Quality Plan.

- Viamed is certified to ISO 9001/2000 & ISO 13485/2003, and these standards are the basis of quality assurance & control throughout the company.
  Manufacturing procedures are in place for all goods manufactured by Viamed and can be found in:

1. ISO Quality manual.
2. ISO Procedures manual.
3. Paperport on network.
4. Word processing documents.
5. Repair procedures can be found as above.
6. Suppliers are continually monitored for quality of goods and delivery reliability.

- All products not manufactured by Viamed and not from a company with a proven 100% record for quality or specification compliance are either individually tested or batch tested.
- All parties currently involved have ISO9001 minimum, or have been historically approved by Viamed for product quality etc.
- All component parts, pre-manufactured by suppliers, will be Inspected and tested by that particular supplier.
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- Final complete package will have final QA by Viamed.

Quality procedures are used for:

- The Testing of the Microstim VM3COP36.01
- Packaging of final Microstim kit. As above