

Pharma-Bio Serv, Inc.
Form 10-K
January 29, 2013

UNITED STATES

SECURITIES AND EXCHANGE COMMISSION
WASHINGTON, D.C. 20549

FORM 10-K

(Mark One)

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

For the fiscal year ended October 31, 2012

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

For the transition period from _____ to _____

Commission File No. 000-50956

PHARMA-BIO SERV, INC.
(Exact Name of Registrant as Specified in Its Charter)

Delaware
(State or Other Jurisdiction of
Incorporation or Organization)

20-0653570
(IRS Employer Identification No.)

Pharma-Bio Serv Building,
#6 Road 696
Dorado, Puerto Rico
(Address of Principal Executive
Offices)

00646
(Zip Code)

787-278-2709
(Registrant's Telephone Number, Including Area Code)

Securities registered pursuant to Section 12(b) of the Act: None

Securities registered pursuant to Section 12(g) of the Act: Common Stock, par value \$0.0001 per share

Indicate by check mark if the registrant is a well-known seasoned issuer, as defined in Rule 405 of the Securities Act.
Yes No

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the Act. Yes No

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was required to file such reports), and (2) has been subject to such filing requirements for the past 90 days. Yes No

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

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K (§229.405 of this chapter) is not contained herein, and will not be contained, to the best of the registrant's knowledge, in definitive proxy or information statements incorporated by reference in Part III of this Form 10-K or any amendment to this Form 10-K.

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

Large accelerated filer	<input type="checkbox"/>	Accelerated filer	<input type="checkbox"/>
Non-accelerated filer	<input type="checkbox"/>	Smaller reporting company	<input type="checkbox"/>

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

The approximate aggregate market value of common stock held by non-affiliates of the registrant, based on the closing price for the registrant's common stock on April 30, 2012 (the last business day of the second quarter of the registrant's current fiscal year), was \$7,991,615.

The number of shares of the registrant's common stock outstanding as of January 28, 2013 was 20,758,695.

DOCUMENTS INCORPORATED BY REFERENCE

Portions of the registrant's Proxy Statement relative to the 2013 Annual Meeting of Stockholders are incorporated by reference in Part III hereof.

PHARMA-BIO SERV, INC.
FORM 10-K
FOR THE YEAR ENDED OCTOBER 31, 2012

TABLE OF CONTENTS

		Page
PART I		
ITEM 1	BUSINESS	1
ITEM 1A	RISK FACTORS	5
ITEM 1B	UNRESOLVED STAFF COMMENTS	11
ITEM 2	PROPERTIES	11
ITEM 3	LEGAL PROCEEDINGS	11
ITEM 4	MINE SAFETY DISCLOSURES	11
PART II		
ITEM 5	MARKET FOR REGISTRANT’S COMMON EQUITY, RELATED STOCKHOLDER MATTERS AND ISSUER PURCHASES OF EQUITY SECURITIES	12
ITEM 6	SELECTED FINANCIAL DATA	12
ITEM 7	MANAGEMENT’S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS	12
	QUANTITATIVE AND QUALITATIVE DISCLOSURES ABOUT MARKET RISK	19
ITEM 7A	FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA	19
ITEM 8	(See F-1)	19
	CHANGES IN AND DISAGREEMENTS WITH ACCOUNTANTS ON ACCOUNTING AND FINANCIAL DISCLOSURE	19
ITEM 9		19
ITEM 9A	CONTROLS AND PROCEDURES	20
ITEM 9B	OTHER INFORMATION	20
PART III		
ITEM 10	DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE	21
ITEM 11	EXECUTIVE COMPENSATION	21
ITEM 12	SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT AND RELATED STOCKHOLDER MATTERS	21
	CERTAIN RELATIONSHIPS AND RELATED TRANSACTIONS, AND DIRECTOR INDEPENDENCE	21
ITEM 13		21
ITEM 14	PRINCIPAL ACCOUNTING FEES AND SERVICES	21

PART IV

ITEM 15	EXHIBITS, FINANCIAL STATEMENT SCHEDULES	22
SIGNATURES		25
FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA		F-1

PART I

ITEM 1. BUSINESS.

GENERAL

Pharma-Bio Serv, Inc. is a Delaware corporation, organized in 2004 under the name Lawrence Consulting Group, Inc. In February 2006, our corporate name was changed to Pharma-Bio Serv, Inc.

On January 25, 2006, pursuant to an agreement and plan of merger among us, Plaza Acquisition Corp., Pharma-Bio Serv PR, Inc. (then known as Plaza Consulting Group, Inc. and referred to as “Pharma-PR”), and the then sole stockholder of Pharma-PR, Plaza Acquisition Corp. was merged into Pharma-PR, with the result that Pharma-PR became our wholly-owned subsidiary and our sole business became the business of Pharma-PR.

Pharma-PR business was established as a sole proprietorship in 1993 and incorporated in 1997 to offer compliance consulting services to the pharmaceutical industry. The business operations provide services to the pharmaceutical, biotechnology, medical device and chemical manufacturing companies principally in Puerto Rico, the United States and Europe.

Our executive offices are located at Pharma-Bio Serv Building, #6 Road 696, Dorado, Puerto Rico 00646. Our telephone number is (787) 278-2709. The financial information about our reporting segments appear in Note K to our Consolidated Financial Statements included in this Annual Report on Form 10-K.

Our website is www.pharmabioserv.com. Information on our website or any other website is not part of this Annual Report on Form 10-K.

References to “we,” “us,” “our” and similar words in this Annual Report on Form 10-K refer to Pharma-Bio Serv, Inc. and its subsidiaries.

OVERVIEW

We are a compliance and technology transfer services consulting firm with a laboratory testing facility with headquarters in Puerto Rico, servicing the Puerto Rico, United States and Europe markets. The compliance consulting service sector in those markets consists of local compliance and validation consulting firms, United States dedicated validation and compliance consulting firms and large publicly traded and private domestic and foreign engineering and consulting firms. We provide a broad range of compliance related consulting services. We also provide microbiological testing services and chemical testing services through our laboratory testing facility (“Lab”) in Puerto Rico. We also provide information technology consulting services and technical training/seminars, which services are not currently significant to our operating results. We market our services to pharmaceutical, chemical, biotechnology and medical devices, and allied products companies in Puerto Rico, the United States and Europe. Our team includes more than 275 experienced engineering and life science professionals, and includes former quality assurance managers or directors, and experienced and trained professionals with bachelors, masters and doctorate degrees in health sciences and engineering.

We have a well-established and consistent relationship with the major pharmaceutical, biotechnology, medical device and chemical manufacturing companies in Puerto Rico and the United States, which provides us access to affiliated companies in other markets. We seek opportunities in markets that could yield profitable margins using our professional consulting force and also provide services such as those performed by our microbiological testing laboratory facility, our information technology service division, Integratek, and our technical training division,

Pharma Serv Academy.

Integratek provides a variety of information technology services such as web pages and portals development, digital art design, intranets, extranets, software development including database integration, Windows and web applications development, software technical training and learning management systems, technology project management, and compliance consulting services, among others. Our Pharma Serv Academy division, through a network of leading industry professional experts in their field, which include resources of our own, provides technical seminars/training that incorporates the latest regulatory trends and standards as well as other related areas. Although these services are not currently significant to our operating results, our goal is to broaden the portfolio of services that we can provide to our customer base and also target other potential customers in other industries.

We believe the most significant factors to achieving future business growth includes our ability to: (i) continue to provide quality value-added compliance services to our clients; (ii) recruit and retain highly educated and experienced professionals; (iii) further expand our products and services to address the expanding needs of our clients; and (iv) expand our market presence in the United States, Europe and possibly other emerging pharmaceutical markets in order to respond to the international compliance needs of our clients. Our business is affected to the extent the current economic downturn affects the decision of our clients and potential clients to establish operations or to continue or expand their existing operations.

Our revenue is derived from (i) time and materials contracts (representing approximately 95% of total revenues), where the clients are charged for the time, materials and expenses incurred on a particular project or service, (ii) fixed-fee contracts or from “not to exceed” contracts (approximately 2% of total revenues), which are generally short-term contracts, in which the value of the contract cannot exceed a stated amount, and (iii) laboratory testing (representing approximately 3% of total revenues) which generally is completed and certified within days of sample receipt. For time and materials contracts, our revenue is principally a function of the number of resources and the number of hours billed per professional. To the extent that our revenue is based on fixed-fee or “not to exceed” contracts, our ability to operate profitably is dependent upon our ability to estimate accurately the costs that we will incur on a project and to manage and monitor the project. If we underestimate our costs on any contract, we could sustain a loss on the contract or its profitability might be reduced.

The principal components for our consulting costs of services are resource compensation (salaries and wages, independent contractors’ fees, taxes and benefits) and expenses relating to the performance of the services. In order to ensure that our pricing is competitive yet minimize the impact on our margins, we manage increasing labor costs by (i) selecting resources according to our cost for specific projects, (ii) negotiating, where applicable, rates with the resource, (iii) subcontracting labor and (iv) negotiating and passing rate increases to our customers, as applicable. Although this strategy has been successful in the past, we cannot give any assurance that such strategy will continue to be successful. As for our testing laboratory operation, the major costs of services components are salaries and wages, occupancy and depreciation expenses, plus consumable goods usage.

We have established quality systems for our employees which include:

Training Programs - including a Current Good Manufacturing Practices exam prior to recruitment and periodic refreshers;

Recruitment Full Training Program - including employee manual, dress code, time sheets and good project management and control procedures, job descriptions, and firm operating and administration procedures;

Safety Program - including OSHA, Environmental Health and Safety; and

Code of Ethics and Business Conduct - a code of ethics and business conduct is used and enforced as one of the most significant company controls on personal behavior.

In addition, we have implemented procedures to respond to client complaints and customer satisfaction survey procedures. As part of our employee performance appraisal annual process, our clients receive an evaluation form for employee project performance feedback, including compliance with our code of ethics.

BUSINESS STRATEGY AND OBJECTIVES

We are actively pursuing new markets as part of our growth strategy. We have a well-established and consistent relationship with the major pharmaceutical, biotechnology, medical device and chemical manufacturing companies in Puerto Rico and the United States which provides us access to affiliated companies in other markets. We seek opportunities in markets that could yield profitable margins using our professional consulting force and also provide new services such as those performed by our microbiological testing laboratory facility and our information technology service division.

Our business strategy is based on a commitment to provide premium quality and professional consulting services and reliable customer service to our customer base. Our business strategy and objectives are as follow:

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

Continue growth in consulting services in each technical service, quality assurance, regulatory compliance, technology transfer, validation, engineering, laboratory testing and manufacturing departments by achieving greater market penetration from our marketing and sales efforts;

Continue to enhance our technical consulting services through internal growth and acquisitions that provide solutions to our customers' needs;

Motivate our professionals and support staff by implementing a compensation program which includes both individual performance and overall company performance as elements of compensation;

 Create a pleasant corporate culture and emphasize operational quality safety and timely service;

 Continue to maintain our reputation as a trustworthy and highly ethical partner; and

 Efficiently manage our operating and financial costs and expenses.

2006 U.S. Validation Compliance Service Business Acquisition

In January 2006, we acquired a validation compliance service business which serves mainly the United States market. We host our U.S. market expansion plans from this organization.

2007 Entrance to Ireland Market

In September 2007, through the formation of an Irish subsidiary, we entered into the Ireland market. Currently we provide the same consulting services we are currently providing in the Puerto Rico and United States markets.

2008 Integratek Acquisition

In December 2008, we acquired through one of our subsidiaries the operations and assets of Integratek Corp. ("Integratek"), an information technology services and consulting firm based in Puerto Rico. With this acquisition, we broaden the portfolio of services to our customer base and also target other potential customers in other industries.

2009 Laboratory Testing Facility

Our laboratory testing facility ("Lab") located in Puerto Rico, with an investment of \$1.5 million for microbiology, chemical and environmental testing, commenced operations in early fiscal 2009. The Lab is U.S. Food and Drug Administration ("FDA") registered and ISO 9001 certified. The Lab incorporates the latest technology and test methodologies meeting pharmacopoeia industry standards and regulations. It offers testing and related services to our core industries already serviced as well as the cosmetic and food industries.

2012 Minority Controlled Company Certification

In line with the strategy to penetrate the United States market, on September 1, 2012 we obtained the renewal of the certification as a "minority-controlled company" as defined by the National Minority Supplier Development Council and Growth Initiative ("NMSDC"). The certification allows us to participate in corporate diversity programs available from various potential customers in the United States and Puerto Rico. The certification is subject to renewal on September 1, 2013.

2012 Entrance to Spain Market

In January 2012, we initiated a consulting services operation to the Spanish market. Currently, these services are covered under the Irish subsidiary umbrella. Consulting services provided in the Spanish market are similar to those covered in the other markets we serve.

TECHNICAL CONSULTING SERVICES

We have established a reputation as a premier technical consulting services firm to the pharmaceutical, biotechnology, medical device and chemical manufacturing industries in various markets. These services include regulatory compliance, validation, technology transfer, engineering, project management and process support. We have approximately 25 clients that are among the largest pharmaceutical, chemical manufacturing, medical device and biotechnology companies. We are actively participating in exhibitions, conferences, conventions and seminars as either exhibitors, sponsors or conference speakers.

MARKETING

We conduct our marketing activities in Puerto Rico, United States, Europe and other marketplaces. We actively utilize our project managers and leaders who are currently managing consulting service contracts at various client locations to also market consulting and laboratory testing services to their existing and past client relationships. Our senior management is also actively involved in the marketing process, especially in marketing to major accounts. Our senior management and staff also concentrate on developing new business opportunities and focus on the larger customer

accounts (by number of professionals or dollar volume) and responding to prospective customers' requests for proposals.

PRINCIPAL CUSTOMERS

We provide a substantial portion of our services to three customers, each of whom accounted for 10% or more of our revenues in the years ended October 31, 2012 and 2011. During the years ended October 31, 2012 and 2011, these customers accounted for, in the aggregate, 32% and 38% of total revenue, respectively. In spite of the fact that just a few customers represent a significant source of revenue, our functions are not a continuous process, accordingly, the client base for which our services are typically rendered, on a project-by-project basis, changes regularly. Therefore, in any given year a small number of customers could represent a significant source of our revenue for that year. The loss of, or significant reduction in the scope of work performed for any major customer or our inability to replace customers upon completion of contracts could adversely affect our revenue and impair our ability to operate profitably.

COMPETITION

We are engaged in a highly competitive and fragmented industry. Some of our competitors are, on an overall basis, larger than we are or are subsidiaries of larger companies, and therefore may possess greater resources than we do. Furthermore, because the technical professional aspects of our consulting business do not usually require large amounts of capital, there is relative ease of market entry for a new entrant possessing acceptable professional qualifications. Accordingly, we compete with regional, national, and international firms. Within the Puerto Rico, United States and Europe markets, certain competitors, including local competitors, may possess greater resources than we do as well as better access to clients and potential clients.

Competition for validation and consulting services used to be primarily based on reputation, track record, experience, and quality of service. However, given the economic recession and our clients' strategies to reduce costs, price of service has become a major factor in sourcing our services. We believe we enjoy competitive advantages over other consulting service firms because of our historical market share within Puerto Rico (20 years), brand name, reputation and track record with many of the major pharmaceutical, biotechnology, medical device and chemical manufacturing companies which have a presence in the markets we are pursuing.

The market of qualified and experienced professionals that are capable of providing technical consulting services is very competitive and consists primarily of our competitors as well as companies in the pharmaceutical, chemical, biotechnology and medical device industries who are our clients and potential clients. In seeking qualified personnel, we market our name recognition in the Puerto Rico market, our reputation with our clients and salary and benefit packages.

RAW MATERIALS

We require the use of various raw materials, including culture media, DNA reagents, LAL reagents and biological indicators, in our testing laboratory facility. We purchase these raw materials from various suppliers. At times, we concentrate orders among a few suppliers in order to strengthen our supplier relationships and receive quantity discounts. Raw materials are generally available from multiple suppliers at competitive prices, and amounts kept in stock are not significant.

ENVIRONMENTAL REGULATIONS

Activities in our microbiological testing laboratory facility are regulated under Puerto Rico and U.S. federal laws designed to protect workers and the environment. Some of these laws include the Occupational Safety and Health Act and the Resource Conservation and Recovery Act. These laws apply to the use, handling and disposal of various biological and chemical substances used in our processes. We believe we are in material compliance with these laws and that continued compliance will not have a materially adverse effect on our business. No specific accounting for environmental compliance has been maintained or projected by us at this time.

INTELLECTUAL PROPERTY RIGHTS

We have no proprietary software or products. We rely on non-disclosure agreements with our employees to protect the proprietary software and other proprietary information of our clients. Any unauthorized use or disclosure of this information could harm our business.

EMPLOYEES

We employ approximately 220 employees, all of whom are full time employees. None of our employees are represented by a labor union, and we consider our employee relations to be good.

EXECUTIVE OFFICERS

The following table sets forth certain information with respect to our executive officers.

Name	Age	Position
Nélida Plaza	45	Acting President and Chief Executive Officer, President of Puerto Rico Operations and Secretary Chief Financial Officer and Vice President -
Pedro J. Lasanta	53	Finance and Administration

Nélida Plaza has served as our Acting President and Chief Executive Officer since January 1, 2013. She has also served as the Vice President of Operations of Pharma-PR since January 2004, our Secretary since January 25, 2006, and our President of Puerto Rico Operations since December 31, 2009, in charge of Scienza Labs, Pharma Academy and Pharma-PR. Ms. Plaza served as our Vice President from January 25, 2006 to December 31, 2009. In July 2000, Ms. Plaza joined Pharma-PR as a project management consultant. In the past, Ms. Plaza was a Unit Operations Leader and Safety Manager at E.I. DuPont De Nemours where she was in charge of all manufacturing operations and was involved with the development, support and audit of environmental, safety and occupational health programs. Ms. Plaza holds a M.S. in Environmental Management from the University of Houston in Clear Lake and a B.S. in Chemical Engineering from the University of Puerto Rico. Nélida Plaza was recognized by Casiano Communications as one of the 40 under 40 distinguished executives in Puerto Rico.

Pedro J. Lasanta has served as our Chief Financial Officer and Vice President - finance and administration since November 2007. From 2006 until October 2007, Mr. Lasanta was in private practice as an accountant, tax and business counselor. From 1999 until 2006, Mr. Lasanta was the Chief Financial Officer for Pearle Vision Center PR, Inc. In the past, Mr. Lasanta was also an audit manager for Ernst & Young, formerly Arthur Young & Company. He is a cum laude graduate in business administration (accounting) from the University of Puerto Rico. Mr. Lasanta is a Certified Public Accountant. In 2012, he was awarded the Puerto Rico Manufacturers Association (North Region) Service Manager of the Year.

Effective January 1, 2013, Elizabeth Plaza, our President and Chief Executive Officer, stepped down and assumed a new role as Senior Strategic Consultant to the Company for a term of one year, though she continues to serve as the Company's Chairman. Nelida Plaza our President of Puerto Rico Operations has assumed the position as Acting President and Chief Executive Officer for the Company.

ITEM 1A. RISK FACTORS.

This Annual Report on Form 10-K includes “forward-looking statements” within the meaning of Section 21E of the Securities Exchange Act of 1934, as amended, including, in particular, certain statements about our plans, strategies and prospects. Although we believe that our plans, intentions and expectations reflected in or suggested by such forward-looking statements are reasonable, we cannot assure you that such plans, intentions or expectations will be achieved. Important factors that could cause our actual results to differ materially from our forward-looking statements include those set forth in this Risk Factors section.

If any of the following risks, or other risks not presently known to us or that we currently believe to not be significant, develop into actual events, then our business, financial condition, results of operations, cash flows or prospects could be materially adversely affected.

Risks That Relate to our Business

Because our business is concentrated in the pharmaceutical industry in Puerto Rico, the United States and Europe, any changes in that industry or in those markets could impair our ability to generate revenue and realize a profit.

Since most of our business is performed in Puerto Rico, the United States and Europe, for pharmaceutical, biotechnology, medical device and chemical manufacturing companies, our ability to generate revenue and realize a profit could be impaired by factors impacting those markets. For example, changes in tax laws or regulatory, political or economic conditions, which discourage businesses from operating in the markets we serve, which affect the need for services such as those provided by us, could impair our ability to generate revenue and realize a profit.

Puerto Rico government enacted ACT 154 of October 22, 2010 which may adversely affect the willingness of our customers to do business in Puerto Rico and consequently adversely affect our business.

On October 22, 2010, Act No. 154 was enacted by the Puerto Rico government. The act primarily affects the industry we serve and consequently our customer base. Act 154 extends the circumstances under which a nonresident alien individual or a non resident corporation or partnership can be treated as doing business in Puerto Rico and is deriving income from sources within Puerto Rico for purposes of income tax. It also provides for the imposition of a temporary excise tax on some acquisitions by non-resident individuals, corporations or partnerships, of products totally or partially manufactured or produced in Puerto Rico and of related services to said products of affiliated entities with the buyer. It basically adopts a modified income sourcing rule and a temporary excise tax that will be enforced for a period of six (6) years and will decrease gradually during this time.

The impact of the act, if any, over the industry and its willingness to do business in Puerto Rico continues to be uncertain. Consequently, our ability to generate revenue in Puerto Rico may be impaired.

Changes in tax benefits may affect the willingness of companies to continue or expand their operations in Puerto Rico.

Until 1996, the Internal Revenue Code provided certain tax benefits to pharmaceutical companies operating in Puerto Rico by enabling their Puerto Rico operations to operate free from federal income taxes. Partly as a result of the tax benefits, numerous pharmaceutical companies established facilities in Puerto Rico. In 1996, this tax benefit was eliminated, although companies that had facilities in Puerto Rico could continue to receive these benefits for ten years, at which time the benefits were set to expire. In order to promote business activities in Puerto Rico, in May 2008 the Puerto Rico government enacted a tax incentive law ("Act 73"). Act 73 provides tax exemption from various taxes, including income tax, and investment credits for activities similar to those of our customers and our company. The change in the tax laws may affect favorably or unfavorably the willingness of pharmaceutical companies to continue or to expand their Puerto Rico operations. To the extent that pharmaceutical companies choose to develop and manufacture products outside of Puerto Rico, our ability to generate new business may be adversely impaired.

Puerto Rico's economy, including its governmental financial crisis, may affect the willingness of businesses to commence or expand operations in Puerto Rico.

As a result of Puerto Rico's governmental financial crisis, businesses may be reluctant to establish or expand their operations in Puerto Rico. Further, since Puerto Rico's economy is petroleum-based, the fluctuating price of oil, combined with Puerto Rico's high level of debt, may make Puerto Rico a less attractive place to expand existing operations or commence new business activities. To the extent that companies in the pharmaceutical and related industries decide not to commence new operations or not to expand their existing operations in Puerto Rico, the market for our services may decline.

Other factors, including economic factors, may affect the decision of businesses to continue or expand their operations in the markets we serve.

Companies in the pharmaceutical and related industries for which we perform service are subject to economic pressures, which affect their global operations and which may influence the decision to reduce or increase the scope of their operations in the markets we serve. These companies consider a wide range of factors in making such a decision, and may be influenced by a need to consolidate operations, to reduce expenses, to increase their business in geographical regions where there are large customer bases, tax, regulatory and political considerations and many other factors. We cannot assure you that our customers and potential customers will not make extensive reductions or terminate their operations in the markets we serve entirely, which could significantly impair our ability to generate revenue.

Because our business is dependent upon a small number of clients, the loss of a major client could impair our ability to operate profitably.

Our business has been dependent upon a small number of clients. During the years ended October 31, 2012 and 2011, a very small number of clients accounted for a disproportionately large percentage of our revenue. In the years ended October 31, 2012 and 2011, three customers accounted for, in aggregate, approximately 32% and 38% of total revenue, respectively.

The loss of, or significant reduction in the scope of work performed for, or any significant change in the financial terms related to, any major customer, could impair our ability to operate profitably. We cannot assure that we will not sustain significant decreases in revenue from our major customers or that we will be able to replace any major customers or the resulting decline in revenue.

Customer procurement and sourcing practices intended to reduce costs could have an adverse affect on our margins and profitability.

In an effort to reduce their costs, many of our customers are establishing or extending the scope of their procurement departments to include consulting and project management services, such as ours. As a result, we have less interaction with the end user of our services (typically labs or production units) when bidding on a project, which we believe decreases the focus on the quality of service provided and increases the emphasis on cost of the service. This may cause us to lower the price of our bids, which would reduce the margins in a given project. Also, some customers have established vendor management/vendor neutral-programs with third-parties (some of whom are also our competitors). Because these vendor management programs may receive a percentage of our fees, without a corresponding increase in the fee itself, our margins would decline. In addition, where a vendor management program is a competitor for a particular service we provide, we may have difficulty securing that particular project, which would adversely impact revenue. Some of these vendor neutral programs intend to limit our interaction with our direct end user, and our interaction is limited to the representative of the vendor neutral agency. This limitation impairs our ability to establish and maintain our relationships with our customers and recognition of the value added in the service.

Since our business is dependent upon the development and enhancement of patented pharmaceutical products or processes by our clients, the failure of our clients to obtain and maintain patents could impair our ability to operate profitably.

Companies in the pharmaceutical industry are highly dependent on their ability to obtain and maintain patents for their products or processes. We are aware of some pharmaceutical companies with operations in Puerto Rico whose patent rights may expire in the near future. The inability to obtain new patents and the expiration of active patents may reduce the need for our services and thereby impair our ability to operate profitably.

We may be unable to pass on increased labor costs to our clients.

The principal components of our cost of revenues are employee compensation (salaries, wages, taxes and benefits) and expenses relating to the performance of the services we provide. We face increasing labor costs which we seek to pass on to our customers through increases in our rates. To remain competitive, we may not be able to pass these increased costs on to our clients, and, to the extent that we are not able to pass these increased costs on to our clients, our gross margin will be reduced.

Consolidation in the pharmaceutical industry may have a harmful effect on our business.

In recent years, the pharmaceutical industry has undergone consolidation, and may in the future undergo further substantial consolidation which may reduce the number of our existing and potential customers. The consolidation in the pharmaceutical industry may have a harmful effect on our business and or ability to maintain and replace customers.

Because the pharmaceutical industry is subject to government regulations, changes in government regulations relating to this industry may affect the need for our services.

Because government regulations affect all aspects of the pharmaceutical, biotechnology, medical device and chemical manufacturing industries, including regulations relating to the testing and manufacturing of pharmaceutical products and the disposal of materials which are or may be considered toxic, any change in government regulations could have a profound effect upon not only these companies but companies, such as ours, that provide services to these industries. If we are not able to adapt and provide necessary services to meet the requirements of these companies in response to changes in government regulations, our ability to generate business may be impaired.

Our reputation and divisions may be impacted by regulatory standards impacting our customer products.

We provide microbiological and chemical testing services to our customer products that are distributed worldwide. Any concerns regarding our customer products complying with regulatory standards could result in inquiries regarding our laboratory or testing services. As a result, we may need to participate in litigation or regulatory audits requiring the investment of significant time and funds. In addition, the reputation of our Lab and technical services, as well as our technical expertise, might suffer. Furthermore, any such inquiries or any adverse impact of such inquiries may financially impact our Lab and other divisions, including our consulting division.

If we are unable to protect our clients' intellectual property, our ability to generate business will be impaired.

Our services either require us to develop intellectual property for clients or provide our personnel with access to our clients' intellectual property. Because of the highly competitive nature of the pharmaceutical, biotechnology, medical device and chemical manufacturing industries and the sensitivity of our clients' intellectual property rights, our ability to generate business would be impaired if we fail to protect those rights. Although all of our employees and contractors are required to sign non-disclosure agreements, any disclosure of a client's intellectual property by an employee or contractor may subject us to litigation and may impair our ability to generate business either from the affected client or other potential clients. In addition, we are required to enter into confidentiality agreements and our failure to protect the confidential information of our clients may impair our business relationship.

We may be subject to liability if our services or solutions for our clients infringe upon the intellectual property rights of others.

It is possible that in performing services for our clients, we may inadvertently infringe upon the intellectual property rights of others. In such event, the owner of the intellectual property may commence litigation seeking damages and an injunction against both us and our client, and the client may bring a claim against us. Any infringement litigation would be costly, regardless of whether we ultimately prevail. Even if we prevail, we will incur significant expenses and our reputation would be hurt, which would affect our ability to generate business and the terms on which we would be engaged, if at all.

We may be held liable for the actions of our employees or contractors when on assignment.

We may be exposed to liability for actions taken by our employees or contractors while on assignment, such as damages caused by their errors, misuse of client proprietary information or theft of client property. Due to the nature of our assignments, we cannot assure you that we will not be exposed to liability as a result of our employees or contractors being on assignment.

To the extent that we perform services pursuant to fixed-price or incentive-based contracts, our cost of services may exceed our revenue on the contract.

Some of our revenue is derived from fixed-price contracts. Our costs of services may exceed revenue of these contracts if we do not accurately estimate the time and complexity of an engagement. Further, we are seeking contracts by which our compensation is based on specified performance objectives, such as the realization of cost savings, quality improvements or other performance objectives. Our failure to achieve these objectives would reduce our revenue and could impair our ability to operate profitably.

Our profit margin is largely a function of the rates we are able to charge and collect for our services and the utilization rate of our professionals. Accordingly, if we are not able to maintain our pricing for our services or an appropriate utilization rate for our professionals without corresponding cost reductions, our profit margin and profitability will suffer. The rates we are able to charge for our services are affected by a number of factors, including:

Our clients' perception of our ability to add value through our services;

Our ability to complete projects on time;

Pricing policies of competitors;

Our ability to accurately estimate, attain and sustain engagement revenues, margins and cash flows over increasingly longer contract periods; and

General economic and political conditions.

Our utilization rates are also affected by a number of factors, including:

Our ability to move employees and contractors from completed projects to new engagements; and

Our ability to manage attrition of our employees and contractors.

Because most of our contracts may be terminated on little or no advance notice, our failure to generate new business could impair our ability to operate profitably.

Most of our contracts can be terminated by our clients with little or no advance notice. Our clients typically retain us on a non-exclusive, engagement-by-engagement basis, and the client may terminate, cancel or delay any engagement or the project for which we are engaged, at any time and on no advance notice. As a result, the termination, cancellation, expiration or delay of contracts could have a significant impact on our ability to operate profitably.

Because of the competitive nature of the pharmaceutical, biotechnology, medical device and chemical manufacturing consulting market, we may not be able to compete effectively if we cannot efficiently respond to changes in the structure of the market and developments in technology.

Because of recent consolidations in the pharmaceutical, biotechnology, medical device and chemical manufacturing consulting business, we are faced with an increasing number of larger companies that offer a wider range of services and have better access to capital than we have. We believe that larger and better-capitalized competitors have enhanced abilities to compete for both clients and skilled professionals. In addition, one or more of our competitors may develop and implement methodologies that result in superior productivity and price reductions without adversely affecting their profit margins. We cannot assure you that we will be able to compete effectively in an increasingly competitive market.

Because we are dependent upon our management, our ability to develop our business may be impaired if we are not able to engage skilled personnel.

Our success in the past has depended in large part on the skills and efforts of Elizabeth Plaza, our former President, Chief Executive Officer and founder. Effective January 1, 2013, Elizabeth Plaza entered into a new role with the Company as Senior Strategic Consultant for a term of one year. Nelida Plaza, our President of Puerto Rico Operations, assumed the position as Acting President and Chief Executive Officer for the Company. It is uncertain if this change

will have a material adverse effect on the development and success of our business. In addition, although we have contracts with Elizabeth Plaza and Nelida Plaza, these agreements do not guarantee that they will continue to be employed by us.

Our future success will depend in part upon our ability to attract and retain additional qualified management and technical personnel. Competition for such personnel is intense and we compete for qualified personnel with numerous other employers, including consulting firms, some of which have greater resources than we have, as well as pharmaceutical companies, most of which have significantly greater financial and other resources than we do. We may experience increased costs in order to retain and attract skilled employees. Our failure to attract additional personnel or to retain the services of key personnel and independent contractors could have a material adverse effect on our ability to operate profitably.

We may not be able to continue to grow unless we consummate acquisitions or enter markets outside of Puerto Rico, the United States, Ireland and Spain.

An important part of our growth strategy is (i) to acquire other businesses which can increase the range of services and products that we can offer and (ii) to establish offices in places where we do not presently operate, either by acquisition or by internal growth. If we fail to make any acquisitions or otherwise expand our business, our future growth may be limited. The success in any market will be dependent on such factors as regulatory, tax, political or economic conditions, our abilities to penetrate the market, hire qualified personnel in a timely manner, obtain and maintain reasonable labor costs, generate service revenue volume and profitable margins.

Any acquisitions we make may be made with cash or our securities or a combination of cash and securities. To the extent that we require cash, we may have to borrow the funds or sell equity securities. We have no commitments from any financing source and we may not be able to raise any cash necessary to complete an acquisition. If we seek to expand our business internally, we will incur significant start-up expenses without any assurance of our ability to penetrate the market.

If we make any acquisitions, they may disrupt or have a negative impact on our business.

If we make acquisitions or establish operations in locales outside of Puerto Rico, we could have difficulty integrating the acquired companies' personnel and operations with our own. In addition, the key personnel of the acquired business may not be willing to work for us. We cannot predict the effect an expansion may have on our core business. Regardless of whether we are successful in making an acquisition, the negotiations could disrupt our ongoing business, distract our management and employees and increase our expenses. In addition to the risks described above, acquisitions are accompanied by a number of inherent risks, including, without limitation, the following:

the difficulty of integrating acquired products, services or operations;

the potential disruption of the ongoing businesses and distraction of our management and the management of acquired companies;

the potential loss of contracts from clients of acquired companies;

the difficulty of maintaining profitability due to increased labor and expenses from acquired companies;

difficulties in complying with regulations in other countries that relate to both the pharmaceutical or other industries to which we provide services as well as our own operations;

difficulties in maintaining uniform standards, controls, procedures and policies;

the potential impairment of relationships with employees and customers as a result of any integration of new management personnel;

the potential inability or failure to achieve additional sales and enhance our customer base through cross-marketing of the products to new and existing customers;

the effect of any government regulations which relate to the business acquired;

potential unknown liabilities associated with acquired businesses or product lines, or the need to spend significant amounts to retool, reposition or modify the marketing and sales of acquired products or the defense of any litigation, whether of not successful, resulting from actions of the acquired company prior to our acquisition;

difficulties in disposing of the excess or idle facilities of an acquired company or business and expenses in maintaining such facilities; and

potential expenses under the labor, environmental and other laws of other countries.

Our business could be severely impaired if and to the extent that we are unable to succeed in addressing any of these risks or other problems encountered in connection with an acquisition. Further, the commencement of business in locales where we have no current operations may be subject to additional significant risks.

Risks Concerning our Securities

Because there is a limited market in our common stock, stockholders may have difficulty in selling our common stock and our common stock may be subject to significant price swings.

There is a very limited market for our common stock. Since trading commenced in December 2006, there has been little activity in our common stock and on some days there is no trading in our common stock. Because of the limited market for our common stock, the purchase or sale of a relatively small number of shares may have an exaggerated effect on the market price for our common stock. We cannot assure stockholders that they will be able to sell common stock or, that if they are able to sell their shares, that they will be able to sell the shares in any significant quantity at the quoted price.

Our revenues, operating results and profitability will vary from quarter to quarter, which may result in increased volatility of our stock price.

Our quarterly revenues, operating results and profitability have varied in the past and are likely to vary significantly from quarter to quarter, making them difficult to predict. This may lead to volatility in our share price. The factors that are likely to cause these variations are:

Seasonality, including number of workdays and holiday and summer vacations;

The business decisions of clients regarding the use of our services;

Periodic differences between clients' estimated and actual levels of business activity associated with ongoing engagements, including the delay, reduction in scope and cancellation of projects;

The stage of completion of existing projects and their termination;

Our ability to move employees quickly from completed projects to new engagements and our ability to replace completed contracts with new contracts with the same clients or other clients;

The introduction of new services by us or our competitors;

Changes in pricing policies by us or our competitors;

Our ability to manage costs, including personnel compensation, support-services and severance costs;

Acquisition and integration costs related to possible acquisitions of other businesses;

Changes in estimates, accruals and payments of variable compensation to our employees or contractors; and

Global economic and political conditions and related risks, including acts of terrorism.

The issuance of securities, whether in connection with an acquisition or otherwise, may result in significant dilution to our stockholders.

If we are required to issue securities either as payment of all or a portion of the purchase price of an acquisition or in order to obtain financing for the acquisition or for other corporate purposes could result in dilution to our stockholders. The amount of such dilution will be dependent upon the terms on which we issue securities. The issuance of securities at a price which is less than the exercise price of warrants or the conversion price of securities could result in additional dilution if we are required to reduce the exercise price or conversion price of the then outstanding options or warrants or other convertible securities.

ITEM 1B. UNRESOLVED STAFF COMMENTS.

Not applicable.

ITEM 2. PROPERTIES.

In February 2012, the Company automatically renewed a lease agreement for our main resource facilities in Dorado, Puerto Rico with an affiliate of Elizabeth Plaza, our Chairman of the Board. These facilities accommodate our testing laboratory, our customer-specialized training facilities, and our Puerto Rico consulting and headquarters offices. The renewal is for a term of five years with monthly rental payments of \$23,930, \$25,127, \$26,383, \$27,702 and \$29,087 for each of the five years remaining under the lease. The agreement also requires the payment of utilities, property taxes, insurance and a portion of expenses incurred by the affiliate in connection with the maintenance of common areas.

In November 2011, the Company entered into a lease agreement for the U.S. office facilities located in Plymouth, Pennsylvania. The lease is for a five-year term with monthly rental payments of \$6,282 for the first three years. Thereafter, the lease will increase four percent every year, including a five-year renewal option, if executed.

The Company maintains office facilities in Cork, Ireland and Madrid, Spain. Both facilities are under month-to-month leases with monthly payments of approximately \$900 and \$1,200, respectively.

We believe that our present facilities are adequate to meet our needs and that, if we require additional space, it will be available on commercially reasonable terms.

ITEM 3. LEGAL PROCEEDINGS.

From time to time, we may be a party to legal proceedings incidental to our business. We do not believe that there are any proceedings threatened or pending against us, which, if determined adversely to us, would have a material effect on our financial position or results of operations and cash flows.

ITEM 4. MINE SAFETY DISCLOSURES

Not applicable.

PART II

ITEM 5. MARKET FOR REGISTRANT'S COMMON EQUITY, RELATED STOCKHOLDER MATTERS AND ISSUER PURCHASES OF EQUITY SECURITIES.

Our common stock has been quoted on the Over the Counter Bulletin Board under the trading symbol PBSV since December 4, 2006. The table below presents the closing high and low bid prices for our common stock for each quarter during the two most recent fiscal years. These prices reflect inter-dealer prices, without retail markup, markdown, or commission, and may not represent actual transactions.

Quarter Ending	High Bid	Low Bid
January 31, 2011	\$ 0.35	\$ 0.26
April 30, 2011	0.40	0.30
July 31, 2011	0.43	0.20
October 31, 2011	0.78	0.37
January 31, 2012	0.78	0.65
April 30, 2012	0.90	0.64
July 31, 2012	0.89	0.65
October 31, 2012	0.93	0.76

On January 25, 2013, the closing price of our common stock on the Over the Counter Bulletin Board was \$0.83 per share and there were approximately 75 holders of record of our common stock.

Prior to the acquisition of Pharma-PR in 2006, Pharma-PR was taxed as an N Corporation under the Puerto Rico Internal Revenue Code, which is similar to that of an S Corporation under the Internal Revenue Code. As a result, all of the income from Pharma-PR was taxed to our then sole stockholder. Other than the distributions to our then sole stockholder which were made during the period that we were an N Corporation, we have not paid dividends on our common stock. We plan to retain future earnings, if any, for use in our business. We do not anticipate paying dividends on our common stock in the foreseeable future.

Equity Compensation Plan Information

The following table summarizes the equity compensation plans under which our securities may be issued as of October 31, 2012.

Plan Category	Number of securities to be issued upon exercise of outstanding options and warrants	Weighted-average exercise price per share of outstanding options and warrants	Number of securities remaining available for future issuance under equity compensation plans
Equity compensation plans approved by security holders	1,740,000	\$ 0.6965	760,000
E Equity compensation plans not approved by security holders	1,830,991	\$ 0.0600	16,500

The securities issuable pursuant to the equity plan that was approved by security holders is the 2005 long-term incentive plan, which was approved by stockholders in April 2006, and amended by stockholder approval in April 2007.

The equity compensation plans not approved by security holders are (i) warrants to purchase 1,830,991 shares of common stock issued to San Juan Holdings for services relating to the acquisition of Pharma-PR and (ii) approximately 16,500 shares of common stock underlying options issuable to employees.

ITEM 6. SELECTED FINANCIAL DATA.

Not Applicable.

ITEM 7. MANAGEMENT'S DISCUSSION AND ANALYSIS OF FINANCIAL CONDITION AND RESULTS OF OPERATIONS.

The following discussion of our results of operations and financial condition should be read in conjunction with Part I, including matters set forth in the "Risk Factors" section of this Annual Report on Form 10-K, and our Consolidated Financial Statements and notes thereto included elsewhere in this Annual Report on Form 10-K.

Overview

We are a compliance services consulting firm with a laboratory testing facility with headquarters in Puerto Rico, servicing the Puerto Rico, United States and Europe markets. The compliance consulting service sector in those markets consists of local compliance and validation consulting firms, United States dedicated validation and compliance consulting firms and large publicly traded and private domestic and foreign engineering and consulting firms. We provide a broad range of compliance related consulting services. We also provide microbiological testing services and chemical testing services through our laboratory testing facility (“Lab”) in Puerto Rico. We also provide information technology consulting services and technical training/seminars, which services are not currently significant to our operating results. We market our services to pharmaceutical, chemical, biotechnology and medical devices, and allied products companies in Puerto Rico, the United States and Europe. Our team includes more than 275 experienced engineering and life science professionals, and includes former quality assurance managers or directors, and experienced and trained professionals with bachelors, masters and doctorate degrees in health sciences and engineering.

We actively operate in Puerto Rico, the United States, Ireland and Spain and pursue to further expand these markets by strengthening our business development infrastructure and by constantly realigning our business strategies as new opportunities and challenges arise.

We market our services with an active presence in industry trade shows, professional conventions, industry publications and company provided seminars to the industry. Our senior management is also actively involved in the marketing process, especially in marketing to major accounts. Our senior management and staff also concentrate on developing new business opportunities and focus on the larger customer accounts (by number of professionals or dollar volume) and responding to prospective customers’ requests for proposals.

While our core business is FDA and international agencies regulatory compliance related services, we feel that our clients are in need of other services that we can provide and allow us to present the company as a global solution provider with a portfolio of integrated services that will bring value added solutions to our customers. Accordingly, our portfolio of services include a laboratory testing facility, an information technology consulting practice and a training center that provides seminars/training to the industry.

The Lab incorporates the latest technology and test methodologies meeting pharmacopoeia industry standards and regulations. It currently offers services to our core industries already serviced as well as the cosmetic and food industries.

We also provide technical seminars/training that incorporate the latest regulatory trends and standards as well as other related areas. A network of leading industry professional experts in their field, which include resources of our own, provide these seminars/training to the industry through our “Pharma Serv Academy” division. These services are provided in the markets we currently serve, as well as others, and position our Company as a key leader in the industry.

Our information technology services and consulting division based in Puerto Rico (“Integratek”) provide a variety of information technology services such as web pages and portals development, digital art design, intranets, extranets, software development including database integration, Windows and web applications development, software technical training and learning management systems, technology project management, and compliance consulting services, among others. Integratek is a Microsoft Certified Partner and a reseller for technology products from leading vendors in the market.

In line with the strategy to further penetrate the United States and Puerto Rico markets, we submit annually for renewal the certification as a "minority-controlled company" as defined by the National Minority Supplier Development Council and Growth Initiative ("NMSDC"). This certification allows us to participate in corporate diversity programs available from various potential customers in the United States and Puerto Rico.

In June 2011, Pharma-Bio, Pharma-PR and Pharma-Serv obtained a new Grant of Industrial Tax Exemption pursuant to the terms and conditions set forth in Act No. 73 of May 28, 2008 ("the Grant") issued by the Puerto Rico Industrial Development Company ("PRIDCO"). The Grant provides relief on various Puerto Rico taxes, including income tax, with certain limitations for most of the activities carried on within Puerto Rico, including those that are for services to parties located outside of Puerto Rico.

Industry consolidations, the pharmaceutical regulatory environment, changes in tax laws, customers' price sensitive procurement processes, and the local and global economies recession continue to be factors and uncertainties that affect our business. As such, we are constantly realigning our business strategies as new opportunities and challenges arise.

We have been able to capitalize on the related challenges and opportunities in the Puerto Rico and United States Consulting markets, in which revenues for the year ended October 31, 2012, increased by \$6.2 million and \$3.3 million, respectively, in these markets as compared to last year. Accordingly, we have aligned and increased our business development and operations support in these markets to follow the favorable revenue trend in the consulting business. In order to diversify the European market, in January 2012, we opened an office in Spain. Our operations in Spain generated \$0.4 million in revenues since January 2012, which was offset by our operations in Ireland which suffered a \$0.5 million loss in revenue and 7% gross margin decline, compared to last year. The revenue decline in Ireland is mainly attributable to contract renegotiations during the first quarter of 2012 with the Ireland division's main customer. Other Company divisions realized minor revenue gains or remained constant, compared to last year. In addition, we continue our efforts to broaden the Lab's customer base.

Our total net revenues for the year ended in October 31, 2012 increased by approximately \$9.3 million, or 46.6%, when compared to last year. The revenue growth, offset by the increase in operational support expenses, and the income tax savings related to the Puerto Rico Grant, resulted in net income of approximately \$4.7 million for the year ended October 31, 2012, an increase of \$1.5 million, or 48.4%, when compared with last year.

The following table sets forth information as to our revenue for the years ended October 31, 2012 and 2011, by geographic regions (dollars in thousands).

Revenues by Region	Year ended October 31,			
	2012		2011	
Puerto Rico	\$ 16,856	57.7%	\$ 10,743	53.9%
United States	9,159	31.3%	5,868	29.4%
Europe	3,212	11.0%	3,322	16.7%
	\$ 29,227	100.0%	\$ 19,933	100.0%

Weak economies where we do business and worldwide industry consolidations will continue to be unfavorable factors going forward. These factors, and the impact on the industry, if any, of the recently enacted U.S. health care reform (Patient Protection and Affordable Care Act) and Puerto Rico Act 154 which imposed temporary excise taxes to the industry we serve, remain as industry uncertainties that might adversely affect our future performance. We believe that our future profitability and liquidity will be highly dependent on the effect the global economy, changes in tax laws and worldwide lifescience manufacturing industry consolidations will have over our operations, and our ability to seek service opportunities and adapt to the current industry trends.

Effective January 1, 2013, Elizabeth Plaza, our President and Chief Executive Officer, stepped down and assumed a new role as Senior Strategic Consultant to the Company for a term of one year, though she continues to serve as the Company's Chairman. Nelida Plaza our President of Puerto Rico Operations has assumed the position as Acting President and Chief Executive Officer for the Company.

Results of Operations

The following table sets forth our statements of operations for the years ended October 31, 2012 and 2011, (dollars in thousands) and as a percentage of revenue:

	Year ended October 31,			
	2012		2011	
Revenues	\$ 29,227	100.0%	\$ 19,933	100.0%
Cost of services	19,355	66.2%	13,072	65.6%
Gross profit	9,872	33.8%	6,861	34.4%
Selling, general and administrative expenses	4,138	14.2%	3,409	17.1%
Other income, net	22	0.1%	12	0.0%
Income before income taxes	5,756	19.7%	3,464	17.3%
Income tax expense	1,069	3.7%	305	1.5%
Net income	4,687	16.0%	3,159	15.8%

Revenues. Revenues for the year ended October 31, 2012 were \$29.2 million, an increase of approximately \$9.6 million, or 46.6%, when compared to last year. This improvement is mainly attributable to \$6.2 and \$3.3 million gain in the Puerto Rico and United States consulting markets, respectively. The additional increase is distributed almost evenly between the European market and the other divisions.

Service revenue for the Ireland and Integratek operations are mostly attributable to volume acquired from one customer in each of these divisions.

Cost of Services; gross profit. The overall gross profit for the year ended in October 31, 2012 reflected a gross profit net decrease of 0.6 percentage points, when compared to last year. The net decrease is mostly driven by the loss in the Ireland operation gross profit, Lab's low gross profit yield as a function of billings versus fixed costs of services, offset by project gains attained in the consulting business (mostly influenced by major customers).

Selling, General and Administrative Expenses. Selling, general and administrative expenses for the year ended in October 31, 2012 were approximately \$4.1 million, a net increase in expenses of approximately \$0.7 million as compared to last year. Business development and operations support expenses were increased to follow the favorable revenue trend in the consulting business.

Income Taxes Expense. Savings on the effective income tax rate are attributable to the Puerto Rico tax grant, which significantly reduced the effective income tax rates for the Puerto Rico operation. In addition, the tax Grant triggered favorable non-recurring adjustments booked in the year ended in October 31, 2011 in the aggregate amount of \$0.2 million which is attributable to the year ended October 31, 2010.

Net Income. Our net income for year ended October 31, 2012 was approximately \$4.7 million, an increase of \$1.5 million, or an increase of 48.4%, when compared to last year.

For the year ended October 31, 2012, earnings per common share basic and diluted were \$0.226 and \$0.204, respectively, an increase of \$0.074 and \$0.064 per share, respectively, when compared to last year.

The non-recurring adjustment booked in the year ended October 31, 2011, related to the adoption of the Act 73 Grant, had the aggregate effect of increasing earnings per share basic and diluted by \$0.010.

Our net income improvement is attributable mainly to the increase in overall gross profit, the savings obtained by the Grant, offset by the increase in selling general and administrative expenses to support the favorable revenue trend.

Liquidity and Capital Resources

Liquidity is a measure of our ability to meet potential cash requirements, including planned capital expenditures. For the year ended October 31, 2012, we generated a working capital increase of approximately \$4.7 million.

Our primary cash needs consist of the payment of compensation to our professional staff, overhead expenses, and statutory taxes. Management believes that based on the current level of operations and cash flows from operations, the collectibility of high quality customer receivables will be sufficient to fund anticipated expenses and satisfy other possible long-term contractual commitments for the next twelve months.

To the extent that we pursue possible opportunities to expand our operations, either by acquisition or by the establishment of operations in a new locale, we will incur additional overhead, and there may be a delay between the period we commence operations and our generation of net cash flow from operations.

While uncertainties relating to the current local and global economic condition, competition, the industries and geographical regions served by us and other regulatory matters exist within the consulting services industry, as described above, management is not aware of any trends or events likely to have a material adverse effect on liquidity or its financial statements.

Off-Balance Sheet Arrangements

We were not involved in any significant off-balance sheet arrangements during the fiscal year ended October 31, 2012.

Critical Accounting Policies and Estimates

The discussion and analysis of our financial condition and results of operations are based upon our consolidated financial statements, which have been prepared in accordance with generally accepted accounting principles (“GAAP”) in the United States. We believe the following are the critical accounting policies that impact the consolidated financial statements, some of which are based on management’s best estimates available at the time of preparation. Actual experience may differ from these estimates.

Consolidation - The accompanying consolidated financial statements include the accounts of all of our wholly owned subsidiaries. All intercompany transactions and balances have been eliminated in consolidation.

Use of Estimates - The preparation of consolidated financial statements in conformity with accounting principles generally accepted in the United States requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the consolidated financial statements and the reported amounts of revenues and expenses during the reporting periods. Actual results may differ from these estimates.

Fair Value of Financial Instruments - Accounting standards have established a fair value hierarchy that requires an entity to maximize the use of observable inputs and minimize the use of unobservable inputs when measuring fair value. A financial instrument's categorization within the fair value hierarchy is based upon the lowest level of input that is significant to the fair value measurement. Accounting standards have established three levels of inputs that may be used to measure fair value:

Level 1: Quoted prices in active markets for identical assets and liabilities.

Level 2: Observable inputs other than Level 1 prices such as quoted prices for similar assets or liabilities, quoted prices in markets with insufficient volume or infrequent transactions (less active markets), or model-derived valuations in which all significant inputs are observable or can be derived principally from or corroborated by observable market data for substantially the full term of the assets or liabilities.

Level 3: Prices or valuation techniques that require inputs that are both significant to the fair value measurement and unobservable (supported by little or no market activity).

Marketable securities consist of an obligation from the Puerto Rico Government Development Bank valued using quoted market prices in active markets with no valuation adjustment. Accordingly, this security is categorized in Level 1.

The carrying value of the Company's financial instruments (excluding marketable securities and obligations under capital leases), cash and cash equivalents, accounts receivable, accounts payable and accrued liabilities, are considered reasonable estimates of fair value due to their liquidity or short-term nature. Management believes, based on current rates, that the fair value of its obligations under capital leases approximates the carrying amount.

Revenue Recognition - Revenue is primarily derived from: (1) time and materials contracts (representing approximately 95% of total revenues), which is recognized by applying the proportional performance model, whereby revenue is recognized as performance occurs, (2) short-term fixed-fee contracts or "not to exceed" contracts (representing approximately 2% of total revenues), which revenue is recognized similarly, except that certain milestones also have to be reached before revenue is recognized, and (3) laboratory testing revenue (representing approximately 3% of total revenues) which is mainly recognized as the testing is completed and certified (normally within days of sample receipt from customer). If we determine that a contract will result in a loss, we recognize the estimated loss in the period in which such determination is made.

Cash Equivalents - For purposes of the consolidated statements of cash flows, cash equivalents include investments in a money market obligations trust that is registered under the U.S. Investment Company Act of 1940 and liquid investments with original maturities of three months or less.

Marketable Securities - We consider our marketable security investment portfolio and marketable equity investments available-for-sale and, accordingly, these investments are recorded at fair value with unrealized gains and losses generally recorded in other comprehensive income; whereas realized gains and losses are included in earnings and determined based on the specific identification method.

Accounts Receivable - Accounts receivable are recorded at their estimated realizable value. Accounts are deemed past due when payment has not been received within the stated time period. Our policy is to review individual past due amounts periodically and write off amounts for which all collection efforts are deemed to have been exhausted. Due to the nature of our customers, bad debts are mainly accounted for using the direct write-off method whereby an expense is recognized only when a specific account is determined to be uncollectible. The effect of using this method

approximates that of the allowance method.

Income Taxes - We follow an asset and liability approach method of accounting for income taxes. This method measures deferred income taxes by applying enacted statutory rates in effect at the balance sheet date to the differences between the tax basis of assets and liabilities and their reported amounts on the financial statements. The resulting deferred tax assets or liabilities are adjusted to reflect changes in tax laws as they occur. A valuation allowance is provided when it is more likely than not that a deferred tax asset will not be realized.

The Company follows guidance from the Financial Accounting Standards Board (“FASB”) related to Accounting for Uncertainty in Income Taxes, which includes a two-step approach to recognizing, de-recognizing and measuring uncertain tax positions. As of October 31, 2012, the Company had no significant uncertain tax positions that would be reduced as a result of a lapse of the applicable statute of limitations.

Property and equipment - Owned property and equipment, and leasehold improvements are stated at cost. Equipment and vehicles under capital leases are stated at the lower of fair market value or net present value of the minimum lease payments at the inception of the leases.

Depreciation and amortization of owned assets are provided for, when placed in service, in amounts sufficient to relate the cost of depreciable assets to operations over their estimated service lives, using straight-line basis. Assets under capital leases and leasehold improvements are amortized, over the shorter of the estimated useful lives of the assets or lease term. Major renewals and betterments that extend the life of the assets are capitalized, while expenditures for repairs and maintenance are expensed when incurred.

We evaluate for impairment our long-lived assets to be held and used, and long-lived assets to be disposed of, whenever events or changes in circumstances indicate that the carrying amount of an asset may not be recoverable. Based on management estimates, no impairment of the operating properties was present.

Intangible assets - Definite-lived intangible assets, such as customer lists and covenants not to compete, are amortized on a straight-line basis over their estimated useful lives. We continually evaluate the reasonableness of the useful lives of these assets.

Stock-based Compensation - Stock-based compensation expense is recognized in the consolidated financial statements based on the fair value of the awards granted. Stock-based compensation cost is measured at the grant date based on the fair value of the award and is recognized as expense over the requisite service period, which generally represents the vesting period, and includes an estimate of awards that will be forfeited. We calculate the fair value of stock options using the Black-Scholes option-pricing model at grant date. Excess tax benefits related to stock-based compensation are reflected as cash flows from financing activities rather than cash flows from operating activities. We have not recognized such cash flow from financing activities since there has been no tax benefit related to the stock-based compensation.

Income Per Share of Common Stock - Basic income per share of common stock is calculated by dividing net income by the weighted average number of shares of common stock outstanding. Diluted income per share includes the dilution of common stock equivalents. The diluted weighted average shares of common stock outstanding were calculated using the treasury stock method for the respective periods.

Foreign Operations - The functional currency of our foreign subsidiary is its local currency. The assets and liabilities of our foreign subsidiary are translated into U.S. dollars at exchange rates in effect at the balance sheet date. Income and expense items are translated at the average exchange rates prevailing during the period. The cumulative translation effect for subsidiaries using a functional currency other than the U.S. dollar is included as a cumulative translation adjustment in stockholders' equity and as a component of comprehensive income.

Our intercompany accounts are typically denominated in the functional currency of the foreign subsidiary. Gains and losses resulting from the remeasurement of intercompany receivables that we consider to be of a long-term investment nature are recorded as a cumulative translation adjustment in stockholders' equity and as a component of comprehensive income, while gains and losses resulting from the remeasurement of intercompany receivables from those international subsidiaries for which we anticipate settlement in the foreseeable future are recorded in the consolidated statements of operations. The net gains and losses recorded in the consolidated statements of income were not significant for the periods presented.

New Accounting Standards

Recently issued FASB guidance and Securities Exchange Commission Staff Accounting Bulletins have either been implemented, with no significant effect, or are not applicable to the Company.

Forward-Looking Statements

Our business, financial condition, results of operations, cash flows and prospects, and the prevailing market price and performance of our common stock, may be adversely affected by a number of factors, including the matters discussed below. Certain statements and information set forth in this Annual Report on Form 10-K, as well as other written or oral statements made from time to time by us or by our authorized executive officers on our behalf, constitute "forward-looking statements" within the meaning of the Federal Private Securities Litigation Reform Act of 1995. These statements include all statements other than those made solely with respect to historical fact and identified by words such as "believes", "anticipates", "expects", "intends" and similar expressions, but such words are not the exclusive means of identifying such statements. We intend for our forward-looking statements to be covered by the safe harbor provisions for forward-looking statements contained in the Private Securities Litigation Reform Act of 1995, and we set forth this statement and these risk factors in order to comply with such safe harbor provisions. You should note

that our forward-looking statements speak only as of the date of this Annual Report on Form 10-K or when made and we undertake no duty or obligation to update or revise our forward-looking statements, whether as a result of new information, future events or otherwise. Although we believe that the expectations, plans, intentions and projections reflected in our forward-looking statements are reasonable, such statements are subject to known and unknown risks, uncertainties and other factors that may cause our actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by the forward-looking statements. The risks, uncertainties and other factors that our stockholders and prospective investors should consider include, but are not limited to, the following:

Because our business is concentrated in the pharmaceutical industry any changes in that industry or in the markets we serve could impair our ability to generate revenue and realize a profit.

Puerto Rico government enacted ACT 154 of October 22, 2010 which may affect adversely the willingness of our customers to do business in Puerto Rico and consequently affect our business.

Changes in tax benefits may affect the willingness of companies to continue or expand their operations in Puerto Rico.

Puerto Rico's economy, including its governmental financial crisis, may affect the willingness of businesses to commence or expand operations in Puerto Rico.

Other factors, including economic factors, may affect the decision of businesses to continue or expand their operations in the markets we serve.

Because our business is dependent upon a small number of clients, the loss of a major client could impair our ability to operate profitably.

Customer procurement and sourcing practices intended to reduce costs could have an adverse affect on our margins and profitability.

Since our business is dependent upon the development and enhancement of patented pharmaceutical products or processes by our clients, the failure of our clients to obtain and maintain patents could impair our ability to operate profitably.

We may be unable to pass on increased labor costs to our clients.

Consolidation in the pharmaceutical industry may have a harmful effect on our business.

Because the pharmaceutical industry is subject to government regulations, changes in government regulations relating to this industry may affect the need for our services.

Our reputation and divisions may be impacted by regulatory standards impacting our customer products.

If we are unable to protect our clients' intellectual property, our ability to generate business will be impaired.

We may be subject to liability if our services or solutions for our clients infringe upon the intellectual property rights of others.

We may be held liable for the actions of our employees or contractors when on assignment.

To the extent that we perform services pursuant to fixed-price or incentive-based contracts, our cost of services may exceed our revenue on the contract.

Because most of our contracts may be terminated on little or no advance notice, our failure to generate new business could impair our ability to operate profitably.

Because we are dependent upon our management, our ability to develop our business may be impaired if we are not able to engage skilled personnel.

We may not be able to continue to grow unless we consummate acquisitions or enter markets outside of Puerto Rico, the United States, Ireland and Spain.

If we identify a proposed acquisition, we may require substantial cash to fund the cost of the acquisition.

If we make any acquisitions, they may disrupt or have a negative impact on our business.

Because there is a limited market in our common stock, stockholders may have difficulty in selling our common stock and our common stock may be subject to significant price swings.

Our revenues, operating results and profitability will vary from quarter to quarter, which may result in increased volatility of our stock price.

The issuance of securities, whether in connection with an acquisition or otherwise, may result in significant dilution to our stockholders.

ITEM 7A. QUANTITATIVE AND QUALITATIVE DISCLOSURE ABOUT MARKET RISK.

Not applicable.

ITEM 8. FINANCIAL STATEMENTS AND SUPPLEMENTARY DATA.

Our Consolidated Financial Statements, together with the report of our independent registered public accounting firm are included herein immediately following the signature page of this report. See Index to Consolidated Financial Statements on page F-1.

ITEM 9. CHANGES IN AND DISAGREEMENTS WITH ACCOUNTANTS ON ACCOUNTING AND FINANCIAL DISCLOSURE.

None.

ITEM 9A. CONTROLS AND PROCEDURES.

Management's Annual Report on Internal Control Over Financial Reporting

Our management is responsible for establishing and maintaining adequate "internal control over financial reporting," as defined in Rules 13a-15(f) and 15d-15(f) under the Exchange Act, for the Company. This rule defines internal control over financial reporting as a process designed by, or under the supervision of, a company's chief executive officer and chief financial officer, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles. Our internal control over financial reporting includes those policies and procedures that:

pertain to the maintenance of records that, in reasonable detail, accurately and fairly reflect the transactions and dispositions of the assets of the company;

provide reasonable assurance that transactions are recorded as necessary to permit preparation of financial statements in accordance with generally accepted accounting principles, and that receipts and expenditures of the company are being made only in accordance with authorizations of management and directors of the company; and

provide reasonable assurance regarding prevention or timely detection of unauthorized acquisition, use or disposition of the company's assets that could have a material effect on the financial statements.

Because of its inherent limitations, our internal control systems and procedures may not prevent or detect misstatements. An internal control system, no matter how well conceived and operated, can provide only reasonable, not absolute, assurance that the objectives of the control system are met. Because of the inherent limitations in all control systems, no evaluation of controls can provide absolute assurance that all control issues and instances of fraud, if any, have been detected. Also, projections of any evaluation of effectiveness to future periods are subject to the risks that controls may become inadequate because of changes in conditions, or that the degree of compliance with the policies and procedures may deteriorate.

We, under the supervision of and with the participation of our management, including the Chief Executive Officer and Chief Financial Officer, assessed the effectiveness of the Company's internal control over financial reporting as of October 31, 2012, based on criteria for effective internal control over financial reporting described in "Internal Control — Integrated Framework" issued by the Committee of Sponsoring Organizations of the Treadway Commission. Based on this assessment, our Chief Executive Officer and Chief Financial Officer concluded that the Company maintained effective internal control over financial reporting as of October 31, 2012, based on the specified criteria.

Disclosure Controls and Procedures.

We carried out an evaluation, under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, of the effectiveness of our disclosure controls and procedures (as defined in Exchange Act Rules 13a-15(e) and 15d-15(e)) as of the end of the period covered by this Annual Report. Based upon that evaluation, our Chief Executive Officer and Chief Financial Officer concluded that our disclosure controls and procedures were effective as of the end of the period covered by this Annual Report.

Changes in Internal Control Over Financial Reporting

Based on an evaluation, under the supervision and with the participation of our management, including our Chief Executive Officer and Chief Financial Officer, there has been no change in our internal control over financial reporting during our last fiscal quarter identified in connection with that evaluation that has materially affected, or is

reasonably likely to materially affect, our internal control over financial reporting.

ITEM 9B. OTHER INFORMATION.

None.

20

PART III

ITEM 10. DIRECTORS, EXECUTIVE OFFICERS AND CORPORATE GOVERNANCE.

The information required by this Item will be included in a definitive proxy statement, pursuant to Regulation 14A, to be filed not later than 120 days after the close of our fiscal year. Such information is incorporated herein by reference.

Information with respect to our executive officers is included in Part I.

ITEM 11. EXECUTIVE COMPENSATION.

The information required by this Item will be included in a definitive proxy statement, pursuant to Regulation 14A, to be filed not later than 120 days after the close of our fiscal year. Such information is incorporated herein by reference.

ITEM 12. SECURITY OWNERSHIP OF CERTAIN BENEFICIAL OWNERS AND MANAGEMENT AND RELATED STOCKHOLDER MATTERS.

The information required by this Item will be included in a definitive proxy statement, pursuant to Regulation 14A, to be filed not later than 120 days after the close of our fiscal year. Such information is incorporated herein by reference.

ITEM 13. CERTAIN RELATIONSHIPS AND RELATED TRANSACTIONS, AND DIRECTOR INDEPENDENCE.

The information required by this Item will be included in a definitive proxy statement, pursuant to Regulation 14A, to be filed not later than 120 days after the close of our fiscal year. Such information is incorporated herein by reference.

ITEM 14. PRINCIPAL ACCOUNTING FEES AND SERVICES.

The information required by this Item will be included in a definitive proxy statement, pursuant to Regulation 14A, to be filed not later than 120 days after the close of our fiscal year. Such information is incorporated herein by reference.

PART IV

ITEM 15. EXHIBITS, FINANCIAL STATEMENT SCHEDULES.

The following documents are filed as a part of this Annual Report on Form 10-K:

1. All Financial Statements: Consolidated Financial Statements are included herein immediately following the signature page of this report. See Index to Consolidated Financial Statements on page F-1.

2. Financial Statement Schedules: None.

3. Exhibits: The following exhibits are filed herewith or are incorporated by reference to exhibits previously filed with the Commission, as indicated in the description of each.

Exhibit Number	Exhibit Description	Form	Incorporated By Reference		
			File Number	Exhibit	Filing Date
3.1	Restated Certificate of Incorporation	8-K	000-50956	99.1	5/1/2006
3.2	By-laws	10-SB12G	000-50956	3.2	9/24/2004
3.3	Amendment No. 1 to the By-laws	8-K	000-50956	3.1	6/6/2008
4.1	Form of warrant issued to Investors in January 2006 private placement	8-K	000-50956	4.2	1/31/2006
4.2	Form of warrant held by initial warrant holders	8-K	000-50956	4.3	1/31/2006
4.3	Form of warrant held by San Juan Holdings	8-K	000-50956	4.4	1/31/2006
4.4	Form of warrants issued to broker-dealers in January 2006 private placement	8-K	000-50956	4.5	1/31/2006
4.5	Form of First Amendment to Series C Common Stock Purchase Warrant.	8-K	000-50956	4.1	1/29/2009
10.1	Form of subscription agreement for January 2006 private placement	8-K	000-50956	99.1	1/31/2006
10.2	Registration rights provisions for the subscription agreement relating to January 2006 private placement	8-K	000-50956	99.2	1/31/2006
10.3	Registration rights provisions for Elizabeth Plaza and San Juan Holdings, Inc.	8-K	000-50956	99.3	1/31/2006

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

10.4	Employment Agreement dated January 2, 2008 between the Registrant and Elizabeth Plaza	10-KSB	000-50956	10.5	1/31/2008
10.5	Amendment to Employment Agreement dated June 9, 2008 between the Registrant and Elizabeth Plaza	10-K	000-50956	10.5	1/29/2009
10.6	Second Amendment to Employment Agreement, dated March 11, 2009, by and between the Company and Elizabeth Plaza.	8-K	000-50956	10.1	3/17/2009
10.7	Third Amendment to Employment Agreement, dated March 11, 2009, by and between the Company and Elizabeth Plaza.	8-K	000-50956	10.2	3/17/2009

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

10.8	Employment Agreement Amendment, effective as of January 1, 2010, by and between the Company and Elizabeth Plaza.	8-K	000-50956	10.1	1/07/2010
10.9	Employment Agreement Amendment, effective as of July 1, 2010, by and between the Company and Elizabeth Plaza	8-K	000-50956	10.1	7/8/2010
10.10	Sixth Employment Agreement Amendment, effective as of August 23, 2010, by and between the Company and Elizabeth Plaza	8-K	000-50956	10.1	8/27/10
10.11	Employment Agreement dated January 25, 2006 between the Registrant and Nélica Plaza	8-K	000-50956	99.5	1/31/2006
10.12	Amendment to Employment Agreement, dated March 11, 2009, by and between the Company and Nelida Plaza.	8-K	000-50956	10.4	3/17/2009
10.13	Employment Agreement, dated as of December 31, 2009, by and between Pharma-Bio Serv PR, Inc. and Nelida Plaza.	8-K	000-50956	10.3	1/07/2010
10.14	Employment Agreement dated November 5, 2007 between the Registrant and Pedro Lasanta	10-K	000-50956	10.8	1/29/2009
10.15	Amendment to Employment Agreement dated December 17, 2008 between the Registrant and Pedro Lasanta	8-K	000-50956	99.1	12/23/2008
10.16	Amendment to Employment Agreement, dated March 11, 2009, by and between the Company and Pedro Lasanta.	8-K	000-50956	10.3	3/17/2009
10.17	Employment Agreement Amendment, effective as of January 1, 2010, by and between the Company and Pedro Lasanta.	8-K	000-50956	10.2	1/07/2010
10.18	2005 Long-term incentive plan, as amended	DEF 14A	000-50956	Appendix C	3/26/2007
10.19	Lease dated March 16, 2004 between Plaza Professional Center, Inc. and the Registrant	SB-2	333-132847	10.9	3/30/2006
10.20		SB-2	333-132847	10.10	3/30/2006

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

Lease dated November 1, 2004
between Plaza Professional Center,
Inc. and the Registrant

10.21	Vendor Agreement dated May 4, 2006 between the Registrant and Schering-Plough Products, L.L.C.	SB-2/A	333-132847	10.12	11/8/2006
10.22	Agreement dated January 17, 2006 between Lilly del Caribe, Inc. and Plaza Consulting Group, Inc.	SB-2/A	333-132847	10.13	11/8/2006

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

10.23	Agreement effective as of November 1, 2005 between SB Pharmco Puerto Rico Inc. d/b/a GlaxoSmithKline	SB-2/A	333-132847	10.14	10/27/2006
10.24	Seventh Employment Agreement Amendment, dated January 23, 2012 and effective as of January 1, 2012, by and between the Company and Elizabeth Plaza	8-K	000-50956	99.1	1/23/2012
10.25	Employment Agreement Amendment, dated January 31, 2012, by and between the Company and Pedro J. Lasanta	8-K	000-50956	10.1	2/2/2012
10.26	Employment Agreement Amendment, dated December 31, 2012, by and between the Company and Pedro J. Lasanta	8-K	000-50956	10.1	1/7/2013
10.27	Consulting Agreement, dated January 7, 2013, by and between Pharma-Bio Serv, Inc. and Elizabeth Plaza	8-K	000-50956	10.1	1/11/2013
10.28	Employment Agreement Amendment, dated January 7, 2013, by and among Pharma-Bio Serv, Inc., Pharma-Bio Serv PR, Inc. and Nelida Plaza	8-K	000-50956	10.2	1/11/2013
14.1	Code of business conduct and ethics for senior management	10-KSB	000-50956	14.1	2/2/2007
21.1*	List of Subsidiaries				
31.1*	Certification of chief executive officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002				
31.2*	Certification of chief financial officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002				
32.1**	Certification of chief executive officer and chief financial officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002				
101.INS***	XBRL Instance Document				
101.SCH***	XBRL Taxonomy Extension Schema				
101.CAL***	XBRL Taxonomy Extension Calculation Linkbase				
101.DEF***	XBRL Taxonomy Extension Definition Linkbase				
101.LAB***					

XBRL Taxonomy Extension Label
Linkbase

101.PRE*** XBRL Taxonomy Extension
Presentation Linkbase

* Filed herewith

** Furnished herewith

*** Pursuant to Rule 406T of Regulation S-T, these interactive data files are deemed not filed or part of a registration statement or prospectus for purposes of Section 11 or 12 of the Securities Act of 1933, as amended, are deemed not filed for purposes of Section 18 of the Securities Exchange Act of 1934, as amended, and otherwise are not subject to liability under those sections.

Exhibits 10.4 through 10.18 and 10.24 through 10.28 are management contracts or compensatory plans, contracts or arrangements.

SIGNATURES

Pursuant to the requirements of Section 13 or 15(d) of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

PHARMA-BIO SERV, INC.

Dated : January 29, 2013

By: /s/ NELIDA PLAZA
 Name: Nelida Plaza
 Title: Acting President and CEO, and
 President of Puerto Rico Operations and
 Secretary

Pursuant to the requirements of the Securities Exchange Act of 1934, this report has been signed below by the following persons on behalf of the registrant and in the capacities and on the dates indicated.

Signature	Title	Date
/s/ Nelida Plaza Nelida Plaza	Acting President and Chief Executive Officer, and President of Puerto Rico Operations and Secretary (Principal Executive Officer)	January 29, 2013
/s/ Pedro J. Lasanta Pedro J. Lasanta	Chief Financial Officer (Principal Financial and Accounting Officer)	January 29, 2013
/s/ Elizabeth Plaza Elizabeth Plaza	Chairman	January 29, 2013
/s/ Kirk Michel Kirk Michel	Director	January 29, 2013
/s/ Howard Spindel Howard Spindel	Director	January 29, 2013
/s/ Dov Perlysky Dov Perlysky	Director	January 29, 2013

/s/ Irving Wiesen Director
Irving Wiesen

January 29, 2013

25

PHARMA-BIO SERV, INC.

INDEX TO CONSOLIDATED FINANCIAL STATEMENTS

	Page
Report of Independent Registered Public Accounting Firm	F-2
Consolidated Balance Sheets as of October 31, 2012 and 2011	F-3
Consolidated Statements of Income for the Years Ended October 31, 2012 and 2011	F-4
Consolidated Statements of Comprehensive Income for the Years Ended October 31, 2012 and 2011	F-5
Consolidated Statements of Changes in Stockholders' Equity for the Years Ended October 31, 2012 and 2011	F-6
Consolidated Statements of Cash Flows for the Years Ended October 31, 2012 and 2011	F-7
Notes to Consolidated Financial Statements	F-8

F-1

REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Board of Directors and Stockholders of
Pharma-Bio Serv, Inc.
Dorado, Puerto Rico

We have audited the accompanying consolidated balance sheets of Pharma-Bio Serv, Inc. as of October 31, 2012 and 2011, and the related consolidated statements of income, comprehensive income, changes in stockholders' equity, and cash flows for the years then ended. Pharma-Bio Serv, Inc.'s management is responsible for these financial statements. Our responsibility is to express an opinion on these consolidated financial statements based on our audits.

We conducted our audits in accordance with the standards of the Public Company Accounting Oversight Board (United States). Those standards require that we plan and perform the audit to obtain reasonable assurance about whether the financial statements are free of material misstatement. The Company is not required to have, nor were we engaged to perform, an audit of its internal control over financial reporting. Our audit included consideration of internal control over financial reporting as a basis for designing audit procedures that are appropriate in the circumstances, but not for the purpose of expressing an opinion on the effectiveness of the company's internal control over financial reporting. Accordingly, we express no such opinion. An audit also includes examining, on a test basis, evidence supporting the amounts and disclosures in the financial statements, assessing the accounting principles used and significant estimates made by management, as well as evaluating the overall financial statement presentation. We believe that our audits provide a reasonable basis for our opinion.

In our opinion, the financial statements referred to above present fairly, in all material respects, the consolidated financial position of Pharma-Bio Serv, Inc. as of October 31, 2012 and 2011, and the consolidated results of its operations and its cash flows for the years then ended in conformity with accounting principles generally accepted in the United States of America.

/S/ HORWATH VÉLEZ & CO, PSC
San Juan, Puerto Rico

January 29, 2013
Puerto Rico Society of Certified Public Accountants
Stamp number 2669443 was
affixed to the original of this report

PHARMA-BIO SERV, INC.
Consolidated Balance Sheets
October 31, 2012 and 2011

	October 31,	
	2012	2011
ASSETS:		
Current assets		
Cash and cash equivalents	\$ 6,538,113	\$ 4,316,725
Marketable securities	95,000	95,000
Accounts receivable	7,580,167	4,864,616
Other	382,773	331,441
Total current assets	14,596,053	9,607,782
Property and equipment	1,113,371	1,216,111
Other assets	25,592	28,306
Total assets	\$ 15,735,016	\$ 10,852,199
LIABILITIES AND STOCKHOLDERS' EQUITY:		
Current liabilities		
Current portion-obligations under capital leases	\$ 39,436	\$ 31,142
Accounts payable and accrued expenses	2,562,462	1,941,658
Income taxes payable	173,620	550,837
Total current liabilities	2,775,518	2,523,637
Obligations under capital leases	83,912	92,237
Total liabilities	2,859,430	2,615,874
Stockholders' equity:		
Preferred Stock, \$0.0001 par value; authorized 10,000,000 shares; none outstanding	-	-
Common Stock, \$0.0001 par value; authorized 50,000,000 shares; issued and outstanding 20,758,695	2,076	2,076
Additional paid-in capital	678,214	654,550
Retained earnings	12,286,714	7,599,708
Accumulated other comprehensive loss	(91,418)	(20,009)
Total stockholders' equity	12,875,586	8,236,325
Total liabilities and stockholders' equity	\$ 15,735,016	\$ 10,852,199

See notes to consolidated financial statements.

PHARMA-BIO SERV, INC.
Consolidated Statements of Income
For the Years Ended October 31, 2012 and 2011

	Years ended October 31,	
	2012	2011
REVENUES	\$ 29,227,167	\$ 19,933,182
COST OF SERVICES	19,355,440	13,072,231
GROSS PROFIT	9,871,727	6,860,951
SELLING, GENERAL AND ADMINISTRATIVE EXPENSES	4,138,111	3,408,953
INCOME FROM OPERATIONS	5,733,616	3,451,998
OTHER INCOME (EXPENSES):		
Interest expense	(7,425)	(6,605)
Interest income	11,076	19,709
Gain (loss) on disposition of property and equipment	18,345	(1,324)
	21,996	11,780
INCOME BEFORE INCOME TAXES	5,755,612	3,463,778
INCOME TAXES	1,068,606	304,797
NET INCOME	\$ 4,687,006	\$ 3,158,981
BASIC EARNINGS PER COMMON SHARE	\$ 0.226	\$ 0.152
DILUTED EARNINGS PER COMMON SHARE	\$ 0.204	\$ 0.140
WEIGHTED AVERAGE NUMBER OF COMMON SHARES OUTSTANDING – BASIC	20,758,695	20,754,043
WEIGHTED AVERAGE NUMBER OF COMMON SHARES OUTSTANDING – DILUTED	22,939,701	22,554,036

See notes to consolidated financial statements.

PHARMA-BIO SERV, INC.
 Consolidated Statements of Comprehensive Income
 For the Years Ended October 31, 2012 and 2011

	Years ended October 31,	
	2012	2011
NET INCOME	\$ 4,687,006	\$ 3,158,981
OTHER COMPREHENSIVE LOSS:		
Foreign currency translation adjustment, net of tax	(71,409)	(791)
TOTAL OTHER COMPREHENSIVE LOSS	(71,409)	(791)
COMPREHENSIVE INCOME	\$ 4,615,597	\$ 3,158,190

See notes to consolidated financial statements.

PHARMA-BIO SERV, INC.
 Consolidated Statements of Changes in Stockholders' Equity
 For the Years Ended October 31, 2012 and 2011

	Common Stock Shares	Common Stock Amount	Preferred Stock Shares	Preferred Stock Amount	Additional Paid-in Capital	Retained Earnings	Accumulated Other Comprehensive (Loss)	Total
BALANCE AT OCTOBER 31, 2010	20,751,215	\$ 2,075	-	\$ -	\$ 645,886	\$ 4,440,728	\$ (19,218)	\$ 5,069,471
STOCK-BASED COMPENSATION	-	-	-	-	8,664	-	-	8,664
CASHLESS CONVERSION OF WARRANTS TO SHARES OF COMMON STOCK	7,480	1	-	-	-	(1)	-	-
NET INCOME	-	-	-	-	-	3,158,981	-	3,158,981
FOREIGN CURRENCY TRANSLATION ADJUSTMENT	-	-	-	-	-	-	(791)	(791)
BALANCE AT OCTOBER 31, 2011	20,758,695	2,076	-	-	654,550	7,599,708	(20,009)	8,236,325
STOCK-BASED COMPENSATION	-	-	-	-	23,664	-	-	23,664
NET INCOME	-	-	-	-	-	4,687,006	-	4,687,006
FOREIGN CURRENCY TRANSLATION ADJUSTMENT	-	-	-	-	-	-	(71,409)	(71,409)
BALANCE AT OCTOBER 31, 2012	20,758,695	\$ 2,076	-	\$ -	\$ 678,214	\$ 12,286,714	\$ (91,418)	\$ 12,875,586

See notes to consolidated financial statements.

PHARMA-BIO SERV, INC.
Consolidated Statements of Cash Flows
For the Years Ended October 31, 2012 and 2011

	Years ended October 31,	
	2012	2011
CASH FLOWS FROM OPERATING ACTIVITIES:		
Net income	\$ 4,687,006	\$ 3,158,981
Adjustments to reconcile net income to net cash provided by operating activities:		
(Gain) loss on disposition of property and equipment	(18,345)	1,324
Stock-based compensation	23,664	8,664
Depreciation and amortization	321,732	320,861
Increase in accounts receivable	(2,735,046)	(2,274,869)
(Increase) decrease in other assets	(50,486)	(78,713)
Increase in liabilities	230,541	1,004,025
NET CASH PROVIDED BY OPERATING ACTIVITIES	2,459,066	2,140,273
CASH FLOWS FROM INVESTING ACTIVITIES:		
Acquisition of property and equipment	(187,362)	(123,511)
Proceeds from sale of property and equipment	18,836	400
NET CASH USED IN INVESTING ACTIVITIES	(168,526)	(123,111)
CASH FLOW FROM FINANCING ACTIVITIES:		
Payments on obligations under capital lease	(32,826)	(25,162)
NET CASH USED IN FINANCING ACTIVITIES	(32,826)	(25,162)
EFFECT OF EXCHANGE RATE CHANGES ON CASH	36,326	7,557
NET INCREASE IN CASH AND CASH EQUIVALENTS	2,221,388	1,999,557
CASH AND CASH EQUIVALENTS - BEGINNING OF YEAR	4,316,725	2,317,168
CASH AND CASH EQUIVALENTS – END OF YEAR	\$ 6,538,113	\$ 4,316,725
SUPPLEMENTAL DISCLOSURES OF CASH FLOW INFORMATION:		
Cash paid during the period for:		
Income taxes	\$ 1,543,021	\$ 6,025
Interest	\$ 7,425	\$ 6,605
SUPPLEMENTARY SCHEDULES OF NON-CASH INVESTING AND FINANCING ACTIVITIES:		
Property and equipment with accumulated depreciation of \$32,122 and \$1,887 disposed during the years ended October 31, 2012 and 2011, respectively.	\$ 32,613	\$ 3,611
Income tax withheld by clients to be used as a credit in the Company's income tax returns	\$ 39,120	\$ 73,671
Obligations under capital lease incurred for the acquisition of a vehicle	\$ 32,795	\$ 76,475

See notes to consolidated financial statements.

F-7

PHARMA-BIO SERV, INC.
Notes To Consolidated Financial Statements
For the Years Ended October 31, 2012 and 2011

NOTE A - ORGANIZATION AND SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

ORGANIZATION

Pharma-Bio Serv, Inc. (“Pharma-Bio”) is a Delaware corporation organized on January 14, 2004. Pharma-Bio is the parent company of Pharma-Bio Serv PR, Inc. (“Pharma-PR”), Pharma Serv, Inc. (“Pharma-Serv”) both Puerto Rico corporations, Pharma-Bio Serv US, Inc. (“Pharma-US”), a Delaware corporation, and Pharma-Bio Serv Validation & Compliance Limited (“Pharma-IR”), an Irish corporation. Pharma-Bio, Pharma-PR, Pharma-Serv, Pharma-US and Pharma-IR are collectively referred to as the “Company.” The Company operates in Puerto Rico, the United States and in Ireland under the name of Pharma-Bio Serv and is engaged in providing technical compliance consulting service, and microbiological and chemical laboratory testing services primarily to the pharmaceutical, chemical, medical device and biotechnology industries.

During the quarter ended April 30, 2012, based on a prior agreement between the Company and the minority shareholder of Pharma-IR, the Company acquired 100% ownership of Pharma-IR for no consideration. This transaction had no significant impact on the consolidated financial statements

SUMMARY OF SIGNIFICANT ACCOUNTING POLICIES

Consolidation

The accompanying consolidated financial statements include the accounts of the Company and all of its wholly owned subsidiaries. All intercompany transactions and balances have been eliminated in consolidation.

Use of Estimates

The preparation of consolidated financial statements in conformity with accounting principles generally accepted in the United States of America requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the consolidated financial statements and the reported amounts of revenues and expenses during the reporting periods. Actual results may differ from these estimates.

Fair Value of Financial Instruments

Accounting standards have established a fair value hierarchy that requires an entity to maximize the use of observable inputs and minimize the use of unobservable inputs when measuring fair value. A financial instrument’s categorization within the fair value hierarchy is based upon the lowest level of input that is significant to the fair value measurement. Accounting standards have established three levels of inputs that may be used to measure fair value:

Level 1: Quoted prices in active markets for identical assets and liabilities.

Level 2: Observable inputs other than Level 1 prices such as quoted prices for similar assets or liabilities, quoted prices in markets with insufficient volume or infrequent transactions (less active markets), or model-derived valuations in which all significant inputs are observable or can be derived principally from or corroborated by observable market data

for substantially the full term of the assets or liabilities.

Level 3: Prices or valuation techniques that require inputs that are both significant to the fair value measurement and unobservable (supported by little or no market activity).

Marketable securities consist of an obligation from the Puerto Rico Government Development Bank valued using quoted market prices in active markets with no valuation adjustment. Accordingly, this security is categorized in Level 1.

The carrying value of the Company's financial instruments (excluding marketable securities and obligations under capital leases): cash and cash equivalents, accounts receivable, accounts payable and accrued liabilities, are considered reasonable estimates of fair value due to their liquidity or short-term nature. Management believes, based on current rates, that the fair value of its obligations under capital leases approximates the carrying amount.

Revenue Recognition

Revenue is primarily derived from: (1) time and materials contracts (representing approximately 95% of total revenues), which is recognized by applying the proportional performance model, whereby revenue is recognized as performance occurs, (2) short-term fixed-fee contracts or "not to exceed" contracts (representing approximately 2% of total revenues), which revenue is recognized similarly, except that certain milestones also have to be reached before revenue is recognized, and (3) laboratory testing revenue (representing approximately 3% of total revenues) which is mainly recognized as the testing is completed and certified (normally within days of sample receipt from customer). If the Company determines that a contract will result in a loss, the Company recognizes the estimated loss in the period in which such determination is made.

Cash Equivalents

For purposes of the consolidated statements of cash flows, cash equivalents include investments in a money market obligations trust that is registered under the U.S. Investment Company Act of 1940 and liquid investments with original maturities of three months or less.

Marketable Securities

We consider our marketable security investment portfolio and marketable equity investments available-for-sale and, accordingly, these investments are recorded at fair value with unrealized gains and losses generally recorded in other comprehensive income; whereas realized gains and losses are included in earnings and determined based on the specific identification method.

Accounts Receivable

Accounts receivable are recorded at their estimated realizable value. Accounts are deemed past due when payment has not been received within the stated time period. The Company's policy is to review individual past due amounts periodically and write off amounts for which all collection efforts are deemed to have been exhausted. Due to the nature of the Company's customers, bad debts are mainly accounted for using the direct write-off method whereby an expense is recognized only when a specific account is determined to be uncollectible. The effect of using this method approximates that of the allowance method.

Income Taxes

The Company follows an asset and liability approach method of accounting for income taxes. This method measures deferred income taxes by applying enacted statutory rates in effect at the balance sheet date to the differences between the tax basis of assets and liabilities and their reported amounts on the financial statements. The resulting deferred tax assets or liabilities are adjusted to reflect changes in tax laws as they occur. A valuation allowance is provided when it is more likely than not that a deferred tax asset will not be realized.

The Company follows guidance from the FASB related to Accounting for Uncertainty in Income Taxes, which includes a two-step approach to recognizing, de-recognizing and measuring uncertain tax positions. As of October 31, 2012, the Company had no significant uncertain tax positions that would be reduced as a result of a lapse of the applicable statute of limitations.

Property and Equipment

Owned property and equipment, and leasehold improvements are stated at cost. Equipment and vehicles under capital leases are stated at the lower of fair market value or net present value of the minimum lease payments at the inception of the leases. Depreciation and amortization of owned assets are provided for, when placed in service, in amount sufficient to relate the cost of depreciable assets to operations over their estimated service lives, using straight-line basis. Assets under capital leases and leasehold improvements are amortized, over the shorter of the estimated useful lives of the assets or initial lease term. Major renewals and betterments that extend the life of the assets are capitalized, while expenditures for repairs and maintenance are expensed when incurred.

The Company evaluates for impairment its long-lived assets to be held and used, and long-lived assets to be disposed of, whenever events or changes in circumstances indicate that the carrying amount of an asset may not be recoverable. Based on management estimates, no impairment of the operating properties was present.

Intangible Assets

Definite-lived intangible assets, such as customer lists and covenants not to compete, are amortized on a straight-line basis over their estimated useful lives. The Company continually evaluates the reasonableness of the useful lives of these assets.

Stock-based Compensation

Stock-based compensation expense is recognized in the consolidated financial statements based on the fair value of the awards granted. Stock-based compensation cost is measured at the grant date based on the fair value of the award and is recognized as expense over the requisite service period, which generally represents the vesting period, and includes an estimate of awards that will be forfeited. The Company calculates the fair value of stock options using the Black-Scholes option-pricing model at grant date. Excess tax benefits related to stock-based compensation are reflected as cash flows from financing activities rather than cash flows from operating activities. The Company has not recognized such cash flow from financing activities since there has been no tax benefit related to the stock-based compensation.

Income Per Share of Common Stock

Basic income per share of common stock is calculated dividing net income by the weighted average number of shares of common stock outstanding. Diluted income per share includes the dilution of common stock equivalents.

The diluted weighted average shares of common stock outstanding were calculated using the treasury stock method for the respective periods.

Foreign Operations

The functional currency of the Company's foreign subsidiary is its local currency. The assets and liabilities of the Company's foreign subsidiary are translated into U.S. dollars at exchange rates in effect at the balance sheet date. Income and expense items are translated at the average exchange rates prevailing during the period. The cumulative translation effect for subsidiaries using a functional currency other than the U.S. dollar is included as a cumulative translation adjustment in stockholders' equity and as a component of comprehensive income.

The Company's intercompany accounts are typically denominated in the functional currency of the foreign subsidiary. Gains and losses resulting from the remeasurement of intercompany receivables that the Company considers to be of a long-term investment nature are recorded as a cumulative translation adjustment in stockholders' equity and as a component of comprehensive income, while gains and losses resulting from the remeasurement of intercompany receivables from those international subsidiaries for which the Company anticipates settlement in the foreseeable future are recorded in the consolidated statements of operations. The net gains and losses recorded in the consolidated statements of income were not significant for the periods presented.

Subsequent Events

The Company has determined that there are no events occurring in this period that required disclosure in or adjustment to the accompanying consolidated financial statements.

Reclassifications

Certain reclassifications have been made to the October 31, 2011 consolidated financial statements to conform them to the October 31, 2012 consolidated financial statements presentation. Such reclassifications do not have effect on net income as previously reported.

Recent Accounting Pronouncements

Recent issued FASB guidance and Securities and Exchange Commission ("SEC") Staff Accounting Bulletins have either been implemented, with no significant effect, or are not applicable to the Company.

NOTE B – MARKETABLE SECURITIES AVAILABLE FOR SALE

At October 31, 2012 and 2011, the marketable securities of \$95,000 consisted of a 5.4% Puerto Rico Commonwealth Government Development Bank Bond, purchased at par and maturing in August 2019. The bond balance approximates its fair market value, therefore no realized or unrealized gains or losses have been recorded.

The primary objectives of the Company's investment portfolio are liquidity and safety of principal. Investments are made with the objective of achieving the highest rate of return consistent with these two objectives. Our investment policy limits investments to certain types of debt and money market instruments issued by institutions primarily with

investment grade credit ratings and places restrictions on maturities and concentration by type and issuer.

We review our available-for-sale securities for other-than-temporary declines in fair value below their cost basis on a quarterly basis and whenever events or changes in circumstances indicate that the cost basis of an asset may not be recoverable. This evaluation is based on a number of factors including, the length of time and extent to which the fair value has been less than our cost basis and adverse conditions specifically related to the security including any changes to the rating of the security by a rating agency. As of October 31, 2012, we believe that the cost base for our available-for-sale securities is recoverable in all material respects.

F-10

NOTE C - PROPERTY AND EQUIPMENT

The balance of property and equipment at October 31, 2012 and 2011 consisted of the following:

	Useful life (years)	October 31,	
		2012	2011
Vehicles	5	\$ 269,267	\$ 250,617
Leasehold improvements	5-8	598,040	598,040
Computers	3	522,792	420,482
Equipment	3-7	1,126,415	1,036,981
Furniture and fixtures	10	137,215	137,215
Projects in progress	-	-	23,524
Total		2,653,729	2,466,859
Less: Accumulated depreciation and amortization		(1,540,358)	(1,250,748)
Property and equipment, net		\$ 1,113,371	\$ 1,216,111

NOTE D - INCOME TAXES

In June 2011, Pharma-Bio, Pharma-PR and Pharma-Serv obtained a Grant of Industrial Tax Exemption pursuant to the terms and conditions set forth in Act No. 73 of May 28, 2008 (“the Grant”) issued by the Puerto Rico Industrial Development Company (“PRIDCO”). The Grant is effective as of November 1, 2009 and covers a fifteen year period. The Grant provides relief on various Puerto Rico taxes, including income tax, with certain limitations, for most of the activities carried on within Puerto Rico, including those that are for services to parties located outside of Puerto Rico. The Grant establishes a threshold (“Baseline”) on the Industrial Development Income (“IDI”) subject to the favorable income tax rates. The Baselines will be gradually reduced to zero within a four-year term. Certain activities covered under the Grant are not subject to a Baseline and are allowed a four year gradual phase-in from the maximum income tax rate of 30%, as provided by the 2011 Puerto Rico Internal Revenue Code, to the favorable fixed Act 73 income tax rate of 4%. In addition, IDI earnings distributions accumulated since November 1, 2009 are exempt from Puerto Rico earnings distribution tax.

For the years ended October 31, 2012 and 2011 the various activities covered by the Act 73 Grant were subject to different reduced effective income tax rates for a net aggregate effective income tax rate of 5.4% and 5.5%, respectively. Prospectively, beginning with our fiscal year ended October 31, 2013, all activities covered under the Act 73 Grant will be subject to the Act 73 reduced fixed income tax rate of 4%, since the gradual phase-in of the fixed income tax rate will be completed and the Baselines will have been reduced to zero.

Effective with our fiscal year ended October 31, 2012, Puerto Rico operations not covered in the exempt activities of the Grant are subject to Puerto Rico income tax at a maximum tax rate of 30%. Previously, the maximum income tax rate under the Puerto Rico Internal Revenue Code was 39%. The operations carried out in the United States by the Company’s subsidiary are taxed in the United States at a maximum regular federal income tax rate of 35%.

Distribution of earnings by the Puerto Rican subsidiaries to its parent are taxed at the federal level; however, the parent is able to receive a credit for the taxes paid by the subsidiary on its operations in Puerto Rico, to the extent of the federal taxes that result from those earnings. As a result, the income tax expense of the Company, under its present corporate structure, would normally be the Puerto Rico taxes on operations in Puerto Rico, federal and state taxes on operations in the United States, plus the earnings distribution tax in Puerto Rico from dividends paid to the Puerto Rican subsidiaries’ parent, and the parent’s federal income tax, if any, incurred upon the subsidiary’s earnings

distribution.

Deferred income tax assets and liabilities are computed for differences between the consolidated financial statements and tax bases of assets and liabilities that will result in taxable or deductible amounts in the future, based on enacted tax laws and rates applicable to the periods in which the differences are expected to affect taxable income.

The adoption of the Act 73 Grant in the fiscal year ended October 31, 2011 triggered a favorable non-recurring adjustment to the year ended October 31, 2011 income tax expense in the aggregate amount of approximately \$200,000, which is attributable to the fiscal year ended October 31, 2010 because of the retroactive nature of Act 73 Grant effective date to November 1, 2009. The non-recurring adjustment had the aggregate effect in the fiscal year ended October 31, 2011 of increasing earnings per share basic and diluted by \$0.010 and \$0.009, respectively.

F-11

The reconciliation between the United States federal statutory rate and our effective tax rate for the years ended October 31, 2012, and 2011 is as follows:

	October 31,	
	2012	2011
United States federal statutory rate	35.0%	35.0%
Non United States earnings invested indefinitely, and Puerto Rico Act 73 Tax Grant effect	(17.0)%	(18.6)%
Puerto Rico Act 73 Tax Grant effective date backdating for the year ended October 31, 2010	-%	(5.9)%
Other, net	0.6%	(1.7)%
Effective tax rate	18.6%	8.8%

As of October 31, 2012 and 2011, the Company has not recognized deferred income taxes on \$10.5 million and \$7.1 million of undistributed earnings of its Puerto Rican subsidiaries, respectively, since such earnings are considered to be reinvested indefinitely. If the earnings were distributed in the form of dividends, the Company would be subject to Puerto Rico earnings distribution tax and United States federal income tax for the aggregate amount of approximately \$1.6 million and \$0.6 million at October 31, 2012 and 2011, respectively.

At October 31, 2012, Pharma-IR has unused operating losses of approximately \$586,000 (with no expiration) after considering various timing differences for income tax purposes, which result in a potential deferred tax asset of approximately \$73,000. However, an allowance has been provided covering the total amount of such balance since it is uncertain whether the net operating losses can be used to offset future taxable income. Realization of future tax benefits related to a deferred tax asset is dependent on many factors, including the company's ability to generate taxable income. Accordingly, the income tax benefit will be recognized when realization is determined to be more probable than not. These net operating losses are available to offset future taxable income indefinitely.

The Company files income tax returns in the United States (federal and various states jurisdictions), Puerto Rico and Ireland. Tax years ending October 31, 2007 through October 31, 2012 are open and may be subject to potential examination in one or more jurisdictions. In the current fiscal year ended October 31, 2012, Pharma-Bio's federal income tax return for year ended October 31, 2008 was examined by the United States Internal Revenue Service, no deficiencies were assessed. Currently, the Company has no federal, state, Puerto Rico or foreign income tax examination.

NOTE E – COMMITMENTS AND CONTINGENCIES

Capitalized lease obligations - The Company leases vehicles under non-cancelable capital lease agreements with a cost of \$200,158 and \$167,363 (accumulated amortization of \$87,174 and \$52,608) as of October 31, 2012 and 2011, respectively. Amortization expense for vehicles under non-cancelable lease agreements amounted to \$34,566 and \$27,100 in the years ended October 31, 2012 and 2011, respectively.

The following is a schedule, by year, of future minimum lease payments under the capitalized leases together with the present value of the net minimum lease payments at October 31, 2012:

Twelve months ending October 31,	Amount
2013	\$ 46,647
2014	36,402
2015	24,293
2016	24,242

Edgar Filing: Pharma-Bio Serv, Inc. - Form 10-K

2017	6,266
Total future minimum lease payments	137,850
Less: Amount of imputed interest	(14,502)
Present value of future minimum lease payments	123,348
Current portion of obligation under capital leases	(39,436)
Long-term portion	\$ 83,912

F-12

Operating facilities - The Company conducts its administrative operations in office facilities which are leased under four different rental agreements.

In February 2012, the Company automatically renewed a lease agreement, with an affiliate of our former chief executive officer, for the headquarters and laboratory testing facilities in Dorado, Puerto Rico. The renewal is for a term of five years with monthly rental payments of \$23,930, \$25,127, \$26,383, \$27,702 and \$29,087 for each of the five years remaining under the lease. The agreement also requires the payment of utilities, property taxes, insurance and a portion of expenses incurred by the affiliate in connection with the maintenance of common areas.

In November 2011, the Company entered into a lease agreement for the U.S. office facilities located in Plymouth, Pennsylvania. The lease is for a five-year term with monthly rental payments of \$6,282 for the first three years. Thereafter, the lease will increase four percent every year, including a five-year renewal option, if executed.

The Company maintains office facilities in Cork, Ireland and Madrid, Spain. Both facilities are under month-to-month leases with monthly payments of approximately \$900 and \$1,200, respectively.

The Company leases certain apartments as dwellings for employees. The leases are under short-term lease agreements and usually are cancelable upon 30-day notification.

Minimum future rental payments under non-cancelable operating leases having remaining terms in excess of one year as of October 31, 2012 are as follows:

	Amount
2013	\$ 373,314
2014	388,211
2015	406,867
2016	426,427
2017	87,262
Total minimum lease payments	\$ 1,682,081

Rent expense during the years ended October 31, 2012 and 2011 was \$411,162 and \$293,687, respectively.

Contingencies - In the ordinary course of business, the Company may be a party to legal proceedings incidental to the business. These proceedings are not expected to have a material adverse effect on the Company's business or financial condition.

NOTE F – WARRANTS

At October 31, 2012 and 2011, the Company had outstanding warrants to purchase shares of the Company's common stock as follows:

	Exercise Price	Expiration Date	Outstanding Warrants
Original Warrants A	\$ 0.06	January 16, 2014	240,800
Broker Warrants B	\$ 0.06	January 24, 2014	1,830,991
Warrants Total			2,071,791

NOTE G – EARNINGS PER SHARE

The following data show the amounts used in the calculations of basic and diluted earnings per share.

	Years ended October 31,	
	2012	2011
Net income available to common equity holders - used to compute basic and diluted earnings per share	\$ 4,687,006	\$ 3,158,981
Weighted average number of common shares - used to compute basic earnings per share	20,758,695	20,754,043
Effect of warrants to purchase common stock	1,919,849	1,777,681
Effect of options to purchase common stock	261,157	22,312
Weighted average number of shares - used to compute diluted earnings per share	22,939,701	22,554,036

Options for the purchase of 374,585 shares of common stock for the year ended in October 31, 2011 were not included in computing diluted earnings per share because their effects were antidilutive.

NOTE H - STOCK OPTIONS AND STOCK BASED COMPENSATION

In October 2005, the Company's board of directors adopted, and on April 25, 2006, the Company's stockholders approved, the 2005 Long-Term Incentive Plan, covering 2,500,000 shares of common stock. The 2005 plan provides for the grant of incentive and non-qualified options, stock grants, stock appreciation rights and other equity-based incentives to employees, including officers, consultants and directors. The 2005 plan is to be administered by a committee of independent directors. In the absence of a committee, the plan is administered by the board of directors. Options intended to be incentive stock options must be granted at an exercise price per share which is not less than the fair market value of the common stock on the date of grant and may have a term which is not longer than ten years. If the option holder holds at least 10% of the Company's common stock, the exercise price must be at least 110% of the fair market value on the date of grant and the term of the option cannot exceed five years.

The Company recognizes stock-based compensation based on the fair value of the awards. Stock-based compensation cost is measured at the grant date based on the fair value of the award and is recognized as expense over the requisite service period, which generally represents the vesting period, and includes an estimate of awards that will be forfeited.

The 2005 Plan stock options activity and status for the years ended October 31, 2012 and 2011 was as follows:

	Year ended October 31,			
	2012		2011	
	Number of	Weighted-Average	Number of	Weighted-Average
	Shares	Option Exercise Price	Shares	Option Exercise Price
Outstanding at beginning of year	454,585	\$ 0.5804	1,267,882	\$ 0.6942
Granted	1,355,000	\$ 0.7372	40,000	\$ 0.2600
Exercised	-		-	
Expired and/or forfeited	(69,585)	\$ 0.7317	(853,297)	\$ 0.7344
Total outstanding at end of year	1,740,000	\$ 0.6965	454,585	\$ 0.5804

Outstanding exercisable stock options at end of year	430,000	\$	0.5702	424,585	\$	0.5974
--	---------	----	--------	---------	----	--------

	October 31,	October 31,
	2012	2011

Weighted average remaining years in contractual life
for:

Total outstanding options	3.4 years	1.7 years
Outstanding exercisable options	1.0 years	1.5 years
Shares of common stock available for issuance pursuant to future stock option grants	760,000	2,045,415

F-14

The following table presents the stock-based compensation included in the Company's consolidated statement of income and the effect in earnings per share:

	Year ended October 31,	
	2012	2011
Stock-based compensation expense:		
Cost of services	\$ 10,000	\$ -
Selling, general and administrative	13,664	8,664
Stock-based compensation before tax	23,664	8,664
Income tax benefit	-	-
Net stock-based compensation expense	\$ 23,664	\$ 8,664
Effect on earnings per share:		
Basic earnings per share	\$ (0.001)	\$ (0.001)
Diluted earnings per share	\$ (0.001)	\$ (0.001)

As of October 31, 2012, estimated stock based compensation expense to be recognized in future periods for granted nonvested stock options amounted to approximately \$60,000. These nonvested stock options compensation expense will be recognized in a weighted average period of approximately 1.2 years.

The fair value of stock-based awards to employees is calculated using the Black-Scholes option pricing model. The Black-Scholes model requires subjective assumptions, including future stock price volatility and expected time to exercise, which greatly affect the calculated values. The expected term of the option has been estimated using the "simplified" method as provided in ("SEC") Staff Accounting Bulletin No. 107. Under this method, the expected term equals the arithmetic average of the vesting term and the contractual term of the option. The risk-free rate is based on the U.S. Treasury rates in effect during the corresponding period of grant. The expected volatility is based on the historical volatility of the Company's stock price. These factors could change in the future, which would affect fair values of stock options granted in such future periods, and could cause volatility in the total amount of the stock-based compensation expense reported in future periods.

The following weighted average assumptions were used to estimate the fair value of stock options granted for the years ended October 31, 2012 and 2011:

	Year ended October 31,	
	2012	2011
Expected dividend yield	0.0%	0.0%
Expected stock price volatility	40.9%	74.3%
Risk free interest rate	0.3%	1.0%
Expected life of options	3.5 years	3.2 years
Weighted average fair value of options granted	\$ 0.2227	\$ 0.1299

As of October 31, 2012 and 2011, the aggregate intrinsic value of options outstanding was approximately \$180,000 and \$91,000, respectively. The aggregate intrinsic value represents the difference between the Company's stock price at year end and the exercise price, multiplied by the number of in-the money options had all option holders exercised their options. This amount changes based on the fair market value of the Company's stock. For the years ended October 31, 2012 and 2011, no stock options were exercised.

NOTE I - CONCENTRATION OF RISKS

Cash and cash equivalents

The Company maintains domestic cash deposits in an FDIC insured bank and in a money market obligations trust registered under the US Investment Company Act of 1940, as amended. The domestic deposit balances usually exceed federally insured limits. In the foreign markets we serve, we also maintain cash deposits in foreign banks, which tend to be not significant and have no specific insurance. No losses have been experienced or are expected on these accounts.

Accounts receivable and revenues

Management deems all its accounts receivable to be fully collectible, and, as such, does not maintain any allowances for uncollectible receivables.

The Company's revenues, and the related receivables, are concentrated in the pharmaceutical industry in Puerto Rico, the United States of America and Ireland. Although few customers represent a significant source of revenue, the Company's functions are not a continuous process, accordingly, the client base for which the services are typically rendered, on a project-by-project basis, changes regularly.

The Company provided a substantial portion of its services to three customers, who accounted for 10% or more of its revenues in either of the years ended October 31, 2012 or 2011. During the year ended October 31, 2012 revenues from these customers were 12%, 10% and 10%, or a total of 32%, as compared to the same period last year for 5%, 18% and 15%, or a total of 38%, respectively. At October 31, 2012 and 2011 amounts due from these customers represented 33% and 29% of total accounts receivable balance, respectively.

The major customer information in the above paragraph is based on revenues earned from said customers at the segment level because in management's opinion contracts by segments are totally independent of each other, and therefore such information is more meaningful to the reader. These revenues pertain to a global group of affiliated companies which in the aggregate represented 54% and 52% of total Company's revenues for the years ended October 31, 2012 and 2011, respectively. At October 31, 2012 and 2011 amounts due from this global group of affiliated companies represented 51% and 37% of total accounts receivable balance, respectively.

NOTE J - RETIREMENT PLAN

Pharma-PR has a qualified profit sharing plan in accordance with the provision of Section 1165(a)(3)(A) of the Puerto Rico Code, for employees who meet certain age and service period requirements. The Company makes contributions to this plan as required by the provisions of the plan document. Following plan provisions, since the year ended October 31, 2009 the Company temporarily suspended contributions to the plan. During the year ended October 31, 2012 the Company started again to make contributions. Contributions for the year ended October 31, 2012 were \$75,144.

NOTE K - SEGMENT DISCLOSURES

The Company's segments are based on the organizational structure for which financial results are regularly evaluated by the Company's chief operating decision maker to determine resource allocation and assess performance. Each reportable segment is managed by its own management team and reports to executive management. The Company has four reportable segments: (i) Puerto Rico technical compliance consulting, (ii) United States technical compliance consulting, (iii) Ireland technical compliance consulting, and (iv) a Puerto Rico microbiological and chemical laboratory testing division ("Lab"). These reportable segments provide services primarily to the pharmaceutical, chemical, medical device and biotechnology industries in their respective markets.

The following table presents information about the reported revenue from services and earnings from operations of the Company for the year ended in October 31, 2012 and 2011. There is no intersegment revenue for the mentioned periods. Corporate expenses that support the operating units have been allocated to the segments. Asset information by reportable segment is not presented, since the Company does not produce such information internally, nor does it use such data to manage its business.

	Year ended October 31,	
	2012	2011
REVENUES:		
Puerto Rico consulting	\$ 15,213,662	\$ 9,050,694
United States consulting	9,159,539	5,868,049
Ireland consulting	3,211,758	3,322,126
Lab (microbiological and chemical testing)	857,948	754,814
Other segments ¹	784,260	937,499
Total consolidated revenues	\$ 29,227,167	\$ 19,933,182
INCOME (LOSS) BEFORE TAXES:		
Puerto Rico consulting	\$ 3,664,238	\$ 2,040,994
United States consulting	2,407,190	1,258,077
Ireland consulting	(245,431)	(71,049)
Lab (microbiological and chemical testing)	(381,406)	(299,348)
Other segments ¹	311,021	535,104
Total consolidated income before taxes	\$ 5,755,612	\$ 3,463,778

¹ Other segments represent activities that fall below the reportable threshold and are carried out in Puerto and United States. These activities include a technical seminars/training division, an information technology services and consulting division, and corporate headquarters, as applicable.

Long lived assets (property and equipment and intangible assets) and related depreciation and amortization expense for the year ended October 31, 2012 and 2011, were concentrated in the domestic markets (Puerto Rico and United States). The aggregate amount of long lived assets for the international operations (Ireland) is considered insignificant.

