Dr. Junko Sato is an Office Director of Office of International Program at Pharmaceuticals and Medical Devices Agency (PMDA).
She started to work in regulatory agency in 1998. She became a review director in Office of New Drug, anti-infective drug area and moved to Office of Safety to introduce a new risk management system through life cycle of drugs. During the period, she worked in U.S. FDA as a guest reviewer from September 2002 to March 2003. From May 2012 to April 2014, she was dispatched to EMA as the MHLW/PMDA Liaison. She enhanced collaboration between EMA and MHLW/PMDA and brought a huge success as the liaison.
She contributes to some global harmonization activities, for example, ICH, CIOMS etc. Currently she is the chair of IPRP. She is also keen for scientific activities. She works for Japanese Society of Clinical Pharmacology and Therapeutics as a board member, and Japanese Society of Chemotherapy as a member of many committees, and more than 10 Societies.