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
Green Valley Affiliates, Inc.

Legal status of issuer

Form
Corporation

Jurisdiction of Incorporation/Organization
Nevada

Date of organization
February 8, 2018

Physical address of issuer
2764 N. Green Valley Parkway, Suite #345, Henderson, NV 89014

Website of issuer
www.cannabiscope.com

Current number of employees
0

	Most recent fiscal year-end	Prior fiscal year-end
Total Assets	$157,923.00	$6,372.00
Cash & Cash Equivalents	$137,923.00	$6,372.00
Accounts Receivable	$0.00	$0.00
Short-term Debt	$36,575.00 *	$0.00
Long-term Debt	$138,928.00	$34,420.00
Revenues/Sales	$3,125.00	$12,375.00
Cost of Goods Sold	$0.00	$5,811.00
Taxes Paid	$0.00	$0.00
Net Income	-$27,493.00	-$18,491.00

* This debt is disclosed as Note D under the attached Financial Statements. The debt was paid back in full in January 2020.

April 20, 2020

FORM C-AR

Green Valley Affiliates, Inc.

This Form C-AR (including the cover page and all exhibits attached hereto, the "Form C-AR) is being furnished by Green Valley Affiliates, Inc., a Nevada 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 www.cannabiscope.com 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 April 20, 2020.

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.

Green Valley Affiliates, Inc. (the "Company") is a Nevada Corporation incorporated on August 27, 2019, upon its conversion from a Nevada limited liability company. The Company was originally formed on February 8, 2018, as Green Valley Affiliates, LLC. The Company is currently also conducting business under the name of Cannabiscope.

The Company is located at 2764 N. Green Valley Parkway, Suite #345, Henderson, NV 89014.

The Company's website is www.cannabiscope.com.

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

The Business

Cannabiscope is a SaaS menu platform that integrates with THC/CBD retailers POS systems to present their inventory live onsite and an e-commerce site, while collecting industry data. Customers sign annual contracts and either pay a set monthly fee or give us 1% of their total revenue, whichever is higher for us to provide this service.

RISK FACTORS

Risks Related to the Company's Business and Industry

The development and commercialization of our products and 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 and services and thus may be better equipped than us to develop and commercialize products and 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 and 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.

As a distributor of a SaaS platform for E-Commerce of CBD Products, our business depends on developing and maintaining close and productive relationships with our vendors.

We depend on our vendors to sell us quality products at favorable prices. Many factors outside our control, including, without limitation, raw material shortages, inadequate manufacturing capacity, labor disputes, transportation disruptions or weather conditions, could adversely affect our vendors' ability to deliver to us quality merchandise at favorable prices in a timely manner. Furthermore, financial or operational difficulties with a particular vendor could cause that vendor to increase the cost of the products or decrease the quality of the products we purchase from it. Vendor consolidation could also limit the number of suppliers from which we may purchase products and could materially affect the prices we pay for these products. We would suffer an adverse impact if our vendors limit or cancel the return privileges that currently protect us from inventory obsolescence.

We 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.

Through our operations, we collect and store certain personal information that our customers provide to purchase products or services, enroll in promotional programs, register on our web site, or otherwise communicate and interact with us.

We may share information about such persons with vendors that assist with certain aspects of our business. Security could be compromised and confidential customer or business information misappropriated. Loss of customer or business information could disrupt our operations, damage our reputation, and expose us to claims from customers, financial institutions, payment card associations and other persons, any of which could have an adverse effect on our business, financial condition and results of operations. In addition, compliance with tougher privacy and information security laws and standards may result in significant expense due to increased investment in technology and the development of new operational processes.

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 following best practices of Data Protection. 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 David Schacter who is CEO of the Company. The Company has or intends to enter into employment agreements with David Schacter 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 Paul Shockley 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 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.

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 Paul Shockley in order to conduct its operations and execute its business plan, however, the Company has not purchased any insurance policies with respect to this individual in the event of their death or disability. Therefore, if Paul Shockley dies or becomes 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.

Medical-use cannabis remains illegal under federal law, and therefore, strict enforcement of federal laws regarding medical-use cannabis would prevent us from executing our business plan.

Cannabis is a Schedule I controlled substance under the Controlled Substance Act ("CSA"). Even in those jurisdictions in which the manufacture and use of medical cannabis has been legalized at the state level, the possession, use and cultivation all remain violations of federal law that are punishable by imprisonment and substantial fines. Moreover, individuals and entities may violate federal law if they intentionally aid and abet another in violating these federal controlled substance laws or conspire with another to violate them. In United States v. Oakland Cannabis Buyers' Cooperative and Gonzales v. Raich, the U.S. Supreme Court ruled that the federal government has the right to regulate and criminalize cannabis, even for medical purposes. We would be unable to execute our business plan if the federal government were to strictly enforce federal law regarding cannabis.

The U.S. Department of Justice, under the Obama administration, issued memoranda, including the so-called "Cole Memo" on August 29, 2013, characterizing enforcement of federal cannabis prohibitions under the CSA to prosecute those complying with state regulatory systems allowing the use, manufacture and distribution of medical cannabis as an inefficient use of federal investigative and prosecutorial resources when state regulatory and enforcement efforts are effective with respect to enumerated federal enforcement priorities under the CSA. In the Cole Memo, the U.S. Department of Justice provided guidance to all federal prosecutors indicating that federal enforcement of the CSA against cannabis-related conduct should be focused on eight priorities, which are to prevent: (1) distribution of cannabis to minors; (2) revenue from sale of cannabis to criminal enterprises, gangs and cartels; (3) transfer of cannabis from states where it is legal to states where it is illegal; (4) cannabis activity from being a pretext for trafficking of other illegal drugs or illegal activity; (5) violence or use of firearms in cannabis cultivation and distribution; (6) drugged driving and adverse public health consequences from cannabis use; (7) growth of cannabis on federal lands; and (8) cannabis possession or use on federal property.

In addition, Congress enacted an omnibus spending bill for fiscal year 2016 including a provision prohibiting the U.S. Department of Justice (which includes the DEA) from using funds appropriated by that bill to prevent states from implementing their medical-use cannabis laws. This provision, however, is effective only until April 28, 2017 and must be renewed by Congress. In United States vs. McIntosh, the United States Court of Appeals for the Ninth Circuit held that this provision prohibits the U.S. Department of Justice from spending funds from relevant appropriations acts to prosecute individuals who engage in conduct permitted by state medical-use cannabis laws and who strictly comply with such laws. However, the Ninth Circuit's opinion, which only applies to the states of Alaska, Arizona, California, Hawaii, and Idaho, also held that persons who do not strictly comply with all state laws and regulations regarding the distribution, possession and cultivation of medical-use cannabis have engaged in conduct that is unauthorized, and in such instances the U.S. Department of Justice may prosecute those individuals.

On January 4, 2018, Attorney General Jeff Sessions issued a memorandum to all Unites States Attorneys concerning marijuana enforcement, stating "Given the Department's well-established general principles, previous nationwide guidance specific to marijuana enforcement including the Cole Memo is unnecessary and is rescinded, effective immediately". However, Donald Trump subsequently signaled that he would support congressional efforts to protect those states that had legalized marijuana. Sessions resigned as Attorney General on November 7, 2018.

In April 2019, the Strengthening the Tenth Amendment Through Entrusting States ("STATES") Act was reintroduced in the United States Senate by a bipartisan group of lawmakers. The STATES Act would, in part, amend the federal Controlled Substances Act such that its provisions would not apply to any individual acting in accordance with state or tribal marijuana laws.

Furthermore, financial transactions involving proceeds generated by cannabis-related conduct can form the basis for prosecution under the federal money laundering statutes, unlicensed money transmitter statutes and the Bank Secrecy Act. However, supplemental guidance from the U.S. Department of Justice directed federal prosecutors to consider the federal enforcement priorities enumerated in the "Cole Memo" when determining whether to charge institutions or individuals with any of the financial crimes described above based upon cannabis-related activity. Accordingly, the Treasury Department's Financial Crimes Enforcement Network ("FinCEN") established suspicious activity reporting guidelines for those banks serving marijuana related businesses that complemented the Cole Memo's eight priorities.

Sessions' rescission of the Cole Memo has not resulted in an increase in financial crimes enforcement against cannabis-related businesses, likely because FinCEN did not rescind its marijuana banking guidance.

Federal prosecutors have significant discretion, and there is no guarantee that the federal prosecutor in those judicial districts in which we conduct business will not choose to strictly enforce federal laws governing cannabis production or distribution. At this time, it is unknown if the STATES Act will become law or if the Trump administration will change the federal government's current enforcement posture with respect to state-licensed medical-use cannabis. Any such change in the federal government's current enforcement posture with respect to state-licensed cultivation of medical-use cannabis would result in our inability to execute our business plan and we would suffer significant losses and be required to cease operations.

Any changes in state or local laws that reduce or eliminate the ability to cultivate and produce medical-use cannabis would have a material negative impact on our business.

The SEC is monitoring the cannabis industry and may halt or prevent the Offering or sale of our securities due to the bad acts of others.

On May 16, 2014 and again on September 5, 2018, the SEC's Office of Investor Education and Advocacy issued an Investor Alert to warn investors about potential risks involving investments in marijuana-related companies. The SEC has noted an increase in the number of investor complaints regarding marijuana-related investments and has issued temporary trading suspensions for the common stock of various different marijuana-related companies. Due to the stigma created by the bad acts of others in the industry, the SEC may halt trading and offerings in all marijuana-related companies which would have a material adverse effect on our ability to raise capital and our business.

Our ability to grow our business depends on state laws pertaining to the cannabis industry. Continued development of the medical-use cannabis industry depends upon continued legislative authorization of cannabis at the state level. The status quo of, or progress in, the regulated medical-use cannabis industry is not assured and any number of factors could slow or halt further progress in this area. While there may be ample public support for legislative action permitting the manufacture and use of cannabis, numerous factors impact the legislative process. For example, states that voted to legalize medical and/or adult-use cannabis in the November 2016 election cycle have seen significant delays in the drafting and implementation of regulations related to the industry. In addition, burdensome regulation at the state level could slow or stop further development of the medical-use cannabis industry, such as limiting the medical conditions for which medical cannabis can be recommended by physicians for treatment, restricting the form in which medical cannabis can be consumed, imposing significant registration requirements on physicians and patients or imposing significant taxes on the growth, processing and/or retail sales of cannabis, which could have the impact of dampening growth of the cannabis industry and making it difficult for cannabis businesses to operate profitably in those states.

FDA regulation of medical-use cannabis and the possible registration of facilities where medical-use cannabis is grown could negatively affect the medical-use cannabis industry and our financial condition.

Should the federal government legalize cannabis for medical use, it is possible that the U.S. Food and Drug Administration, or the FDA, would seek to regulate it under the Food, Drug and Cosmetics Act of 1938. Additionally, the FDA may issue rules and regulations including certified good manufacturing practices, or cGMPs, related to the growth, cultivation, harvesting and processing of medical cannabis. Clinical trials may be needed to verify efficacy and safety. It is also possible that the FDA would require that facilities where medical-use cannabis is grown register with the FDA and comply with certain federally prescribed regulations. In the event that some or all of these regulations are imposed, we do not know what the impact would be on the medical-use cannabis industry, including what costs, requirements and possible prohibitions may be enforced. If we are unable to comply with the regulations or registration as prescribed by the FDA, we may be unable to continue to operate.

We may have difficulty accessing the service of banks, which may make it difficult to contract for real estate needs.

Financial transactions involving proceeds generated by cannabis-related conduct can form the basis for prosecution under the federal money laundering statutes, unlicensed money transmitter statute and the Bank Secrecy Act. Recent guidance issued by FinCen, a division of the U.S. Department of the Treasury, clarifies how financial institutions can provide services to cannabis-related businesses consistent with their obligations under the Bank Secrecy Act. Furthermore, supplemental guidance from the U.S. Department of Justice directs federal prosecutors to consider the federal enforcement priorities enumerated in the "Cole Memo" when determining whether to charge institutions or individuals with any of the financial crimes described above based upon cannabis-related activity. Nevertheless, banks remain hesitant to offer banking services to cannabis-related businesses. Consequently, those businesses involved in the regulated medical-use cannabis industry continue to encounter difficulty establishing banking relationships. Our inability to maintain our current bank accounts would make it difficult for us to operate our business, increase our operating costs, and pose additional operational, logistical and security challenges and could result in our inability to implement our business plan.

The Internal Revenue Code provides a higher tax rate for cannabis industry businesses.
Internal Revenue Code section 280E prohibits medical-marijuana businesses from deducting their ordinary and necessary business expenses, forcing them to contend with higher effective federal tax rates than similar companies in other industries. Our effective tax rate depends on how large our ratio of nondeductible expenses is to our total revenues, but it can be as high as 90%. This relatively higher tax rate will affect our future profitability and could cause us to perform worse than investments in different industries.

Investors risk criminal liability and the cannabis business's assets are subject to forfeiture.
Because marijuana is federally illegal, investing in cannabis businesses could be found to violate the CSA. Investors and company directors or management could be indicted under federal law, and all of the assets contributed to the Company, including real property, cash, equipment and other goods, could be subject to asset forfeiture.

The SEC is monitoring the cannabis industry and may halt or prevent the Offering or sale of our securities due to the bad acts of others.
On May 16, 2014, the SEC's Office of Investor Education and Advocacy issued an Investor Alert to warn investors about potential risks involving investments in marijuana-related companies. The SEC noted an increase in the number of investor complaints regarding marijuana-related investments. The SEC issued temporary trading suspensions for the common stock of five different marijuana-related companies. Due to the stigma created by the bad acts of others in the industry, the SEC may halt trading and offerings in all marijuana-related companies which would have a material adverse effect on our ability to raise capital and our business.

Laws and regulations affecting the regulated cannabis industry are constantly changing, which could materially adversely affect our proposed operations, and we cannot predict the impact that future regulations may have on us.
Local, state and federal cannabis laws and regulations are broad in scope and subject to evolving interpretations, which could require us to incur substantial costs associated with compliance or alter our business plan. In addition, violations of these laws, or allegations of such violations, could disrupt our business and result in a material adverse effect on our operations. It is also possible

that regulations may be enacted in the future that will be directly applicable to our proposed business. We cannot predict the nature of any future laws, regulations, interpretations or applications, nor can we determine what effect additional governmental regulations or administrative policies and procedures, when and if promulgated, could have on our business.

Applicable state laws may prevent us from maximizing our potential income.
Depending on the laws of each particular state, we may not be able to fully realize our potential to generate profit. Colorado and Washington have residency requirements for those directly involved in the medical-use cannabis industry, which may impede our ability to contract with cannabis businesses in those states. Furthermore, cities and counties are being given broad discretion to ban certain cannabis activities. Even if these activities are legal under state law, specific cities and counties may ban them.

Assets leased to cannabis businesses may be forfeited to the federal government.
Any assets used in conjunction with the violation of federal law are potentially subject to federal forfeiture, even in states where cannabis is legal. If the federal government decides to initiate forfeiture proceedings against cannabis businesses, we may lose all of our assets.

The lending industry is highly regulated and changes in regulations or in the way regulations are applied to our business could adversely affect our business.
The regulatory environment in which lending institutions operate has become increasingly complex, and following the financial crisis of 2008, supervisory efforts to enact and apply relevant laws, regulations and policies have become more intense. Changes in laws or regulations or the regulatory application or judicial interpretation of the laws and regulations applicable to us could adversely affect our ability to operate in the manner in which we currently conduct business or make it more difficult or costly for us to originate or otherwise make additional loans, or for us to collect payments on loans by subjecting us to additional licensing, registration and other regulatory requirements in the future or otherwise. For example, if our loans were determined for any reason not to be commercial loans, we would be subject to many additional requirements, and our fees and interest arrangements could be challenged by regulators or our customers. A material failure to comply with any such laws or regulations could result in regulatory actions, lawsuits and damage to our reputation, which could have a material adverse effect on our business and financial condition and our ability to originate and service loans and perform our obligations to investors and other constituents.

A proceeding relating to one or more allegations or findings of our violation of such laws could result in modifications in our methods of doing business that could impair our ability to collect payments on our loans or to acquire additional loans or could result in the requirement that we pay damages and/or cancel the balance or other amounts owing under loans associated with such violation.

Business development corporations are subject to the Investment Company Act of 1940.
The Investment Company Act of 1940, as amended (1940 Act), imposes numerous constraints on the operations of business development corporations (BDC's). For example, BDC's are required to invest at least 70% of their total assets in qualifying assets, primarily securities of "eligible portfolio companies" (as defined under the 1940 Act), cash, cash equivalents, U.S. government

securities, and other high quality debt investments that mature in one year or less. Our regulatory requirements may hinder our ability to take advantage of attractive investment opportunities and, as a result, achieve our investment objective. In addition, we rely upon several exemptive orders from the SEC permitting us to consolidate our financial reporting and operate our business as presently conducted. Our failure to satisfy the conditions set forth in those exemptive orders could result in our inability to rely upon such orders or to cause the SEC to revoke the orders which could result in material changes in our financial reporting or the way in which we conduct our business. Furthermore, any failure to comply with the requirements imposed on BDC's by the 1940 Act could have material adverse consequences to us or our investors, including possible enforcement action by the SEC and the possible loss of our ability to qualify as a RIC that is exempt from corporate-level income tax under the Code. If we do not remain a BDC, we might be regulated as a closed-end investment company under the 1940 Act, which would further significantly decrease our operating flexibility.

We are subject to the Dodd-Frank Wall Street Reform and Consumer Protection Act.
The Dodd-Frank Wall Street Reform and Consumer Protection Act, or the Dodd-Frank Act, was signed into law on July 21, 2010. The Dodd-Frank Act significantly changes federal financial services regulation and affects, among other things, the lending, deposit, investment, trading, and operating activities of financial institutions and their holding companies. The Dodd-Frank Act requires extensive rulemaking by various regulatory agencies. The Dodd-Frank Act rulemaking process is ongoing and any changes resulting from such process, as well as any other changes in the laws or regulations applicable to us more generally, may negatively impact the profitability of our business activities, require us to change certain of our business practices, materially affect our business model, limit the activities in which we may engage, affect retention of key personnel, require us to raise additional regulatory capital, increase the amount of liquid assets that we hold, otherwise affect our funding profile or expose us to additional costs (including increased compliance costs). Any such changes may also require us to invest significant management attention and resources to make any necessary changes and may adversely affect our ability to conduct our business as previously conducted or our results of operations or financial condition. As such, we cannot predict and may not be able to anticipate all the effects of the Dodd-Frank Act on our financial condition or operations.

We operate in a highly regulated environment, and if we are found to be in violation of any of the federal, state, or local laws or regulations applicable to us, our business could suffer.
We are also subject to a wide range of federal, state, and local laws and regulations, such as local licensing requirements, and retail financing, debt collection, consumer protection, environmental, health and safety, creditor, wage-hour, anti-discrimination, whistleblower and other employment practices laws and regulations and we expect these costs to increase going forward. The violation of these or future requirements or laws and regulations could result in administrative, civil, or criminal sanctions against us, which may include fines, a cease and desist order against the subject operations or even revocation or suspension of our license to operate the subject business. As a result, we have incurred and will continue to incur capital and operating expenditures and other costs to comply with these requirements and laws and regulations.

The collection, processing, storage, use and disclosure of personal data could give rise to liabilities as a result of governmental regulation, conflicting legal requirements or differing views of personal privacy rights.

We receive, collect, process, transmit, store and use a large volume of personally identifiable information and other sensitive data from customers and potential customers. There are federal, state and foreign laws regarding privacy, recording telephone calls and the storing, sharing, use, disclosure and protection of personally identifiable information and sensitive data. Specifically, personally identifiable information is increasingly subject to legislation and regulations to protect the privacy of personal information that is collected, processed and transmitted. Any violations of these laws and regulations may require us to change our business practices or operational structure, address legal claims and sustain monetary penalties and/or other harms to our business.

The regulatory framework for privacy issues in the United States and internationally is constantly evolving and is likely to remain uncertain for the foreseeable future. The interpretation and application of such laws is often uncertain, and such laws may be interpreted and applied in a manner inconsistent with our current policies and practices or require changes to the features of our platform. If either we or our third-party service providers are unable to address any privacy concerns, even if unfounded, or to comply with applicable laws and regulations, it could result in additional costs and liability, damage our reputation and harm our business.

The Consumer Financial Protection Bureau or CFPB is a new agency, and there continues to be uncertainty as to how the agency's actions or the actions of any other new agency could impact our business or that of our issuing banks.

The CFPB, which commenced operations in July 2011, has broad authority over the businesses in which we engage. This includes authority to write regulations under federal consumer financial protection laws, such as the Truth in Lending Act and the Equal Credit Opportunity Act, and to enforce those laws against and examine large financial institutions, such as our issuing banks, for compliance. The CFPB is authorized to prevent "unfair, deceptive or abusive acts or practices" through its regulatory, supervisory and enforcement authority. To assist in its enforcement, the CFPB maintains an online complaint system that allows consumers to log complaints with respect to various consumer finance products, including the loan products we facilitate. This system could inform future CFPB decisions with respect to its regulatory, enforcement or examination focus.

We are subject to the CFPB's jurisdiction, including its enforcement authority, as a SaaS E-Commerce Platform. The CFPB may request reports concerning our organization, business conduct, markets and activities. The CFPB may also conduct on-site examinations of our business on a periodic basis if the CFPB were to determine, through its complaint system, that we were engaging in activities that pose risks to consumers.

There continues to be uncertainty as to how the CFPB's strategies and priorities, including in both its examination and enforcement processes, will impact our businesses and our results of operations going forward. Our compliance costs and litigation exposure could increase materially if the CFPB or other regulators enact new regulations, change regulations that were previously adopted, modify, through supervision or enforcement, past regulatory guidance, or interpret existing regulations in a manner different or stricter than have been previously interpreted.

Our earnings may decrease because of changes in prevailing interest rates.

Our profitability is directly affected by changes in prevailing interest rates. The following are certain material risks we face related to changes in prevailing interest rates:

* an increase in prevailing interest rates could generate an increase in delinquency, default and foreclosure rates resulting in an increase in both operating expenses and interest expense and could cause a reduction in the value of our assets;

* an increase in prevailing interest rates could adversely affect our loan originations volume because refinancing an existing loan would be less attractive for homeowners and qualifying for a purchase money loan may be more difficult for consumers;

* an increase in prevailing interest rates would increase the cost of servicing our outstanding debt, including our ability to finance servicing advances and loan originations;

* a decrease in prevailing interest rates may increase prepayment speeds causing our servicing fees to decline more rapidly than anticipated and we may record a decrease in the value of our MSRs;

* a decrease in prevailing interest rates may lead to higher compensating interest expense and increased amortization expense as we revise downward our estimate of total expected income as prepayment speeds increase; and a decrease in prevailing interest rates could reduce our earnings from our custodial deposit accounts.

Fluctuations in interest rates could negatively affect transaction volume.
All personal and nearly all small business loans facilitated through our marketplace are issued with fixed interest rates, and education and patient finance loans facilitated by our platform are issued with fixed or variable rates, depending on the type of loan. If interest rates rise, investors who have already committed capital may lose the opportunity to take advantage of the higher rates. Additionally, potential borrowers could seek to defer loans as they wait for interest rates to settle, and borrowers of variable rate loans through may be subject to increased interest rates. If interest rates decrease after a loan is made, borrowers through our marketplace may prepay their loans to take advantage of the lower rates. Investors through our marketplace would lose the opportunity to collect the above-market interest rate payable on the corresponding loan and may delay or reduce future loan investments. As a result, fluctuations in the interest rate environment may discourage investors and borrowers from participating in our marketplace and may reduce our loan originations, which may adversely affect our business.

As a registered investment advisor, our failure to comply with investment guidelines set by our clients or the provisions of the management agreement and other agreements to which we are a party could result in damage awards against us and a loss of assets under management (AUM), either of which could have an adverse effect on us.
As an investment adviser, each adviser has a fiduciary duty to its clients. When clients retain an adviser to manage assets on their behalf, they may specify certain guidelines regarding investment allocation and strategy that such adviser is required to observe in the management of its portfolios. In addition, such adviser is required to comply with the obligations set forth in the management agreements and other agreements to which it is a party. Although each adviser utilizes procedures, processes and the services of experienced advisors to assist it in adhering to these guidelines and agreements, we cannot assure you that such precautions will protect us from potential liabilities. An adviser's failure to comply with these guidelines or the terms of these agreements could have a material adverse effect on us.

Negative public opinion could damage our reputation and adversely affect our business.
Reputation risk, or the risk to our business from negative public opinion, is inherent in our business. Negative public opinion can result from our actual or alleged conduct in any number of activities, including lending and debt collection practices, corporate governance, and actions taken by government regulators and community organizations in response to those activities. Negative public opinion can also result from media coverage, whether accurate or not. Negative public opinion can adversely affect our ability to attract and retain customers and employees and can expose us to litigation and regulatory action.

Our business and operating results may be impacted by adverse economic conditions.
General economic factors and conditions in the United States or worldwide, including the general interest rate environment, unemployment rates and residential home values, may affect borrower willingness to seek loans and investor ability and desire to invest in loans. For example, during the 2008 financial crisis, banks severely constrained lending activities, which caused a decline in loan issuances. A similar crisis could negatively impact the willingness of investors and borrowers to participate on our marketplace. Although the U.S. and global economies have shown improvement, the recovery remains modest and uncertain. If present U.S. and global economic uncertainties persist, many of our investors may delay or reduce their investment in the loans facilitated through our marketplace. Adverse economic conditions could also reduce the number of individuals seeking to invest in loans facilitated on our marketplace, reduce the number of qualified borrowers seeking loans on our marketplace and result in borrowers being unable to make payments. Should any of these situations occur, our revenue and transactions on our marketplace would decline and our business would be negatively impacted.

Our business depends on our ability to successfully manage our credit risk, and failing to do so may result in high charge-off rates.
Our success depends on our ability to manage our credit risk while attracting new customers with profitable usage patterns. We select our customers, manage their accounts and establish terms and credit limits using proprietary scoring models and other analytical techniques that are designed to set terms and credit limits to appropriately compensate us for the credit risk we accept, while encouraging customers to use their available credit. The models and approaches we use to manage our credit risk may not accurately predict future charge-offs for various reasons discussed in the preceding risk factor. There can be no assurance that our credit underwriting and risk management strategies will enable us to avoid high charge-off levels or delinquencies, or that our allowance for loan losses will be sufficient to cover actual losses.

Our collection operations may not compete effectively to secure more of customers' diminished cash flow than our competitors. In addition, we may not identify customers who are likely to default on their payment obligations to us and reduce our exposure by closing credit lines and restricting authorizations quickly enough, which could have an adverse effect on our business. Our ability to manage credit risk also may be adversely affected by legal or regulatory changes (such as bankruptcy laws and minimum payment regulations) and collection regulations, competitors' actions and consumer behavior, as well as inadequate collections staffing, techniques, models and performance of vendors such as collection agencies.

Our microlending loan book exposes us to credit risk and our allowance for doubtful finance loans receivable may not be sufficient to absorb future write-offs.

Our microlending loan book is approximately $138.928 at 9/10/2019. The majority of these finance loans made are for a period of six months or less and we are in the process of determining and understanding the impairment risk of the book. We have created an allowance for doubtful finance loans receivable related to this book. However, this is a new allowance and management considered factors including the period of the loan outstanding, creditworthiness of the customers and the past payment history and trends of our established lending book. We consider this policy to be appropriate taking into account factors such as historical bad debts, current economic trends and changes in our customer payment patterns. However, additional allowances may be required should the ability of our customers to make payments when due deteriorate in the future. A significant amount of judgment is required to assess the ultimate recoverability of these finance loan receivables, including on-going evaluation of the creditworthiness of each customer.

Our epay and money transfer businesses may be susceptible to fraud and/or credit risks occurring at the retailer and/or consumer level, which could adversely affect our business.

We contract with retailers that accept payment on our behalf, which we then transfer to a trust or other operating account for payment to content providers. In the event a retailer does not transfer to us payments that it receives for prepaid content sales, whether as a result of fraud, insolvency, billing delays or otherwise, we are responsible to the content provider for the cost of the product sold. We can provide no assurance that retailer fraud or insolvency will not increase in the future or that any proceeds we receive under our credit enhancement insurance policies will be adequate to cover losses resulting from retailer fraud, which could have a material adverse effect on our business, financial condition and results of operations.

Hedging instruments often involve counterparty risks and costs.

We will be subject to credit risk with respect to our counterparties to the derivative contracts (whether a clearing corporation in the case of exchange-traded instruments or our hedge counterparty in the case of uncleared over-the-counter instruments) and other instruments entered into directly by us or held by special purpose or structured vehicles in which we invest. Counterparty risk is the risk that the other party in a derivative transaction will not fulfill its contractual obligation. Changes in the credit quality of our counterparties with respect to their derivative transactions may affect the value of those instruments. By entering into derivatives, we assume the risk that these counterparties could experience financial hardships that could call into question their continued ability to perform their obligations. As a result, concentrations of such derivatives in any one counterparty would subject our funds to an additional degree of risk with respect to defaults by such counterparty.

Our regulatory compliance programs and other enterprise risk management efforts cannot eliminate all systemic risk.

We have devoted significant time and energy to develop our enterprise risk management program, including substantially expanded regulatory compliance policies and procedures. We expect to continue to do so in the future. The goal of enterprise risk management is not to eliminate all risk, but rather to identify, assess and rank risk. The goal of regulatory compliance policies is to have formal written procedures in place that are intended to reduce the risk of inadvertent regulatory violations. Nonetheless, our efforts to identify, monitor and manage risks may not be fully effective. Many of our methods of managing risk and exposures depend upon the implementation of federal and state regulations and other policies or procedures affecting our customers or employees. Management of operational, legal and regulatory risks requires, among other things,

policies and procedures, and these policies and procedures may not be fully effective in managing these risks.

While many of the risks that we monitor and manage are described in this Risk Factors section of this Memorandum, our business operations could also be affected by additional factors that are not presently described in this section or known to us or that we currently consider immaterial to our operations.

We may face competition from other companies that offer smart card technology, other innovative payment technologies and payment processing, which could result in loss of our existing business and adversely impact our ability to successfully market additional products and services.

Our primary competitors in the payment processing market include other independent processors, as well as financial institutions, independent sales organizations, and, potentially card networks. Many of our competitors are companies who are larger than we are and have greater financial and operational resources than we have. These factors may allow them to offer better pricing terms or incentives to customers, which could result in a loss of our potential or current customers or could force us to lower our prices as well. Either of these actions could have a significant effect on our revenues and earnings.

In addition to competition that our system faces from the use of cash, checks, credit and debit cards, existing payment systems and the providers of financial services and low cost bank accounts, there are a number of other products that use smart card technology in connection with a funds transfer system. During the past several years, smart card technology has become increasingly prevalent. We believe that the most competitive product in this marketplace is EMV, a system that is promoted by most of the major card companies such as Visa, MasterCard, JCB and American Express. Also, governments and financial institutions are, to an increasing extent, implementing general-purpose reloadable prepaid cards as a low-cost alternative to provide financial services to the unbanked population. Moreover, while we see the acceptance over time of using a mobile phone to facilitate financial services as an opportunity, there is a risk that other companies will be able to introduce such services to the marketplace successfully and that customers may prefer those services to ours, based on technology, price or other factors.

If our payment processors and disbursement partners experience an interruption in service, our business and revenue would be harmed.

Our payment processors and disbursement partners have experienced service outages or an inability to connect with our processing systems and this may reoccur in the future. If a payment processor experiences a service outage or service interruption that results in our being unable to collect funds from customers, our liquidity could be harmed and we may not meet our capital requirements. We do not directly access the ACH system or payment card networks such as Visa and MasterCard, which systems enable our acceptance of bank account-funded transactions, credit cards and debit cards. As a result, we rely on banks and other payment processors and disbursement partners to process transactions. In the event of service outages in the payment card or ACH networks, or if our payment processors or disbursement partners were unable to access the payment card or ACH networks, our business would be harmed.

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 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.

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 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 zero percent of the products sold to our customers at our location. As a result, we are dependent upon the uninterrupted and efficient operation of our manufacturing facility 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, and 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 GRN Gummies Lime 100mg CBD, 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 2019 accounted for 0% 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.

Because our business is seasonal, with the highest volume of net sales during the fourth quarter, adverse events during the fourth quarter could materially affect our financial statements as a whole.

We generally recognize our highest volume of net sales during the holiday selling season, which occurs in the fourth quarter of our fiscal year. In anticipation of this holiday, we purchase substantial amounts of seasonal inventory. Adverse events, such as deteriorating economic conditions, higher unemployment, higher gas prices, public transportation disruptions, or unanticipated adverse weather could result in lower-than-planned sales during the holiday season. An excess of seasonal merchandise inventory could result if our net sales during the holiday selling season fall below seasonal norms or expectations. If our fourth quarter sales results were substantially below expectations, our financial performance and operating results could be adversely affected by unanticipated markdowns, especially in seasonal merchandise.

The seasonality of our business places increased strain on our operations.

A disproportionate amount of our sales normally occur during our fourth quarter. If we do not stock or are otherwise unable to source products sufficient to meet customer demand, our business would be adversely affected. If we liquidate products, as we have in the past, we may be required to take significant inventory markdowns or write-offs, which could reduce profits. We may experience an increase in our shipping cost due to complimentary upgrades, split-shipments, and additional long-zone shipments necessary to ensure timely delivery for the holiday season. If too many customers access our Website within a short period of time due to increased holiday demand, we may experience system interruptions that make our Website unavailable or prevent us from efficiently fulfilling orders, which may reduce the volume of goods we sell and the attractiveness of our products and services. In addition, we may be unable to adequately staff our fulfillment and customer service centers during peak periods, and delivery services and other fulfillment companies and customer service providers may be unable to meet the seasonal demand.

Our profitability may be negatively affected by inventory shrinkage.

We are subject to the risk of inventory loss and theft. We experience significant inventory shrinkage and cannot be sure that incidences of inventory loss and theft will decrease in the future or that the measures we are taking will effectively reduce the problem of inventory shrinkage. Although some level of inventory shrinkage is an unavoidable cost of doing business, if we were to experience higher rates of inventory shrinkage or incur increased security costs to combat inventory theft, our business and results of operations could be affected adversely.

Failure to execute our opportunistic buying could adversely affect our business.

We purchase the majority of our inventory opportunistically, with our buyers purchasing close to need. Establishing the "treasure hunt" nature of the off-price buying experience to drive traffic to our stores requires us to offer changing assortments of merchandise in our stores. While opportunistic buying provides our buyers the ability to buy at desirable times and prices, in the quantities we need and into market trends, it places considerable discretion in our buyers, subjecting us to risks related to the pricing, quantity, nature and timing of inventory flowing to our stores. If we are unable to provide frequent replenishment of fresh, high quality, attractively priced merchandise in our stores, it could adversely affect traffic to our stores as well as our sales and margins. We base our purchases of inventory, in part, on our sales forecasts. If our sales forecasts do not match customer demand, we may experience higher inventory levels and need to markdown excess or slow-moving inventory, leading to decreased profit margins, or we may have insufficient

inventory to meet customer demand, leading to lost sales, either of which could adversely affect our financial performance.

We need to purchase inventory sufficiently below conventional retail to maintain our pricing differential to regular department and specialty store prices and to attract customers and sustain our margins, which we may not achieve at various times and which could adversely affect our results.

Failure to execute our inventory management process could adversely affect our business.
We must also properly execute our inventory management strategies by appropriately allocating merchandise among our stores, timely and efficiently distributing inventory to stores, maintaining an appropriate mix and level of inventory in stores, appropriately changing the allocation of floor space of stores among product categories to respond to customer demand and effectively managing pricing and markdowns, and there is no assurance we will be able to do so. Failure to effectively execute our inventory management strategies could adversely affect our performance and our relationship with our customers.

Our business could suffer if we are unsuccessful in making, integrating, and maintaining commercial agreements, strategic alliances, and other business relationships.
We provide e-commerce and other services to businesses through commercial agreements, strategic alliances, and business relationships. Under these agreements, we enable sellers to offer products or services through our websites. These arrangements are complex and require substantial infrastructure capacity, personnel, and other resource commitments, which may limit the amount of business we can service. We may not be able to implement, maintain, and develop the components of these commercial relationships, which may include web services, fulfillment, customer service, inventory management, tax collection, payment processing, hardware, content, and third-party software, and engaging third parties to perform services. The amount of compensation we receive under certain of our commercial agreements is partially dependent on the volume of the other company's sales. Therefore, if the other company's offering is not successful, the compensation we receive may be lower than expected or the agreement may be terminated. Moreover, we may not be able to enter into additional commercial relationships and strategic alliances on favorable terms. We also may be subject to claims from businesses to which we provide these services if we are unsuccessful in implementing, maintaining, or developing these services.

As our agreements terminate, we may be unable to renew or replace these agreements on comparable terms, or at all. We may in the future enter into amendments on less favorable terms or encounter parties that have difficulty meeting their contractual obligations to us, which could adversely affect our operating results.

Our present and future e-commerce services agreements, other commercial agreements, and strategic alliances create additional risks such as: disruption of our ongoing business, including loss of management focus on existing businesses; impairment of other relationships; variability in revenue and income from entering into, amending, or terminating such agreements or relationships; and difficulty integrating under the commercial agreements.

Our business may be adversely affected by catastrophic events and extreme or unseasonable weather conditions.

Unforeseen events, including war, terrorism and other international conflicts, public health issues and natural disasters such as earthquakes, hurricanes or tornadoes, whether occurring in the United States or abroad, could disrupt our supply chain operations, international trade or result in political or economic instability. Any of the foregoing events could result in property losses, reduce demand for our products or make it difficult or impossible to obtain merchandise from our suppliers.

Extreme weather conditions in the areas in which our stores are located, particularly in markets where we have multiple stores, could adversely affect our business. For example, heavy snowfall, rainfall or other extreme weather conditions over a prolonged period might make it difficult for our customers to travel to our stores and thereby reduce our sales and profitability. Our business is also susceptible to unseasonable weather conditions. For example, extended periods of unseasonably warm temperatures during the winter season or cool weather during the summer season could render a portion of our inventory incompatible with those unseasonable conditions. Reduced sales from extreme or prolonged unseasonable weather conditions could adversely affect our business.

We may not timely identify or effectively respond to consumer trends or preferences, whether involving physical retail, e-commerce retail or a combination of both retail offerings, which could negatively affect our relationship with our customers and the demand for our products and services.

It is difficult to predict consistently and successfully the products and services our customers will demand. The success of our business depends in part on how accurately we predict consumer demand, availability of merchandise, the related impact on the demand for existing products and the competitive environment, whether for customers purchasing products at our stores and clubs, through our e-commerce businesses or through the combination of both retail offerings. A critical piece of identifying consumer preferences involves price transparency, assortment of products, customer experience and convenience. These factors are of primary importance to customers and they continue to increase in importance, particularly as a result of digital tools and social media available to consumers and the choices available to consumers for purchasing products online, at physical locations or through a combination of both retail offerings. Failure to timely identify or effectively respond to changing consumer tastes, preferences (including the key factors described above) and spending patterns, whether for our physical retail offerings, e-commerce offerings or through a combination of these retail offerings, could negatively affect our relationship with our customers and the demand for our products and services.

Decreases in discretionary consumer spending may have an adverse effect on us.

A substantial portion of the products and services we offer are products or services that consumers may view as discretionary items rather than necessities. As a result, our results of operations are sensitive to changes in macroeconomic conditions that impact consumer spending, including discretionary spending. Difficult macroeconomic conditions, particularly high levels of unemployment, also impact our customers' ability to obtain consumer credit. Other factors, including consumer confidence, employment levels, interest rates, tax rates, consumer debt levels, and fuel and energy costs could reduce consumer spending or change consumer purchasing habits. Slowdowns in the U.S. or global economy, or an uncertain economic outlook, could adversely affect consumer spending habits and our results of operations.

If we do not continue to source new products, our ability to compete will be undermined, and we may be unable to implement our business plan.

Our ability to compete in the direct marketing industry and to expand into the traditional retail environment depends to a great extent on our ability to develop or acquire new innovative products under particular brands and to complement these products with related families of products under those brands. If we do not source new products as our existing products mature through their product life cycles, or if we do not develop related families of products under our brands, we will not be able to implement our business plan, and the value of your investment may decrease.

Our business and results of operations may be adversely affected if we are unable to maintain our customer experience or provide high quality customer service.

The success of our business largely depends on our ability to provide superior customer experience and high quality customer service, which in turn depends on a variety of factors, such as our ability to continue to provide a reliable and user-friendly website interface for our customers to browse and purchase our products, reliable and timely delivery of our products, and superior after sales services. Our sales may decrease if our website services are severely interrupted or otherwise fail to meet our customer requests. Should we or our third-party delivery companies fail to provide our product delivery and return services in a convenient or reliable manner, or if our customers are not satisfied with our product quality, our reputation and customer loyalty could be negatively affected. In addition, we also depend on our call center and online customer service representatives to provide live assistance to our customers. If our call center or online customer service representatives fail to satisfy the individual needs of customers, our reputation and customer loyalty could be negatively affected and we may lose potential or existing customers and experience a decrease in sales. As a result, if we are unable to continue to maintain our customer experience and provide high quality customer service, we may not be able to retain existing customers or attract new customers, which could have an adverse effect on our business and results of operations.

We depend upon designers, vendors and other sources of merchandise, goods and services.

Our business could be affected by disruptions in, or other legal, regulatory, political or economic issues associated with, our supply network. Our relationships with established and emerging designers have been a significant contributor to our past success. Our ability to find qualified vendors and access products in a timely and efficient manner is often challenging, particularly with respect to goods sourced outside the United States. Our procurement of goods and services from outside the United States is subject to risks associated with political or financial instability, trade restrictions, tariffs, currency exchange rates, transport capacity and costs and other factors relating to foreign trade. In addition, our procurement of all our goods and services is subject to the effects of price increases, which we may or may not be able to pass through to our customers. All of these factors may affect our ability to access suitable merchandise on acceptable terms, are beyond our control and could negatively affect our business and results of operations.

Our advertising and marketing efforts may be costly and may not achieve desired results.

We incur substantial expense in connection with our advertising and marketing efforts. Although we target our advertising and marketing efforts on current and potential customers who we believe are likely to be in the market for the products we sell, we cannot assure you that our advertising and marketing efforts will achieve our desired results. In addition, we periodically adjust our

advertising expenditures in an effort to optimize the return on such expenditures. Any decrease in the level of our advertising expenditures, which may be made to optimize such return could adversely affect our sales.

We may be required to collect sales tax on our direct marketing operations.
With respect to the direct sales, sales or other similar taxes are collected primarily in states where we have retail stores, another physical presence or personal property. However, various states or foreign countries may seek to impose sales tax collection obligations on out-of-state direct mail companies. A successful assertion by one or more states that we or one or more of our subsidiaries should have collected or should be collecting sales taxes on the direct sale of our merchandise could have an adverse effect on our business.

Government regulation is evolving and unfavorable changes could harm our business.
We are subject to general business regulations and laws, as well as regulations and laws specifically governing the Internet, e-commerce, electronic devices, and other services. Existing and future laws and regulations may impede our growth. These regulations and laws may cover taxation, privacy, data protection, pricing, content, copyrights, distribution, mobile communications, electronic device certification, electronic waste, energy consumption, environmental regulation, electronic contracts and other communications, competition, consumer protection, web services, the provision of online payment services, information reporting requirements, unencumbered Internet access to our services, the design and operation of websites, the characteristics and quality of products and services, and the commercial operation of unmanned aircraft systems. It is not clear how existing laws governing issues such as property ownership, libel, and personal privacy apply to the Internet, e-commerce, digital content, and web services. Jurisdictions may regulate consumer-to-consumer online businesses, including certain aspects of our seller programs. Unfavorable regulations and laws could diminish the demand for our products and services and increase our cost of doing business.

Changes in federal, state or local laws and regulations could increase our expenses and adversely affect our results of operations.
Our business is subject to a wide array of laws and regulations. The current political environment, financial reform legislation, the current high level of government intervention and activism and regulatory reform may result in substantial new regulations and disclosure obligations and/or changes in the interpretation of existing laws and regulations, which may lead to additional compliance costs as well as the diversion of our management's time and attention from strategic initiatives. If we fail to comply with applicable laws and regulations we could be subject to legal risk, including government enforcement action and class action civil litigation that could disrupt our operations and increase our costs of doing business. Changes in the regulatory environment regarding topics such as privacy and information security, product safety or environmental protection, including regulations in response to concerns regarding climate change, collective bargaining activities, minimum wage laws and health care mandates, among others, could also cause our compliance costs to increase and adversely affect our business and results of operations.

Our profitability is vulnerable to cost increases, inflation and energy prices.
Future increases in our costs, such as the cost of merchandise, shipping rates, freight and fuel costs, and store occupancy costs, may reduce our profitability. The minimum wage has increased or is

scheduled to increase in multiple states and local jurisdictions, and there is a possibility Congress will increase the federal minimum wage. These cost changes may be the result of inflationary pressures, which could further reduce our sales or profitability. Increases in other operating costs, including changes in energy prices, wage rates and lease and utility costs, may increase our costs of sales or operating expenses and reduce our profitability.

Fluctuations in the mix of customer demand for our various types of solution offerings could impact our financial performance and ability to forecast performance.

Due to fluctuations in customer needs, changes in customer industries, and general economic conditions, customer demand for the range of our offerings varies from time to time and is not predictable. Hemp Derived CBD is newly legal and the government's laws regarding these products can greatly impact the legality around our ability to sell these products. See the Farm Bill from 2018. In addition, our gross margins vary by customer and by segment and the mix of services provided to our customers could impact our results of operations as certain of our customers and segments have different gross margin profiles. Generally, the profitability of an account increases over time. As a result, the mix of solutions we provide to our customers varies at any given time, both within a quarter and from quarter-to-quarter. These variations in service mix impact gross margins and the predictability of gross margins for any period. You should not rely on the results of any one quarter as an indication of our future performance.

Our operating results may fluctuate due to factors that are difficult to forecast and not within our control.

Our past operating results may not be accurate indicators of future performance, and you should not rely on such results to predict our future performance. Our operating results have fluctuated significantly in the past, and could fluctuate in the future. Factors that may contribute to fluctuations include:

* changes in aggregate capital spending, cyclicality and other economic conditions, or domestic and international demand in the industries we serve;

* our ability to effectively manage our working capital;

* our ability to satisfy consumer demands in a timely and cost-effective manner;

* pricing and availability of labor and materials;

* our inability to adjust certain fixed costs and expenses for changes in demand;

* shifts in geographic concentration of customers, supplies and labor pools; and

* seasonal fluctuations in demand and our revenue.

If we fail to attract and retain enough sufficiently trained customer service associates and other personnel to support our operations, our business and results of operations will be seriously harmed.

We rely on customer service associates, and our success depends to a significant extent on our ability to attract, hire, train and retain qualified customer service associates. Companies in our

industry, including us, experience high employee attrition. Our attrition rate for our customer service associates who remained with us following a 90-day training and orientation period was on average approximately 5% per month. A significant increase in the attrition rate among our customer service associates could decrease our operating efficiency and productivity. Our failure to attract, train and retain customer service associates with the qualifications necessary to fulfill the needs of our existing and future clients would seriously harm our business and results of operations.

Our ability to sell our products and services is dependent on the quality of our technical support services, and our failure to offer high quality technical support services would have a material adverse effect on our sales and results of operations.

Once our products are deployed within our end-customers' operations, end-customers depend on our technical support services to resolve any issues relating to these products. If we do not effectively assist our customers in deploying these products, succeed in helping our customers quickly resolve post-deployment issues, and provide effective ongoing support, our ability to sell additional products and services to existing customers would be adversely affected and our reputation with potential customers could be damaged. As a result, our failure to maintain high quality support services would have an adverse effect on our business and results of operations.

We may be adversely affected by cyclicality, volatility or an extended downturn in the United States or worldwide economy, or in or related to the industries we serve.

Our revenues are generated primarily from servicing customers seeking to hire qualified professionals in the technology, healthcare, hospitality and finance sectors and the energy industry. Demand for these professionals tends to be tied to economic and business cycles. Increases in the unemployment rate, specifically in the technology, healthcare, finance and other vertical industries we serve, cyclicality or an extended downturn in the economy could cause our revenues to decline. Therefore, our operating results, business and financial condition could be significantly harmed by an extended economic downturn or future downturns, especially in regions or industries where our operations are heavily concentrated. Further, we may face increased pricing pressures during such periods as customers seek to use lower cost or fee services, which may adversely affect our financial condition and results of operations.

We are subject to rapid technological change and dependence on new product development.

Our industry is characterized by rapid and significant technological developments, frequent new product introductions and enhancements, continually evolving business expectations and swift changes. To compete effectively in such markets, we must continually improve and enhance our products and services and develop new technologies and services that incorporate technological advances, satisfy increasing customer expectations and compete effectively on the basis of performance and price. Our success will also depend substantially upon our ability to anticipate, and to adapt our products and services to our collaborative partner's preferences. There can be no assurance that technological developments will not render some of our products and services obsolete, or that we will be able to respond with improved or new products, services, and technology that satisfy evolving customers' expectations. Failure to acquire, develop or introduce new products, services, and enhancements in a timely manner could have an adverse effect on our business and results of operations. Also, to the extent one or more of our competitors introduces products and services that better address a customer's needs, our business would be adversely affected.

Failure to obtain new clients or renew client contracts on favorable terms could adversely affect results of operations.

We may face pricing pressure in obtaining and retaining our clients. Our clients may be able to seek price reductions from us when they renew a contract, when a contract is extended, or when the client's business has significant volume changes. They may also reduce services if they decide to move services in-house. On some occasions, this pricing pressure results in lower revenue from a client than we had anticipated based on our previous agreement with that client. This reduction in revenue could result in an adverse effect on our business and results of operations.

Further, failure to renew client contracts on favorable terms could have an adverse effect on our business. Our contracts with clients generally run for several years and include liquidated damage provisions that provide for early termination fees. Terms are generally renegotiated prior to the end of a contract's term. If we are not successful in achieving a high rate of contract renewals on favorable terms, our business and results of operations could be adversely affected.

We derive significant revenue and profit from commercial and federal government contracts awarded through competitive bidding processes, including renewals, which can impose substantial costs on us.

Many of these contracts are extremely complex and require the investment of significant resources in order to prepare accurate bids and proposals. Competitive bidding imposes substantial costs and presents a number of risks, including: (i) the substantial cost and managerial time and effort that we spend to prepare bids and proposals for contracts that may or may not be awarded to us; (ii) the need to estimate accurately the resources and costs that will be required to implement and service any contracts we are awarded, sometimes in advance of the final determination of their full scope and design; (iii) the expense and delay that may arise if our competitors protest or challenge awards made to us pursuant to competitive bidding, and the risk that such protests or challenges could result in the requirement to resubmit bids, and in the termination, reduction, or modification of the awarded contracts; and (iv) the opportunity cost of not bidding on and winning other contracts we might otherwise pursue. Adverse events or developments in any of these bidding risks and uncertainties could materially and negatively impact our business and results of operations.

We may rely on subcontractors and partners to provide customers with a single-source solution or we may serve as a subcontractor to a third-party prime contractor.

From time to time, we may engage subcontractors, teaming partners or other third parties to provide our customers with a single-source solution for a broader range of service needs. Similarly, we are and may in the future be engaged as a subcontractor to a third-party prime contractor. Subcontracting arrangements pose unique risks to us because we do not have control over the customer relationship, and our ability to generate revenue under the subcontract is dependent on the prime contractor, its performance and relationship with the customer and its relationship with us. While we believe that we perform appropriate due diligence on our prime contractors, subcontractors and teaming partners and that we take adequate measures to ensure that they comply with the appropriate laws and regulations, we cannot guarantee that those parties will comply with the terms set forth in their agreements with us (or in the case of a prime contractor, their agreement with the customer), or that they will be reasonable in construing their contractual rights and obligations, always act appropriately in dealing with us or customers, provide adequate service, or remain in compliance with the relevant laws, rules or regulations. We may have disputes with our

prime contractors, subcontractors, teaming partners or other third parties arising from the quality and timeliness of work being performed, customer concerns, contractual interpretations or other matters. We may be exposed to liability if we lose or terminate a subcontractor or teaming partner due to a dispute, and subsequently have difficulty engaging an appropriate replacement or otherwise performing their functions in-house, such that we fail to fulfill our contractual obligations to our customer. In the event a prime contract, under which we serve as a subcontractor, is terminated, whether for non-performance by the prime contractor or otherwise, then our subcontract will similarly terminate and we could face contractual liability and the resulting contract loss could adversely affect our business and results of operations.

Our business and financial condition may be impacted by military actions, global terrorism, natural disasters and political unrest.

Military actions in Iraq, Afghanistan and elsewhere, global terrorism, natural disasters and political unrest in the Middle East and other countries are among the factors that may adversely impact regional and global economic conditions and our clients' ability, capacity and need to invest in our services. Additionally, hurricanes or other unanticipated catastrophes, both in the U.S. and globally, could disrupt our operations and negatively impact our business as well as disrupt our clients' businesses, which may result in a further adverse impact on our business. As a result, significant disruptions caused by such events could materially and adversely affect our business and financial condition.

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 CBD.io and CBD.co, 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

Cannabiscope is a SaaS menu platform that integrates with THC/CBD retailers POS systems to present their inventory live onsite and an e-commerce site, while collecting industry data. Customers sign annual contracts and either pay a set monthly fee or give us 1% of their total revenue, whichever is higher for us to provide this service.

Business Plan

The Company is committed to bringing the best user experience to its customers through its innovative software and services. The Company's business strategy leverages its unique ability to design and develop its own application software and services to provide its customers products and solutions with innovative design, superior ease-of-use and seamless integration. As part of its strategy, the Company continues to expand its platform for the discovery and delivery of third-party digital content and applications. Our goal is to grow our product offering in phases to generate as much revenue as possible. Phase 1 of our revenue plan is to provide CBD retailers an e-commerce solution that will empower payment processing. Phase 2 will focus on Point of Sale integration with THC businesses that will allow them to show their inventory live both onsite and online. We will be ready to provide THC businesses e-commerce as soon as the federal government legalizes cannabis at a national level. Phase 3 of our growth will focus on collecting and presenting

relevant data to our customers to help them drive business decisions and generate sales. Phase 3 will look to diversify revenue streams through unique data offerings based on our POS integrations and unique insights derived from the interactive wheel.

History of the Business

On August 27, 2019, the company restructured from a Nevada limited liability company to a Nevada corporation.

The Company's Products and/or Services

Product / Service	Description	Current Market
Cannabiscope Interactive Wheel	An interactive shopping tool where people search for THC/CBD products based on what they want out of the experience.	THC Dispensary THC Delivery THC Brands CBD Dispensaries CBD Brands

We plan to build a complete platform that will provide our customers an interactive shopping wheel, a content management system, and an e-commerce cart that can scale nationally in the U.S. The platform will empower businesses in the cannabis industry to accelerate revenue both onsite and online. Once the platform is to scale, we plan to collect and organize industry data to sell to our clients as data dashboards that will provide insights to drive our customers business decisions.

We currently have direct relationships with our customers that have been built through networking. We are not conducting any outbound or inbound marketing yet as our product is not ready to scale.

Competition

The Company's primary competitors are Leafly, Weedmaps, Dutchie, Potguide, iHeart Jane, Baker, Massroots.

The cannabis tech space is competitive and dynamic. Each of the competitors provide front end menus, which all have API integration with the Point of Sale systems. The competitors are well funded and have market penetration. However the market is still new, the dust hasn't settled. There is room for new and innovative ideas. What differentiates Cannabiscope from all other online menus is the wheel. The user interface is one of a kind. There's nothing like it as far as customer experience. Cannabiscope is set to be the standard in classifying cannabis products. An incredible amount of information is packed in an easy to use device. Ordering online is becoming more and more common. The other menu providers' strategy is to show the retailer's location and what they have in stock. Cannabiscope lets customers find exactly what they are looking for and then lets them know where it's available. The cannabis community has embraced the wheel as an educational tool. Doctors and nurses use it to consult patients. Budtenders love it as a sales tool when talking to customers to further explain products. Cannabiscope's interactive menu is simply the best way to discover cannabis, get information about products and sell items

Customer Base

The Company's customers are primarily in retailers in the cannabis and hemp space.

Intellectual Property

Trademarks

Application or Registration #	Goods / Services	Mark	File Date	Registration Date	Country
5177251	IC 035 Advertising and commercial information services, via the Internet; Advertising and directory services, namely, promoting the services of others by providing a web page featuring links to the websites of others; Advertising and marketing services, namely, promoting the goods and services of others; Advertising on the Internet for others; Advertising	Cannabiscope	August 18, 2015	April 4, 2017	United States

	services, namely, promoting the brands, goods and services of others. IC 044 Computerized and Internet-based medical information services concerning cannabis and related products thereof; medical information services concerning cannabis and products thereof using an interactive mapping system; medical information services providing cannabis strain classification.				

Governmental/Regulatory Approval and Compliance

We are heavily dependent on state laws and federal laws regarding medicinal and recreational cannabis. While our business does not directly touch the plant, our customers do and it will affect our ability to provide services.

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 2764 N. Green Valley Parkway, Suite #345, Henderson, NV 89014.

The Company conducts business in California.

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

David Schacter

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

January 2017 - Present CEO

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

Cannabiscope January 2019-Present General Manager Cannabiscope is a SaaS Menu Platform that accelerates revenue through on-site and online sales while collecting valuable data insights across the industry. This platform looks to empower CBD online retailers, THC Dispensary and delivery services to transition into the e-commerce age of cannabis while finding unique industry insights to create growth opportunities. As general manager Paul has reworked the business plan and pitch deck to attract series A capital. He is working closely with the founders to update the product road map and data offering to keep Cannabsicope a cutting edge solution for an explosive industry. He will be instrumental in building the team that will take the platform to its next stage of growth.
The Guru Experience www.TheGuru.co June 2015 - December 2018: Co-Founder & Chief Innovation Officer At Guru we are working to fundamentally change how people learn about culture and transform the way cultural institutions drive visitor acquisition. This SaaS solution provides our partners with a cutting edge location aware mobile platform that drives experience through digital storytelling. Audio overviews, pictures, videos, augmented reality and virtual reality experiences allow people to learn in entirely new ways. These experiences allow our partners to gather critical data around how people move, engage and learn in their respective institutions. As the Chief Innovation Officer at Guru it is my job to push the boundaries of what is possible in mobile education. I manage incredible teams of developers and creatives to construct an affordable platform approach that solves our partner's specific needs. I work to provide users with unforgettable experiences that drive the insights for strategic planning and growth in the cultural institution space.

Officers

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

David Schacter

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

January 2017 - Present CEO

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

Cannabiscope January 2019-Present General Manager Cannabiscope is a SaaS Menu Platform that accelerates revenue through on-site and online sales while collecting valuable data insights across the industry. This platform looks to empower CBD online retailers, THC Dispensary and delivery services to transition into the e-commerce age of cannabis while finding unique industry insights to create growth opportunities. As general manager Paul has reworked the business plan and pitch deck to attract series A capital. He is working closely with the founders to update the product road map and data offering to keep Cannabsicope a cutting edge solution for an explosive industry. He will be instrumental in building the team that will take the platform to its next stage of growth.
The Guru Experience www.TheGuru.co June 2015 - December 2018: Co-Founder & Chief Innovation Officer At Guru we are working to fundamentally change how people learn about culture and transform the way cultural institutions drive visitor acquisition. This SaaS solution provides our partners with a cutting edge location aware mobile platform that drives experience through digital storytelling. Audio overviews, pictures, videos, augmented reality and virtual reality experiences allow people to learn in entirely new ways. These experiences allow our partners to gather critical data around how people move, engage and learn in their respective institutions. As the Chief Innovation Officer at Guru it is my job to push the boundaries of what is possible in mobile education. I manage incredible teams of developers and creatives to construct an affordable platform approach that solves our partner's specific needs. I work to provide users with unforgettable experiences that drive the insights for strategic planning and growth in the cultural institution space.

Indemnification

Indemnification is authorized by the Company to directors, officers or controlling persons acting in their professional capacity pursuant to Nevada 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 0 employees.

CAPITALIZATION AND OWNERSHIP

Capitalization

The Company has issued the following outstanding Securities:

Type of security	Common Stock
Amount outstanding	4,117,646
Voting Rights	One vote per share
Anti-Dilution Rights	N/A

Type of security	Crowd Notes
Amount outstanding	138,928
Voting Rights	N/A
Anti-Dilution Rights	N/A

The Company has the following debt outstanding:

Type of debt	Convertible Notes
Name of creditor	Regulation CF Investors
Amount outstanding	$138,928
Interest rate and payment schedule	N/A
Amortization schedule	N/A
Describe any collateral or security	None
Maturity date	None
Other material terms	All Notes have a Valuation Cap of $2.5M. Upon the occurrence of a sale by the Company of its Preferred Units following the from which the Company receives gross proceeds of not less than $1,000,000 (the "Qualified

	Financing") the Note will convert into Preferred Stock pursuant to the following: (a) if the Investor is not a Major Investor (an investor with a purchase price of $25,000 or greater), the Note will convert upon the earlier of (i) the Company's election or (ii) a change of control transaction or IPO; and (b) if the Investor is a Major Investor, the Company will convert the Note prior to the closing of the Qualified Equity Financing.

The total amount of outstanding debt of the company is $138,928.00

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
Crowd Notes	138,928	$138,928	Platform Development	September 25, 2019	Regulation CF

Ownership

A majority of the Company is owned by David Schacter, his family, the initial co-founders of Cannabiscope. Paul Shockley has 14.28% ownership as a co-founder and GM.

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
David Schacter	47.1%

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.

Operations

The first major financial milestone will be to get monthly recurring revenue in Q3 of 2020 while testing the stability of the platform with CBD products. In Q4 we plan to launch a more complete CBD offering that will allow for an increase in monthly recurring revenue. At this point, we will be looking to put people on salaries while scaling up our marketing efforts. By Q1 2021, we hope to have enough revenue to support both marketing and R&D efforts.

Cannabiscope will be spending the proceeds from this offering on product development, general operation costs, and a small go to market campaign. The product development will be broken into three phases. Phase 1 will focus on getting the foundation of the platform built with an integration with a payment processor for CBD business. It should take around 6 months to finish phase 1 where a soft launch will occur where 5-10 CBD companies will pay 10% of all transactions per month for our service. Phase 2 will work to expand the platform offering to include the "Interactive Wheel" and will make sure the current CBD customers are receiving effective service from the platform. Phase 2 should take 6 months. Phase 3 will begin the POS integration process for CBD businesses as well as working to make the "Transaction Data Dashboard" can go live at the end of this phase. At the end of Phase 3, which should take 3 months, the company will have spent all the proceeds set aside for the product. We will have had 5 months of revenue and hope this will support hiring a set of employees to empower further growth. For the small go to market campaign, one month before the end of Phase 2 of product, we will begin our larger scale marketing efforts to get as many CBD online retailers on the platform as possible. We hope that this major push will generate enough monthly recurring revenue that we will be able sustain a marketing push from that point on. We will also want to invest in research and development around the data side of our offering.

Liquidity and Capital Resources

On September 25, 2019 the Company conducted an offering pursuant to Regulation CF and raised $138,928.

The Company does not have any additional sources of capital other than the proceeds from the Regulation CF Offering.

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 are transferred: 1) to the Company, 2) to an accredited investor, as

defined by Rule 501(d) of Regulation D promulgated under the Securities Act, 3) as part of an IPO 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 member of the family 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.

In addition to the foregoing restrictions, prior to making any transfer of the Securities or any Securities into which they are convertible, such transferring Investor must either make such transfer pursuant to an effective registration statement filed with the SEC or provide the Company with an opinion of counsel stating that a registration statement is not necessary to effect such transfer.

In addition, the Investor may not transfer the Securities or any Securities into which they are convertible to any of the Company's competitors, as determined by the Company in good faith.

Furthermore, upon the event of an IPO, the capital stock into which the Securities are converted will be subject to a lock-up period and may not be sold for up to 180 days following such IPO.

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 conducted the following transactions with related persons:

Loans

Related Person/Entity	Beverly Katz and Benjamin Shacter
Relationship to the Company	Immediate family members
Total amount of money involved	$34,420.00
Benefits or compensation received by related person	Compensated with Shares
Benefits or compensation received by Company	During the restructuring of the company to a corporation, Green Valley Affiliates converted the debt into common stock in the company. The holders of the debt received a combined 588,236 shares, or 10% of the company, in the conversion.
Description of the transaction	Note Payable

Related Person/Entity	David Schacter and Paul Shockley
Relationship to the Company	Officer/Director/Co-Founder of the Company (Schacter); Co-Founder of the Company (Shockley)
Total amount of money involved	$36,575.00
Benefits or compensation received by related person	N/A
Benefits or compensation received by Company	The Related Persons loaned an aggregate of $36,575.00 to the Company for expenses incurred in 2019 related to the Reg CF offering (marketing, legal, etc.). The note did not bear any interest. The full amount was repaid to the Related Persons in January 2020, and no obligations remain outstanding from the parties.
Description of the transaction	Note Payable

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.

/s/David Schacter
(Signature)

David Schacter
(Name)

CEO/Founder
(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/David Schacter
(Signature)

David Schacter
(Name)

CEO/Founder
(Title)

April 20,2020
(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.

I, David Schacter, being the Chief Executive Officer of Green Valley Affiliates, Inc., a Nevada corporation (the "Company"), hereby certify as of the date listed below that:

(i) the accompanying unaudited financial statements of the Company, which comprise the balance sheet as of December 31, 2019 and the related statements of income (deficit), stockholder's equity and cash flows for the year ended December 31, 2018, and the related notes to said financial statements (collectively, the "Financial Statement"), are true and complete in all material respects; and

(ii) The tax return information of the Company included in this Form C reflects accurately the information reported on the tax return for the Company filed for the fiscal year ended December 31, 2019.

/s/David Schacter
(Signature)

David Schacter
(Name)

Chief Executive Officer
(Title)

April 20, 2020
(Date)

EXHIBITS

Exhibit A Financial Statements

EXHIBIT A

Financial Statements

GREEN VALLEY AFFILIATES, INC.

Financial Statements For The Years Ended December 31, 2019 and 2018 (Reviewed)

March 18, 2020

GREEN VALLEY AFFILIATES, INC.
BALANCE SHEET
DECEMBER 31, 2019 AND 2018

	2019	2018
ASSETS		
CURRENT ASSETS		
Cash	$ 137,923	$ 6,372
Prepaid Expense	20,000	
TOTAL CURRENT ASSETS	157,923	6,372
TOTAL ASSETS	157,923	6,372
LIABILITIES AND SHAREHOLDERS' EQUITY		
CURRENT LIABILITIES		
Deferred Revenue	-	3,125
Related Party Loan	36,575	
TOTAL CURRENT LIABILITIES	36,575	3,125
NON-CURRENT LIABILITIES		
Related Party Loan	-	34,420
Convertible Note	138,928	
TOTAL LIABILITIES	175,503	37,545
SHAREHOLDERS' EQUITY		
Common Stock (10,000,000 shares authorized; 3,529,410 issued and outstanding at 12/31/2018; 4,117,646 issued and outstanding at 12/31/2019; $0.0001 par value)	412	353
Additional Paid-in Capital	41,027	
Retained Earnings (Deficit)	(59,019)	(31,526)
TOTAL SHAREHOLDERS' EQUITY	(17,580)	(31,173)
TOTAL LIABILITIES AND SHAREHOLDERS' EQUITY	$ 157,923	$ 6,372

GREEN VALLEY AFFILIATES, INC.
INCOME STATEMENT
FOR THE YEARS ENDED DECEMBER 31, 2019 AND 2018

	2019	2018
Operating Income		
Sales, Net	$ 3,125	$ 12,375
Cost of Goods Sold		5,811
Gross Profit	3,125	6,564
Operating Expense		
General & Administrative	5,946	11,310
Equipment	51	1,429
Advertisement	10,250	1,260
Professional Services	14,371	11,056
	30,618	25,055
Net Income from Operations	(27,493)	(18,491)
Net Income	$ (27,493)	$ (18,491)

GREEN VALLEY AFFILIATES, INC.
STATEMENT OF CHANGES IN SHAREHOLDERS' EQUITY
FOR THE YEARS ENDED DECEMBER 31, 2019 AND 2018

	Common Stock Number	Common Stock Amount	Additional Paid-in Capital	Retained Earnings	Total Stockholders' Equity
Balance at December 31, 2017	3,529,410	$ 353	$ -	$ (13,032)	$ (12,679)
Issuance of Stock					
Net Income				(18,491)	(18,491)
Balance at December 31, 2018	$ 3,529,410	353	$ -	$ (31,523)	$ (31,173)
Issuance of Stock	588,236	59	41,027		41,086
Net Income				(27,493)	(27,493)
Balance at December 31, 2019	$ 4,117,646	412	$ 41,027	$ (59,019)	$ (17,580)

GREEN VALLEY AFFILIATES, INC.
STATEMENT OF CASH FLOWS
FOR THE YEARS ENDED DECEMBER 31, 2019 AND 2018

	2019	2018
Cash Flows From Operating Activities		
Net Income (Loss) For The Period	$ (27,493)	$ (18,491)
Change in Prepaid Expense	(20,000)	440
Change in Deferred Revenue	(3,125)	3,125
Net Cash Flows From Operating Activities	(50,618)	(14,926)
Cash Flows From Financing Activities		
Proceeds from Issuance of Related Party Loan	43,241	20,790
Proceeds from Issuance of Convertible Note	138,928	-
Net Cash Flows From Financing Activities	182,169	20,790
Cash at Beginning of Period	6,372	508
Net Increase (Decrease) In Cash	131,551	5,864
Cash at End of Period	$ 137,923	$ 6,372

NOTE A- ORGANIZATION AND NATURE OF ACTIVITIES

Green Valley Affiliates, Inc. ("the Company") is a corporation organized under the laws of Nevada. The Company operates as a software for service menu platform that helps businesses in the cannabis space educate consumers and sell products in-store and online.

NOTE B- GOING CONCERN MATTERS

The financial statements have been prepared on the going concern basis, which assumes that the Company will continue in operation for the foreseeable future. However, management has identified the following conditions and events that created an uncertainty about the ability of the Company to continue as a going concern. The company sustained net operating losses in 2019 and 2018.

The following describes management's plans that are intended to mitigate the conditions and events that raise substantial doubt about the Company's ability to continue as a going concern. The Company plans to raise additional capital through a Regulation CF offering. The Company's ability to meet its obligations as they become due is dependent upon the success of management's plans, as described above.

These conditions and events create an uncertainty about the ability of the Company to continue as a going concern through March 18, 2021 (one year after the date that the financial statements are available to be issued). The financial statements do not include any adjustments that might be necessary should the Company be unable to continue as a going concern.

NOTE C- SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

Basis of Presentation

The accompanying financial statements have been prepared in accordance with accounting principles generally accepted in the United States of America ("US GAAP").

Reclassifications
Certain amounts in the 2018 financial statements have been reclassified to conform to the 2019 presentation with no effect on the Company's previously reported net income or capital and surplus.

Significant Risks and Uncertainties

The Company is subject to customary risks and uncertainties associated with the dependence on key personnel, costs of services provided by third parties, the need to obtain additional financing, and limited operating history.

Use of Estimates

The preparation of financial statements in conformity with GAAP requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent

NOTE C- SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES (continued)

Use of Estimates (continued)

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 balances, and highly liquid investments with maturities of three months or less when purchased.

Revenue

The Company recognizes revenue when persuasive evidence of an arrangement exists, delivery has occurred, or services have been rendered, the fee for the arrangement is fixed or determinable and collectability is reasonably assured.

Advertising

The Company records advertising expenses in the year incurred.

Debt Issuance Costs

The Company expenses costs of issuing debt in the year incurred.

Equity Based Compensation

The Company accounts for stock options issued to employees under ASC 718 (Stock Compensation). Under ASC 718, share-based compensation cost to employees is measured at the grant date, based on the estimated fair value of the award, and is recognized as an item of expense ratably over the employee's requisite vesting period. The Company has elected early adoption of ASU 2018-07, which permits measurement of stock options at their intrinsic value, instead of their fair value. An option's intrinsic value is defined as the amount by which the fair value of the underlying stock exceeds the exercise price of an option. In certain cases, this means that option compensation granted by the Company may have an intrinsic value of $0.

The Company measures compensation expense for its non-employee stock-based compensation under ASC 505 (Equity). The fair value of the option issued or committed to be issued is used to measure the transaction, as this is more reliable than the fair value of the services received. The fair value is measured at the value of the Company's common stock on the date that the commitment for performance by the counterparty has been reached or the counterparty's performance is complete. The fair value of the equity instrument is charged directly to expense and credited to additional paid-in capital.

Income Taxes

The Company applies ASC 740 Income Taxes ("ASC 740"). Deferred income taxes are recognized for the tax consequences in future years of differences between the tax bases of assets and liabilities and their financial statement reported amounts at each period end, based on enacted tax laws and statutory tax

NOTE C- SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES (continued)

Income taxes (continued)

rates applicable to the periods in which the differences are expected to affect taxable income. Valuation allowances are established, when necessary, to reduce deferred tax assets to the amount expected to be realized. The provision for income taxes represents the tax expense for the period, if any and the change during the period in deferred tax assets and liabilities. ASC 740 also provides criteria for the recognition, measurement, presentation and disclosure of uncertain tax positions. A tax benefit from an uncertain position is recognized only if it is "more likely than not" that the position is sustainable upon examination by the relevant taxing authority based on its technical merit.

The Company is subject to tax filing requirements as a corporation in the federal jurisdiction of the United States. The Company sustained net operating losses during fiscal years 2018 and 2019. Net operating losses will be carried forward to reduce taxable income in future years. Due to management's uncertainty as to the timing and valuation of any benefits associated with the net operating loss carryforwards, the Company has elected to recognize an allowance to account for them in the financial statements, but has fully reserved it. Under current law, net operating losses may be carried forward indefinitely.

The Company is subject to modified business tax filing requirements in the State of Nevada.

Recently Adopted Accounting Pronouncements

From time to time, new accounting pronouncements are issued by the Financial Accounting Standards Board, or FASB, or other standard setting bodies and adopted by the Company as of the specified effective date. Unless otherwise discussed, the Company believes that the impact of recently issued standards that are not yet effective will not have a material impact on its financial position or results of operations upon adoption.

In November 2015, the FASB issued ASU (Accounting Standards Update) 2015-17, *Balance Sheet Classification of Deferred Taxes*, or ASU 2015-17. The guidance requires that all deferred tax assets and liabilities, along with any related valuation allowance, be classified as noncurrent on the balance sheet. For all entities other than public business entities, the guidance becomes effective for financial statements issued for annual periods beginning after December 15, 2017, and interim periods within annual periods beginning after December 15, 2018. Early adoption is permitted for all entities as of the beginning of an interim or annual reporting period. The adoption of ASU 2015-17 had no material impact on the Company's financial statements and related disclosures.

In November 2016, the FASB issued ASU 2016-18, *Statement of Cash Flows (Topic 230), Restricted Cash*, or ASU 2016-18. The amendments of ASU 2016-18 were issued to address the diversity in classification and presentation of changes in restricted cash and restricted cash equivalents on the statement of cash flows which is currently not addressed under Topic 230. ASU 2016-18 would require an entity to include amounts generally described as restricted cash and restricted cash equivalents with cash and cash equivalents when reconciling the beginning of period and end of period total amounts on the statement of cash flows. This guidance is effective for annual reporting periods, and interim periods within those years, beginning after December 15, 2018 for non-public entities. Early adoption is permitted, and the standard must be applied retrospectively. The adoption of ASU 2016-18 had no material impact on the Company's financial statements and related disclosures.

NOTE C- SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES (continued)

Recently Adopted Accounting Pronouncements (continued)

In May 2014, the FASB issued ASU, 2014-09—*Revenue from Contracts with Customers (Topic 606)*, or ASU 2014-09, and further updated through ASU 2016-12, or ASU 2016-12, which amends the existing accounting standards for revenue recognition. ASU 2014-09 is based on principles that govern the recognition of revenue at an amount to which an entity expects to be entitled to when products are transferred to customers. This guidance is effective for annual reporting periods, and interim periods within those years, beginning December 15, 2018 for non-public entities. The new revenue standard may be applied retrospectively to each prior period presented or retrospectively with the cumulative effect recognized as of the date of adoption. The adoption of ASU 2014-09 had no material impact on the Company's financial statements and related disclosures.

In February 2016, the FASB issued ASU 2016-02, *Leases (Topic 842)*, or ASU 2016-02, which supersedes the guidance in ASC 840, *Leases*. The new standard requires lessees to apply a dual approach, classifying leases as either finance or operating leases based on the principle of whether or not the lease is effectively a financed purchase by the lessee. This classification will determine whether lease expense is recognized based on an effective interest method or on a straight-line basis over the term of the lease. A lessee is also required to record a right-of-use asset and a lease liability for all leases with a term of greater than 12 months regardless of their classification. Leases with a term of 12 months or less will be accounted for similar to existing guidance for operating leases today. This guidance is effective for annual reporting periods beginning after December 15, 2019 for non-public entities. The adoption of ASU 2016-02 had no material impact on the Company's financial statements and related disclosures.

In March 2016, the FASB issued ASU 2016-09, *Improvements to Employee Share-based Payment Accounting*, or ASU 2016-09. ASU 2016-09 simplifies several aspects of the accounting for share-based payment transactions, including the income tax consequences, classification of awards as either equity or liabilities, and classification on the statement of cash flows. Some of the areas of simplification apply only to non-public companies. This guidance was effective on December 31, 2016 for public entities. For entities other than public business entities, the amendments are effective for annual periods beginning after December 15, 2017, and interim periods within annual periods beginning after December 15, 2018. Early adoption is permitted for an entity in any interim or annual period for which financial statements have not been issued or made available for issuance. An entity that elects early adoption must adopt all amendments in the same period. The adoption of ASU 2016-09 had no material impact on the Company's financial statements and related disclosures.

In May 2017, the FASB issued ASU 2017-09, *Compensation—Stock Compensation (Topic 718): Scope of Modification Accounting*, or ASU 2017-09, which clarifies when to account for a change to the terms or conditions of a share-based payment award as a modification. Under the new guidance, modification accounting is required only if the fair value, the vesting conditions, or the classification of the award (as equity or liability) changes as a result of the change in terms or conditions. This guidance is effective for annual reporting periods, and interim periods within those years, beginning after December 15, 2017, for both public entities and non-public entities. Early adoption is permitted. The adoption of ASU 2017-09 had no material impact on the Company's financial statements and related disclosures.

NOTE D- DEBT

In 2017, the company issued a note payable to a related party in exchange for cash for the purpose of funding continuing operations ("the Related Party Note"). The note does not accrue interest and is payable at a future date to be determined by management. In 2019, the note payable was converted into 588,236 shares of common stock.

In 2019, the Company issued a note payable to a related party in exchange for cash for the purpose of funding continuing operations ("the Related Party Note"). The note does not accrue interest and is payable at a future date to be determined by management.

In 2019, the Company entered into a Form C/A and issued a convertible note to unrelated parties in exchange for cash for the purpose of funding continuing operations ("the Crowdfunding Note"). The note does not accrue interest and is convertible to Preferred Stock or payable by the Company upon the earlier of (a) a Qualified Equity Financing of gross proceeds not less than $1,000,000 in exchange for Preferred Stock, or (b) a Corporate Transaction. Costs associated with the issuance of debt totaled $1,000 for 2019.

NOTE E- FAIR VALUE MEASUREMENTS

Fair value is an exit price, representing the price that would be received to sell an asset or paid to transfer a liability in an orderly transaction between market participants based on the highest and best use of the asset or liability. As such, fair value is a market-based measurement that should be determined based on assumptions that market participants would use in pricing an asset or liability. The Company uses valuation techniques to measure fair value that maximize the use of observable inputs and minimize the use of unobservable inputs. These inputs are prioritized as follows:

Level 1 - Observable inputs, such as quoted prices for identical assets or liabilities in active markets;
Level 2 - Inputs, other than the quoted prices in active markets, that are observable either directly or indirectly, such as quoted prices for similar assets or liabilities, or market-corroborated inputs; and
Level 3 - Unobservable inputs for which there is little or no market data which require the reporting entity to develop its own assumptions about how market participants would price the assets or liabilities.

The valuation techniques that may be used to measure fair value are as follows:

Market approach - Uses prices and other relevant information generated by market transactions involving identical or comparable assets or liabilities.
Income approach - Uses valuation techniques to convert future amounts to a single present amount based on current market expectations about those future amounts, including present value techniques, option-pricing models, and excess earnings method.
Cost approach - Based on the amount that currently would be required to replace the service capacity of an asset (replacement cost).

NOTE F- CONCENTRATIONS OF RISK

Financial instruments that potentially subject the Company to credit risk consist of cash and cash equivalents. The Company places its cash and cash equivalents with a limited number of high-quality financial institutions and at times may exceed the amount of insurance provided on such deposits.

NOTE G – CORRECTION OF PRIOR PERIOD ERRORS

Management has made an adjustment to decrease Common Stock and increase Retained Earnings as of December 31, 2018. The error occurred due to a miscommunication in the date of certain stock issuance related to the extinguishment of related party loan. The adjustment was made as follows:

Retained Earnings	$59	
Common Stock		$59

NOTE H- SUBSEQUENT EVENTS

In December 2019, a new coronavirus known as COVID-19 was first detected in China and has now spread globally. The Secretary of Health and Human Services (HHS) declared a public health emergency on January 31, 2020, and effective March 1, 2020, the President of the United States declared the COVID-19 outbreak a national emergency. On March 11, 2020, the World Health Organization characterized the COVID-19 outbreak as a pandemic. Management cannot anticipate all of the ways in which health pandemics, such as COVID-19, could adversely impact the Company.

Management has evaluated subsequent events through March 18, 2020, the date the financial statements were available to be issued, and no events, except as noted above, that met recognition or disclosure criteria were identified.