

The Pharmacotherapy Preparatory Review & Recertification Course Pediatrics Kirsten H. Ohler, PharmD, BCPS University of Illinois at Chicago

#### Conflict of Interest Disclosure

 The speaker, Kirsten Ohler, has no real or potential conflicts of interest related to the subject matter in this presentation.

#### Agenda

Discuss the pharmacological management of the following pediatric disease states:

- Pediatric and neonatal sepsis/meningitis
- Respiratory syncytial virus (RSV)
- Otitis media
- Immunizations
- Pediatric seizure disorders
- Attention deficit hyperactivity disorder (ADHD)

#### Case 1

Neonate born at 36 week's gestational age develops respiratory distress, hypotension, and mottling at 5 hours of life. Witnessed seizure in the NICU. Mother is GBS positive; three doses of penicillin given before delivery

#### Best empiric antibiotic regimen?

- a. Ampicillin + gentamicin
- b. Cefuroxime
- c. Ceftriaxone + vancomycin
- d. Rifampin

#### Sepsis/Meningitis - Pathogens

| <u>Age</u><br>0 - 1 month | <u>Organism</u><br>Group B β Streptococcus, E. coli,<br>Listeria, viral, nosocomial |
|---------------------------|---|
| 1 - 3 months              | Neonatal pathogens, H. influenzae,<br>N. meningitidis, Strep pneumoniae             |
| 3 mo - 12 yr              | H. influenzae, N. meningitidis, Strep<br>pneumoniae                                 |
| > 12 yr                   | N. meningitidis, Strep pneumoniae   |
|                           | Page 1- 5   |

#### Case 1

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Best empiric antibiotic regimen?

- 🕨 a. Vancomycin
  - b. Ampicillin + gentamicin
  - c. Ampicillin + ceftriaxone
  - d. Ceftazidime + gentamicin

Page 1-5

#### Case 2

Culture results reveal gram negative rods in the cerebral spinal fluid.

Which recommendation regarding antibiotic prophylaxis is best?

- a. 5-month old stepsister is at high risk and should receive rifampin
- b. The patient should receive rifampin to eliminate nasal carriage
- c. Antibiotic prophylaxis is not indicated
- d. All close contacts should receive rifampin

## Chemoprophylaxis

- Purpose: prevent the spread of Haemophilus influenzae and Neisseria meningitidis
- High risk groups: household contacts, nursery or child care center contacts, direct contact with patient's secretions
- Drug of choice: rifampin

Page 1-6

#### Case 2 Culture results reveal gram negative rods in the cerebral spinal fluid. Which recommendation regarding antibiotic prophylaxis is best? a. 5-month old stepsister is at high risk and should receive rifampin b. The patient should receive rifampin to eliminate nasal carriage c. Antibiotic prophylaxis is not indicated d. All close contacts should receive rifampin

d. All close contacts should receive rifampin Page 1-6

#### Case 3

6-year-old boy presents to the ED with fever, altered mental status & petechiae. No trauma. Tox screen negative. Elevated WBC with a left shift. Cultures are pending.

#### Best empiric antibiotic regimen?

- a. Ampicillin + gentamicin
- b. Cefuroxime
- c. Ceftriaxone + vancomycin
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#### Page 1-6

## Age Organism 0 - 1 month Group B β Streptococcus, E. coli, Listeria, viral, nosocomial 1 - 3 months Neonatal pathogens, H. influenzae, N. meningitidis, Strep pneumoniae 3 mo - 12 yr H. influenzae, N. meningitidis, Strep pneumoniae > 12 yr N. meningitidis, Strep pneumoniae

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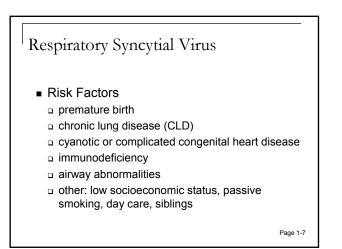
#### Case 4

You are screening babies during RSV season for risk factors associated with the development of severe RSV infection.

Which of the following is the best recommendation to make regarding the use of palivizumab for RSV prophylaxis?

Page 1-7

## Case 4 Palivizumab should <u>NOT</u> be prescribed for: a. A 34 weeks' gestation baby with a cyanotic congenital heart defect b. A 21-day-old, 31 weeks' gestation baby, only child, non-smoking parents, will not attend day care c. A 5-month-old, 29 weeks' gestation infant, history of CLD, no O<sub>2</sub> or meds d. An 18-month-old, 26 weeks' gestation infant

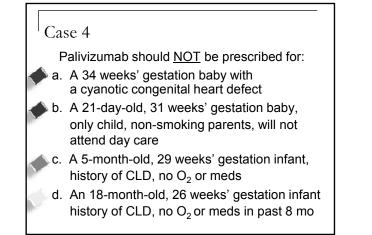


#### Respiratory Syncytial Virus

- AAP recommendations for prophylaxis
  - □ infants born < 32 weeks who are ≤ 6 mo at the beginning of RSV season</p>

history of CLD, no O2 or meds in past 8 mo

- infants with CLD who are < 2 yo and require medical management of CLD w/in last 6 months
- □ infants between 32 34 weeks, 6 days gestation who are ≤ 3 mo at the beginning of RSV season with risk factors may benefit
- □ infants ≤ 24 months of age with hemodynamically significant congenital heart disease Page 1-8



#### Case 5

18-month-old with history of premature birth and CLD is admitted to the PICU with respiratory distress requiring intubation, fever, and a 3-day history of cold-like symptoms. A nasal swab is positive for respiratory syncytial virus.

#### Case 5

Which is the best intervention?

- 🕨 a. Palivizumab
- b. Corticosteroids
- c. Cefuroxime
- d. Intravenous fluids and supportive care

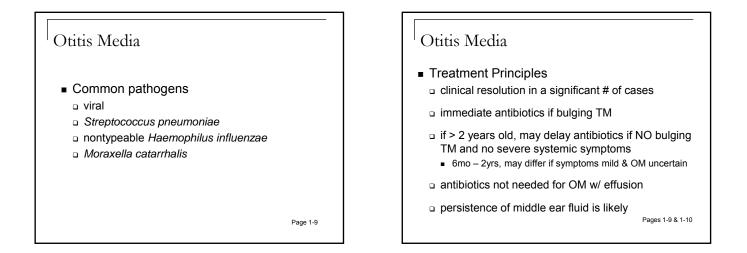
#### Case 6

A 5-month-old infant, born at term, healthy is treated for her first case of otitis media with amoxicillin 45 mg/kg/day for 7 days. Follow-up exam shows fullness of middle ear, cloudy TM. Afebrile and eating well.

Best treatment recommendation?

- a. No antibiotics are warranted at this time
- b. High-dose (90 mg/kg/day) amoxicillin x 7 days
- c. Decongestant & antihistamine daily
- d. Azithromycin

#### Page 1-10



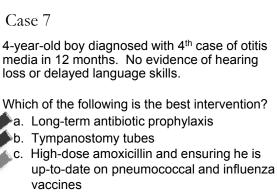
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Page 1-10



d. No antibiotic therapy is warranted

#### Case 8

1-year-old boy with history of Kawasaki disease treated 4 months ago with IVIG. At well-child check-up, due for MMR and varicella. Mother has several concerns regarding immunizations.

Best reason to defer administration of vaccines? a. Association between MMR & autism

- b. Allergic reaction to MMR if patient has egg allergy
- c. Many concurrent vaccines can overload immune system
- d. Decreased vaccine efficacy because of previous IVIG administration Page 1-11

#### Immunizations

- Barriers to routine immunization contraindications
  - anaphylactic reaction to the vaccine
  - acute moderate severe febrile illness
  - immunodeficiency, pregnancy, recent IVIG
  - encephalopathy w/in 7 days of previous DTaP

misconceptions regarding contraindications

mild acute illness, current antibiotics, etc.

Page 1-13

#### Immunizations Special populations Preterm infants immunize based on chronologic age Immunocompromised children no live vaccines Patients receiving corticosteroids recommendations depend on steroid dose / duration Patients who recently received IVIG affects live vaccines (ex. MMR, varicella) recommendations depend on indication / dose of IVIG HIV-infected patients recommendations depend on degree of immunocompromise Page 1-14



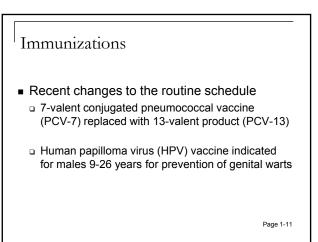
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#### Case 9 For which of the following patients would it be best to recommend deferring immunizations? a. 12-month-old boy who recently completed a cycle chemotherapy for ALL b. 6-month-old girl on amoxicillin for otitis media c. 12-month-old, HIV-positive boy with CD4 >1000 d. 12-year-old girl completing a prednisone

"burst" (1 mg/kg/day) for asthma exacerbation Page 1-14



| chedule (Figure 3))                         |       |       |        | *      |        |         |             | '        | _          | fall behind             |       |           |                         |
|---|-------|-------|--------|--------|--------|---------|-------------|----------|------------|-------------------------|-------|-----------|-------------------------|
|   |       | 1     | 2      | 4      |        | 9       | 12          |          |            | 19-23                   |       |           |                         |
| Vaccine V 🛛 Age 🕨                           | Birth | month | months | months | nonths | montins | months      | months   | months     | months                  | years | years     | Range of                |
| Hepatits B'                                 | Hep B | He    | pB     |        |        |         | НерВ        |          |            |                         |       |           | examed<br>scentral      |
| Rotevirus'                                  |       |       | RV     | RV     | R₩     |         |             |          |            |                         |       |           | dige                    |
| Diphtheria, tetanus, pertussis <sup>a</sup> |       |       | DTaP   | DTaP   | DīaP   |         | see bohole" | וס       | a?         |                         |       | DTaP      |                         |
| Haemophilus influenzae type b'              |       |       | Hib    | Hib    | Hib'   |         | H           | ib       |            |                         |       |           | fance di                |
| Pheumococcař                                |       |       | PCV    | PCV    | PCV    |         | P           | X        |            |                         | Př    | SV 🛛      | econnect<br>spector set |
| inactivated poliovirus <sup>1</sup>         |       |       | PV     | IPV    |        |         | IPV         |          |            |                         |       | IPV       | hgh tek<br>groups       |
| influenza <sup>7</sup>                      |       |       |        |        |        |         |             | Influenz | a (Yearly) |                         |       |           | 111                     |
| Neasles, mumps, rubella <sup>y</sup>        |       |       |        |        |        |         | M           | /R       |            | sec bahale"             |       | MMR       | (///                    |
| Varicella <sup>r</sup>                      |       |       |        |        |        |         | Vario       | cella    |            | see bahdit <sup>a</sup> |       | Varicelia | Range of<br>examined    |
| Hepatitis A <sup>ra</sup>                   |       |       |        |        |        |         |             | Dos      | e1º        |                         | /HepÅ | Series//  | chiber and<br>estamligh |
| Neniroccocca/*                              |       |       |        |        |        |         |             | NCV4     | - see foo  | tnote"                  |       |           | ist grups               |

| Vaccine <b>V</b>           | Age⊮                       | 7-10 years                 | 11-12 years               | 13-18 years            |                             |
|----------------------------|----------------------------|----------------------------|---------------------------|------------------------|-----------------------------|
| Tetanus, diphtheri         | ia, pertussis <sup>1</sup> | 1 dose (if indicated)      | 1 dose                    | 1 dose (if indicated)  | Range of<br>recomme         |
| Human papilloma            | winus <sup>2</sup>         | See footnote <sup>2</sup>  | 3 doses                   | Complete 3-dase series | agestoral<br>children       |
| Meningococcal <sup>8</sup> |                            | See faatn ote <sup>3</sup> | Dose 1                    | Boosterat age 16 yea   | 5                           |
| Influenza <sup>4</sup>     |                            |                            | Influenza (yearly)        |                        |                             |
| Pheumococcal <sup>5</sup>  |                            |                            | See footnote <sup>5</sup> |                        | large of                    |
| Hepatitis A <sup>6</sup>   |                            |                            | Complete 2-dose series    |                        | ages for ca                 |
| Hepatitis B'               |                            |                            | Complete 3-dose series    |                        |                             |
| Inactivated policy         | rirus <sup>8</sup>         |                            | Complete 3-dose series    |                        |                             |
| Measles, mumps, i          | rubella <sup>9</sup>       |                            | Complete 2-dose series    |                        | Rapped                      |
| Varicella <sup>10</sup>    |                            |                            | Complete 2+dose series    |                        | ages for ce<br>high rest of |

#### Immunizations Special populations Preterm infants immunize based on chronologic age Immunocompromised children no live vaccines Patients receiving corticosteroids recommendations depend on steroid dose / duration Patients who recently received IVIG affects live vaccines (ex. MMR, varicella) recommendations depend on indication / dose of IVIG HIV-infected patients recommendations depend on degree of

immunocompromise Pages 1-14 & 1-15

### Case 9

For which of the following patients would it be best to recommend deferring immunizations?

- a. 12-month-old boy who recently completed a cycle chemotherapy for ALL
- b. 6-month-old girl on amoxicillin for otitis media
- c. 12-month-old, HIV-positive boy with CD4 >1000
- d. 12-year-old girl completing a prednisone "burst" (1 mg/kg/day) for asthma exacerbation Page 1-14

### Case 10 14-year-old moderately obese girl complains of oxcarbazepine three weeks ago for partial seizures. Sexually active + contraception. a. Change to carbamazepine



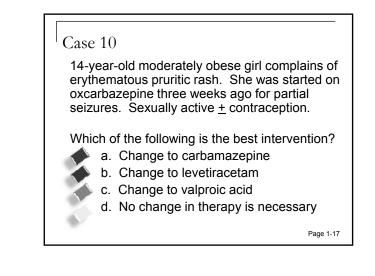
erythematous pruritic rash. She was started on

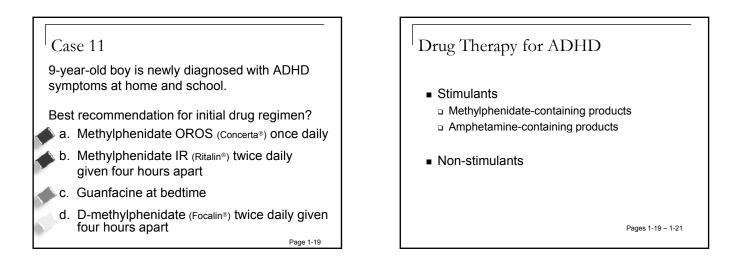
Which of the following is the best intervention?

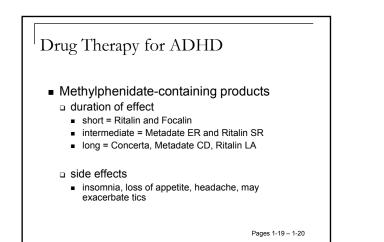
- b. Change to levetiracetam
- c. Change to valproic acid
- d. No change in therapy is necessary

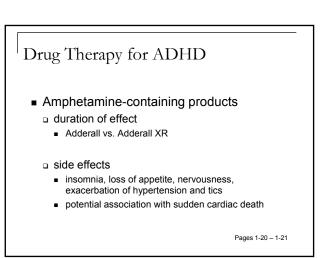
| Seizure type     | Drugs of Choice                 | Alternatives  |
|------------------|---------------------------------|---|
| Partial          | VPA, CBZ, PHT                   | PB, Gabapentin, Lamotrigine,<br>Tiagabine, Topiramate,<br>Oxcarbazepine, Zonisamide,<br>Levetiracetam |
| Generalized      |                                 |   |
| Tonic-clonic     | VPA, CBZ, PHT                   | Lamotrigine, Topiramate,<br>Zonisamide, Levetiracetam   |
| Myoclonic        | VPA                             | Topiramate, Zonisamide,<br>Levetiracetam  |
| Absence          | Ethosuximide, VPA               | Lamotrigine, Zonisamide,<br>Levetiracetam   |
| Lennox-Gastaut   | VPA, Topiramate,<br>Lamotrigine | Felbamate, Zonisamide   |
| Infantile spasms | ACTH                            | Lamotrigine, tiagabine,<br>topiramate, VPA, zonisamide  |

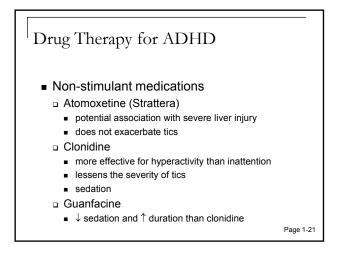
| Pediatric Seizures   |  |
|--|--|
| Rash• Carbamazepine• Oxcarbazepine• Lamotrigine• Phenytoin• Phenobarbital• ZonisamideMenstrual irregularities• Valproic acid | <ul> <li>Weight gain</li> <li>Valproic acid</li> <li>Gabapentin</li> <li>Weight loss</li> <li>Topiramate</li> <li>Zonisamide</li> <li>Cognitive/CNS effects</li> <li>Phenobarbital</li> <li>Topiramate</li> <li>Levetiracetam</li> </ul> |

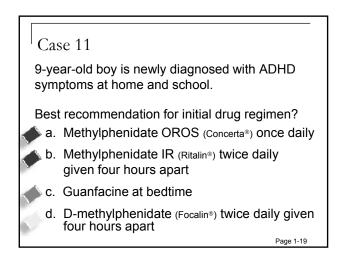












Case 12 The patient is started on methylphenidate OROS (Concerta®); symptoms well-controlled, but complaining of insomnia. Best modification to treatment regimen?

- a. Administer Concerta later in dayb. Change to methylphenidate modified
- release (Metadate CD) once a day.
- d. Change to atomoxetine at bedtime

Page 1-22

#### 2012 Updates in Therapeutics:

The Pharmacotherapy Preparatory Review and Recertification Course

#### Geriatrics

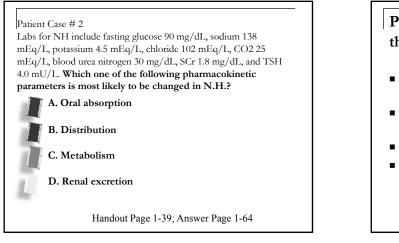
Jennifer Dugan, PharmD, BCPS Kaiser Permanente Colorado

#### **Conflict of Interest Disclosure**

Questions

 The speaker, Jennifer Dugan, has no real or potential conflicts of interest related to the subject matter in this presentation.

#### Patient Case 1 Patient Case # 1cont. Which of the following functional assessments NH is an 85 yo woman in a nursing facility. is most important? Type 2 DM, HTN, moderate dementia due to A. IADLS CVA, s/p hip fracture. B. Assessment for depression Glyburide 10 mg/d, lisinopril 10 mg/d, metformin 500 mg BID, donepezil 10 mg/d, C. Assessment for gait and balance aspirin 81 mg/d, MVI, zolpidem 5 mg QHS PRN, Meclizine 12.5 mg TID PRN, bowel D. Assessment for pressure sores regimen Handout Page 1-39; Answer Page 1-64 Handout Page 1-39; Answer Page 1-64



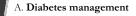
## Physiologic Changes in the Elderly Pearls

- Absorption from transdermal patches may be reduced if insufficient subcutaneous fat
- Distribution may be increased for highly protein-bound meds
- Metabolism impacts benzodiazepine choices
- Elimination is not just about Serum Creatinine

#### Patient Case # 3

Based on your assessment of age- and diseaserelated changes in N.H., which one of the following areas of pharmacotherapy is best to

#### address first?

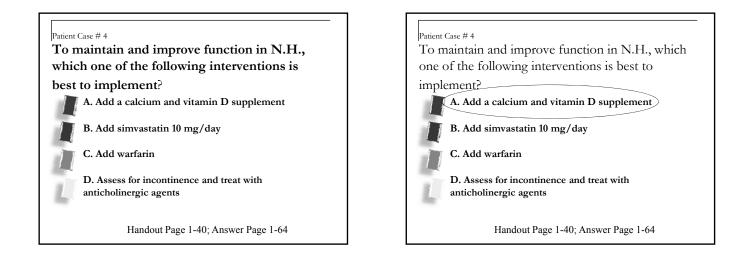


- B. Alzheimer disease treatment
- C. Hypertension treatment
- D. Stroke prevention

Handout Page 1-39; Answer Page 1-64

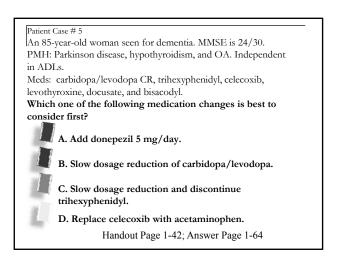
#### N.H. meds

- Glyburide 10 mg/day
- Lisinopril 10 mg/day
- Metformin 500 mg BID
- Donepezil 10 mg/day
- Aspirin 81 mg/day
- MVI



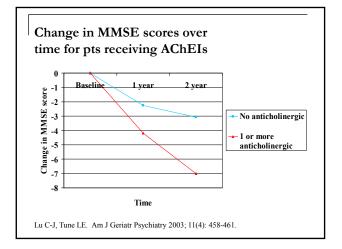
#### Common Drug Related Problems in Elderly

- Overuse
- Underuse
  - ACE inhibitors in CHF, anticoagulation in A fib, drug therapy post MI, untreated depression
- Medication Adherence
   Intentional nonadherence related to perceived overmedication, ADRs, cost
- Use of inappropriate medications
- Adverse drug events



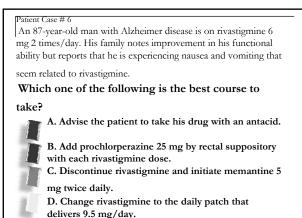
#### Potentially Inappropriate Medications

- Common offenders
  - Diphenhydramine
  - Long Acting Benzos (Diazepam, Chlordiazepoxide)
  - Skeletal Muscle Relaxants
  - Amitriptyline, Doxepin, Imipramine
  - GI antispasmodics and other anticholinergics
  - Indomethacin, Piroxicam
  - Promethazine
  - Butalbital compounds
  - Proposyphene



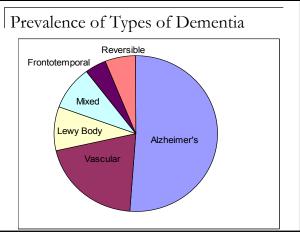
## Treating Adverse Effects with New Med

- Watch for prescribing cascade:
  - $\hfill\square$  Metoclopramide  ${\rightarrow} \textsc{Parkinsonian}$  sxs  ${\rightarrow}$  Levodopa
  - $\ \ \, \square \ \, \text{Donepezil} \rightarrow \text{Incontinence} \rightarrow \text{Oxybutynin}$
  - □ Diphenhydramine  $\rightarrow$  Urinary Retention  $\rightarrow$  Terazosin
  - $\Box \text{ Dihydropyridine CCB} \rightarrow \text{Edema} \rightarrow \text{Furosemide}$



vers 9.5 mg/day. Handout Page 1-44; Answer Page 1-64

# Symptoms of Dementia • Functional disability • Cognitive impairments • Behavioral and psychological symptoms



| Differentiating Dementias |   |
|---------------------------|---|
| Diagnosis                 | Key Symptoms  |
| Dementia with Lewy bodies | Visual hallucinations,<br>Parkinsonian sx, fluctuating<br>alertness |
| Vascular Dementia         | Acute onset, stepwise<br>deterioration, focal neurologic<br>signs   |
| Alzheimer's Disease       | Slow onset, progressive decline                                     |

#### Delirium

- Disturbance of consciousness and difficulty with attention
- Change in cognition (eg, memory deficit, disorientation, language disturbance, perceptual disturbance)
- The disturbance develops over a short period (usually hours to days) and tends to fluctuate during the course of the day.
- Evidence from the history, physical examination, or laboratory findings is present that indicates the disturbance is caused by a direct physiologic consequence of a general medical condition, an intoxicating substance, medication use, or more than one cause.

Adapted from: American Psychiatric Association. Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR). 4th ed. Washington, DC: American Psychiatric Association; 2000.

#### Therapy for Dementia

- Acetylcholinesterase Inhibitors
  - Donepezil
  - Galantamine
- Rivastigmine
- Memantine
- Efficacy and Safety Pearls

#### GI effects from AChEIs

|          | Donepezil | Galantamine | Rivastigmine<br>po | Rivastigmine<br>patch |
|----------|-----------|-------------|--------------------|-----------------------|
| Nausea   | 19%       | 24%         | 47%                | 7%                    |
| Vomiting | 8%        | 13%         | 31%                | 6%                    |
| Diarrhea | 15%       | 12%         | 19%                | 6%                    |

#### Patient Case # 7 RA is 75 yo woman with Alzheimer disease on donepezil 10 mg/day for 3 years. MMSE $21/30 \rightarrow 17/30$ . RA is at home with husband- can't do IADLs but can do ADLs with cueing.

Which one of the following is the best course of action?

A. Change her treatment from donepezil to rivastigmine.

B. Stop donepezil.

C. Add memantine 5 mg/day.

D. Add vitamin E 400 units 2 times/day.

Handout Page 1-45; Answer Page 1-64

#### Evaluating Efficacy

- Evaluate patient in 3-6 months to determine need for continued treatment
- Utilize caregiver reports, MMSE/SLUMS, and/or ADLs
- No change or mild improvement at 6 months→continue treatment
- Continued decline on therapy → consider discontinuation or changing medication
- 4 points/year is average decline without treatment

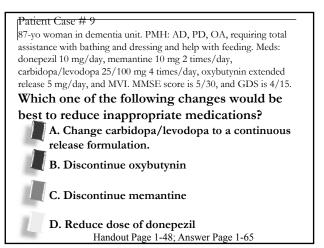
#### Patient Case # 8

87-yo woman in dementia unit. PMH: AD, PD, OA, requiring total assistance with bathing and dressing and help with feeding. Meds: donepezil 10 mg/day, memantine 10 mg 2 times/day, carbidopa/levodopa 25/100 mg 4 times/day, oxybutynin extended release 5 mg/day, and MVI. MMSE score is 5/30, and GDS is 4/15. Patient crying out "Help me, help me." Which one of the following additional assessment tools is most necessary in assessing this patient? **A. Brief Psychiatric Rating Scale** 

B. Functional Assessment Staging

C. An evaluation of incontinence

D. Framingham Risk Assessment Handout Page 1-48; Answer Page 1-64



Patient Case # 10

This same patient (MMSE 5/30, GDS 4/15) is medically assessed, and reversible causes of her hyper-vocalization are ruled out. Which one of the following represents the best approach to treating her behavioral symptoms? A. Implement a behavioral approach

B. Add valproic acid

- C. Add quetiapine
- D. Add citalopram
  - Handout Page 1-48; Answer Page 1-65

## General Approach to Behaviors in Dementia

- Define target agitated behavior
- Consider contributing causes
- Address ALL causes
- Non-pharmacologic measures
- Pharmacologic interventions

Lyketsos et al, Am J Geriatr Psychiatry July 2006;14:7

|           | ples of Non-pha<br>ventions               | rmacologic  |
|-----------|---|---|
| Behavior  | Causes                                    | Management  |
| Agitation | Discomfort, pain                          | Assess/manage pain, constipation, infection                     |
| I         | Physical illness (UTI)                    | Evaluate medically, treat                                       |
|           | Overstimulation-noise,<br>TV, people, etc | Reduce noise, stress; limit TV, crowding                        |
| Paranoia  | Forgot where placed object                | Offer to help find; have more than one of same object           |
|           | Misinterpreting actions<br>or words       | Do not argue or try to reason, do not take personally, distract |
| 1         | Change in environment                     | Familiarize, reassure, set routine                              |
| Insomnia  | Depression                                | Treat with antidepressant                                       |
| 1         | Less need for sleep                       | Later bedtime, more exercise                                    |

Adapted from Sutor B et al. Mayo Clin Proc. 2001;76:540-550

When should we consider pharmacologic treatment of BPSD?

 Behavior is dangerous, distressing, damaging to social relationships and persistent

AND

 Has not responded to comprehensive nonpharmacologic treatment plan, including removal of possibly offending drugs

OR

 Requires emergency treatment to allow proper investigation of underlying problems

#### Pharmacologic Treatment

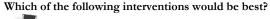
- Cochrane review suggests best evidence is with risperidone and olanzapine for psychosis and aggression
- Start at low doses
- Use quetiapine if patient has comorbid Parkinson's disease or Lewy Body Dementia
- Use for shortest duration possible
- Adverse effects include increased mortality; recent cohort study\* suggests worse with haloperidol, less with quetiapine

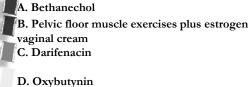
\*BMJ 2012;344:e977

#### Patient Case #11

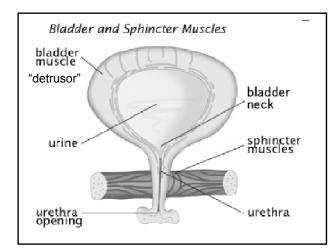
A 75-year-old woman reports urinary urgency, frequency, and loss of urine when she cannot make it to the bathroom in time. She wears a pad at night that she changes 2 or 3 times. PMH: Alzheimer disease (MMSE 23), osteoarthritis, and hypothyroidism.

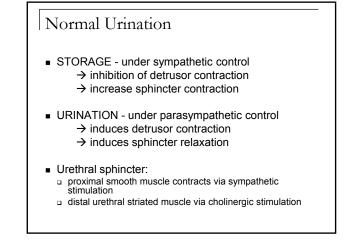
UA negative, exam WNL, PVR normal.





Handout Page 1-53; Answer Page 1-65





#### Types of Urinary Incontinence

- Functional
- Urge (Bladder overactivity)
- Stress (Urethral underactivity)
- Overflow (Urethral overactivity/Bladder underactivity)
- Mixed

#### Nonpharmacologic Interventions

- Pelvic floor exercises (Kegel exercises)
- Bladder training
- Biofeedback
- Scheduled/Timed Voiding
- Avoid aspartame, spicy/citrus foods, caffeine, carbonated beverages
- Pessaries/bladder neck support prostheses

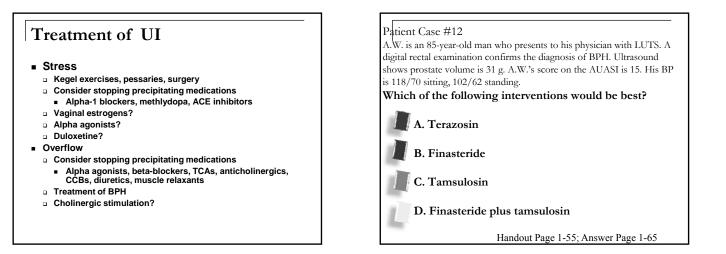
#### Treatment of UI

- Functional
  - Assist with functional disabilities
  - Scheduled bathroom visits
  - Bedside commode
  - Stop precipitating drugs
- Urge
  - Nonpharmacologic interventions
  - Anticholinergics (generally equivalent efficacy)

#### Anticholinergic Adverse Effects

| Drug         | Dry mouth % | Constipation % | Dizziness% |
|--------------|-------------|----------------|------------|
| Oxybutynin   | 88          | 32             | 38         |
| Oxy ER/XL    | 68          | 9              | 11         |
| Oxy TDS      | 10          | 5              | 4          |
| Oxy gel      | 8           | 1              | 3          |
| Tolterodine  | 50, 39      | 10, 10         | 4, 3       |
| Fesoterodine | 99          | 14             | 2          |
| Trospium     | 33          | 11             | ?          |
| Solifenacin  | 34          | 19             | 1          |
| Darifenacin  | 59          | 28             | 0          |

Treatment of overactive bladder in women. AHRQ Publication No. 09-E017. 8/09

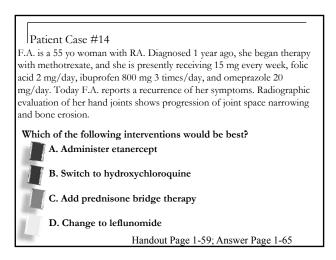


#### Patient Case #13 WF is an 85-year-old man with pain from hip OA. He also has hypertension, coronary artery disease, and BPH. For his OA, W.F. has been taking acetaminophen 650 mg 3 times/day. W.F. reports that acetaminophen helps, but he still experiences pain that limits his ability to Alpha Blockers walk. Which of the following interventions would be best? Alpha Reductase Inhibitors Combination Therapy A. Change the analgesic to celecoxib May be needed in men with LUTS, a larger B. Add hydrocodone prostate size (>40g), and an elevated PSA C. Change the analgesic to ibuprofen D. Add glucosamine Handout Page 1-58; Answer Page 1-65

#### **Osteoarthritis**

**BPH** 

- Nonpharmacologic Treatment
- Acetaminophen dosing
- NSAIDs vs Opioids
- Preventing adverse effects
- Glucosamine

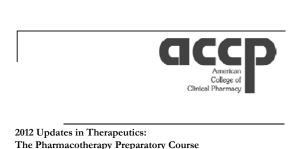


#### **Rheumatoid Arthritis**

- DMARDs first line
  - □ MTX
  - Hydroxychloroquine
  - Sulfasalazine
  - □ Leflunomide
- Biologic Treatments
- NSAIDs and Corticosteroids
  - Short term
  - No effect on disease progression

#### Questions

?????



The Pharmacotherapy Preparatory Course Gastrointestinal Disorders Brian A. Hemstreet, Pharm.D., BCPS University of Colorado Skaggs School of Pharmacy and Pharmaceutical Sciences

#### Conflict of Interest Disclosures

Dr. Hemstreet has conducted research sponsored by Astra Zeneca.

#### Learning Objectives

- 1. Review and apply national guideline treatment strategies for the following gastrointestinal (GI) disorders: gastroesophageal reflux disease (GERD), peptic ulcer disease (PUD), ulcerative colitis, Crohn's disease, viral hepatitis, chronic liver disease, upper GI bleeding, constipation, diarrhea, irritable bowel syndrome (IBS), nausea, vomiting, pancreatitis, prevention of stress related mucosal disease (SRMD).
- 2. Recommend appropriate pharmacologic and nonpharmacologic interventions for the treatment of GERD.
- 3. Differentiate between clinical signs, symptoms, risk factors, and treatment of both *Helicobacter pylori* and nonsteroidal anti-inflammatory drug (NSAID)-associated PUD.

#### Learning Objectives

- 4. Discuss the role of pharmacologic intervention in the treatment of nonvariceal upper GI bleeding.
- 5. Review the clinical differences in signs, symptoms, and treatment of Crohn's disease and ulcerative colitis.
- 6. Identify the common manifestations of chronic liver disease and their treatment.
- 7. Review the treatment of both acute and chronic viral hepatitis.

#### Learning Objectives

- 8. Recognize pertinent information for educating patients and prescribers regarding the appropriate use of pharmacologic agents for various GI disorders.
- 9. Recommend appropriate pharmacologic and nonpharmacologic interventions for diarrhea and constipation.
- 10. Review recommendations for the treatment and prevention of nausea and vomiting.

#### Learning Objectives

- 11. Discuss the clinical and treatment differences between acute and chronic pancreatitis.
- 12. Discuss the role of pharmacologic intervention in the treatment of IBS.
- 13. Understand commonly encountered statistical tests and concepts using GI disorders as examples.

#### Patient Case # 1

- HPI: 55 year old man with 8 month history of GERD symptoms 4-5 days/week. Prescriber wishes to initiate esomeprazole 20 mg/day.
- PMH: GERD, MI, HF, Hypothyroidism
- MEDS: Ranitidine + Calcium Carbonate, Metoprolol, Furosemide, Lisinopril, Aspirin

Handout Page 1-81; Answer Page 1-137

#### Patient Case # 1

- Which one of the following baseline tests is best to perform in this patient today before initiating his esomeprazole therapy?
- A. Peripheral bone mineral density screening.
- B. Serum magnesium.
- C. Serum potassium.
- D. Chest radiograph.

Handout Page 1-81; Answer Page 1-137

#### Treatment of GERD

- Nonpharmacologic/Lifestyle modifications
   Targeted
- Antacids
- Acid suppression (as needed or scheduled)
- Proton Pump Inhibitors
- Histamine-2 Receptor Antagonists
- Promotility Agents
- Proper patient education
- Surgical intervention

#### PPI Safety Concerns

| Adverse Effect                              | Prevention and Management  |  |  |
|---|--|--|--|
| Risk of Fracture<br>(Hip, wrist, spine)     | Re-evaluate need     Limit dose and duration     Ensuring adequate Calcium and Vitamin D     BMD screening if at risk for low bone mass     Weight bearing Exercise                                    |  |  |
| Hypomagnesemia                              | Re-evaluate need     Limit dose and duration     Consider baseline testing (diuretics, digoxin)     Supplementation  |  |  |
| Clostridium difficle<br>associated diarrhea | Re-evaluate need     Limit dose and duration     Evaluate for C. difficle if patient receiving PPI has diarrhea     that is not improving. Have patients report diarrhea.     Report cases to Medwatch |  |  |

#### Patient Case # 1

- Which one of the following baseline tests is best to perform in this patient today before initiating his esomeprazole therapy?
  - A. Peripheral bone mineral density screening.

#### B. Serum magnesium.

- C. Serum potassium.
- D. Chest radiograph.

Handout Page 1-81; Answer Page 1-137

#### Patient Case # 2

- HPI: 68 year old female with heme positive stools anemia and abdominal pain. Use of OTC ketoprofen for 2 months.
- PMH: Type 2 DM, Peripheral neuropathy, Hypertension
- MEDS: metformin, aspirin, gabapentin, lisinopril
- Diagnostics: endoscopy reveals 1 cm gastric ulcer with an intact clot, *H. pylori* negative via CLO Test

Handout Page 1-87; Answer Page 1-137

#### Patient Case # 2

- Which one of the following treatments is best for this patient's ulcer?
- A. Ranitidine 150 mg 2 times/day for 4 weeks
- B. Lansoprazole 30 mg 2 times/day plus amoxicillin 1000 mg 2 times/day plus clarithromycin 500 mg 2 times/day for 10 days.
- C. Lansoprazole 30 mg/day for 8 weeks
- D. Misoprostol 200 mcg 4 times/day for 8 weeks.

Handout Page 1-87; Answer Page 1-137

#### Peptic Ulcer Disease (PUD)

- Classification
   Duodenal ulcer
  - Gastric ulcer
- Etiologies
  - Helicobacter pylori (carcinogen)
  - NSAIDs
- Symptoms
  - Epigastric pain, nausea, anorexia, belching
  - May be temporally related to food intake

#### NSAID Associated PUD

- NSAIDs have topical and systemic adverse GI effects
   COX-2 vs. COX-1 effects
- Risk Factors
  - □ Age >60, History of PUD +/- complications
  - $\hfill\square$  Corticosteroids, anticoagulants, low dose aspirin, aspirin,  $\uparrow$  NSAID dose
- Contributing factors
  - □ H. pylori, Smoking, CVD, RA, SSRIs

#### Management of NSAID-Associated PUD

- Remove and reevaluate need for NSAID and/or aspirin
  - □ Test for *H. pylori* and treat if positive
- Acid suppression
   PPI for 8-12 weeks
- Misoprostol
- COX-2 Inhibitors
- Cardiovascular risks
- Use with aspirin

## Patient Case # 2 Which one of the following treatments is best for this patient's ulcer? A. Ranitidine 150 mg 2 times/day for 4 weeks B. Lansoprazole 30 mg 2 times/day plus amoxicillin 1000 mg 2 times/day plus clarithromycin 500 mg 2 times/day for 10 days. C. Lansoprazole 30 mg/day for 8 weeks D. Misoprostol 200 mcg 4 times/day for 8 weeks. Handout Page 1-87; Answer Page 1-137

#### Patient Case #3

- HPI: 42 year old male with sharp epigastric pain for 6 weeks. Pain is worse with eating and is present approximately 5 days per week. Some relief with OTC antacids.
- MEDS: antacids as needed
- Allergies: Penicillin (severe rash)
- UBT for *H. pylori* is positive

Handout Page 1-87; Answer Page 1-137

#### Patient Case #3

- Which one of the following treatments for *H. pylori* is best?
- A. Amoxicillin, clarithromycin, omeprazole for 10 days
- B. Cephalexin, clarithromycin, omeprazole for 10 days
- c. Bismuth, tetracycline, metronidazole, omeprazole for 14 days
- D. Levofloxacin, metronidazole, omeprazole for 10 days

Handout Page 1-87; Answer Page 1-137

#### Diagnosis of H. pylori

- Invasive testing (endoscopic)
  - Histology
  - □ Rapid urease (affected by antisecretory agents)
  - □ Culture
- Non-invasive testing
  - □ Serologic (IgG)
  - Urea breath test (affected by antisecretory agents)
  - □ Fecal antigen (affected by antisecretory agents)

#### Treatment of H. pylori

- Triple therapy
  - □ PPI + amoxicillin or metronidazole + clarithromycin
  - □ 10-14 days of treatment (14 preferred)
  - □ Efficacy affected by previous macrolide exposure
- Quadruple Therapy
  - □ PPI + Bismuth + Metronidazole + Tetracycline
  - Ist line, PCN allergy, previous macrolide exposure, failure of triple therapy
  - □ 10-14 days of treatment

#### Patient Case #3

• Which one of the following treatments for *H. pylori* is best?

Amoxicillin, clarithromycin, omeprazole for 10 days

Cephalexin, clarithromycin, omeprazole for 10 days

Bismuth, tetracycline, metronidazole, omeprazole for 14 days

Levofloxacin, metronidazole, omeprazole for 10 days

Handout Page 1-87; Answer Page 1-137

#### Patient Case #4

- HPI: 35 year old male with ulcerative colitis (majority of colon). Experiences 5-6 bloody bowel movements per day when prednisone is reduced to less than 40mg/day.
- MEDS: Balsalazide 6.75 g/day x 2 years, prednisone 40 mg/day x 1 year

Handout Page 1-100; Answer Page 1-137

#### Patient Case #4

- What would be an appropriate modification of his drug regimen at this time?
- A. Change balsalazide to sulfasalazine 6g/day
- B. Initiate therapy with methotrexate IM weekly
- c. Initiate infliximab and taper prednisone
- D. Add mesalamine suppository daily

Handout Page 1-100; Answer Page 1-137

| Clinical Findings        | Ulcerative Colitis | Crohn's Disease  |
|--------------------------|--------------------|------------------|
| Bowel Involvement        | Rectum/Colon       | Mouth to Anus    |
| Perianal Involvement     | No                 | Yes              |
| Depth                    | Superficial        | Submucosa/deeper |
| Pattern of inflammation  | Continuous         | Patchy           |
| Histology                | Crypt abscesses    | Granulomas       |
| Fistula, perforation, or | No                 | Yes              |
| Strictures               |                    |                  |
| Toxic megacolon          | Yes                | No               |
| Colorectal cancer        | Yes                | Uncommon         |
| Malnutrition             | Rare               | Yes              |
| Pseudopolyps             | Common             | Fairly Common    |

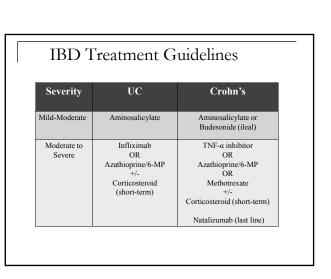
| Drug Treatmen  | nt Options   |
|--|--|
| <ul> <li>5-Aminosalicylates</li> <li>Sulfasalazine</li> <li>Mesalamine</li> <li>Olsalazine</li> <li>Balsalazide</li> <li>Antibiotics</li> <li>Metronidazole</li> <li>Ciprofloxacin</li> <li>Corticosteroids</li> </ul> | <ul> <li>Immunomodulators</li> <li>Azathioprine</li> <li>6-Mercaptopurine</li> <li>Methotrexate</li> <li>Cyclosporine</li> <li>Tacrolimus</li> <li>Biologics</li> <li>Infliximab</li> <li>Adalimumab</li> <li>Certolizumab</li> <li>Natalizumab</li> </ul> |
|  |  |

#### Approach to the Treatment of IBD

- 1. Indentify disease: UC vs. CD
- 2. Severity: Active (mild to fulminant) or remission
- 3. Determine extent and location of disease

#### 4. Pick drug(s) based on

- Onset of action
- □ Formulation (Oral, Topical, Parenteral)
- Effectiveness
- Potential adverse effects or contraindications



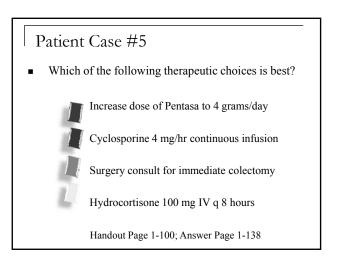
| Drug(s)                  | Adverse Effects   |  |  |
|--------------------------|---|--|--|
| TNF-alpha<br>antagonists | Risk of infection (screen for TB and Viral hepatitis)     Risk of Heart Failure and/or exacerbation     Hepatosplenic Tccell lymphoma when used with azathioprine or     6-MP in young male patients     Antibody formation |  |  |
| Antimotility<br>agents   | Risk of toxic megacolon in active disease   |  |  |
| Azathioprine/6MP         | Bone marrow suppression, pancreatitis, hypersensitivity     Need to check TPMT activity   |  |  |
| Methotrexate             | Bone marrow suppression, pulmonary and hepatic toxicity   |  |  |
| Corticosteroids          | Adrenal suppression, metabolic effects, infection   |  |  |
| Natalizumab              | Progressive mutilfocal leukoencephalopathy  |  |  |

# Patient Case #4 What would be an appropriate modification of his drug regimen at this time? Change balsalazide to sulfasalazine 6g/day Initiate therapy with methotrexate IM weekly Initiate infliximab and taper prednisone Add mesalamine suppository daily Handout Page 1-100; Answer Page 1-137

### Patient Case #5 HPI: 25 year old woman with Crohn's disease. Presents with a 2 day history of crampy abdominal pain, fever, fatigue, and 10-12 bloody stools per day.

- MEDS: Pentasa 250mg #4 caps 2 times/day
- PMH: Crohn's Disease x 5 years
- Vitals: Temp 101F, HR=110, RR=18, BP = 118/68

Handout Page 1-100; Answer Page 1-138



#### Patient Case #6

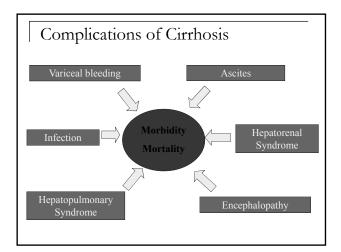
- HPI: 47 year old woman with nausea, abdominal pain, fever. Abdominal distention with tenderness and shifting dullness.
- PMH: Cirrhosis (Class C)
- MEDS: Furosemide, spironolactone
- Diagnostics: Paracentesis ( albumin 0.9 g/dl, WBC 1000/mm<sup>3</sup>)

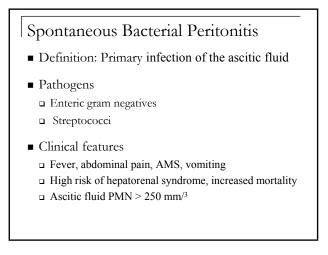
Handout Page 1-107; Answer Page 1-138

#### Patient Case #6

- Which recommendation is best at this time for treatment of this patient's hepatic encephalopathy?
- A. Intravenous albumin
- B. Intravenous vancomycin plus tobramycin
- c. Intravenous cefotaxime plus albumin
- D. Oral trimethoprim/sulfamethoxazole DS daily

Handout Page 1-107; Answer Page 1-138





#### SBP Treatment and Prevention

- Treatment: 3<sup>rd</sup> gen Cephalosporin + albumin
- Primary Prevention
  - During setting of an acute GI bleed
  - Ascitic fluid protein < 1.5 g/dl + Scr > 1.2 mg/dl or BUN > 25 mg/dl or Na < 130 mEq/L, or CP > 9 with bilirubin > 3 mg/dl
- Secondary Prevention: any patient with prior episode
- Hospital: Ceftriaxone/Cefotaxime, Fluoroquinolone
- Outpatient: TMP/SMX, Norfloxacin/ciprofloxacin

# Patient Case #6 • Which recommendation is best at this time for treatment of this patient's SBP? • Intravenous albumin • Intravenous vancomycin plus tobramycin • Intravenous cefotaxime plus albumin • Oral trimethoprim/sulfamethoxazole DS daily • Handout Page 1-107; Answer Page 1-138

#### Patient Case #7

- HPI: 36 year old female with 36 hours of hematemesis, fatigue, dizziness, black tarry stools.
- PMH: Cirrhosis, alcohol abuse, MI (2 years ago)
- Diagnostics: EGD several large esophageal varices that are banded.

Handout Page 1-107; Answer Page 1-138

#### Patient Case #7

- In addition to the endoscopic band ligation which of the pharmacologic interventions is best?
- A. Nadolol 20mg orally once a day x 3 days
- B. Vasopressin continuous infusion x 2 days
- c. Octreotide 50 ug bolus, then 50 ug/hr for 5 days
- D. Pantoprazole 80mg bolus, then 8mg/hr x 72 hours

Handout Page 1-107; Answer Page 1-138

#### Variceal Bleeding

- Varices: Collateral vessels formed secondary to increased resistance to blood flow within the liver
- Bleeding risk
  - □ 25-35% of patient with cirrhosis
  - □ 30-50% mortality per bleed
- High recurrence rate
   ~70% within first month of bleed

#### Treatment of Variceal Bleeding

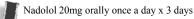
- Stabilization + IV fluids
- Endoscopic interventions
  - □ Sclerotherapy
  - Band ligation
- Medical Management
  - Vasopressin + nitroglycerin
  - Octreotide x 3-5 days
  - □ Antibiotics (3<sup>rd</sup> Gen Ceph or Fluoroquinolone)

#### Prevention of Variceal Bleeding

- Pharmacologic +/- endoscopic
- Primary prevention
  - Small varices + high bleeding risk
  - Medium/Large varices
  - Non selective beta blockers
- Secondary prevention
  - □ All patients with history of bleeding
  - Non selective beta blockers
  - Endoscopic (band ligation)

#### Patient Case #7

• In addition to the endoscopic band ligation which of the pharmacologic interventions is best?



Vasopressin continuous infusion x 2 days

Octreotide 50 ug bolus, then 50 ug/hr for 5 days

Pantoprazole 80mg bolus, then 8mg/hr x 72 hours

Handout Page 1-107; Answer Page 1-138

#### Patient Case #8

- HPI: 45-year old woman with history of IVDA. Diagnosed 8 months ago with HBV. Treatment naive. No ascites or encephalopathy.
- Diagnostics:
  - □ AST 650 IU/ml, ALT 850 IU/ml
  - □ HBSAg (+), HBeAg (+), YMDD mutation
  - $\square~$  HBV DNA 107, 000 IU/ml
  - □ Biopsy: severe necroinflammation/bridging fibrosis

Handout Page 1-121; Answer Page 1-138

#### Patient Case #8

- What is the most appropriate course of action at this time?
- A. No treatment; Recheck HBV DNA in 6 months
- в. Initiate PEG-IFN + ribavirin
- c. Initiate lamivudine 100 mg/day
- D. Initiate tenofovir 300 mg/day

Handout Page 1-121; Answer Page 1-138

#### Hepatitis B

- DNA Virus, Genotypes A-H
- Transmission
   Parenteral, bodily fluids, sexual contact, perinatal
- Detect via serologies, symptoms, LFTs
   Patients with active disease will be HBsAg (+)
- Treat patients with chronic disease (> 6 months)
   > 2 x ALT, HBV DNA > 20,000 IU/ml

#### Chronic Hepatitis B Treatment

- Need to distinguish if HBV:
  - · is HBeAg positive or negative
  - Harbors the "YMDD mutation" of the DNA polymerase
- Difficult patient populations
  - · Decompensated liver disease
  - •Co-infection
  - Treatment experienced

| HBV<br>Population | Preferred Treatment<br>Options   | Duration             | Comments   |
|-------------------|--|----------------------|--|
| HBeAg<br>positive | Entecavir and tenofovir are<br>preferred oral agents<br>Use of the other oral reverse<br>transcriptase inhibitors is<br>possible but not preferred | Minimum of 1<br>year | Preferred if contraindications or ·<br>nonresponse to INFα         |
|                   | INFα<br>PEG-INFα   | 16 weeks<br>48 weeks | If contraindication or no response,<br>use entecavir and tenofovir |
| HBeAg<br>negative | Entecavir and tenofovir are<br>preferred oral agents<br>Use of the other oral reverse<br>transcriptase inhibitors is<br>possible but not preferred | > 1 year             | Preferred if contraindications or no response to INFα              |
|                   | INFα<br>PEG-INFα   | ≥ 1 year             | If contraindication or nonresponse,<br>use entecavir and tenofovir |

#### Nucleoside Analog Adverse Effects

- Class effects
- Rebound hepatitis upon discontinuation
- GI Effects (N/V/D/Abdominal pain)
- HIV resistance
   Lactic Acidosis (rare)
- Lactic Acidosis (rare)
   Reductions in bone mineral density
- Nephrotoxicity (adefovir)
- Telbivudine
   Elevations in CK
- Elevations in CK
   Peripheral neuropathy
- Renally dose all medications

#### Patient Case #8

• What is the most appropriate course of action at this time?

No treatment; Recheck HBV DNA in 6 months

Initiate PEG-IFN + ribavirin

Initiate lamivudine 100 mg/day

Initiate tenofovir 300 mg/day

Handout Page 1-121; Answer Page 1-138

#### Patient Case #9

- HPI: 38 year old male with chronic hepatitis C (genotype 1) currently undergoing treatment Evaluated at 12 week follow up appointment after starting treatment.
- MEDS: Pegylated interferon + ribavirin
- LABS:
   AST 90 IU/ml (350 IU/ml), ALT 64 IU/ml (420 IU/ml)
   HCV RNA 3500 IU/ml (450,000 IU/ml)

Handout Page 1-121; Answer Page 1-138

#### Patient Case #9

- What is the most appropriate course of action at this time?
- A. Discontinue therapy and monitor for symptoms
- B. Continue treatment for an additional 12 weeks
- c. Add boceprevir for an additional 12 weeks
- D. Continue treatment for an additional 72 weeks

Handout Page 1-121; Answer Page 1-138

#### Hepatitis C

- RNA Virus
  - Genotypes 1-6 (1-3 most common is US)
    Several subtypes
  - Genotype 1 most resistant to drug treatment
  - Transfusion, IV drug abuse, transplant
- Major cause of chronic liver disease
   60-80% progression following acute infection
  - □ #1 reason for transplant

#### Treatment of Chronic Hepatitis C

- First line:
  - Genotype 1 :Pegylated interferon + ribavirin + telaprevir OR boceprevir
  - Genotypes 2 and 3 :Pegylated interferon + ribavirin

#### Pegylated Interferon Dosing:

- Degasys: 180ug SQ Weekly
- □ Peg Intron: 1-1.5 ug/kg/week SQ
- Ribavirin orally in 2 divided doses:
- Dose differs based on genotype, weight, and interferon product

#### Direct Acting Antivirals (DAAs)

|                         | Telapravir (Incivek®)   | Boceprevir (Victrelis®)   |
|-------------------------|---|---|
| FDA Approved indication | Chronic HCV therapy<br>(genotype 1) in <u>combination</u><br>with PEG-INf affa and ribavirin<br>in patients with compensated<br>liver disease | <ul> <li>chronic HCV genotype 1 infection,<br/>in combination with peginterferon<br/>alfa and ribavirin, in adult<br/>patients (≥18 years of age) with<br/>compensated liver disease,<br/>including cirrhosis, who are</li> </ul> |
|                         | Not studied in Child-Pugh class     B or C  | previously untreated or who have<br>failed previous interferon and<br>ribavirin therapy.  |

#### Direct Acting Antivirals (DAAs)

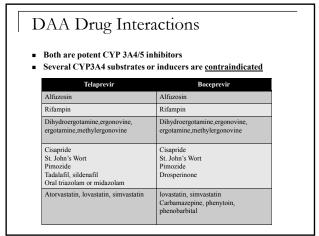
|      |   | Telapravir (Incivek®)   |   | Boceprevir (Victrelis®)  |
|------|---|---|---|--|
| Dose | • | 750 mg three times daily for 12 weeks<br>followed by PEG-INF and ribavirin x<br>12 weeks if undetectable HCV RNA<br>at week 4 and 12. | • | 800 mg orally three times daily starting<br>after 4 weeks of PEG-INF and ribavirin |
|      | • | 375 mg tablets  | · | 200 mg capsules  |
|      | • | Give doses 7-9 hours apart; give with<br>meal that has at least 20 g fat ingested<br>20 minutes prior                                 | • | Give doses 7-9 hours apart; give with meal or light snack                          |
|      | • | Take missed doses if within 4 hours   | • | Take missed doses if within 2 hours  |

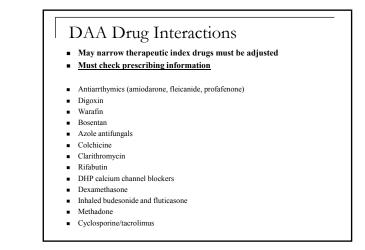
#### DAA Safety

Both contraindicated in pregnancy and in male partners of pregnant women

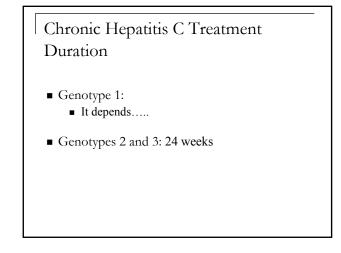
#### Telaprevir

- □ Rash (up to 56%) maculopapular/eczematous
- DRESS, Stevens Johnson Syndrome
- Anemia, pruritis, nausea
- Boceprevir
  - Anemia, neutropenia, fatigue, dysgeusia

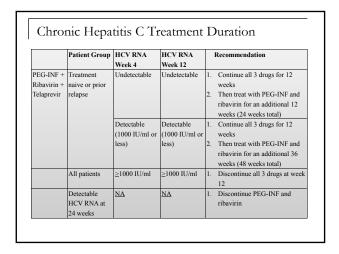


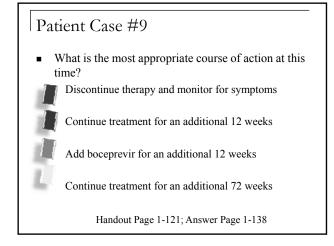


| Parameter                               | Definition  |
|---|---|
| Rapid Virological<br>Response (RVR)     | Negative HCV RNA at week 4 of treatment   |
| Early Virological<br>Response (EVR)     | > 2 log decline in HCV RNA<br>compared to baseline or negative<br>HCV RNA at 12 weeks |
| End of Treatment<br>Response (ETR)      | Negative HCV RNA at the end of a<br>24 or 48 week course depending on<br>genotype     |
| Sustained Virological<br>Response (SVR) | Negative HCV RNA 24 weeks after<br>finishing treatment                                |



| Regimen  | Patient Group   | HCV RNA<br>Week 4 | HCV RNA<br>Week 8  | Recommendation   |
|--|---|-------------------|--|--|
| PEG-INF+                                       | Previously  | Undetectable      | Undetectable   | Continue all 3 drugs for 28 weeks total  |
| Ribavirin +<br>Boceprevir                      | untreated   | Detectable        | Undetectable   | <ol> <li>Continue all 3 drugs for a total of 36<br/>weeks.</li> <li>Then continue PEG-INF and ribavirin for<br/>through week 48</li> </ol> |
| Previous partial<br>responders or<br>relapsers | Previous partial  | Undetectable      | Undetectable   | Continue all 3 drugs for 36 weeks total  |
|  | Detectable  | Undetectable      | <ol> <li>Continue all 3 drugs for a total of 36<br/>weeks.</li> <li>Then continue PEG-INF and ribavirin for<br/>through week 48</li> </ol> |  |
|  | Patients with HCV<br>RNA > 100 IU/ml<br>at week 12 or<br>detectable HCV<br>RNA at week 24 | NA                | NA   | 1. Discontinue all 3 drugs   |

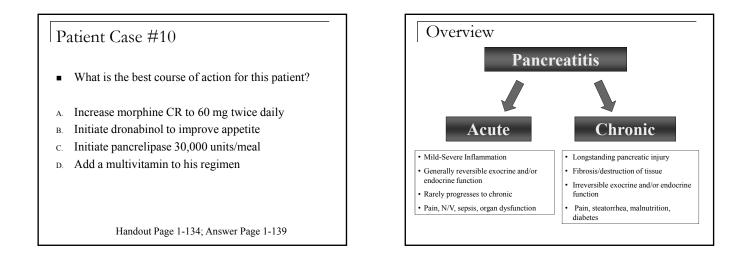




#### Patient Case #10

- HPI: 55 year old man with chronic alcohol abuse and chronic pancreatitis. Steatorrhea and weight loss (now 135 lb)
- LABS: Albumin 2.1 g/dl, Fecal fat 20g/day
- Medications: morphine CR, oxycodone IR as needed

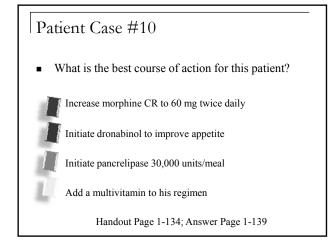
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#### Acute Pancreatitis

- Largely supportive Care
- Pain management
- Antiemetics
- Nutritional support
  - Enteral
  - Hyperglycemia
- Antibiotics
  - □ Infection, abscess, or necrosis

| Complication                     | Targeted<br>Therapies | Comments   |  |
|----------------------------------|-----------------------|--|--|
| Pain                             | Narcotic +/ non-      | <ul> <li>Acetaminophen and/or NSAIDs</li> </ul>  |  |
|                                  | narcotic therapies    | <ul> <li>Long acting narcotic preparations + IR<br/>breakthrough</li> </ul>                  |  |
|                                  | Pancreatic enzymes    | <ul> <li>Caution with acetaminophen and narcotics<br/>if alcohol use is continued</li> </ul> |  |
| Maldigeston and<br>Malabsorption | Pancreatic enzymes    | • Start around 30,000-40,000 lipase units per meal; <sup>1/2</sup> dose for snacks           |  |
|                                  |                       | <ul> <li>Do not crush or chew</li> </ul>   |  |
|                                  |                       | <ul> <li>Max 2500 u/kg/dose; 10,000 u/kg/day</li> </ul>                                      |  |
|                                  |                       | <ul> <li>Titrate to steatorrhea + weight gain</li> </ul>                                     |  |
|                                  |                       | <ul> <li>Porcine based so avoid if pork allergy</li> </ul>                                   |  |
|                                  | Fat soluble vitamins  | • ADEK   |  |
| Diabetes                         | Insulin               | Long acting + short acting   |  |
|                                  |                       | <ul> <li>Oral intake may be variable</li> </ul>  |  |



#### Patient Case #11

- HPI: 32 year old woman with crampy abdominal pain, bloating and constipation for 6 months. Not food related. Diagnosed with IBS-C.
- LABS: within normal limits
- Medications and allergies: none

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#### Patient Case #11

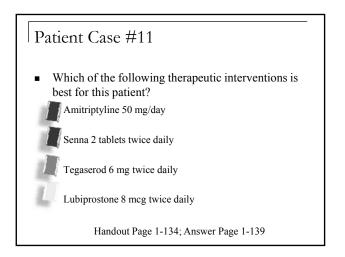
- A. Which of the following therapeutic interventions is best for this patient?
- B. Amitriptyline 50 mg/day
- c. Senna 2 tablets twice daily
- D. Tegaserod 6 mg twice daily
- D. Lubiprostone 8 mcg twice daily

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#### Irritable bowel syndrome

- Categories
  - Diarrhea Predominant (IBS-D)
  - □ Constipation Predominant (IBS-C)
  - Mixed Pattern (IBS-M)
- Features
  - □ Change in frequency and/or stool appearance
  - De Pain, bloating, Relief with defecation
- Target main symptoms and comorbidities

| Therapies   | Comments  |
|---|---|
| Hyoscyamine,<br>dicyclomine   | Target pain due to spasm and also treat diarrhea     Initial or adjunctive therapy for IBS-D or IBS-M     |
| Tricyclic<br>antidepressants  | Target pain and diarrhea     Generally reserved for IBS-D     Low doses                                   |
| SSRIs, SNRIs  | Target pain and often have promotility action in IBS-D     Can also treat comorbid depression and anxiety |
| Lubiprostone  | Indicated for IBS-C in women > 18 years     Main adverse effect is nausea, more expensive option          |
| Loperamide  | Adjunctive for IBS-D, but does not treat pain   |
| Probiotics  | Some potential improvement in global symptoms and pain  |
| Alosetron Indicted for IBS-D in women > 18 years failing other therapies<br>• Must be enrolled in prescribing program<br>• Risk of ischemic colitis |   |
| Tegaserod   | • Indication: IBS-C; available on emergency use only due to CV risk                                       |
| Rifaximin   | Some data to support improvement in bloating  |



#### Patient Case #12

- HPI: 30 year old pregnant woman (14 weeks) with myalgias, watery diarrhea (4-5), vomiting x 1.
- LABS: influenza (-), WBC 8000 x 10<sup>3</sup>
- Medications: prenatal vitamin
- Allergies: none

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#### Patient Case #12

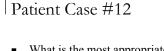
- What is the most appropriate course of action at this time for this patients diarrhea?
- A. Loperamide
- в. Bismuth subsalicylate
- c. Lactase
- D. Pyridoxine

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#### Management of Diarrhea

- Remove correct underlying cause
   Identify drug-induced causes
- Rehydration
  - □ ORS
  - Parenteral
- Dietary modification

| Therapies                                     | Comments   |  |  |
|---|--|--|--|
| Loperamide                                    | • OTC and prescription products, tablet and liquid<br>• OTC indicated in age > 6<br>• Pregnancy category B   |  |  |
| Opioids (diphenoxylate,<br>tincture of opium) | Generally reserved for more severe cases     Increased risk of CNS adverse effects   |  |  |
| Bismuth subsalicylate                         | OTC tablet and liquid preparations     Avoid:         Patients < 12 years of age         Pregnancy         Salicylate allergy         Signs/symptoms of bleeding or mucous     Stool and tongue discoloration     Chelation interactions |  |  |
| Lactase                                       | Suspected or diagnosed lactose intolerance   |  |  |
| Probiotics                                    | Data in AAD, IBD, IBS, radiation induced   |  |  |



• What is the most appropriate course of action at this time for this patients diarrhea?

Loperamide

Bismuth subsalicylate

Lactase

Pyridoxine

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