Dissolution Rate Testing

In accordance with Ph. Eur. 2.9.3 and USP <711> harmonized in ICH Q4B Annex 7R2 as well as Ph. Eur. 2.9.42, Ph. Eur. 2.9.43 and USP <724>

provided by GBA Pharma

Your Experts for Pharmaceutical Analytics and Solutions
Our Services

- Great number of semiautomated DR testing systems according to USP I and USP II (basket and paddle)
- DR testing systems according to USP IV (flow-through cells)
- DR testing of highly potent substances in the safety lab
- Monitoring according to your needs by LC, UPLC, CE, GC as well as spectrophotometer
- Great expertise in method development and validation as well as batch control for all dosage forms

Further information about our service portfolio can be found at www.gba-pharma.de

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