

Background

Polypharmacy in elderly patients

Associated with a **high potential for drug-drug interactions**

Different cytochrome P450 (CYP) isoforms and transporters may be involved and variously disturbed

The aim of our study was to identify frequently disturbed metabolic or transport pathways in geriatric patients and to identify involved drugs.

Settings and method

From a multicentric cohort of hospitalized elderly patients

Analysis of drug prescriptions on a given day

Following inhibitors and inducers of metabolic and transport pathways were investigated

Drugs involved were identified

Were excluded: dosage forms with low systemic absorption (ointments, drops and inhalants)

CYP 1A2 CYP 2C19
CYP 2B6 CYP 2D6
CYP 2C8 CYP 2E1
CYP 2C9 CYP 3A4
P-gP



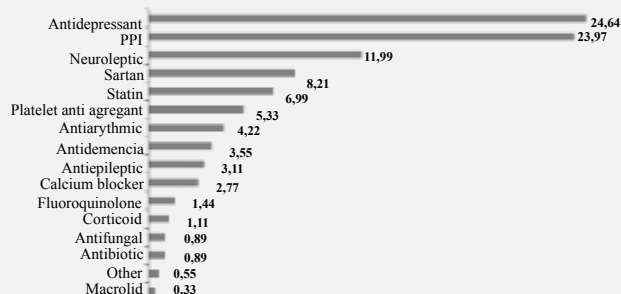
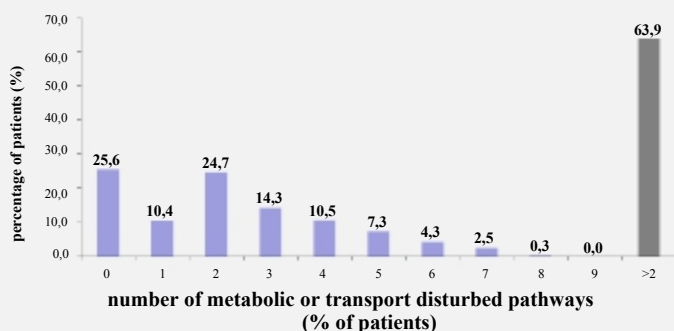
Results

Number of prescriptions analysed	672	
Sex ratio (M/F)	200/472	
	Mean	Standard deviation
Age (years)	85,9	7,5
Mean number of drugs	7,8	3,2

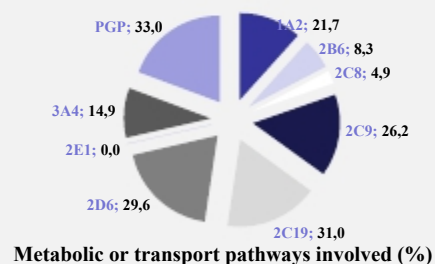
Mean metabolic or transport pathways potentially disturbed

2.3 (SD= 1.9)

Two disrupted pathways for 63.9% of patients



Drugs involved (%)



Metabolic or transport pathways involved (%)

Discussion

Prescribed drugs for elderly people

significant number of metabolic or transport pathways disrupted

New prescription in elderly:

high risk of potential drug- drug interactions

could lead to overdose, side effects or ineffective treatment

Frequent used of most involved drugs

questions about the **safety** of their use in the elderly

introduction of a new drug in these frail and polymedicated patients:

should always be **carefully** considered in terms of potentially disturbed metabolic or transport pathways