

ANALYTICAL SERVICES & FORMULATION DEVELOPMENT

Bilcare provides innovative solutions for all of your processing needs. Our dedicated personnel provide technical support during each phase of your product's development and manufacturing. GMP qualified scientists oversee comprehensive development and testing of your materials. We take extra steps to ensure that your products comply with international regulatory standards.



CLINICAL MANUFACTURING, PACKAGING, AND LABELING

Once a drug has been formulated, Bilcare has a manufacturing facility that can provide fast, efficient and agile manufacturing, packaging and labeling of the test product. Our unique approach enables us to support customer requirements for short-term studies as well as large-volume ongoing projects.



GLOBAL LOGISTICS AND SUPPLY CHAIN MANAGEMENT

The logistics of dealing with effective storage, shipping and regulatory compliance can be a complex and complicated process. However, with multiple global facilities for distribution, Bilcare provides you with the unique capability of a real-time distribution process using a simple and effective local management model.



INTERACTIVE VOICE/WEB RESPONSE SYSTEM (IVRS) SOLUTIONS

Bilcare provides a state-of-the-art Interactive Voice and Web Response System (IVRS/IWRS) which is an innovative, value-based product for managing data, including automatic patient enrollment and randomization, capturing data, patient enrollment management and drug supply management as well as patient diaries.



IN-SOURCING: OFFSITE CLINICAL SUPPLIES OPERATIONS

Bilcare's major strength lies in its ability to fully integrate innovative solutions and services throughout a drug's clinical trial life cycle – Onsite or Offsite. We bring all of our global expertise and experience to you. If your clinical trial needs can not come to us, we can come to you to achieve the success that you are striving for in all of your clinical supplies needs.



QP SERVICES

Bilcare GCS offers QP Services in accordance with the EU requirement that all investigational, medicinal products manufactured or imported by organizations holding a MA IMP have available a Qualified Person to approve and release each batch of IMP for clinical trial use.

