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[E6\(R2\) Good Clinical Practice: Integrated Addendum To ICH ...](#)

E6(R2) Good Clinical Practice: Integrated Addendum To ICH E6(R1) Guidance For Industry . U.S. Department Of Health And Human Services . Food And Drug Administration

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ICH Guideline Q7 On Good Manufacturing Practice For Active Pharmaceutical Ingredients â€” Questions And Answers EMA/CHMP/ICH/468930/2015 Page 2/37

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International Council For Harmonisation Of Technical Requirements For Pharmaceuticals For Human Use (ich) Ich Harmonised Guideline Integrated Addendum To Ich E6(r1): Guideline For Good Clinical Practice

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- 2 - Analysis Guideline For The RoHS Directive Ver. 4.0 1 Objective 1.1 Main Purpose Of Revision In Response To The European RoHS Directive (2002/95/EC) Published In January 2003

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