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the asq certified pharmaceutical gmp professional handbook assists candidates preparing for the certified pharmaceutical good manufacturing practices professional cpmp examination and serves as a handy reference guide for practitioners in the field this handbook covers compliance with good manufacturing practices gmps as regulated and guided by national and international agencies for the pharmaceutical industry

the purpose of this handbook is to assist individuals for the certified pharmaceutical good manufacturing practices professional cpmp examination and provide a reference for the practitioner the second edition reflects the body of knowledge which was updated in 2015 this edition has also incorporated additional information including updated references the updates reflect the current trends and expectations of the evolving pharmaceutical industry driven by consumer expectations and regulatory oversight this handbook covers compliance with good manufacturing practices gmps as regulated and guided by national and international agencies for the pharmaceutical industry it covers finished human and veterinary drugs and biologics and combination devices as well as their component raw materials including active pharmaceutical ingredients apis and excipients and packaging and labeling operations

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no other area of regulatory compliance receives more attention and scrutiny by regulatory authorities than the regulation of sterile products for obvious reasons with the increasing number of potent products particularly the new line of small protein products joining the long list of proven sterile products the technology of manufacturing ster

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