ARNET PHARMACEUTICAL CORPORATION

GENERAL INFORMATION

- Established in 1972
- Arnet is dedicated to researching, manufacturing and selling natural products in the USA & worldwide
- Arnet provides private labeling services as well as its own brands (for the international market)
- Located in Davie, Florida, Arnet occupies 85,000 square feet including:
  - Offices
  - Laboratories
  - Production
  - Warehouse
  - Pilot Plant
- Divisions: Domestic & International

OVERVIEW

VISION

We are committed to becoming the preferred manufacturer of natural supplements for our customers by delivering orders on time and according to their specifications.

MISSION

To promote wellness by producing high quality natural supplements with the finest ingredients.

VALUES

Quality: We achieve quality by doing things right the first time.
Integrity: We achieve integrity by being honest and truthful in our duties.
Customer Service: We strive to deliver on time and to always being available to fulfill our customer's request.

THE COMPANY

- cGMP Certified
  - Certification issued by the Florida Department of Agriculture
  - NSF
  - SGS cGMP Certification
  - Eurofins
  - KSA SFDA
- The Quality Control Department works hand in hand with our customers and vendors (VQP) to ensure and guarantee the local requirements of production and documentation where products are being marketed.
- IN-HOUSE laboratories
  - Our QA/QC team consists of 42 members:
    - Chemistry
    - Physical & Microbiology
- cGMP Compliant Pilot Plant
  - Blending
  - Compression
  - DRX Encapsulation
  - Liquid Encapsulation
  - Coating
  - Flavor Development
  - Powder

PACKAGING CAPABILITIES & SERVICES

- Bulk Packaging
- Bottle Packaging
- Packaging Solutions & Graphic Design Assistance
- Special Packaging Services (Blisters, packets, pouches, sachets)

PRODUCT DEVELOPMENT SERVICES

- Non-Disclosure & Material transfer agreements
- Formula & Flavor Development
- Branded & Private Label
- A variety of delivery forms (Liquid Capsules, Capsules, Tablets, & Powders)
- Extensive experience in developing unique custom formulation including:
  - Chewable products for adults & children
  - Flavored green powder
  - Meal replacement shakes
  - Daily energy powders
  - Enteric coating
  - Timed Release
  - Quick-Dissolve Tablet
- KM and OU Kosher and Halal Certified

THE LINE OFFERS

An attractive Product Portfolio that provides safe and simple alternatives for common ailments for the family.

- Single Vitamins
- Multi Vitamins
- Minerals
- Herbal & Food Supplements
- Joint & Bone Health
- Natural Stimulants
- Fitness-Sport & Beauty

ARNET EXPANDED WORLDWIDE (1996)

Arnet Contract Manufacturer Services are in 69 Markets

- **Middle East:**
  Bahrain, Egypt, Iran, Jordan, Kingdom of Saudi Arabia, Kuwait, Lebanon, Libya, Qatar, United Arab Emirates, Palestine, Syria, Yemen, Israel.
- **Far East:**
  Hong Kong, Myanmar, Malaysia, Philippines, Singapore, Taiwan, Vietnam, South Korea, Macau, Thailand, China, Indonesia, Cambodia, Japan.
- **Africa:**
  Nigeria, Tunisia, Morocco, Algeria, Ghana.
- **Europe:**
  Italy, Lithuania, Portugal, Russia, Spain, Turkey, Azerbaijan, Poland, Kazakhstan.
- **North America:**
  Canada, United States of America, Puerto Rico.
- **Caribbean:**
  Aruba, Barbados, Curacao, Haiti, Trinidad & Tobago, Antigua, Dominican Republic.
- **Latin America:**
  Bolivia, Brazil, Colombia, Chile, Costa Rica, Ecuador, El Salvador, Belice, Guyana, Suriname, Guatemala, Honduras, Mexico, Panama, Paraguay, Peru, Venezuela.

REGISTRATION DEPARTMENT

We assist our clients with the product registration processing in their country if required by the local health authorities and legalization or apostille of documents. One of our key strengths is our ability to customize product ingredients and labeling to comply with the requirements of the health departments of individual countries.

- We provide documentation such as:
  - GMP Certificate
  - FSC Certificate
  - Product's dossier including:
    - Quali-Quantitative Formula
    - Stability Studies
    - Product Specification
    - Certificate of Analysis for raw materials
    - Certificate of Analysis for finished products
    - Samples (raw materials and finished goods)
  - CTD Format dossiers

RECENT AUDITS

- FDA- June 2018: No remarks and no 483 Form Issued
- NSF GMP Certified - Approved member since 2009
- NHPD GMP Certification
- SGS GMP Certified March 2016
- Eurofins GMP Certification June 2018
- KSA SFDA September 2018

SUMMARY

- Compliance with FDA and cGMP standards
- Products Manufactured Following World Wide GMP Standards Beyond 21 CFR 111
- NSF, SGS, NHPD & KSA GMP Facility
- In-House Laboratory
- A state-of-the-art facility
- Product development
- Packaging solutions
- Documentation and registration support for export
- Strategically located in South Florida facilitating shipping effectiveness and efficiency superior service