Author Index Volume 9 (1996)

The issue number is given in front of the page numbers.

0924-6479/96/\$8.00 © 1996 - IOS Press. All rights reserved

International Conference on National Medicinal Drug Policies Statement of the International Working Group on transparency and accountability in drug	(1) 47- 59
regulation Editorial: Time to lift the veil of secrecy	(3) 211–217 (3) 131–132
Amschler, U., Drug safety regulations of the 'Drug Commission of the Counties'	(2) 91- 93
Bankowski, Z., The monitoring of drug safety and reporting of adverse drug reactions Bardelay, D., An ISDB survey to assess the degree of transparency of drug regulatory agencies Beckmann, J., Drug safety regulations and strategies of the German authority for medicinal	(2) 115–120 (3) 151–155
products Beppu, H., The case of contaminated blood products in Japan Bertelsmann, A., Statutory regulations governing the reporting of adverse reactions Bixler, E.O., see Kales, A.	(2) 75- 81 (3) 157-159 (2) 101-106 (1) 7- 27
Dinnendahl, V., Drug safety regulations of the 'Drug Commission of the German Pharmacists' Dukes, G., Editorial: Drug safety: can more be done? Dukes, M.N.G., Contaminated Human Growth hormone as a cause of Creutzfeldt–Jakob	(2) 87- 90 (2) 71- 73
disease Dukes, M.N.G., Drug regulation and the tradition of secrecy	(1) 41– 46 (3) 143–149
Helsing, E., Health promoting policies: strong, weak,, and sometimes harmful? Hemminki, E., see Ollila, E. Herxheimer, A., Side effects: freedom of information and the communication of doubt Hodgkin, C., International harmonisation – the need for transparency	(1) 1- 6 (3) 161-172 (3) 201-210 (3) 195-199
Jensen, L.P., see Levi, N.	(1) 29- 31
Kales, A., A.N. Vgontzas and E.O. Bixler, A reassessment of triazolam	(1) 7- 27
Lee, R., Drug safety regulations in the CPMP Levi, N., L.P. Jensen and T.V. Schroeder, Audit of the Danish national vascular database	(2) 107–113 (1) 29– 31
Medawar, C., Secrecy and medicines Munter, KH., Drug safety regulations of the 'Drug Commission of the German Physicians'	(3) 133–141 (2) 83– 86
Ollila, E. and E. Hemminki, Secrecy in drug regulation	(3) 161–172
Razak, D.A., Access to regulatory information in Malaysia - a preliminary study	(3) 179–186

Sammut, M. and C. Savona-Ventura, Petrol lead in a small island environment	(1) 33- 40
Savona-Ventura, C., see Sammut, M.	(1) 33- 40
Schroeder, T.V., see Levi, N.	(1) 29- 31
Tempelaar, A.F., Risk and safety in practice	(1) 65- 69
Tempelaar, A.F., Safety and risk in practice	(2) 121 - 129
Thiele, A., The German 'Graduated Plan' - criteria, administration, regulations	(2) 95–100
Vgontzas, A.N., see Kales, A.	(1) 7- 27
Vrhovać, B., Access to information on drug regulation in the countries of Central and Eastern	
Europe	(3) 173 - 178
Weerasuriya, K., Globalisation of drug regulation and drug regulators in developing countries	(3) 187–193