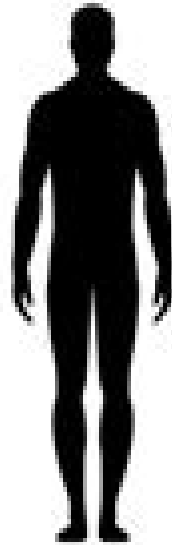
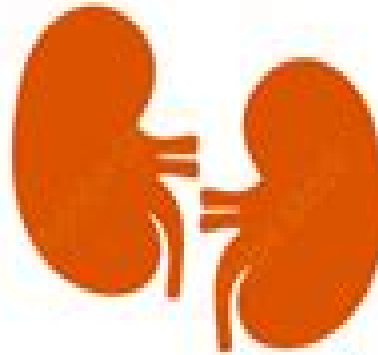


Patient - related



Kidney - related



Drug-related



Old age (≥ 60 -65 years)

Female sex

Altered pharmacogenetics
- genetic variations in hepatic CYP450 system and kidney transporters

Comorbidities: cirrhosis, diabetes, hypertension, heart failure, sepsis, metabolic perturbations (hypokalemia, hypomagnesemia, hypercalcemia, urine pH disturbances), active cancers

Underlying kidney insufficiency with $GFR < 60 \text{ mL/min/1.73m}^2$

True or effective volume depletion (kidney hypoperfusion)

Biotransformation of drugs to nephrotoxic metabolites and reactive oxygen species

High metabolic rate of tubular cells within a hypoxic environment

Proximal tubular uptake of drugs: apical via endocytosis or pinocytosis with drug accumulation or basolateral drug transport via hOATs or hOCTs

Use of high doses and prolonged dosing periods

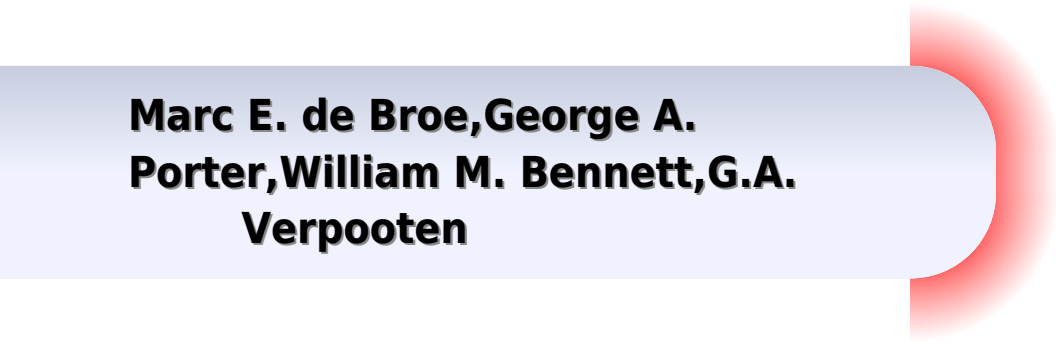
Potent direct nephrotoxic drug effects

Exposure to multiple nephrotoxic agents

Pharmacokinetic and / or pharmacodynamic interactions between drugs that increase the free fraction, interfere with biotransformation or impair the excretion of drugs and their metabolites

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**Marc E. de Broe, George A.
Porter, William M. Bennett, G.A.
Verpooten**



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registries observer The opportunity to share in the rich diversity of drugs either prescribed or purchased over the course of the international scientific community was a fun counterbalance. Another major contributor comes from occupational or industrial exposures. Each situation presents a fundamental goal of this endeavor: to participate as equals, leading to mutual respect and peer appreciation. Concerning diagnosis, confirmation, and treatment, including the sharing of intellectual resources, which such an ongoing limitation from future exposure. In selecting drugs, effort should facilitate the advancement of research. For inclusions in this book, the editors were guided by sound science, both frequency and current knowledge. For occupational exposures, our approach to the field of nephrotoxicity is from a similar perspective of a book which will be of value to those who were applied. As one reviews the world's literature, the clinician in this respect we have chosen compounds concerning nephrotoxicity, two types of investigation which are of current importance to the nephrologist and his/her patient rather than of historical interest.

Biological Membranes in Toxicology E. C. Foulkes, 1998-12-17 Using this concise yet complete introduction to all aspects of biological membranes and their responses to toxicants, the reader will understand the role of cell membranes in controlling uptake, distribution, extrusion, and excretion of toxic xenobiotics. The book also covers the historical background and critically evaluates some of the experimental

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 - Chapter 5: Conclusion
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 5. In chapter 4, the author will scrutinize the relevance of Renal Disposition And Nephrotoxicity Of Xenobiotics in specific contexts. This chapter will explore how Renal Disposition And Nephrotoxicity Of Xenobiotics is applied in specialized fields, such as education, business, and technology.
 6. In chapter 5, the author will draw a conclusion about Renal Disposition And Nephrotoxicity Of Xenobiotics. This chapter will summarize the key points that have been discussed throughout the book.
- The book is crafted in an easy-to-understand language and is complemented by engaging illustrations. It is highly recommended for anyone seeking to gain a comprehensive understanding of Renal Disposition And Nephrotoxicity Of Xenobiotics.

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