products and technologies. To accomplish this, we must commit substantial efforts, funds, and other resources to research and development. A high rate of failure is inherent in the research and development of new products and technologies. Any such expenditures that we make will be made without any assurance that our efforts will be successful. Failure can occur at any point in the process, including after significant funds have been invested.

Regardless of whether our clinical trials are deemed to be successful, promising new product candidates may fail to reach the market or may only have limited commercial success because of efficacy or safety concerns, failure to achieve positive clinical outcomes, inability to obtain necessary regulatory approvals or satisfy regulatory criteria, limited scope of approved uses, excessive costs to manufacture, the failure to establish or maintain intellectual property rights, or infringement of the intellectual property rights of others.

Even if we successfully develop new products or enhancements, they may be quickly rendered obsolete by changing customer preferences, changing industry standards, or competitors' innovations. Innovations may not be quickly accepted in the marketplace because of, among

other things, entrenched patterns of clinical practice or uncertainty over third-party reimbursement. We cannot state with certainty when or whether any of our products under development will be launched, whether we will be able to develop, license, or otherwise acquire drug candidates or products, or whether any products will be commercially successful. Failure to launch successful new products or new indications for existing products may cause our products to become obsolete, which may limit our ability to achieve profitability.

Even if we are able to obtain regulatory approvals for our new pharmaceutical products, generic or branded, the success of those products is dependent upon market acceptance.

Levels of market acceptance for our new products could be impacted by several factors, including but not limited to: i) the availability of alternative products from our competitors, ii) the price of our products relative to that of our competitors, iii) the timing of our market entry, iv) the ability to market our products effectively to the retail level and v) the acceptance of our products by government and private entities. Some of these factors are not within our control. Additionally, continuing studies of the proper utilization, safety and efficacy of pharmaceutical products are being conducted by the industry, government agencies and others. Such studies, which increasingly employ sophisticated methods and techniques, can call into question the utilization, safety and efficacy of previously marketed products. In some cases, studies have resulted, and may in the future result, in the discontinuance of product marketing or other risk management programs such as the need for a patient registry. These situations, should they occur, could have a material adverse effect on our profitability, business, financial position and results of operations.

Our manufacturing activity is subject to certain risks.

[We manufacture approximately [ ] percent of the products sold to our customers at our [name of facility] location. As a result, we are dependent upon the uninterrupted and efficient operation of our manufacturing facility in [name of location] and our distribution facilities throughout the country. Our manufacturing facilities and distribution facilities are subject to the risk of catastrophic loss due to, among other things, earthquake, fire, flood, terrorism or other natural or man-made disasters, as well as occurrence of significant equipment failures. If any of these facilities were to experience a catastrophic loss, it would be expected to disrupt our operations and could result in personal injury or property damage, damage relationships with our customers or result in large expenses to repair or replace the facilities or systems, as well as result in other liabilities and adverse impacts.]

[In addition,] we contract with third-party manufacturers to produce [some of] our products in accordance with our specifications and standards. [These contract manufacturers are subject to the same risks as our manufacturing facility as noted above.] While we have implemented stringent quality control procedures to verify that our contract manufacturers comply with our specifications and standards, we do not have full control over their manufacturing activities. Any difficulties, delays and defects in our products resulting from the activities of our contract manufacturers may have an adverse effect on our business and results of operations.

In addition, the occurrence of manufacturing-related compliance issues could require subsequent withdrawal of the drug approval, reformulation of the drug product, additional testing or changes

in labeling of the finished product. Any delay, interruption or cessation of production by our third-party manufacturers or strategic partners of our commercial products or product candidates, or their respective materials and components, as a result of any of the above factors or otherwise, may limit our ability to meet demand for commercial products and/or delay ongoing clinical trials, either of which could have a material adverse effect on our business, results of operations and financial condition.

We could experience difficulties and delays in the manufacturing, distribution and sale of our products.

Our product supply and related patient access could be negatively impacted by, among other things: (i) product seizures or recalls or forced closings of manufacturing plants; (ii) disruption in supply chain continuity including from natural or man-made disasters at one of our facilities or at a critical supplier, as well as our failure or the failure of any of our suppliers to comply with Current Good Manufacturing Practices and other applicable regulations or quality assurance guidelines that could lead to manufacturing shutdowns, product shortages or delays in product manufacturing; (iii) manufacturing, quality assurance/quality control, supply problems or governmental approval delays; (iv) the failure of a sole source or single source supplier to provide us with the necessary raw materials, supplies or finished goods within a reasonable timeframe; (v) the failure of a third-party manufacturer to supply us with bulk active or finished product on time; (vi) construction or regulatory approval delays for new facilities or the expansion of existing facilities, including those intended to support future demand for our biologics products; (vii) the failure to meet new and emerging regulations requiring products to be tracked throughout the distribution channels using unique identifiers to verify their authenticity in the supply chain; and (viii) other manufacturing or distribution issues, including limits to manufacturing capacity due to regulatory requirements, and changes in the types of products produced, such as biologics, physical limitations or other business interruptions, any of which could have a negative effect on our business and results of operations.

Increased concerns over the safety of our products may result in negative publicity or increased regulatory controls on our products.

The Company's reputation is the foundation of our relationships with physicians, patients and other customers. If we are unable to effectively manage real or perceived issues, which could negatively impact sentiments toward the Company, our business could suffer. Pharmaceuticals and medical devices are perceived to be dangerous products and our customers may have a number of concerns about the safety of our products whether or not such concerns have a basis in generally accepted science or peer-reviewed scientific research. These concerns may be increased by negative publicity, even if the publicity is inaccurate. [In addition, government investigations related to the use of our products, but not the efficacy of the products themselves, may cause reputational harm to the Company.] Negative publicity could also result in an increased number of product liability claims, whether or not these claims have a basis in scientific fact.

[We are also subject to adverse event reporting regulations that require us to report to the FDA or similar bodies in other countries if our products are associated with a death or serious injury, even if there is no available evidence of a causal relationship between the adverse event and the

product. Such reports may be publicly released by the FDA and other authorities.] Furthermore, any adverse publicity associated with adverse events for our products, and related post-marketing actions, could cause consumers to seek alternatives to our products, and thereby cause our sales to decline, even if our products are ultimately determined not to have been the primary cause of the adverse event.

Pharmaceutical products can develop unexpected safety or efficacy concerns, which could have a material adverse effect on our business.

Pharmaceutical products receive regulatory approval based on data obtained in controlled clinical trials of limited duration. After approval, the products are used for longer periods of time by much larger numbers of patients; we and others (including regulatory agencies and private payers) collect extensive information on the efficacy and safety of our marketed products by continuously monitoring the use of our products in the marketplace. In addition, we or others may conduct post-marketing clinical studies on efficacy and safety of our marketed products. New safety or efficacy data from market surveillance, post-marketing clinical studies or general use may result in product label changes, product recalls, withdrawals, or declining sales, as well as product liability, consumer fraud and/or other claims, including potential civil or criminal governmental actions.

Product labeling changes for our marketed products could result in a negative impact on revenues.

We or regulatory authorities may need to change the labeling for any pharmaceutical product, including after a product has been marketed for several years. These changes are often the result of additional data from post-marketing studies, head-to-head trials, adverse events reports, studies that identify biomarkers (objective characteristics that can indicate a particular response to a product or therapy) or other studies or post-marketing experience that produce important additional information about a product. New information added to a product's label can affect its risk-benefit profile, leading to potential recalls, withdrawals, or declining revenue, as well as product liability claims. Sometimes additional information from these studies identifies a portion of the patient population that may be non-responsive to a medicine or would be at higher risk of adverse reactions and labeling changes based on such studies may limit the patient population. The studies providing such additional information may be sponsored by us, but they could also be sponsored by competitors, insurance companies, government institutions, managed care organizations, scientists, investigators, or other interested parties. While additional safety and efficacy information from such studies assist us and healthcare providers in identifying the best patient population for each product, it can also negatively impact our revenues due to inventory returns and a more limited patient population going forward. Additionally, certain study results, especially from head-to-head trials, could affect a product's formulary listing, which could also adversely affect our revenues.

We are dependent on our collaborative agreements for the development of products and business development, which exposes us to the risk of reliance on the viability of third parties.

In conducting our research and development activities, we currently rely, and will in the future rely, on collaborative agreements with third parties such as manufacturers, contract research

organizations, commercial partners, universities, governmental agencies and not-for-profit organizations for both strategic and financial resources. The loss of, or failure to perform by us or our partners under, any applicable agreements or arrangements, or our failure to secure additional agreements for other products in development, would substantially disrupt or delay our research and development and commercialization activities. Any such loss would likely increase our expenses and materially harm our business, financial condition and results of operation.

We extensively outsource our clinical trial activities and usually perform only a small portion of the start-up activities in-house.

We rely on independent third-party contract research organizations (CROs) to perform most of our clinical studies, including document preparation, site identification, screening and preparation, pre-study visits, training, program management and bioanalytical analysis. Many important aspects of the services performed for us by the CROs are out of our direct control. If there is any dispute or disruption in our relationship with our CROs, our clinical trials may be delayed. Moreover, in our regulatory submissions, we rely on the quality and validity of the clinical work performed by third-party CROs. If any of our CROs' processes, methodologies or results were determined to be invalid or inadequate, our own clinical data and results and related regulatory approvals could be adversely impacted.

Reliance on third-party relationships and outsourcing arrangements could adversely affect our business.

We utilize third parties, including suppliers, alliances with other pharmaceutical and biotechnology companies, and third-party service providers, for selected aspects of product development, the manufacture and commercialization of certain products, support for information technology systems, and certain financial transactional processes. For example, we outsource [the day-to-day management and oversight of our clinical trials to contract research organizations] [the manufacture of certain of our products]. Outsourcing these functions involves the risk that the third parties may not perform to our standards or legal requirements, may not produce reliable results, may not perform in a timely manner, may not maintain the confidentiality of our proprietary information, or may fail to perform at all. Failure of these third parties to meet their contractual, regulatory, confidentiality, or other obligations to us could have a material adverse effect on our business.

Product liability claims could harm our business.

The development, manufacture, testing, marketing and sale of pharmaceutical products are associated with significant risks of product liability claims. Side effects or adverse events known or reported to be associated with, or manufacturing defects in, the products sold by us could exacerbate a patient's condition, or could result in serious injury or impairments or even death. This could result in product liability. Some of our products, including [name product], have boxed warnings in their labels. Product liability claims may be brought by individuals seeking relief for themselves, or by groups seeking to represent a class of injured patients. Further, third party payors, either individually or as a putative class, may bring actions seeking to recover monies spent on one of our products. As sales of our products increase, the risk that product

liability claims will be made against us increases. The risk of product liability claims may also increase if a company receives a warning letter from a regulatory agency. We cannot predict the frequency, outcome or cost to defend any such claims.

Product liability insurance coverage is expensive, can be difficult to obtain and may not be available to us in the future on acceptable terms, or at all. Our product liability insurance may not cover all of the future liabilities we might incur in connection with the development, manufacture or sale of our products. In addition, we may not continue to be able to obtain insurance on satisfactory terms or in adequate amounts. A successful claim or claims brought against us in excess of available insurance coverage could subject us to significant liabilities and could have a material adverse effect on our business, financial condition, results of operations and growth prospects. Such claims whether meritorious or not could also harm our reputation and the reputation of our products, adversely affecting our ability to market our products successfully. In addition, defending a product liability lawsuit is expensive and can divert the attention of key employees from operating our business.

In addition, product liability claims could result in an investigation of the safety or efficacy of our products, our manufacturing processes and facilities, or our marketing programs conducted by the FDA, the EMA, or the competent authorities of the EU member states. Such investigations could also potentially lead to a recall of our products or more serious enforcement actions, limitations on the indications for which they may be used, or suspension, variation, or withdrawal of approval, any of which would adversely affect our business.

Limited reimbursement or insurance coverage of our approved products, if any, by third party payors may render our products less attractive to patients and healthcare providers.

Market acceptance and sales of any approved products will depend significantly on reimbursement or coverage of our products by third party payors and may be affected by existing and future healthcare reform measures or the prices of related products for which third party reimbursement applies. Coverage and reimbursement by a third party payor may depend upon a number of factors, including the third party payor's determination that use of a product is: a covered benefit under its health plan; safe, effective and medically necessary; appropriate for the specific patient; cost-effective; and neither experimental nor investigational.

Obtaining coverage and reimbursement approval for a product from a government or other third party payor is a time consuming and costly process that could require us to provide supporting scientific, clinical and cost-effectiveness data for the use of our products to the payor, which we may not be able to provide. Furthermore, the reimbursement policies of third party payors may significantly change in a manner that renders our clinical data insufficient for adequate reimbursement or otherwise limits the successful marketing of our products. Even if we obtain coverage for our product candidates, third party payors may not establish adequate reimbursement amounts, which may reduce the demand for, or the price of, our products. If reimbursement is not available or is available only to limited levels, we may not be able to commercialize certain of our products.

Publication of discounts by third party payors or authorities may lead to further pressure on the prices or reimbursement levels within the country of publication and other countries. If

reimbursement of our products is unavailable or limited in scope or amount, or if pricing is set at unacceptable levels, we or our partner may elect not to commercialize our products, and our business and financial condition could be adversely affected.

If we are unable to negotiate and maintain satisfactory arrangements with group purchasing organizations with respect to the purchase of our products, our business could be adversely affected.

Our ability to sell our products to hospitals in the United States depends in part on our relationships with group purchasing organizations, or GPOs. Many existing and potential customers for our products become members of GPOs. GPOs negotiate pricing arrangements and contracts, sometimes on an exclusive basis, with medical supply manufacturers and distributors. These negotiated prices are then made available to a GPO's affiliated hospitals and other members. If we are not one of the providers selected by a GPO, affiliated hospitals and other members may be less likely to purchase our products, and if the GPO has negotiated a strict sole source, market share compliance or bundling contract for another manufacturer's products, we may be precluded from making sales to members of the GPO for the duration of the contractual arrangement. Our failure to renew contracts with GPOs may cause us to lose market share and could have a material adverse effect on our sales, financial condition and results of operations. We cannot assure you that we will be able to renew these contracts at the current or substantially similar terms. If we are unable to keep our relationships and develop new relationships with GPOs, our competitive position may suffer.

We are subject to complex government healthcare legislation and reimbursement programs, as well as other cost-containment pressures.

Many of our products are purchased or reimbursed by federal and state government authorities, private health insurers and other organizations, including heath maintenance and managed care organizations. These third-party payors increasingly challenge pharmaceutical and medical device product pricing, which could result in lower reimbursement rates and a reduction in demand for our products.

In addition, legislative and regulatory proposals and enactments to reform healthcare insurance programs could significantly influence the manner in which pharmaceutical products, biologic products and medical devices are prescribed and purchased. Individual states have also become increasingly aggressive in passing legislation and implementing regulations designed to control pharmaceutical product pricing, including price or patient reimbursement constraints, discounts, restrictions on certain product access, and to encourage importation from other countries and bulk purchasing. Furthermore, regional healthcare authorities and individual hospitals are increasingly using bidding procedures to determine what pharmaceutical products and which suppliers will be included in their prescription drug and other healthcare programs. Any legally mandated price controls or utilization of bidding procedures could negatively and materially impact our revenues, results of operations and financial condition.

Sales of our products may be adversely affected by the continuing consolidation of our customer base.

A significant proportion of our sales is made to relatively few U.S. retail drug chains, wholesalers, managed care purchasing organizations, mail order distributors and hospitals. These customers are continuing to undergo significant consolidation. Net sales to one such customer in [year] accounted for [ ]% of our total consolidated sales. Such consolidation has provided and may continue to provide them with additional purchasing leverage, and consequently may increase the pricing pressures that we face. Additionally, the emergence of large buying groups representing independent retail pharmacies, and the prevalence and influence of managed care organizations and similar institutions, enable those groups to extract price discounts on our products, which could have a material adverse effect on our business, financial condition and results of operations.

Increased pricing pressure and other restrictions in the U.S. and abroad from managed care organizations, institutional Investors, and government agencies and programs, among others, could negatively affect our revenues and profit margins.

Our products continue to be subject to increasing pressures from market access, pricing and rebates and other restrictions in the U.S.[, the EU and other regions around the world], including from (i) rules and practices of managed care organizations and institutional and governmental Investors; (ii) judicial decisions and governmental laws and regulations for Medicare, Medicaid and U.S. healthcare reform, including the 2010 Patient Protection and Affordable Care Act; (iii) the potential impact of pharmaceutical reimbursement, Medicare Part D Formularies and product pricing in general; (iv) delays in gaining reimbursement; (v) government price erosion mechanisms across Europe and in other countries, resulting in deflation for pharmaceutical product pricing; (vi) collection delays in government-funded public hospitals outside the U.S. (vii) the impact on pricing from parallel trade across borders; (viii) other developments in technology and/or industry practices that could impact the reimbursement policies and practices of third-party payers; and (ix) limited or blocked market access due to real or perceived differences in value propositions for our products compared to competing products.

The illegal importation of counterfeit products and pharmaceutical and medical device products from countries where government price controls or other market dynamics result in lower prices may adversely affect our sales and profitability in the U.S. and other countries in which we operate.

Foreign imports are illegal under current U.S. law, with the sole exception of limited quantities of prescription drugs imported for personal use. However, the volume of illegal imports continues to rise as the ability of patients and other customers to obtain these lower priced imports has grown significantly. In addition, U.S. policy makers may expand consumers' ability to import lower priced versions of our products and competing products from Canada, where there are government price controls. Any future legislation or regulations that increase consumer access to lower priced medicines from outside the U.S. may lower the prices we receive for our products, which could adversely impact our revenues.

Illegal imports and counterfeit products may reduce demand for our products.

The illegal importation of counterfeit products and pharmaceutical products from countries where government price controls or other market dynamics result in lower prices may adversely

affect our sales and profitability in the United States and other countries in which we operate. Foreign imports are illegal under current U.S. law, with the sole exception of limited quantities of prescription drugs imported for personal use. However, the volume of illegal imports continues to rise as the ability of patients and other customers to obtain these lower priced imports has grown significantly. In addition, U.S. policy makers may expand consumers' ability to import lower priced versions of our products and competing products from Canada, where there are government price controls. Any future legislation or regulations that increase consumer access to lower priced medicines from outside the United States could adversely impact our revenues.

In addition, third parties may illegally distribute and sell counterfeit versions of our products, which do not meet our rigorous manufacturing and testing standards. A patient who receives a counterfeit drug may be at risk for a number of dangerous health consequences. Our reputation and business could suffer harm as a result of counterfeit drugs sold under our brand name. In addition, thefts of inventory at warehouses, plants or while in-transit, which are then not properly stored and are later sold through unauthorized channels, could adversely impact patient safety, our reputation and our business.

The Company could be negatively impacted if found to have infringed on intellectual property rights.

Technology companies, including many of the Company's competitors, frequently enter into litigation based on allegations of patent infringement or other violations of intellectual property rights. In addition, patent holding companies seek to monetize patents they have purchased or otherwise obtained. As the Company grows, the intellectual property rights claims against it will likely increase. The Company intends to vigorously defend infringement actions in court and before the U.S. International Trade Commission. The plaintiffs in these actions frequently seek injunctions and substantial damages. Regardless of the scope or validity of such patents or other intellectual property rights, or the merits of any claims by potential or actual litigants, the Company may have to engage in protracted litigation. If the Company is found to infringe one or more patents or other intellectual property rights, regardless of whether it can develop non-infringing technology, it may be required to pay substantial damages or royalties to a third-party, or it may be subject to a temporary or permanent injunction prohibiting the Company from marketing or selling certain products. In certain cases, the Company may consider the desirability of entering into licensing agreements, although no assurance can be given that such licenses can be obtained on acceptable terms or that litigation will not occur. These licenses may also significantly increase the Company's operating expenses.

Regardless of the merit of particular claims, litigation may be expensive, time-consuming, disruptive to the Company's operations and distracting to management. In recognition of these considerations, the Company may enter into arrangements to settle litigation. If one or more legal matters were resolved against the Company's consolidated financial statements for that reporting period could be materially adversely affected. Further, such an outcome could result in significant compensatory, punitive or trebled monetary damages, disgorgement of revenue or profits, remedial corporate measures or injunctive relief against the Company that could adversely affect its financial condition and results of operations.

Indemnity provisions in various agreements potentially expose us to substantial liability for intellectual property infringement and other losses.

Our agreements with advertisers, advertising agencies, customers and other third parties may include indemnification provisions under which we agree to indemnify them for losses suffered or incurred as a result of claims of intellectual property infringement, damages caused by us to property or persons, or other liabilities relating to or arising from our products, services or other contractual obligations. The term of these indemnity provisions generally survives termination or expiration of the applicable agreement. Large indemnity payments would harm our business, financial condition and results of operations. In addition, any type of intellectual property lawsuit, whether initiated by us or a third party, would likely be time consuming and expensive to resolve and would divert management's time and attention.

We rely heavily on our technology and intellectual property, but we may be unable to adequately or cost-effectively protect or enforce our intellectual property rights, thereby weakening our competitive position and increasing operating costs.

To protect our rights in our services and technology, we rely on a combination of copyright and trademark laws, patents, trade secrets, confidentiality agreements with employees and third parties, and protective contractual provisions. We also rely on laws pertaining to trademarks and domain names to protect the value of our corporate brands and reputation. Despite our efforts to protect our proprietary rights, unauthorized parties may copy aspects of our services or technology, obtain and use information, marks, or technology that we regard as proprietary, or otherwise violate or infringe our intellectual property rights. In addition, it is possible that others could independently develop substantially equivalent intellectual property. If we do not effectively protect our intellectual property, or if others independently develop substantially equivalent intellectual property, our competitive position could be weakened.

Effectively policing the unauthorized use of our services and technology is time-consuming and costly, and the steps taken by us may not prevent misappropriation of our technology or other proprietary assets. The efforts we have taken to protect our proprietary rights may not be sufficient or effective, and unauthorized parties may copy aspects of our services, use similar marks or domain names, or obtain and use information, marks, or technology that we regard as proprietary. We may have to litigate to enforce our intellectual property rights, to protect our trade secrets, or to determine the validity and scope of others' proprietary rights, which are sometimes not clear or may change. Litigation can be time consuming and expensive, and the outcome can be difficult to predict.

We rely on agreements with third parties to provide certain services, goods, technology, and intellectual property rights necessary to enable us to implement some of our applications.

Our ability to implement and provide our applications and services to our clients depends, in part, on services, goods, technology, and intellectual property rights owned or controlled by third parties. These third parties may become unable to or refuse to continue to provide these services, goods, technology, or intellectual property rights on commercially reasonable terms consistent with our business practices, or otherwise discontinue a service important for us to continue to operate our applications. If we fail to replace these services, goods, technologies, or intellectual

property rights in a timely manner or on commercially reasonable terms, our operating results and financial condition could be harmed. In addition, we exercise limited control over our third-party vendors, which increases our vulnerability to problems with technology and services those vendors provide. If the services, technology, or intellectual property of third parties were to fail to perform as expected, it could subject us to potential liability, adversely affect our renewal rates, and have an adverse effect on our financial condition and results of operations.

We depend on profitable royalty-bearing licenses of our technology, and if we are unable to maintain and generate such license agreements, then we may not be able to sustain existing levels of revenue or increase revenue.

We depend upon the identification, investment in and license of new patents for our revenues. If we are unable to maintain such license agreements and to continue to develop new license arrangements, then we may not have the resources to identify new technology-based opportunities for future patents and inventions in order to maintain sustainable revenue and growth.

Our current or future license agreements may not provide the volume or quality of royalty revenue to sustain our business. In some cases, other technology sources may compete against us as they seek to license and commercialize technologies. These and other strategies may reduce the number of technology sources and potential clients to whom we can market our services. Our inability to maintain current relationships and sources of technology or to secure new licensees, may have a material adverse effect on our business and results of operations.

If we fail to maintain or expand our relationships with our suppliers[, in some cases single-source suppliers,] we may not have adequate access to new or key technology necessary for our products, which may impair our ability to deliver leading-edge products.

In addition to the technologies we develop, our suppliers develop product innovations at our direction that are requested by our customers. Further, we rely heavily on our component suppliers, such as [name of suppliers], to provide us with leading-edge components that conform to required specifications or contractual arrangements on time and in accordance with a product roadmap. If we are not able to maintain or expand our relationships with our suppliers or continue to leverage their research and development capabilities to develop new technologies desired by our customers, our ability to deliver leading-edge products in a timely manner may be impaired and we could be required to incur additional research and development expenses. Also, disruption in our supply chain or the need to find alternative suppliers could impact the costs and/or timing associated with procuring necessary products, components and services. Similarly, suppliers have operating risks that could impact our business. These risks could create product time delays, inventory and invoicing problems, staging delays, and other operational difficulties.

We must acquire or develop new products, evolve existing ones, address any defects or errors, and adapt to technology change.

Technical developments, client requirements, programming languages, and industry standards change frequently in our markets. As a result, success in current markets and new markets will depend upon our ability to enhance current products, address any product defects or errors,

acquire or develop and introduce new products that meet client needs, keep pace with technology changes, respond to competitive products, and achieve market acceptance. Product development requires substantial investments for research, refinement, and testing. We may not have sufficient resources to make necessary product development investments. We may experience technical or other difficulties that will delay or prevent the successful development, introduction, or implementation of new or enhanced products. We may also experience technical or other difficulties in the integration of acquired technologies into our existing platform and applications. Inability to introduce or implement new or enhanced products in a timely manner could result in loss of market share if competitors are able to provide solutions to meet customer needs before we do, give rise to unanticipated expenses related to further development or modification of acquired technologies as a result of integration issues, and adversely affect future performance.

Our failure to deliver high quality server solutions could damage our reputation and diminish demand for our products, and subject us to liability.

Our customers require our products to perform at a high level, contain valuable features and be extremely reliable. The design of our server solutions is sophisticated and complex, and the process for manufacturing, assembling and testing our server solutions is challenging. Occasionally, our design or manufacturing processes may fail to deliver products of the quality that our customers require. For example, a vendor may provide us with a defective component that failed under certain heavy use applications. As a result, our product would need to be repaired. The vendor may agree to pay for the costs of the repairs, but we may incur costs in connection with the recall and diverted resources from other projects. New flaws or limitations in our products may be detected in the future. Part of our strategy is to bring new products to market quickly, and first-generation products may have a higher likelihood of containing undetected flaws. If our customers discover defects or other performance problems with our products, our customers' businesses, and our reputation, may be damaged. Customers may elect to delay or withhold payment for defective or underperforming products, request remedial action, terminate contracts for untimely delivery, or elect not to order additional products. If we do not properly address customer concerns about our products, our reputation and relationships with our customers may be harmed. In addition, we may be subject to product liability claims for a defective product. Any of the foregoing could have an adverse effect on our business and results of operations.

Cyclical and seasonal fluctuations in the economy, in internet usage and in traditional retail shopping may have an effect on our business.

Both cyclical and seasonal fluctuations in internet usage and traditional retail seasonality may affect our business. Internet usage generally slows during the summer months, and queries typically increase significantly in the fourth quarter of each year. These seasonal trends may cause fluctuations in our quarterly results, including fluctuations in revenues.

The products we sell are advanced, and we need to rapidly and successfully develop and introduce new products in a competitive, demanding and rapidly changing environment.

To succeed in our intensely competitive industry, we must continually improve, refresh and expand our product and service offerings to include newer features, functionality or solutions,

and keep pace with price-to-performance gains in the industry. Shortened product life cycles due to customer demands and competitive pressures impact the pace at which we must introduce and implement new technology. This requires a high level of innovation by both our software developers and the suppliers of the third-party software components included in our systems. In addition, bringing new solutions to the market entails a costly and lengthy process, and requires us to accurately anticipate customer needs and technology trends. We must continue to respond to market demands, develop leading technologies and maintain leadership in analytic data solutions performance and scalability, or our business operations may be adversely affected.

We must also anticipate and respond to customer demands regarding the compatibility of our current and prior offerings. These demands could hinder the pace of introducing and implementing new technology. Our future results may be affected if our products cannot effectively interface and perform well with software products of other companies and with our customers' existing IT infrastructures, or if we are unsuccessful in our efforts to enter into agreements allowing integration of third-party technology with our database and software platforms. Our efforts to develop the interoperability of our products may require significant investments of capital and employee resources. In addition, many of our principal products are used with products offered by third parties and, in the future, some vendors of non-Company products may become less willing to provide us with access to their products, technical information and marketing and sales support. As a result of these and other factors, our ability to introduce new or improved solutions could be adversely impacted and our business would be negatively affected.

Industry consolidation may result in increased competition, which could result in a loss of customers or a reduction in revenue.

Some of our competitors have made or may make acquisitions or may enter into partnerships or other strategic relationships to offer more comprehensive services than they individually had offered or achieve greater economies of scale. In addition, new entrants not currently considered to be competitors may enter our market through acquisitions, partnerships or strategic relationships. We expect these trends to continue as companies attempt to strengthen or maintain their market positions. The potential entrants may have competitive advantages over us, such as greater name recognition, longer operating histories, more varied services and larger marketing budgets, as well as greater financial, technical and other resources. The companies resulting from combinations or that expand or vertically integrate their business to include the market that we address may create more compelling service offerings and may offer greater pricing flexibility than we can or may engage in business practices that make it more difficult for us to compete effectively, including on the basis of price, sales and marketing programs, technology or service functionality. These pressures could result in a substantial loss of our customers or a reduction in our revenue.

Our business could be negatively impacted by cyber security threats, attacks and other disruptions.

Like others in our industry, we continue to face advanced and persistent attacks on our information infrastructure where we manage and store various proprietary information and sensitive/confidential data relating to our operations. These attacks may include sophisticated

malware (viruses, worms, and other malicious software programs) and phishing emails that attack our products or otherwise exploit any security vulnerabilities. These intrusions sometimes may be zero-day malware that are difficult to identify because they are not included in the signature set of commercially available antivirus scanning programs. Experienced computer programmers and hackers may be able to penetrate our network security and misappropriate or compromise our confidential information or that of our customers or other third-parties, create system disruptions, or cause shutdowns. Additionally, sophisticated software and applications that we produce or procure from third-parties may contain defects in design or manufacture, including "bugs" and other problems that could unexpectedly interfere with the operation of the information infrastructure. A disruption, infiltration or failure of our information infrastructure systems or any of our data centers as a result of software or hardware malfunctions, computer viruses, cyber attacks, employee theft or misuse, power disruptions, natural disasters or accidents could cause breaches of data security, loss of critical data and performance delays, which in turn could adversely affect our business.

If we do not respond to technological changes or upgrade our websites and technology systems, our growth prospects and results of operations could be adversely affected.

To remain competitive, we must continue to enhance and improve the functionality and features of our websites and technology infrastructure. As a result, we will need to continue to improve and expand our hosting and network infrastructure and related software capabilities. These improvements may require greater levels of spending than we have experienced in the past. Without such improvements, our operations might suffer from unanticipated system disruptions, slow application performance or unreliable service levels, any of which could negatively affect our reputation and ability to attract and retain customers and contributors. Furthermore, in order to continue to attract and retain new customers, we are likely to incur expenses in connection with continuously updating and improving our user interface and experience. We may face significant delays in introducing new services, products and enhancements. If competitors introduce new products and services using new technologies or if new industry standards and practices emerge, our existing websites and our proprietary technology and systems may become obsolete or less competitive, and our business may be harmed. In addition, the expansion and improvement of our systems and infrastructure may require us to commit substantial financial, operational and technical resources, with no assurance that our business will improve.

We currently obtain components from single or limited sources, and are subject to significant supply and pricing risks.

Many components, including those that are available from multiple sources, are at times subject to industry-wide shortages and significant commodity pricing fluctuations. While the Company has entered into agreements for the supply of many components, there can be no assurance that we will be able to extend or renew these agreements on similar terms, or at all. A number of suppliers of components may suffer from poor financial conditions, which can lead to business failure for the supplier or consolidation within a particular industry, further limiting our ability to obtain sufficient quantities of components. The follow-on effects from global economic conditions on our suppliers, also could affect our ability to obtain components. Therefore, we remain subject to significant risks of supply shortages and price increases.

Our products often utilize custom components available from only one source. Continued availability of these components at acceptable prices, or at all, may be affected for any number of reasons, including if those suppliers decide to concentrate on the production of common components instead of components customized to meet our requirements. The supply of components for a new or existing product could be delayed or constrained, or a key manufacturing vendor could delay shipments of completed products to us adversely affecting our business and results of operations.

The Company depends on the performance of distributors, carriers and other resellers.
The Company distributes its products through [cellular network carriers,] wholesalers, national and regional retailers, and value-added resellers, many of whom distribute products from competing manufacturers. The Company also sells its products and third-party products in most of its major markets directly to education, enterprise and government customers, and consumers and small and mid-sized businesses through its online and retail stores.

[Carriers providing cellular network service for iPhone typically subsidize users' purchases of the device. There is no assurance that such subsidies will be continued at all or in the same amounts upon renewal of the Company's agreements with these carriers or in agreements the Company enters into with new carriers.]

Many resellers have narrow operating margins and have been adversely affected in the past by weak economic conditions. Some resellers have perceived the expansion of the Company's direct sales as conflicting with their business interests as distributors and resellers of the Company's products. Such a perception could discourage resellers from investing resources in the distribution and sale of the Company's products or lead them to limit or cease distribution of those products. The Company has invested and will continue to invest in programs to enhance reseller sales, including [staffing selected resellers' stores with Company employees and contractors, and] improving product placement displays. These programs could require a substantial investment while providing no assurance of return or incremental revenue. The financial condition of these resellers could weaken, these resellers could stop distributing the Company's products, or uncertainty regarding demand for the Company's products could cause resellers to reduce their ordering and marketing of the Company's products.

In addition to the risks listed above, businesses are often subject to risks not foreseen or fully appreciated by the management. It is not possible to foresee all risks that may affect us. Moreover, the Company cannot predict whether the Company will successfully effectuate the Company's current business plan. Each prospective Purchaser is encouraged to carefully analyze the risks and merits of an investment in the Securities and should take into consideration when making such analysis, among other, the Risk Factors discussed above.

BUSINESS

Description of the Business

Genobank allows users to track and establish ownership of their genetic information as private property through the use of blockchain technology. We sell a white-label (patent pending) anonymous DNA/RNA extraction kit that is represented by a non-fungible token based in the ethereum standard ERC721 to allow users/patients establish ownership and control over their biosamples and residual biological datasets.

Business Plan

History of the Business

The Company's Products and/or Services

Product / Service	Description	Current Market
Incrypto Privacy-preserving Saliva DNA Extraction Kit	First ever incognito saliva DNA/RNA extraction kit	Direct to Consumer Genetic Test, Clinical Trial Recruitment Services, Privacy-preserving Ancestry Companies
Incrypto Privacy-preserving Family Saliva DNA extraction Kit	First ever incognito saliva DNA/RNA extraction kit for Families, it includes a minimum of 3 sample tubes or swabs.	Direct to Consumer Genetic Test, Clinical Trial Recruitment Services, Privacy-preserving Ancestry Companies.
Incrypto Privacy-preserving DNA extraction kits Vending Machine (ATM).	A vending machine or kiosk to purchase DNA test kits using only QR codes and a pin code to preserve user's privacy.	Direct to Consumer Genetic Test, Clinical Trial Recruitment Services, Privacy-preserving Ancestry, Nutrigenomics Companies.

We have no new products in development.

We offer our platform services to Laboratories and Researchers through our website and our API that allow them to exchange DNA/RNA biosamples and biological data in a secure, private and transparent way with their users/patients using blockchain technology to record every permission in the form of a tokenized consent.

Competition

The Company's primary competitors are 23&me, AncestryDNA, MyHeritage, Nebula Genomics, and LunaDNA.

Genobank's DNA extraction kit allows consumers to retain the rights to their BioData, dictate how that BioData is used and benefit from any sales of their BioData.

Supply Chain and Customer Base

The Company relies on outside suppliers for all of its manufacturing supplies, parts, and components. These component suppliers can be replaced within a reasonable period of time.

The products are focused on demographics which demonstrate concern about privacy but still want to be a part of the genomic revolution. Products are specifically designed for users who were considering a DNA test, but decided against it because of privacy concerns and how their private BioData may be used, exploited, and stored. Usually Diagnostic Laboratories and Genomic Research entities.

Intellectual Property

Patents

Application or Registration #	Title	Description	File Date	Grant Date	Country

16428700	Anonymous DNA/RNA bio specimen tracking process, anonymous DNA/RNA bio specimen tracking system, and an anonymous DNA/RNA bio specimen extraction kit that utilizes public key infrastructure, asymmetric encryption and blockchain to relate, claim ownership, consent to use, and keep track of correspondin g data sets.	Patent	May 31, 2018		US
62878585	Privacy-protecting DNA/RNA/ Microbiome test kit kiosk and locker that pairs to and stores results data in private digital wallet.	Provisional patent	June 25, 2019		US

62936166	Anonymous DNA/RNA bio specimen tracking process for human families to use and an anonymous DNA/RNA bio specimen extraction kit that utilizes public key infrastructure (PKI), asymmetric encryption, non-fungible-tokens (NFT) on a public blockchain to create a self-sovereign digital DNA fingerprint for each DNA donor, relate, claim ownership, consent to use, establish biological relationship between donors, and keep track of correspondin g mulitomics data sets.	Provisional Patent	November 15, 2019		US

Trademarks

Application or Registration #	Goods / Services	Mark	File Date	Registration Date	Country
88344176	Service Mark	Incrypto Saliva DNA Extraction Kit	March 18, 2019	May 29, 2019	US
87750068	Trademark	Genobank.io	January 2, 2019	July 16, 2019	US

Governmental/Regulatory Approval and Compliance

We are subject to extensive federal, state and local laws and regulations, including the recently enacted comprehensive health care reform legislation with respect health care for our employees, those relating to building and zoning requirements and those relating to the preparation and sale of food. Such laws and regulations are subject to change from time to time. Typically, licenses, permits and approvals under such laws and regulations must be renewed annually and may be revoked, suspended or denied renewal for cause at any time if governmental authorities determine that our conduct violates applicable regulations.

Litigation

There are no existing legal suits pending, or to the Company's knowledge, threatened, against the Company.

Other

The Company's principal address is 55 E Third Ave, San Mateo, CA 94401

The Company has the following additional addresses:

The Company conducts business in United States.

DIRECTORS, OFFICERS AND EMPLOYEES

Directors

The directors or managers of the Company are listed below along with all positions and offices held at the Company and their principal occupation and employment responsibilities for the past three (3) years and their educational background and qualifications.

Name

Daniel Uribe

All positions and offices held with the Company and date such position(s) was held with start and ending dates

Chief Executive Officer, President, Secretary

Principal occupation and employment responsibilities during at least the last three (3) years with start and ending dates

Chief Executive Officer, President, Secretary, Genobank.io, 2018-Present CEO, United IT Consultants, Inc., 2004-2018

Education

MBA, IPADE Business School; BS in Electronic and Communications Engineering

Officers of the Company

The officers of the Company are listed below along with all positions and offices held at the Company and their principal occupation and employment responsibilities for the past three (3) years and their educational background and qualifications.

Name

Daniel Uribe

All positions and offices held with the Company and date such position(s) was held with start and ending dates

Chief Executive Officer, President, Secretary

Principal occupation and employment responsibilities during at least the last three (3) years with start and ending dates

Chief Executive Officer, President, Secretary, Genobank.io, 2018-Present CEO, United IT Consultants, Inc., 2004-2018

Education

MBA, IPADE Business School; BS in Electronic and Communications Engineering

Name

Candy Bellau

All positions and offices held with the Company and date such position(s) was held with start and ending dates

CFO, Treasurer

Principal occupation and employment responsibilities during at least the last three (3) years with start and ending dates

CFO, Treasurer, Genobank.io, 2020-Present

Education

Dowling Collegue, BBA, Accounting. Dowling Collegue, Finance, MBA

Indemnification

Indemnification is authorized by the Company to directors, officers or controlling persons acting in their professional capacity pursuant to California law. Indemnification includes expenses such as attorney's fees and, in certain circumstances, judgments, fines and settlement amounts actually paid or incurred in connection with actual or threatened actions, suits or proceedings involving such person, except in certain circumstances where a person is adjudged to be guilty of gross negligence or willful misconduct, unless a court of competent jurisdiction determines that such indemnification is fair and reasonable under the circumstances.

Employees

The Company currently has 2 employees in .

CAPITALIZATION AND OWNERSHIP

Capitalization

The Company has issued the following outstanding Securities:

Type of security	Common Stock
Amount outstanding	7,100,000
Voting Rights	One vote per share.

Anti-Dilution Rights	N/A
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	50,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	10,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	25,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	10,000
Voting Rights	None.
Anti-Dilution Rights	None.

How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	10,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	25,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	10,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	50,000
Voting Rights	None.
Anti-Dilution Rights	None.

How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

Type of security	SAFE (Simple Agreement for Future Equity)
Amount outstanding	25,000
Voting Rights	None.
Anti-Dilution Rights	None.
How this Security may limit, dilute or qualify the Notes/Bonds issued pursuant to Regulation CF	N/A

The Company has the following debt outstanding:

Type of debt	Convertible Notes
Name of creditor	Paul Grotowski
Amount outstanding	$25,000.00
Interest rate and payment schedule	2%
Amortization schedule	
Describe any collateral or security	None.
Maturity date	August 7, 2020
Other material terms	

Type of debt	Notes
Name of creditor	CrowdSAFE
Amount outstanding	$473,708.00
Interest rate and payment schedule	0
Amortization schedule	0
Describe any collateral or security	security
Maturity date	
Other material terms	

The total amount of outstanding debt of the company is $0.00

The Company has conducted the following prior Securities offerings in the past three years:

Security Type	Number Sold	Money Raised	Use of Proceeds	Offering Date	Exemption from Registration Used or Public Offering
SAFE (Simple Agreement for Future Equity)	9	$215,000.00	Develop DNAWallet & ATM prototypes, Develop the 3 patents, crowdfunding campaign and Marketing Materials	February 24, 2019	Rule 506(b)

Ownership

A majority of the Company is owned by Daniel Uribe.

Below the beneficial owners of 20% percent or more of the Company's outstanding voting equity securities, calculated on the basis of voting power, are listed along with the amount they own.

Name	Percentage Owned
Daniel Uribe	71.8%

FINANCIAL INFORMATION

Please see the financial information listed on the cover page of this Form C-AR and attached hereto in addition to the following information. Financial statements are attached hereto as Exhibit A.

Recent Tax Return Information

Total Income	Taxable Income	Total Tax
$0.00	$0.00	$0.00

Operations

The Company will complete its seed round of financing on July 2021. Following the Offering, we should have enough liquidity to execute our business plan until July 2022. We intend to be profitable by July 2022. Our significant challenges are developing and marketing a viable product in a competitive environment to ensure we have sufficient billable transactions in our secure network.

The Company intends to achieve profitability in the next 12 months by integrating our platform to 100 to 150 Laboratories and Research institutions by July 2022.

Liquidity and Capital Resources

On August 1st, 2020 the Company conducted an offering pursuant to Regulation CF and raised $464,420.00.

The Company has the following sources of capital in addition to the proceeds from the Regulation CF Offering:
Additional Capital raises.

Capital Expenditures and Other Obligations

The Company does not intend to make any material capital expenditures in the future.

Material Changes and Other Information

Trends and Uncertainties

The financial statements are an important part of this Form C-AR and should be reviewed in their entirety. The financial statements of the Company are attached hereto as Exhibit A.

Restrictions on Transfer

Any Securities sold pursuant to Regulation CF being offered may not be transferred by any Investor of such Securities during the one-year holding period beginning when the Securities were issued, unless such Securities were transferred: 1) to the Company, 2) to an accredited investor, as defined by Rule 501(d) of Regulation D of the Securities Act of 1933, as amended, 3) as part of an Offering registered with the SEC or 4) to a member of the family of the Investor or the equivalent, to a trust controlled by the Investor, to a trust created for the benefit of a family member of the Investor or the equivalent, or in connection with the death or divorce of the Investor or other similar circumstances. "Member of the family" as used herein means a child, stepchild, grandchild, parent, stepparent, grandparent, spouse or spousal equivalent, sibling, mother/father/daughter/son/sister/brother-in-law, and includes adoptive relationships. Remember that although you may legally be able to transfer the Securities, you may not be able to find another party willing to purchase them.

TRANSACTIONS WITH RELATED PERSONS AND CONFLICTS OF INTEREST

Related Person Transactions

From time to time the Company may engage in transactions with related persons. Related persons are defined as any director or officer of the Company; any person who is the beneficial owner of 10 percent or more of the Company's outstanding voting equity securities, calculated on the basis of voting power; any promoter of the Company; any immediate family member of any of the foregoing persons or an entity controlled by any such person or persons.

The Company has the following transactions with related persons:

Loans

Related Person/Entity	Daniel Uribe
Relationship to the Company	CEO
Total amount of money involved	$1,619.00
Benefits or compensation received by related person	none.
Benefits or compensation received by Company	none.
Description of the transaction	As of December 31, 2019, the company has a receivable from its founder and CEO Daniel Uribe Benitez

Conflicts of Interest

To the best of our knowledge the Company has not engaged in any transactions or relationships, which may give rise to a conflict of interest with the Company, its operations or its security holders.

OTHER INFORMATION

The Company has not failed to comply with the ongoing reporting requirements of Regulation CF § 227.202 in the past.

Bad Actor Disclosure

The Company is not subject to any Bad Actor Disqualifications under any relevant U.S. securities laws.

SIGNATURE

Pursuant to the requirements of Sections 4(a)(6) and 4A of the Securities Act of 1933 and Regulation Crowdfunding (§ 227.100 et seq.), the issuer certifies that it has reasonable grounds to believe that it meets all of the requirements for filing on Form C-AR and has duly caused this Form to be signed on its behalf by the duly authorized undersigned.

The issuer also certifies that the attached financial statements are true and complete in all material respects.

/s/Daniel Uribe

(Signature)

Daniel Uribe

(Name)

CEO, President, Secretary

(Title)

Pursuant to the requirements of Sections 4(a)(6) and 4A of the Securities Act of 1933 and Regulation Crowdfunding (§ 227.100 et seq.), this Form C-AR has been signed by the following persons in the capacities and on the dates indicated.

/s/Daniel Uribe

(Signature)

Daniel Uribe

(Name)

Chief Executive Officer, President, Secretary

(Title)

04-30-2021

(Date)

/s/Candy Bellau

(Signature)

Candy Bellau

(Name)

CFO, Treasurer

(Title)

04-30-2021

(Date)

Instructions.

1. The form shall be signed by the issuer, its principal executive officer or officers, its principal financial officer, its controller or principal accounting officer and at least a majority of the board of directors or persons performing similar functions.

2. The name of each person signing the form shall be typed or printed beneath the signature.

Intentional misstatements or omissions of facts constitute federal criminal violations. See 18 U.S.C. 1001.

EXHIBITS

Exhibit A Financial Statements

EXHIBIT A

Financial Statements

GENOBANK. IO INC.

FINANCIAL STATEMENTS AND INDEPENDENT ACCOUNTANT'S AUDIT REPORT
YEAR ENDED DECEMBER 31, 2019 AND 2018

(Expressed in United States Dollars)

INDEX TO FINANCIAL STATEMENTS

(UNAUDITED)

INDEPENDENT ACCOUNTANT'S REVIEW REPORT

To the Board of Directors of
Genobank.io Inc.
San Jose, California.

We have audited the accompanying financial statements of Genobank.io Inc. (the "Company"), which comprise the balance sheets as of December 31, 2019 and December 31, 2018, and the related statement of operations, statement of shareholders' equity (deficit), and cash flows for the years ending December 31, 2019 and December 31, 2018, and the related notes to the financial statements.

Management's Responsibility for the Financial Statements

Management is responsible for the preparation and fair presentation of these financial statements in accordance with accounting principles generally accepted in the United States of America; this includes the design, implementation, and maintenance of internal control relevant to the preparation and fair presentation of financial statements that are free from material misstatement, whether due to fraud or error.

Accountant's Responsibility

Our responsibility is to express an opinion on these financial statements based on our audit. We conducted our audit in accordance with auditing standards generally accepted in the United States of America. Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free from material misstatement.

An audit involves performing procedures to obtain audit evidence about the amounts and disclosures in the financial statements. The procedures selected depend on the auditor's judgment, including the assessment of the risks of material misstatement of the financial statements, whether due to fraud or error. In making those risk assessments, the auditor considers internal control relevant to the Company's preparation and fair presentation of the financial statements in order to design audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the Company's internal control. Accordingly, we express no such opinion. An audit also includes evaluating the appropriateness of accounting policies used and the reasonableness of significant accounting estimates made by management, as well as evaluating the overall presentation of the combined financial statements.

We believe that the audit evidence we have obtained is sufficient and appropriate to provide a basis for our audit opinion.

Opinion

In our opinion, the financial statements referred to above present fairly, in all material respects, the financial position of Genobank.io, Inc. as of December 31, 2019 and December 31, 2018, and the results of its operations and its cash flows for the years ended December 31, 2019 and December 31, 2018 in accordance with accounting principles generally accepted in the United States of America.

Going Concern

As discussed in Note 9, certain conditions indicate that the Company may be unable to continue as a going concern. The accompanying financial statements do not include any adjustments that might be necessary should the Company be unable to continue as a going concern.

SetApart FS

November 13, 2020
Los Angeles, California

GENOBANK.IO INC.
BALANCE SHEET

As of December 31,	2019	2018
(USD $ in Dollars)		
ASSETS		
Current Assets:		
Cash & cash equivalents	$ 1,027	$ 36
Other current assets	1,619	-
Total current assets	**2,646**	**36**
Non current assets		
Property and equipment, net	**3,832**	**-**
Total assets	**$ 6,478**	**$ 36**
LIABILITIES AND STOCKHOLDERS' EQUITY		
Non Current Liabilities:		
SAFEs note	165,000	-
Total liabilities	**165,000**	**-**
STOCKHOLDERS' EQUITY		
Common stock	71	71
Subscription receivable	(71)	(71)
Additional paid in capital	50	50
Retained earnings/(Accumulated Deficit)	(158,572)	(14)
Total Stockholders' equity	**(158,522)**	**36**
Total liabilities and stockholders' equity	**$ 6,478**	**$ 36**

See accompanying notes to financial statements.

GENOBANK.IO INC.

STATEMENTS OF OPERATIONS

For Fiscal Year Ended December 31,	2019	2018
(USD $ in Dollars)		
Net revenue	$ -	$ -
Cost of goods sold	-	-
Gross profit	-	-
Operating expenses		
General and administrative	94,143	14
Research and development	56,312	-
Sales and marketing	8,103	-
Total operating expenses	158,558	14
Operating income/(loss)	(158,558)	(14)
Interest expense	-	-
Income/(Loss) before provision for income taxes	(158,558)	(14)
Provision/(Benefit) for income taxes	-	-
Net income/(Net Loss)	**$ (158,558)**	**$ (14)**

See accompanying notes to financial statements.

GENOBANK.IO INC.

STATEMENTS OF CHANGES IN STOCKHOLDERS' EQUITY

(in , $US)	Common StocK Shares	Common StocK Amount	Subscription Receivable	Additional Paid In Capital	Accumulated Deficit	Total Shareholder Equity
Inception (May 29, 2019)	-	-	-	-	-	-
Net income/(loss)	-	-	-	-	(14)	(14)
Issuance of Common Stock	7,100,000	71	(71)	50	-	50
Contribution	-	-	-	-	-	
Balance—December 31, 2018	**7,100,000**	71	**$ (71)**	**$ 50**	**$ (14)**	36
Net income/(loss)	-	-	-	-	(158,558)	(158,558)
Balance—December 31, 2019	**7,100,000**	**$ 71**	**$ (71)**	**$ 50**	**$ (158,572)**	**$ (158,522)**

See accompanying notes to financial statements.

GENOBANK.IO INC.

STATEMENTS OF CASH FLOWS

For Fiscal Year Ended December 31,	2019	2018
(USD $ in Dollars)		
CASH FLOW FROM OPERATING ACTIVITIES		
Net income/(loss)	$ (158,558)	$ (14)
Depreciation	14	
Adjustments to reconcile net income to net cash provided/(used) by operating activities:		
Changes in operating assets and liabilities:		
Other current assets	(1,619)	
Net cash provided/(used) by operating activities	**(160,163)**	**(14)**
CASH FLOW FROM INVESTING ACTIVITIES		
Purchase of property and equipment	(3,846)	-
Net cash provided/(used) by investing activities	**(3,846)**	**-**
CASH FLOW FROM FINANCING ACTIVITIES		
Shareholder contribution	-	50
Issuance of SAFE notes	165,000	-
Net cash provided/(used) by financing activities	**165,000**	**50**
Change in cash, cash equvalents and restricted cash	991	36
Cash, cash equivalents and restricted cash —beginning of year	36	-
Cash, cash equivalents and restricted cash —end of year	**$ 1,027**	**$ 36**
SUPPLEMENTAL DISCLOSURE OF CASH FLOW INFORMATION		
Cash paid during the year for interest	$ -	$ -
Cash paid during the year for income taxes	$ -	$ -
OTHER NONCASH INVESTING AND FINANCING ACTIVITIES AND SUPPLEMENTAL DISCLOSURES		
Purchase of property and equipment not yet paid for	$ -	$ -
Conversion of debt into equity	$ -	$ -

See accompanying notes to financial statements.

All amounts in these Notes are expressed in thousands of United States dollars ("$" or "US$"), unless otherwise indicated.

1. SUMMARY

Genobank.io Inc., was formed on May 24, 2018 ("Inception") in the State of Delaware. The financial statements of Genobank.io Inc., (which may be referred to as the "Company", "we," "us," or "our") are prepared in accordance with accounting principles generally accepted in the United States of America ("U.S. GAAP"). The Company's headquarters are located in San Francisco, CA.

Genobank.io Inc., is a decentralized platform that enables users to establish ownership, store, process & control of their Genomic data by using our Blockchain enabled and patent pending "In Crypto Saliva DNA extraction Kit". After receiving your saliva sample stored our FDA approved kit, we'll process it in a CLIA certified Lab within US to obtain your DNA data and deposit the resulting files into a personal, safe data container that will be encrypted using your private key, so only the owner of that secret can read the data.

2. SIGNIFICANT ACCOUNTING POLICIES

Basis of Presentation

The accounting and reporting policies of the Company conform to accounting principles generally accepted in the United States of America ("US GAAP").

Use of Estimates

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

Cash and Cash Equivalents

Cash and cash equivalents include all cash in banks. The Company's cash are deposited in demand accounts at financial institutions that management believes are creditworthy.

Restricted cash

Restricted cash is that portion of cash that is set aside for a specific purpose and is not available for general business use on an immediate basis. This cash is held on a special account (escrow account) and it pertains to fundraising from crowdfunding campaign, and it remain separate from the rest of the company's cash and cash equivalent.

Income Taxes

Genobank.io. Inc. is a C corporation for income tax purposes. The Company accounts for income taxes under the liability method, and deferred tax assets and liabilities are recognized for the future tax consequences attributable to differences between the financial statement carrying values of existing assets and liabilities and their respective tax bases. Deferred tax assets and liabilities are measured using enacted tax rates in effect for the year in which those temporary differences are expected to be recovered or settled. A valuation allowance is provided on deferred tax assets if it is determined that it is more likely than not that the deferred tax asset will not be realized. The Company records interest, net of any applicable related income tax benefit, on potential income tax contingencies as a component of income tax expense. The Company records tax positions taken or expected to be taken in a tax return based upon the amount that is more likely than not to be realized or paid, including in connection with the resolution of any related appeals or other legal processes. Accordingly, the Company recognizes liabilities for certain unrecognized tax benefits based on the amounts that are more likely than not to be settled with the relevant taxing authority. The Company recognizes interest and/or penalties related to unrecognized tax benefits as a component of income tax expense.

Software Development Costs

The Company recognized software development costs based on the guidance of Accounting Standards Codification (ASC) 985, *Software.* The software has not reached technological feasibility, and therefore, the Company has expensed all costs incurred to date.

Concentration of Credit Risk

The Company maintains its cash with a major financial institution located in the United States of America which it believes to be creditworthy. Balances are insured by the Federal Deposit Insurance Corporation up to $250,000. At times, the Company may maintain balances in excess of the federally insured limits.

Revenue Recognition

The Company will recognize revenues primarily from sale of our products when (a) persuasive evidence that an agreement exists; (b) the service has been performed; (c) the prices are fixed and determinable and not subject to refund or adjustment; and (d) collection of the amounts due is reasonably assured.

Operating Leases

Operating leases relate to office space. Rent expense for operating leases is recognized on a straight-line basis over the term of the lease.

Fair Value of Financial Instruments

The carrying value of the Company's financial instruments included in current assets and current liabilities (such as cash and cash equivalents, restricted cash and cash equivalents, accounts receivable, accounts payable and accrued expenses approximate fair value due to the short-term nature of such instruments.

The inputs used to measure fair value are based on a hierarchy that prioritizes observable and unobservable inputs used in valuation techniques. These levels, in order of highest to lowest priority, are described below:

Level 1—Quoted prices (unadjusted) in active markets that are accessible at the measurement date for identical assets or liabilities.

Level 2—Observable prices that are based on inputs not quoted on active markets, but corroborated by market data.

Level 3—Unobservable inputs reflecting the Company's assumptions, consistent with reasonably available assumptions made by other market participants. These valuations require significant judgment.

Subsequent Events

The Company considers events or transactions that occur after the balance sheets date, but prior to the issuance of the financial statements to provide additional evidence relative to certain estimates or to identify matters that require additional disclosure. Subsequent events have been evaluated through November 13, 2020, which is the date the financial statements were issued.

Recently Issued and Adopted Accounting Pronouncements

In February 2019, FASB issued ASU No. 2019-02, Leases, that requires organizations that lease assets, referred to as "lessees", to recognize on the balance sheet the assets and liabilities for the rights and obligations created by those leases with lease terms of more than 12 months. ASU 2019-02 will also require disclosures to help investors and other financial statement users better understand the amount, timing, and uncertainty of cash flows arising from leases and will include qualitative and quantitative requirements. The new standard for nonpublic entities will be effective for fiscal years beginning after December 15, 2019, and interim periods within fiscal years beginning after December 15, 2020, and early application is permitted. We are currently evaluating the effect that the updated standard will have on the financial statements and related disclosures.

In June 2019, FASB amended ASU No. 2019-07, Compensation – Stock Compensation, to expand the scope of Topic 718, Compensation – Stock Compensation, to include share-based payment transactions for acquiring goods and services from nonemployees. The new standard for nonpublic entities will be effective for fiscal years beginning after December 15, 2019, and interim periods within fiscal years beginning after December 15, 2020, and early application is permitted. We are currently evaluating the effect that the updated standard will have on the financial statements and related disclosures.

In August 2019, amendments to existing accounting guidance were issued through Accounting Standards Update 2019-15 to clarify the accounting for implementation costs for cloud computing arrangements. The amendments specify that existing guidance for capitalizing implementation costs incurred to develop or obtain internal-use software also applies to implementation costs incurred in a hosting arrangement that is a service contract. The guidance is effective for fiscal years beginning after December 15, 2020, and interim periods within fiscal years beginning after December 15, 2021, and early application is permitted. We are currently evaluating the effect that the updated standard will have on the financial statements and related disclosures.

The FASB issues ASUs to amend the authoritative literature in ASC. There have been a number of ASUs to date, including those above, that amend the original text of ASC. Management believes that those issued to date either (i)

provide supplemental guidance, (ii) are technical corrections, (iii) are not applicable to us or (iv) are not expected to have a significant impact our financial statements.

3. DETAILS OF CERTAIN ASSETS AND LIABILITIES

Other current liabilities consist of the following items:

As of Year Ended December 31,	2019	2018
Other Current Assets consist of:		
Due from Daniel Uribe Benitez	$ 1,619	$ -
Total Other Current Assets	**$ 1,619**	**$ -**

4. CAPITALIZATION AND EQUITY TRANSACTIONS

Common Stock

We have authorized the issuance of 10,000,000, shares of our common stock with par value of $0.00001. As of December 31, 2019, 7,100,000 shares of common stock are issued and outstanding for a consideration of $71.

DEBT

Future Equity Obligations (SAFEs)

During 2019 , the Company entered into four Simple Agreements for Future Equity ("SAFE") for an aggregate purchase amount of $165,000.

The SAFE notes carry different discount rates. The following is the breakout:

As of December 31, 2019

Insturment	Amount	Valuation Cap	Discount
SAFE	$ 50,000	N/A	60%
SAFE	10,000	N/A	70%
SAFE	85,000	N/A	75%
SAFE	20,000	N/A	80%
Total	**$ 165,000**		

If there is an Equity Financing before the expiration or termination of this instrument, the Company will automatically issue to the Investor a number of shares of Safe Preferred Stock equal to the Purchase Amount divided by the Discount Price.

If there is a Liquidity Event before the termination of this Safe, the company will automatically issue to the investor a number of Safe Preferred Stock equal to the purchase amount divided by the discounted price. As of December 31, 2019, the SAFEs have not yet converted to equity. The notes have been classified as non-current liability as the Company might settle them in cash in certain cases.

If there is a Dissolution Event before this instrument expires or terminates, the Company will pay an amount equal to the Purchase Amount, due and payable to the Investor immediately prior to, or concurrent with, the consummation of the Dissolution Event.

As of December 31, 2019, the SAFE has not yet converted to equity. The entire notes have been classified as non-current liability as the Company might settle them in cash in certain cases.

5. INCOME TAXES

The provision for income taxes for the year ended December 31, 2019 and December 31, 2018 consists of the following:

As of Year Ended December 31,	2019	2018
Net Operating Loss	$ (44,789)	$ (4)
Valuation Allowance	44,789	4
Net Provision for income tax	$ -	$ -

Significant components of the Company's deferred tax assets and liabilities at December 31, 2019, and December 31, 2018 are as follows:

As of Year Ended December 31,	2019	2018
Net Operating Loss	$ (44,797)	$ (4)
Valuation Allowance	44,797	4
Total Deferred Tax Asset	$ -	$ -

Management assesses the available positive and negative evidence to estimate if sufficient future taxable income will be generated to use the existing deferred tax assets. On the basis of this evaluation, the Company has determined that it is more likely than not that the Company will not recognize the benefits of the federal and state net deferred tax assets, and, as a result, full valuation allowance has been set against its net deferred tax assets as of December 31, 2019. The amount of the deferred tax asset to be realized could be adjusted if estimates of future taxable income during the carryforward period are reduced or increased.

As of December 31, 2019, the Company had net operating loss ("NOL") carryforwards of approximately $158,572. Utilization of some of the federal and state NOL carryforwards to reduce future income taxes will depend on the Company's ability to generate sufficient taxable income prior to the expiration of the carryforwards. Under the provisions of the Internal Revenue Code, the NOLs and tax credit carryforwards are subject to review and possible

adjustment by the IRS and state tax authorities. NOLs and tax credit carryforwards may become subject to an annual limitation in the event of certain cumulative changes in the ownership interest of significant stockholders over a three-year period in excess of 50%, as defined under Sections 382 and 383 of the Internal Revenue Code, as well as similar state provisions. This could limit the amount of tax attributes that can be utilized annually to offset future taxable income or tax liabilities. The amount of the annual limitation is determined based on the value of the Company immediately prior to the ownership change. The Company has not performed a comprehensive Section 382 study to determine any potential loss limitation with regard to the NOL carryforwards and tax credits.

The Company recognizes the impact of a tax position in the financial statements if that position is more likely than not of being sustained on a tax return upon examination by the relevant taxing authority, based on the technical merits of the position. As of December 31, 2019, and December 31, 2018 the Company had no unrecognized tax benefits.

The Company recognizes interest and penalties related to income tax matters in income tax expense. As of December 31, 2019, and December 31, 2018 the Company had no accrued interest and penalties related to uncertain tax positions.

The Company is subject to examination for its US federal and California jurisdictions for each year in which a tax return was filed.

6. RELATED PARTY

As of December 31, 2019, the company has a receivable from its founder and CEO Daniel Uribe Benitez in the amount of $1,619.

7. COMMITMENTS AND CONTINGENCIES

Contingencies

The Company's operations are subject to a variety of local and state regulation. Failure to comply with one or more of those regulations could result in fines, restrictions on its operations, or losses of permits that could result in the Company ceasing operations.

Operating lease

During 2019, the company entered into a 6 months rental contract with CColab44 America Inc. for shared workplace for $333 per month. In 2019, rental expense incurred was in the amount of $ 1,310.

During 2019, the company entered into a month to month rental contract with Ace Parking for parking lots. In 2019, rental expenses incurred in the amount of $ 840.

Litigation and Claims

From time to time, the Company may be involved in litigation relating to claims arising out of operations in the normal course of business. As of December 31, 2019, there were no pending or threatened lawsuits that could reasonably be expected to have a material effect on the results of the Company's operations.

8. SUBSEQUENT EVENTS

The Company has evaluated subsequent events through November 13, 2020, the date the financial statements were available to be issued.

On April 21, 2020, the Company received an equity free $50,000 grant in order to incentivize the use of EOSIO Blockchain Technology in the next 6 to 12 months with no ownership conditions.

On July 31, 2020, the Company completed up to $1,070,000 of (the "Crowdfunded Offering") in Simple Agreements for Future Equity (SAFEs) through OpenDeal Portal LLC (the "Intermediary" aka "Republic" or "Republic.co"). The Intermediary will be entitled to receive a 6% commission fee and 2% of the securities issued in this offering. The company received a total disbursement of $423,787 excluding commission and fees in the amount of $40,643.

On August 7, 2020, the Company entered a convertible promissory note agreement with a certain investor in the amount of $25,000. The note carries an interest rate of 2% and matures after 24 months from the issuance date.

There have been no other events or transactions during this time which would have a material effect on these financial statements.

9. GOING CONCERN

The accompanying financial statements have been prepared on a going concern basis, which contemplates the realization of assets and the satisfaction of liabilities in the normal course of business. The Company has a net operating loss of $158,558, an operating cash flow loss of $160,163 and an accumulated deficit of $158,522 as of December 31, 2019. The Company's situation raises a substantial doubt on whether the entity can continue as a going concern in the next twelve months.

The Company's ability to continue as a going concern in the next twelve months following the date the financial statements were available to be issued is dependent upon its ability to produce revenues and/or obtain financing sufficient to meet current and future obligations and deploy such to produce profitable operating results.

Management has evaluated these conditions and plans to generate revenues and raise capital as needed to satisfy its capital needs. During the next twelve months, the Company intends to fund its operations through debt and/or equity financing.

There are no assurances that management will be able to raise capital on terms acceptable to the Company. If it is unable to obtain sufficient amounts of additional capital, it may be required to reduce the scope of its planned development, which could harm its business, financial condition, and operating results. The accompanying financial statements do not include any adjustments that might result from these uncertainties.