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standards and recommendations that apply to thousands of highly specific tasks processes and roles in the conduct of clinical research Interpreting and implementing GCP standards continue to represent challenges for pharmaceutical biotechnology and medical device industries This publication sets out to systematically collect catalog and answer the most important emerging and difficult questions regarding the interpretation and implementation of GCP standards today This text s question and answer format address a vast array of these questions some of which have persisted for many years without definitive answers and some of which have emerged in response to new clinical trials or regulatory related developments

~Aœ Guide to Good Clinical Practice Alex D. Kanarek,2005 **Guide to Good Clinical Practice** Mary Bernadette Ott,1999 **Guide to Good Clinical Practice** Mary Bernadette Ott,1993 **Good Clinical Practice** Munish Mehra,2007

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