Shun Jin is responsible for regional regulatory activities such as the registration and approval of new innovative/generic pharmaceutical products. In the drug regulatory affairs field in the pharmaceutical industry for over 25 years, his extensive experience covers new and generic drug applications, life cycle maintenance, regulatory intelligence, policy and GxP related regulatory activities.

Prior to joining Sandoz, he led Asian regulatory affairs teams at Abbott, Takeda, Parexel and GE Healthcare. Shun also served as regulatory policy head in AbbVie for Asia and Japan. Currently the chairman of DIA Singapore steering committee and regional editor of DIA *global forum*, Mr. Jin is a pharmacist by training with a business degree.