

pharmacokinetics and pharmacodynamics of atorvastatin

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The standard cholesterol diet was used to induce hyperlipidemia in Wistar rats. *Journal of Pharmacy Research*. Nifedipine, a hypertensive calcium channel blocker, is commonly administered to subjects with coronary heart disease who often exhibit hyperlipidemia. In vitro anti-Candida activity of *Calotropis gigantea*. Forgot Password Forgot Username. Average reductions from baseline were Meropenem is a broad-spectrum antibacterial that is usually used in the treatment of serious lower respiratory tract infections LRTIs. Response to Lin and Ito. Narsimha Reddy Yellu E-mail: Cite article How to cite? Pharmacodynamics and pharmacokinetic-pharmacodynamic relationships of atorvastatin, an HMG-CoA reductase inhibitor. Stern RH(1), Yang BB, Hounslow NJ, MacMahon M, Abel RB, Olson SC. Author information: (1)Department of Clinical Pharmacology, Parke-Davis Pharmaceutical Research Division of. **STUDY OBJECTIVE:** To investigate the potential effect of atorvastatin 80 mg/day on the pharmacokinetics and pharmacodynamics of the thienopyridines prasugrel and clopidogrel. **DESIGN:** Open-label, randomized, crossover, two-arm, parallel-group study. **SETTING:** Single clinical research center in the United Kingdom. LIPITOR tablets for oral administration contain 10, 20, 40 or 80 mg atorvastatin and the Pharmacodynamics. Atorvastatin as well as some of its metabolites are pharmacologically active in humans. The liver is the primary site of action and the principal site of cholesterol synthesis Pharmacokinetics and Drug Metabolism. Atorvastatin (Lipitor) is a member of the drug class known as statins. It is used for lowering cholesterol. Atorvastatin is a competitive inhibitor of hydroxymethylglutaryl-coenzyme A (HMG-CoA) reductase, the rate-determining enzyme in cholesterol biosynthesis via the mevalonate pathway. HMG-CoA reductase catalyzes the. Geriatric patients (>65 years old) exhibit altered pharmacokinetics of atorvastatin compared to young adults, with mean AUC and C_{max} values that are 40% and 30% higher, respectively. Additionally, healthy elderly patients show a greater pharmacodynamic response to atorvastatin at any dose; therefore, this population. Abstract. The pharmacodynamic effects and pharmacokinetics of atorvastatin, a potent investigational inhibitor of HMG-CoA reductase, were studied in 16 normolipidemic subjects after administration of 40 mg daily for 15 days in the morning or evening. Lipid and apolipoprotein parameters were determined, and plasma. The pharmacodynamic effects and pharmacokinetics of atorvastatin, a potent investigational inhibitor of HMG-CoA reductase, were studied in 16 normolipidemic subjects after administration of 40 mg daily for 15 days in the morning or evening. Lipid and apolipoprotein parameters were determined, and plasma atorvastatin. 1 Feb In Geriatric patients (>65 years old) show altered pharmacokinetics of atorvastatin compared to young adults. The mean AUC and C_{max} values are higher (40% and 30%, respectively) for geriatric patients. Additionally, healthy elderly patients show a greater pharmacodynamic response to atorvastatin at. [20]. This paper aims to provide an update of the chemical and pharmacokinetic properties of statins, as well as reviewing their lipid-modifying and safety profiles. Chemistry and functional properties. Lovastatin, pravastatin and simvastatin are fungal-derived inhibitors of HMG-CoA reductase, while atorvastatin, cerivastatin. In animal models, LIPITOR lowers plasma cholesterol and lipoprotein levels by inhibiting HMG-CoA reductase and cholesterol synthesis in the liver and by increasing the number of hepatic LDL receptors on the cell Pharmacodynamics Effect of Co-administered Drugs on the Pharmacokinetics of Atorvastatin.