We are able to provide contract QP(IMP) and RP services as required to clients.

We have in-depth knowledge of both GDP and GMP requirements for all aspects of the drug supply chain

We have many years experience operating within the GDP / GMP QA environment and can provide a very experienced and knowledgeable approach to all aspects of Quality Assurance (QA).

Whether you require writing of SOPs and Quality Management systems or review of master and post production batch records we are able to help.

We can also fulfil your routine QA requirements as well.

We take a pragmatic view to problem solving and can assist with all aspects of change control, deviation management and CAPA.

We can offer support and guidance in all aspects of dossier writing, compilation and submission, as well as providing any strategic advice.

We ensure that your products and suppliers comply with cGxP by undertaking the following services:

- Input, compilation and review of CMC / IMPD / IND and CTA documentation
- SOP review, training and implementation
- Protocol writing, review and implementation

- Full quality management system review and implementation
- Report compilation and review