Genesis Pharmaceuticals Enterprises, Inc.
Form 10-K/A
April 10, 2009

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

Amendment No. 1
FORM 10-K/A

(Mark One)

x ANNUAL REPORT UNDER SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the fiscal year ended: June 30, 2008

o TRANSITION REPORT UNDER SECTION 13 OR 15(d) OF THE SECURITIES EXCHANGE ACT OF 1934

For the transition period from _____ to _____

Commission file number: 000-53037

GENESIS PHARMACEUTICALS ENTERPRISES, INC.
(Name of small business issuer in its charter)

Florida	65-1130026
(State or other jurisdiction of incorporation or organization)	(IRS Employer Identification No.)

Middle Section, Longmao Street, Area A, Laiyang Waixiangxing Industrial Park
Laiyang City, Yantai, Shandong Province, People's Republic of China 265200
(Address of principle executive offices)

(0086) 535-7282997
(Issuer's telephone number)

Securities registered under Section 12(b) of the Exchange Act:
None

Securities registered under Section 12(g) of the Exchange Act:
Common Stock, $0.001 Par Value

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

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

Indicate by check mark whether the registrant (1) has filed all reports required to be filed by Section 13 or 15(d) of the Securities Exchange Act of 1934 during the preceding 12 months (or for such shorter period that the registrant was

required to file such report(s)), and (2) has been subject to such filing requirements for the past 90 days. Yes x No ¨

Indicate by check mark if disclosure of delinquent filers pursuant to Item 405 of Regulation S-K is not contained herein, and will not be contained, to the best of 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 definitions of "large accelerated filer," "accelerated filer," and smaller reporting companies in Rule 12b-2 of the Exchange Act.

Large accelerated filer	Accelerated filer
Non-accelerated filer (Do not check if a smaller reporting company)	Smaller reporting company x

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

The aggregate market value of the voting and non-voting common equity held by non-affiliates of the registrant, based upon the closing sale price of the registrant's common stock on December 31, 2007 as reported on the OTC Bulletin Board was approximately $50.4 million (5,454,384 shares at $9.24). Approximately 4,887,000 shares of common stock held by each officer and director and by each person who owns 10% or more of the outstanding common stock have been excluded because such persons may be deemed to be affiliates.

The number of outstanding shares of the registrant's common stock on September 29, 2008 was 10,330,344.

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Explanatory Note:

This Annual Report on Form 10-K/A is being filed as Amendment No. 1 to our Annual Report on Form 10-K for the year ended June 30, 2008, which was originally filed with the Securities and Exchange Commission on September 29, 2008. We are amending:

- Part II, Item 7 to provide additional information related to our revenue increase disclosed in the “Management's Discussion and Analysis or Plan of Operations - Comparison of Years Ended June 30, 2008 and June 30, 2007” and “Management's Discussion and Analysis or Plan of Operations - Comparison of Years Ended June 30, 2007 and June 30, 2006”:
- Part II, Item 7 to revise the table of the contractual obligations to include interest related to our contractual obligations disclosed in the “Management's Discussion and Analysis or Plan of Operations - Contractual Obligations and Off-Balanced Sheet Arrangements”;
- Part III Item 13 to revise and provide additional information in the “Certain Relationships and Related Transactions and Director Independence – Related Parties Transactions of Laiyang Jiangbo” regarding certain obligations and filing a lease agreement as Exhibit 10.11; and
- Part IV, Item 15 in the “Exhibit, Financial Statement Schedules” to include Exhibit 10.11 that was omitted from the original Annual Report and to revise and update certain disclosure in the notes to the financial statements including Note 1 – “Organization and business”, Note 2 – “Summary of significant accounting policies” under the sub-headings “Principals of consolidation” and “Investments and restricted investments”,, Note 4- “Supplemental disclosure of cash flow information”, and Note 12 – Taxes payable” under the paragraph sub-heading “ Income Taxes”. In addition, new officer certifications are filed as exhibits to this Amendment No. 1.

The cover page to this Amendment No. 1 has also been revised to correct the file number that was incorrect on the cover page of the original filing due to clerical error. None of these changes rose to the level of a restatement of our financial statements. Except as specifically referenced herein, this Amendment No. 1 to Annual Report on Form 10-K/A does not reflect any event occurring subsequent to September 29, 2008, the filing date of the original report.

CAUTIONARY STATEMENT REGARDING FORWARD-LOOKING INFORMATION

Certain statements in this annual report on Form 10K contain or may contain forward-looking statements that are subject to known and unknown risks, uncertainties and other factors which may cause actual results, performance or achievements to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements. These forward-looking statements were based on various factors and were derived utilizing numerous assumptions and other factors that could cause our actual results to differ materially from those in the forward-looking statements. These factors include, but are not limited to, economic, political and market conditions and fluctuations, government and industry regulation, interest rate risk, global competition, and other factors as relate to our doing business within the People's Republic of China. Most of these factors are difficult to predict accurately and are generally beyond our control. You should consider the areas of risk described in connection with any forward-looking statements that may be made herein. Readers are cautioned not to place undue reliance on these forward-looking statements, which speak only as of the date of this report. Readers should carefully review this annual report in its entirety, including but not limited to our financial statements and the notes thereto and the risks described in "Item 1. Description of Business—Risk Factors." Except for our ongoing obligations to disclose material information under the Federal securities laws, we undertake no obligation to release publicly any revisions to any forward-looking statements, to report events or to report the occurrence of unanticipated events.

When used in this annual report, the terms the "Company," "Genesis," "GNPH," "we," "us," "our," and similar terms refer to Genesis Pharmaceuticals Enterprises, Inc., a Florida corporation, and our subsidiaries. The information which appears on our website www.genesis-china.net is not part of this report.

PART I

ITEM 1. DESCRIPTION OF BUSINESS

We operate, control and beneficially own the pharmaceutical business of Laiyang Jiangbo. Laiyang Jiangbo researches, develops, manufactures, markets and sells pharmaceutical products and health supplements in the PRC. From our inception in 2001 until our acquisition of Karmoya International Ltd. ("Karmoya") in October 2007, we were a business development and marketing firm specializing in advising and providing turn-key solutions for Chinese small and mid-sized companies entering Western markets.

On June 3, 2008, our board of directors and the majority holders of our capital stock approved amendments to our Articles of Incorporation to increase our authorized common stock from 600,000,000 shares to 900,000,000 shares (the "July 2008 Authorized Share Amendment"). The Certificate of Amendment and Certificate of Change to our Articles of Incorporation to effect the July 2008 Authorized Share Amendment was filed with Florida's Secretary of State on July 29, 2008.

On July 27, 2008, our board of directors and the majority holders of our capital stock approved a one-for-forty reverse stock split of our common stock. On August 29, 2008, we received confirmation from the Department of the State of Florida that the Articles of Amendment to the Amended and Restated Articles of Incorporation ("August 2008 Amended Articles of Incorporation") to effect a reverse stock split was duly filed and on September 3, 2008, the reverse stock split was effectuated. Following the reverse stock split, the total number of shares of our common stock outstanding was reduced from 412,986,078 shares to approximately 10,325,000 shares. Pursuant to the August 2008 Amended Articles of Incorporation, the maximum number of shares of common stock that the Company is authorized to issue was also reduced from 900,000,000 to 22,500,000. The financial statements have been retroactively adjusted to reflect the reverse split. Additionally, all share representations are on a post-split basis hereinafter.

Corporate Structure

The following diagram illustrates our current corporate structure:

CONTRACTUAL ARRANGEMENTS WITH LAIYANG JIANGBO AND ITS SHAREHOLDERS

Our relationships with Laiyang Jiangbo and its shareholders are governed by a series of contractual arrangements primarily between two entities associated with our wholly owned subsidiary Karmoya: (1) GJBT, Karmoya's wholly foreign owned enterprise in PRC, and (2) Laiyang Jiangbo, Karmoya's operating company in PRC. Under PRC laws, each of GJBT and Laiyang Jiangbo is an independent legal person and neither of them is exposed to liabilities incurred by the other party. The contractual arrangements constitute valid and binding obligations of the parties of such agreements. Each of the contractual arrangements, as amended and restated, and the rights and obligations of the parties thereto are enforceable and valid in accordance with the laws of the PRC. Other than pursuant to the contractual arrangements described below, Laiyang Jiangbo does not transfer any other funds generated from its operations to any other member of the LJ Group. On September 21, 2007, we entered into the following contractual arrangements (collectively, the " LJ Agreements"):

Consulting Services Agreement. Pursuant to the exclusive consulting services agreement between GJBT and Laiyang Jiangbo, GJBT has the exclusive right to provide to Laiyang Jiangbo general consulting services related to pharmaceutical business operations, as well as consulting services related to human resources and technological research and development of pharmaceutical products and health supplements (the " Services"). Under this agreement, GJBT owns the intellectual property rights developed or discovered through research and development while providing the Services for Laiyang Jiangbo. Laiyang Jiangbo pays a quarterly consulting service fee in Chinese Renminbi (" RMB") to GJBT that is equal to all of Laiyang Jiangbo's revenue for such quarter.

Operating Agreement. Pursuant to the operating agreement among GJBT, Laiyang Jiangbo and the shareholders of Laiyang Jiangbo who collectively hold 100% of the outstanding shares of Laiyang Jiangbo (collectively, the " Laiyang Shareholders"), GJBT provides guidance and instructions on Laiyang Jiangbo's daily operations, financial management and employment issues. The Laiyang Shareholders must appoint the candidates recommended by GJBT as members of Laiyang Jiangbo's board of directors. GJBT has the right to appoint senior executives of Laiyang Jiangbo. In addition, GJBT agrees to guarantee Laiyang Jiangbo's performance under any agreements or arrangements relating to Laiyang Jiangbo's business arrangements with any third party. Laiyang Jiangbo, in return, agrees to pledge its accounts receivable and all of its assets to GJBT. Moreover, Laiyang Jiangbo agrees that without the prior consent of GJBT, Laiyang Jiangbo will not engage in any transactions that could materially affect the assets, liabilities, rights or operations of Laiyang Jiangbo, including, but not limited to, incurrence or assumption of any indebtedness, sale or purchase of any assets or rights, incurrence of any encumbrance on any of its assets or intellectual property rights in favor of a third party, or transfer of any agreements relating to its business operation to any third party. The term of this agreement is ten (10) years from September 21, 2007 unless early termination occurs in accordance with the provisions of the agreement and may be extended only upon GJBT's written confirmation prior to the expiration of the this agreement, with the extended term to be mutually agreed upon by the parties.

Equity Pledge Agreement. Pursuant to the equity pledge agreement among GJBT, Laiyang Jiangbo and the Laiyang Shareholders, the Laiyang Shareholders pledged all of their equity interests in Laiyang Jiangbo to GJBT to guarantee Laiyang Jiangbo's performance of its obligations under the consulting services agreement. If either Laiyang Jiangbo or any of the Laiyang Shareholders breaches its respective contractual obligations, GJBT, as pledgee, will be entitled to certain rights, including the right to sell the pledged equity interests. The Laiyang Shareholders also granted GJBT an exclusive, irrevocable power of attorney to take actions in the place and stead of the Laiyang Shareholders to carry out the security provisions of the equity pledge agreement and take any action and execute any instrument that GJBT may deem necessary or advisable to accomplish the purposes of the equity pledge agreement. The Laiyang Shareholders agreed, among other things, not to dispose of the pledged equity interests or take any actions that would prejudice GJBT's interest. The equity pledge agreement will expire two (2) years after Laiyang Jiangbo obligations under the exclusive consulting services agreement have been fulfilled.

Option Agreement. Pursuant to the option agreement among GJBT, Laiyang Jiangbo and the Laiyang Shareholders, the Laiyang Shareholders irrevocably granted GJBT or its designated person an exclusive option to purchase, to the extent permitted under PRC law, all or part of the equity interests in Laiyang Jiangbo for the cost of the initial contributions to the registered capital or the minimum amount of consideration permitted by applicable PRC law. GJBT or its designated person has sole discretion to decide when to exercise the option, whether in part or in full. The term of this agreement is ten (10) years from September 21, 2007 unless early termination occurs in accordance with the provisions of the agreement and may be extended only upon GJBT's written confirmation prior to the expiration of the this agreement, with the extended term to be mutually agreed upon by the parties.

Proxy Agreement. Pursuant to the proxy agreement among GJBT and the Laiyang Shareholders, the Laiyang Shareholders agreed to irrevocably grant and entrust all the rights to exercise their voting power to the person(s) appointed by GJBT. GJBT may from time to time establish and amend rules to govern how GJBT shall exercise the powers granted to it by the Laiyang Shareholders, and GJBT shall take action only in accordance with such rules. The Laiyang Shareholders shall not transfer their equity interests in Laiyang Jiangbo to any individual or company (other than GJBT or the individuals or entities designated by GJBT). The Laiyang Shareholders acknowledged that they will continue to perform this agreement even if one or more than one of them no longer hold the equity interests of Laiyang Jiangbo. This agreement may not be terminated without the unanimous consent of all of the parties, except that GJBT may terminate this agreement by giving thirty (30) days prior written notice to the Laiyang Shareholders.

Recent Financings

May 2008 Financing

On May 30, 2008, we entered into a Securities Purchase Agreement (the "May 2008 Securities Purchase Agreement"), pursuant to which, on May 30, 2008, we sold to investors $30,000,000 principal amount of our 6% Notes and Class A Warrants to purchase 1,875,000 shares of our common stock, in transactions exempt from registration under the Securities Act. Pursuant to the terms of the May 2008 Securities Purchase Agreement, we will use the net proceeds for working capital purposes.

The Notes are due May 30, 2011 and are convertible into shares of our common stock at a conversion price of $8.00 per share, subject to adjustment pursuant to customary anti-dilution provisions and automatic downward adjustments in the event of certain sales or issuances by us of common stock at a price per share less than $8.00 on a post-split basis. Interest on the outstanding principal balance of the notes is payable at the rate of 6% per annum, in semi-annual installments payable on November 30th and May 30th of each year, with the first interest payment due on November 30, 2008. The Class A Warrants are exercisable for a five-year period beginning on May 30, 2008 at an initial exercise price of $10.00 per share.

In connection with May 2008 financing, Mr. Wubo Cao, our chief executive officer and chairman of the board, placed 3,750,000 shares of our common stock owned by him into an escrow account pursuant to a Make Good Escrow Agreement, dated as of May 30, 2008. In the event that either (i) our adjusted 2008 earnings before taxes is less than $26,700,000 ("2008 Guaranteed EBT") or (ii) our 2008 adjusted fully diluted earnings before taxes per share is less than $1.60 per share ("2008 Guaranteed Diluted EBT"), 1,500,000 of such shares (the "2008 Make Good Shares") are to be released pro rata to the investors. In the event that either (i) our adjusted 2009 earnings before taxes is less than US $38,400,000 ("2009 Guaranteed EBT") or (ii) our adjusted fully diluted earnings before taxes per share is less than $2.32 (or US $2.24, if the 500,000 shares of common stock held in escrow, in connection with the November 2007 financing have been released from escrow)("2009 Guaranteed Diluted EBT"), 2,250,000 of such shares, (the "2009 Make Good Shares") are to be released pro rata to the Investors. Should we successfully satisfy these respective financial milestones, the 2008 Make Good Shares and 2009 Make Good Shares will be returned to Mr. Cao. In addition, Mr. Cao is required to deliver shares of common stock owned by him to the investors on a pro rata basis equal to the number of shares (the "Settlement Shares") required to satisfy all costs and expenses associated with the settlement of all legal and other matters pertaining to the Company prior to or in connection with the completion of the our October 2007 share exchange in accordance with formulas set forth in the Securities Purchase Agreement.

In connection with the May 2008 financing, we entered into a Registration Rights Agreement dated as of May 30, 2008 with the investors. Pursuant to the Registration Rights Agreement, we agreed to file a registration statement covering the resale of (i) the shares of common stock underlying the Notes and Class A Warrants that are being registered in this offering, (ii) the 2008 Make Good Shares, (iii) the 2009 Make Good Shares, and (iv) the Settlement Shares. We were required to file an initial registration statement covering the shares of common stock underlying the notes and warrants no later than 45 days from the closing of the May 2008 financing and to have such registration statement declared effective no later than 180 days from the closing of May 2008 financing. If we do not timely file such registration statement or cause it to be declared effective by the required dates, then we will be required to pay liquidated damages to the investors equal to 1.0% of the aggregate purchase price paid by such investors for each month that we do not file the registration statement or cause it to be declared effective. Notwithstanding the foregoing, in no event shall liquidated damages exceed 10% of the aggregate amount of the purchase price. We filed an initial registration statement on July 14, 2008 to satisfy our obligations under the registration rights agreement. In connection with the May 2008 financing, we and the purchaser of our Debentures and November Warrants, agreed that such securities shall be included in this registration statement. See "November 2007 Financing" below.

In connection with the May 2008 financing, Mr. Cao entered into a Lock-Up Agreement dated May 30, 2008 with us, pursuant to which he agreed not to transfer any shares of our common stock owned by him until 18 months after the effective date of the registration statement discussed above.

November 2007 Financing

On November 6, 2007, we entered into a Securities Purchase Agreement (the "November Securities Purchase Agreement") with Pope Investments, LLC, pursuant to which, on November 7, 2007, we issued and sold to Pope Investments (i) $5,000,000 principal amount of our debentures and (ii) warrants to purchase 250,000 shares of common stock at an exercise price of 512.80 per share, subject to adjustment as provided therein. The exercise price and number of shares for which the warrants are exercisable were adjusted to 400,000 post-split shares at $8.00 per share, in connection with the May 2008 financing.

The debentures bear interest at the rate of 6% per annum, payable in semi-annual installments on May 31 and November 30 of each year, with the first interest payment being due on May 31, 2008. The initial conversion price of the debentures was $10.00 per share. If we issue common stock at a price that is less than the effective conversion price, or common stock equivalents with an exercise or conversion price less than the then effective conversion price, the conversion price of the debenture and the exercise price of the warrant will be reduced to such price. The exercise price of the debentures was reduced to $8.00 per share in connection with the May 2008 financing. The debentures may not be prepaid without the prior written consent of the holder.

Pursuant to the November Securities Purchase Agreement, the we entered into a Registration Rights Agreement (the "November Registration Rights Agreement"), pursuant to which we must file on each Filing Date (as defined therein) a registration statement to register the portion of the Registrable Securities (as defined therein) as permitted by the SEC's guidance.

Pursuant to the November Registration Rights Agreement, the initial registration statement with respect to the shares of common stock issuable upon conversion of the Debentures and exercise of the November Warrants was required to be filed within 90 days of the November 7, 2007 closing date and declared effective within 180 days following such closing date. Any subsequent registration statements that are required to be filed on the earliest practical date on which we are permitted by the SEC's guidance to file such additional registration statement. Such additional registration statements must he effective 90 days following the date on which it is required to be filed. In the event that the registration statement was not timely filed or declared effective, we were required, pursuant to the November registration rights agreement to pay liquidated damages. Such liquidated damages shall be, at the investor's option, either $81,643.83 or 165 shares of our common stock per day that the registration statement is not timely filed or declared effective as required pursuant to the November registration rights agreement, subject to an amount of liquidated damages not exceeding either $600,000, 60,000 shares of common stock, or a combination thereof based upon 12% liquidated damages in the aggregate. In connection with the May 2008 financing, Pope Investments waived the initial filing and effectiveness deadlines set forth in the November registration rights agreement and agreed that we would be required to include the Registrable Securities covered by the November Registration Rights Agreement in the Registration Rights Agreement executed in connection with the May 2008 financing.

LAIYANG JIANGBO PHARMACEUTICAL CO., LTD.

As discussed above, our operations are conducted through Laiyang Jiangbo Pharmaceutical Co., Ltd., ("Laiyang Jiangbo") a limited liability company headquartered in the PRC and organized under the laws of PRC (" Laiyang Jiangbo"). Laiyang Jiangbo was organized on August 18, 2003, and its fiscal year end is June 30.

PRINCIPAL PRODUCTS OR SERVICES

Laiyang Jiangbo is engaged in research, development, production, marketing and sales of pharmaceutical products. It is located in Northeast China in an Economic Development Zone in Laiyang City, Shandong province, and is one of the major pharmaceutical companies in China producing tablets, capsules, and granules for both Western medical drugs and Chinese herbal-based medical drugs. Laiyang Jiangbo is also a major manufacturer of liquid chemical supply for medical use in China. Approximately 20% of its current products are Chinese herbal-based drugs and 80% are Western medical drugs. Laiyang Jiangbo has several Certificates of Good Manufacturing Practices for Pharmaceutical Products (GMP Certificates) issued by the Shandong State Drug Administration (SDA) and currently produces six types of drugs.

Laiyang Jiangbo's top four products in fiscal 2008 include Clarithromycin sustained-release tablets, Itopride Hydrochloride granules, Ciprofloxacin Hydrochloride tablets, and Baobaole Chewable tablets.

Drug Development and Production

Development and production of pharmaceutical products is Laiyang Jiangbo's largest and most profitable business. Its principal pharmaceutical products include:

Clarithromycin sustained-release tablets

Clarithromycin sustained-release tablets, Chinese Drug Approval Number H20052746, are semi-synthetic antibiotics for curing Clarithromycin sensitive microorganism infections. Laiyang Jiangbo is one of only two domestic Chinese pharmaceutical companies that has the technology to manufacture and actively produce and sell this drug. The Company's sales of this drug were approximately RMB 337.2 million (US $46.4 million) with gross margin over 75% in fiscal 2008, which is approximately 50% of the market share in China for this type of drug.

Clarithromycin is the second generation of macrolide antibiotic and replaces the older generation of Erythromycin. Clarithromycin first entered the pharmaceutical market in Ireland in 1989, and as of 2007, it is one of thirty medicines which generate the greatest sales revenue all over the world. Chemically, Clarithromycin has a wider antimicrobial spectrum and longer duration of acid resistance. Its activity is 2 to 4 times better than Erythromycin, but the toxicity is 2-12 times lower.

Clarithromycin sustained-release tablets utilize sustained-release technology, which requires a high degree of production technology. Because of the high degree of technology required to produce this product, PRC production requirements are very strict and there are very few manufacturers who gain permission to produce this product. Therefore, there is a significant barrier to entry in the PRC market. Currently, our Clarithromycin sustained-release tablets are the leading product in the PRC domestic antibiotic sustained-release tablets market. Our goal is to maintain our current market share for this product.

Itopride Hydrochloride granules

Itopride Hydrochloride granules, Chinese Drug Approval Number H20050932, are a stomach and intestinal drug for curing digestive system-related diseases. The Company's sales for this drug reached RMB 258.1 million (US $35.5

million) with gross margin over 85% in fiscal 2008, and the Company has approximately 10-12% of the market share in China for this type of drug. This product is widely regarded for its pharmacological properties, i.e., rapid absorption, positive clinical effects, and few side effects. Based on clinical observation, it has been shown that Itopride Hydrochloride granules can improve 95.1% of gastrointestinal indigestion symptoms.

Itopride Hydrochloride granules are the fourth generation of gastrointestinal double dynamic medicines, which are used for curing most symptoms due to functional indigestion. The older generations are Metoclopramide Paspertin, Domperidone and Cisapride.

Itopride Hydrochloride granules are SDA-approved and entered the PRC pharmaceutical market in June 2005. Since 2005, Laiyang Jiangbo has seized the opportunity presented by this product by rapidly establishing a domestic sales network and developing the market for this product. Currently, this product has competition from two other famous stomach medicines, namely Dompendone Tablets and Vitamin U Belladonna and Aluminum Capsules II. Itopride Hydrochloride granules are a new product for Laiyang Jiangbo, but it already has a nationwide sales network in China. The Company’s goal is to maintain the current market share and profit margin for this product.

Ciprofloxacin Hydrochloride tablets

Ciprofloxacin Hydrochloride tablets, Chinese Drug Approval Number H37022737, are an antibiotic drug used to cure infection caused by bacteria. The Company’s sales for this drug was approximately RMB 18.1 million (US $2.5 million) in fiscal 2008.

Due to a stoppage in production of raw material manufacturing in PRC in 2004, the price of certain raw materials which are used to produce Ciprofloxacin Hydrochloride tablets rose rapidly and Laiyang Jiangbo seized this opportunity by using its stored raw materials to produce a significant amount of Ciprofloxacin Hydrochloride tablets. As a result, Laiyang Jiangbo's sales of this product won a large percentage of the market in PRC from 2004 to 2006. However, other companies resumed production in 2007, which has lead to stronger competition and a decrease in Laiyang Jiangbo's profits for this product. As both the sales volume and profit decreased for this product, the Company is not actively promoting this product and only continues to produce Ciprofloxacin Hydrochloride tablets to support the Company's product variety and brand name.

Paracetamol tablets

Paracetamol tablets, Chinese Drug Approval Number H37022733, are a nonprescription analgesic drug, mainly used for curing fever due to common flu or influenza. It is also used for relief of aches and pains. The Company's sales for this drug was approximately RMB 3.6 million ($500,000) in fiscal 2008. Laiyang Jiangbo is authorized by the PRC Ministry of Health to be an appointed producer of common antibiotics in Jiangsu Province, Guangdong Province, Zhejiang Province, Fujian Province, Shandong Province and Guangxi Province. Paracetamol tablets are one of PRC's national A-level Medicare medicines. This product entered the Chinese market in July 2004. As the sales volume and profit both significantly decreased in recent years, the Company plans to gradually exit the market for this product in fiscal 2009.

Baobaole Chewable tablets

Baobaole Chewable tablets, Chinese Drug Approval Number Z20060294, are a new product of Laiyang Jiangbo and formally entered the market in November 2007. Baobaole Chewable tablets are nonprescription over-the counter drugs for gastric cavity aches. This drug stimulates the appetite and promotes digestion. Baobaole is used to cure deficiencies in the spleen and stomach, abdomen aches, loss of appetite, and loose bowels. Its effects are mild and lasting. The drug has quickly gained its popularity in the market and the sales for this drug has grown at a fast pace since its initial introduction.

The Company's sales for Baobaole Chewable tablets was approximately RMB 95.1 million (US $13.1 million) with gross margin over 80% in fiscal 2008. The Company plans to continue expanding the distribution network for this product and actively promote the drugs to sustain the product's sales growth.

Radix Isatidis Disperable Tablet

Radix Isatidis Disperable Tablets, Chinese Drug Approval Number Z20080142, nonprescription Traditional Chinese Medicine, is used to cure virus influenza and sour throat. Laiyang Jiangbo recently obtained the approval for this drug and is the only company that owns this manufacturing technology in China. It clears away heat, detoxifies and promote pharynx. The research study indicates Radix Isatidisthe's ingredients included Indole, hapoxanthineuraci, quina-alkaloids, amino acid, etc., have anti-inflammation and anti-virus effects.

Compared with similar existing Radix Isatidis products, Radix Isatidis Disperable Tablet utilizes the new disperable tablet formula, which is convenient to take and fast to dissolve. It is also easy to absorb and has high stability. The product was first introduced to the market in September 2008. The Company plans to formally sell this product in early October 2008 and will heavily promote this drug through advertising and various promotional activities.

RESEARCH AND DEVELOPMENT

For the fiscal year ended June 30, 2008, Laiyang Jiangbo spent approximately US $3.2 million or approximately 3.3% of its fiscal 2008 revenue on research and development of products. For the fiscal year ended June 30, 2007, Laiyang

Jiangbo spent approximately US $11 million or approximately 14.6% of its fiscal 2007 revenue on research and development of various pharmaceutical products.

Laiyang Jiangbo places great emphasis on product research and development and maintains strategic relationships with many research institutions in PRC developing new drugs, such as Pharmaceutical Institute of Shandong University, The Institute of Microbiology and Shandong Chinese Traditional Medicine Technical School. These relationships help to ensure that Laiyang Jiangbo maintains a continuing pipeline of high quality drugs with market potential into the future. Other than a number of potential R&D projects that are currently under evolution and yet to be locked in, the Company currently has three products pending on PRC SFDA approval in the pipeline for commercialization in China. Additionally, the Company also is negotiating to purchase a Class I drug that is currently being developed. The products are-

Drug Name	Target Treatment/Drug Type	Status
Felodipine Sustained Release Tablets	Treat high blood pressure and arteriosclerosis/Western Drug	(A) Expected approval date - second quarter of fiscal year 2009
Yuandu Hanbi Capsules	Relieve arthritis pain /Traditional Chinese Medicine	(A) Expected approval date - second quarter of fiscal year 2009
Bezoar Yijin Tablets	Cures inflammations such as pharyngitis/Traditional Chinese Medicine	(A) Expected approval date - second quarter of fiscal year 2009

(A) Subject to SFDA. Pending administrative protection and approval.

Ligustrazine Ferulic Acid Acetate (LFAA)

LFAA is a Cardiac Cerebral Vascular innovative medicine, researched by Pharmaceutical Institute of Shandong University that is undergoing Phase III (final) clinical studies and for which the Company is in the process of negotiating a drug purchase contract. LFAA is protected by the patent of invention in China. Its PRC invention patent application number is 02135989X, publication number is CN1424313A and patent number is ZL02135989X filed in December 2005.

LFAA is a synthetic innovation medicine based on Liqustrazine. It is the successor of Liqustrazine, which has independent intellectual property rights. LFAA helps to reduce blood clotting and prevent platelets in the blood from clumping together. Based on clinical studies, LFAA’s artery endothelium cell proliferation stimulating function is better than Liqustrazine in a number of measures. Laiyang Jiangbo is currently in the process of finalizing the drug patent and manufacturing right purchase agreement with Pharmaceutical Institute of Shandong University. The Company anticipates being able to start producing LFAA in fiscal 2010.

DISTRIBUTION METHODS OF THE PRODUCTS OR SERVICES AND OUR CUSTOMERS

Laiyang Jiangbo has a well-established sales network across China. It has a distribution network covering over 30 provinces and regions in the PRC. Currently, Laiyang Jiangbo has approximately 1,100 distribution agents and salespeople throughout the PRC. Laiyang Jiangbo will continue to establish more representative offices and engage additional distribution agents in order to strengthen its distribution network.

Laiyang Jiangbo recognizes the importance of branding as well as packaging. All of Laiyang Jiangbo’s products bear a uniform brand but have specialized designs to differentiate the different categories of Laiyang Jiangbo's products.

Laiyang Jiangbo conducts promotional marketing activities to publicize and enhance its image as well as to reinforce the recognition of its brand name including:

1. publishing advertisements and articles in national as well as specialized and provincial newspapers, magazines, and in other media, including the Internet;

2. participating in national meetings, seminars, symposiums, exhibitions for pharmaceutical and other related industries;

3. organizing cooperative promotional activities with distributors; and

4. sending direct mail to major physician offices and laboratories.

CUSTOMERS

Currently, Laiyang Jiangbo has approximately 1,200 terminal clients. Terminal clients are hospitals and medical institutions which purchase large supplies of pharmaceutical drugs. Laiyang Jiangbo is also authorized by the PRC Ministry of Health as an appointed Medicare medication supplier in six provinces, namely Jiangsu Province, Shandong Province, Zhejiang Province, Fujian Province, Guangdong Province and Guangxi Province.

For the fiscal years ended June 30, 2008, 2007 and 2006, five customers accounted for approximately 18.1%, 33.3% and 30.5%, respectively, of Laiyang Jiangbo's sales. These five customers represented 11.8% and 24.6% of Laiyang Jiangbo's total accounts receivable as of June 30, 2008 and 2007, respectively.

COMPETITION

As a pharmaceutical manufacturing and distribution company in PRC, overall, Laiyang Jiangbo has two major competitors in the PRC: Zhuhai Lizhu and Beijing Nohua. These companies have number of popular pharmaceutical products, strong financial position and a large market share in the industry. Laiyang Jiangbo is able to compete with these competitors because of its favorable geographic position, strong R&D capability, unique products, extensive sales network, and lower prices.

Our major competitors in China on individual product basis are Jiangsu Hengrui Pharmaceuticals (Clarithromycin sustained release tablets), Xi'an Yangsen (Itopride Hydrochloride Granules) and Jiangzhong Pharmaceuticals (Baobaole Chewable tablets), respectively. We are able to compete with Jiangsu Hengrui Pharmaceuticals because of our extensive sales network as well as flexible and favorable incentive policy. Compared with Motihium of Xi'an Yangsen, a gastro dynamic only drug, our Itopride Hydrochloride Granules has better efficacy due to its gastro-intestinal dynamic characteristic, higher security and less side effects. Referring to Children Jiangwei Xiaoshi Tablets of Jiangzhong Pharmaceutcials, our Baobaole Chewable tablet is able to significantly stimulate appetite and fundamentally nurse children's gastro-intestinal system. Also, it is very convenient for children to take. As such, we believe we have competitive advantages for those products.

SOURCES AND AVAILABILITY OF RAW MATERIALS AND THE PRINCIPAL SUPPLIERS

Laiyang Jiangbo has strategic relationships with many research institutions in PRC developing new drugs, such as Jiangsu Drug Research Institute, Pharmaceutical Institute of Shandong University, Chinese Traditional Medicine Institute, Shandong Chinese Traditional Medicine Technical School, and the Institute for Drug Control Departments. These relationships help to ensure that Laiyang Jiangbo maintains a continuing pipeline of high quality drugs into the future. Laiyang Jiangbo’s own production facilities supply most of the raw materials used to manufacture its products. Laiyang Jiangbo designs, creates prototypes and manufactures its products at its manufacturing facilities located in Laiyang City, Shandong province. Its principal raw materials include Ciprofloxacin Hydrochloride tablets. The prices for these raw materials are subject to market forces largely beyond our control, including energy costs, organic chemical prices, market demand, and freight costs. The prices for these raw materials have varied significantly in the past and may vary significantly in the future.

INTELLECTUAL PROPERTY

Laiyang Jiangbo relies on a combination of trademarks copyright and trade secret protection laws in the PRC and other jurisdictions, as well as confidentiality procedures and contractual provisions to protect its intellectual property and brand. Laiyang Jiangbo has been issued design patents in the PRC for drug packaging and drug containers, each valid for 10 years, and it intends to apply for more patents to protect its core technologies. Laiyang Jiangbo is currently in the process of acquiring the rights to a new Class I drug recently patented and made available to Laiyang Jiangbo through its relationship with the Pharmaceutical Institute of Shandong University. This is a Class I drug which means that all PRC national hospitals and other major medical facilities must carry this drug. Laiyang Jiangbo also enters into confidentiality, non-compete and invention assignment agreements with its employees and consultants and nondisclosure agreements with third parties. “Jiangbo” and a certain circular design affiliated with our brand are our registered trademarks in the PRC.

Pharmaceutical companies are at times involved in litigation based on allegations of infringement or other violations of intellectual property rights. Furthermore, the application of laws governing intellectual property rights in the PRC and abroad is uncertain and evolving and could involve substantial risks to us.

GOVERNMENT REGULATION

General PRC Government Approval

The Drug Administration Law of the PRC governs Laiyang Jiangbo and its products. The State Food & Drug Administration of the PRC regulates and implements PRC drug laws. The State FDA has granted Laiyang Jiangbo government permits to produce the following products: Clarithromycin sustained-released tablets, Itopride Hydrochloride granules, Ciprofloxacin Hydrochloride tablets, Paracetamol tablets, Baobaole Chewable tablets, Compound Sufamethoxazole tablets, and Vitamin C tablets.

The drug approval process takes about two years: including local SFDA approval, Local SFDA test, State SFDA processing, state SFDA expert valuation, clinical trial, and final approval.

No enterprise may start production at its facilities until it receives approval from the PRC Ministry of Agriculture to begin operations. Laiyang Jiangbo currently has obtained the requisite approval and licenses from the Ministry of Agriculture in order to operate its production facilities.

Circular 106 Compliance and Approval

On May 31, 2007, the PRC State Administration of Foreign Exchange (“SAFE”) issued an official notice known as "Circular 106," which requires the owners of any Chinese companies to obtain SAFE’s approval before establishing any offshore holding company structure for foreign financing as well as subsequent acquisition matters in China.

In early September 2007, the three owners of 100% of the equity in Laiyang Jiangbo, Cao Wubo, Xun Guihong and Zhang Yihua, submitted their application to SAFE. On September 19, 2007, SAFE approved their application, permitting these Chinese citizens to establish an offshore company, Karmoya International Ltd., as a “special purpose vehicle” for any foreign ownership and capital raising activities by Laiyang Jiangbo.

After SAFE’s approval, Cao Wubo, Xun Guihong and Zhang Yihua became the majority owners of Karmoya International Ltd. on September 20, 2007.

COSTS AND EFFECTS OF COMPLIANCE WITH ENVIRONMENTAL LAWS

In compliance with PRC environmental regulations, Laiyang Jiangbo spent approximately $2,750 in fiscal 2008, $2,000 in fiscal 2007, and approximately $1,600 in fiscal 2006, mainly for the wastewater treatment in connection with its production facilities.

EMPLOYEES

Laiyang Jiangbo currently has more than 1,430 employees, including 320 production crew, 440 full-time salespersons and 620 part-time salespersons. Approximately 200 of these employees are represented by Laiyang City Jiangbo Pharmaceuticals Union, which is governed by the City of Laiyang. Laiyang Jiangbo has not experienced a work stoppage since inception and does not anticipate any work stoppage in the foreseeable future. Management believes that its relations with its employees and the union are good.

CORPORATE INFORMATION

Laiyang Jiangbo's principal executive offices are located at Middle Section, Longmao Street, Area A, Laiyang Waixiangxing Industrial Park, Laiyang City, Yantai, Shandong Province, PRC 265200.

ITEM 1A. RISK FACTORS

Investing in our securities involves a great deal of risk. Careful consideration should be made of the following factors as well as other information included in this prospectus before deciding to purchase our common stock. You should pay particular attention to the fact that we conduct all of our operations in China and are governed by a legal and regulatory environment that in some respects differs significantly from the environment that may prevail in other countries. Our business, financial condition or results of operations could be affected materially and adversely by any or all of these risks.

THE FOLLOWING MATTERS MAY HAVE A MATERIAL ADVERSE EFFECT ON OUR BUSINESS, FINANCIAL CONDITION, LIQUIDITY, RESULTS OF OPERATIONS OR PROSPECTS, FINANCIAL OR OTHERWISE. REFERENCE TO THIS CAUTIONARY STATEMENT IN THE CONTEXT OF A FORWARD-LOOKING STATEMENT OR STATEMENTS SHALL BE DEEMED TO BE A STATEMENT THAT ANY ONE OR MORE OF THE FOLLOWING FACTORS MAY CAUSE ACTUAL RESULTS TO DIFFER MATERIALLY FROM THOSE IN SUCH FORWARD-LOOKING STATEMENT OR STATEMENTS.

Risks Relating to Our Business

Our limited operating history makes it difficult to evaluate our future prospects and results of operations.

We have a limited operating history. Laiyang Jiangbo commenced operations in 2003 and first achieved profitability in the fiscal year ended June 30, 2005. Accordingly, you should consider our future prospects in light of the risks and uncertainties experienced by early stage companies in evolving industries such as the pharmaceutical industry in China. Some of these risks and uncertainties relate to our ability to:

- maintain our market position in the pharmaceuticals business in China;
- offer new and innovative products to attract and retain a larger customer base;
- attract additional customers and increase spending per customer;

- increase awareness of our brand and continue to develop user and customer loyalty;
- respond to competitive market conditions;
- respond to changes in our regulatory environment;
- manage risks associated with intellectual property rights;
- maintain effective control of our costs and expenses;
- raise sufficient capital to sustain and expand our business;

- attract, retain and motivate qualified personnel; and
- upgrade our technology to support additional research and development of new products.

If we are unsuccessful in addressing any of these risks and uncertainties, our business may be materially and adversely affected.

We may need additional financing to execute our business plan.

The revenues from the production and sale of pharmaceutical products and the projected revenues from these products may not be adequate to support our expansion and product development programs. We may need substantial additional funds to build our new production facilities, pursue further research and development, obtain regulatory approvals, market our products, and file, prosecute, defend and enforce our intellectual property rights. We will seek additional funds through public or private equity or debt financing, strategic transactions and/or from other sources. We could enter into collaborative arrangements for the development of particular products that would lead to our relinquishing some or all rights to the related technology or products.

There are no assurances that future funding will be available on favorable terms or at all. If additional funding is not obtained, we will need to reduce, defer or cancel development programs, planned initiatives or overhead expenditures, to the extent necessary. The failure to fund our capital requirements would have a material adverse effect on our business, financial condition and results of operations.

Our success depends on collaborative partners, licensees and other third parties over whom we have limited control.

Due to the complexity of the process of developing pharmaceuticals, our core business depends on arrangements with pharmaceutical institutes, corporate and academic collaborators, licensors, licensees and others for the research, development, clinical testing, technology rights, manufacturing, marketing and commercialization of our products. We have several research collaborations. Our license agreements could obligate us to diligently bring potential products to market, make milestone payments and royalties that, in some instances, could be substantial, and incur the costs of filing and prosecuting patent applications. There are no assurances that we will be able to establish or maintain collaborations that are important to our business on favorable terms, or at all.

A number of risks arise from our dependence on collaborative agreements with third parties. Product development and commercialization efforts could be adversely affected if any collaborative partner:

- terminates or suspends its agreement with us
- causes delays
- fails to timely develop or manufacture in adequate quantities a substance needed in order to conduct clinical trials
- fails to adequately perform clinical trials
- determines not to develop, manufacture or commercialize a product to which it has rights or
- otherwise fails to meet its contractual obligations.

Our collaborative partners could pursue other technologies or develop alternative products that could compete with the products we are developing.

The profitability of our products will depend in part on our ability to protect proprietary rights and operate without infringing the proprietary rights of others.

The profitability of our products will depend in part on our ability to obtain and maintain patents and licenses and preserve trade secrets, and the period our intellectual property remains exclusive. We must also operate without infringing the proprietary rights of third parties and without third parties circumventing our rights. The patent positions of pharmaceutical enterprises, including ours, are uncertain and involve complex legal and factual questions for which important legal principles are largely unresolved. The pharmaceutical patent situation outside the US is uncertain, is currently undergoing review and revision in many countries, and may not protect our intellectual property rights to the same extent as the laws of the US. Because patent applications are maintained in secrecy in some cases, we cannot be certain that we or our licensors are the first creators of inventions described in our pending patent applications or patents or the first to file patent applications for such inventions.

Most of our drug products have been approved by the PRC's Food and Drug Administration (SFDA) but have not received patent protection. For instance, Clarithromycin sustained-release tablets, one of our most profitable products, are produced by other companies in China. If any other company were to obtain patent protection for Clarithromycin sustained-release tablets in China, or for any of our other drug products, it would have a material adverse effect on our revenue.

Other companies may independently develop similar products and design around any patented products we develop. We cannot assure you that:

- any of our patent applications will result in the issuance of patents;
- we will develop additional patentable products;
- the patents we have been issued will provide us with any competitive advantages;
- the patents of others will not impede our ability to do business; or
- third parties will not be able to circumvent our patents.

A number of pharmaceutical, research, and academic companies and institutions have developed technologies, filed patent applications or received patents on technologies that may relate to our business. If these technologies, applications or patents conflict with ours, the scope of our current or future patents could be limited or our patent applications could be denied. Our business may be adversely affected if competitors independently develop competing technologies, especially if we do not obtain, or obtain only narrow, patent protection. If patents that cover our activities are issued to other companies, we may not be able to obtain licenses at a reasonable cost, or at all; develop our technology; or introduce, manufacture or sell the products we have planned.

Patent litigation is becoming widespread in the pharmaceutical industry. Such litigation may affect our efforts to form collaborations, to conduct research or development, to conduct clinical testing or to manufacture or market any products under development. There are no assurances that our patents would be held valid or enforceable by a court or that a competitor's technology or product would be found to infringe our patents in the event of patent litigation. Our business could be materially affected by an adverse outcome to such litigation. Similarly, we may need to participate in interference proceedings declared by the U.S. Patent and Trademark Office or equivalent international authorities to determine priority of invention. We could incur substantial costs and devote significant management resources to defend our patent position or to seek a declaration that another company's patents are invalid.

Much of our know-how and technology may not be patentable, though it may constitute trade secrets. There are no assurances that we will be able to meaningfully protect our trade secrets. We cannot assure you that any of our existing confidentiality agreements with employees, consultants, advisors or collaborators will provide meaningful protection for our trade secrets, know-how or other proprietary information in the event of any unauthorized use or disclosure. Collaborators, advisors or consultants may dispute the ownership of proprietary rights to our technology, for example by asserting that they developed the technology independently.

We may encounter difficulties in manufacturing our products.

Before our products can be profitable, they must be produced in commercial quantities in a cost-effective manufacturing process that complies with regulatory requirements, including GMP, production and quality control regulations. If we cannot arrange for or maintain commercial-scale manufacturing on acceptable terms, or if there are delays or difficulties in the manufacturing process, we may not be able to conduct clinical trials, obtain regulatory approval or meet demand for our products. Production of our products could require raw materials which are scarce or

which can be obtained only from a limited number of sources. If we are unable to obtain adequate supplies of such raw materials, the development, regulatory approval and marketing of our products could be delayed.

We could need more clinical trials or take more time to complete our clinical trials than we have planned.

Clinical trials vary in design by factors including dosage, end points, length, and controls. We may need to conduct a series of trials to demonstrate the safety and efficacy of our products. The results of these trials may not demonstrate safety or efficacy sufficiently for regulatory authorities to approve our products. Further, the actual schedules for our clinical trials could vary dramatically from the forecasted schedules due to factors including changes in trial design, conflicts with the schedules of participating clinicians and clinical institutions, and changes affecting product supplies for clinical trials.

We rely on collaborators, including academic institutions, governmental agencies and clinical research organizations, to conduct, supervise, monitor and design some or all aspects of clinical trials involving our products. Since these trials depend on governmental participation and funding, we have less control over their timing and design than trials we sponsor. Delays in or failure to commence or complete any planned clinical trials could delay the ultimate timelines for our product releases. Such delays could reduce investors' confidence in our ability to develop products, likely causing our share price to decrease.

We may not be able to obtain the regulatory approvals or clearances that are necessary to commercialize our products.

The PRC and other countries impose significant statutory and regulatory obligations upon the manufacture and sale of pharmaceutical products. Each regulatory authority typically has a lengthy approval process in which it examines pre-clinical and clinical data and the facilities in which the product is manufactured. Regulatory submissions must meet complex criteria to demonstrate the safety and efficacy of the ultimate products. Addressing these criteria requires considerable data collection, verification and analysis. We may spend time and money preparing regulatory submissions or applications without assurances as to whether they will be approved on a timely basis or at all.

Our product candidates, some of which are currently in the early stages of development, will require significant additional development and pre-clinical and clinical testing prior to their commercialization. These steps and the process of obtaining required approvals and clearances can be costly and time-consuming. If our potential products are not successfully developed, cannot be proven to be safe and effective through clinical trials, or do not receive applicable regulatory approvals and clearances, or if there are delays in the process:

- the commercialization of our products could be adversely affected;
- any competitive advantages of the products could be diminished; and
- revenues or collaborative milestones from the products could be reduced or delayed.

Governmental and regulatory authorities may approve a product candidate for fewer indications or narrower circumstances than requested or may condition approval on the performance of post-marketing studies for a product candidate. Even if a product receives regulatory approval and clearance, it may later exhibit adverse side effects that limit or prevent its widespread use or that would force us to withdraw the product from the market.

Any marketed product and its manufacturer will continue to be subject to strict regulation after approval. Results of post-marketing programs may limit or expand the further marketing of products. Unforeseen problems with an approved product or any violation of regulations could result in restrictions on the product, including its withdrawal from the market and possible civil actions.

In manufacturing our products we will be required to comply with applicable good manufacturing practices regulations, which include requirements relating to quality control and quality assurance, as well as the maintenance of records and documentation. If we cannot comply with regulatory requirements, including applicable good manufacturing practice requirements, we may not be allowed to develop or market the product candidates. If we or our manufacturers fail to comply with applicable regulatory requirements at any stage during the regulatory process, we may be subject to sanctions, including fines, product recalls or seizures, injunctions, refusal of regulatory agencies to review pending market approval applications or supplements to approve applications, total or partial suspension of production, civil penalties, withdrawals of previously approved marketing applications and criminal prosecution.

Competitors may develop and market pharmaceutical products that are less expensive, more effective or safer, making our products obsolete or uncompetitive.

Some of our competitors and potential competitors have greater product development capabilities and financial, scientific, marketing and human resources than we do. Technological competition from pharmaceutical companies is intense and is expected to increase. Other companies have developed technologies that could be the basis for competitive products. Some of these products have an entirely different approach or means of accomplishing the desired curative effect than products we are developing. Alternative products may be developed that are more effective, work faster and are less costly than our products. Competitors may succeed in developing products earlier than us, obtaining approvals and clearances for such products more rapidly than us, or developing products that are

more effective than ours. In addition, other forms of treatment may be competitive with our products. Over time, our technology or products may become obsolete or uncompetitive.

Our products may not gain market acceptance.

Our products may not gain market acceptance in the pharmaceutical community. The degree of market acceptance of any product depends on a number of factors, including establishment and demonstration of clinical efficacy and safety, cost-effectiveness, clinical advantages over alternative products, and marketing and distribution support for the products. Limited information regarding these factors is available in connection with our products or products that may compete with ours.

To directly market and distribute our pharmaceutical products, we or our collaborators require a marketing and sales force with appropriate technical expertise and supporting distribution capabilities. We may not be able to further establish sales, marketing and distribution capabilities or enter into arrangements with third parties on acceptable terms. If we or our partners cannot successfully market and sell our products, our ability to generate revenue will be limited.

Our operations and the use of our products could subject us to damages relating to injuries or accidental contamination.

Our research and development processes involve the controlled use of hazardous materials. We are subject to PRC national, provincial and local laws and regulations governing the use, manufacture, storage, handling and disposal of such materials and waste products. The risk of accidental contamination or injury from handling and disposing of such materials cannot be completely eliminated. In the event of an accident involving hazardous materials, we could be held liable for resulting damages. We are not insured with respect to this liability. Such liability could exceed our resources. In the future we could incur significant costs to comply with environmental laws and regulations.

If we were successfully sued for product liability, we could face substantial liabilities that may exceed our resources.

We may be held liable if any product we develop, or any product which is made using our technologies, causes injury or is found unsuitable during product testing, manufacturing, marketing, sale or use. These risks are inherent in the development of agricultural and pharmaceutical products. We currently do not have product liability insurance. We are not insured with respect to this liability. If we choose to obtain product liability insurance but cannot obtain sufficient insurance coverage at an acceptable cost or otherwise protect against potential product liability claims, the commercialization of products that we develop may be prevented or inhibited. If we are sued for any injury caused by our products, our liability could exceed our total assets.

We have limited business insurance coverage.

The insurance industry in China is still at an early stage of development. Insurance companies in China offer limited business insurance products. We do not have any business liability or disruption insurance coverage for our operations in China. Any business disruption, litigation or natural disaster may result in our incurring substantial costs and the diversion of our resources.

Our business depends substantially on the continuing efforts of our executive officers and our ability to maintain a skilled labor force, and our business may be severely disrupted if we lose their services.

Our future success depends substantially on the continued services of our executive officers, especially Wubo Cao our chief executive officer and the chairman of our board. We do not maintain key man life insurance on any of our executive officers. If one or more of our executive officers are unable or unwilling to continue in their present positions, we may not be able to replace them readily, if at all. Therefore, our business may be severely disrupted, and we may incur additional expenses to recruit and retain new officers. In addition, if any of our executives joins a competitor or forms a competing company, we may lose some of our customers.

Our success depends on attracting and retaining qualified personnel.

We depend on a core management and scientific team. The loss of any of these individuals could prevent us from achieving our business objective of commercializing our product candidates. Our future success will depend in large part on our continued ability to attract and retain other highly qualified scientific, technical and management personnel, as well as personnel with expertise in clinical testing and government regulation. We face competition for personnel from other companies, universities, public and private research institutions, government entities and other organizations. If our recruitment and retention efforts are unsuccessful, our business operations could suffer.

We may not be able to manage the expansion of our operations effectively, which may have an adverse effect on our business and results of operations.

The revenues from the production and sale of our current product offerings and the projected revenues from these products may not be adequate to support our expansion and product development programs. We will need substantial additional funds to expand our production facilities, pursue research and development, obtain regulatory approvals; file, prosecute, defend and enforce our intellectual property rights and market our products. We will seek additional funds through public or private equity or debt financing, strategic transactions and/or from other sources. We could enter into collaborative arrangements for the development of particular products that would lead to our relinquishing some or all rights to the related technology or products. There are no assurances that future funding will be available on favorable terms or at all. If additional funding is not obtained, we will need to reduce, defer or cancel development programs, planned initiatives or overhead expenditures, to the extent necessary. The failure to fund our capital requirements would have a material adverse effect on our business, financial condition and results of operations.

Risks Related to Our Corporate Structure

PRC laws and regulations governing our businesses and the validity of certain of our contractual arrangements are uncertain. If we are found to be in violation, we could be subject to sanctions. In addition, changes in such PRC laws and regulations may materially and adversely affect our business.

There are substantial uncertainties regarding the interpretation and application of PRC laws and regulations, including, but not limited to, the laws and regulations governing our business, or the enforcement and performance of our contractual arrangements with our affiliated Chinese entity, Laiyang Jiangbo, and its shareholders. We are considered a foreign person or foreign invested enterprise under PRC law. As a result, we are subject to PRC law limitations on foreign ownership of Chinese companies. These laws and regulations are relatively new and may be subject to change, and their official interpretation and enforcement may involve substantial uncertainty. The effectiveness of newly enacted laws, regulations or amendments may be delayed, resulting in detrimental reliance by foreign investors. New laws and regulations that affect existing and proposed future businesses may also be applied retroactively.

The PRC government has broad discretion in dealing with violations of laws and regulations, including levying fines, revoking business and other licenses and requiring actions necessary for compliance. In particular, licenses and permits issued or granted to us by relevant governmental bodies may be revoked at a later time by higher regulatory bodies. We cannot predict the effect of the interpretation of existing or new PRC laws or regulations on our businesses. We cannot assure you that our current ownership and operating structure would not be found in violation of any current or future PRC laws or regulations. As a result, we may be subject to sanctions, including fines, and could be required to restructure our operations or cease to provide certain services. Any of these or similar actions could significantly disrupt our business operations or restrict us from conducting a substantial portion of our business operations, which could materially and adversely affect our business, financial condition and results of operations.

The PRC government restricts foreign investment in pharmaceutical businesses in China. Accordingly, we operate our business in China through Laiyang Jiangbo. Laiyang Jiangbo holds the licenses and approvals necessary to operate our pharmaceutical business in China. We have contractual arrangements with Laiyang Jiangbo and its shareholders that allow us to substantially control Laiyang Jiangbo. We cannot assure you, however, that we will be able to enforce these contracts.

Although we believe we comply with current PRC regulations, we cannot assure you that the PRC government would agree that these operating arrangements comply with PRC licensing, registration or other regulatory requirements, with existing policies or with requirements or policies that may be adopted in the future. If the PRC government determines that we do not comply with applicable law, it could revoke our business and operating licenses, require us to discontinue or restrict our operations, restrict our right to collect revenues, require us to restructure our operations, impose additional conditions or requirements with which we may not be able to comply, impose restrictions on our business operations or on our customers, or take other regulatory or enforcement actions against us that could be harmful to our business.

We may be adversely affected by complexity, uncertainties and changes in PRC regulation of pharmaceutical business and companies, including limitations on our ability to own key assets.

The PRC government regulates the pharmaceutical industry including foreign ownership of, and the licensing and permit requirements pertaining to, companies in the pharmaceutical industry. These laws and regulations are relatively new and evolving, and their interpretation and enforcement involve significant uncertainty. As a result, in certain circumstances it may be difficult to determine what actions or omissions may be deemed to be a violation of applicable laws and regulations. Issues, risks and uncertainties relating to PRC government regulation of the pharmaceutical industry include the following:

- we only have contractual control over Laiyang Jiangbo. We do not own it due to the restriction of foreign investment in Chinese businesses; and

- uncertainties relating to the regulation of the pharmaceutical business in China, including evolving licensing practices, means that permits, licenses or operations at our company may be subject to challenge. This may disrupt our business, or subject us to sanctions, requirements to increase capital or other conditions or enforcement, or compromise enforceability of related contractual arrangements, or have other harmful effects on us.

The interpretation and application of existing PRC laws, regulations and policies and possible new laws, regulations or policies have created substantial uncertainties regarding the legality of existing and future foreign investments in, and the businesses and activities of, pharmaceutical businesses in China, including our business.

Our contractual arrangements with Laiyang Jiangbo and its shareholders may not be as effective in providing control over these entities as direct ownership.

Since the law of the PRC limits foreign equity ownership in pharmaceutical companies in China, we operate our business through Laiyang Jiangbo. We have no equity ownership interest in Laiyang Jiangbo and rely on contractual arrangements to control and operate such business. These contractual arrangements may not be effective in providing control over Laiyang Jiangbo as direct ownership. For example, Laiyang Jiangbo could fail to take actions required for our business despite its contractual obligation to do so. If Laiyang Jiangbo fails to perform under its agreements with us, we may have to incur substantial costs and resources to enforce such arrangements and may have to rely on legal remedies under the law of the PRC, which may not be effective. In addition, we cannot assure you that Laiyang Jiangbo's shareholders would always act in our best interests.

The chairman of the board of directors of Laiyang Jiangbo has potential conflicts of interest with us, which may adversely affect our business.

Mr. Cao Wubo, our Chairman and Chief Executive Officer, is also the Chairman of the Board of Directors and General Manager of Laiyang Jiangbo. Conflicts of interests between his duties to our company and Laiyang Jiangbo may arise. As Mr. Cao is a director and executive officer of our company, he has a duty of loyalty and care to us under Florida law when there are any potential conflicts of interests between our company and Laiyang Jiangbo. We cannot assure you, however, that when conflicts of interest arise, Mr. Cao will act completely in our interests or that conflicts of interests will be resolved in our favor. In addition, Mr. Cao could violate his legal duties by diverting business opportunities from us to others. If we cannot resolve any conflicts of interest between us and Mr. Cao, we would have to rely on legal proceedings, which could result in the disruption of our business.

Risks Related to Doing Business in China

Failure to comply with PRC regulations relating to the establishment of offshore special purpose companies by PRC residents may subject our PRC resident shareholders to personal liability, limit our ability to acquire PRC companies or to inject capital into our PRC subsidiaries, limit our PRC subsidiaries' ability to distribute profits to us or otherwise materially adversely affect us.

In October 2005, the PRC State Administration of Foreign Exchange, or SAFE, issued the Notice on Relevant Issues in the Foreign Exchange Control over Financing and Return Investment Through Special Purpose Companies by Residents Inside China, generally referred to as Circular 75, which required PRC residents to register with the competent local SAFE branch before establishing or acquiring control over an offshore special purpose company, or SPV, for the purpose of engaging in an equity financing outside of China on the strength of domestic PRC assets originally held by those residents. Internal implementing guidelines issued by SAFE, which became public in June 2007 (known as Notice 106), expanded the reach of Circular 75 by (i) purporting to cover the establishment or acquisition of control by PRC residents of offshore entities which merely acquire "control" over domestic companies or assets, even in the absence of legal ownership; (ii) adding requirements relating to the source of the PRC resident's funds used to establish or acquire the offshore entity; (iii) covering the use of existing offshore entities for offshore financings; (iv) purporting to cover situations in which an offshore SPV establishes a new subsidiary in China or acquires an unrelated company or unrelated assets in China; and (v) making the domestic affiliate of the SPV responsible for the accuracy of certain documents which must be filed in connection with any such registration, notably, the business plan which describes the overseas financing and the use of proceeds. Amendments to registrations made under Circular 75 are required in connection with any increase or decrease of capital, transfer of shares, mergers and acquisitions, equity investment or creation of any security interest in any assets located in China to guarantee offshore obligations, and Notice 106 makes the offshore SPV jointly responsible for these filings. In the case of an SPV which was established, and which acquired a related domestic company or assets, before the implementation date of Circular 75, a retroactive SAFE registration was required to have been completed before March 31, 2006; this date was subsequently extended indefinitely by Notice 106, which also required that the registrant establish that all foreign exchange transactions undertaken by the SPV and its affiliates were in compliance with applicable laws and regulations. Failure to comply with the requirements of Circular 75, as applied by SAFE in accordance with Notice 106, may result in fines and other penalties under PRC laws for evasion of applicable foreign exchange restrictions. Any such failure could also result in the SPV's affiliates being impeded or prevented from distributing their profits and the proceeds from any reduction in capital, share transfer or liquidation to the SPV, or from engaging in other transfers of funds into or out of China.

We believe our shareholders who are PRC residents, as defined in Circular 75, have registered with the relevant branch of SAFE, as currently required, in connection with their equity interests in us and our acquisitions of equity interests in our PRC subsidiaries. However, we cannot provide any assurances that their existing registrations have fully complied with, or that they have made all necessary amendments to their registration to fully comply with, all

applicable registrations or approvals required by Circular 75. Moreover, because of uncertainty over how Circular 75 will be interpreted and implemented, and how or whether SAFE will apply it to us, we cannot predict how it will affect our business operations or future strategies. For example, our present and prospective PRC subsidiaries' ability to conduct foreign exchange activities, such as the remittance of dividends and foreign currency-denominated borrowings, may be subject to compliance with Circular 75 by our PRC resident beneficial holders. In addition, such PRC residents may not always be able to complete the necessary registration procedures required by Circular 75. We also have little control over either our present or prospective direct or indirect shareholders or the outcome of such registration procedures. A failure by our PRC resident beneficial holders or future PRC resident shareholders to comply with Circular 75, if SAFE requires it, could subject these PRC resident beneficial holders to fines or legal sanctions, restrict our overseas or cross-border investment activities, limit our subsidiaries' ability to make distributions or pay dividends or affect our ownership structure, which could adversely affect our business and prospects.

If the PRC enacts regulations which forbid or restrict foreign investment, our ability to grow may be severely impaired.

We intend to expand our business in areas relating to our present business. We may also expand by making acquisitions of companies in related industries. Many of the rules and regulations that we would face are not explicitly communicated, and we may be subject to rules that would affect our ability to grow, either internally or through acquisition of other Chinese or foreign companies. There are also substantial uncertainties regarding the proper interpretation of current laws and regulations of the PRC. New laws or regulations that forbid foreign investment could severely impair our businesses and prospects. Additionally, if the relevant authorities find us in violation of PRC laws or regulations, they would have broad discretion in dealing with such a violation, including, without limitation:

- levying fines;
- revoking our business and other licenses; and
- requiring that we restructure our ownership or operations.

Any deterioration of political relations between the United States and the PRC could impair our operations and your investment in us.

The relationship between the United States and the PRC is subject to sudden fluctuation and periodic tension. Changes in political conditions in the PRC and changes in the state of Sino-U.S. relations are difficult to predict and could adversely affect our operations or cause potential acquisition candidates or their goods and services to become less attractive. Such a change could lead to a decline in our profitability. Any weakening of relations between the United States and the PRC could have a material adverse effect on our operations and your investment in us, particularly in our efforts to raise capital to expand our other business activities.

Adverse changes in economic and political policies of the PRC government could have a material adverse effect on the overall economic growth of China, which could adversely affect our business.

Substantially all of our business operations are conducted in China. Accordingly, our results of operations, financial condition and prospects are subject to a significant degree to economic, political and legal developments in China. China's economy differs from the economies of most developed countries in many respects, including with respect to:

- the amount of government involvement;
- level of development;
- growth rate;
- control of foreign exchange; and
- allocation of resources.

While the PRC economy has experienced significant growth in the past 20 years, growth has been uneven across different regions and among various economic sectors of China. The PRC government has implemented various measures to encourage economic development and guide the allocation of resources. Some of these measures benefit the overall PRC economy, but may also have a negative effect on us. For example, our financial condition and results of operations may be adversely affected by government control over capital investments or changes in tax regulations that are applicable to us. Since early 2004, the PRC government has implemented certain measures to control the pace of economic growth. Such measures may cause a decrease in the level of economic activity in China, which in turn could adversely affect our results of operations and financial condition.

Price controls may affect both our revenues and net income.

The laws of the PRC provide for the government to fix and adjust prices. Although we are not presently subject to price controls in connection with the sale of our products, it is possible that price controls may be imposed in the future. To the extent that we are subject to price control, our revenue, gross profit, gross margin and net income will be affected since the revenue we derive from our sales will be limited and, unless there is also price control on the products that we purchase from our suppliers, we may face no limitation on our costs. Further, if price controls affect both our revenue and our costs, our ability to be profitable and the extent of our profitability will be effectively subject

to determination by the applicable regulatory authorities in the PRC.

Our operations may not develop in the same way or at the same rate as might be expected if the PRC economy were similar to the market-oriented economies of OECD member countries.

The economy of the PRC has historically been a nationalistic, “planned economy,” meaning it functions and produces according to governmental plans and pre-set targets or quotas. In certain aspects, the PRC’s economy has been making a transition to a more market-oriented economy, although the government imposes price controls on certain products and in certain industries. However, we cannot predict the future direction of these economic reforms or the effects these measures may have. The economy of the PRC also differs from the economies of most countries belonging to the Organization for Economic Cooperation and Development (the “OECD”), an international group of member countries sharing a commitment to democratic government and market economy. For instance:

- the level of state-owned enterprises in the PRC, as well as the level of governmental control over the allocation of resources is greater than in most of the countries belonging to the OECD;
- the level of capital reinvestment is lower in the PRC than in other countries that are members of the OECD;
- the government of the PRC has a greater involvement in general in the economy and the economic structure of industries within the PRC than other countries belonging to the OECD;
- the government of the PRC imposes price controls on certain products and our products may become subject to additional price controls; and
- the PRC has various impediments in place that make it difficult for foreign firms to obtain local currency, as opposed to other countries belonging to the OECD where exchange of currencies is generally free from restriction.

As a result of these differences, our business may not develop in the same way or at the same rate as might be expected if the economy of the PRC were similar to those of the OECD member countries.

Because some of our officers and directors reside outside of the United States, it may be difficult for you to enforce your rights against them or enforce United States court judgments against them in the PRC.

Most of our executive officers and directors reside in the PRC and a substantial portion of our assets are located in the PRC. It may therefore be difficult for United States investors to enforce their legal rights, to effect service of process upon our directors or officers or to enforce judgments of United States courts predicated upon civil liabilities and criminal penalties of our directors and officers under federal securities laws. Further, it is unclear if extradition treaties now in effect between the United States and the PRC would permit effective enforcement of criminal penalties of the federal securities laws.

We may have limited legal recourse under Chinese law if disputes arise under contracts with third parties.

Almost all of our agreements with our employees and third parties, including our supplier and customers, are governed by the laws of the PRC. The legal system in the PRC is a civil law system based on written statutes. Unlike common law systems, such as we have in the United States, it is a system in which decided legal cases have little precedential value. The government of the PRC has enacted some laws and regulations dealing with matters such as corporate organization and governance, foreign investment, commerce, taxation and trade. However, their experience in implementing, interpreting and enforcing these laws and regulations is limited, and our ability to enforce commercial claims or to resolve commercial disputes is unpredictable. The resolution of these matters may be subject to the exercise of considerable discretion by agencies of the PRC, and forces unrelated to the legal merits of a particular matter or dispute may influence their determination. Any rights we may have to specific performance or to seek an injunction under Chinese law are severely limited, and without a means of recourse by virtue of the Chinese legal system, we may be unable to prevent these situations from occurring. The occurrence of any such events could have a material adverse effect on our business, financial condition and results of operations.

Because we may not be able to obtain business insurance in the PRC, we may not be protected from risks that are customarily covered by insurance in the United States.

Business insurance is not readily available in the PRC. To the extent that we suffer a loss of a type which would normally be covered by insurance in the United States, such as product liability and general liability insurance, we would incur significant expenses in both defending any action and in paying any claims that result from a settlement or judgment.

Failure to comply with the United States Foreign Corrupt Practices Act could subject us to penalties and other adverse consequences.

We are subject to the United States Foreign Corrupt Practices Act, which generally prohibits United States companies from engaging in bribery or other prohibited payments to foreign officials for the purpose of obtaining or retaining business. Foreign companies, including some that may compete with us, are not subject to these prohibitions. Corruption, extortion, bribery, pay-offs, theft and other fraudulent practices occur from time-to-time in the PRC. We can make no assurance, however, that our employees or other agents will not engage in such conduct for which we might be held responsible. If our employees or other agents are found to have engaged in such practices, we could suffer severe penalties and other consequences that may have a material adverse effect on our business, financial condition and results of operations.

A downturn in the economy of the PRC may slow our growth and profitability.

The growth of the Chinese economy has been uneven across geographic regions and economic sectors. There can be no assurance that growth of the Chinese economy will be steady or that any downturn will not have a negative effect on our business especially if it results in either a decreased use of products such as ours or in pressure on us to lower our prices. The Chinese economy has been transitioning from a planned economy to a more market-oriented economy. Although in recent years the Chinese government has implemented measures emphasizing the utilization of market forces for economic reform, the reduction of state ownership of productive assets and the establishment of sound corporate governance in business enterprises, a substantial portion of the productive assets in China is still owned by the Chinese government. The continued control of these assets and other aspects of the national economy by the Chinese government could materially and adversely affect our business. The Chinese government also exercises significant control over Chinese economic growth through the allocation of resources, controlling payment of foreign currency-denominated obligations, setting monetary policy and providing preferential treatment to particular industries or companies. Efforts by the Chinese government to slow the pace of growth of the Chinese economy could result in reduced demand for our products.

Any adverse change in the economic conditions or government policies in China could have a material adverse effect on the overall economic growth and the level of pharmaceutical investments and expenditures in China, which in turn could lead to a reduction in demand for our products and consequently have a material adverse effect on our businesses.

Downturns in the economies of the U.S. and Europe may affect the PRC economy which could reduce the demand for our products.

The rapid growth of the PRC economy in recent years has been partially related to the U.S. and European countries' demand for goods made in and exported from the PRC. The downturns in the U.S. and European economies may reduce the demand for goods exported by the PRC which could eventually affect the PRC economy as overseas orders decrease. The downturn in the PRC economy may in turn negatively impact the demand for our products.

If certain tax exemptions within the PRC regarding withholding taxes are removed, we may be required to deduct corporate withholding taxes from any dividends we may pay in the future.

Under the PRC's current tax laws, regulations and rulings, companies are exempt from paying withholding taxes with respect dividends paid to stockholders outside of the PRC. However, if the foregoing exemption is removed, we may be required to deduct certain amounts from any dividends we pay to our stockholders.

Laiyang Jiangbo is subject to restrictions on making payments to us.

We are a holding company incorporated in the State of Florida and do not have any assets or conduct any business operations other than our investments in our affiliated entity in China, Laiyang Jiangbo. As a result of our holding company structure, we rely entirely on payments from Laiyang Jiangbo under our contractual arrangements. The PRC government also imposes controls on the conversion of RMB into foreign currencies and the remittance of currencies out of China. We may experience difficulties in completing the administrative procedures necessary to obtain and remit foreign currency. See "Government control of currency conversion may affect the value of your investment." Furthermore, if our affiliated entity in China incurs debt on its own in the future, the instruments governing the debt may restrict its ability to make payments. If we are unable to receive all of the revenues from our operations through these contractual or dividend arrangements, we may be unable to pay dividends on our ordinary shares.

Uncertainties with respect to the PRC legal system could adversely affect us.

We conduct our business primarily through our affiliated Chinese entity, Laiyang Jiangbo. Our operations in China are governed by PRC laws and regulations. We are generally subject to laws and regulations applicable to foreign investments in China and, in particular, laws applicable to wholly foreign-owned enterprises. The PRC legal system is based on written statutes. Prior court decisions may be cited for reference but have limited precedential value.

Since 1979, PRC legislation and regulations have significantly enhanced the protections afforded to various forms of foreign investments in China. However, China has not developed a fully integrated legal system and recently enacted laws and regulations may not sufficiently cover all aspects of economic activities in China. In particular, because these laws and regulations are relatively new, and because of the limited volume of published decisions and their nonbinding nature, the interpretation and enforcement of these laws and regulations involve uncertainties. In addition, the PRC legal system is based in part on government policies and internal rules (some of which are not published on a timely basis or at all) that may have a retroactive effect. As a result, we may not be aware of our violation of these policies and rules until some time after the violation. In addition, any litigation in China may be protracted and result in substantial costs and diversion of resources and management attention.

You may experience difficulties in effecting service of legal process, enforcing foreign judgments or bringing original actions in China based on United States or other foreign laws against us, our management or the experts named in the prospectus.

We conduct substantially all of our operations in China and substantially all of our assets are located in China. In addition, most of our senior executive officers reside within China. As a result, it may not be possible to effect service of process within the United States or elsewhere outside China upon our senior executive officers, including with respect to matters arising under U.S. federal securities laws or applicable state securities laws. Moreover, our PRC counsel has advised us that the PRC does not have treaties with the United States or many other countries providing for the reciprocal recognition and enforcement of judgment of courts.

Governmental control of currency conversion may affect the value of your investment.

The PRC government imposes controls on the convertibility of RMB into foreign currencies and, in certain cases, the remittance of currency out of China. We receive substantially all of our revenues in RMB. Under our current structure, our income is primarily derived from payments from Laiyang Jiangbo. Shortages in the availability of foreign currency may restrict the ability of our PRC subsidiaries and our affiliated entity to remit sufficient foreign currency to pay dividends or other payments to us, or otherwise satisfy their foreign currency denominated obligations. Under existing PRC foreign exchange regulations, payments of current account items, including profit distributions, interest payments and expenditures from trade-related transactions, can be made in foreign currencies without prior approval from the PRC State Administration of Foreign Exchange by complying with certain procedural requirements. However, approval from appropriate government authorities is required where RMB is to be converted into foreign currency and remitted out of China to pay capital expenses such as the repayment of bank loans denominated in foreign currencies. The PRC government may also at its discretion restrict access in the future to foreign currencies for current account transactions. If the foreign exchange control system prevents us from obtaining sufficient foreign currency to satisfy our currency demands, we may not be able to pay dividends in foreign currencies to our shareholders.

Fluctuation in the value of RMB may have a material adverse effect on your investment.

The value of RMB against the U.S. dollar and other currencies may fluctuate and is affected by, among other things, changes in political and economic conditions. Our revenues and costs are mostly denominated in RMB, while a significant portion of our financial assets are denominated in U.S. dollars. We rely entirely on fees paid to us by our affiliated entity in China. Any significant fluctuation in value of RMB may materially and adversely affect our cash flows, revenues, earnings and financial position, and the value of, and any dividends payable on, our stock in U.S. dollars. For example, an appreciation of RMB against the U.S. dollar would make any new RMB denominated investments or expenditures more costly to us, to the extent that we need to convert U.S. dollars into RMB for such purposes. An appreciation of RMB against the U.S. dollar would also result in foreign currency translation losses for financial reporting purposes when we translate our U.S. dollar denominated financial assets into RMB, as RMB is our reporting currency.

We face risks related to health epidemics and other outbreaks.

Our business could be adversely affected by the effects of SARS or another epidemic or outbreak. China reported a number of cases of SARS in April 2004. Any prolonged recurrence of SARS or other adverse public health developments in China may have a material adverse effect on our business operations. For instance, health or other government regulations adopted in response may require temporary closure of our production facilities or of our offices. Such closures would severely disrupt our business operations and adversely affect our results of operations. We have not adopted any written preventive measures or contingency plans to combat any future outbreak of SARS or any other epidemic.

Risks Related to an Investment in Our Securities

We do not anticipate paying any cash dividends.

We presently do not anticipate that we will pay any dividends on any of our capital stock in the foreseeable future. The payment of dividends, if any, would be contingent upon our revenues and earnings, if any, capital requirements, and general financial condition. The payment of any dividends is within the discretion of our Board of Directors. We presently intend to retain all earnings, if any, to implement our business plan; accordingly, we do not anticipate the declaration of any dividends in the foreseeable future.

Because the OTC Bulletin Board is a quotation system, not an issuer listing service, market or exchange, it may be difficult for you to sell your common stock or you may not be able to sell your common stock for an optimum trading price.

The OTC Bulletin Board is a regulated quotation service that displays real-time quotes, last sale prices and volume limitations in over-the-counter securities. Because trades and quotations on the OTC Bulletin Board involve a manual process, the market information for such securities cannot be guaranteed. In addition, quote information, or even firm quotes, may not be available. The manual execution process may delay order processing and intervening price fluctuations may result in the failure of a limit order to execute or the execution of a market order at a significantly different price. Execution of trades, execution reporting and the delivery of legal trade confirmations may be delayed significantly. Consequently, one may not be able to sell shares of our common stock at the optimum trading prices.

The dealer’s spread (the difference between the bid and ask prices) may be large and may result in substantial losses to the seller of securities on the OTC Bulletin Board if the common stock or other security must be sold immediately. Further, purchasers of securities may incur an immediate “paper” loss due to the price spread. Moreover, dealers trading on the OTC Bulletin Board may not have a bid price for securities bought and sold through the OTC Bulletin Board. Due to the foregoing, demand for securities that are traded through the OTC Bulletin Board may be decreased or eliminated.

The application of the “penny stock” rules could adversely affect the market price of our common stock and increase your transaction costs to sell those shares.

In the event the trading price of our common shares reaches below $5 per share, the open-market trading of our common shares will be subject to the “penny stock” rules. The “penny stock” rules impose additional sales practice requirements on broker-dealers who sell securities to persons other than established customers and accredited investors (generally those with assets in excess of $1,000,000 or annual income exceeding $200,000 or $300,000 together with their spouse). For transactions covered by these rules, the broker-dealer must make a special suitability determination for the purchase of securities and have received the purchaser's written consent to the transaction before the purchase. Additionally, for any transaction involving a penny stock, unless exempt, the broker-dealer must deliver, before the transaction, a disclosure schedule prescribed by the SEC relating to the penny stock market. The broker-dealer also must disclose the commissions payable to both the broker-dealer and the registered representative and current quotations for the securities. Finally, monthly statements must be sent disclosing recent price information on the limited market in penny stocks. These additional burdens imposed on broker-dealers may restrict the ability or decrease the willingness of broker-dealers to sell our common shares, and may result in decreased liquidity for our common shares and increased transaction costs for sales and purchases of our common shares as compared to other securities.

Our common shares are thinly traded and, you may be unable to sell at or near ask prices or at all if you need to sell your shares to raise money or otherwise desire to liquidate your shares.

We cannot predict the extent to which an active public market for its common stock will develop or be sustained. However, we do not rule out the possibility of applying for listing on the Nasdaq National Market or other exchanges.

Our common shares have historically been sporadically or "thinly-traded" on the OTC Bulletin Board, meaning that the number of persons interested in purchasing our common shares at or near bid prices at any given time may be relatively small or non-existent. This situation is attributable to a number of factors, including the fact that we are a small company which is relatively unknown to stock analysts, stock brokers, institutional investors and others in the investment community that generate or influence sales volume, and that even if we came to the attention of such persons, they tend to be risk-averse and would be reluctant to follow an unproven company such as ours or purchase or recommend the purchase of our shares until such time as we became more seasoned and viable. As a consequence, there may be periods of several days or more when trading activity in our shares is minimal or non-existent, as compared to a seasoned issuer which has a large and steady volume of trading activity that will generally support continuous sales without an adverse effect on share price. We cannot give you any assurance that a broader or more active public trading market for our common stock will develop or be sustained, or that current trading levels will be sustained.

The market price for our common stock is particularly volatile given our status as a relatively small company with a small and thinly traded “float” and lack of current revenues that could lead to wide fluctuations in our share price. The price at which you purchase our common stock may not be indicative of the price that will prevail in the trading market. You may be unable to sell your common stock at or above your purchase price if at all, which may result in substantial losses to you.

The market for our common shares is characterized by significant price volatility when compared to seasoned issuers, and we expect that our share price will continue to be more volatile than a seasoned issuer for the indefinite future. The volatility in our share price is attributable to a number of factors. First, as noted above, our common shares are sporadically and/or thinly traded. As a consequence of this lack of liquidity, the trading of relatively small quantities of shares by our shareholders may disproportionately influence the price of those shares in either direction. The price for our shares could, for example, decline precipitously in the event that a large number of our common shares are sold on the market without commensurate demand, as compared to a seasoned issuer which could better absorb those

sales without adverse impact on its share price. Secondly, we are a speculative or "risky" investment due to our lack of revenues or profits to date and uncertainty of future market acceptance for our current and potential products. As a consequence of this enhanced risk, more risk-adverse investors may, under the fear of losing all or most of their investment in the event of negative news or lack of progress, be more inclined to sell their shares on the market more quickly and at greater discounts than would be the case with the stock of a seasoned issuer. The following factors may add to the volatility in the price of our common shares: actual or anticipated variations in our quarterly or annual operating results; adverse outcomes; the termination of our contractual agreements with Laiyang Jiangbo; and additions or departures of our key personnel, as well as other items discussed under this "Risk Factors" section, as well as elsewhere in this report. Many of these factors are beyond our control and may decrease the market price of our common shares, regardless of our operating performance. We cannot make any predictions or projections as to what the prevailing market price for our common shares will be at any time, including as to whether our common shares will sustain their current market prices, or as to what effect that the sale of shares or the availability of common shares for sale at any time will have on the prevailing market price. However, we do not rule out the possibility of applying for listing on the Nasdaq National Market or other exchanges.

Shareholders should be aware that, according to SEC Release No. 34-29093, the market for penny stocks has suffered in recent years from patterns of fraud and abuse. Such patterns include (1) control of the market for the security by one or a few broker-dealers that are often related to the promoter or issuer; (2) manipulation of prices through prearranged matching of purchases and sales and false and misleading press releases; (3) boiler room practices involving high-pressure sales tactics and unrealistic price projections by inexperienced sales persons; (4) excessive and undisclosed bid-ask differential and markups by selling broker-dealers; and (5) the wholesale dumping of the same securities by promoters and broker-dealers after prices have been manipulated to a desired level, along with the resulting inevitable collapse of those prices and with consequent investor losses. Our management is aware of the abuses that have occurred historically in the penny stock market. Although we do not expect to be in a position to dictate the behavior of the market or of broker-dealers who participate in the market, management will strive within the confines of practical limitations to prevent the described patterns from being established with respect to our securities. The occurrence of these patterns or practices could increase the volatility of our share price.

The market price for our stock may be volatile and the volatility in our common share price may subject us to securities litigation.

The market price for our stock may be volatile and subject to wide fluctuations in response to factors including the following:

- actual or anticipated fluctuations in our quarterly operating results;
- changes in financial estimates by securities research analysts;
- conditions in pharmaceutical and agricultural markets;
- changes in the economic performance or market valuations of other pharmaceutical companies;
- announcements by us or our competitors of new products, acquisitions, strategic partnerships, joint ventures or capital commitments;
- addition or departure of key personnel;
- fluctuations of exchange rates between RMB and the U.S. dollar;
- intellectual property litigation; and
- general economic or political conditions in China.

In addition, the securities market has from time to time experienced significant price and volume fluctuations that are not related to the operating performance of particular companies. These market fluctuations may also materially and adversely affect the market price of our stock.

The market for our common stock is characterized by significant price volatility when compared to seasoned issuers, and we expect that our share price will continue to be more volatile than a seasoned issuer for the indefinite future. In the past, plaintiffs have often initiated securities class action litigation against a company following periods of volatility in the market price of its securities. We may, in the future, be the target of similar litigation. Securities litigation could result in substantial costs and liabilities and could divert management's attention and resources.

Our corporate actions are substantially controlled by our principal shareholders and affiliated entities.

Our principal shareholders and their affiliated entities own approximately 53% of our outstanding common shares, representing approximately 53% of our voting power. These shareholders, acting individually or as a group, could exert substantial influence over matters such as electing directors and approving mergers or other business combination transactions. In addition, because of the percentage of ownership and voting concentration in these principal shareholders and their affiliated entities, elections of our board of directors will generally be within the control of these shareholders and their affiliated entities. While all of our shareholders are entitled to vote on matters submitted to our shareholders for approval, the concentration of shares and voting control presently lies with these principal shareholders and their affiliated entities. As such, it would be difficult for shareholders to propose and have approved proposals not supported by management. There can be no assurances that matters voted upon by our officers and directors in their capacity as shareholders will be viewed favorably by all shareholders of our company.

The elimination of monetary liability against our directors, officers and employees under Florida law and the existence of indemnification rights to our directors, officers and employees may result in substantial expenditures by us and may

discourage lawsuits against our directors, officers and employees.

Our articles of incorporation contain specific provisions that eliminate the liability of our directors for monetary damages to our company and shareholders, and we are prepared to give such indemnification to our directors and officers to the extent provided by Florida law. We may also have contractual indemnification obligations under our employment agreements with our officers. The foregoing indemnification obligations could result in our company incurring substantial expenditures to cover the cost of settlement or damage awards against directors and officers, which we may be unable to recoup. These provisions and resultant costs may also discourage our company from bringing a lawsuit against directors and officers for breaches of their fiduciary duties, and may similarly discourage the filing of derivative litigation by our shareholders against our directors and officers even though such actions, if successful, might otherwise benefit our company and shareholders.

Legislative actions, higher insurance costs and potential new accounting pronouncements may impact our future financial position and results of operations.

There have been regulatory changes, including the Sarbanes-Oxley Act of 2002, and there may potentially be new accounting pronouncements or additional regulatory rulings that will have an impact on our future financial position and results of operations. The Sarbanes-Oxley Act of 2002 and other rule changes are likely to increase general and administrative costs and expenses. In addition, insurers are likely to increase premiums as a result of high claims rates over the past several years, which we expect will increase our premiums for insurance policies. Further, there could be changes in certain accounting rules. These and other potential changes could materially increase the expenses we report under generally accepted accounting principles, and adversely affect our operating results.

Past activities of Genesis and its affiliates may lead to future liability.

Prior to the Exchange Agreement among Genesis, Karmoya and the Karmoya Shareholders executed on October 1, 2007, we engaged in businesses unrelated to our current operations. Neither Genesis’s prior management nor any of its shareholders prior to the Exchange Transaction are providing indemnifications against any loss, liability, claim, damage or expense arising out of or based on any breach of or inaccuracy in any of their representations and warranties made regarding such acquisition, and any liabilities relating to such prior business against which we are not completely indemnified may have a material adverse effect on our company. For example, we are aware of three lawsuits arising from past activities of Genesis, alleging breach of contract. Please see “Legal Proceedings” for more information.

We may need additional capital, and the sale of additional shares or other equity securities could result in additional dilution to our shareholders.

We believe that our current cash and cash equivalents, anticipated cash flows from operations and the net proceeds from a proposed offering will be sufficient to meet our anticipated cash needs for the near future. We may, however, require additional cash resources due to changed business conditions or other future developments, including any investments or acquisitions we may decide to pursue. If our resources are insufficient to satisfy our cash requirements, we may seek to sell additional equity or debt securities or obtain a credit facility. The sale of additional equity securities could result in additional dilution to our shareholders. The incurrence of indebtedness would result in increased debt service obligations and could result in operating and financing covenants that would restrict our operations. We cannot assure you that financing will be available in amounts or on terms acceptable to us, if at all.

Existing stockholders may experience some dilution as a result of the exercise of warrants.

In the May 2008 financing, we issued notes and, in conjunction with the notes, Class A warrants to purchase, collectively, up to 1,875,,000 shares of our common stock, subject to adjustment. In the November 2007 financing, we issued debentures and, in connection with the debentures, warrants to purchase, collectively, up to 400,000 shares of our common stock. Any issuances of shares upon any exercise of these warrants will cause dilution in the interests of our shareholders.

If we fail to maintain an effective system of internal controls, we may not be able to accurately report our financial results or prevent fraud.

We will be subject to reporting obligations under the U.S. securities laws. The SEC, as required by Section 404 of the Sarbanes-Oxley Act of 2002, adopted rules requiring every public company to include a management report on such company's internal controls over financial reporting in its annual report, which contains management's assessment of the effectiveness of our internal controls over financial reporting. In addition, an independent registered public accounting firm must attest to and report on management's assessment of the effectiveness of our internal controls

over financial reporting. Our management may conclude that our internal controls over our financial reporting are not effective. Moreover, even if our management concludes that our internal controls over financial reporting are effective, our independent registered public accounting firm may still decline to attest to our management's assessment or may issue a report that is qualified if it is not satisfied with our controls or the level at which our controls are documented, designed, operated or reviewed, or if it interprets the relevant requirements differently from us. Our reporting obligations as a public company will place a significant strain on our management, operational and financial resources and systems for the foreseeable future. Effective internal controls, particularly those related to revenue recognition, are necessary for us to produce reliable financial reports and are important to help prevent fraud. As a result, our failure to achieve and maintain effective internal controls over financial reporting could result in the loss of investor confidence in the reliability of our financial statements, which in turn could harm our business and negatively impact the trading price of our stock. Furthermore, we anticipate that we will incur considerable costs and use significant management time and other resources in an effort to comply with Section 404 and other requirements of the Sarbanes-Oxley Act.

We will incur increased costs as a result of being a public company.

As a public company, we will incur significant legal, accounting and other expenses that we did not incur as a private company. In addition, the Sarbanes-Oxley Act and other new rules subsequently implemented by SEC have required changes in corporate governance practices of public companies. We expect these new rules and regulations to increase our legal, accounting and financial compliance costs and to make certain corporate activities more time-consuming and costly. In addition, we will incur additional costs associated with our public company reporting requirements. We are currently evaluating and monitoring developments with respect to these new rules, and we cannot predict or estimate the amount of additional costs we may incur or the timing of such costs.

ITEM 1B. UNRESOLVED STAFF COMMENTS

Not applicable.

ITEM 2. DESCRIPTION OF PROPERTY

Our principal executive offices are located at Middle Section, Longmao Street, Area A, Laiyang Waixiangxing Industrial Park, Laiyang City, Yantai, Shandong Province, PRC 265200, where we have developed approximately 45,356 square meters of production, office, and garage space. Our total building area is 7,172 square meters and our production workshop area is more than 3,132 square meters. This property is owned by us.

On August 13, 2003, the Laiyang Development Planning Agency approved Laiyang Jiangbo's plan to construct garage and office space. On August 18, 2003, the Laiyang Industrial Park Administration certified Laiyang Jiangbo's investment of RMB 10 million ($1.3 million) in Section A of the Industrial Park to build on a 13,000 square meters lot.

In October 2007, the Laiyang Bureau of Land and Resources sold us a 50 years land use right for a 266,664 square meters lot located in Laiyang City to Laiyang Jiangbo. The Company paid approximately RMB 60.8 million ($8.9 million) for the land use right.

ITEM 3. LEGAL PROCEEDINGS

Except as discussed below, we are not a party to any pending legal proceeding, nor are we aware of any legal proceedings being contemplated against us by any governmental authority:

Elizabeth Hiromoto et al v. Telecom Communications, Inc. et al. - Case No. 2:07-cv-07858-PSG-E, United States District Court, Central District of California (Western Division - Los Angeles)

On December 3, 2007, two individuals filed a lawsuit against the Company, its former Chief Executive Officer James Wang, and certain others, alleging breach of contract relating to damages arising from the sale of Telecom Communications, Inc.("TCOM") to Arran Services Limited, in which Mr. Wang acted as the Company's President and Chairman to provide consulting services to TCOM and certain misrepresentations made on behalf of and in conjunction with TCOM's majority shareholder . On July 2, 2008, the Company and the plaintiffs settled the lawsuit with prejudice and claims and plaintiffs have agreed to file a Request for Dismissal with Prejudice of the lawsuit.

Fernando Praca, Plaintiff v.s. EXTREMA, LLC and Genesis Pharmaceuticals Enterprises, Inc.- Case No. 50 2005 CA 005317, Circuit Court of the 15th Judicial Circuit in and for Palm Beach County, Florida

Fernando Praca, former Director and former President of the Company's discontinued subsidiary, Extrema LLC, filed an action in Dade County, Florida against Extrema, LLC and the Company in June 2005 relating to damages arising

from the sale of Extrema LLC to Genesis Technology Group, Inc. Praca had filed a Motion of Temporary Injunction but had not proceeded to move this case forward. The plaintiff has decided to reinitiate the legal action in March 2008. In July 2008, the Company and Praca entered into a Settlement Agreement whereby Praca agreed to dismiss this action against the Company and to surrender to the Company for cancellation, 100,000 shares of common stock in the Company held by him and the Company agreed to provide Praca with a legal opinion of its counsel removing the restrictive legend on the 1,269,607 shares of common stock held by Praca.

Kenneth Clinton vs. Genesis Pharmaceuticals Enterprises, Inc., GNPH Holdings, Capital Growth Financial, Inc., Gary L. Wolfson and Pacific Rim Consultants, Inc. - Case No. 50 2007 CA 023923, Circuit Court of the 15th Judicial Circuit in and for Palm Beach County, Florida

On December 21, 2007, Kenneth Clinton, a former director and former President of the Company, filed a lawsuit against the Company and certain entities and persons related to our predecessor Genesis Technology Group, Inc. The complaint alleged, among other things, breach of contract against the Company for an agreement to pay the plaintiff certain shares of other public companies (collectively, the "Reverse Merger Shares") in connection with reverse merger transactions arranged by our predecessor, and breach of contract against the Company for failure to allow the plaintiff to exercise certain stock options for shares in the Company or exchange such options for new shares in the Company. The plaintiff sought relief in the form of (1) delivery of the Reverse Merger Shares, or in the alternative damages in the amount of those shares, (2) a judgment against the Company to allow the plaintiff to exchange and exercise his stock options for shares in the Company, or in the alternative damages in the amount of those shares, and (3) a declaratory judgment regarding a pledge and escrow agreement with defendant Capital Growth Financial.

In February 2008, the Company entered into a settlement agreement and general release with Mr. Clinton whereby the Company agreed to allow Mr. Clinton to exercise 1.5 million stock options issued under the Company's 2007 stock option plan for shares in the Company and released and discharged Mr. Clinton from any and all claims, demands or obligations. Mr. Clinton agreed to waive and release the Company from any and all claims, demands or obligations.

CRG Partners, Inc. and Genesis Technology Group, Inc., n/k/a Genesis Pharmaceuticals Enterprises, Inc. (ARBITRATION) - Case No. 32 145 Y 00976 07, American Arbitration Association, Southeast Case Management Center

On December 4, 2007, CRG Partners, Inc. ("CRG"), a former consultant of the Company, filed a demand for arbitration against the Company alleging breach of contract and seeking damages of approximately $10 million as compensation for consulting services rendered to the Company. The amount of damages sought by the claimant is equal to the dollar value of 29,978,900 shares of the Company's common stock (Pre 40 to 1 reverse split) on November 2, 2007 which the claimant alleges are due and owing to CRG. On December 5, 2007, we gave notice of termination of our relationship with CRG under the consulting agreement. The arbitration is scheduled to be conducted in Miami Dade County, Florida. We plan to vigorously defend our position.

China West II, LLC and Genesis Technology Group, Inc., n/k/a Genesis Pharmaceuticals Enterprises, Inc. (ARBITRATION)

In June 2008, China West II, LLC ("CW II") filed a Demand For Arbitration with the American Arbitration Association the case of CW II and Genesis Technology Group, Inc. n/k/a Genesis Pharmaceuticals Enterprises, Inc. and Joshua Tan. In that matter, CW II seeks breach of contract damages in connection with the Company's October 2007 reverse merger from the Company and Joshua Tan, jointly and severally for approximately $6.7 million estimated by CW II. As of the date of this report, the Company is unable to estimate a loss, if any, the Company may incur related expenses to this lawsuit. The Company believes CW II's demand was without merit and plans to vigorously defend its position.

ITEM 4. SUBMISSION OF MATTERS TO A VOTE OF SECURITY HOLDERS

On June 13, 2008, the holders of 52.4% of the Company's issued and outstanding common stock (216,677,951) approved an increase in the Company's number of authorized shares of the Company's Common Stock to 900,000,000. The share increase became effective on July 29, 2008.

PART II

Item 5. MARKET FOR COMMON EQUITY, RELATED SHAREHOLDER MATTERS AND ISSUER PURCHASES OF EQUITY SECURITIES

Market Information

Our common stock is not listed on any stock exchange. Our common stock is traded over-the-counter on the Over-the-Counter Electronic Bulletin Board under the symbol “GNPH”. The following table sets forth the high and low bid information for our common stock for each quarter within our last two fiscal years, as reported by the Over-the-Counter Electronic Bulletin Board. The bid prices reflect inter-dealer quotations, do not include retail markups, markdowns or commissions and do not necessarily reflect actual transactions.

	LOW	HIGH
2008		
Quarter ended June 30, 2008	$ 7.50	$ 14.40
Quarter ended March 31, 2008	$ 7.04	$ 14.72
Quarter ended December 31, 2007	$ 8.80	$ 14.40
Quarter ended September 30, 2007	$ 3.40	$ 6.00
2007		
Quarter ended June 30, 2007	$ 4.04	$ 7.40
Quarter ended March 31, 2007	$ 4.80	$ 7.40
Quarter ended December 31, 2006	$ 3.20	$ 7.16
Quarter ended September 30, 2006	$ 3.60	$ 7.40

As of September 26, 2008, the closing sales price for shares of our common stock was $8.70 per share on the Over-The-Counter Bulletin Board.

Holders

As of September 25, 2008, there were approximately 955 shareholders of record of our common stock based upon the shareholders' listing provided by our transfer agent. Our transfer agent is Computershare Trust Company, 350 Indiana St., #800, Golden, Colorado 80401, and its telephone number is (303) 262-0600.

Dividend Policy

We have not paid cash dividends on our common stock since the Company became public through reverse merger. We intend to keep future earnings to finance the expansion of our business, and we do not anticipate that any cash dividends will be paid in the foreseeable future. We rely on dividends from Laiyang Jiangbo for our funds and PRC regulations may limit the amount of funds distributed to us from Laiyang Jiangbo, which will affect our ability to declare any dividends. See "Risk Factors - Risks Related to Doing Business in the PRC – Laiyang Jiangbo and GJBT are subject to restrictions on paying dividends and making other payments to us" and " Governmental control of currency conversion may affect the value of your investment."

Our future payment of dividends will depend on our earnings, capital requirements, expansion plans, financial condition and relevant factors that our board of directors may deem relevant. Our retained earnings limits our ability to pay dividends.

Recent Sales of Unregistered Securities

The following private placements of the Company's securities were made in reliance upon the exemption from registration under Section 4(2) of the Securities Act of 1933, as amended, and/or, Rule 506 of Regulation D promulgated under the Securities Act. The Company did not use underwriters in any of the following private placements.

In July 2008, the Company issued 2,500 shares of restricted common stock to its officers and directors for services rendered. The Company valued these common stock at the fair market valueon the date of grant of $8 per share or, $20,000, in total base on the trading price of common stock. Accordingly, the Company recorded stock-based compensation expense of $20,000.

In June 2008, the Company issued 5,875 shares of restricted common stock to its officers and directors for services rendered. The Company valued these common stock at the fair market on the date of grant of $8 per share, or $47,000, in total base on the trading price of common stock. Accordingly, the Company recorded stock-based compensation expense of $47,000. In June 2008, we issued 235,000 shares of restricted common stock

In February 2008, in conjunction with a settlement between the Company and the Company's former officer, Mr. Kenneth Clinton, Mr. Clinton exercised 1,500,000 options and the remaining 941,406 options held by the former officer were cancelled. The Company received $157,500 in cash and the proceed was used for working capital purposes.

Issuer Purchases of Equity Securities.

None.

ITEM 6. SELECTED FINANCIAL DATA

As a smaller reporting company, we are not required to provide the information called for by Item 6 of Form 10-K.

ITEM 7. MANAGEMENT'S DISCUSSION AND ANALYSIS OR PLAN OF OPERATION

The following analysis of our consolidated financial condition and results of operations for the years ended June 30, 2008, 2007 and 2006, should be read in conjunction with our audited consolidated financial statements, including footnotes, and other information presented elsewhere in this annual report on Form 10-K. This discussion contains forward-looking statements that involve significant risks and uncertainties. As a result of many factors, such as those set forth under “Forward Looking Statements” and “Item 1A. Risk Factors” and elsewhere in this Form 10-K, our actual results may differ materially from those anticipated in these forward-looking statements. When used in this section, "fiscal 2008" means our fiscal year ended June 30, 2008 and "fiscal 2007" means our fiscal year ended June 30, 2007.

OVERVIEW

We were originally incorporated on August 15, 2001 in the State of Florida under the name Genesis Technology Group, Inc. On October 12, 2001, we consummated a merger with NewAgeCities.com, an Idaho public corporation originally formed in 1969. We were the surviving entity after the merger with the Idaho public corporation.

On October 1, 2007, we completed a share exchange transaction by and among us, Karmoya International Ltd., a British Virgin Islands company (“Karmoya”), and Karmoya’s shareholders. As a result of the share exchange transaction, Karmoya, a company which was established as a “special purpose vehicle” for the foreign capital raising activities of its Chinese subsidiaries, became our wholly owned subsidiary and our new operating business. Karmoya was incorporated under the laws of the British Virgin Islands on July 17, 2007 and owns 100% of the capital stock of Union Well International Limited, a Cayman Islands company (“Union Well”). Karmoya conducts its business operations through Union Well’s wholly owned subsidiary, Genesis Jiangbo (Laiyang) Biotech Technology Co., Ltd. (“GJBT”). GJBT was incorporated under the laws of the People’s Republic of China ("PRC") on September 16, 2007 and registered as a wholly foreign owned enterprise (WOFE) on September 19, 2007. GJBT has entered into consulting service agreements and equity-related agreements with Laiyang Jiangbo Pharmaceutical Co., Ltd. (“Laiyang Jiangbo”), a PRC limited liability company incorporated on August 18, 2003.

As a result of the share exchange transaction, our primary operations consist of the business and operations of Karmoya and its subsidiaries, which are conducted by Laiyang Jiangbo in the PRC. Laiyang Jiangbo produces and sells western pharmaceutical products in China and focuses on developing innovative medicines to address various medical needs for patients worldwide.

FINANCIAL PERFORMANCE HIGHLIGHTS:

Net Revenues

	2008	2007	2006
Net Revenues (in '000)	$ 99,547	$ 76,194	$ 49,156

Item 5. Other Information

None

Item 6. Exhibits

Exhibit Exhibit

31.1	Certification of the Chief Executive Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
31.2	Certification of the Chief Financial Officer pursuant to Section 302 of the Sarbanes-Oxley Act of 2002.
32.1	Certification of the Principal Executive Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.
32.2	Certification of the Principal Financial Officer pursuant to Section 906 of the Sarbanes-Oxley Act of 2002.

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned thereunto duly authorized.

Sector 10, Inc.

February 14, 2012 Date	By: /s/ Pericles DeAvila Pericles DeAvila, President
February 14, 2012 Date	By: /s/ Laurence A. Madison Laurence A. Madison Chief Financial Officer

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