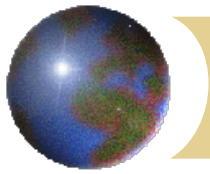


# Opening Remarks

**Tatsuya Kondo, MD, PhD**

**Chief Executive, PMDA**



# 2010 China-Japan Symposium on Global Clinical Trials and Ethnic Factors

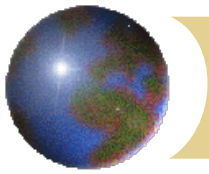
Organized by PMDA/JPMA/CCPIE  
Supported by RDPAC

**主办单位:**  **日本医药品医疗器械综合机构**  
Organizers: **Pharmaceuticals and Medical Devices Agency, Japan**

 **JPMA** **日本制药工业协会**  
**Japan Pharmaceutical Manufacturers Association**

 **中国医药国际交流中心**  
**China Center for Pharmaceutical International Exchange**

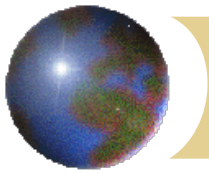
**协办单位:**  **中国外商投资企业协会药品研制和开发行业委员会**  
Supporter: **The China Association of Enterprises with Foreign Investment  
R&D-based Pharmaceutical Association Committee**



# The Purpose of this Symposium

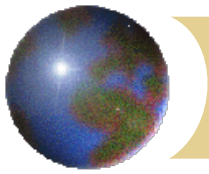
**Global drug development has accelerated in these days. Global Clinical Trials are rapidly increasing for simultaneous global development and application in East Asian countries.**

- 1) To promote and advance Global Clinical Trials in China/Japan.
- 2) To discuss the views of the regulators, reviewers, researchers, and the industry of the two countries on the issues of ethnic factors on MRCT
- 3) To realize speedy development and approval review using data in China/Japan.



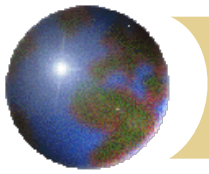
# Theme of the Symposium

- **How to promote Global Clinical Trials**
- **How to tackle ethnic factors in Global Clinical Trial**



## **Program Agenda**

- **Keynote Speech:**
- **Session 1: Government efforts to promote Global Clinical Trials and current situation of Pharmaceutical regulations**
- **Session 2: Current and future trends in clinical research on Ethnic Factors**
- **Session 3: Challenges on Clinical trials and approval reviews in consideration of Ethnic Factors**
  - **Reviewer's Viewpoints**
  - **Industry's Viewpoints**
  - **Panel Discussion**



**中日の豊富な経験と叡智を  
熱心な議論により交換しよう**

**Let's exchange experience and  
wisdom through earnest discussion**

