Titenare GXL Technologies: Global Supply Chain Management

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Integrated Pharma Business, Development and Commercial Solutions:

Highest Quality at affordable Cost & Collaboration Connected

BUSINESS SOLUTIONS AND SERVICES

Access to right Products & Funding	*Access to integrated Services*	*Access to right Markets*
BUSINESS DEVELOPMENT ➢ New Product Strategies ...R&D, Inlicensing, Outlicensing ➢ Global Supply Chain management ➢ Mergers & Acquisitions ➢ Advisory services ➢ Turnkey project solutions ➢ Fund raising support	TECHNICAL SERVICES ➢ Facilities Design & Engineering ➢ Regulatory Affairs – Drug & Device ➢ Quality Management Systems ➢ Development Services – Product/Analytical ➢ API Development Monitoring Support ➢ Technology/Site Transfers ➢ Third Party Audit and Compliance ➢ ISO//CE Mark/UL certifications ➢ Global Supply Chain Management ➢ Intellectual Property	MARKETING & SALES ➢ Market Assessment ➢ Competitor Differentiation ➢ Pipeline Prioritization ➢ Competitor and customer perspectives ➢ Complete organization value maps. ➢ Implementation planning and support ➢ Market access: India, EU, US, RoW

Global Supply Chain Management

- Establish manufacturing strategies integrating current and future requirements
- Establish strategic partnerships:
  > Identify contract manufacturing partners
  > Establish sound techno-commercial agreements
  > Relationships Management
- Implement an integrated plan to manufacture various technologies, dosage forms and therapeutic categories.
- End-to-end supply chain management: from PO placement to delivery
- Fast launch planning
- Technical sourcing: Formulations, API, excipients, key intermediates, electronic items, molds, packing materials.
- Provide inputs on acquisitions, own set up, evaluation of various scenarios, etc.
- Risk assessment & mitigation options
- Bulk supplies with local packing strategies
- Establishment of warehouse/Inventory management system
- Establishment of end-to-end logistics support/management system
- End-to-end remediation project handling

CMO Management

- Manage identification, qualification, and quotes from multiple CMO's. approved by USFDA/EUGMP/1348/UL, including API and device components
- Recommend to Client, help select a CMO, and provide CMO management oversight
- Negotiate supply agreement and quality agreement on behalf of the Client
- Qualification of CMO from screening, facility tours, system audit and review of following:
  > Organization chart and structure
  > List of all current Standard Operating Procedure ("SOP") and policies
  > Experience of regulatory audits, audit findings and responses by CM
  > Training records with curriculum vitae and job responsibilities of investigators and other staff
- Create and facilitate estimation of Cost of Goods, commercial and technical agreements

FLEXIBLE ENGAGEMENT MODELS

- Specific project based
- Rent-a-resource: Monthly retainer based
- Resource placement at the manufacturing sites