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 / or sub-contractors) and can be found in:

1. ISO Quality manual.

2. ISO Procedures manual.

3. Paperport on network.

4. Word processing documents.

5. Every Batch is first piece tested
   Every batch is finished product inspected to ANSI/ASQC Z1-1993 Single Sampling Plan for Normal Inspection

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.

- Final complete package will have final QA by Viamed.