BIO/PHARMA MANUFACTURING AND TESTING SERVICES

BASIC RESEARCH/ DISCOVERY	IND FILING /PRECLINICAL	PHASE I-II	PHASE III	FILING	APPROVAL	POST APPROVAL / MANUFACTURING

- Custom Synthesis

- GMP Manufacturing

- Chemistry, Manufacturing and Controls (CMC) Support

- Method Development
- API, Raw Materials, Excipients Testing and Characterization

- Method Validation, Verification and Method Transfer
- Stability Testing
- Release Testing
- Residual Impurity Testing
- Extractables and Leachables
- Environmental Monitoring, Facilities Qualification / Validation

element