MusclePharm Corp
Form 10-K/A
July 02, 2012

UNITED STATES

SECURITIES AND EXCHANGE COMMISSION

Washington D.C. 20549

FORM 10-K/A (Amendment No. 1)

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

For the fiscal year ended: **December 31, 2011**

¨ **TRANSITION REPORT UNDER SECTION 13 OR 15(D) OF THE SECURITIES EXCHANGE ACT OF 1934**

Commission file number: **000-53166**

MusclePharm Corporation

(Exact name of registrant as specified in its charter)

Nevada	**77-0664193**
(State or Other Jurisdiction of Incorporation or Organization)	(I.R.S. Employer Identification No.)

4721 Ironton Street, Building A

Denver, CO 80239
(Address of principal executive offices)

(303) 396-6100
(Registrant's telephone number, including area code)

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

Securities registered under Section 12(g) of the Exchange Act: **Common Stock, par value $0.001 per share**

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

Indicate by check mark if the registrant is not required to file reports pursuant to Section 13 or Section 15(d) of the Act. Yes ¨ No 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 reports) and (2) has been subject to such filing requirements for the past 90 days. Yes x No ¨

Indicate by checkmark whether the registrant has submitted electronically and posted on its corporate Website, if any, every Interactive Data File required to be submitted and posted pursuant to Rule 405 of Regulation S-T (§232.405 of this chapter) during the preceding 12 months (or for such shorter period that the registrant was required to submit and post such files). Yes 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 the definition of "large accelerated filer," "accelerated filer" and "smaller reporting company" in Rule 12b-2 of the Exchange Act:

Large accelerated filer	¨	Non-accelerated filer	¨
Accelerated filer	¨	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 registrant's voting and non-voting common equity held by non-affiliates (as defined by Rule 12b-2 of the Exchange Act) computed by reference to the average bid and asked price of such common equity on June 30, 2011, was $5,637,180. As of June 29, 2012, the registrant has one class of common equity, and the number of shares outstanding of such common equity was 1,399,074,207.

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Explanatory Note

On May 14, 2012, MusclePharm Corporation's (the "Company") independent registered public accounting firm and the Company's board of directors (the "Board") determined, after consultation with Company management that the

Company's audited financial statements for the period ended December 31, 2011 and December 31, 2010, filed in an annual report on Form 10-K with the Securities and Exchange Commission (the "Commission") on April 16, 2012 and April 1, 2011, respectively, contained certain material misstatements. Our financial statements contained in this amended annual report on Form 10-K/A restate a previous error in accounting for the Company's calculation of net sales and presentation of general and administrative expenses. The Company has determined that advertising related credits that were granted to customers fell within the guidance of ASC No. 605-50-55 ("Revenue Recognition" - Customer Payments and Incentives - Implementation Guidance and Illustrations).

The foregoing guidance indicates that, absent evidence of benefit to the vendor, appropriate U.S. GAAP treatment requires netting these types of payments against revenues and not expensing as advertising expense. The Company also noted other credits and discounts that, upon further review, had been previously classified as advertising expense as a component of general and administrative expense that require a reallocation of presentation as amounts to be netted against gross revenues. Additionally, for the year ended December 31, 2011, the Company reclassified certain items classified as samples originally included as an advertising expense to cost of sales. There were no such reclassifications for 2010.

The Company previously deducted certain credits and promotions as general and administrative expenses. After a thorough analysis and review as noted above, the Company has determined to net any credits and promotions directly against gross sales instead of classifying the same amounts as an advertising expense. Because this accounting change is a reclassification of expenses in the Company's Consolidated Statements of Operations, the Company's net loss will not be affected by the restatement, nor does the restatement affect the net loss amounts reported in its unaudited quarterly financial statements during 2011 or audited Consolidated Balance Sheets at December 31, 2011 and December 31, 2010 or Consolidated Statement of Equity at December 31, 2011 and December 31, 2010 or Consolidated Statements of Cash Flows for the year ended December 31, 2011 and December 31, 2010.

PART I

Item 1. Business.

General

Headquartered in Denver, Colorado, MusclePharm Corporation ("MusclePharm" or the "Company") is an expanding healthy lifestyle company that develops and manufactures a full line of scientifically approved (NSF International approved) nutritional supplements that are 100% free of any banned substances. Based on years of research, MusclePharm products are created through an advanced six-stage research protocol involving the expertise of top nutritional scientists. These products are field tested by more than 100 elite professional athletes from various sporting organizations including the National Football League, mixed martial arts, and Major League Baseball. The Company's award-winning proprietary products address all categories of an active lifestyle including, muscle building, weight loss and maintaining general fitness through a daily nutritional supplement regimen. MusclePharm is a marketing and branding company. It does not directly manufacturer or ship to end user customers. We extend and market the brand while innovating and distributing new products. Our customers are re-sellers of our products.

The Company's headquarters in Denver, CO, features has a state-of-the-art exercise and weight lifting facility, with a full size octagon UFC™ fighting cage, an indoor football field, cardio work-out equipment and a state-of-the-art on-site medical department, complete with equipment for measuring and conducting clinical studies and supporting athletes. A staff team of medical and clinical professionals is on hand to assist with training. MusclePharm products are sold in over 120 countries and available in over 10,000 U.S. retail outlets, including Wal-Mart, Dicks Sporting Goods, GNC, Vitamin Shoppe, and Vitamin World. The Company also sells its products in over 100 online stores, including bodybuilding.com, amazon.com and vitacost.com.

Business Strategy

Our primary focus at the current time is on the following:

(1) Increase our distribution and sales through domestic and international growth and market penetration;

(2) Conduct additional testing of the safety and efficacy of our products and create new products; and

(3) Create marketing and branding opportunities through endorsements, sponsorships and brand extensions to increase brand awareness.

The Sports Nutrition and High Energy Supplement Market

The sports nutrition and high energy supplement market is comprised of sports beverages, sports food and sports supplements. According to BCC Research's 2008 Global Research Report, sports beverages maintain the largest market share, with approximately $24.9 billion in annual sales in 2007. The sports food segment had approximately $1.2 billion in annual sales and the sports supplement segment saw 2007 annual sales of approximately $1.1 billion. BCC projected that the sports supplement market would reach $2.3 billion by 2013.

According to BCC Research, the United States is the largest consumer market for sports nutrition products, with annual sales reaching approximately $22 billion in 2007, and projected sales of $29 billion in 2013. Western Europe and Japan are the second and third largest consumers. The key market drivers for sports nutrition products are taste, price, variety and brand loyalty. In recent years, the consumption of sports nutrition products has shifted to mainstream consumers who have become the key drivers of growth within the industry.

Current Products

We currently offer seventeen (17) high-quality, specially-formulated, athlete-focused supplement products. These include: Assault™, Armor-V, Battle Fuel™, Bullet Proof®, Combat Powder®, MuscleGel®, Shred Matrix®, Re-con®, BCAA 3:1:2™, Glutamine™, Creatine™, Casein™, CLA Core™, ZMA Max™, Hybrid - NO™. We also offer the private lab products Recover Elite™ and Perform Elite™ under the MMA Elite™ name. These are distributed directly to Wal-Mart and Walgreens. Our products are comprised of amino acids, herbs and proteins scientifically tested and proven as safe and effective for the overall health of athletes. These nutritional supplements were created to enhance the effects of workouts, repair muscles, and nourish the body for optimal physical fitness. The following is a brief description of each of our products:

Assault™

Pre-Performance Amplifier

- Fuel power for long-lasting energy;
- Enhance focus; and
- Build lean muscle mass.

Assault™ helps fight fatigue, boost performance, build muscle, increase intensity, hydrate muscles and feed them valuable, clinically-proven nutrients like ConCrete, Beta Alanine, BCAAs and Cinnulin. Our team of sports medicine specialists worked with top professional athletes to create a safe pre-workout that increases strength, aerobic and anaerobic performance, reduces stomach fat and meets all regulations when it comes to being free of banned substances. Assault™ is specifically designed for performance-boosting pre-workout power.

Battle Fuel™

Maximizes Workout Performance with No Side Effects

- Increases aggression and focus;
- Boosts testosterone and feeds anabolism; and
- promotes cellular health and recovery.

Battle Fuel™ helps athletes increase lean mass and strength, improve endurance and energy levels, naturally detoxify and enhance aggressive mental focus. Battle Fuel™ is an herbal formula that improves testosterone levels to drive strength, power and lean muscle mass development. An intense combination of cleansing agents and natural elements reduce fatigue and improve cellular immunity.

Bullet Proof™

Advanced Nighttime Recovery System

- Promotes deep sleep to maximize repair;
- Optimizes anabolic/anti-catabolic environment; and
- Stimulates growth hormone/testosterone output.

Bullet Proof™ helps increase recovery effectiveness and hormonal up-regulation, improve lean muscle tissue growth and help relieve some forms of pain. Deep nourishing sleep is an athlete's best friend for the long-term building of strength, mass and speed. During this rest period, key ingredients like our proprietary blend of essential amino acids, Beta Alanine and zinc magnesium aspartate (ZMA) are hard at work repairing tissue and staving off muscle breakdown. Other ingredients boost your immune system and reduce swelling, preparing the body for that next hard workout.

Combat™

Feeds Muscle Up To 8 Hours

- Technologically advanced protein super-food;
- Enhances digestion of nutrients; and
- Maximizes adaptive response to hard training.

Combat™ helps you: receive 25 grams of high quality protein, fuel fat loss, support healthy body composition, nourish lean muscle and speed recovery. Combat™ is designed to help fill that gap in nutrition many athletes and super-active people experience, to ensure their bodies are growing and recovering. The staggered absorption rate of the five different protein components guarantees a complete 8-hour nutrient infusion.

Re-con®

Post-Workout Recharger

~~O~~ptimize an athlete's "anabolic window";
~~Pr~~omote post workout growth & repair; and
~~Re~~plenish vital nutrients.

Re-con ® helps athletes recover quicker and more effectively, repair muscle cells, feed the body nutrients and grow stronger with ingredients like base-change amino acids (BCAAs), essential amino acid complexes (EAAs), cellular detoxifiers, muscle-loading carbohydrates and stress hormone regulators. This maximizes an athlete's anabolic window, the post-workout phase where the body repairs and rebuilds tissue. Re-con® nourishes and promotes growth from every angle, delivering proteins and nutritious elements in their ideal forms. Recon® provides muscle reconstruction nutrition.

MuscleGel®

Delicious On-The-Go Protein and Nutrition

~~St~~ay leaner and be healthier;
~~Pr~~oteins absorbs into the athlete's body easier; and
~~N~~utritious and easy to enjoy.

MuscleGel® helps athletes receive more of the nutrients the body needs every day, shed pounds and fat and enjoy the convenience of the ready-to-eat packs. Packed full of different proteins like "building block" amino acids, MuscleGel's® patented gel format yields a fast-absorbing, highly bio-available source of next generation fitness food. For protein, carbohydrates and vitamins, MuscleGel® delivers. It works on-the-go, fills you up quickly and streams right to those parts of the athlete's body where nutrients are needed most.

SHRED Matrix®

Multi-Level Weight Loss System

- Ramps up your metabolism;
- Suppresses hunger and cravings; and
- Burns fat through all-natural herbs.

SHRED Matrix ® helps burn fat naturally, counteract mood swings and help athletes stay focused on weight loss and results. This 8-Stage Weight Loss System is for people who exercise regularly. As a total body diet, it sheds pounds, burns fat cells and attacks fat loss from every angle. Proven ingredients like Sugar Stop™ and the enzyme aid matrix also keep your appetite in check. The formula is tuned so athletes won't experience "jitters" or a crash.

Armor-V™

Advanced Multi-Vitamin Complex

- Complete source of vitamins and minerals;
- Total immune system support; and
- Added B vitamins and probiotics.

Armor-V™ helps athletes receive a full dose of important vitamins and minerals, keep vital organs like the liver clean of toxins, recover faster and keep the body's hormones balanced. This system was designed to meet the standards of professional athletes, who need a dedicated source of vitamins and minerals. Loaded with anti-oxidants and system optimizers derived from fruits and vegetables, Armor V™ brings together organic, herbal and natural ingredients into a multi-nutrient complex that benefits active bodies.

BCAA™

Rapidly Absorbed Branched Chain Amino Acid Complex

- Delivers BCAAs before and after workout;
- Minimizes muscle damage; and
- 100% pharmaceutical grade.

BCAA™ helps you: receive ideal amounts of BCAAs, Leucine, Isoleucine and Valine from this patented ratio of 3:1:2, promote muscle development and maintenance, increase lean body mass and spur weight loss. BCAAs are part of the group of essential amino acids a body needs. Our patented 3:1:2 ratio is designed to release the ideal amounts of each amino acid both before and after a workout. This prevents muscle breakdown and leads to gains in body mass without losing weight.

Creatine

Five Superior Blends of Creatine

- Promote strength, power and endurance;
- No loading; and
- 100% pharmaceutical quality.

MusclePharm MP Core Series Creatine™ increases creatine status by enhancing uptake and bioavailability and also fuels stamina, strength and lean muscle growth. Many athletes who engage in high-intensity/short duration exercises like weightlifting use creatine. The clinically-proven ingredient Cinnulin heightens absorption, which assists our five pure and diverse creatine complexes, delivering a range of benefits launched directly into muscles. MP Creatine™ increases explosive energy, ATP energy and overall power.

ZMA Max™

Anabolic Mineral Support Formula with Fenugreek®

- Increases testosterone;
- Promotes deep, healing sleep; and
- Supports healthy libido function

MusclePharm ZMA Max™ supports muscle growth and recovery, promotes deeper and more efficient sleep to maximize healing, tissue repair, anabolic hormone production and testosterone levels. It delivers the benefits of precise dosages and ZMA ingredient ratios, and adds the synergistic effects of clinically-proven Fenugreek ® to support the balance of cholesterol levels as well as increase of healthy libido function in women and men.

CLA Core™

Supports Healthy Body Composition

- Aids in weight loss;
- Increases metabolic rates; and
- Reduces body fat

CLA Core™ supports energy sources for hard-training people on low-carb diets, induces muscle gains without buildup of fatty tissue, and helps protect the joints. CLA is a high-quality linoleic acid that is naturally occurring and helps you feel healthy and energized. Many researchers believe that this fatty acid also helps reduce body fat and increase muscle mass. CLA Core is a blend of Omega-6 and medium-chain triglycerides (MCTs).

Casein™

100% Micellar Casein Protein

- Delivers 25g of protein;
- Serves as ideal nighttime protein source; and
- Doubles as digestive enzyme and probiotic blend.

Casein ™ delivers a steady, prolonged release of amino acids, works hard while body is at rest and promotes added nutrient utilization through the natural enzymes. As a slow digesting protein, Casein repairs and rebuilds muscle at night while you are asleep—feeding your muscles even hours after you went to bed. The great-tasting shake also works overtime readying you for that next day's workout.

Hybrid N.O.™

Nitric Oxide Amplifier

- Dramatically enhance muscle pumps
- Increase muscle fullness and vascularity
- Maximize vasodilation

Hybrid N.O.™ promotes muscle pumps through enhanced vasodilation, utilizes GlycoCarn™, a scientifically-researched nitric oxide booster, and encourages blood flow throughout the muscles. Hybrid N.O. promotes higher exercise tolerance by increasing levels of plasma nitrite and nitrate, which increases systemic nitric oxide bioavailability. Hybrid N.O. delivers maximum muscle pumps and increased health.

Glutamine

Rapidly Absorbed Glutamine Complex

- Increase recovery time;
- Enhance muscle growth; and
- 100% pharmaceutical grade.

MusclePharm Core Series MP Glutamine™ supplement increases whole body glutamine status by enhancing an athlete's uptake, bioavailability and digestion. Feeding the body a dedicated source of glutamine ultimately provides optimal muscle-tissue saturation through an exclusive array of three pure, yet diverse nutritional glutamine complexes that deliver a substantial range of benefits. MP Glutamine™ helps athletes rehydrate, rebuild and recover from even the toughest of workouts quicker and more efficiently.

PERFORM Elite™

Pre-Performance Amplifier

- Fuel power for long-lasting energy;
- Increase strength & endurance;
- Enhance focus & intensity; and
- Build lean muscle mass.

Perform Elite™ helps: fight fatigue, boost performance, build muscle, amp up intensity, hydrate muscles and feed muscle cells the nutrients they need. People need a pre-workout supplement because it prepares them to stay focused, energetic and fully-powered from start to finish. In Perform Elite™, ingredients like creatine and beta alanine feed muscle cells the nutrients they need to grow bigger and stronger. Without a pre-workout like Perform Elite™, muscles don't have the building blocks they need to gain size and strength.

RECOVER Elite™ (540g)

Post-Workout Recharger

- Optimize athletes' "muscle building";
- Promote post workout growth & repair;
- Replenish vital nutrients; and
- Speed recovery time.

Recover Elite™ helps athletes recover quicker and more effectively, repair muscle cells, maintain nutrient levels and grow bigger and stronger. When people work out, their muscle cells break down. In that post-workout period, it is important to feed these cells a supply of the right nutrients like the ones in Recover Elite™, so they can rebuild properly. When this is done correctly, muscles become both stronger and larger. If you don't use a post-workout supplement, these cells don't have the building blocks they need to recover and you won't increase size or strength.

Future Products

We have trademarked and registered an energy product for active lifestyles and competitive athletes, under the name Energel®. This product will be distributed primarily via internet sales, convenience stores, cycling shops, ski shops, and fitness and runner shoes retail sales stores by the end of the second quarter of 2012.

Sales & Distribution

We sell our products both domestically and internationally. With respect to domestic sales, we have three traditional distribution systems:

1) Approximately 40% of sales are through a domestic Internet site named Bodybuilding.com ("Body Building"), which is the largest online retailer of sports nutrition products in the United States. Body Building awarded MusclePharm the title of the "Breakout Brand," "Best Packaging" and "Best New Product for Assault" in 2011 and MusclePharm is now the number two best-selling brand on BB.com, and has two products in their top ten best sellers, and eight products in the top fifty selling products out of over 14,000 sku's. In addition to Body Building.com, we also sell domestically through several distributors and over 100 Internet sites;

2) We sell through traditional brick and mortar stores, and are carried in approximately 450 The Vitamin Shoppes outlets and we sell our products into over 5000 GNC stores, and we are in 400 Vitamin World retail stores;

3) We have regional salesmen that support wholesale distributors like Europa, selling in up to 15,000 smaller retail or regional stores. We also work with other large distributors who have begun to place the Company's product in small retail stores and gyms across the United States. Internationally, we are expanding rapidly into Central America, Mexico, Brazil, the Middle East, Europe, Russia, and the UK, and have Sportika Export as our international distributor that services over 120 countries.

In addition, we just recently launched a partnership with Eurpac to distribute our line of products to U.S. military bases and stores all over the world.

Marketing Strategy

Our core marketing strategy is to brand MusclePharm as the "must have" nutritional supplement line for high performance athletes. We want to be known as the athlete's company, run by athletes who create their products for other athletes both professional and otherwise. We have endorsements from over 50 UFC fighters, well-known NFL players, as well as top X-Game and fitness athletes. Athletes are considered role models and many people strive to emulate their fitness and well-being regimen. Athlete sponsorships are the most logical tactic for our business. The objective of these athletic endorsements is to build both consumer awareness and confidence and to drive consumer demand for our products.

MusclePharm in 2011 became the Official Supplement Provider and Sponsor of the Ultimate Fighting Championship ("UFC"). Our agreement includes prominent logo placement on the mat, and our branding can be seen on Fox and pay-per-view worldwide.

The fighters we sponsor feature our brand on their uniforms and we also extensively advertise at the Ultimate Fighting Championship events. In 2011, we launched a state-of-the-art website that will tap into the social networking world, further expanding our brand and consumer awareness.

The Company is also currently engaged in various in-store promotions, including point-of-purchase stands, aisle displays in our retail outlets, as well as sample demonstrations and athlete appearances in Wal-Mart, GNC, Vitamin World and Vitamin Shoppe locations.

Research and Development

Each and every product sold by MusclePharm is the end result of a long development process involving leading nutrition scientists, doctors, and top professional athletes.

Manufacturing and Product Quality

We are committed to producing and selling highly efficacious products that are trusted for their quality and safety. To date, our products have been outsourced to a third party manufacturer where the products are manufactured in full compliance with the Good Manufacturing Practice standards set by the Food & Drug Administration.

Trademarks and Patents

We regard our trademarks and other proprietary rights as valuable assets and believe that protecting our key trademarks is crucial to our business strategy of building strong brand name recognition. These trademarks are crucial elements of our business, and have significant value in the marketing of our products.

Our policy is to pursue registrations for all of the trademarks associated with our products. Federally registered trademarks have a perpetual life, provided that they are maintained and renewed on a timely basis and used correctly as trademarks, subject to the rights of third parties to attempt to cancel a trademark if priority is claimed or there is confusion of usage. We rely on common law trademark rights to protect our unregistered trademarks. Common law trademark rights generally are limited to the geographic area in which the trademark is actually used, while a United States federal registration of a trademark enables the registrant to stop the unauthorized use of the trademark by any third party anywhere in the United States. Furthermore, the protection available, if any, in foreign jurisdictions may not be as extensive as the protection available to us in the United States.

Although we seek to ensure that we do not infringe on the intellectual property rights of others, there can be no assurance that third parties will not assert intellectual property infringement claims against us.

Competition

The sports nutrition business is highly competitive. Competition is based primarily on quality and assortment of products, marketing support, and availability of new products. Currently, our main competitors are three private companies: Optimum Nutrition, Inc. (“Optimum”), Iovate Health Sciences, Inc. (“IHS”), and Bio-Engineered Supplements and Nutrition, Inc. (“BSN”). Optimum is a wholly owned subsidiary of Glanbia Nutritionals, Inc., an international nutritional ingredients group. Optimum owns and operates two brands of nutritional supplements (Optimum Nutrition and American Body Building), providing a line of products across multiple categories. IHS is a nutritional supplement company that delivers a range of products to the nutritional marketplace. Headquartered in Oakville, Ontario, Canada, IHS’s line of products can be found in major retail stores and include such brands as Hydroxy-Cut™, Cell-Tech™, Six Star Nutrition™. BSN is also a sports nutrition leader whose top products include No-Explode™ and Syntha Six Protein™.

MusclePharm intends to compete by aggressively marketing our brand, emphasizing our relationships with professional athletes, and maximizing our relationships with those athletes, retail outlets and industry publications that align with our vision. We also tout the strength of the science behind MusclePharm products, as this is a key point of difference.

Regulatory Matters

The manufacture, packaging, labeling, advertising, promotion, distribution and sale of our products are subject to regulation by numerous governmental agencies. Our products are subject to regulation by, among other regulatory entities, the Consumer Product Safety Commission (CPSC), the U.S. Department of Agriculture (USDA), the Environmental Protection Agency (EPA) and the U.S. Food and Drug Administration (FDA). Advertising and other forms of promotion and methods of marketing are subject to regulation primarily by the U.S. Federal Trade Commission (FTC), which regulates these activities under the Federal Trade Commission Act (FTCA). The manufacture, labeling and advertising of our products are also regulated by various state and local agencies as well as those of each foreign country to which we distribute our products.

The Dietary Supplement Health and Education Act of 1994 (DSHEA) revised the provisions of the Federal Food, Drug, and Cosmetic Act (FFDC Act) concerning the regulation of dietary supplements. All of the products we market are regulated as dietary supplements under the FFDC Act.

Under the current provisions of the FFDC Act, there are four categories of claims that pertain to the regulation of dietary supplements. Health claims are claims that describe the relationship between a nutrient or dietary ingredient and a disease or health related condition and can be made on the labeling of dietary supplements if supported by significant scientific agreement and authorized by the FDA in advance via notice and comment rulemaking. Nutrient content claims describe the nutritional value of the product and may be made if defined by the FDA through notice and comment rulemaking and if one serving of the product meets the definition. Statements of nutritional support or product performance, which are permitted on labeling of dietary supplements without FDA pre-approval, are defined to include statements that: (i) claim a benefit related to a classical nutrient deficiency disease and disclose the prevalence of such disease in the United States; (ii) describe the role of a nutrient or dietary ingredient intended to affect the structure or function in humans; (iii) characterize the documented mechanism by which a dietary ingredient acts to maintain such structure or function; or (iv) describe general well-being from consumption of a nutrient or dietary ingredient. In order to make a nutritional support claim, the marketer must possess adequate substantiation to demonstrate that the claim is not false or misleading and if the claim is for a dietary ingredient that does not provide traditional nutritional value, prominent disclosure of the lack of FDA review of the relevant statement and notification to the FDA of the claim is required. Drug claims are representations that a product is intended to diagnose, mitigate, treat, cure or prevent a disease. Drug claims are prohibited from use in the labeling of dietary supplements.

Claims made for our dietary supplement products may include statements of nutritional support and health and nutrient content claims when authorized by the FDA or otherwise allowed by law. The FDA’s interpretation of what constitutes an acceptable statement of nutritional support may change in the future, thereby requiring that we revise our labeling. In addition, a dietary supplement that contains a new dietary ingredient (i.e., one not on the market before October 15, 1994) must have a history of use or other evidence of safety establishing that it is reasonably expected to be safe. The manufacturer must notify the FDA at least 75 days before marketing products containing new dietary ingredients and provide the FDA the information upon which the manufacturer based its conclusion that the product has a reasonable expectation of safety. There is no assurance that the FDA will accept the evidence of safety

for any new dietary ingredients that we may wish to market, and the FDA's refusal to accept that evidence could prevent the marketing of the new dietary ingredients and dietary supplements containing a new dietary ingredient.

Our dietary supplements must comply with the Dietary Supplement and Nonprescription Drug Consumer Protection Act, which became effective on December 22, 2007. This Act amends the FFDC Act to mandate the reporting of serious adverse events received by us to the FDA.

The FDA has also announced its intention to promulgate new GMPs specific to dietary supplements, to fully enforce DSHEA and monitor compliance with the Bioterrorism Act of 2002.

Our failure to comply with applicable FDA regulatory requirements could result in, among other things, injunctions, product withdrawals, recalls, product seizures, fines and criminal prosecutions. We intend to comply with the new GMPs once they are adopted. The new GMPs, predicted to be finalized shortly, would be more detailed and stringent than the GMPs that currently apply to dietary supplements and may, among other things, require dietary supplements to be prepared, packaged, produced and held in compliance with regulations similar to the GMP regulations for drugs. There can be no assurance that, if the FDA adopts GMP regulations for dietary supplements, we will be able to comply with the new regulations without incurring a substantial expense.

As a result of our efforts to comply with applicable statutes and regulations in the United States and elsewhere, we have from time to time reformulated, eliminated or relabeled certain of our products and revised certain advertising claims. We cannot predict the nature of any future laws, regulations, interpretations or applications, nor can we determine what effect additional governmental regulations or administrative orders, when and if promulgated, would have on our business in the future. They could, however, require the reformulation of certain products to meet new standards, the recall or discontinuance of certain products not capable of reformulation, additional record keeping, expanded documentation of the properties of certain products, expanded or different labeling, and/or scientific substantiation. Any or all of such requirements could have a material adverse effect on our business, financial condition and results of operations.

Our advertising of dietary supplement products is subject to regulation by the FTC under the FTCA. Section 5 of the FTCA prohibits unfair methods of competition and unfair or deceptive acts or practices in or affecting commerce. Section 12 of the FTCA provides that the dissemination or the causing to be disseminated of any false advertisement pertaining to drugs or foods, which would include dietary supplements, is an unfair or deceptive act or practice. Under the FTC's Substantiation Doctrine, an advertiser is required to have a "reasonable basis" for all objective product claims before the claims are made. Failure to adequately substantiate claims may be considered either deceptive or unfair practices. Pursuant to this FTC requirement, we are required to have adequate substantiation for all material advertising claims made for our products.

On November 18, 1998, the FTC issued "Dietary Supplements: An Advertising Guide for Industry." This guide provides marketers of dietary supplements with guidelines on applying FTC law to dietary supplement advertising. It includes examples of the principles that should be used when interpreting and substantiating dietary supplement advertising. Although the guide provides additional explanation, it does not substantively change the FTC's existing policy that all supplement marketers have an obligation to ensure that claims are presented truthfully and to verify the adequacy of the support behind such claims. Our outside counsel reviews our advertising claims for compliance with FTC requirements.

The FTC has a variety of processes and remedies available to it for enforcement, both administratively and judicially, including compulsory process, cease and desist orders and injunctions. FTC enforcement can result in orders requiring, among other things, limits on advertising, corrective advertising, consumer redress, divestiture of assets, rescission of contracts and such other relief as may be deemed necessary. A violation of such orders could have a material adverse effect on our business, financial condition and results of operations.

Advertising and labeling for dietary supplements and conventional foods are also regulated by state, county and other local governmental authorities. Some states also permit these laws to be enforced by private attorney generals. These private attorney generals may seek relief for consumers, seek class action certifications, seek class-wide damages, seek class-wide refunds and product recalls of products sold by us. There can be no assurance that state and local authorities will not commence regulatory action, which could restrict the permissible scope of our product advertising claims, or products that can be sold in the future.

Governmental regulations in foreign countries where we plan to or expand sales may prevent or delay entry into the market or prevent or delay the introduction, or require the reformulation, of certain of our products. Compliance with such foreign governmental regulations is generally the responsibility of our distributors for those countries. These distributors are independent contractors over whom we have limited control.

Number of Total Employees and Number of Full Time Employees

We believe that our success will depend greatly on our ability to identify, attract, and retain capable employees. As of April 13, 2012, we had 24 full time employees. Our employees are not represented by any collective bargaining unit, and we believe our relations with our employees are good. We have recently completed staffing for the in-house medical and physiology center on-site in our work-out, fight training and training facilities.

Item 1A. Risk Factors.

Risks Related to Our Business and Industry

OUR INDEPENDENT AUDITORS HAVE EXPRESSED SUBSTANTIAL DOUBT ABOUT OUR ABILITY TO CONTINUE AS A GOING CONCERN, WHICH MAY HINDER OUR ABILITY TO CONTINUE AS A GOING CONCERN AND OUR ABILITY TO OBTAIN FUTURE FINANCING.

As reflected in the accompanying restated financial statements, the Company had a net loss of $23,280,950 and net cash used in operations of $5,801,761 for the year ended December 31, 2011, and a working capital deficit and stockholders' deficit of $13,693,267 and $12,971,212, respectively, at December 31, 2011. These factors raise substantial doubt about the Company's ability to continue as a going concern.

The ability of the Company to continue its operations is dependent on management's plans, which include the raising of capital through debt and/or equity markets with some additional funding from other traditional financing sources, including term notes, until such time that funds provided by operations are sufficient to fund working capital requirements. The Company may need to incur liabilities with certain related parties to sustain the Company's existence.

The Company will require additional funding to finance the growth of its current and expected future operations as well as to achieve its strategic objectives. The Company believes its current available cash along with anticipated revenues may be insufficient to meet its cash needs for the near future. There can be no assurance that financing will be available in amounts or terms acceptable to the Company, if at all.

In response to these problems, management has taken the following actions:

- seeking additional third party debt and/or equity financing;
- execute a plan to recapitalize the company;
- continue with the implementation of the business plan;

- generate new sales from international customers; and

- allocate sufficient resources to continue with advertising and marketing efforts.

In their report dated April 13, 2012, our independent auditors stated that our financial statements for the period ended December 31, 2011, were prepared assuming that we would continue as a going concern, which contemplates the realization of assets and the satisfaction of liabilities in the normal course of business. These financial statements do not include any adjustments relating to the recovery of the recorded assets or the classification of the liabilities that might be necessary should the Company be unable to continue as a going concern.

WE WILL NEED TO RAISE ADDITIONAL CAPITAL TO CARRY OUT OUR BUSINESS PLAN.

We will need to raise additional capital to fund the growth of our business. There is no guarantee that we will be able to access additional capital at rates and on terms which are attractive to us, if at all. Without the additional funding needed to fund our growth we may not be able to grow as planned.

OUR FAILURE TO APPROPRIATELY RESPOND TO COMPETITIVE CHALLENGES, CHANGING CONSUMER PREFERENCES AND DEMAND FOR NEW PRODUCTS COULD SIGNIFICANTLY HARM OUR CUSTOMER RELATIONSHIPS AND PRODUCT SALES.

The nutritional sports supplement industry is characterized by intense competition for product offerings and rapid and frequent changes in consumer demand. Our failure to accurately predict product trends could negatively impact our products and inventory levels and cause our revenues to decline.

Our success with any particular product offering (whether new or existing) depends upon a number of factors, including our ability to:

- deliver products in a timely manner in sufficient volumes;
- accurately anticipate customer needs and forecast accurately to the manufacturer in a rapidly expanding business;
- differentiate our product offerings from those of our competitors;
- competitively price our products; and
- develop and/or acquire new products.

Products often have to be promoted heavily in stores or in the media to obtain visibility and consumer acceptance. Acquiring distribution for products is difficult and often expensive due to slotting and other promotional charges mandated by retailers. Products can take substantial periods of time to develop consumer awareness, consumer acceptance and sales volume. Accordingly, some products fail to gain or maintain sufficient sales volume and as a result have to be discontinued. In a highly completive marketplace it may be difficult to have retailers open stock-keeping units (sku's) for new products.

OUR INDUSTRY IS HIGHLY COMPETITIVE, AND OUR FAILURE TO COMPETE EFFECTIVELY COULD ADVERSELY AFFECT OUR MARKET SHARE, FINANCIAL CONDITION AND FUTURE GROWTH.

The sports supplement industry is highly competitive with respect to:

- price;
- shelf space and store placement;
- brand and product recognition;
- new product introductions; and
- raw materials.

Several of our competitors are larger, more established and possess greater financial, personnel, distribution and other resources. We face competition in the health food channel from a limited number of large nationally known manufacturers, private label brands and many smaller manufacturers of dietary supplements.

WE RELY ON A LIMITED NUMBER OF CUSTOMERS FOR A SUBSTANTIAL PORTION OF OUR SALES, AND THE LOSS OF OR MATERIAL REDUCTION IN PURCHASE VOLUME BY ANY OF THESE CUSTOMERS WOULD ADVERSELY AFFECT OUR SALES AND OPERATING RESULTS.

In 2011, two customers accounted for approximately 55% of net sales. Our largest customer in 2011 represented 41% of our sales. In 2010, three customers accounted for approximately 67% of our sales. The largest customer in 2010 accounted for 45% of our sales. The loss of any of our major customers, a significant reduction in purchases by any major customer, or, any serious financial difficulty of a major customer, could have a material adverse effect on our sales and results of operations.

Customer	2011		2010	
A	41	%	45	%
B	14	%	7	%
C	-	%	15	%

ADVERSE PUBLICITY OR CONSUMER PERCEPTION OF OUR PRODUCTS AND ANY SIMILAR PRODUCTS DISTRIBUTED BY OTHERS COULD HARM OUR REPUTATION AND ADVERSELY AFFECT OUR SALES AND REVENUES.

We are highly dependent upon positive consumer perceptions of the safety and quality of our products as well as similar products distributed by other sports nutrition supplement companies. Consumer perception of sports nutrition supplements and our products in particular can be substantially influenced by scientific research or findings, national media attention and other publicity about product use. Adverse publicity from such sources regarding the safety, quality or efficacy of dietary supplements and our products could harm our reputation and results of operations. The mere publication of reports asserting that such products may be harmful or questioning their efficacy could have a material adverse effect on our business, financial condition and results of operations, regardless of whether such reports are scientifically supported or whether the claimed harmful effects would be present at the dosages recommended for such products.

IF WE ARE UNABLE TO RETAIN KEY PERSONNEL, OUR ABILITY TO MANAGE OUR BUSINESS EFFECTIVELY AND CONTINUE OUR GROWTH COULD BE NEGATIVELY IMPACTED.

Key management employees include Brad J. Pyatt, Cory Gregory, Jeremy DeLuca, Larry Meer, John H. Bluher, and certain other individuals. These key management employees are primarily responsible for our day-to-day operations, and we believe our success depends in large part on our ability to retain them and to continue to attract additional qualified individuals to our management team. Currently, we have executed employment agreements with our key management employees. We anticipate having all key executives under new performance based contracts by the end of the second quarter of 2012. The loss or limitation of the services of any of our key management employees or the inability to attract additional qualified personnel could have a material adverse effect on our business and results of operations. We may obtain key man insurance on one or more key executives.

OUR OPERATING RESULTS MAY FLUCTUATE, WHICH MAKES OUR RESULTS DIFFICULT TO PREDICT AND COULD CAUSE OUR RESULTS TO FALL SHORT OF EXPECTATIONS.

Our operating results may fluctuate as a result of a number of factors, many outside of our control. As a result, comparing our operating results on a period-to-period basis may not be meaningful, and you should not rely on our past results as an indication of our future performance. Our quarterly, year-to-date, and annual expenses as a percentage of our revenues may differ significantly from our historical or projected rates. Our operating results in future quarters may fall below expectations. Each of the following factors may affect our operating results:

- our ability to deliver products in a timely manner in sufficient volumes;

- our ability to recognize product trends;
- our loss of one or more significant customers;
- the introduction of successful new products by our competitors; and
- adverse media reports on the use or efficacy of sports nutrition supplements.

Because our business is changing and evolving, our historical operating results may not be useful to you in predicting our future operating results.

THE EFFECTS OF THE RECENT GLOBAL ECONOMIC CRISIS MAY IMPACT OUR BUSINESS, OPERATING RESULTS, OR FINANCIAL CONDITION.

The recent global economic crisis has caused disruptions and extreme volatility in global financial markets and increased rates of default and bankruptcy, and has impacted levels of consumer spending. These macroeconomic developments could negatively affect our business, operating results, or financial condition. For example, if consumer spending continues to decrease, this may result in lower sales.

OUR BUSINESS AND OPERATIONS ARE EXPERIENCING RAPID GROWTH. IF WE FAIL TO EFFECTIVELY MANAGE OUR GROWTH, OUR BUSINESS AND OPERATING RESULTS COULD BE HARMED.

We have experienced and expect to continue to experience rapid growth in our operations, which has placed, and will continue to place, significant demands on our management, operational and financial infrastructure. If we do not effectively manage our growth, we may fail to timely deliver products to our customers in sufficient volume or the quality of our products could suffer, which could negatively affect our operating results. To effectively manage this growth, we will need to hire additional persons, particularly in sales and marketing, and we will need to continue to improve our operational, financial and management controls and our reporting systems and procedures. These additional employees, systems enhancements and improvements will require significant capital expenditures and management resources. Failure to implement these improvements could hurt our ability to manage our growth and our financial position.

WE MAY BE EXPOSED TO MATERIAL PRODUCT LIABILITY CLAIMS, WHICH COULD INCREASE OUR COSTS AND ADVERSELY AFFECT OUR REPUTATION AND BUSINESS.

As a marketer and distributor of products designed for human consumption, we are subject to product liability claims if the use of our products is alleged to have resulted in injury. Our products consist of vitamins, minerals, herbs and other ingredients that are classified as dietary supplements and in most cases are not subject to pre-market regulatory approval in the United States or internationally. Previously unknown adverse reactions resulting from human consumption of these ingredients could occur.

We have not had any product liability claims filed against us, but in the future we may be, subject to various product liability claims, including among others that our products had inadequate instructions for use, or inadequate warnings concerning possible side effects and interactions with other substances. The cost of defense can be substantially higher than the cost of settlement even when claims are without merit. The high cost to defend or settle product liability claims could have a material adverse effect on our business and operating results.

OUR INSURANCE COVERAGE OR THIRD PARTY INDEMNIFICATION RIGHTS MAY NOT BE SUFFICIENT TO COVER OUR LEGAL CLAIMS OR OTHER LOSSES THAT WE MAY INCUR IN THE FUTURE.

We maintain insurance, including property, general and product liability, and workers' compensation to protect ourselves against potential loss exposures. In the future, insurance coverage may not be available at adequate levels or on adequate terms to cover potential losses, including on terms that meet our customer's requirements. If insurance

coverage is inadequate or unavailable, we may face claims that exceed coverage limits or that are not covered, which could increase our costs and adversely affect our operating results.

OUR INTELLECTUAL PROPERTY RIGHTS ARE VALUABLE, AND ANY INABILITY TO PROTECT THEM COULD REDUCE THE VALUE OF OUR PRODUCTS AND BRAND.

We have invested significant resources to protect our brands and intellectual property rights. However, we may be unable or unwilling to strictly enforce our intellectual property rights, including our trademarks, from infringement. Our failure to enforce our intellectual property rights could diminish the value of our brands and product offerings and harm our business and future growth prospects.

WE MAY BE SUBJECT TO INTELLECTUAL PROPERTY RIGHTS CLAIMS, WHICH ARE COSTLY TO DEFEND, COULD REQUIRE US TO PAY DAMAGES AND COULD LIMIT OUR ABILITY TO SELL SOME OF OUR PRODUCTS.

As a marketer and distributor of products designed for human consumption, we are subject to product liability claims if the use of our products is alleged to have resulted in injury. Our products consist of vitamins, minerals, herbs and other ingredients that are classified as dietary supplements and in most cases are not subject to pre-market regulatory approval in the United States or internationally. Previously unknown adverse reactions resulting from human consumption of these ingredients could occur (See Item 3. Legal Proceedings).

WE RELY ON HIGHLY SKILLED PERSONNEL AND, IF WE ARE UNABLE TO RETAIN OR MOTIVATE KEY PERSONNEL, HIRE QUALIFIED PERSONNEL, WE MAY NOT BE ABLE TO GROW EFFECTIVELY.

Our performance largely depends on the talents and efforts of highly skilled individuals. Our future success depends on our continuing ability to identify, hire, develop, motivate and retain highly skilled personnel for all areas of our organization, particularly sales and marketing. Competition in our industry for qualified employees is intense. In addition, our compensation arrangements, such as our equity award programs, may not always be successful in attracting new employees and retaining and motivating our existing employees. Our continued ability to compete effectively depends on our ability to attract new employees and to retain and motivate our existing employees.

AN INCREASE IN PRODUCT RETURNS COULD NEGATIVELY IMPACT THE COMPANY'S OPERATING RESULTS AND PROFITABILITY.

The Company permits the return of damaged or defective products and accepts limited amounts of product returns in certain instances. Accordingly, the Company provides allowances for the estimated amounts of these returns at the time of revenue recognition based on historical experience. While such returns have historically been nominal and within management’s expectations and the provisions established, future return rates may differ from those experienced in the past. Any significant increase in damaged or defective products or expected returns could have a material adverse effect on the Company’s operating results for the period or periods in which such returns materialize.

A SHORTAGE IN THE SUPPLY OF KEY RAW MATERIALS COULD INCREASE OUR COSTS OR ADVERSELY AFFECT OUR SALES AND REVENUES.

We obtain all of our raw materials from third-party suppliers with whom we do not have significant long-term supply contracts. Since all of the ingredients in our products are commonly used, we have not experienced any shortages or delays in obtaining raw materials. If things changed, shortages could result in materially higher raw material prices or adversely affect our ability to manufacture a product. Price increases from a supplier would directly affect our profitability if we are not able to pass price increases on to customers. Our inability to obtain adequate supplies of raw materials in a timely manner or a material increase in the price of our raw materials could have a material adverse effect on our business, financial condition and results of operations.

BECAUSE WE ARE SUBJECT TO NUMEROUS LAWS AND REGULATIONS, AND WE MAY BECOME INVOLVED IN LITIGATION FROM TIME TO TIME, WE COULD INCUR SUBSTANTIAL JUDGMENTS, FINES, LEGAL FEES AND OTHER COSTS.

Our industry is highly regulated. The manufacture, labeling and advertising for our products are regulated by various federal, state and local agencies as well as those of each foreign country to which we distribute. These governmental authorities may commence regulatory or legal proceedings, which could restrict the permissible scope of our product claims or the ability to manufacture and sell our products in the future. The FDA regulates our products to ensure that the products are not adulterated or misbranded. Failure to comply with FDA requirements may result in, among other things, injunctions, product withdrawals, recalls, product seizures, fines and criminal prosecutions. Our advertising is subject to regulation by the FTC under the FTCA. In recent years the FTC has initiated numerous investigations of dietary supplement and weight loss products and companies. Additionally, some states also permit advertising and labeling laws to be enforced by private attorney generals, who may seek relief for consumers, seek class action certifications, seek class wide damages and product recalls of products sold by us. Any of these types of adverse actions against us by governmental authorities or private litigants could have a material adverse effect on our business, financial condition and results of operations.

Other Risks Factors

WE MAY, IN THE FUTURE, ISSUE ADDITIONAL COMMON SHARES, WHICH WOULD REDUCE INVESTORS' PERCENT OF OWNERSHIP AND MAY DILUTE OUR SHARE VALUE.

Our Articles of Incorporation authorize the issuance of 2,500,000,000 shares of common stock, 5,000,000 shares of Series A Convertible Preferred Stock, 51 shares of Series B Preferred Stock, 500 shares of Series C Convertible Preferred Stock. The Company currently has 9,999,449 shares of blank-check preferred stock authorized but undesignated. The future issuance of common stock may result in substantial dilution in the percentage of our common stock held by our then existing shareholders. We may value any common stock issued in the future on an arbitrary basis. The issuance of common stock for future services or acquisitions or other corporate actions may have the effect of diluting the value of the shares held by our investors, and might have an adverse effect on any trading market for our common stock.

OUR COMMON STOCK IS QUOTED ON THE OTCBB, WHICH MAY HAVE AN UNFAVORABLE IMPACT ON OUR STOCK PRICE AND LIQUIDITY.

Our common stock is quoted on the OTCBB. The OTCBB is a significantly more limited market than the New York Stock Exchange or NASDAQ system. The quotation of our shares on the OTCBB may result in a less liquid market available for existing and potential stockholders to trade shares of our common stock, could depress the trading price of our common stock and could have a long-term adverse impact on our ability to raise capital in the future.

OUR COMMON SHARES ARE SUBJECT TO THE "PENNY STOCK" RULES OF THE SEC AND THE TRADING MARKET IN OUR SECURITIES IS LIMITED, WHICH MAKES TRANSACTIONS IN OUR STOCK CUMBERSOME AND MAY REDUCE THE VALUE OF AN INVESTMENT IN OUR STOCK.

The Securities and Exchange Commission has adopted Rule 15g-9 which establishes the definition of a "penny stock," for the purposes relevant to us, as any equity security that has a market price of less than $5.00 per share or with an exercise price of less than $5.00 per share, subject to certain exceptions.

For any transaction involving a penny stock, unless exempt, the rules require:

(a) that a broker or dealer approve a person's account for transactions in penny stocks; and

(b) the broker or dealer receive from the investor a written agreement to the transaction, setting forth the identity and quantity of the penny stock to be purchased.

In order to approve a person's account for transactions in penny stocks, the broker or dealer must: (a) obtain financial information and investment experience objectives of the person; and (b) make a reasonable determination that the transactions in penny stocks are suitable for that person and the person has sufficient knowledge and experience in financial matters to be capable of evaluating the risks of transactions in penny stocks.

The broker or dealer must also deliver, prior to any transaction in a penny stock, a disclosure schedule prescribed by the Commission relating to the penny stock market, which, in highlight form: (a) sets forth the basis on which the broker or dealer made the suitability determination, and (b) that the broker or dealer received a signed, written agreement from the investor prior to the transaction. Generally, brokers may be less willing to execute transactions in securities subject to the "penny stock" rules. This may make it more difficult for investors to dispose of our Common shares and cause a decline in the market value of our stock.

Disclosure also has to be made about the risks of investing in penny stocks in both public offerings and in secondary trading and about the commissions payable to both the broker-dealer and the registered representative, current quotations for the securities and the rights and remedies available to an investor in cases of fraud in penny stock transactions. Finally, monthly statements have to be sent disclosing recent price information for the penny stock held in the account and information on the limited market in penny stocks.

LIABILITY OF DIRECTORS FOR BREACH OF DUTY OF CARE IS LIMITED.

According to Nevada law (NRS 78.138(7)), all Nevada corporations limit the liability of directors and officers, including acts not in good faith. Our stockholders' ability to recover damages for fiduciary breaches may be reduced by this statute. In addition, we are obligated to indemnify our directors and officers regarding stockholder suits which they successfully defend (NRS 78.7502).

BECAUSE WE DO NOT INTEND TO PAY ANY CASH DIVIDENDS ON OUR COMMON STOCK, OUR STOCKHOLDERS WILL NOT BE ABLE TO RECEIVE A RETURN ON THEIR SHARES UNLESS THEY SELL THEM.

We intend to retain any future earnings to finance the development and expansion of our business. We do not anticipate paying any cash dividends on our common stock in the foreseeable future. Unless we pay dividends, our stockholders will not be able to receive a return on their shares unless they sell them. There is no assurance that stockholders will be able to sell shares when desired.

WE WILL INCUR ONGOING COSTS AND EXPENSES FOR SEC REPORTING AND COMPLIANCE, AND WITHOUT REVENUE WE MAY NOT BE ABLE TO REMAIN IN COMPLIANCE WITH THE SEC, MAKING IT DIFFICULT FOR INVESTORS TO SELL THEIR SHARES, IF AT ALL.

To remain eligible for quotation on the OTCBB, issuers must remain current in their filings with the SEC. Market Makers are not permitted to begin quotation of a security whose issuer does not meet this filing requirement. Securities already quoted on the OTCBB that become delinquent in their required filings will be removed following a 30 or 60 day grace period if they do not make their required filing during that time. In order for us to remain in compliance we will require future revenues to cover the cost of these filings, which could comprise a substantial portion of our available cash resources. If we are unable to generate sufficient revenues to remain in compliance it may be difficult for you to resell any shares you may purchase, if at all.

WE MAY ISSUE ADDITIONAL SHARES OF PREFERRED STOCK IN THE FUTURE THAT MAY ADVERSELY IMPACT YOUR RIGHTS AS HOLDERS OF OUR COMMON STOCK.

Our articles of incorporation authorize us to issue up to issue up to 15,000,000 shares of preferred stock in various classes. To date, the Company has issued 51 shares of Series B preferred stock and 190 shares of Series C preferred stock. The Series C preferred stock is convertible into shares of the Company’s common stock. Our board of directors will have the authority to fix and determine the relative rights and preferences of preferred shares, as well as the authority to issue additional shares, without further stockholder approval. As a result, our board of directors could authorize the issuance of a series of preferred stock that would grant to holders preferred rights to our assets upon liquidation, the right to receive dividends before dividends are declared to holders of our common stock, and the right to the redemption of such preferred shares, together with a premium, prior to the redemption of the common stock. To the extent that we do issue such additional shares of preferred stock, your rights as holders of common stock could be impaired thereby, including, without limitation, dilution of your ownership interests in us. In addition, shares of preferred stock could be issued with terms calculated to delay or prevent a change in control or make removal of management more difficult, which may not be in your interest as a holder of common stock.

Item 1B. Unresolved Staff Comments.

Not applicable.

Item 2. Properties.

MusclePharm's corporate headquarters is located in Denver, Colorado. This commercial office building is 30,320 sq. ft. with 5,000 sq. ft. being used for offices and the other 25,000 sq. ft. utilized for research and development. The space includes a full performance training center, medical laboratory, and a 96 seat theatre room. The term of the lease is 65 months, expiring on December 31, 2015. We currently pay approximately $13,500 in lease payments per month.

MusclePharm is leasing a small office and distribution warehouse in Boise, Id. The lease expires in February 2013 and the Company pays approximately $3,500 per month in rent fees.

Item 3. Legal Proceedings.

From time to time we may become involved in legal proceedings which could adversely affect us. We are currently not involved in any litigation, other than litigation in the ordinary course of business, that we believe could have a materially adverse effect on our financial condition or results of operations. There is no action, suit, proceeding, inquiry or investigation before or by any court, public board, government agency, self-regulatory organization or body pending or, to the knowledge of the executive officers of our company or any of our subsidiaries, threatened against or affecting our company, our common stock, any of our subsidiaries or of our company's or our company's subsidiaries' officers or directors in their capacities as such, in which an adverse decision could have a material adverse effect.

Item 4. Mine Safety Disclosures.

Not applicable.

PART II

Item 5. Market for Common Equity, Related Stockholder Matters and Issuer Purchases of Equity Securities.

(a) Market Information

Our shares of common stock were cleared for trading under the symbol "TTWZ:OB" on the OTCBB on November 24, 2008, and later began trading on the OTCBB under the symbol "MSLP:OB" on April 27, 2010. Prior to this period, there was minimal trading in our common stock. The high and low prices for our common stock during the calendar quarters ended were:

Quarter ended	High	Low
December 31, 2011	$0.026	$0.007
September 30, 2011	$0.039	$0.014
June 30, 2011	$0.081	$0.025
March 31, 2011	$0.130	$0.036
December 31, 2010	$0.900	$0.050
September 30, 2010	$1.030	$0.410
June 30, 2010	$1.180	$0.950
March 31, 2010	$-	$-
December 31, 2009	$-	$-
September 30, 2009	$-	$-
June 30, 2009	$-	$-
March 31, 2009	$-	$-

Quotations on the OTCBB reflect bid and ask quotations, may reflect inter-dealer prices, without retail markup, markdown or commission, and may not represent actual transactions. In periods prior to June 30, 2010, there was no volume in the Company's common stock.

(b) Holders

As of April 13, 2012, we estimate that there were approximately 3,750 holders of record of our common stock. This figure does not take into account those shareholders whose certificates are held in the name of broker-dealers, "street name," or other nominees.

(c) Dividends

We have not paid any dividends to the holders of our common stock and we do not expect to pay any such dividends in the foreseeable future as we expect to retain our future earnings for use in the operation and expansion of our business.

(d) Securities Authorized for Issuance under Equity Compensation Plan

As of December 31, 2011, we had an employee stock option plan under which 5,000,000 shares had been reserved for issuance. The following table shows information with respect to this plan as of the fiscal year ended December 31, 2011.

Equity Compensation Plan Information

Plan category	Number of securities to be issued upon exercise of outstanding options, warrants and rights (a)	Weighted-average exercise price of outstanding options, warrants and rights(b)	Number of securities remaining available for future issuance under equity compensation plans (excluding securities reflected in column (a)) (c)
Equity compensation plans approved by security holders	1,617,500	$ 0.50	3,382,500
Equity compensation plans not approved by security holders	-	-	-
Total	1,617,500	$ 0.50	3,382,500

Unregistered Sales of Equity Securities

During the year ended December 31, 2011, the Company issued 487,281,174 shares of its common stock

Item 6. Selected Financial Data.

Not applicable.

Item 7. Management's Discussion and Analysis of Financial Condition and Results of Operations.

This report and other reports filed by our Company from time to time with the United States Securities and Exchange Commission (collectively the "Filings") contain or may contain forward-looking statements and information that are based upon beliefs of, and information currently available to, our management as well as estimates and assumptions made by our management. Readers are cautioned not to place undue reliance on these forward-looking statements, which are only predictions and speak only as of the date hereof. When used in the filings, the words "anticipate," "believe," "estimate," "expect," "future," "intend," "plan," or the negative of these terms and similar expressions as they relate us or our management identify forward-looking statements. Such statements reflect our current view with respect to future events and are subject to risks, uncertainties, assumptions, and other factors, including those set forth in the Risk Factors on page 10. Should one or more of these risks or uncertainties materialize, or should the underlying assumptions prove incorrect, actual results may differ significantly from those anticipated, believed, estimated, expected, intended, or planned.

Although we believe that the expectations reflected in the forward-looking statements are reasonable, we cannot guarantee future results, levels of activity, performance, or achievements. Except as required by applicable law, including the securities laws of the United States, we do not intend to update any of the forward-looking statements to conform these statements to actual results.

Our financial statements are prepared in accordance with accounting principles generally accepted in the United States ("GAAP"). These accounting principles require us to make certain estimates, judgments and assumptions. We believe that the estimates, judgments and assumptions upon which we rely are reasonable based upon information available to us at the time that these estimates, judgments and assumptions are made. These estimates, judgments and assumptions can affect the reported amounts of assets and liabilities as of the date of the financial statements as well as the reported amounts of revenues and expenses during the periods presented. Our financial statements would be affected to the extent there are material differences between these estimates and actual results. In many cases, the accounting treatment of a particular transaction is specifically dictated by GAAP and does not require management's judgment in its application. There are also areas in which management's judgment in selecting any available alternative would not produce a materially different result. The following discussion should be read in conjunction with our consolidated financial statements and notes thereto appearing elsewhere in this report.

Plan of Operation

Headquartered in Denver, Colorado, MusclePharm is a rapidly expanding healthy life-style company that develops and distributes a full line of National Sanitation Foundation International and scientifically approved, nutritional supplements that are 100% free of any banned substances. Based on years of research, MusclePharm products are created through an advanced six-stage research protocol involving the expertise of top nutritional scientists and field tested by more than 100 elite professional athletes from various sports including the National Football League, mixed martial arts, and Major League Baseball. The Company's propriety and award winning products address all categories of an active lifestyle including muscle building, weight loss, and maintaining general fitness through a daily nutritional supplement regimen. MusclePharm is sold in over 120 countries and available in over 5,000 U.S. retail outlets, including GNC, Vitamin Shoppe, and Vitamin World. The Company also sells its products in over 100 online stores, including bodybuilding.com, amazon.com and vitacost.com.

Our primary focus at the current time is on the following:

(1) Increase our distribution and sales;

(2) Continue aggressive marketing campaign to further build upon our brand and market awareness;

(3) Conduct additional testing of the safety and efficacy of our products; and

(4) Hire additional key employees to continue to strengthen the Company.

Results of Operations

For the Year Ended December 31, 2011(As Restated) Compared to the Year Ended December 31, 2010 (As Restated)

Revenues

Net revenues from the sale of products were approximately $17 million for the year ended December 31, 2011, as compared to revenue from the sale of products of approximately $3.2 million for the year ended December 31, 2010.

Sales activities during the year ended December 31, 2011, increased due to the increase in advertising and promotion efforts, as well as the change in the Company's manufacturers, which provided more consistent shipments to customers. The increase is also related to the significant capital spent on marketing with distributors and marketing and brand recognition with endorsements and sponsorships.

Cost of Sales

Cost of sales for the year ended December 31, 2011, were approximately $14.8 million or 86% of revenue, as compared to approximately $2.8 million or 88% of revenue for the year ended December 31, 2010. The cost of sales as a percentage of revenues were not consistent from December 31, 2010 to December 31, 2011. The cost of sales as a percentage of revenues increased due primarily to a reclassification of advertising expense to cost of sales in the amount of $374,455 during 2011 due to our restatement. There were no such reclassifications in 2010.

Operating Expenses

Operating expenses for the year ended December 31, 2011, were approximately $18.6 million, as compared to approximately $18.7 million for the year ended December 31, 2010.

The approximate $.1 million decrease is primarily due to an increase in stock based compensation – of approximately $3.7 million, an increase in depreciation expense of approximately $0.2 million and an increase in travel, meetings and entertainment of approximately $0.3 million, offset by a decrease in investment advisory services of approximately $2.4 million, a decrease in research and development costs of approximately $1.2 million and the decrease of advertising expense of $0.9 million .

Operating Loss

Operating loss for the year ended December 31, 2011 was approximately $16.2 million, as compared to approximately $18.3 million for the year ended December 31, 2010.

Interest Expense

Interest expense for the year ended December 31, 2011, was approximately $3.7 million, as compared to approximately $0.5 million for the year ended December 31, 2010. The increase in interest expense primarily relates to amortization of the debt discounts and debt issue costs of $3.5 million and interest charges incurred on our debt instruments of approximately $0.2 million.

Other Expenses

Other expenses for the year ended December 31, 2011, were approximately $7 million, as compared to approximately $1.3 million for the year ended December 31, 2010. The $5.7 million increase in other expenses is primarily due to an increase in derivative expense of approximately $4.7 million, an increase in interest expense of approximately $3.2 million and increases in the losses on settlement of accounts payable of approximately $3.4 million, offset by changes in the fair value of derivative liabilities of approximately $5.3 million and licensing income of approximately $0.2 million.

Net Loss

Net loss for the year ended December 31, 2011, was approximately $23.3 million, or loss per share of $0.08, as compared to the net loss of approximately $19.6 million or loss per share of $0.48 for the year ended December 31, 2010.

Inflation did not have a material impact on the Company’s operations for the period. Other than the foregoing, management knows of no trends, demands, or uncertainties that are reasonably likely to have a material impact on the Company’s results of operations.

Liquidity and Capital Resources

The following table summarizes total current assets, liabilities and working capital at December 31, 2011, compared to December 31, 2010.

	December 31, 2011	December 31, 2010	Increase/Decrease
Current Assets	$4,016,833	$2,494,441	$ 1,522,392
Current Liabilities	$17,710,100	$4,215,648	$ 13,494,452
Working Capital (Deficit)	$(13,693,267)	$(1,721,207)	$ (11,972,060)

Our primary source of operating cash has been through the sale of equity and through the issuance of convertible secured promissory notes and other short term debt as discussed below.

At December 31, 2011, the Company had cash of $659,764 and working capital deficit of approximately $13.7 million, compared to cash of $43,704 and a working capital deficit of approximately $1.7 million at December 31, 2010. The working capital deficit increase of approximately $12.0 million is primarily due to an increase in accounts payable and accrued liabilities of approximately $6.1 million, an increase in derivative liabilities of approximately $6.4 million and an increase of debt, net of discounts of approximately $1 million, offset by an increase to prepaid sponsorships of approximately $0.2 million and an increase in accounts receivable of approximately $2.1 million.

Cash used in operating activities was approximately $5.8 million for the year ended December 31, 2011, as compared to cash used in operating activities of approximately $3.8 million for the year ended December 31, 2010. The increase in cash used in operating activities of approximately $2.0 million for the year ended December 31, 2011, compared to the year ended December 31, 2010, was primarily due to an increase of accounts receivable of approximately $2.1 million, an increase in the change in fair value of derivative liabilities of approximately $5.3 million, a decrease of stock issued for services of approximately $7.6 million, offset by an increase in warrants issued for services of approximately $2.0 million, an increase in the amortization of debt discount and debt issue costs of approximately $3.0 million, an increase in the loss on settlement of accounts payable of approximately $1.7 million, the increase on the loss on conversion of debt of approximately $1.7 million and the increase in derivative expense of approximately $4.7 million.

Cash used in investing activities was $831,511 for the year ended December 31, 2011, as compared to cash used in investing activities of $117,303 for the year ended December 31, 2010. The increase in cash used in investing activities represents purchases of property and equipment, and the build out of our gym facility. We also maintain a website (www.musclepharm.com), designed for customers and investors. Future investments in property and equipment, as well as further development of our Internet presence will largely depend on available capital resources.

Cash flows provided by financing activities were approximately $7.2 million for the year ended December 31, 2011, as compared to cash flows provided by financing activities of approximately $4.0 million for the year ended December 31, 2010. The approximately $3.3 million increase is due to the $4.5 million increase in proceeds from issuance of debt, offset by decreases in the proceeds from issuance of debt – related party of approximately $0.4 million, the decrease in the proceeds from the issuance of common stock and warrants – net of recapitalization payment of $0.6 million and the increase in the cash paid for debt issue costs of approximately $0.3 million.

Cash Flows From Financing Activities: For the Years Ended	December 31 2011	December 31 2010
Cash (cash overdraft)	-	(17,841)
Due to related party	-	(27,929)
Proceeds from issuance of debt	6,612,900	2,140,608
Proceeds from issuance of debt - related party	-	358,077
Repayment of debt	(75,285)	-
Debt issuance costs	(263,283)	-
Proceeds from issuance of preferred stock	100,000	-
Losses incurred in connection with debt conversion prior to maturity	875,000	1,503,569
Net Cash Provided By Financing Activities	7,249,332	3,956,484

Off-Balance Sheet Arrangements

Other than the operating leases, as of December 31, 2011, MusclePharm did not have any off-balance sheet arrangements.

Critical Accounting Policies

The preparation of financial statements in conformity with U.S. generally accepted accounting principles requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period.

Making estimates requires management to exercise significant judgment. It is at least reasonably possible that the estimate of the effect of a condition, situation or set of circumstances that existed at the date of the financial statements, which management considered in formulating its estimate could change in the near term due to one or more future non-conforming events. Accordingly, the actual results could differ significantly from estimates.

Restatement

On May 14, 2012, the Company determined that a material misstatement exists in the Company's 2011 quarterly and 2011 and 2010 annual financial statements. The Company concluded that the following financial statements contained material misstatements: (i) the Company's audited financial statements for the year ended December 31, 2011, filed in an annual report on Form 10-K with the U.S. Securities and Exchange Commission (the "SEC") on April 16, 2012; (ii) the Company's audited financial statements for the year ended December 31, 2010, filed in an annual report on Form 10-K with the U.S. Securities and Exchange Commission (the "SEC") on April 1, 2011; (iii) the Company's unaudited financial statements for the period ended September 30, 2011, filed in a quarterly report on Form 10-Q with the SEC on November 14, 2011; (iv) the Company's unaudited financial statements for the period ended June 30, 2011, filed in a quarterly report on Form 10-Q with the SEC on August 16, 2011; and (v) the Company's unaudited financial statements for the period ended March 31, 2011, filed in a quarterly report on Form 10-Q with the SEC on May 23, 2011.

The foregoing financial statements contained material misstatements pertaining to the Company's calculation of net sales and presentation of general and administrative expenses and cost of sales. The Company has determined that advertising related credits that were granted to customers fell within the guidance of ASC No. 605-50-55 (" *Revenue Recognition" – Customer Payments and Incentives – Implementation Guidance and Illustrations)* . The guidance indicates that, absent evidence of benefit to the vendor, appropriate treatment requires netting these types of payments against revenues and not expensing as advertising expense. The Company also noted other credits and discounts that, upon further review, had been previously classified as advertising expense as a component of general and administrative expense that require a reallocation of presentation as amounts to be netted against revenues. Additionally, for the year ended December 31, 2011, the Company reclassified certain advertising expenses relating to sample items to cost of sales. There were no such reclassifications for 2010. The Company's net loss and loss per share will not be affected by this reallocation in the statement of operations.

Promotions, credits and non-specific advertising with its customers have been reclassified from general and administrative expenses to revenues.

Samples shipped to customers not clearly identifiable were reclassified from general and administrative expense to cost of sales.

	Year Ended December 31, 2011 As Restated	Adjustments	Year Ended December 31, 2011 As Issued	Year Ended December 31, 2010 As Restated	Adjustments	Year Ended December 31, 2010 As Issued
Sales - net	$ 17,212,636	$ (3,625,701)	$ 20,838,337	$ 3,202,687	$ (844,608)	$ 4,047,295
Cost of sales	14,845,069	374,455	14,470,614	2,804,274	-	2,804,274
Gross profit	2,367,567	(4,000,156)	6,367,723	398,413	(844,608)	1,243,021
General and administrative expenses	18,587,727	(4,000,156)	22,587,883	18,650,249	(844,608)	19,494,857
Loss from operations	(16,220,160)	-	(16,220,160)	(18,251,836)	-	(18,251,836)
Other income (expense)						
Derivative expense	(4,777,654)	-	(4,777,654)	(93,638)	-	(93,638)
Change in fair value of derivative liabilities	5,162,100	-	5,162,100	(149,306)	-	(149,306)
Loss on settlement of accounts payable and debt	(3,862,458)	-	(3,862,458)	(433,400)	-	(433,400)
Interest expense	(3,711,278)	-	(3,711,278)	(480,589)	-	(480,589)
Other expense	(121,500)	-	(121,500)	(160,568)	-	(160,568)
Licensing income	250,000	-	250,000	-	-	-
Total other income	(7,060,790)	-	(7,060,790)	(1,317,501)	-	(1,317,501)

(expense) - net						
Net loss	$ (23,280,950)	$ -	$ (23,280,950)	$ (19,569,337)	$ -	$ (19,569,337)
Net loss available to common stockholders						
Net loss	$ (23,280,950	$ -	$ (23,280,950	$ (19,569,337)	$ -	$ (19,569,337)
Series C preferred stock dividend	(293)	-	(293)	-	-	-
Net loss available to common stockholders	$ (23,280,657)	$ -	$ (23,280,657)	$ (19,569,337)	$ -	$ (19,569,337)
Net loss per common share available to common stockholders - basic and diluted	$ (0.08)	$ -	$ (0.08)	$ (0.48)	$ -	$ (0.48)
Weighted average number of common shares outstanding during the year – basic and diluted	281,484,658	-	281,484,658	41,141,549	-	41,141,549

Accounts Receivable

MusclePharm performs ongoing evaluations of its customer's financial condition and generally does not require collateral. Management reviews accounts receivable periodically and reduces the carrying amount by a valuation allowance that reflects management's best estimate of amounts that may not be collectible. Allowances, if any, for uncollectible accounts receivable are determined based upon information available and historical experience.

Property and Equipment

Property and equipment are stated at cost less accumulated depreciation. Included in property and equipment are website development costs which represent capitalized costs of design, configuration, coding, installation, and testing of the Company's website. Depreciation is computed on the straight-line method over the asset's useful lives which range from three to five years. Maintenance and repairs are charged to expense as incurred; improvements and betterments are capitalized.

Long-Lived Assets

MusclePharm's primary long-lived assets are property and equipment. The Company assesses the recoverability of its long-lived assets whenever events and circumstances indicate the carrying value of an asset or asset group may not be recoverable from estimated future cash flows expected to result from its use and eventual disposition.

Fair Value Measurements

The Company follows guidance for fair value measurements which defines fair value, establishes a framework for using fair value to measure financial assets and liabilities on a recurring basis, and expands disclosures about fair value measurements. The Company also applies the guidance to non-financial assets and liabilities measured at fair value on a non-recurring basis, which includes goodwill and intangible assets. The guidance establishes a hierarchy for inputs used in measuring fair value that maximizes the use of observable inputs and minimizes the use of unobservable inputs by requiring that the most observable inputs be used when available. Observable inputs are inputs that market participants would use in pricing the asset or liability developed based on market data obtained from sources independent of the Company. Unobservable inputs are inputs that reflect the Company's assumptions of what market participants would use in pricing the asset or liability developed based on the best information available in the circumstances. The hierarchy is broken down into three levels based on the reliability of the inputs as follows:

- Level 1 – Valuation is based upon unadjusted quoted market prices for identical assets or liabilities in active markets that the Company has the ability to access.

- Level 2 – Valuation is based upon quoted prices for similar assets or liabilities in active markets; quoted prices for identical or similar assets or liabilities in inactive markets; or valuations based on models where the significant inputs are observable in the market.

- Level 3 – Valuation is based on models where significant inputs are not observable. The unobservable inputs reflect the Company's own assumptions about the inputs that market participants would use.

Financial instruments consist of cash, accounts receivable, prepaid expenses, accounts payable and accrued expenses. The carrying amount of these financial instruments approximates fair value due to their short-term nature or the current rates at which the Company could borrow funds with similar remaining maturities. Unless otherwise noted, it is management's opinion that the Company is not exposed to significant interest, currency or credit risks arising from these financial statements.

Derivative Financial Instruments

Fair value accounting requires bifurcation of embedded derivative instruments such as conversion features in convertible debt or equity instruments, and measurement of their fair value for accounting purposes. In determining the appropriate fair value, the Company uses the Black-Scholes option-pricing model. In assessing the convertible debt instruments, management determines if the convertible debt host instrument is conventional convertible debt and further if there is a beneficial conversion feature requiring measurement. If the instrument is not considered conventional convertible debt, the Company will continue its evaluation process of these instruments as derivative financial instruments.

Revenue Recognition (As Restated)

The Company records revenue when all of the following have occurred: (1) persuasive evidence of an arrangement exists, (2) product has been shipped or delivered by the third party manufacturer, (3) the sales price to the customer is fixed or determinable, and (4) collectability is reasonably assured.

Depending on individual customer agreements, sales are recognized either upon shipment of products to customers or upon delivery. For one of our largest customers, which represent 14% of total revenue in 2011, revenue is recognized upon delivery.

The Company has determined that advertising related credits that were granted to customers fell within the guidance of ASC No. 605-50-55 (“ *Revenue Recognition” – Customer Payments and Incentives – Implementation Guidance and Illustrations)* . The guidance indicates that, absent evidence of benefit to the vendor, appropriate treatment requires netting these types of payments against revenues and not expensing as advertising expense.

The Company records store support, give aways, sales allowances and discounts as a direct reduction of sales. The Company recorded reductions to gross revenues totaling approximately $4,000,000 and $1,000,000 for the years ended December 31, 2011 and 2010, respectively.

The Company grants volume incentive rebates to certain customers based on contractually agreed percentages ranging from 2.5% - 5.5% as a percentage of sales once a certain threshold has been met. The credits are recorded as a direct reduction to sales. Included in the reductions to revenues above are volume incentive rebates. Total volume incentive rebates granted for the years ended December 31, 2011 and 2010 were approximately $500,000 and $0, respectively.

The Company has an informal 7-day right of return for products. There were nominal returns in 2011 and 2010.

During the years ended December 31, 2011 and 2010, the Company had the following concentrations of revenues with customers:

Customer	2011 (As Restated)		2010 As Restated)	
A	41	%	45	%

B	14	%	7	%
C	-	%	15	%

The Company does not manufacture or physically hold any inventory. Inventory is held and distributed by the Company's third party manufacturer.

Sponsorship and Endorsement Agreements

As a component of its marketing strategy, the Company enters into sponsorship and endorsement agreements with prominent athletes, trainers, and other high profile individuals that provide the Company ongoing sources of exposure to its products. The agreements sometimes specify certain contingencies that must be met to receive payments; others may require regular or periodic payments with no specified service or events that trigger payments under an agreement, or a combination of both. Agreements that are contingent upon the successful completion of an event prior to payment are considered unearned until the completion of the triggering event, and as such, no expense or liability is recorded until the successful completion of the triggering event. Where agreements are based on time and not on specific triggering events, the services are considered to be earned ratably over the period of the agreement, and as such expenses and liabilities are recorded ratably over the term of the agreement.

On September 1, 2011, the Company signed a sponsorship agreement with Zuffa Marketing, LLC, the owner of Ultimate Fighting Championship™ ("UFC™"). The Company is one of five primary sponsors with Dodge Trucks, Harley-Davidson, Bud Light and Tapout. The Company is the "Official Supplement Company of the UFC." The Company receives brand placement at UFC ™ events. Our agreement includes prominent logo placement on the mat, and our branding can be seen on Fox and pay-per-view worldwide.

Stock-Based Compensation

MusclePharm measures and recognizes compensation expense for all share-based awards made to employees and directors, including stock options and stock purchase warrants, based on estimated fair values. The Company must estimate the fair value of share-based awards on the grant date using an option pricing model. MusclePharm values share-based awards using the Black-Scholes option pricing model. The Black-Scholes model is highly complex and dependent on key estimates by management.

Recent Accounting Pronouncements

In May 2011, the FASB issued ASU No. 2011-04, which amended ASC Topic 820 to achieve common fair value measurements and disclosure requirements in U.S. GAAP and International Financial Reporting Standards ("IFRS"). The amendments in ASU No. 2011-05 result in common fair value measurement and disclosure requirements in U.S. GAAP and IFRSs. Consequently, the amendments change the wording used to describe many of the requirements in U.S. GAAP for measuring fair value and for disclosing information about fair value measurements. This amendment is effective for fiscal years, and interim periods within those years, beginning after December 15, 2011. The Company does not anticipate this amendment will have a material impact on its financial statements.

Item 7A. Quantitative and Qualitative Disclosures about Market Risk

Not applicable.

Item 8. Financial Statements and Supplementary Data.

Our financial statements are contained in pages F-1 through F-23, which appear at the end of this Annual Report.

Item 9. Changes in and Disagreements with Accountants on Accounting and Financial Disclosure.

Not applicable.

ITEM 9A. CONTROLS AND PROCEDURES

(a) EVALUATION OF DISCLOSURE CONTROLS AND PROCEDURES

Based on their evaluation as of the end of the period covered by this Annual Report on Form 10-K, our Chief Executive Officer and Chief Financial Officer have concluded that our disclosure controls and procedures (as defined

in Rules 13a-15(c) and 15d-15(e) under the Exchange Act) are not effective to ensure that information required to be disclosed by us in report that we file or submit under the Exchange Act is recorded, processed, summarized and reported within the time periods specified in the Securities and Exchange Commission's rules and forms and to ensure that information required to be disclosed by us in the reports that we file or submit under the Exchange Act is accumulated and communicated to our management, including our chief executive officer and chief financial officer, as appropriate to allow timely decisions regarding required disclosure.

(b) CHANGES IN INTERNAL CONTROL OVER FINANCIAL REPORTING

This company's management is responsible for establishing and maintaining internal controls over financial reporting and disclosure controls. Internal Control Over Financial Reporting is a process designed by, or under the supervision of, the Company's principal executive and principal financial officers, or persons performing similar functions, and effected by the issuer's board of directors, management and other personnel, to provide reasonable assurance regarding the reliability of financial reporting and the preparation of financial statements for external purposes in accordance with generally accepted accounting principles and includes those policies and procedures that:

1. Pertain to the maintenance of records that in reasonable detail accurately and fairly reflect the transactions and dispositions of the assets of the issuer;

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

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

Disclosure controls and procedures are designed to ensure that information required to be disclosed in reports filed under the Securities Exchange Act of 1934, as amended, is appropriately recorded, processed, summarized and reported within the specified time periods.

Management has conducted an evaluation of the effectiveness of our internal control over financial reporting for the year ended December 31, 2011 based on the framework established in Internal Control-Integrated Framework issued by the Committee of Sponsoring Organizations of the Treadway Commission ("COSO").

Based on this assessment, management concluded that as of May 2012 it had material weaknesses in its internal control procedures.

The Company’s assessment identified certain material weaknesses which are set forth below:

Functional Controls and Segregation of Duties

Because of the company's limited resources, there are limited controls over information processing, and insufficient internal controls over the accuracy, completeness and authorization of transactions.

There is an inadequate segregation of duties consistent with control objectives. Our company's management is composed of a small number of individuals resulting in a situation where limitations on segregation of duties exist. In order to remedy this situation we would need to hire additional staff to provide greater segregation of duties. Currently, it is not feasible to hire additional staff to obtain optimal segregation of duties. Management will reassess this matter in the following year to determine whether improvement in segregation of duty is feasible.

Accordingly, as the result of identifying the above material weaknesses we have concluded that these control deficiencies resulted in a reasonable possibility that a material misstatement of the annual or interim financial statements would not be prevented or detected on a timely basis by the company's internal controls.

Management believes that the material weaknesses set forth above were the result of the scale of our operations and are intrinsic to our small size. Management believes these weaknesses did not have a material effect on our financial results and is actively working to take remedial actions.

We are committed to improving our financial organization. As part of this commitment, we will create a position to segregate duties consistent with control objectives and will increase our personnel resources and technical accounting expertise within the accounting function when funds are available to us by preparing and implementing sufficient written policies and checklists which will set forth procedures for accounting and financial reporting with respect to the requirements and application of US GAAP and SEC disclosure requirements.

(1) We will continue to update the documentation of our internal control processes, including formal risk assessment of our financial reporting processes.

(2) Retain staff to assist in financial and accounting controls.

Subsequent to December 31 2011, we have undertaken the following steps to address the deficiencies stated above:

- Continued the development and documentation of internal controls and procedures surrounding the financial reporting process, primarily through the use of account reconciliations, and supervision.

We carried out an evaluation, under the supervision and with the participation of our management, including our PEO and PFO, of the effectiveness of the design and operation of our disclosure controls and procedures as of the end of the period covered by this Annual Report. Based upon that evaluation, the PEO and PFO concluded that the Company's disclosure controls and procedures were ineffective.

(c) Inherent Limitations on Effectiveness of Controls

Our management, including the Chief Executive Officer and Chief Financial Officer, does not expect that our disclosure controls or internal control over financial reporting will prevent all errors and all fraud. A control system, no matter how well designed and operated, can provide only reasonable, not absolute, assurance that the control system's objectives will be met. Further, the design of a control system must reflect the fact that there are resource constraints, and the benefits of controls must be considered relative to their costs. Because of the inherent limitations in all control systems, no evaluation of controls can provide absolute assurance that all control issues and instances of fraud, if any, within MusclePharm have been detected.

Item 9B. Other Information.

Not applicable.

PART III

Item 10. Directors, Executive Offices and Corporate Governance.

Directors and Executive Officers

The following table and text sets forth the names and ages of all our directors and executive officers and our key management personnel as of April 13, 2012. All of our directors serve until the next annual meeting of stockholders and until their successors are elected and qualified, or until their earlier death, retirement, resignation or removal. Executive officers serve at the discretion of the Board of Directors, and are elected or appointed to serve until the next Board of Directors meeting following the annual meeting of stockholders. Also provided is a brief description of the business experience of each director and executive officer and the key management personnel during the past five years and an indication of directorships held by each director in other companies subject to the reporting requirements under the Federal securities laws.

Name	Age	Position
Brad J. Pyatt	32	Chief Executive Officer and Director
Cory Gregory	33	Senior President and Director
Jeremy DeLuca	33	President and Chief Marketing Officer
Lawrence S. Meer	51	Chief Financial Officer
John H. Bluher	54	Chief Operating Officer

The biographies of each of our executive officers and directors are as follows:

Brad J. Pyatt, age 32, Chief Executive Officer, Director

Mr. Pyatt has served as the Chief Executive Officer and Director of the Company since February 18, 2010, and as President and Chief Executive Officer of Muscle Pharm, LLC, since its inception in April 2008. His background includes seven years of experience as a professional athlete, and more than five years of experience in the sports nutrition arena. Mr. Pyatt played in National Football League (NFL) for the Indianapolis Colts during the 2003, 2004,

and 2005 NFL seasons as well for the Miami Dolphins during the 2006 NFL season. Mr Pyatt also played in the Arena Football League (AFL) for the Colorado Crush during the 2007 and 2008 AFL seasons. Mr. Pyatt attended the University of Kentucky from 1999 to 2002, where he studied kinesiology exercise science, as well the University of Northern Colorado, from 2002 to 2003.

The Company believes that Mr. Pyatt's experience in the sports nutrition sector over the past five plus years, along with his background as a former professional athlete, give him an unique perspective on the nutrition industry as a whole and makes him a valuable member to the Company's board of directors.

Cory Gregory, age 33, Senior President, Director

Mr. Gregory is currently the Senior President and member of the Company's board of directors, roles he has served in since May 2010. Prior to joining the Company, Mr. Gregory served as the President, managing member, and owner of T3 Personal Training LLC ("T3") from April 2009 until November 2000. T3 was a personal training service that managed and oversaw over 40 clients using 7 trainers over a ten year period. During the same period, Mr. Gregory served as President of the Ohio Natural Bodybuilding Federation, a federation founded by Mr. Gregory in 2004 which hosted 14 bodybuilding competitions over a six year period. In 2004, Mr. Gregory purchased the Old School Gym, located in Pataskala, OH, which he continues to own at present day.

The Company believes that Mr. Gregory's extensive bodybuilding and personal training experience provide him with the insight necessary to understand the ongoing demands and changes to the nutrition industry and as such, makes him a valuable member to the Company's board of directors.

Jeremy DeLuca, age 33, President and Chief Marketing Officer

Mr. DeLuca is the Company's President and Chief Marketing Officer. Prior to joining the Company, from April 1999 to November 2010, Mr. DeLuca served as the President of Bodybuilding.com, an online sports nutrition and supplements company which he co-founded in 1999 ("Bodybuilding.com"). As President, Mr. DeLuca was actively involved in all aspects of Bodybuilding.com's business, with a focus on marketing, sales, and e-commerce. Mr. DeLuca's responsibilities also included managing all vendor relations, marketing strategies, sales promotions, store content and store site development. During Mr. DeLuca's tenure, Bodybuilding.com grew tremendously, achieving annual sales of over $200,000,000 in 2010.

Lawrence S. Meer, age 51, Chief Financial Officer

Mr. Meer has served as Chief Financial Officer of the Company since July 2010. Prior to becoming the Chief Financial Officer he was the Director of Finance at Muscle Pharm, LLC from October 2009 to July 2010. His other past experience includes daily cash management and treasury functions, including the establishment of credit and collection procedures to maximize cash flow, reduce corporate debt and enhance shareholder value. He previously served as President and Chief Financial Officer in Miami, FL, at Color It, Inc., a textile finishing business, from March 2002 to December 2008. Mr. Meer also previously served as Executive Vice President at Customer Assets in Denver, CO, an India-based call center, from 2000 to 2002. Prior to joining Customer Assets, he was Chief Financial Officer and Chief Operating Officer at GS Sportswear in Denver, CO, a sportswear promotional company, from 1998 to 2000. Mr. Meer also served as Chief Financial Officer at Davis Audio-Visual, Inc., a retailer of audio-visual equipment, from 1996 to 1998; and Vice President of Finance at Pacer Cats in Englewood, CO., a ticketing and concession software provider from 1991 to 1996. Mr. Meer earned a BS in accounting from the University of Colorado at Boulder.

John H. Bluher, age 54, Chief Operating Officer

Mr. Bluher is a specialist in corporate governance for growing companies. He is also a specialist in investment management, capital structuring, merger and acquisition, private equity and valuations of public and private companies. He has significant experience working with corporate structuring, corporate boards and committees, risk management, and public company corporate governance. His experience also includes negotiating transactions and purchases, and sales of assets and properties on a global basis. He has deep experience in creating and implementing corporate governance plans, working in the corporate board room, and as director of risk, developing internal audit programs and insurance programs for public companies. During 2010, Mr. Bluher provided consulting services to a leading financial advisory and management consultant firm. Mr. Bluher was responsible for managing transactions, business development, developing corporate governance standards and corporate structuring for companies. Since December 2009, Mr. Bluher assisted in raising capital, marketing and co-managed Coachman Energy Funds at Caddis Capital, LLC, a private equity portfolio focused on oil and gas investments. From February 2010 to August 2010, Mr.

Bluher acted as investment banker and special financial advisor to the AARP Mutual Fund Board of Trustees in a platform divestiture. From December 2007 to May 2009, Mr. Bluher served as managing director and general counsel at Lehman Brothers, Inc.'s (NYSE:LEH) investment management division. Mr. Bluher also served as global chief legal and compliance officer and managing director of Neuberger Berman during this period. From August 2004 to June 2007, Mr. Bluher served as general counsel and director of risk and Janus Capital, Inc. (NYSE:JNS). From June 2002 to July 2004, Mr. Bluher served as executive vice president, general counsel and corporate secretary and director of risk management of Knight Trading Group (NASDAQ:NITE). From January 2001 to May 2002, Mr. Bluher served as senior vice president and global chief compliance officer for Prudential Securities, Inc. (NYSE:PRU). From October 1997 to January 2001, Mr. Bluher served as general counsel and chief compliance officer of Sun America, Inc. (NYSE:SAI) later (NYSE:AIG). From 1992 – 1997, Mr. Bluher served as senior vice president, regional and divisional Counsel at Prudential Securities, Inc. From 1987 to 1992, Mr. Bluher was senior counsel for the Division of Enforcement at the Securities and Exchange Commission. Mr. Bluher holds a Bachelor of Science and a J.D. degree from the University of Wyoming and holds FINRA Series 7, Series 24 and Series 14 licenses. He has served on the boards of ICI Mutual Insurance Company, the NASDAQ Chairman's Advisory Board, Cherry Hills Founders Group, Inc., Targeted Medical Pharma, Inc., Arete Industries, Inc., and Safe Communications, Inc., and the University of Wyoming Foundation Board, and College of Law Advisory Board. Mr. Bluher is a frequent speaker at financial services industry meetings and conferences.

Advisory Board

We have established an Advisory Board currently consisting of nine members, which serves to advise management with respect to product formulations, product ideas, marketing and related matters. Members of the Advisory Board do not meet on a formal or regular basis. Our management team consults with one or more members of the Advisory Board as needed, from time to time, by means of meetings or telephone conference calls.

Following is a brief description of the background of our advisory board members:

Dr. Eric Serrano **– Chief Medical Advisor .** Dr. Serrano has been practicing medicine in the State of Ohio for over 12 years and is considered one of the leading sports nutrition doctors in the country. His clients include a wide array of athletes from the NFL, NHL, and MLB, in addition to many elite amateur athletes. Dr. Serrano was a professor of family practice medicine at Ohio State University, where he was awarded Professor of The Year and Preceptor of The Year. Dr. Serrano currently lectures across the country to universities, medical groups and health & fitness conferences on the topics of sports nutrition, performance enhancement, and injury prevention. Dr. Serrano's expertise in blood analysis, sports nutrition, and injury prevention gives athletes the advantage over the competition. He has formulated numerous nutritional supplements for some of the leading nutritional companies on the market and also been a contributing writer for some of the leading health and fitness magazines. Dr. Serrano has been involved in the final formulations for each of our products. Dr. Serrano received his B.A. from Kansas State University in Biology, his M.A. from Kansas State University in Exercise Physiology, and his M.D. from the University of Kansas Medical School.

Roscoe M. Moore, Jr. **– Chief Scientific Director.** A Former U.S. Assistant Surgeon General, Dr. Roscoe M. Moore, Jr. served with the United States Department of Health and Human Services (HHS) and was for the last twelve years of his career the principal person responsible for global development support within the Office of the Secretary, HHS, with primary emphasis on Continental Africa and other less developed countries of the world (e.g. Indonesia, Malaysia, and Vietnam). He was the principal liaison person between the HHS and Ministries of Health in Africa with regard to the development of infrastructure and technical support for the delivery of preventive and curative health needs for the continent. Dr. Moore represented the HHS in cooperative international efforts with African nations in addressing continued health and human resource problems. Dr. Moore received his undergraduate and Doctor of Veterinary Medicine degrees from Tuskegee Institute; his Master of Public Health degree in Epidemiology from the University of Michigan; and his Doctor of Philosophy degree in Epidemiology from the Johns Hopkins University. He was awarded the Doctor of Science degree (Honoris Causa) in recognition of his distinguished public health career by Tuskegee University. Dr. Moore was a career officer within the Commissioned Corps of the United States Public Health Service (USPHS) entering with the U.S. National Institutes of Health and rising to the rank of Assistant United States Surgeon General (Rear Admiral, USPHS) within the Immediate Office of the Secretary, HHS. He was selected as Chief Veterinary Medical Officer, USPHS, by Surgeon General C. Everett Koop.

Dr. Richard Ogden PHD, (CSCS) **– Medical Advisor**

Dr. Odgen's career in clinical research and development spans nearly forty years. After earning a Ph.D. from Cambridge University, his career started with postdoctoral research studying RNA transcription and processing. Following that, he undertook independent research, funded by the National Science Foundation. In 1984, he joined Agouron Pharmaceuticals, Inc. as one of its founding scientists. Following Agouron's merger with Pfizer, he served as a Senior Director and was the scientific liaison for the Agouron/Pfizer commercial and corporate organizations. In this role, he worked with organizations all over the world. In 2006, Dr. Ogden, co-founded RORR Inc., a medical, scientific Consulting and Education company with clients in the U.S. and Europe. In addition to publication in numerous medical journals, he is co-editor of two books relating to AIDS therapy.

Dr. Michael Ray Stevens **– Advisor.** Dr. Stevens has over twenty years of well diversified experience in the healthcare and pharmaceutical industry. Dr. Stevens spent 17 years at Bristol-Myers Squibb, where he held positions of increasing responsibility in the areas of Market Research (Oncology and HIV), Marketing (Oncology), and Medical Affairs (HIV). In addition served as a member of the Executive Council for the Forum for Collaborative HIV Research — a public-private partnership facilitating discussion on emerging issues in HIV clinical research and working to translate research results into patient care. He has also served on 15 Protocol Committees within the Adult AIDS Clinical Trials Group (ACTG). Michael received his BS Pharmacy and Doctor of Pharmacy degrees from Purdue University.

***Dr. Ron Sekura* – Director of Therapeutic Research.** Dr. Sekura is the former Chief of the Pharmaceutical and Regulatory Affairs Branch of the Division of AIDS at The National Institute of Allergy and Infectious Diseases (NIAID) of the National Institute of Health (NIH) as well as a former Research Chemist at The National Institute of Child Health and Human Development (NICHD) at the NIH and the Center for Biologics Evaluation and Research (CBER), and FDA. He received his Bachelor of Science and Master of Science in Biochemistry degrees at Pennsylvania State University and his PhD at Cornell University. Dr. Sekura is the author of over sixty scientific publications.

***Mariel Selbovitz* – Director of Global Therapeutics Product Procurement Development.** Ms. Selbovitz is a graduate of Cornell University and received her Master's in Public Health at the Johns Hopkins University Bloomberg School of Health. She worked as the Client Intake Specialist at Positive Health Project and Syringe Exchange Program Coordinator at the Foundation for Research on Sexually Transmitted Diseases and is a partner in BioEquity Partners. Selbovitz is a member of the Cornell AIDS Clinical Trials Group Community Advisory Board and AIDS Treatment Advocacy Coalition. She presented at the 5th European Conference on Clinical and Social Research on AIDS and Drugs, International Conference on Antiviral Research, 5th IAS Conference on HIV Pathogenesis, Treatment and Prevention and XVIII International AIDS Conference.

***Louie Simmons* – Chief Strength Advisor.** Mr. Simmons is a strength consultant for the New England Patriots, Green Bay Packers, Seattle Seahawks, Cleveland Browns, and numerous Football Bowl Subdivision college football teams. Mr. Simmons is the owner of the West Side Barbell, located in Columbus, Ohio.

***Greg Jackson* – Director of Fight Development.** Mr. Jackson is an expert in mixed martial arts, representing a combination of basic Judo and wrestling. He has trained and developed top-ranked fight teams, with several fights appearing on spike TV's Ultimate Fighter.

***Paul Dillet* – Chief Bodybuilding Advisor.** Mr. Paul Dillet is one of the most influential bodybuilders and a legend in the bodybuilding world. He has been instrumental in creating a new era in fitness and bodybuilding for the everyday athlete.

Legal Proceedings

None of the members of the board of directors or other executives has been involved in any bankruptcy proceedings, criminal proceedings, any proceeding involving any possibility of enjoining or suspending members of our board of directors or other executives from engaging in any business, securities or banking activities, and have not been found to have violated, nor been accused of having violated, any Federal or State securities or commodities laws.

Director Independence

On an annual basis, each director and executive officer will be obligated to disclose any transactions with our Company and any of its subsidiaries in which a director or executive officer, or any member of his or her immediate family, have a direct or indirect material interest. Following completion of these disclosures, our board of directors will make an annual determination as to the independence of each director using the current standards for "independence" that satisfy both the criteria for the Nasdaq and the NYSE Amex Equities.

As of December 31, 2011, the board of directors determined that the Company does not currently have any directors that are considered "independent" under the aforementioned standards.

Committees of the Board of Directors

Concurrent with having sufficient members and resources, the board of directors intends to establish an audit committee and a compensation committee. The audit committee will review the results and scope of the audit and other services provided by the independent auditors and review and evaluate the system of internal controls. The compensation committee will review and recommend compensation arrangements for the officers and employees. No final determination has yet been made as to the memberships of these committees or when we will have sufficient members to establish committees. We believe that we will need a minimum of three (3) independent directors to have effective committee systems.

Item 11. Executive Compensation.

Summary Compensation Table

The following summary compensation tables sets forth all compensation awarded to, earned by, or paid to the named executive officers and directors by us during the period ended December 31, 2011, 2010, and 2009.

Name and Principal Position	Year	Salary ($)	Bonus ($)	Stock Awards ($)	Option Awards ($)	Non-Equity Incentive Plan Compensation ($)	All Other Compen-sation ($)	Total ($)
Brad J. Pyatt	2011	$250,000	$170,410	$1,555,921	$0	$ 0	$ 0	$1,976,331
Chief Executive Officer	2010	$194,821	$0	$2,650,000	$0	$ 0	$ 0	$2,844,821
	2009	$133,992	$0	$0	$0	$ 0	$ 0	$133,992
Cory Gregory	2011	$150,000	$170,410	$1,555,921	$0	$ 0	$ 0	$1,876,331
Senior President	2010	$78,892	$0	$2,650,000	$0	$ 0	$ 0	$2,728,892
	2009	$17,846	$0	$0	$0	$ 0	$ 0	$17,846
Lawrence S. Meer	2011	$74,400	$0	$0	$0	$ 0	$ 0	$74,400
Chief Financial Officer	2010	$75,493	$0	$0	$228,000(1)	$ 0	$ 0	$303,493
Leonard K. Armenta (2)	2011	$86,400	$0	$0	$0	$ 0	$ 0	$86,400
Former Executive Vice President	2010	$83,215	$0	$0	$228,000(1)	$ 0	$ 0	$311,215
	2009	$54,799	$0	$0	$0	$ 0	$ 0	$54,799
Jeremy DeLuca	2011	$65,833	$170,410	$1,555,921	$0	$ 0	$ 0	$1,792,164
President, Chief Marketing Officer								
John H. Bluher	2011	$36,458	$50,000	$00	$0	$ 0	$ 0	$86,458
Chief Operating Officer								

Explanatory Information Relating to 2011 Summary Compensation Table

Please note the following points in connection with the information in the 2011 Summary Compensation Table:

The compensation of the executive officers of the Company is reviewed on an annual basis by the board of directors. Each year, the Company considers whether to adjust the base salaries of senior management, including the executive officers, in order to reward individual performance, keep pace with cost of living increases and respond to competitive considerations.

2011 OUTSTANDING EQUITY AWARDS AT FISCAL YEAR-END

	OPTION AWARDS					STOCK AWARDS			
Name (a)	Number of Securities Underlying Unexercised Options Exercisable (#) (b)	Number of Securities Underlying Unexercised Options Unexercisable (#) (c)	Equity Incentive Plan Awards: Number of Securities Underlying Unexercised Unearned Options (#) (d)	Option Exercise Price ($) (e)	Option Expiration Date (f)	Number of Shares or Units of Stock That Have Not Vested (#) (g)	Market Value of Shares or Units of Stock That Have Not Vested ($) (h)	Equity Incentive Plan Awards: Number of Unearned Shares, Units or Other Rights That Have Not Vested (#) (i)	Equity Incentive Plan Awards: Market or Payout Value of Unearned Shares, Units or Other Rights That Have Not Vested (#) (j)
Brad J. Pyatt *Chief Executive Officer*	-	-	-	-	-	-	-	-	-
Cory Gregory *Senior President*	-	-	-	-	-	-	-	-	-
Lawrence S. Meer *Chief Financial Officer*	1,000,000 (1)	-	-	$ 0.50	4/2/2015	-	-	-	-
Leonard K. Armenta (2) *Former Executive VP*	-	-	-	-	-	-	-	-	-
Jeremy DeLuca *Chief Marketing Officer* *President*	-	-	-	-	-	-	-	-	-
John H. Bluher *Chief Operating Officer*	-	-	-	-	-	-	-	-	-

(1) Represents 1,000,000 options issued, valued on the date of grant, April 2, 2010.

(2) Resigned on September 16, 2011.

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Director Compensation

The following summary compensation table sets forth all compensation awarded to, earned by, or paid to the named directors by us during the years ended December 31, 2011, 2010 and 2009.

Name And Principal Position (a)	Year (b)	Salary ($) (c)	Bonus ($) (d)	Stock Awards ($) (e)	Option Awards ($) (f)	Non-Equity Incentive Plan Compensation ($) (g)	All Other Compensation ($) (h)	Total ($) (i)
Brad J. Pyatt	2011	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0
Director	2010	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0
	2009	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0
Cory Gregory	2011	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0
Director	2010	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0
	2009	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0	$ 0

Employment Agreements

Brad J. Pyatt, Chief Executive Officer

On August 15, 2011, the Company entered into an employment agreement (the “Pyatt Employment Agreement”) with Brad J. Pyatt, individually, pursuant to which Mr. Pyatt will serve as the Company’s Chief Executive Officer (the “CEO”). The term of the Pyatt Employment Agreement is for a period of sixty (60) months, commencing retroactively on January 1, 2011, and expiring on December 31, 2015 (the “Pyatt Term”). Pursuant to the terms of the Employment Agreement, the CEO is to receive a base salary of $250,000 for the 2011 calendar year; $350,000 for the 2012 calendar year; $400,000 for the 2013 calendar year; $450,000 for the 2014 calendar year; and $500,000 for the 2015 calendar year. Further, the CEO shall receive, upon execution of the Pyatt Employment Agreement, 31 shares of the Company’s Series B Preferred Stock. In addition, upon the three year anniversary of the Pyatt Employment Agreement, the CEO shall receive 10,000 shares of the Company’s Series A Preferred Stock.

During the Pyatt Term, the CEO’s responsibilities will include all aspects of the day to day business operations of the Company. The CEO shall also be responsible for determining necessary strategic partnerships and investment opportunities relating to the Company, both nationally and internationally, and shall have wide discretion in implementing the vision, strategic goals and operational mission of the Company. The CEO shall, on a full time and

exclusive basis, devote all of his business time, attention and energies to the operations of the Company and other duties as required by the Pyatt Employment Agreement, and shall use his best efforts to advance the best interests of the Company.

On November 14, 2011, the Company entered into an amended and restated employment agreement with Mr. Pyatt. The parties amended the Pyatt Employment Agreement in order to amend section 3(c) as it relates to Mr. Pyatt's bonus payment. The amended Pyatt Employment Agreement now provides that, for each one million dollars ($1,000,000) in revenue growth achieved by the Company from the revenue figure reported for the prior fiscal year, Mr. Pyatt shall receive (i) ten thousand dollars ($10,000) and (ii) one hundred thousand dollars ($100,000) worth of the Company's common stock, such stock to be valued based on the average closing price for the twenty (20) trading days prior to the date of issuance of such stock. The aforementioned payments to Mr. Pyatt shall be made within 90 days after the end of the Company's fiscal year.

Cory Gregory, Senior President

On August 15, 2011, the Company entered into an employment agreement (the "Gregory Employment Agreement") with Cory Gregory, individually, pursuant to which Mr. Gregory will serve as the Company's Senior President (the "Senior President"). The term of the Gregory Employment Agreement is for a period of sixty (60) months, commencing retroactively on January 1, 2011, and expiring on December 31, 2015 (the "Gregory Term"). Pursuant to the terms of the Gregory Employment Agreement, the Senior President is to receive a base salary of $150,000 for the 2011 calendar year; $200,000 for the 2012 calendar year; $250,000 for the 2013 calendar year; $300,000 for the 2014 calendar year; and $350,000 for the 2015 calendar year. Further, the Senior President shall receive, upon execution of the Gregory Employment Agreement, 20 shares of the Company's Series B Preferred Stock. In addition, upon the three year anniversary of the Gregory Employment Agreement, the Senior President shall receive 10,000 shares of the Company's Series A Preferred Stock.

During the Gregory Term, the Senior President's responsibilities will include, but shall not be limited to, on a full time and exclusive basis, devoting all of his business time, attention and energies to the operations of the Company and other duties as required by the Gregory Employment Agreement and as directed by the Board of Directors, and shall use his best efforts to advance the best interests of the Company.

On November 14, 2011, the Company entered into an amended and restated employment agreement with Mr. Gregory. The parties amended the Gregory Employment Agreement in order to amend section 3(c) as it relates to Mr. Gregory's bonus payment. The amended Gregory Employment Agreement now provides that, for each one million dollars ($1,000,000) in revenue growth achieved by the Company from the revenue figure reported for the prior fiscal year, Mr. Gregory shall receive (i) ten thousand dollars ($10,000) and (ii) one hundred thousand dollars ($100,000) worth of the Company's common stock, such stock to be valued based on the average closing price for the twenty (20) trading days prior to the date of issuance of such stock. The aforementioned payments to Mr. Gregory shall be made within 90 days after the end of the Company's fiscal year.

John H. Bluher, Chief Operating Officer

On September 16, 2011, the Company entered into an employment agreement (the "Bluher Employment Agreement") with John H. Bluher, individually ("Bluher"), appointing Bluher as the Company's Chief Operating Officer.

Pursuant to the terms of the Bluher Employment Agreement, Bluher is to serve as the Company's Chief Operating Officer from September 16, 2011 (the "Bluher Effective Date"), until September 15, 2013 (the "Bluher Term"). Upon expiration of the Bluher Term, the Bluher Employment Agreement shall be automatically renewed unless either the Company or Bluher provides the other party with written notice at least sixty (60) days prior to the last date of the respective term. During the Bluher Term, Bluher's responsibilities will include general oversight and management of the Company's daily operations, as well as any responsibilities delegated to him by the Company's Chief Executive Officer or board of directors (the "Bluher Duties").

In consideration for performance of the Bluher's Duties during the Term, Bluher is to receive an initial base salary of one hundred and seventy five thousand dollars ($175,000) per year (the "Bluher Base Salary"), any increases to such salary during the Bluher Term to be determined at the discretion of the Company. Bluher is also eligible to receive an annual performance bonus based on certain goals and performances levels mutually established by the parties.

Bluher was also entitled to receive, beginning on December 31, 2012, and on each successive calendar year end thereafter, stock options to purchase shares of the Company's common stock in the amount of five hundred thousand dollars ($500,000) (the "2012 Options"). The 2012 Options were to be exercisable into shares of the Company's common stock at an exercise price equal to the average of the high and low reported selling prices of the Company's common

stock on the date of grant and vest in accordance with the schedule outlined in the Bluher Employment Agreement.

On March 13, 2012, the Company and Bluher executed an amendment to the Bluher Employment Agreement, whereby Bluher waived his rights to the equity based compensation in both the Bluher Employment Agreement and a consulting agreement with Endion Capital, LLC, and now is to receive (i) 20,000,000 shares of the Company's common stock with piggy-back registration rights, subject to a lock-up period of one year and (ii) a warrant to purchase 10,000,000 shares of the Company's common stock at an exercise price of $0.008 per share, subject to a lock-up period of six months.

Jeremy DeLuca, President, Chief Marketing Officer

On November 14, 2011 (the "DeLuca Execution Date"), the Company entered into an employment agreement (the "DeLuca Employment Agreement") with Jeremy DeLuca, the Company's President and Chief Marketing Officer (the "President"). The term of the DeLuca Employment Agreement commences on the DeLuca Execution Date and expires on December 31, 2014 (the "DeLuca Term"). Pursuant to the terms of the DeLuca Employment Agreement, the President is to receive a base salary of $125,000 for the 2011 calendar year; $175,000 for the 2012 calendar year; $225,000 for the 2013 calendar year; and $300,000 for the 2014 calendar year. In addition, upon the three year anniversary of the DeLuca Employment Agreement, the President shall receive 5,000 shares of the Company's Series A Preferred Stock.

During the DeLuca Term, the President's responsibilities will include all aspects of the day to day business operations of the Company. The President shall also be responsible for determining necessary strategic partnerships and investment opportunities relating to the Company, both nationally and internationally, and shall have wide discretion in implementing the vision, strategic goals and operational mission of the Company. The President shall, on a full time and exclusive basis, devote all of his business time, attention and energies to the operations of the Company and other duties as required by the DeLuca Employment Agreement, and shall use his best efforts to advance the best interests of the Company.

Item 12. Security Ownership of Certain Beneficial Owners and Management and Related Stockholder Matters.

The following table sets forth information known to MusclePharm with respect to the beneficial ownership of MusclePharm's common stock as of April 13, 2012, unless otherwise noted, by:

- each stockholder known to MusclePharm to own beneficially more than 5% of MusclePharm's common stock;
- each of MusclePharm's directors;
- each of MusclePharm's executive officers; and
- all of MusclePharm's current directors and executive officers as a group.

Beneficial ownership is determined in accordance with the rules of the SEC and generally includes voting or dispositive power with respect to securities. Common shares relating to options or warrants currently exercisable, or exercisable within 60 days of April 12, 2012, are deemed outstanding for computing the percentage of the person holding such securities but are not deemed outstanding for computing the percentage of any other person. Except as indicated by footnote, and subject to the community property laws where applicable, the persons or entities named in the tables have sole voting and dispositive power with respect to all shares shown as beneficially owned by them.

Name and Address of Beneficial Owner	Amount and Nature of Beneficial Ownership	Percentage of Beneficial Ownership (1)
Brad J. Pyatt 4721 Ironton St	166,962,288	11.6 %

Denver, CO 80239			
Cory Gregory 4721 Ironton St Denver, CO 80239	158,665,986	11.0	%
Lawrence S. Meer 4721 Ironton St Denver, CO 80239	0	0	%
Jeremy DeLuca 4721 Ironton St Denver, CO 80239	148,182,972	10.2	%
John H. Bluher 4721 Ironton St Denver, CO 80239	0	0	%
All executive officers and directors as a group (5 persons)	473,811,226	32.8	%
Drew Ciccarelli	105,000,000(2)	7.27	%
All executive officers, directors and 5% holders as a group	578,811,226	40.07	%

(1) Percent of class based on 1,444,521,399 common shares outstanding as of April 12, 2012. This percentage does not include preferred stock ownership or other ownership of convertible securities.

(2) These shares are held by Mr. Ciccarelli individually, and through (i) TSX Ventures, LLC, a South Carolina limited liability company and (ii) Five Star Consulting, LLC, a Nevada limited liability company, entities of which Mr. Ciccarelli maintains sole dispositive voting power.

Changes in Control

We are not aware of any arrangements that may result in “changes in control” as that term is defined by the provisions of Item 403(c) of Regulation S-K.

Item 13. Certain Relationships and Related Transactions, and Director Independence.

Any future transactions or loans between us and our officers, directors, principal stockholders or affiliates will be on terms no less favorable to us than could be obtained from an unaffiliated third party, and will be approved by a majority of disinterested directors.

On February 18, 2010, the Company issued a total of 26,000,000 shares of its common stock to the 12 former owners of Muscle Pharm, LLC in reliance on the exemption from registration provided by Section 4(2) of the Securities Act of 1933, as amended.

On November 18, 2010, Brad Pyatt, the Company’s Chief Executive Officer, loaned the Company $100,000 and received an 8% Convertible Promissory Note exchange. On November 23, 2010, Mr. Pyatt loaned the Company $256,250 and received an 8% Convertible Promissory Note in exchange. On December 14, 2010, Mr. Pyatt converted all principal and accrued interest underlying the notes ($358,077.40) into 7,161,548 shares of the Company’s common stock.

Muscle Pharm, LLC was formed as a Colorado limited liability company on April 22, 2008. The initial owners of Muscle Pharm LLC were Brad J. Pyatt and Cory Gregory. Mr. Pyatt received a 60% membership interest in exchange for his contribution of formulations for potential products, contacts with GNC Canada and other potential customers, and contacts with professional athletes. Mr. Gregory received a 40% membership interest in exchange for his contacts with Dr. Serrano, Louie Simmons, potential distributors, professional athletes and potential investors. Neither Mr. Pyatt nor Mr. Gregory contributed any cash and no value was placed on their respective contributions.

Other than as set forth above, there are no transactions since our inception, or proposed transactions, to which we were or are to be a party, in which any of the following persons had or is to have a direct or indirect material interest:

(a) Any director or executive officer of the Company;

(b) Any majority security holder; and

(c) Any member of the immediate family (including spouse, parents, children, siblings, and in-laws) of any of the persons in the above.

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Item 14. Principal Accountant Fees and Services.

Summary of Principal Accountant Fees for Professional Services Rendered

The following table presents the aggregate fees for professional audit services and other services rendered by Berman & Co., P.A., our independent registered public accountant in 2011 and 2010, respectively.

	Fiscal Year Ended December 31, 2011	Fiscal Year Ended December 31, 2010
Audit and Audit Related Fees	$ 211,328	$ 110,000
Tax Fees	$ 0	$ 0
All Other Fees	$ 0	$ 0

PART IV

Item 15. Exhibits, Financial Statement Schedules.

Exhibit No.	Description
2.1	Share Exchange Agreement, dated February 1, 2010, by and between Tone in Twenty, Inc. and Muscle Pharm LLC (as filed as Exhibit 2.1 on Form 8-K on February 2, 2010)
3.1	Tone In Twenty Articles of Incorporation, dated August 4, 2006 (as filed as Exhibit 3.1 to Company's Form SB-2 Registration Statement, filed November 2, 2007)
3.2	Bylaws of MusclePharm Corporation, dated August 5, 2006 (as filed as Exhibit 3.2 to Company's Form SB-2 Registration Statement, filed November 2, 2007)
3.3	Amendment to the Articles of Incorporation, dated February 23, 2007 (as filed as Exhibit 3.3 to Company's Form SB-2 Registration Statement, filed November 2, 2007)
3.4	Series A Certificate of Designation (as filed as Exhibit 3.4 to the Company's Current Report on Form 8-K, filed on February 24, 2010)
3.5	Amendment to the Articles of Incorporation (as filed as Exhibit 3.1 to the Company's Quarterly Report on Form 10-Q, filed on May 23, 2011)
3.6	Series B Certificate of Designation (as filed as Exhibit 3.1 to the Company's Quarterly Report on Form 10-Q, filed on August 16, 2011)
3.7	Series C Certificate of Designation, filed with the Secretary of State of the State of Nevada on October 25, 2011 (as filed as Exhibit 3.1 to the Company's Current Report on Form 8-K, filed on November 4, 2011)
3.8	Amendment to the Articles of Incorporation, dated November 17, 2011 (as filed as Exhibit 3.1 to the Company's Current Report on Form 8-K, filed on November 23, 2011)
3.9	Amendment to the Articles of Incorporation, dated January 18, 2012 (as filed as Exhibit 3.1 to the Company's Current Report on Form 8-K, filed on January 27, 2012)
3.10	Amendment to the Articles of Incorporation, dated March 26, 2012 (as filed as Exhibit 3.1 to the Company's Current Report on Form 8-K, filed on March 29, 2012)
4.1	$900,000 Convertible Promissory Note, dated June 7, 2011, issued in favor of JMJ Financial (as filed as Exhibit 4.1 to the Company's Current Report on Form 8-K, filed on July 8, 2011)
4.2	

	$2,651,000 Senior Secured Convertible Promissory Note, dated June 29, 2011, issued in favor of Inter-Mountain Capital Corp. (as filed as Exhibit 4.1 to the Company's Quarterly Report on Form 10-Q, filed on November 14, 2011)
4.3	Common Stock Purchase Warrant, dated June 29, 2011, issued in favor of Inter-Mountain Capital Corp. (as filed as Exhibit 4.2 to the Company's Quarterly Report on Form 10-Q, filed on November 14, 2011)
4.4	Form of Promissory Note (as filed as Exhibit 4.1 to the Company's Current Report on Form 8-K, filed on December 12, 2011)
4.5	Form of Common Stock Purchase Warrant (as filed as Exhibit 4.2 to the Company's Current Report on Form 8-K, filed on December 12, 2011)
10.1	Sponsorship Agreement, dated January 18, 2011, by and between MusclePharm Corporation, and The Cincinnati Reds LLC (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on January 24, 2011)

10.2	Registration Rights Agreement, dated June 7, 2011, by and between the Company and JMJ Financial (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on July 8, 2011)
10.3	Stock Purchase Agreement, dated July 7, 2011, by and between MusclePharm Corporation and Carriage Group, LLC (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on July 19, 2011)
10.4	Endorsement Agreement, dated July 20, 2011, by and between MusclePharm Corporation and Michael Vick, individually (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on July 22, 2011)
10.5	Note and Warrant Purchase Agreement, dated June 29, 2011, by and between MusclePharm Corporation and Inter-Mountain Capital Corp. (as filed as Exhibit 10.1 to the Company's Quarterly Report on Form 10-Q, filed on November 14, 2011)
10.6	Employment Agreement, dated September 16, 2011, by and between MusclePharm Corporation and John H. Bluher, individually (as filed as Exhibit 10.4 to the Company's Quarterly Report on Form 10-Q, filed on November 14, 2011)
10.7	Employment Agreement, dated November 14, 2011, by and between MusclePharm Corporation and Jeremy DeLuca, individually (as filed as Exhibit 10.5 to the Company's Quarterly Report on Form 10-Q, filed on November 14, 2011)
10.8	Amended and Restated Employment Agreement, dated November 14, 2011, by and between MusclePharm Corporation and Brad Pyatt, individually (as filed as Exhibit 10.8 to the Company's Annual Report on Form 10-K, filed on April 16, 2012)
10.9	Amended and Restated Employment Agreement, dated November 14, 2011, by and between MusclePharm Corporation and Cory Gregory, individually (as filed as Exhibit 10.9 to the Company's Annual Report on Form 10-K, filed on April 16, 2012)
10.10	Stock Purchase Agreement, dated December 2, 2011, by and between MusclePharm Corporation and TSX Holdings, LLC (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on December 8, 2011)
10.11	Amendment No. 1 to Stock Purchase Agreement, dated December 8, 2011, by and between MusclePharm Corporation and Carriage Group, LLC (as filed as Exhibit 10.2 to the Company's Current Report on Form 8-K, filed on December 9, 2011)
10.12	Form of Subscription Agreement (as filed as Exhibit 10.1 to the Company's Current Report on Form 8-K, filed on December 12, 2011)
10.13	Equity Purchase Agreement, dated November 4, 2011, by and between MusclePharm Corporation and Southridge Partners II, LP (as filed as Exhibit 10.8 to the Company's Registration Statement on Form S-1/A, filed on December 29, 2011)
10.14	Registration Rights Agreement, dated November 4, 2011, by and between MusclePharm Corporation and Southridge Partners II, LP (as filed as Exhibit 10.9 to the Company's Registration Statement on Form S-1/A, filed on December 29, 2011)

31.1	Certification by the Principal Executive Officer of Registrant pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 (Rule 13a-14(a) or Rule 15d-14(a)) *
31.2	Certification by the Principal Financial Officer of Registrant pursuant to Section 302 of the Sarbanes-Oxley Act of 2002 (Rule 13a-14(a) or Rule 15d-14(a)) *

32.1	Certification by the Principal Executive Officer pursuant to 18 U.S.C. 1350 as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 *
32.2	Certification by the Principal Financial Officer pursuant to 18 U.S.C. 1350 as adopted pursuant to Section 906 of the Sarbanes-Oxley Act of 2002 *

*Filed herewith

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SIGNATURES

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

MUSCLEPHARM CORPORATION

Dated: June 29, 2012 By: */s/ Brad J. Pyatt*
Brad J. Pyatt
Chief Executive Officer

Principal Executive Officer

Dated: June 29, 2012 By: */s/ Lawrence S. Meer*
Lawrence S. Meer
Chief Financial Officer

Principal Financial Officer

Principal Accounting Officer

PURSUANT TO THE REQUIREMENTS OF THE SECURITIES EXCHANGE ACT OF 1934, AS AMENDED, THIS REPORT HAS BEEN SIGNED BY OR ON BEHALF OF THE FOLLOWING PERSONS IN THE CAPACITIES AND ON THE DATES INDICATED:

Signature	**Title**	**Date**
/s/ Brad J. Pyatt Brad J. Pyatt	Chairman, Chief Executive Officer, Principal Executive Officer	June 29, 2012
/s/ Cory Gregory Cory Gregory	Senior President, Director	June 29, 2012
/s/ Lawrence S. Meer Lawrence S. Meer	Chief Financial Officer, Principal Financial Officer, Principal Accounting Officer	June 29, 2012

/s/ Jeremy DeLuca Jeremy DeLuca	President and Chief Marketing Officer	June 29, 2012
/s/ John H. Bluher John H. Bluher	Chief Operating Officer	June 29, 2012

MusclePharm Corporation

Consolidated Financial Statements

December 31, 2011 (Restated) and 2010 (Restated)

CONTENTS

REPORT OF INDEPENDENT REGISTERED PUBLIC ACCOUNTING FIRM

To the Board of Directors and Stockholders of:

MusclePharm Corporation

We have audited the accompanying consolidated balance sheets of MusclePharm Corporation and Subsidiary as of December 31, 2011 and 2010, and the related consolidated statements of operations, stockholders' deficit and cash flows for the years then ended. These financial statements are the responsibility of the Company's management. Our responsibility is to express an opinion on these financial statements based on our audits.

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

In our opinion, the financial statements referred to above present fairly, in all material respects, the financial position of MusclePharm Corporation and Subsidiary as of December 31, 2011 and 2010, and the results of its operations and its cash flows for the years then ended, in conformity with accounting principles generally accepted in the United States of America.

The accompanying financial statements have been prepared assuming that the Company will continue as a going concern. As discussed in Note 2 to the financial statements, the Company has a net loss of $23,280,950 and net cash used in operations of $5,801,761 for the year ended December 31, 2011; and has a working capital deficit of $13,693,267, and a stockholders' deficit of $12,971,212 at December 31, 2011. These factors raise substantial doubt about the Company's ability to continue as a going concern. Management's plan in regards to these matters is also described in Note 2.

Berman & Company, P.A.

Boca Raton, Florida

April 13, 2012 except for note 1 as to which the date is June 28, 2012

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Registered with the PCAOB Ÿ *Member* AICPA *Center For Audit Quality*

Member American Institute of Certified Public Accountants

Member Florida Institute of Certified Public Accountants

MusclePharm Corporation and Subsidiary

Consolidated Balance Sheets

	December 31, 2011	December 31, 2010
Assets		
Current Assets		
Cash	$659,764	$43,704
Accounts receivable - net	2,569,092	426,761
Prepaid stock compensation	534,456	1,965,911
Prepaid sponsorship fees	203,333	-
Other	50,188	58,065
Total Current Assets	4,016,833	2,494,441
Property and equipment - net	907,522	138,551
Debt issue costs - net	68,188	34,404
Other assets	53,585	53,585
Total Assets	$5,046,128	$2,720,981
Liabilities and Stockholders' Deficit		
Current Liabilities:		
Accounts payable and accrued liabilities	$9,359,073	$3,227,483
Customer deposits	8,047	75,733
Debt - net	1,281,742	289,488
Derivative liabilities	7,061,238	622,944
Total Current Liabilities	17,710,100	4,215,648
Long Term Liabilities:		
Debt - net	307,240	250,000
Total Liabilities	18,017,340	4,465,648
Stockholders' Deficit		
Series A, Convertible Preferred Stock, $0.001 par value; 5,000,000 shares authorized, none issued and outstanding	-	-
Series B, Preferred Stock, $0.001 par value; 51 shares authorized, 51 and none, respectively, issued and outstanding	-	-
Series C, Convertible Preferred Stock, $0.001 par value; 500 shares authorized, 190 and none, respectively, issued and outstanding	-	-

Common Stock, $0.001 par value; 2,500,000,000 shares authorized,605,930,613 and 118,649,439 issued and outstanding	605,931	118,649
Additional paid-in capital	31,579,538	20,012,122
Accumulated deficit	(45,156,681)	(21,875,438)
Total Stockholders' Deficit	(12,971,212)	(1,744,667)
Total Liabilities and Stockholders' Deficit	$5,046,128	$2,720,981

See accompanying notes to consolidated financial statements

MusclePharm Corporation and Subsidiary

Consolidated Statements of Operations

	For The Year Ended December 31,	
	2011 As Restated	2010 As Restated
Sales - net	$ 17,212,636	$ 3,202,687
Cost of sales	14,845,069	2,804,274
Gross profit	2,367,567	398,413
General and administrative expenses	18,587,727	18,650,249
Loss from operations	(16,220,160)	(18,251,836)
Other income (expense)		
Derivative expense	(4,777,654)	(93,638)
Change in fair value of derivative liabilities	5,162,100	(149,306)
Loss on settlement of accounts payable and debt	(3,862,458)	(433,400)
Interest expense	(3,711,278)	(480,589)
Other expense	(121,500)	(160,568)
Licensing income	250,000	-
Total other income (expense) - net	(7,060,790)	(1,317,501)
Net loss	$ (23,280,950)	$ (19,569,337)
Net loss available to common stockholders		
Net loss	$ (23,280,950	$ (19,569,337)
Series C preferred stock dividend	(293)	-
Net loss available to common stockholders	$ (23,280,657)	$ (19,569,337)
Net loss per share available to common stockholders - basic and diluted	$ (0.08)	$ (0.48)
Weighted average number of common shares outstanding during the year – basic and diluted	281,484,658	41,141,549

See accompanying notes to consolidated financial statements

MusclePharm Corporation and Subsidiary

Consolidated Statement of Stockholders' Deficit

Years Ended December 31, 2011 and 2010

	Series A, Convertible Preferred Stock		Series B, Preferred Stock		Series C, Convertible Preferred Stock		Common Stock		Additional Paid-	Accumulated	Total Stockholde
	Shares	Amount	Shares	Amount	Shares	Amount	Shares	Amount	in Capital	Deficit	Deficit
Balance - December 31, 2009	-	$-	-	$-	-	$-	26,000,000	$26,000	$1,099,508	$(2,306,101)	$(1,180,59
Recapitalization and deemed issuance	83,333	83	-	-	-	-	70,838	71	(25,261)	-	(25,107
Issuance of common stock:											
Conversion of preferred stock to common stock	(83,333)	(83)	-	-	-	-	16,666,600	16,667	(16,584)	-	-
Conversion of convertible debt to common stock	-	-	-	-	-	-	7,708,906	7,709	1,025,791	-	1,033,500
Stock and warrants	-	-	-	-	-	-	4,167,767	4,168	1,524,508	-	1,528,676
Services - third parties	-	-	-	-	-	-	22,457,214	22,457	4,532,158	-	4,554,615
Services - third parties - future services	-	-	-	-	-	-	10,545,200	10,545	2,724,003	-	2,734,548
Services - related parties	-	-	-	-	-	-	10,000,000	10,000	5,290,000	-	5,300,000
Services paid with previously issued stock to related parties	-	-	-	-	-	-	-	-	1,039,500	-	1,039,500
Settlement of debt - third parties	-	-	-	-	-	-	4,165,571	4,166	1,186,898	-	1,191,064
Settlement of debt - related party	-	-	-	-	-	-	7,161,548	7,161	350,916	-	358,077
Settlement of accounts payable	-	-	-	-	-	-	9,014,286	9,014	424,386	-	433,400
	-	-	-	-	-	-	50,000	50	30,450	-	30,500

Debt offering - additional interest expense											
Extension of debt maturity date	-	-	-	-	-	-	130,000	130	95,370	-	95,500
Contract settlement in connection with lawsuit	-	-	-	-	-	-	511,509	511	99,489	-	100,000
Share based payments	-	-	-	-	-	-	-	-	630,990	-	630,990
Net loss	-	-	-	-	-	-	-	-	-	(19,569,337)	(19,569,3
Balance - December 31, 2010	-	-	-	-	-	-	118,649,439	118,649	20,012,122	(21,875,438)	(1,744,66
Issuance of common and preferred stock:											
Conversion of convertible debt	-	-	-	-	-	-	254,061,743	254,062	4,014,795	-	4,268,857
Conversion of secured/unsecured debt	-	-	-	-	-	-	40,277,378	40,277	817,675		857,952
Cash	-	-	-	-	-	-	82,000,000	82,000	793,000	-	875,000
Cash	-	-	-	-	100	-	-	-	100,000	-	100,000
Services - third parties	-	-	-	-	-	-	46,521,157	46,522	1,153,322	-	1,199,844
Services - third parties	-	-	-	-	90	-	-	-	90,000	-	90,000
Services - third parties - future services	-	-	-	-	-	-	4,000,000	4,000	210,250	-	214,250
Extension of debt maturity date	-	-	-	-	-	-	9,375,000	9,375	151,875	-	161,250
Settlement of accounts payable	-	-	-	-	-	-	54,545,896	54,546	3,592,173	-	3,646,719
Cancellation of shares	-	-	-	-	-	-	(3,500,000)	(3,500)	3,500	-	-
Share based payments - related parties	-	-	51	-	-	-	-	-	-	-	-
Dividends on series C preferred stock - related parties	-	-	-	-	-	-	-	-	-	(293)	(293
Reclassification of derivative liability to additional paid in capital	-	-	-	-	-	-	-	-	640,826		640,826

Net loss	-	-	-	-	-	-	-	-	-	(23,280,950)	(23,280,9
Balance - December 31, 2011	-	$-	51	$-	190	$-	605,930,613	$605,931	$31,579,538	$(45,156,681)	$(12,971,2

See accompanying notes to consolidated financial statements

MusclePharm Corporation and Subsidiary

Consolidated Statements of Cash Flows

	Year Ended December 31, 2011	December 31, 2010
Cash Flows From Operating Activities:		
Net loss	$(23,280,950)	$(19,569,337)
Adjustments to reconcile net loss to net cash used in operating activities:		
Depreciation	171,587	18,567
Bad debt	120,477	119,468
Warrants issued for services - third parties	1,989,982	-
Stock issued for services - third parties	1,289,844	4,554,615
Stock issued for services - related parties	-	5,300,000
Services paid with previously issued stock to related parties	-	1,039,500
Stock issued to extend maturity date of debt	161,250	95,500
Stock issued as settlement in connection with lawsuit	-	100,000
Stock issued with unsecured debt offering-additional interest expense	-	30,500
Share based payments	-	630,990
Amortization of prepaid stock compensation	1,745,705	768,637
Amortization of debt discount and debt issue costs	3,466,718	485,689
Loss on settlement of accounts payable	2,123,129	433,400
Loss on conversion of debt	1,739,329	-
Derivative expense	4,777,654	93,638
Change in fair value of derivative liabilities	(5,162,100)	149,306
Changes in operating assets and liabilities:		
(Increase) decrease in:		
Accounts receivable	(2,262,808)	(434,753)
Prepaid sponsorship fees	(203,333)	-
Inventory	-	4,245
Deposits	-	32,116
Other	7,877	(66,703)
Increase (decrease) in:		
Accounts payable and accrued liabilities	7,581,564	2,358,430
Customer deposits	(67,686)	60,715
Net Cash Used In Operating Activities	(5,801,761)	(3,795,477)
Cash Flows From Investing Activities:		
Purchase of property and equipment	(831,511)	(117,303)
Net Used In Investing Activities	(831,511)	(117,303)
Cash Flows From Financing Activities:		
Cash overdraft	-	(17,841)
Due to related party	-	(27,929)
Proceeds from issuance of debt	6,612,900	2,140,608

Proceeds from issuance of debt - related party		358,077
Repayment of debt	(75,285)	-
Cash paid for debt issue costs	(263,283)	-
Proceeds from issuance of preferred stock	100,000	-
Proceeds from issuance of common stock and warrants-net of recapitalization payment	875,000	1,503,569
Net Cash Provided By Financing Activities	7,249,332	3,956,484
Net increase in cash	616,060	43,704
Cash at beginning of year	43,704	-
Cash at end of year	$659,764	$43,704
Supplemental disclosures of cash flow information:		
Cash paid for interest	$28,806	$15,882
Supplemental disclosure of non-cash investing and financing activities:		
Stock issued for future services - third parties	$214,250	$2,734,548
Non cash increase in accounts payable related to future services to be paid for with common stock	$100,000	$-
Debt discount recorded on convertible and unsecured debt accounted for as a derivative liability	$5,473,291	$380,000
Conversion of convertible debt and accrued interest for common stock	$3,387,480	$1,033,500
Stock issued to settle debt - third parties	$-	$1,191,064
Stock issued to settle debt - related party	$-	$358,077
Stock issued to settle accounts payable and due to factor	$1,440,779	$433,400
Reclassification of derivative liability to additional paid in capital	$640,826	$-
Conversion of preferred stock to common stock	$-	$83
Stock issued to acquire equipment	$82,811	$-
Auto acquired through financing	$26,236	$-
Dividends on series C preferred stock - related parties	$293	$-
Original issue discount	$-	$37,500

See accompanying notes to consolidated financial statements

MusclePharm Corporation and Subsidiary

Consolidated Notes to Financial Statements

December 31, 2011(Restated) and 2010 (Restated)

Note 1 Nature of Operations and Summary of Significant Accounting Policies (Restated)

Nature of Operations

MusclePharm Corporation (the "Company", "We", "Our" or "MP"), was organized as a limited liability company in the State of Colorado on April 22, 2008. On February 18, 2010, the Company executed a reverse recapitalization with Tone in Twenty, Inc. and changed its name to MP (See Note 3).

The Company markets branded sports nutrition products.

Restatement

On May 14, 2012, the Company determined that a material misstatement exists in the Company's 2011 quarterly and 2011 and 2010 annual financial statements. The Company concluded that the following financial statements contained material misstatements: (i) the Company's audited financial statements for the year ended December 31, 2011, filed in an annual report on Form 10-K with the U.S. Securities and Exchange Commission (the "SEC") on April 16, 2012; (ii) the Company's audited financial statements for the year ended December 31, 2010, filed in an annual report on Form 10-K with the SEC on April 1, 2011; (iii) the Company's unaudited financial statements for the period ended September 30, 2011, filed in a quarterly report on Form 10-Q with the SEC on November 14, 2011; (iv) the Company's unaudited financial statements for the period ended June 30, 2011, filed in a quarterly report on Form 10-Q with the SEC on August 16, 2011; and (v) the Company's unaudited financial statements for the period ended March 31, 2011, filed in a quarterly report on Form 10-Q with the SEC on May 23, 2011.

The foregoing financial statements contained material misstatements pertaining to the Company's calculation of net sales and presentation of general and administrative expenses and cost of sales. The Company has determined that advertising related credits that were granted to customers fell within the guidance of ASC No. 605-50-55 (" *Revenue Recognition" – Customer Payments and Incentives – Implementation Guidance and Illustrations)* . The guidance indicates that, absent evidence of benefit to the vendor, appropriate treatment requires netting these types of payments

against revenues and not expensing as advertising expense. The Company also noted other credits and discounts that, upon further review, had been previously classified as advertising expense as a component of general and administrative expense that require a reallocation of presentation as amounts to be netted against revenues. The Company's net loss and loss per share will not be affected by this reallocation in the statement of operations.

Promotions, credits and non-specific advertising with its customers have been reclassified from general and administrative expenses to revenues.

Samples shipped to customers not clearly identifiable were reclassified from general and administrative expense to cost of sales.

	Year Ended December 31, 2011 As Restated	Adjustments	Year Ended December 31, 2011 As Issued	Year Ended December 31, 2010 As Restated	Adjustments	Year Ended December 31, 2010 As Issued
Sales - net	$ 17,212,636	$ (3,625,701)	$ 20,838,337	$ 3,202,687	$ (844,608)	$ 4,047,295
Cost of sales	14,845,069	374,455	14,470,614	2,804,274	-	2,804,274
Gross profit	2,367,567	(4,000,156)	6,367,723	398,413	(844,608)	1,243,021
General and administrative expenses	18,587,727	(4,000,156)	22,587,883	18,650,249	(844,608)	19,494,857
Loss from operations	(16,220,160)	-	(16,220,160)	(18,251,836)	-	(18,251,836)
Other income (expense)						
Derivative expense	(4,777,654)	-	(4,777,654)	(93,638)	-	(93,638)
Change in fair value of derivative liabilities	5,162,100	-	5,162,100	(149,306)	-	(149,306)
Loss on settlement of accounts payable and debt	(3,862,458)	-	(3,862,458)	(433,400)	-	(433,400)
Interest expense	(3,711,278)	-	(3,711,278)	(480,589)	-	(480,589)
Other expense	(121,500)	-	(121,500)	(160,568)	-	(160,568)
Licensing income	250,000	-	250,000	-	-	-
Total other income (expense) - net	(7,060,790)	-	(7,060,790)	(1,317,501)	-	(1,317,501)
Net loss	$ (23,280,950)	$ -	$ (23,280,950)	$ (19,569,337)	$ -	$ (19,569,337)
Net loss available to common stockholders						
Net loss	$ (23,280,950)	$ -	$ (23,280,950)	$ (19,569,337)	$ -	$ (19,569,337)
	(293)	-	(293)	-	-	-

Series C preferred stock dividend						
Net loss available to common stockholders	$ (23,280,657)	$ -	$ (23,280,657)	$ (19,569,337)	$ -	$ (19,569,337)
Net loss per share available to common stockholders - basic and diluted	$ (0.08)	$ -	$ (0.08)	$ (0.48)	$ -	$ (0.48)
Weighted average number of common shares outstanding during the year – basic and diluted	281,484,658	-	281,484,658	41,141,549	-	41,141,549

Risks and Uncertainties

The Company operates in an industry that is subject to rapid change and intense competition. The Company's operations will be subject to significant risk and uncertainties including financial, operational, technological, regulatory and other risks, including the potential risk of business failure.

Use of Estimates

The preparation of financial statements in conformity with U.S. generally accepted accounting principles requires management to make estimates and assumptions that affect the reported amounts of assets and liabilities and disclosure of contingent assets and liabilities at the date of the financial statements and the reported amounts of revenues and expenses during the reporting period.

Making estimates requires management to exercise significant judgment. It is at least reasonably possible that the estimate of the effect of a condition, situation or set of circumstances that existed at the date of the financial statements, which management considered in formulating its estimate could change in the near term due to one or more future non-conforming events. Accordingly, the actual results could differ significantly from estimates.

Principles of Consolidation

All inter-company accounts and transactions have been eliminated in consolidation.

Cash and Cash Equivalents

The Company considers all highly liquid instruments purchased with an original maturity of three months or less and money market accounts to be cash equivalents. At December 31, 2011 and 2010, the Company had no cash equivalents.

The Company minimizes its credit risk associated with cash by periodically evaluating the credit quality of its primary financial institution. The balance at times may exceed federally insured limits. At December 31, 2011 there was one account that had a balance that exceeded the federally insured limit by approximately $378,000. In 2010, there were no balances that exceeded the federally insured limit.

Accounts Receivable and Allowance for Doubtful Accounts

Accounts receivable represent trade obligations from customers that are subject to normal trade collection terms. The Company periodically evaluates the collectability of its accounts receivable and considers the need to establish an allowance for doubtful accounts based upon historical collection experience and specific customer information. Accordingly, the actual amounts could vary from the recorded allowances.

The Company does not charge interest on past due receivables. Receivables are determined to be past due based on the payment terms of the original invoices.

MusclePharm Corporation and Subsidiary

Consolidated Notes to Financial Statements

December 31, 2011(Restated) and 2010 (Restated)

Accounts receivable at December 31, 2011 and 2010 were as follows:

Accounts receivable	$2,766,776	$542,863
Less: allowance for doubtful accounts	(197,684)	(116,102)
Accounts receivable – net	$2,569,092	$426,761

As of December 31, 2011 and 2010, the Company had the following concentrations of accounts receivable with customers:

Customer	2011	2010
A	36 %	24 %
B	12 %	2 %
C	10 %	- %
D	7 %	40 %
E	5 %	11 %

Property and Equipment

Property and equipment are stated at cost and depreciated to their estimated residual value over their estimated useful lives. When assets are retired or otherwise disposed of, the assets and related accumulated depreciation are relieved from the accounts and the resulting gains or losses are included in operating income in the statements of operations. Repairs and maintenance costs are expensed as incurred. Depreciation is provided using the straight-line method for all property and equipment.

Website Development Costs

Costs incurred in the planning stage of a website are expensed, while costs incurred in the development stage are capitalized and amortized over the estimated useful life of the asset.

Long-Lived Assets

The Company reviews long-lived assets for impairment whenever events or changes in circumstances, such as service discontinuance or technological obsolescence, indicate that the carrying amount of the long-lived asset may not be recoverable. When such events occur, the Company compares the carrying amount of the asset to the undiscounted expected future cash flows related to the asset. If the comparison indicates that impairment is present, the amount of impairment is calculated as the difference between the excess of the carrying amount over the fair value of the asset. If a readily determinable market price does not exist, fair value is estimated using discounted expected cash flows attributable to the asset.

Fair Value of Financial Instruments

The Company measures assets and liabilities at fair value based on an expected exit price as defined by the authoritative guidance on fair value measurements, which represents the amount that would be received on the sale of an asset or paid to transfer a liability, as the case may be, in an orderly transaction between market participants. As such, fair value may be based on assumptions that market participants would use in pricing an asset or liability. The authoritative guidance on fair value measurements establishes a consistent framework for measuring fair value on either a recurring or nonrecurring basis whereby inputs, used in valuation techniques, are assigned a hierarchical level.

The following are the hierarchical levels of inputs to measure fair value:

- Level 1: Observable inputs that reflect quoted prices (unadjusted) for identical assets or liabilities in active markets.

- Level 2: Inputs reflect quoted prices for identical assets or liabilities in markets that are not active; quoted prices for similar assets or liabilities in active markets; inputs other than quoted prices that are observable for the assets or liabilities; or inputs that are derived principally from or corroborated by observable market data by correlation or other means.

- Level 3: Unobservable inputs reflecting the Company’s assumptions incorporated in valuation techniques used to determine fair value. These assumptions are required to be consistent with market participant assumptions that are reasonably available.

MusclePharm Corporation and Subsidiary

Consolidated Notes to Financial Statements

December 31, 2011(Restated) and 2010 (Restated)

The following are the major categories of liabilities measured at fair value on a recurring basis as of December 31, 2011 and 2010, using quoted prices in active markets for identical liabilities (Level 1); significant other observable inputs (Level 2); and significant unobservable inputs (Level 3):

		2011	2010
Derivative liabilities	Level 2	$7,061,238	$622,944

The Company's financial instruments consisted primarily of accounts receivable, prepaids, accounts payable and accrued liabilities, derivative liabilities and debt. The carrying amounts of the Company's financial instruments generally approximated their fair values as of December 31, 2011 and 2010, respectively, due to the short-term nature of these instruments.

Revenue Recognition (Restated)

The Company records revenue when all of the following have occurred: (1) persuasive evidence of an arrangement exists, (2) product has been shipped or delivered by the third party manufacturer, (3) the sales price to the customer is fixed or determinable, and (4) collectability is reasonably assured.

Depending on individual customer agreements, sales are recognized either upon shipment of products to customers or upon delivery. For one of our largest customers, which represent 14% of total revenue in 2011, revenue is recognized upon delivery.

The Company has determined that advertising related credits that were granted to customers fell within the guidance of ASC No. 605-50-55 (“ *Revenue Recognition” – Customer Payments and Incentives – Implementation Guidance and Illustrations)* . The guidance indicates that, absent evidence of benefit to the vendor, appropriate treatment requires netting these types of payments against revenues and not expensing as advertising expense.

The Company records store support, give aways, sales allowances and discounts as a direct reduction of sales. The Company recorded reductions to gross revenues totaling approximately $4,000,000 and $1,000,000 for the years ended December 31, 2011 and 2010, respectively.

The Company grants volume incentive rebates to certain customers based on contractually agreed percentages ranging from 2.5% - 5.5% as a percentage of sales once a certain threshold has been met. The credits are recorded as a direct reduction to sales. Included in the reductions to revenues above are volume incentive rebates. Total volume incentive rebates granted for the years ended December 31, 2011 and 2010 were approximately $500,000 and $0, respectively.

The Company has an informal 7-day right of return for products. There were nominal returns in 2011 and 2010.

During the years ended December 31, 2011 and 2010, the Company had the following concentrations of revenues with customers:

Customer	2011		2010	
A	41	%	45	%
B	14	%	7	%
C	-	%	15	%

The Company does not manufacture or physically hold any inventory. Inventory is held and distributed by the Company’s third party manufacturer.

Licensing Income and Royalty Revenue

On May 5, 2011, the Company granted an exclusive indefinite term license to a third party for $250,000. The licensee may market, manufacture, design and sell the Company’s existing apparel line. The licensee will pay the Company a 10% net royalty based on its net income at the end of each fiscal year. To date, no royalty revenue has been earned.

Cost of Sales (Restated)

Cost of sales represents costs directly related to the production and third party manufacturing of the Company’s products.

In 2011, cost of sales increased due to a reclassification from advertising expense in the amount of $374,454.

See discussion of restatement

MusclePharm Corporation and Subsidiary

Consolidated Notes to Financial Statements

December 31, 2011(Restated) and 2010 (Restated)

Shipping and Handling

Product sold is typically shipped directly to the customer from the manufacturer. Any freight billed to customers is offset against shipping costs and included in cost of sales.

Freight billed to customers for the years ended December 31, 2011 and 2010 was $309,690 and $71,983, respectively.

Advertising (Restated)

The Company expenses advertising costs when incurred.

Advertising for the years ended December 31, 2011 and 2010 are as follows:

	2011 As Issued	Adjustments	2011 Restated	2010 As Issued	Adjustments	2010 Restated
Advertising	$ 9,241,741	$ (4,000,156)	$ 5,241,585	$ 7,084,955	$ (844,608)	$ 6,240,347

See discussion of restatement

Income Taxes

Through February 18, 2010, the Company was taxed as a pass-through entity (LLC) under the Internal Revenue Code and was not subject to federal and state income taxes; ac