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Guidance

- ANVISA: Resolution- RDC Nº 27, Requirements for the validation of bioanalytical methods.

- CDSCO: Guidelines for bioavailability and bioequivalence studies

- CFDA: Technical guideline for human bioavailability an bioequivalence studies on chemical drug products
  http://www.cde.org.cn/attachmentout.do?mothed=list&id=167

- EMA: Guideline on the investigation of Bioequivalence

- EMA: Guideline on the validation of bioanalytical methods

- EMA: Reflection Paper for Laboratories that perform the analysis or evaluation of clinical trial samples
FDA: Guidance for Industry: Bioanalytical Method Validation

FDA: Guidance for Industry - Safety Testing of Drug Metabolites

HPFB: Conduct and Analysis of Comparative BA Studies

ICH: E6(R1) Guideline for Good Clinical Practice

ICH: Q2(R1): Validation of Analytical Procedures: Text and Methodology

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MHLW: Draft Guideline on Bioanalytical Method Validation in Pharmaceutical Development

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