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905 UNIFORMITY OF DOSAGE UNITS USP34 - US ...

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2.9.3. DISSOLUTION TEST FOR SOLID DOSAGE FORMS

EUROPEAN PHARMACOPOEIA 6.0 2.9.3. Dissolution test for solid dosage forms Assemble the apparatus, equilibrate the dissolution medium to 37 ± 0.5 °C, and remove the thermometer. The test may also be carried out with the thermometer in place, provided it is shown that results equivalent to those obtained without the thermometer are obtained.

2.9.31. PARTICLE SIZE ANALYSIS BY LASER LIGHT ...

2.9.31. Particle size analysis by laser light diffraction EUROPEAN PHARMACOPOEIA 6.0 particles in the light beam. Hence, the continuous angular intensity distribution is converted into a discrete spatial intensity distribution on a set of detector elements. It is assumed that the measured scattering pattern of the

RECOMMENDED METHODS OF ANALYSIS AND SAMPLING ...

European Pharmacopoeia 2.5.5 Tritrimetry (Colorimetric) I . CXS 234-1999 10 Fats and oils and related products Fish oils Phospholipids USP-FCC 12 2S (Krill oil – phospholipids), Nuclear Magnetic Resonance Spectroscopy I Fish oils P-Anisidine value European Pharmacopoeia 2.5.36/ AOCS Cd 18-90/ ...