CONTENTS

Vol.44 No.4 July 2013

Original Article		
Lack of Pharmacokinetic Interaction between Pilsicainide and		
Rifampicin in Healthy Volunteers	Tsuyoshi SHIGA, et al.	301
Bioequivalence in Safety and Pharmacokinetics of Fentanyl Tape for 3 Days		
after a Single Application in Japanese Healthy Male Subjects Co-Administrated Naltrexone Hydrochloride	Ken SHIMADA, et al.	307
Survey on "Renal Function" Information in the Package Inserts of Prescription Drugs	Ai KANDA, et al.	313
2 a	,,, _,, _	
Case Report		
A Narrative Approach in Psychological Aspects of Four Participants in a		
Placebo-Controlled Double-Blind Trial of Cancer Chemotherapy: A Case Report	Takako FUJITANI, et al.	319
Proceedings of the 33rd Annual Meeting of the Japanese Society of Clinical		
Pharmacology and Therapeutics		
Contents		323
"How Can We Assess Efficacy and Safety of Approved Drugs?"		325
Symposium 5 "Efficient Supply of Post-Marketing Information of Drugs"		339
Symposium 11 "Pharmacokinetics Study on Patients"		349
Symposium 19 "Personalized Medicine Based on Phamaceutical Care"		359
Symposium 20 "What is Needed for Early Phase Clinical Studies in Japan to be International Studies in Japan to be Internat	onally Recognized?"	367
		207
Report		
Report of Trainee of the Foreign Clinical Pharmacology Training Program: No.4	Atsushi SAKURABA	379
Report of Trainee of the Foreign Clinical Pharmacology Training Program: No.1	Mihoko YAMADE	383
Report of CRC Trainee of the Foreign Clinical Pharmacology	Mitsuo KIMATA, et al.	385
Drug Information		121E
Information for Authors		125E
		1075
Announcements		127E

324 Proceedings

Contents of Proceedings of the 33rd Annual Meeting of the Japanese Society of Clinical Pharmacology and Therapeutics

"How Can We Assess Efficacy and Safety of Approved Drugs?"1. Clinical Epidemiology of Adverse Drug Events2. Evaluation of Safety and Efficacy of Medicine in the Registry after Coronary Revascularization	Takeshi MORIMOTO Masahiro NATSUAKI, et al.	325 329
3. Safety and Efficacy Evaluation of Cardiovascular Drugs after Approval	Reiko SATO	334
4. Registry Based Comparative Effectiveness Research	Shinichiro UEDA	335
Symposium 5 "Efficient Supply of Post-Marketing Information of Drugs"		
Summary	Masahiro NOMOTO, et al.	339
1. Outline of Risk Management Plan	Shinichi WATANABE	341
2. Supply and Utilization of Useful Post Marketing Information in Medical Field		
From a Physician's Viewpoint	Hideki MOCHIZUKI	343
3. Problem in Post-Marketing Supply of InformationMainly in Package Leaflet	Hiroaki ARAKI	345
4. Providing Proactive and Timely Information Resulting from Ongoing		010
Post-Marketing Studies to Medical Institutes	Yasuhiko KAI, et al.	347
Tost Markening Statios to Moureal Institutes		011
Symposium 11 "Pharmacokinetics Study on Patients"		
Summary	Yukiko MARUYAMA, et al.	349
1. The Implementation System in Ehime University Hospital for		
Pharmacokinetic Studies of Patients	Risako YAMASHITA, et al.	351
2. Problems in Performing Pharmacokinetic Examinations in Men with Pathological Conditio	ns Kazuhiro HARADA	353
3. PK/PD Study in Patients in Phase 1 Unit	Hinako UCHIMARU, et al.	355
4. Request to Japanese Medical Institution to Promote Clinical PK/PD Trial in Patients	Shingo MATSUI	357
Symposium 19 "Personalized Medicine Based on Phamaceutical Care"		
1. An Approach to Personalized Medicine	Hiroki ITO	359
2. Application of Gene Analysis and TDM in Personalized Drug Therapy	Tomonori NAKAMURA, et al.	361
3. Adverse Event Monitoring	Toshiaki NAKAMURA, et al.	363
4. Expectation for Clinical Pharmacology from Bed Side	Tohru HASHIDA	365
Symposium 20 "What is Needed for Early Phase Clinical Studies in Japan to be International	y Recognized?"	
Summary	Masako NAKANO, et al.	367
1. FIH Study: In Order to Conduct Successful Global Exploratory Studies	Kazuo UMEMURA	369
2. Proof of Concept Trials	Masahiro NOMOTO	371
3. Conducting Phase 1 Study in Patient Subjects: Points to Consider	Masanari SHIRAMOTO, et al.	373
4. FIH (First-in-Human) Trial of Oncology Drug Products:		
How to Enhance International Competitiveness of Japanese Academic Hospitals	Yasuhiro FUJIWARA	375
5. Promotion of Early Stage Clinical Trials: From the Viewpoint of MHLW	Masanobu YAMADA	377