



12.02.03

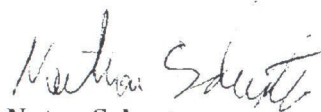
To whom it may concern

Subject: Quality System of Fermantek Ltd.

The Quality System of this company was upgraded and includes all requirements of GMP API complying with Standard Q7A, U.S. Department of Health and Human Services – FDA.
(Active Pharmaceutical Ingredients).

The quality manual and work instructions refer to all subjects in the above requirements.

In the last ISO audit the implementation of this requirement was reviewed, all production and QA activities that were sampled were performed according to those requirements.


Natan Schuster,

Lead Auditor

Cc: Jacob Jarosinski
Head of Quality Systems
Certification Directorate