Alice has more than 29 years' experience in lifecycle management of pharma facilities inclusive of regulatory compliance, facility & process design, technology transfer, commissioning, qualification, and validation. Previously she worked for Sandoz/Novartis in Basel, Kundl and Ireland, before leading the Global Regulatory, Commissioning and Qualification Group for a Global Engineering Company for 13 years. Alice has global project experience of green field facilities including fill finish, biotech, gene therapy, medical devices, small and large molecule API and OSD from design through to operational readiness.

A recipient of ISPE's Robert F. Sherwood Article of the Year Award, she was a committee member on the Biotech and Disposables CoPs and Lead on the Knowledge Assets Committee. Alice currently serves on the ISPE C&Q CoP Steering Committee, is a former co-chair of this and RQHC/RSC committees and is on the ISPE Commissioning and Qualification Baseline® Guide Revision Team. She participated in the inaugural Women in Pharma forum at ISPE Annual Meeting 2016. Alice holds a PhD in Cell Culture from Dublin City University, an MBS in Project Management from University of Limerick and a Degree in Biotechnology from Dublin City University.