

SECURITY PROFILE OF PATIENTS TREATED WITH PHENYTOIN IN A HOSPITAL

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Background

The determination of the plasma concentration of free phenytoin (CpFL) could improve the seizure control and prevention of adverse effects.

Purpose

To evaluate security profile of patients treated with phenytoin using CpFL.

Material and methods

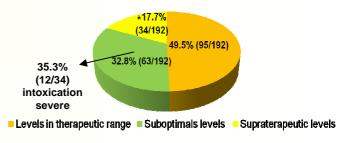
Prospective study (2013-2014) in a hospital. Phenytoin therapeutic range, C_pFL: 1-2.5 mcg/mL. Moderate intoxication, C_pFL: 2.5-3.0 mcg/mL and severe, C_pFL > 3.0 mcg/mL. To determine the renal clearance was used CKD-EPI. It was defined moderate renal impairment as Cl_{cr}: 20-50 mL/min. Polymedicated patients: > 5 drugs. Statistical analysis: Spearman correlation and χ^2 test.

Results

Table1. Patients treated with pheytoin (2013-2014)

	Monitored patients: 93 (192 cases) (range of cases/patient: 1-6)
Men / Age (years)	48/93 (51.6%) / 58 (range: 27- 84)
Daily dose (mg)	299
C _p FL (mcg/mL)	1.1 (range: 0.5-5.7)
Cl _{cr} (mL/min)	51.7
Serum albumin (g/dL)	3.6

Intoxicated patients treated with phenytoin (2013-2014)



*Plasma concentration of free phenytoin: 3.8 mcg/mL; range: 2.6-5.7 mcg/mL. Age: 71 years, creatinine clearance: 38.9 mL/min, serum albumin 3.4 g/dL.

CpFL: plasma concentration of free phenytoin; Clcr: creatinine clearance.

Three patients were hospitalised. Polymedicated patients: 71% vs. 50% the rest. Patients with drugs that bind over 70% to plasma proteins: 48%. Patients > 70 years had a higher risk of intoxication (p=0.033). We observed an inverse correlation between C_pFL and C_{lcr} (Spearman rho: -0.562; p=0.04) or with albumin (Spearman rho: -0.623; p<0.01). In relation to moderate intoxication, concentration of phenytoin plasmatic presented a value 23% higher than C_pFL .

Conclusion

Elderly patients, polymedicated and with moderate renal insufficiency and hypoalbuminemia presented a higher risk of phenytoin toxicity. It would be advisable to be careful with these patients because in our study efficacy / toxicity is correlated better with CpFL.

Keywords: phenytoin, safety, pharmacokinetics.



