Drug Use in the Elderly

Therapeutic Considerations in the Elderly

- Elderly patients are those 65 years of age and older.
- The health characteristics of those 65-74 years of age are different from those who are 85 years of age and older.
- Institutionalized individuals are also different from those living in the community.
- Age-related changes in physiology can affect the pharmacokinetics and pharmacodynamics of drugs.

Therapeutic Considerations in the Elderly

- Drug-related problems in older adults are common and cause significant morbidity.
- Common medical conditions in the elderly include: hypertension, diabetes mellitus, bronchial asthma, COPD, cancer, arthritis, heart diseases, Alzheimer's disease and cognitive dysfunction, and stroke.
- The most common sensory impairments are: difficulties in hearing and vision.

Human Aging & Changes in Drug Pharmacokinetics and Pharmacodynamics

Clinical manifestations of normal aging include:

- 1. Changes in biochemical makeup of tissues.
- 2. Reduced <u>capacity</u> of body systems.
- 3. Reduced <u>ability to adapt</u> to physiological stress.
- 4. Increased <u>vulnerability</u> to disease.
- 5. Frailty (weakness, fatigue, weight loss and functional decline)
- Individuals experience aging at different rates.

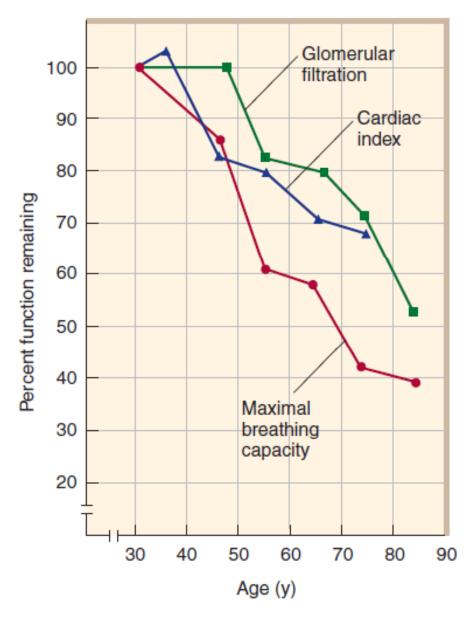


FIGURE 60–1 Effect of age on some physiologic functions. (Modified and reproduced, with permission, from Kohn RR: *Principles of Mammalian Aging*. Prentice-Hall, 1978.)

TABLE 60-1 Some changes related to aging that affect pharmacokinetics of drugs.

| Variable | Young Adults (20-30 years) | Older Adults (60-80 years) | |
|---------------------------------------|-------------------------------|-------------------------------|--|
| Body water (% of body weight) | 61 | 53 | |
| Lean body mass (% of body weight) | 19 | 12 | |
| Body fat (% of body weight) | 26-33 (women) | 38–45 | |
| | 18–20 (men) | 36–38 | |
| Serum albumin (g/dL) | 4.7 | 3,8 | |
| Kidney weight (% of young adult) | (100) | 80 | |
| Hepatic blood flow (% of young adult) | (100) | 55–60 | |

Common Physiological Changes Associated with Aging

- These changes may be associated with <u>reduced</u> <u>functional reserve capacity</u> and <u>reduced ability</u> <u>to maintain homeostasis</u>, making them susceptible to <u>de-compensation</u> in stressful situations.
- Examples of such impaired homeostatic mechanisms: postural or gait stability, orthostatic blood pressure responses, thermoregulation, cognitive reserve, and bowel or bladder function.

Absorption:

- Absorption of drugs may be affected by age-related changes in GIT physiology, drug-food interactions, concurrent medication, and co-morbidities affecting GI function.
- The bioavailability of drugs absorbed by passive diffusion may <u>not</u> be affected significantly.
- Drugs absorbed by active transport (vitamin B₁₂, calcium, iron, magnesium) may have impaired absorption.

- First-pass effect is decreased, bioavailability and plasma concentration are increased for drugs such as propranolol and labetolol. There is reduced bioavailability of pro-drugs such as enalapril and codeine.
- In atrophic gastritis, or in patients taking gastric acid-lowering agents, extent of absorption of some drugs may be reduced (ketoconazole, iron, digoxin, and atazanavir). These require an acidic environment for absorption.

Distribution:

- The distribution of drugs in the body depends on blood flow, body composition and protein binding.
- The volume of distribution of water-soluble drugs (ethanol, gentamicin, digoxin, and cimetidine) is reduced.
- Lipophilic drugs (benzodiazepines, metronidazole, and rifampin) exhibit an increased volume of distribution.

- Changes in the volume of distribution affect loading doses of drugs.
- Reduction of tissue perfusion with aging slows the distribution of dugs to their site of action.
- Decreased serum albumin and increased α_1 -acid glycoprotein with aging may affect the distribution of dugs that have high hepatic extraction ratio, extensive protein binding, especially when administered intravenously.

 The brain of elderly patients may be exposed to higher concentrations of drugs and toxins because of age-related changes in the bloodbrain-barrier.

Metabolism:

- Hepatic metabolism of drugs depends on liver perfusion, activity and capacity of drug metabolizing enzymes, and protein binding.
- All of these factors are affected by the aging process.
- For drugs that have high intrinsic clearance (high hepatic extraction ratio), hepatic clearance depends on hepatic blood flow (flow-limited metabolism).

- For drugs that have low intrinsic clearance (low hepatic extraction ratio), clearance depends on hepatic enzyme activity (capacity-limited metabolism).
- Age-related decreases in hepatic blood flow (20-50%) can decrease significantly the metabolism of high extraction ratio drugs (propranolol, amitriptyline, diltiazem, lidocaine, metoptolol, morphine and verapamil).

- Generally, liver size and its enzyme content are reduced in the elderly.
- Hepatic metabolism of warfarin, piroxicam and lorazepam is reduced with aging.
- Metabolism of phenytoin, ibuprofen, and naproxen is increased with aging.
- Metabolism of diazepam, temazepam, and valproic acid is not affected with aging.

- Serum albumin concentration declines with age.
- For capacity-limited drugs with extensive protein binding, the fraction of the drug unbound will increase, leading to increased total hepatic clearance (naproxen).
- Generally, phase II drug metabolism, in contrast to phase I, is preserved in the elderly.
- Frail older adults may experience reduced phase
 Il drug metabolism as well.

Elimination:

- Age-related reductions in GFR are well documented.
- Serum creatinine is a poor indicator of renal function in the elderly. Creatinine is produced by muscles and there may be reduction in muscle mass in the elderly. (i.e. reduced creatinine production).

 Cockcroft and Gault equation may be used to calculate creatinine clearance:

Creatinine clearance =
$$\frac{(140 - \text{Age}) \text{ (Actual body weight)}}{72 \text{ (Serum creatinine concentration)}}$$

Multiply the result by 0.85 for females.

You should measure CLcr accurately when you plan dose adjustment in patients with reduced renal function.

- Dosing guidelines of drugs that are eliminated by the kidney are based on creatinine clearance.
- Some drugs that should be avoided when CLcr <
 30 mL/min: colchicine, co-trimoxazole,
 glyburide, nitrofurantoin, probenecid,
 spironolactone, triamterene.
- Some drugs that need dose reduction in reduced renal function: acyclovir, amantadine, ciprofloxacin, gabapentin, ranitidine.

- Less understood than altered pharmacokinetics.
- Proposed changes include:
- 1. Changes in drug concentration at the receptor.
- 2. Changes in receptor numbers.
- 3. Changes in receptor affinity.
- 4. Post-receptor changes.
- 5. Age-related changes in homeostatic mechanisms.

- Older adults are sensitive to the CNS effects of drugs:
- 1. Changes in size and weight of brain.
- 2. Changes in the neurotransmitter systems.
- 3. Drugs penetrate CNS easier than in young adults.
- For example, in elderly there is decreased levels of dopamine transporters, decreased number of dopaminergic neurons, and decreased density of dopamine receptors; leading to increased sensitivity to the adverse effects of antipsychotic drugs.

- There is also increased sensitivity to benzodiazepines, opioids, general anesthtecis.
 Antipsychotics, lithium and anticholinergic drugs.
- The elderly are more likely to develop orthostatic hypotension as an adverse effect of some drugs.

- There is increased hypotensive and bradycardic effect to calcium channel blockers.
- Reduced blood pressure response to β-blockers.
- Reduced effectiveness of diuretics.
- Increased risk of bleeding with warafarin.

- Include 3 important, potentially preventable, negative outcomes:
- 1. Withdrawal effects.
- 2. Therapeutic failure.
- 3. Adverse drug reactions.

Risk Factors:

- 1. Polypharmacy including prescription and nonprescription drugs, herbal medicines, supplements and unnecessary drugs.
- Polypharmacy has been strongly associated with ADRs, risk of geriatric syndromes (falls, cognitive impairment), non-adherance, diminished functional status, and increased health care costs.

- 2. Inappropriate Prescribing, which includes:
- a. Wrong dose and duration.
- b. Duplication.
- c. Drug interaction problem.
- d. Prescription of drugs that should be avoided in the elderly.
- 3. Underuse:
- Omission of drug therapy that is indicated in prevention or treatment of disease.

4. Medication non-adherence:

Causes:

- a. Adverse effects.
- b. Complex regimens.
- c. Misunderstanding of information about prescribed medications.
- d. Cost.
- e. Dys-mobiliy (arthritis, ..).
- f. Social factors (living alone).
- g. Dementia.

1. Compare the patient's problem list with drug list:

A drug may be considered unnecessary if:

- a. It does not have indication per the problem list.
- b. Is not effective.
- c. The risk of its use outweigh the benefits.
- d. There is therapeutic duplication.

- 2. Determine if the patient is having a chronic condition but is not receiving an evidence-based medication to improve outcome.
- 3. Monitor efficacy and toxicity of drugs by clinical assessment and lab tests.
- Examples:

Amiodarone hepatic function tests

Antiepileptics Drug level

ACEi & ARBs K+ level

Antipsychotics Extrapyramidal ADRs

Diuretics K+ level

Hypoglycemics Glucose and glycated HB

Lithium Serum level

Warfarin PT or INR

etc..

- 4. Documenting problems and Formulating a Therapeutic Plan:
- A reasonable outcome for a 40-year-old patient may not be reasonable for an 80-year-old patient.
- Take into account remaining life expectancy, time until therapeutic benefit, treatment target, medication regimen complexity, and goals of care when deciding on prescribing rationale.

- 5. Implement a team-based management approach and develop strategies to avoid prescribing errors.
- 6. Take measures to enhance adherence to medications:
- a. Modify medication schedule to fit patient's lifestyle.
- b. Prescribe generic agents to reduce cost.

- 5. Offer easy-to-open bottles.
- a. Offer easy-to-swallow dosage forms.
- b. Provide both written and oral drug information.
- c. Involve caregivers stressing the importance of adherence.

Assessing the presence of drug-disease interactions:

Anticholinergics: benign prostatic hyperplasia & dementia or cognitive impairment.

Antipsychotics: history of falls & Parkinson's disease.

Aspirin: peptic ulcer disease.

Calcium Channel blockers: heart failure.

Metoclopramide: Parkinson's disease.

NSAIDs: peptic ulcer disease, heart failure, renal failure.

etc..

Table 2. 2015 American Geriatrics Society Beers Criteria for Potentially Inappropriate Medication Use in Older Adults

| Organ System, Therapeutic Category, | Rationale | Recommendation | Quality of Evidence | Strength of Recommendation | Evidence |
|---|---|----------------|------------------------|-------------------------------|--|
| Drug(s) | | | Evidence | Recommendation | |
| Anticholinergics | <u> </u> | | l | | |
| First-generation antihistamines: Brompheniramine Carbinoxamine Chlorpheniramine Clemastine Cyproheptadine Dexchlorpheniramine Dimenhydrinate Dimenhydrinate Diphenhydramine (oral) Doxylamine Hydroxyzine Meclizine Promethazine Triprolidine | Highly anticholinergic; clearance reduced with advanced age, and tolerance develops when used as hypnotic; risk of confusion, dry mouth, constipation, and other anticholinergic effects or toxicity Use of diphenhydramine in situations such as acute treatment of severe allergic reaction may be appropriate | Avoid | Moderate | Strong | 2015 Criteria: Duran 2013 Fox 2014 Kalisch Ellet 2014 From previous criteria: Agostini 2001 Boustani 2007 Guaiana 2010 Han 2001 Rudolph 2008 |
| Antiparkinsonian agents Benztropine (oral) Trihexyphenidyl | Not recommended for prevention of extrapyramidal symptoms with antipsychotics; more- effective agents available for treatment of Parkinson disease | Avoid | Moderate | Strong | Rudolph 2008 |
| Antispasmodics: Atropine (excludes ophthalmic) Belladonna alkaloids Clidinium- | Highly anticholinergic, uncertain effectiveness | Avoid | Moderate | Strong | Lechevallier- Michel 2005 Rudolph 2008 |

- 1. Anticholinergics + other drugs with anticholinergic activity such as antihistamines:
- Rationale: elimination reduced in older adults
- Risk: confusion, dry mouth, constipation, urine retention
- Quality of evidence: moderate.
- Strength of recommendation: strong.

2. Nitrofurantoin:

- Rationale: potential for pulmonary toxicity, hepatotoxicity, and peripheral neuropathy.
- Quality of evidence: low.
- Strength of recommendation: strong.
- 3. Peripheral and central α -blockers:
- Rationale: High risk of adverse effects, orthostatic hypotension, and CNS adverse effects respectively.
- Quality of evidence: moderate low.
- Strength of recommendation: strong.

- 4. Immediate-release nifedipine:
- Rationale: potential for hypotension and myocardial ischemia.
- Quality of evidence: high.
- Strength of recommendation: strong.
- 5. Amiodaraone:
- Rationale: High risk of many adverse effects.
- Quality of evidence: high.
- Strength of recommendation: strong.

6. Antidepressants:

- Rationale: highly anticholinergic, sedating, orthostatic hypotension and myocardial ischemia.
- Quality of evidence: high.
- Strength of recommendation: strong.
- 7. Antipsychotics:
- Rationale: increased risk of CVA, cognitive decline, dementia, and mortality.
- Quality of evidence: moderate.
- Strength of recommendation: strong.

- 7. Barbiturates & benzodiazepines:
- Rationale: highly rate of dependence, tolerance, sedation, cognitive impairment, delirium, falls, fractures.
- Quality of evidence: high moderate.
- Strength of recommendation: strong.
- 8. Insulin, sliding scale:
- Rationale: increased risk of hypoglycemia.
- Quality of evidence: moderate.
- Strength of recommendation: strong.

- 9. Long-acting sulfonylureas:
- Rationale: increased risk of hypoglycemia.
- Quality of evidence: high.
- Strength of recommendation: strong.
- 10. Metoclopramide:
- Rationale: increased risk of extrapyramidal adverse effects, dyskinesia.
- Quality of evidence: moderate.
- Strength of recommendation: strong.

11. Proton pump inhibitors:

- Rationale: risk of Clostridium difficile infection.
- Quality of evidence: high.
- Strength of recommendation: strong.

12. Meperidine (pethidine):

- Rationale: high risk of neurotoxicity, including delirium.
- Quality of evidence: moderate.
- Strength of recommendation: strong.

13. NSAIDs:

- Rationale: Increased risk of peptic ulcer disease, cardiovascular disease, renal failure.
- Quality of evidence: moderate.
- Strength of recommendation: strong.
- 14. Central muscle relaxants (chlorzoxazone, cyclobenzaprine, orphenadrine):
- Rationale: poorly tolerated because of anticholinergic effects, sedation, increased risk of fractures.
- Quality of evidence: moderate.
- Strength of recommendation: strong.