

This Consultancy was established in 2001 to provide a service to the Pharmaceutical Industry that is highly professional, pays close attention to detail and is up to date with current legislation for registration of medicines.

It is founded on 30 years experience in Regulatory Affairs and Production Management.

All tasks are undertaken with unquestionable integrity, building quality into performance, and optimising time-to-completion.

The following list of Services is a guide. Whatever your requirements, please give us a call:

- ✚ Act 101 medicines - Human and Veterinary
- ✚ New application dossiers and resolutions
- ✚ Data screening
- ✚ Registered medicines' dossier updates
- ✚ MBR1 conversion to MRF1
- ✚ Package inserts
- ✚ Fluent in English & Afrikaans

It will be a pleasure to assist you.