

## Lecture 13 – DISEASE RELATED DIETS: NUTRITION & DIET THERAPY

1. For each of the following conditions: celiac disease, inflammatory bowel diseases, short-gut syndrome and re-feeding:
  - a. Describe the effects of the condition on dietary intakes & nutrition status.
  - b. List & describe the rationale for specific nutrient needs.
  - c. Identify the goal(s) of nutrition & diet therapy.
  - d. Describe the therapeutic diet & the strategies to support the achievement of the goal(s)
  - e. Describe some patient nutrition education & counselling needs.
  
2. Identify the rationale & indications for vitamin & mineral supplementation in selected diseases/conditions: celiac disease, inflammatory bowel diseases, short-gut syndrome, re-feeding syndrome.
  
3. Identify the indications for nutrition support in the management of inflammatory bowel diseases & short-gut syndrome (enteral nutrition support (EN) & parenteral nutrition support (PN)).

### **Celiac Disease:**

Nutrition Status	<ul style="list-style-type: none"> <li>• Protein – energy malnutrition (BMI, growth)</li> <li>• Anemia – iron, folate and/or vitamin B12 deficiencies</li> <li>• Metabolic bone disease – bone density, serum vitamin D</li> <li>• Abnormal bowel function – acquired lactate deficiency, constipation, diarrhea</li> <li>• Nutrition status doesn't appear to correlate well with degree of villous atrophy</li> </ul>	
Nutrition goals	<ul style="list-style-type: none"> <li>• Reverse malnutrition, optimize nutrition status &amp; restore normal growth</li> <li>• Resolve nutrient deficiencies</li> <li>• Complete healing of the GI mucosa, skin lesions</li> </ul>	
Diet	Recommended dietary pattern	<ul style="list-style-type: none"> <li>• Gluten free for life (should only be initiated after a confirmatory diagnosis)</li> </ul>
	Supplementation	<ul style="list-style-type: none"> <li>• Correct micronutrient deficiencies</li> </ul>
Patient education	<ul style="list-style-type: none"> <li>• Work &amp; effort required to change long-established eating &amp; food selection patterns + ongoing vigilance to maintain a gluten-free diet</li> <li>• Achieving an adequate and balanced dietary intake is challenging               <ul style="list-style-type: none"> <li>○ Most gluten-free grains aren't fortified (iron, thiamine, niacin, riboflavin and folate)</li> <li>○ Tendency to be lower in dietary fibre</li> </ul> </li> <li>• Significant number experience difficulty purchasing &amp; preparing gluten-free foods, eating with family &amp; friends, in restaurants, school, work or during travel</li> <li>• Relief &amp; acceptance of diagnosis, but also frustration &amp; sense of isolation (appears to improve over time)</li> <li>• Social, peer &amp; NPOs (ex// Canadian Celiac Association) have considerable influence on successful self-management</li> </ul>	

### **Inflammatory bowel disease:**

Nutrition Status	<ul style="list-style-type: none"> <li>• Malnutrition affects 20 – 85% (weight loss, growth failure, loss of appetite)</li> <li>• Malabsorption (esp. Crohn's) – vitamin B12, vitamin A, zinc, magnesium</li> <li>• Decreased dietary intake               <ul style="list-style-type: none"> <li>○ Increased needs (energy, protein, micronutrients – esp. folate, calcium)</li> <li>○ Increased nutrient losses (iron)</li> </ul> </li> <li>• Food intolerance, food aversions, self-imposed dietary restrictions</li> </ul>	
Nutrition goals	<ul style="list-style-type: none"> <li>• Reverse malnutrition, optimize nutrition status &amp; restore normal growth</li> <li>• Resolve nutrient deficiencies</li> <li>• Induce remission, prevent flare</li> </ul>	

Diet	Recommended diet during flare	<ul style="list-style-type: none"> <li>• Adequate energy, protein, vitamins &amp; minerals (some above DRIs) from a variety of foods (including B-carotene)</li> <li>• Reduced dietary fibre (esp. insoluble)</li> <li>• Frequent meals &amp; snacks</li> <li>• Limit sugar &amp; dietary fats (individualize)</li> <li>• Separate solids from fluids (individualize)</li> <li>• Lactose restriction (individualize)</li> </ul>
	Supplementation	<ul style="list-style-type: none"> <li>• Iron, calcium, vit. D, vit. B12, folate, zinc, magnesium</li> <li>• Probiotics (strain specific) – possible role in shortening time to &amp; maintaining remission (esp. in Ulcerative Colitis – UC)</li> <li>• Psyllium – possible benefit for maintenance of remission in UC</li> </ul>
Nutrition Support	EN (oral, polymeric)	<ul style="list-style-type: none"> <li>• To address inadequate macronutrient intakes from food</li> <li>• Crohn's: taken to meet 50% of energy needs; shortens time to remission (esp. children)</li> </ul>
	PN	<ul style="list-style-type: none"> <li>• For achievement of gut rest &amp; restoration of nutrition status in Crohn's with complicated fistulas or post-operative complications</li> </ul>
Patient Education		<ul style="list-style-type: none"> <li>• Disease/illness knowledge to promote self-management <ul style="list-style-type: none"> <li>○ Adequate, balanced dietary intakes +/- supplementation</li> </ul> </li> <li>• Optimal health-related quality of life <ul style="list-style-type: none"> <li>○ Avoidance of unnecessary dietary restrictions (2/3 pt restrict food during remission)</li> </ul> </li> <li>• Crohn's &amp; Colitis Canada</li> </ul>

### Short gut syndrome

Nutrition Status		<ul style="list-style-type: none"> <li>• Post-operative dependence on PN for adequate macro/micro-nutrient &amp; hydration <ul style="list-style-type: none"> <li>○ Colon resection – greater risk of dehydration due to high volume GI fluid losses</li> <li>○ Small bowel resection – greater risk of inadequate macronutrient absorption &amp; energy intake due to fat malabsorption, mono- &amp; di- saccharide malabsorption</li> </ul> </li> <li>• Vitamin B12 and essential fatty acid deficiency</li> <li>• Osteoporosis</li> </ul>
Nutrition goals		<ul style="list-style-type: none"> <li>• Reverse malnutrition, optimize nutrition status &amp; normalize normal growth</li> <li>• Support adequate hydration</li> <li>• Augment intestinal adaptation</li> </ul>
Diet	Recommended dietary pattern	<ul style="list-style-type: none"> <li>• Multiple (small) meals per day – goal 50% more than estimated energy needs</li> <li>• Emphasis on complex carbohydrates</li> <li>• Limit sugars</li> <li>• Dietary fats should not be restricted (emphasize essential FAs) <ul style="list-style-type: none"> <li>○ Unless colon is intact – then moderate to control steatorrhea to minimize fluid, electrolyte losses</li> </ul> </li> <li>• Emphasis on soluble fibre – up to 1000 kcal derived from SCFA produced by bacterial fermentation; increase transit time</li> </ul>
	Supplementation	<ul style="list-style-type: none"> <li>• Protein supplementation generally not required</li> <li>• Oral rehydration solutions</li> <li>• Oral vitamin &amp; mineral supplementation (above DRIs) <ul style="list-style-type: none"> <li>○ Magnesium &amp; zinc due to high output</li> <li>○ Calcium citrate to minimize risk of kidney stone formation</li> <li>○ Vitamin ADEK</li> <li>○ Intramuscular vit B12 (ileal resections)</li> <li>○ Thiamine</li> </ul> </li> </ul>
Nutrition Support	EN	<ul style="list-style-type: none"> <li>• Continuous tube feeding may augment rate of intestinal adaptation</li> <li>• Usually polymeric unless malabsorption is severe</li> </ul>
	PN	<ul style="list-style-type: none"> <li>• To restore &amp; maintain adequate nutrition status &amp; hydration, until sufficient bowel adaptation has occurred to enable maintenance of adequate nutrition status</li> </ul>

Patient Education	<ul style="list-style-type: none"> <li>• Strong knowledge of personal gastrointestinal structure &amp; function</li> <li>• Strategies to support optimal nutrient status; maintain normal hydration; intestinal adaptation</li> <li>• Optimal health-related quality of life <ul style="list-style-type: none"> <li>○ Poor sleep – frequent night time urination, bowel movements</li> <li>○ PN home management</li> <li>○ Very high “health work” load</li> </ul> </li> </ul>
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**Re-feeding Syndrome:**

Nutrition Status	<ul style="list-style-type: none"> <li>• Under-recognized complication of nutrition support among hospitalized patients <ul style="list-style-type: none"> <li>○ Under-recognition of malnutrition in the acute care setting</li> <li>○ Lack of universally accepted definition and means for detecting risk</li> <li>○ A carefully conducted nutrition assessment will detect under-nutrition, including increased risk of refeeding syndrome due to underlying disease, illness, and therapies</li> </ul> </li> <li>• Malnutrition in acute-care is costly, delays illness recover, and causes re-feeding complications</li> </ul>	
Nutrition goals	<ul style="list-style-type: none"> <li>• Nutritional repletion</li> <li>• Prevention of re-feeding complications <ul style="list-style-type: none"> <li>○ Malnourished patients do NOT tolerate over-zealous re-feeding!</li> <li>○ Nutritional rehabilitation and the correction of deficiencies takes time</li> <li>○ Failure to detect S/S can lead to fatality</li> </ul> </li> <li>• Gradual, slow incremental increase in the indicated EN/PN according to tolerance (highly individualized)</li> </ul>	
Diet	Recommended dietary pattern	<ul style="list-style-type: none"> <li>• Minimal or no nutritional intake for &gt; 5 days: begin nutrition support at ≤ 50% of goal energy intake <ul style="list-style-type: none"> <li>○ Patients at high-risk of re-feeding syndrome: 10 kcal/kg/24h</li> <li>○ Severe malnutrition (BMI ≤ 14): 5 kcal/kg/24h</li> </ul> </li> </ul>
	Supplementation	<ul style="list-style-type: none"> <li>• Vitamins – begin before &amp; continue for first 10 days of re-feeding</li> <li>• Minerals – replete low levels before initiating nutrition support</li> <li>• Fluid – restore &amp; maintain adequate fluid volume</li> </ul>

4. List some methods for assessing & monitoring nutrition status among patients receiving EN or PN.

- Nutrition status indicators: macro, micro & hydration status
- Fluid overload
- Tolerance/acceptability

5. Explain the risks and benefits of popular diets:

a. Gluten-free diet (in the absence of celiac disease)

???

b. Low-FODMAPS (Fermentable carbohydrates; oligo-, di-, mono-saccharides and polyols)

- Small, short duration, prospective & randomized trials – low FODMAPs may reduce sx of irritable bowel syndrome
- Meta-analysis – efficacy in management of functional GI sx (of irritable bowel syndrome)
- No long-term studies on efficacy or safety
  - Reduced bacteria count (bifidobacteria)

**LECTURE 14 – DISEASE-RELATED DIETS; NUTRITION & DIET THERAPY PART 2**

1. For each of the following conditions: heart failure and renal disease/dysfunction:
  - a. Describe the effects of the condition on dietary intakes & nutrition status.
  - b. List & describe the rationale for specific nutrient needs.
  - c. Identify the goal(s) of nutrition & diet therapy.
  - d. Describe the therapeutic diet & the strategies to support the achievement of the goal(s)
  - e. Describe some patient nutrition education & counselling needs.

*Answered below under sub-topics.*

**Heart Failure:**

2. Discuss key elements of the nutrition assessment that are focused on for heart failure patients.

Anthropometric	<ul style="list-style-type: none"> <li>• Weights – UBW, IBW, % losses/gain, weight history, fluid weight vs. lean mass, BMI</li> <li>• Patients weigh themselves daily</li> </ul>
Biochemical	<ul style="list-style-type: none"> <li>• Blood work: lipids, FBG, HbA1c, CRP, pre-albumin, albumin, electrolytes, Mg, PO<sub>4</sub>, Ca, iron, Hb</li> <li>• Stool tests – C. diff</li> <li>• Bone density</li> </ul>
Clinical	<ul style="list-style-type: none"> <li>• BP, EF%, NHYA class</li> <li>• Physical activity, smoking, alcohol/drug, family hx</li> <li>• Number of HF hospital admissions, meds, comorbidities</li> </ul>
Diet-related hx	<ul style="list-style-type: none"> <li>• FFQ, 3 day food record, 24 hour recall, previous knowledge</li> <li>• Goals, beliefs, stage of change, motivations, ability</li> </ul>

3. What are key nutrients to consider for heart failure management.

Sodium	<ul style="list-style-type: none"> <li>• Inadequate blood flow to kidneys → aldosterone &amp; ADH secretion → fluid (&amp; Na) retention</li> <li>• Sodium restriction: 2 g/day</li> </ul>
Magnesium	<ul style="list-style-type: none"> <li>• Deficiency common as a result of poor dietary intake &amp; diuretics (increase Mg excretion)               <ul style="list-style-type: none"> <li>○ Aggravates changes in electrolyte balance - +ve sodium and -ve potassium balance</li> </ul> </li> <li>• Should be measured in HF patients</li> </ul>
Thiamin	<ul style="list-style-type: none"> <li>• At risk of thiamin deficiency because of poor food intake, use of loop diuretics &amp; advanced age</li> <li>• Thiamin supplementation can improve LVEF and symptoms</li> </ul>
Iron	

4. Understand the current lifestyle management recommendations for persons with heart failure.

- Sodium restriction to 2g/d
- Fluid restriction: 1.5 – 2 L/d

5. Understand nutrient interactions and food considerations with HF medications.

6. Describe counselling tips to help with patient education.

Sodium restriction	<ul style="list-style-type: none"> <li>• Identify sources of sodium and to choose lower sodium options</li> <li>• Read labels for sodium content in foods</li> <li>• Write down food record for analysis</li> <li>• Phone apps; handouts</li> <li>• Decrease processed foods; reduce sodium while eating out; prepare low sodium food at home               <ul style="list-style-type: none"> <li>○ Low sodium meal delivery – <i>Better Meals</i></li> </ul> </li> <li>• If finances are an issue and patient on PWD, apply for funding</li> </ul>
Fluid restriction	<ul style="list-style-type: none"> <li>• Monitor all fluids (anything liquid at room temperature) throughout the day</li> <li>• Teachings around what is a fluid, counting fruits, helping to manage thirst</li> </ul>

7. Understand what cardiac cachexia is, diagnosis and current management recommendations.

Definitions	<ul style="list-style-type: none"> <li>• Multi-factorial disease affecting 5-15% of HF patients</li> <li>• Characterized by protein-calorie malnutrition with muscle wasting &amp; peripheral edema</li> <li>• Complex imbalance of catabolic &amp; anabolic body systems             <ul style="list-style-type: none"> <li>○ Caused by a series of immunological, metabolic and neuro-hormonal processes</li> <li>○ Mediated by sympathetic nervous system activation, pro-inflammatory cytokines, ATII, glucocorticoids and TGF-B</li> </ul> </li> </ul>
Diagnosis	<ul style="list-style-type: none"> <li>• 5-6% edema-free weight loss in <math>\geq 6</math> months</li> <li>• SGA and MUST assessment tools for malnutrition not sensitive enough or validated for cachexia</li> </ul>
Management	<ul style="list-style-type: none"> <li>• High protein, high calorie, low sodium             <ul style="list-style-type: none"> <li>○ Energy: 30 – 35 kcal/kg/d or 1.3 – 1.5 REE</li> <li>○ Protein: 1.3 – 1.5 g/kg/d</li> </ul> </li> <li>• Optimize nutrient intake to prevent further muscle/weight loss             <ul style="list-style-type: none"> <li>○ High calorie, high protein diet</li> <li>○ Size &amp; frequency of meals</li> <li>○ Supplement diet with multivitamin</li> <li>○ Assess for deficiencies</li> <li>○ Decrease in pro-inflammatory lifestyle factors</li> <li>○ Possible tube feed or TPN to supplement</li> </ul> </li> </ul>

8. Understand nutrition management for a person with a Ventricular Assist Device (VAD).

- Adequate protein intake
- Anorexia/cachexia
- Early satiety
- Fluid retention or dehydration
- Nutrition post-surgery (wound management)
- Delayed gastric emptying
- Nausea
- Chronic inflammation (CRP levels)
- Stress – GERD, indigestion
- Warfarin & vit K – consistency!!

9. Describe the nutrition considerations for post-heart transplant care.

- Malnourished?
- Medication interactions
- Food safety
- Blood sugar control
- Fluid/salt directly post-transplant
- K, Mg, Ca (bone health), Zn, supplements
- Protein wk 4-6: 1.3 – 1.5 g/kg
  - > 6 wk: 1 – 1.3 g/kg
- Weight management
- Cholesterol management

10. Understand what cardiac allograft vasculopathy (CAV) is, and lifestyle prevention.

Definition	<ul style="list-style-type: none"> <li>• Accelerated form of CAD</li> <li>• Major factor in limiting long-term survival post-heart transplant</li> <li>• Characterized by concentric fibrous intimal hyperplasia along the length of coronary vessels</li> <li>• Both immunologic and non-immunologic risk factors contribute to development</li> <li>• Difficult to treat → prevention is critical</li> </ul>
Lifestyle prevention	<ul style="list-style-type: none"> <li>• Obesity, dyslipidemia, diabetes, hypertension, metabolic syndrome             <ul style="list-style-type: none"> <li>○ Hyperlipidemia &amp; insulin resistance most significant non-immunologic factors</li> </ul> </li> </ul>

Goals of Medical Nutrition Therapy:

- Reduced readmission to hospital
- Reduced length of stay
- Improved compliance/adherence to sodium and fluid
- Address nutrient deficiencies
- Address muscle wasting, cachexia
- Improve quality of life

## Renal Disease:

### 11. What are 4 major manifestations of renal disease (brief overview)?

- Kidney stones; acute kidney injury; chronic kidney disease; end-stage renal disease

### 12. Understand what kidney stones are the most common kidney stones (calcium oxalate stones) medical nutrition therapy.

Kidney stones	<ul style="list-style-type: none"> <li>• Calcium stones are most common: calcium oxalate (60%); calcium phosphate (10%); 10% both</li> <li>• Uric acid (5-10%); struvite (5-10%); cystine (1%)</li> </ul>
MNT for calcium oxalate stones	<ul style="list-style-type: none"> <li>• Drinking adequate fluids</li> <li>• Reducing sodium</li> <li>• Reducing animal protein (meat, eggs, fish)</li> <li>• Getting enough calcium from food or taking calcium supplements with food</li> <li>• Avoid foods high in oxalate (spinach, rhubarb, nuts, wheat bran)</li> <li>• Vitamin D supplements not recommended</li> </ul>

### 13. Understand what acute kidney injury is and medical nutrition therapy. What are the key nutrients to monitor?

AKI	<ul style="list-style-type: none"> <li>• Sudden reduction in GFR; amount of filtrate per unit in nephrons</li> <li>• Altered ability of kidney to excrete the daily production of metabolic waste</li> </ul>
Causes	<ul style="list-style-type: none"> <li>• Prerenal – inadequate renal perfusion</li> <li>• Intrinsic – diseases within the renal parenchyma</li> <li>• Postrenal – urinary tract obstruction</li> </ul>
MNT	<ul style="list-style-type: none"> <li>• Issue of balancing protein &amp; energy needs with treatment of acidosis and excessive nitrogenous waste is complex <ul style="list-style-type: none"> <li>○ Protein amount is influenced by underlying cause &amp; presence of other conditions <ul style="list-style-type: none"> <li>▪ Non-dialysis: 0.5 – 0.8 g/kg; dialysis: 1-2 g/kg; CCRT: 1.5 – 2.5 g/kg</li> <li>▪ When stable: 0.8 – 1 g/kg → 60 – 70% high biological value (HBV)</li> </ul> </li> </ul> </li> <li>• Other nutrients: sodium, potassium</li> <li>• Mortality rate increases in malnourished patients</li> </ul>

### 14. What are the stages of chronic kidney disease (CKD)?

1	90-130 mL/min	Kidney damage but normal to increased kidney function
2	60-89 mL/min	Mild decrease in function
3	30-59 mL/min	Moderate decrease in function
4	15-29 mL/min	Severe decrease in function
5	<15 mL/min	Kidney failure with treatment necessary (ESRD)

#### a. Understand what different nutritional therapy may be proposed at each stage.

Goals of MNT	<ul style="list-style-type: none"> <li>• Manage sx (edema, hypoalbuminemia, hyperlipidemia)</li> <li>• Decrease risk of progression to renal failure</li> <li>• Maintain nutritional stores</li> <li>• Each stage of CKD has different therapy</li> </ul>
Protein reduction	<ul style="list-style-type: none"> <li>• Reduction to 0.8 g/kg/day may decrease proteinuria without adversely affecting albumin <ul style="list-style-type: none"> <li>○ 60% from HBV</li> </ul> </li> <li>• Preserves renal function in mild-mod renal failure; should restrict if GFR &lt;30</li> </ul>
Sodium	<ul style="list-style-type: none"> <li>• 2-3 g/day</li> </ul>
Potassium	<ul style="list-style-type: none"> <li>• Monitored especially in stage 4 CKD; may need to be reduced/restricted</li> </ul>
Phosphorus	<ul style="list-style-type: none"> <li>• Serum PO<sub>4</sub> elevates as GFR decreases <ul style="list-style-type: none"> <li>○ Early initiation of PO<sub>4</sub> reduction therapies is advantageous for delaying hyperparathyroidism and bone disease</li> <li>○ GFR &lt;60 should be monitored</li> </ul> </li> <li>• Phosphate binders may be needed (calcium carbonate or acetate, sevelamer, lanthanum)</li> </ul>
Vitamins	<ul style="list-style-type: none"> <li>• Multivitamin (with water soluble vitamins) – custom supplement for renal pts</li> </ul>

15. Understand what cardio-renal syndrome is.

- Cardiorenal syndrome = bidirectional interaction between kidney disease and heart disease
  - Mortality increased in pts with HF who have reduced GFR
  - CKD patients are at risk of atherosclerotic CVD & HF
  - Acute or chronic systemic disorders can cause both cardiac & renal dysfunction
- CRS defined in National Heart, Lung & Blood Institute = therapy to relieve congestive sx of HF is limited by a decline in renal function manifested by reduced GFR

16. What is end stage renal disease (ESRD)?

- ESRD = inability to excrete waste products, maintain fluid & electrolyte balance and produce hormones
  - Circulating nitrogenous waste → uremia (malaise, weakness, N, V, muscle cramps, itching, metallic taste, neurologic impairment)
- 90% of ESRD patients have chronic DM, HTN, and/or glomerulonephritis

a. What are the different options for dialysis (review)?

Hemodialysis	<ul style="list-style-type: none"> <li>• Fluid &amp; electrolyte content is similar to normal plasma</li> <li>• Waste products &amp; electrolytes move by diffusion, ultrafiltration and osmosis from the blood into the dialysate, and then removed</li> <li>• 3-5 hours 3x per week</li> </ul>
Peritoneal dialysis	<ul style="list-style-type: none"> <li>• Makes use of body's own semipermeable membrane – peritoneum</li> <li>• High dextrose dialysate instilled into peritoneum, diffusion carries waste products from blood through peritoneal membrane and into dialysate, water moves by osmosis</li> <li>• Tissue weight gain can occur due to absorbing 400-800 cal/d from glucose in the dialysate</li> </ul>

17. Understand the goals of medical nutrition therapy in the management of ESRD.

- Prevent deficiency and maintain good nutrition status
- Control edema and electrolyte imbalance by controlling sodium, potassium and fluid intake
- Prevent or slow the development of renal osteodystrophy by controlling Ca, PO<sub>4</sub>, vit D and PTH
- Enable patient to enjoy their food as much as possible with the restrictions
- Coordinate patient with good care team management
- Provide ongoing counseling and long-term monitoring

a. Understand what nutrients are monitored for nutrition intervention.

Protein	<ul style="list-style-type: none"> <li>• Dialysis is a drain on body protein → protein intake must be increased               <ul style="list-style-type: none"> <li>○ PD: 1.2 – 1.5 g/kg/d (at least 50% from HBV)                   <ul style="list-style-type: none"> <li>▪ Protein losses of 20-30 g/24h occur in PD (average 1g/hr)</li> </ul> </li> <li>○ HD 3x/wk: 1.2 g/kg/d</li> </ul> </li> <li>• Patients on dialysis with low albumin levels have much higher mortality rates</li> <li>• Serum BUN, creatinine levels, uremic symptoms and weight should be monitored               <ul style="list-style-type: none"> <li>○ Albumin more routinely used as nutritional marker rather than pre-albumin (which is metabolized by kidney so is routinely elevated in renal failure)</li> </ul> </li> </ul>
Fluid/sodium	<ul style="list-style-type: none"> <li>• Assessed with looking at weight, edema, serum sodium level, dietary intake &amp; BP</li> </ul>
Phosphorus	<ul style="list-style-type: none"> <li>• Large molecular weight = not easily removed by dialysis → restriction &amp; binders required               <ul style="list-style-type: none"> <li>○ NOTE: food sources are about 60% absorbed; additives can be 100% absorbed</li> </ul> </li> <li>• SEs of binders: GI distress, diarrhea, gas, severe constipation/impaction</li> <li>• Phosphate restriction MAY need to be lifted at times to allow more dairy products to meet protein needs</li> </ul>
Potassium	<ul style="list-style-type: none"> <li>• Usually needs restriction</li> </ul>

Calcium and PTH	<ul style="list-style-type: none"> <li>Balance between phosphate and calcium is difficult for body to maintain             <ol style="list-style-type: none"> <li>Decreased ability of kidney to convert vit D to active form</li> <li>Decreased calcium can cause over-secretion of PTH → resorption of bone</li> <li>Need for serum calcium increases as serum phosphate increases</li> </ol> </li> </ul>
Iron & EPO	<ul style="list-style-type: none"> <li>Anemia caused by inability of kidney to produce EPO (stimulates bone to produce RBCs)             <ul style="list-style-type: none"> <li>Synthetic form of EPO used to treat anemia</li> <li>Iron IV supplementation also used when hematocrit rises (oral iron not effective)</li> </ul> </li> <li>Serum ferritin = accurate indicator of iron status in renal failure</li> </ul>
Vitamins	<ul style="list-style-type: none"> <li>Water soluble vitamins are rapidly lost during dialysis; also PO<sub>4</sub> and K restriction             <ul style="list-style-type: none"> <li>Require supplementation</li> </ul> </li> </ul>

**LECTURE 19 – DISEASE RELATED DIETS: NUTRITION & DIET THERAPY PART 3**

**Diabetes:**

**1. Understand what prediabetes is and what are at least 3 ways lifestyle can play a role to prevent diabetes.**

Pre-diabetes	<ul style="list-style-type: none"> <li>BG higher than normal; Hb 5.7 – 6.4%</li> <li>50% will develop DM2</li> </ul>
Lifestyle	<ul style="list-style-type: none"> <li>Clinical trials comparing lifestyle interventions to control group reported RRR 29-67% for DM2</li> </ul>
MNT goals	<ul style="list-style-type: none"> <li>Healthy food choices for moderate weight loss (5%)             <ul style="list-style-type: none"> <li>Obesity has an impact on insulin sensitivity</li> </ul> </li> <li>Encouraging fiber intake improves insulin sensitivity</li> <li>Meal patterns &amp; portion sizes</li> </ul>

**2. Understand what role carbohydrates play in diabetes (DM) management.**

- Blood glucose levels after eating are primarily determined by rate of appearance of glucose from carb digestion & absorption into blood stream + ability of insulin to clear glucose from circulation
- Balancing amount of carbs, overall nutrition & BG levels
  - Day-to-day consistency important, especially if on MNT alone, glucose lowering meds, or fixed insulin regimens
- In persons with DM1 or DM2 who adjust meal-time insulin, or are on insulin pump therapy, insulin can be adjusted to match carb intake (insulin:carbohydrate ratios)

**3. Understand 2 key strategies to monitoring carbohydrates and overall meal balance for DM management.**

**a. Plate method and carb counting**

Carb counting	Plate method
<ul style="list-style-type: none"> <li>Food portions contributing 15 g of carbs = 1 carb serving</li> </ul>	<ul style="list-style-type: none"> <li>½ vegetables (at least 2 kinds)</li> <li>¼ starch and ¼ protein</li> </ul>

**b. Understand when and why each may be used**

- Need to determine what the patient is able to handle
- What variation in their diet and eating patterns do they have?
- What might give the person the most success with their blood sugar levels?

**4. Understand how the Glycemic index is used and what it is.**

- Developed to compare the physiologic effects of carbs on glucose
- Low GI foods often increase satiety, which may also decrease the amount one consumes
  - Limitation: variability of response to a specific carb food
  - Note: if a carb food is combined with fats & protein, it changes how fast that food is absorbed
- Glycemic load is also important



5. Understand carbohydrate counting and how to apply it to a meal plan.

- Insulin:carbohydrate ratios discussed with RD, RN and endocrinologist → allows for self adjustment
- Patients often keep a cheat sheet of how much carbs different food servings have
- Number of carb servings/meal depends on goals for a patient, and whether a patient is sedentary or active
  - Typical amounts: 2-5 servings
  - Most women/less active men = 3-4 servings (45-60 g)
  - Most men/more active women = 4-5 servings (60-75 g)
  - If weight loss is a goal, this may be adjusted safely with patient by RD
- Servings include dairy/alt, fruits/veggies, starches

6. Understand how fibre is accounted for on a label with carbohydrate counting.

- Fibre doesn't raise blood glucose levels (therefore subtract from carbohydrate counting)
  - Ex// whole wheat bread = 36 g carbs / 2 slices; fiber = 6g → 30 g carb / 2 slices = 15 g carb / slice

7. Understand how sugar alcohols are accounted for in carbohydrates/calories.

- Only partly absorbed, have fewer calories than sugar and have no major effect on blood glucose
- >10 g/day → SEs (gas, bloating, diarrhea)

8. Be familiar with different types of sugar substitutes and their role for diabetic patients.

- Acesulfame potassium, aspartame, cyclamate, saccharin, sucralose, steviol glycosides
- Evidence for sugar substitutes in weight loss is mixed, possible reasons cited are:
  - Sugar substitutes are sweet and increase preference for sweet foods
  - Hormones not released to tell body it is full and could lead to increased calories later
  - Psychological – those that knew they were eating sugar substitutes tended to compensate by increasing calories other ways, while those that didn't know tended to decrease calories overall
  - More studies needed to look at long-term weight loss

9. Understand what are the protein and dietary fat recommendations for DM management

Protein	<ul style="list-style-type: none"> <li>• Minimal acute effects on glycemic response &amp; no long-term effect on insulin requirements</li> <li>• Usual recommendation: 15 – 20% of total calories                             <ul style="list-style-type: none"> <li>○ Including protein with each meal can help patients with satiety and portion sizes</li> </ul> </li> <li>• Short-term, small subject numbers: protein &gt; 20% of total energy may improve glucose &amp; insulin concentrations, reduce appetite and improve satiety → assist with weight loss for some pts</li> </ul>
Dietary Fat	<ul style="list-style-type: none"> <li>• 2-3 servings of fish per week recommended (omega 3 content)</li> <li>• If diet = 20% protein, 45-50% carbs, then fat is 25-30%                             <ul style="list-style-type: none"> <li>○ Canadian Cardiovascular Society: 9% saturated (7% if dyslipidemia), therefore 21-23% unsaturated (MUFA and PUFA)</li> </ul> </li> <li>• Encouraging more unsaturated fats (unsalted nuts &amp; seeds, avocado, fish, extra virgin olive oil) can be beneficial for satiety at meals, nutrient absorption and CV health</li> </ul>
Ketogenic Diet	<ul style="list-style-type: none"> <li>• Diet 10% carbs, 70% fat, 20% protein</li> <li>• Current studies are short-term, small number of subjects</li> </ul>

10. Understand what needs to be considered for DM patients and exercise.

Hypoglycemia	<ul style="list-style-type: none"> <li>• Concern for DM1, and to a lesser extent in DM2 using insulin or insulin secretagogues</li> <li>• If pre-exercise BG levels &lt;5.5 mmol/L → ingest 15 – 30 g carb before exercise                             <ul style="list-style-type: none"> <li>○ Amount depends on injected insulin, exercise duration &amp; intensity, results of BG monitoring</li> </ul> </li> <li>• If diabetes is controlled by lifestyle or oral hypoglycemic agents that don't increase insulin levels, risk of developing hypoglycemia during exercise is minimal                             <ul style="list-style-type: none"> <li>○ Most will not need to monitor BG levels or require supplements with carbs for exercise &lt;1h</li> </ul> </li> </ul>
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Hyperglycemia	<ul style="list-style-type: none"> <li>• In DM1 who are severely insulin deficient (ex// insulin omission, illness) hyperglycemia can be worsened by exercise</li> <li>• Blood or urine ketone bodies can be tested to assess best course of action</li> <li>• Important to ensure proper hydration &amp; monitor for S/S, especially for exercise in the heat</li> </ul>
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11. Understand why a whole family approach to counselling is beneficial for children with obesity and/or type 2 diabetes.

- Fostering a healthy relationship with food for the child and the family is important
- Family-based lifestyle interventions with a behavioral component aimed at changes in diet and physical activity patterns have shown to result in significant weight reduction in both children and adolescents

12. Name 5 factors that could increase a child's risk for developing type 2 DM.

- Being overweight or inactive
- Ethnicities at high risk for DM2 (African, Hispanic, Asian, South Asian, Aboriginal)
- Family history of DM2
- Mother with gestational diabetes
- High fat or cholesterol in blood
- High levels of fatty deposits in liver
- High blood pressure
- Polycystic ovarian syndrome
- Certain meds for mental health conditions

13. What are important considerations for children and youth diagnosed with type 1 diabetes.

- Nutrition management of pediatric diabetes needs to consider both optimal growth & development + tight BG control
- Important to note that infants & toddlers often have erratic eating schedules
  - Amount of carbs they eat at a meal may differ (esp. picky eaters)
  - Helping parents & children to understand carb counting is important (in foods AND meal plan)
- Insulin pumps has allowed more flexibility in the diet (pumps covered for pediatric patients)
- Higher risk of eating disorders in youth DM1
  - Often associated with insulin omission to lose weight & impaired metabolic control
  - With increased independence in adolescence (incl. food choices & influences), positive relationship with food is very important
- Celiac disease in 4-9% of children with DM1 (although 60-70% of these children are asymptomatic)

14. List 3 types of supplement beverages that are available for diabetic patients and 2 key characteristics of their components.

- Glucerna bars and beverages
  - Diabetic resources - tetras and tube feed
  - Boost Glucose Control
- ➔ Lower amount of carbohydrates and added sugars, and higher in fibre

**LECTURE 20 - DISEASE RELATED DIETS: NUTRITION & DIET THERAPY**

1. Understand what the current CCS (Canadian Cardiovascular Society) nutrition guidelines are for 2016.

Fat (unsaturated & saturated)	<ul style="list-style-type: none"> <li>• Limit total fat to &lt; 30% of daily energy</li> <li>• Emphasize PUFA (in particular omega-3 from fish) and MUFA</li> <li>• Saturated fats (SFA) &lt; 7% (dyslipidemia) or &lt; 9% (no dyslipidemia) of daily energy</li> </ul>
Dietary patterns	<ul style="list-style-type: none"> <li>• Recommended diets: Mediterranean, Portfolio or DASH diets</li> </ul>
Cholesterol	<ul style="list-style-type: none"> <li>• Dyslipidemia or high risk of CVD: limit cholesterol to 200 mg daily</li> <li>• Others: no specific restricted</li> </ul>
Key foods to lower cholesterol	<ul style="list-style-type: none"> <li>• Increase phytoestersols, soluble fiber, soy, nut intake</li> </ul>
Other	<ul style="list-style-type: none"> <li>• Increase fiber to &gt; 30 g</li> <li>• Limit alcohol to ≤ 30 g (1-2 drinks)</li> <li>• Smoking cessation</li> <li>• Engage in regular activity</li> </ul>

2. The World Health Organization's recommendations for added sugar.

- < 5-10% added sugar per day

3. What are 3 ways diet can affect heart health and what can they modify?

Diet can modify..	By affecting ...
Lipid concentration	<ul style="list-style-type: none"> <li>• Cholesterol absorption</li> <li>• Chylomicron and VLDL production</li> <li>• Regulation of LDL receptors</li> </ul>
Progression of CVD	<ul style="list-style-type: none"> <li>• Oxidation of LDL cholesterol</li> <li>• Thrombogenesis</li> <li>• LDL particle size</li> </ul>
Plays a role in inflammation and oxidative stress	

4. Understand the importance of what is replacing saturated fats or other nutrients in decreasing risk.

SFA controversy	<ul style="list-style-type: none"> <li>• Believed that if SFA increased total cholesterol, they must increase heart disease risk</li> </ul>
Evidence of dietary changes	<ul style="list-style-type: none"> <li>• Replacing SFA with carbs (particularly refined carbs) <u>exacerbated</u> atherogenic dyslipidemia</li> <li>• Replacement of SFA with PUFA reduces CHD Risk</li> </ul>
Questions to consider	<ul style="list-style-type: none"> <li>• What are we replacing the SFA or carbs with?</li> <li>• How does changing macronutrient percentages in our diet affect health?</li> <li>• What about dairy SFA vs. meat/poultry SFA?</li> <li>• What are other factors from diet affecting heart disease? Serum cholesterol profiles are an important risk factor but there are others.</li> <li>• Is it possible that fats express differently in the context of other components of a diet?</li> </ul>
Current recommendations	<ul style="list-style-type: none"> <li>• Fat 20-35%    CHO 45-65%    Protein 10-35%</li> <li>• Linoleic acid (omega-6) 5-10%    alpha Linolenic acid (ALA) (omega 3) 0.6 - 1.2%</li> </ul>

5. Understand why a balance of omega 3s and omega 6s are important in the PUFA intake profile and why it is important in heart health.

- When unsaturated fats are substituted for SFA, it reduces LDL cholesterol by up-regulating LDL receptors
- Increased PUFA intake relative to saturated fats results in a decrease in Apo B production → reduce CHD risk
  - HOWEVER, need to consider the balance between omega-3 and omega-6 because studies found increased risk of death with replacing SFA with mostly omega-6

6. Understand the role of omega 3 supplements.

- Benefit for very high triglyceride levels
  - 2-4 g of fish oil lowered high TG levels by 25-30%

7. List 4 types of saturated fats and what type affects lipids more favourably.

Lauric acid (C-12)	Coconut oil, palm oil	Increases LDL cholesterol by down-regulating LDL receptors
Myristic (C-14)	Coconut oil, dairy, palm oil	
Palmitic (C-16)	Palm oil, meat, cheese	
Stearic (C-18)	Coco butter, meat	Foods higher in stearic acid appear to affect lipids more favourably

8. Be able to calculate how much fat in grams is 7% (dyslipidemia) or 9% (no dyslipidemia) or 30% (total fat) of a pt's diet.

- 1 tsp = 4 g of fat = 36 kcal
- Ex// total daily intake 1500 kcal = 166 g fat → total fat (30%) = 50 g; SFA (7%) = 11 g; UFA (total - SFA) = 39 g
  - For most people SFA < 20 g/d

9. Be able to give examples of two types of foods that are high in monounsaturated fats and polyunsaturated fats.

- **MUFA:** olive oil, canola, avocado, nuts
- **PUFA:** essential fatty acids (omega 3 and 6); safflower, sunflower, corn, soybean oils; nuts, seeds; fatty fish

10. Understand the pros and cons of coconut oil for heart health.

Pros	Cons
<ul style="list-style-type: none"> <li>• Solid at room temperature</li> <li>• Good shelf-life</li> <li>• Moderate heat point</li> </ul>	<ul style="list-style-type: none"> <li>• Found to raise total LDL to a greater extent than unsaturated oils BUT lesser extent than butter</li> <li>• Need to consider dietary pattern and if this is the primary fat source or main SFA source</li> </ul>
RECOMMENDATION: use a variety of oils; consider total SFA in day + balance of PUFAs and MUFAs	

11. Understand what smoke point is and its importance in choosing a vegetable oil to use.

- **Smoke point** = temperature at which an oil starts to burn and smoke (often more refined = higher smoke point)
  - Phytochemicals can be destroyed and can cause free radicals
- High to low smoke point: avocado > grapeseed > canola (refined) > butter > EVO > flaxseed

12. Be able to give 3 practical tips in regards to fat intake.

- Trim fat off meats, remove skin off chicken; marinating longer can increase meat tenderness and flavour
- Limit butter intake and replace with unsaturated fats (avocado, unsalted nut butters, non-hydrogenated margarines, olive oil and other vegetable oils)
- Use unsalted nuts as snacks (up to ¼ cup per day)
- Consume fish 2-3 times per week and eat vegetarian at least 1 time per week

13. Understand the role carbohydrates plays in heart health and CCS recommendations.

- Low GI dietary patterns recommended
- If saturated fats are replaced with MUFAs and carbs, then people should choose plant sources of MUFAs (olive oil, canola oil, nuts and seeds) and high-quality sources of carbs (whole grains and low GI carbs)

14. What are the 3 key dietary patterns recommended by the CCS?

Mediterranean Diet	<ul style="list-style-type: none"> <li>• Shown to improve lipid profiles and reduce CVD mortality</li> <li>• Servings/day: veggies 2-4 cups; fruits 1-2 cups; dairy 1-3; whole grains/starch veg 4-6; nuts/legumes 1-3</li> </ul>
Portfolio Diet	<ul style="list-style-type: none"> <li>• 4 dietary components: nuts 30 g; soluble fiber &gt; 10 g; soy 25 g; phytosterols 2 g                             <ul style="list-style-type: none"> <li>○ Each have been shown independently to lower cholesterol</li> </ul> </li> </ul>
DASH diet	LO # 19

15. Understand the role of soluble fiber in heart health and the therapeutic amount recommended for dyslipidemia.

- Fiber (especially soluble) reduces LDL cholesterol by reducing cholesterol absorption, increase bile excretion of cholesterol and up-regulate LDL receptors
- Choose whole-grain, high-fiber foods
  - Aim 20-40 g fiber/day (range due to age, weight)
  - Aim 10-25 g of viscous (soluble) fiber /day
- Total fiber = soluble + insoluble fibers
  - **Soluble:** oats, beans, lentils, barley, psyllium, some fruits & veggies, pectin
  - **Insoluble:** wheat bran, whole grain foods, skins/seeds, nuts & nut butters, fruits & veggies

16. What are phytosterols and how do they therapeutically benefit heart health?

- Work by blocking cholesterol absorption in the intestines
  - 2g of phytosterols/day can lower cholesterol by up to 9%
- Found in plant foods that contain high fat or fiber content (wheat germ, nuts, seeds, legumes, soybeans, vegetable oils and fruits/vegetables)

17. Understand what the therapeutic lifestyle changes (TLC) are and what their cumulative benefit can be.

	Dietary component	Dietary change	Approx.. LDL-C reduction
Major interventions	Saturated fat	< 7% of calories	8-10%
	Dietary cholesterol	< 200 mg	3-5%
	Weight reduction	Lose 10 pounds	5-8%
Other options	Viscous fiber	5-10 g/d	3-5%
	Phytosterols	2 g/d	6-15%
<b>Cumulative estimate</b>			<b>20-30%</b>

18. What are 3 practical tips to implement a healthy heart dietary pattern?

- How do we keep track as to whether we are following a healthy dietary pattern?
- Start tracking how many time you dine out?
- Take a look at your grocery cart/bill, how much of it is fresh produce (fruits, veg, fresh meats, fish, etc)?
- How many fruits and vegetables do you eat in a day?
- What canned/packaged foods are you using? Are you adding fresh to frozen, canned or packaged?
- Start observing what you are doing daily and weekly
- Dietary pattern is built over time

19. Understand what the DASH diet is and what are key points of how it can help to reduce hypertension.

- DASH diet in adults with HTN was associated with reduced risk of stroke & reduced all-cause mortality
- Servings/d: whole grains 6-8; fruits 4-5; vgs 4-5; low-fat milk/alt 2-3; fish & lean meat <6 oz; nuts, fats & oils 2-3 tsp
- Servings/wk: nuts, seeds, legumes 4-5; sweets ≤ 5 tbsp (1 tbsp = 15 g)

20. Be able to make 3 suggestions to make a diet more heart healthy.

- Decreasing saturated fats from meats and butter (in particular) has been shown to be beneficial for reducing LDL-C
- Increasing fruits & veggies to at least 7 servings/day can achieve at least 7g of soluble fiber beneficial to heart health
- Reducing processed foods and dining out can assist in decreasing sodium intake & increasing fiber + other nutrients
- Be carb conscious – reduce added sugars to < 5-10% and choose low GI, high-fiber starches

21. Have an understanding of the burden comorbidities can be on a patient in regards to their diet.

- Warfarin – consistency with vitamin K intake
- Many medications – avoid grapefruit
- Diuretics - ↑↓ potassium
- Salt – restrict or decrease
- Diabetes
- Reducing saturated fats

**LECTURE 16 – MULTIPLE VITAMINS AND MINERALS**

1. To discuss appropriateness of daily multivitamins in otherwise healthy individuals.

- Why are you interested in taking a supplement? Is there a condition they want to treat/prevent?
- Evidence vs. safety issues?
- Current diet; current medications; allergies?
- If starting a supplement: dosing, duration, potential SEs, etc

2. To discuss the evidence with respect to use of ocular vitamins and multivitamins.

Vitamins for age-related macular degeneration	<ul style="list-style-type: none"> <li>• Antioxidants + Zn 80 mg, copper 2 mg, for intermediate or advanced AMD ONLY: <ul style="list-style-type: none"> <li>○ Fewer pts progressed to a more advanced AMD or wet AMD</li> <li>○ Pts on treatment had reduced rate of 15 letter visual acuity loss (vs. placebo)</li> </ul> </li> <li>• Modifications to above regimen: <ul style="list-style-type: none"> <li>○ Omega-3s, xanthophylls: no additional benefit</li> <li>○ Xantophylls: safer than beta-carotene</li> <li>○ Removing beta-carotene: didn't decrease benefit <ul style="list-style-type: none"> <li>▪ B-carotene increases yellowing of skin, and lung cancer in smokers</li> </ul> </li> </ul> </li> </ul>
Antioxidants	<ul style="list-style-type: none"> <li>• Vit A, vit C, vit E, selenium vs. placebo <ul style="list-style-type: none"> <li>○ Increased mortality – specifically w/ b-carotene, vit E, high-dose vit A (&gt;5000 IU)</li> </ul> </li> </ul>
Multivitamins in healthy populations	<ul style="list-style-type: none"> <li>• No effect on overall mortality, CVD/cancer mortality, CVD events, cancer incidence</li> </ul>
Multivitamins for children	<ul style="list-style-type: none"> <li>• Generally not needed if eating well (according to dietary guidelines) and are growing</li> </ul>
Multivitamins for teens	<ul style="list-style-type: none"> <li>• Time of increased bone formation</li> <li>• Increased iron requirements</li> <li>• Time of changes in diet (vegetarian, calorie cutting, increased junk food, etc)</li> <li>• In Canada, the nutrients with lowest intakes are vit A, D, Mg, PO<sub>4</sub>, Ca</li> </ul>

3. To compare and contrast available products.

- Ocular vitamins: Ocuville, Vitalux, I-Care
  - Multivitamins: Centrum (Regular, Forte, Silver [Canadian closest to Select Essentials 50+]), Super VitaVim
  - Children's: Centrum Junior, Flintstones Multiple Vitamins
- ➔ Comparisons to DRIs in slides

**LECTURE 17 – CALCIUM AND VITAMIN D (BONE HEALTH)**

1. Describe the clinical risk factors for fracture.

Age < 50 year	Age 50 – 64 years	Modifiable factors
<ul style="list-style-type: none"> <li>• Fragility fractures</li> <li>• Use of high-risk medications</li> <li>• Hypogonadism</li> <li>• Malabsorption syndromes</li> <li>• Chronic inflammatory conditions</li> <li>• Primary hyperparathyroidism</li> <li>• Other disorders associated w/ rapid bone loss/fractures</li> </ul>	<ul style="list-style-type: none"> <li>• Fragility fracture after age 40</li> <li>• Parental hip fracture</li> <li>• Vertebral fracture or osteopenia (radiography)</li> <li>• Prolonged use of glucocorticoids /other high-risk meds</li> <li>• High alcohol intake or current smoking</li> <li>• Low body weight (&lt;60 kg) or major weight loss (&gt;10% of body weight at age 25)</li> <li>• Other disorders strongly associated with osteoporosis</li> </ul>	<ul style="list-style-type: none"> <li>• Low calcium (and vit D) intake</li> <li>• Propensity to fall</li> <li>• Increased dietary pattern?</li> <li>• Sedentary lifestyle</li> <li>• Increased caffeine/alcohol intake</li> <li>• Current tobacco use</li> <li>• Increased carbonated beverage use?</li> </ul>
NOTE: all men and women > 65 years should be investigated (per CPG)		

2. Estimate a person's 10-year risk of fracture.

- FRAX tool

3. Determine an individual patient's daily elemental calcium and vit D requirements (based on the Health Canada/IOM standards), his/her average intake through foods and supplements, and appropriate recommendations to make up any deficit between current intake and requirements.

Age group	Calcium Intakes		Vit D Intakes	
	RDA (mg)	UTL (mg)	RDA (IU)	UTL (IU)
0 - 6 months	200	1000	400	1000
7 - 12 months	260	1500	400	1500
1 - 3 years	700	2500	600	2500
4 - 8 years	1000	2500		300
9 - 18 years	1300	3000		4000
19 - 50 years	1000	2500		
51 - 70 years	Males: 1000 Females: 1200	2000		
> 70 years	1200	2000	800	4000
Pregnancy/Lactation			600	4000
14-18 years	1300	3000		
19-50 years	1000	2500		

- a. Know the elemental calcium and vit D content of the listed foods, supplements, and different calcium salts.

Food and serving size	Calcium (mg)	Vitamin D (IU)
Milk, 1 cup	300	90
Hard cheese, 50 g	370	
Egg yolk, 1		25
Cottage cheese, 1 cup	150	
Yogurt, fruit bottom, ¾ cup	230	
Orange, 1 medium	50	
Tofu (calcium sulphate), 150 g	350	
White beans, ¾ cup	120	
Almonds, ¼ cup	90	
Atlantic sardines (with bones), 75 g	280	70
Pacific sardines canned, 75 g		585
Salmon canned (with bones), 75 g	200	585
Tuna canned, 75 g		44
Rice or soy beverage (fortified), 1 cup	300	90
Regular soy beverage, 1 cup	100	
Fortified orange juice, ½ cup	185	50
Broccoli, ½ cup	30	

Product	Calcium Salt	Calcium (mg)	Vitamin D (IU)
Caltrate	Carbonate	600	
Caltrate 600 + D		600	800
Caltrate Select		600	400
Caltrate Plus		600	800
Caltrate Mini-Cals		300	800
Tums regular		200	
Tums Extra		300	
Tums Ultra		400	
Citracal + D		Citrate	315
Citracal petites + D	200		200
Novo-calcium gluconate 650 mg	Gluconate	60	
Gramacal	Gluconolactate, carbonate	1000	
Baby D-drops			400
Booster D-drops			600
D-drops			1000

- b. Briefly discuss the evidence that opposes these recommendations.

- Inconsistent results with respect to:
  - Dosing; combo of calcium and vit D, or vit D alone
  - Absolute benefit for fractures, fall reductions, other conditions vs. potential harms
- Evidence discussed in LO #6

4. Compare and contrast calcium citrate and carbonate with respect to clinical use, side effects, etc.

Calcium carbonate (40% elemental calcium)	Calcium citrate (21% of elemental calcium)
<ul style="list-style-type: none"> <li>• Inexpensive</li> <li>• Contains highest amount of elemental calcium</li> <li>• Well-absorbed by most of population</li> <li>• To enhance absorption: <ul style="list-style-type: none"> <li>○ Take with food (comparable absorption to calcium citrate)</li> <li>○ Take in divided doses</li> </ul> </li> <li>• SEs: flatulence, bloating, constipation</li> </ul>	<ul style="list-style-type: none"> <li>• More expensive than carbonate in general</li> <li>• pH independent</li> <li>• Less carbon dioxide production</li> <li>• Recommended for patients with: achlorhydria, kidney stones, intolerable gas with carbonate</li> <li>• Avoid in chronic renal failure <ul style="list-style-type: none"> <li>○ Binds aluminum in dialysis solution</li> </ul> </li> </ul>

5. Discuss the evidence for the use of calcium and/or vitamin D in fracture reduction, falls prevention, mortality reduction.

Fracture reduction	<ul style="list-style-type: none"> <li>• Calcium or calcium + vit D: 10% RRR</li> <li>• Dietary calcium: most <u>cohort</u> studies show neutral relationship; RCTs too small to show SS</li> </ul>						
Glucocorticoid-induced osteoporosis	<ul style="list-style-type: none"> <li>• Calcium + vit D more effective at retarding lumbar &amp; forearm bone loss than placebo OR calcium alone</li> </ul>						
Falls (in community)	<ul style="list-style-type: none"> <li>• Vit D supplements MAY reduce falls in older community pts (but only those who have low blood levels pre-treatment)</li> </ul>						
Falls (nursing homes or hospital)	<ul style="list-style-type: none"> <li>• Vit D (800 IU) + Ca (1000 mg): reduced rate of falls, but not risk of falling <ul style="list-style-type: none"> <li>◦ Pts in study had low or very low vit D levels</li> </ul> </li> </ul>						
Does calcium need to be taken with vit D?  Review of meta-analyses	<table border="1"> <tr> <td>Fall prevention  (9 analyses)</td> <td> <ul style="list-style-type: none"> <li>• 2 concluded vit D only effective with calcium</li> <li>• 2 concluded vit D effective alone</li> <li>• Others concluded vit D only effective under certain circumstances <ul style="list-style-type: none"> <li>◦ Dose <math>\geq</math> 700 IU/day or people with low serum levels</li> </ul> </li> </ul> </td> </tr> <tr> <td>Fracture reduction  (10 analyses)</td> <td> <ul style="list-style-type: none"> <li>• 5 concluded vit D only effective with calcium</li> <li>• 3 concluded vit D alone effective at <math>\geq</math> 482 IU/day</li> <li>• 2 concluded vit D had no beneficial effect on fracture risk</li> </ul> </td> </tr> <tr> <td>All-cause mortality  (5 analyses)</td> <td> <ul style="list-style-type: none"> <li>• 3 concluded only beneficial if calcium &amp; vit D are combined</li> <li>• 2 concluded vit D alone is beneficial</li> </ul> </td> </tr> </table>	Fall prevention  (9 analyses)	<ul style="list-style-type: none"> <li>• 2 concluded vit D only effective with calcium</li> <li>• 2 concluded vit D effective alone</li> <li>• Others concluded vit D only effective under certain circumstances <ul style="list-style-type: none"> <li>◦ Dose <math>\geq</math> 700 IU/day or people with low serum levels</li> </ul> </li> </ul>	Fracture reduction  (10 analyses)	<ul style="list-style-type: none"> <li>• 5 concluded vit D only effective with calcium</li> <li>• 3 concluded vit D alone effective at <math>\geq</math> 482 IU/day</li> <li>• 2 concluded vit D had no beneficial effect on fracture risk</li> </ul>	All-cause mortality  (5 analyses)	<ul style="list-style-type: none"> <li>• 3 concluded only beneficial if calcium &amp; vit D are combined</li> <li>• 2 concluded vit D alone is beneficial</li> </ul>
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6. Discuss the evidence for the potential cardiovascular harm of calcium and/or vitamin D.

- Calcium (w/ or w/o vit D): no relationship to risk of CVD or mortality
- High calcium: decreased risk of incident atherosclerosis but may increase risk for incident coronary artery calcification

7. Discuss the role of calcium in the formation of kidney stones & counsel on ways to reduce risk of developing kidney stones.

- Most kidney stones are made of calcium oxalate (precipitation of crystals in urine)
  - LOW calcium diets increase oxalate in urine (and this is likely cause of kidney stones)
  - Dietary calcium binds oxalates in GIT – limits their absorption and prevents precipitation in kidney
- Patients with calcium oxalate stones can:
  - Limit sources of oxalate (beet greens, rhubarb, sorrel, nuts, berries, spinach, chocolate)
  - Continue with calcium intake – calcium citrate may have a protective effect

8. Correctly advise individual patients on the use of “bone health” nutrition and supplements for general health, treatment or prevention of osteoporosis/bone fracture including treatment regimens and appropriate counselling.

Interactions with Ca	<ul style="list-style-type: none"> <li>• Phytates (bran, Metamucil, soybeans, seeds, etc)</li> <li>• Some antibiotics (quinolones, tetracyclines)</li> <li>• Thiazides, Bisphosphonates, L-thyroxine, iron, etc</li> </ul>
Counselling tips	<ul style="list-style-type: none"> <li>• <u>Carbonate</u>: with food for better absorption</li> <li>• <u>Citrate</u>: may be taken without regard to food</li> <li>• Divided doses separate from other calcium sources (max single dose 500 mg)</li> </ul>
Magnesium	<ul style="list-style-type: none"> <li>• Concurrent ingestion with calcium (no effects on absorption) <ul style="list-style-type: none"> <li>◦ Possible benefits: laxative effect (&gt;350 mg)</li> </ul> </li> <li>• Has a role in bone and teeth formation</li> <li>• DRI: 320 – 420 mg; sources: whole grains, green leafy veggies, nuts, seafood</li> </ul>
Other nutrients	<ul style="list-style-type: none"> <li>• No evidence to recommend additional intakes for prevention/treatment of osteoporosis</li> </ul>



**LECTURE 15, 16 AND 22- MICRONUTRIENT PHARMACOLOGY (PARTS 1 AND 2)**

**1. Define the terms vitamin, mineral, DRI, RDA, AI and UL.**

Micronutrients (vitamins & minerals)	<ul style="list-style-type: none"> <li>• Substances essential in very small quantities for normal metabolic &amp; physiological fxn</li> <li>• Either not formed at all or are formed in insufficient amounts in body                             <ul style="list-style-type: none"> <li>○ Must be supplied from exogenous sources</li> </ul> </li> </ul>
Dietary reference intakes (DRI)	<ul style="list-style-type: none"> <li>• Harmonized US &amp; Canadian guidelines for recommended nutrient intakes in healthy pops</li> <li>• Evidence-based                             <ul style="list-style-type: none"> <li>○ Consider maintenance of good health &amp; prevention of chronic disease</li> <li>○ Also considers possible toxicity of excess intake</li> </ul> </li> </ul>
Recommended dietary allowances (RDA)	<ul style="list-style-type: none"> <li>• Dietary intake level sufficient to meet needs of 97 - 98% individuals in a group</li> <li>• Calculated based on the <b>Estimated Average Requirement (EAR)</b> = the median daily intake estimated to meet the requirement of 50% of healthy individuals in a group</li> </ul>
Adequate intake (AI)	<ul style="list-style-type: none"> <li>• For nutrients for which there isn't enough information to calculate an RDA</li> <li>• Based on the estimated average nutrient intake by a group (or groups) of healthy individual</li> </ul>
Tolerable Upper Intake Levels (UL)	<ul style="list-style-type: none"> <li>• Highest level of daily intake likely to pose no risks or adverse effects to almost all individuals in the general population</li> </ul>

**2. Differentiate between use of vitamins and minerals as dietary supplements and as therapeutic agents.**

Are supplements required?	<ul style="list-style-type: none"> <li>• Vitamin &amp; mineral needs can (and should usually be met) by eating a balanced diet with a wide variety of foods (ex// Canada's Food Guide)</li> <li>• Health Canada also requires the fortification of certain foods in an effort to prevent nutritional deficiencies                             <ul style="list-style-type: none"> <li>○ Vit A &amp; D - added to milk and margarine</li> <li>○ Thiamine, riboflavin, niacin, folic acid, iron -added to enriched white flour</li> <li>○ Iodine - added to table salt</li> </ul> </li> </ul>
When are supplements recommended?	<ol style="list-style-type: none"> <li>1) Requirements are increased such that it's difficult to obtain sufficient amounts from diet                             <ol style="list-style-type: none"> <li>a. Pregnant &amp; lactating women (vitamins, folic acid, calcium + vit D, iron)</li> </ol> </li> <li>2) Intake from diet is likely to be inadequate                             <ol style="list-style-type: none"> <li>a. People on very low calorie diets</li> <li>b. Those who avoid entire food groups (vegetarians, vegans)</li> <li>c. Chronic substance abuse</li> <li>d. Elderly</li> </ol> </li> <li>3) Malabsorption is a risk                             <ol style="list-style-type: none"> <li>a. Following bariatric surgery</li> <li>b. GI diseases known to cause malabsorption (ex// Cystic Fibrosis)</li> </ol> </li> </ol>
When are vitamins & minerals needed as therapeutic agents?	<ol style="list-style-type: none"> <li>1) For correction of deficiency (UL do not apply)</li> <li>2) Use of high doses (generally &gt; 10 times RDA) of one or a few agents in the prevention or treatment of diseases or conditions unrelated to nutritional deficiency</li> </ol>

**3. For each of the vitamins discussed, state the chemical name of the active substances, and list at least 2 major sources from which the vitamin can be obtained.**

Vitamin A	Name	<ul style="list-style-type: none"> <li>• Collective name for a group of compounds having the biological activity of retinol                             <ul style="list-style-type: none"> <li>○ Retinol, retinal, retinoic acid</li> </ul> </li> </ul>
	Sources	<ul style="list-style-type: none"> <li>• Pre-formed, mostly as retinyl esters: from animal sources (liver, eggs, milk)</li> <li>• Pro-vitamin A carotenoids (precursors): from pigmented fruits &amp; veggies                             <ul style="list-style-type: none"> <li>○ B-carotene as largest biological activity; supplements better absorbed than from diet (green, leafy veggies and intensely colored fruits/veggies)</li> </ul> </li> </ul>
	Requirement	<ul style="list-style-type: none"> <li>• Vary between 300 - 1300 RAE/day (depending on age, gender, etc)                             <ul style="list-style-type: none"> <li>○ RAE = retinol activity equivalents (1 RAE = 1 ug all-trans retinol)</li> </ul> </li> </ul>

B-vitamins	Names	<ul style="list-style-type: none"> <li>Thiamine, riboflavin, pyridoxine, niacin, pantothenic acid, biotin, folic acid, cyanocobalamin (B12), choline, inositol, carnitine</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Thiamine (B1): whole grain cereals, legumes (beans/lentils), nuts, lean pork and yeast <ul style="list-style-type: none"> <li>In Canada, wheat flour is fortified with thiamine</li> </ul> </li> <li>Riboflavin (B2): cereal, nuts, milk, eggs, green leafy vegetables, lean meat</li> <li>Niacin (B3): meat, poultry, fish (tuna, salmon), fortified cereals, legumes, and nuts <ul style="list-style-type: none"> <li>Also formed from tryptophan, in a reaction that requires B6</li> <li>Less bioavailable from cereal grains</li> </ul> </li> <li>Pyridoxine (B6): beans, legumes, nuts, eggs, meats, fish breads, cereals</li> </ul>
Vitamin C	Name	<ul style="list-style-type: none"> <li>Ascorbic acid</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Citrus fruits, green peppers, strawberries, tomatoes, broccoli, sweet &amp; white potatoes</li> </ul>
	Requirements	<ul style="list-style-type: none"> <li>RDA = 90 mg/d for adult men and 75 mg/day for adult women</li> <li>RDA increased to 125 (men) and 110 (women) mg/day in smokers</li> </ul>
Vitamin D	Names	<ul style="list-style-type: none"> <li>Ergocalciferol (vitamin D2): from yeasts and fungi</li> <li>Cholecalciferol (vitamin D3): from animal tissues</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Formed in skin on exposure to sunlight</li> <li><u>Dietary</u>: fatty fish (mackerel, salmon, sardines), fish liver oils, fortified milk</li> <li><u>OTC supplements</u>: either vitamin D2 or D3 (50-77% more biological activity than D2)</li> </ul>
	Requirements	<ul style="list-style-type: none"> <li><u>IOM</u>: birth - 1 year = 400 IU; 1-70 yo = 600 IU ; &gt; 70 yo = 800 IU</li> <li><u>CCS</u>: adults = 1000 IU in winter months, or if at high risk of deficiency</li> <li><u>Osteoporosis</u>: low-risk adults &lt;50 yo = 400-1000 IU ; &gt;50+ high risk adults 800-2000 IU</li> </ul>
Vitamin E	Name	<ul style="list-style-type: none"> <li>Includes 8 naturally occurring compounds called tocopherols <ul style="list-style-type: none"> <li>Alpha-tocopherol: found in greatest amounts in circulation and tissues</li> </ul> </li> </ul>
	Sources	<ul style="list-style-type: none"> <li><u>Dietary</u>: vegetable oils, nuts, whole grains (contain the active RRR-<math>\alpha</math>-tocopherol)</li> <li><u>Synthetic</u>: contains all rac-<math>\alpha</math>-tocopherol but only 4/9 stereoisomers are biologically active</li> <li><u>Supplements</u>: sold as esters of either natural or synthetic vitamin E</li> </ul>
	Requirements	<ul style="list-style-type: none"> <li>RDA: 4-19 mg of RRR- <math>\alpha</math>-tocopherol/day <ul style="list-style-type: none"> <li>1 IU natural vitamin E = 0.67 mg RRR- <math>\alpha</math>-tocopherol</li> <li>1 IU synthetic vitamin E = 0.45 mg RRR- <math>\alpha</math>-tocopherol</li> </ul> </li> </ul>
Vitamin K	Name	<ul style="list-style-type: none"> <li>Vitamin K<sub>1</sub> = phytomenadione</li> <li>Vitamin K<sub>2</sub> = menaquinone series</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Vitamin K<sub>1</sub>: plants, especially vegetables; also synthetic</li> <li>Vitamin K<sub>2</sub>: synthesized by intestinal bacteria</li> </ul>

4. Outline the absorption, distribution, and metabolism of vitamins A, C, and D.

Vitamin A	Absorption	<ul style="list-style-type: none"> <li>Vitamin A <math>\rightarrow</math> intestinal mucosa <math>\rightarrow</math> trans-retinol</li> <li>Carotenoids <math>\rightarrow</math> trans-retinol</li> </ul>
	Distribution	<ul style="list-style-type: none"> <li>Trans-retinol = transport form - bound to retinol binding protein</li> <li><math>\rightarrow</math> Retinoic acid - active form in all tissues (except eye)</li> <li><math>\rightarrow</math> Retinal - active form in eye</li> </ul>
	Metabolism	<ul style="list-style-type: none"> <li>Liver via vitamin A esters</li> </ul>
Vitamin C	Absorption	<ul style="list-style-type: none"> <li>Dose-dependent active transport in upper GIT</li> </ul>
	Metabolism	<ul style="list-style-type: none"> <li>Approx. 15% converted to oxalic acid</li> </ul>
	Excretion	<ul style="list-style-type: none"> <li>At low intake, undergoes active tubular reabsorption</li> <li>At doses <math>\geq</math> 1000 mg, excess is excreted in urine</li> </ul>
Vitamin D	Activation	<ol style="list-style-type: none"> <li>7-dehydrocholesterol (skin) <math>\rightarrow</math> SUN EXPOSURE <math>\rightarrow</math> vitamin D<sub>3</sub> (cholecalciferol)</li> <li>Vitamin D<sub>3</sub> (skin) <math>\rightarrow</math> LIVER <math>\rightarrow</math> 25-OHD<sub>3</sub> (calcifediol)</li> <li>25-OHD<sub>3</sub> <math>\rightarrow</math> KIDNEY <math>\rightarrow</math> 1, 25-(OH)<sub>2</sub>D<sub>3</sub> (calcitriol)</li> </ol>

5. State the physiological function of each vitamin, and where discussed, explain their mechanism of action and regulation.

Vitamin A	<ul style="list-style-type: none"> <li>• Retinoic acid (RA) acts by regulating gene expression in target cells: <ul style="list-style-type: none"> <li>○ Circulating retinol is converted to retinoic acid</li> <li>○ Acts through nuclear retinoic acid receptors (RAR)</li> </ul> </li> <li>• Maintenance of normal vision, particularly night vision <ul style="list-style-type: none"> <li>○ <u>Visual cycle</u>: series of enzymatic reactions for recycling retinoids used in photoreceptor cells of the retina for light detection <ol style="list-style-type: none"> <li>1. Retinol circulates to the retina &amp; moves into retinal pigment epithelial cells → isomerized and oxidized to form 11-cis-retinal</li> <li>2. 11-cis-retinal transported to rod cells → binds to opsin to form rhodopsin (visual pigment)</li> <li>3. Absorption of a photon of light catalyzes isomerization of 11-cis-retinal to all-trans-retinal, changing conformation of rhodopsin &amp; triggering cascade of signalling events → optic nerve transmission → vision <ol style="list-style-type: none"> <li>a) AT SAME TIME, all-trans-retinal is released from rhodopsin, converted to all-trans-retinol and transported back to retinal epithelial cell for storage as all-trans retinyl esters → re-entry into cycle</li> </ol> </li> </ol> </li> </ul> </li> <li>• Differentiation and function of epithelial cells</li> <li>• Embryonic development, growth and cell differentiation</li> <li>• Function of the immune system</li> </ul>	
Vitamin B	Thiamine (B1)	<ul style="list-style-type: none"> <li>• Essential coenzyme in several decarboxylation reactions in carb &amp; amino acid metabolism</li> <li>• Required for energy generation</li> </ul>
	Riboflavin (B2)	<ul style="list-style-type: none"> <li>• Converted to FMN (flavin mononucleotide) and FAD (flavin adenine dinucleotide) <ul style="list-style-type: none"> <li>○ Co-enzymes for flavoproteins, act as electron carriers in redox reactions in fatty acid and amino acid metabolism and the TCA cycle</li> <li>○ FAD is part of the electron transport (respiratory) chain – central to energy production</li> </ul> </li> <li>• Involved in metabolism of vit B6, folic acid, and niacin</li> </ul>
	Niacin (B3)	<ul style="list-style-type: none"> <li>• Form part of nicotinamide adenine dinucleotide (NAD) and nicotinamide adenine dinucleotide phosphate (NADP) <ul style="list-style-type: none"> <li>○ Co-enzymes in many oxidation-reduction reactions, also participate in DNA repair</li> </ul> </li> </ul>
	Pyridoxine (B6)	<ul style="list-style-type: none"> <li>• As pyridoxal-phosphate in amino acid and glycogen metabolism, and steroid hormone action</li> <li>• Acts as co-factor in conversion of tryptophan to niacin</li> </ul>
Vitamin C	<ul style="list-style-type: none"> <li>• As an antioxidant in a number of biochemical reactions (ex// conversion of proline to hydroxyproline in collagen synthesis)</li> <li>• Also suggested to protect against free radical-induced damage</li> </ul>	
Vitamin D	<ul style="list-style-type: none"> <li>• Vitamin D acts by regulating gene expression in target cells via nuclear vitamin D receptors (VDR) <ul style="list-style-type: none"> <li>○ Receptors are widespread (brain, prostate, colon, immune cells, etc)</li> </ul> </li> <li>• With calcitonin &amp; PTH, regulates serum Ca and Pi levels within narrow physiological range <ol style="list-style-type: none"> <li>○ Supports normal bone mineralization, neuromuscular function and cell physiology <ol style="list-style-type: none"> <li>1. Increases expression of Ca channels, Ca binding proteins, Pi transporter in intestinal epithelial cells → increases Ca and Pi absorption</li> <li>2. Enhances mobilization of Ca and Pi from bone (if dietary Ca is insufficient)</li> <li>3. Decreases Ca and Pi excretion by the kidney (minor effect)</li> </ol> </li> </ol> </li> <li>• Controls the expression of more than 200 genes, including for cell proliferation, differentiation, apoptosis <ul style="list-style-type: none"> <li>○ Decreases proliferation of normal and cancer cells and induces their differentiation</li> </ul> </li> <li>• ALSO: influences immune cell function, inhibits renin synthesis, increases insulin production &amp; increases myocardial contractility (in studies in cells and animals)</li> </ul>	
Vitamin E	<ul style="list-style-type: none"> <li>• No defined metabolic function</li> <li>• Lipid-soluble antioxidant, suggested to protect cell membranes against oxidative damage by free radicals</li> </ul>	

Vitamin K	<ul style="list-style-type: none"> <li>Required for normal blood clotting <ul style="list-style-type: none"> <li>Co-factor for vit K-dependent carboxylase</li> <li>7 vitamin-K dependent clotting factors must undergo gamma carboxylation to become active</li> <li>Vitamin K becomes oxidized during the reaction, and must be reduced to participate in rxn again</li> <li>Warfarin inhibits vit K reduction → reduces synthesis of vit-K dependent clotting factors</li> </ul> </li> <li>Other vitamin K dependent proteins: <ul style="list-style-type: none"> <li>Osteocalcin: formed by osteoblasts under influence of vit D; may be involved in mineralization</li> <li>Matrix Gla protein: may prevent calcification of soft tissues</li> </ul> </li> </ul>
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6. Describe the condition produced by deficiency of each of the vitamins, including the major symptoms, and explain the circumstances under which this might occur.

Vitamin A	<ul style="list-style-type: none"> <li>Not common in N. America</li> <li>In N. America, most often seen in chronic diseases associated with fat malabsorption or in conditions associated with impaired storage (ex// liver cirrhosis)</li> <li>Develops slowly in adults (liver stores maintain plasma levels for up to 2 years), but much more rapidly in children</li> </ul>							
	Symptoms	<table border="1"> <tr> <td>Eye</td> <td> <ul style="list-style-type: none"> <li>Retinal dysfunction leading to progressive reversible night blindness</li> <li>Xerophthalmia: changes in structure &amp; function of epithelial cells <ul style="list-style-type: none"> <li>Conjunctival &amp; corneal xerosis (dry, wrinkled, hazy) → corneal ulceration, scarring → blindness</li> </ul> </li> </ul> </td> </tr> <tr> <td>Other</td> <td> <ul style="list-style-type: none"> <li>Decreased resistance and increased morbidity &amp; mortality due to infections</li> <li>Loss of functional, mucous-secreting epithelial cells <ul style="list-style-type: none"> <li>Increased susceptibility to respiratory infections, rash, diarrhea</li> </ul> </li> </ul> </td> </tr> </table>	Eye	<ul style="list-style-type: none"> <li>Retinal dysfunction leading to progressive reversible night blindness</li> <li>Xerophthalmia: changes in structure &amp; function of epithelial cells <ul style="list-style-type: none"> <li>Conjunctival &amp; corneal xerosis (dry, wrinkled, hazy) → corneal ulceration, scarring → blindness</li> </ul> </li> </ul>	Other	<ul style="list-style-type: none"> <li>Decreased resistance and increased morbidity &amp; mortality due to infections</li> <li>Loss of functional, mucous-secreting epithelial cells <ul style="list-style-type: none"> <li>Increased susceptibility to respiratory infections, rash, diarrhea</li> </ul> </li> </ul>		
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	Riboflavin (B2)	<ul style="list-style-type: none"> <li>Uncommon and rarely occurs alone due to efficient re-cycling of riboflavin</li> <li>Symptoms: cheilosis and angular stomatitis (cracks &amp; sores on lips &amp; corner of mouth); inflammation and redness of tongue; skin rash; corneal vascularization; anemia</li> </ul>						
Niacin (B3)	<ul style="list-style-type: none"> <li>Known as pellagra</li> <li>Symptoms: dermatitis, diarrhea, dementia</li> </ul>							
Pyridoxine (B6)	<ul style="list-style-type: none"> <li>Uncommon, may occur in alcoholics, malabsorption diseases <ul style="list-style-type: none"> <li>Symptoms: dermatitis, neuropathy, anemia</li> </ul> </li> <li>Drug-induced (ex// isoniazid): associated with sensory neuropathy</li> </ul>							
Vitamin C	Severe (scurvy)	<ul style="list-style-type: none"> <li>Uncommon: elderly, alcoholics, drug addicts and those with very poor diets at risk</li> <li>Symptoms (mild): anorexia, weakness, fatigue, irritability</li> <li>Symptoms (severe): red &amp; swollen gums, loose teeth, anemia, hemorrhage</li> </ul>						
Vitamin D	Deficiency	<ul style="list-style-type: none"> <li><u>Children</u>: Rickets – failure to mineralize new bone &amp; cartilage matrix <ul style="list-style-type: none"> <li>Results in soft bones, deformed joints &amp; increased susceptibility to fracture</li> </ul> </li> <li><u>Adults</u>: <ul style="list-style-type: none"> <li><i>Osteomalacia</i>: ↓ bone density &amp; mineralization, muscle weakness, bone pain</li> <li><i>Osteoporosis</i> &amp; increased risk of fractures</li> </ul> </li> </ul>						

	Risk of deficiency	<ul style="list-style-type: none"> <li>At Vancouver's latitude (49° N) UVB radiation too low for vit D production in skin from mid-October to mid-March <ul style="list-style-type: none"> <li>During remainder of year, short periods of sun exposure should produce adequate vitamin D (depending on factors such as sun intensity, skin color, age)</li> </ul> </li> <li>4% of Canadians considered vit D deficient; 10% vit-D insufficient</li> </ul>
	Risk increased in	<ul style="list-style-type: none"> <li>Exclusively breastfed infants</li> <li>Elderly</li> <li>Covering all exposed skin or using sunscreen (SPF ≥ 8) whenever outside</li> <li>Dark skin pigmentation</li> <li>Fat malabsorption syndromes and inflammatory bowel disease</li> <li>If renal production of 1,25-(OH)D<sub>3</sub> is impaired</li> </ul>
	Impaired production of calcitriol	<ul style="list-style-type: none"> <li>Ex// renal osteodystrophy, hypoparathyroidism, vit D-dependent rickets <ul style="list-style-type: none"> <li>Calcitriol: active form of vitamin</li> <li>Alfacalcidol: already hydroxylated in 1α position (25 position)</li> </ul> </li> </ul>
Vitamin E		<ul style="list-style-type: none"> <li>Normally symptomatic dietary deficiency is not seen</li> <li>Neurological sx and anemia responsive to vitamin E found in premature infants &amp; in children with fat malabsorption syndromes</li> </ul>
Vitamin K		<ul style="list-style-type: none"> <li>Hypoprothrombinemia → increased tendency to hemorrhage</li> </ul>
	Risk of deficiency	<ul style="list-style-type: none"> <li><u>Newborn</u>: minimal stores; no established intestinal flora; breast-fed infants have limited dietary intake <ul style="list-style-type: none"> <li>Administration of a small dose of phytomenadione to newborn infants routine</li> </ul> </li> <li><u>After infancy</u>: although limited storage, dietary deficiency is rare (supplied in diet; conserved by vitamin K cycle and is produced by intestinal bacteria)</li> <li><u>Inadequate absorption</u>: fat malabsorption syndromes</li> </ul>

7. Describe the adverse effects associated with excess intake of each of the vitamins.

Vitamin A		<ul style="list-style-type: none"> <li>Intake &gt; 1500 ug/d associated with decreased bone density &amp; increased risk of fractures in adults</li> <li><u>Chronic toxicity</u> can occur on intake of 10-20 x the RDA for as little as 6 m in adults (less in children) <ul style="list-style-type: none"> <li>Liver damage may occur; dry itchy peeling skin; increased intracranial pressure (children)</li> </ul> </li> <li><u>Acute toxicity</u> (on ingestion of massive doses – 500 x the RDA in adults): severe headache &amp; generalized peeling of skin (after 24 h)</li> <li>Excessive intake of vit A during pregnancy (particularly first 2 months) increases risk of birth defects <ul style="list-style-type: none"> <li>Women who are pregnant or planning to become pregnant should not take more than the RDA</li> </ul> </li> <li>UL for vitamin A: 3000 ug (10,000 IU/day) for adults <ul style="list-style-type: none"> <li>Based on prevention of liver toxicity (decreased bone density not considered)</li> </ul> </li> <li>Toxicity of B-carotene (not associated with vitamin A toxicity or teratogenicity) <ul style="list-style-type: none"> <li>One sign of excess intake is yellow-orange skin</li> <li>Studies found increased risk of lung cancer in smokers and total mortality</li> </ul> </li> </ul>
Vitamin B	Thiamine (B1)	<ul style="list-style-type: none"> <li>No UL; no known toxicity on oral intake</li> </ul>
	Riboflavin (B2)	<ul style="list-style-type: none"> <li>No UL; no known toxicity</li> <li>High-dose colors urine fluorescent yellow</li> </ul>
	Niacin (B3)	<ul style="list-style-type: none"> <li>Nicotinic acid: flushing of skin (primarily face, arms, chest) <ul style="list-style-type: none"> <li>Flushing with nicotinamide is rare</li> <li>UL for niacin (both nicotinic acid &amp; nicotinamide) set at 35 mg/day to avoid it</li> </ul> </li> <li>Other SEs: pruritus; GI distress; abnormal liver function <ul style="list-style-type: none"> <li>Most common when used in high doses for treatment of hyperlipoproteinemia</li> </ul> </li> </ul>
	Pyridoxine (B6)	<ul style="list-style-type: none"> <li>Sensory neuropathy with intakes &gt; 200 mg/day for several months</li> <li>UL = 100 mg/day for adults</li> </ul>

Vitamin C	<ul style="list-style-type: none"> <li>• Generally well-tolerated</li> <li>• UL for adults = 2g/d</li> <li>• High doses (&gt;4 g/d) → osmotic diarrhea &amp; GI disturbances <ul style="list-style-type: none"> <li>○ Avoid high doses if history of renal stones or pre-existing renal disease or failure (may be aggravated by excess oxalate production)</li> </ul> </li> </ul>
Vitamin D	<ul style="list-style-type: none"> <li>• Hypercalcemia → calcification of soft tissues; irreversible liver failure <ul style="list-style-type: none"> <li>○ S/S: weakness and fatigue, headache, nausea, vomiting, diarrhea, impaired renal function</li> </ul> </li> <li>• UL = 4000 IU/day in &gt;9 years old <ul style="list-style-type: none"> <li>○ Toxicity seen most often at doses &gt; 10,000 IU/day</li> </ul> </li> </ul>
Vitamin E	<ul style="list-style-type: none"> <li>• UL = 1,000 mg daily of alpha-tocopherol (any form) <ul style="list-style-type: none"> <li>○ Based on possibility that high doses vit E may have anticoagulant activity &amp; could cause hemorrhage</li> <li>○ Increased risk of bleeding has been observed with co-administration of vit E and warfarin</li> </ul> </li> </ul>
Vitamin K	<ul style="list-style-type: none"> <li>• Phytomenadione generally non-toxic → no UL been established</li> </ul>

8. Explain the therapeutic uses, if any, of each of the vitamins other than in the treatment of vitamin deficiency.

Vitamin A	Only used in the correction of vit A deficiency						
	Retinoids	<ul style="list-style-type: none"> <li>• Natural and synthetic analogs of retinol (not all have vitamin A activity)</li> <li>• Used in dermatological conditions (ex// isotretinoin in severe cystic acne)</li> <li>• Prevention &amp; treatment of epithelial cell cancers</li> </ul>					
Vit B	Thiamine (B1)	<ul style="list-style-type: none"> <li>• Prevention and treatment of thiamine deficiency</li> <li>• Other uses under investigation (diabetes, Alzheimer's)</li> </ul>					
	Riboflavin (B2)	<ul style="list-style-type: none"> <li>• Migraine prophylaxis (based on single small RCT of 400 mg/day) <ul style="list-style-type: none"> <li>○ NOTE: only 25 mg can be absorbed in a single oral dose</li> </ul> </li> </ul>					
	Niacin (B3)	<ul style="list-style-type: none"> <li>• In pharmacological doses, nicotinic acid (NOT nicotinamide) decreases serum cholesterol and triglycerides</li> </ul>					
	Pyridoxine (B6)	<table border="1"> <tr> <td>PMS</td> <td> <ul style="list-style-type: none"> <li>• 50-100 mg/day may be of benefit</li> </ul> </td> </tr> <tr> <td>Morning sickness</td> <td> <ul style="list-style-type: none"> <li>• May be slightly beneficial in reducing N and/or V in pregnancy <ul style="list-style-type: none"> <li>○ 25 mg every 8 hours for 3 days</li> <li>or</li> <li>○ 10 mg every 8 hours for 5 days</li> </ul> </li> </ul> </td> </tr> <tr> <td>No evidence for</td> <td> <ul style="list-style-type: none"> <li>• Prevention of side effects of low-dose oral contraceptives</li> <li>• Carpal tunnel syndrome</li> <li>• Depression</li> <li>• Mood or cognitive function in elderly</li> </ul> </td> </tr> </table>	PMS	<ul style="list-style-type: none"> <li>• 50-100 mg/day may be of benefit</li> </ul>	Morning sickness	<ul style="list-style-type: none"> <li>• May be slightly beneficial in reducing N and/or V in pregnancy <ul style="list-style-type: none"> <li>○ 25 mg every 8 hours for 3 days</li> <li>or</li> <li>○ 10 mg every 8 hours for 5 days</li> </ul> </li> </ul>	No evidence for
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Vitamin C	CV disease	<ul style="list-style-type: none"> <li>• No evidence of benefit in prevention of CVD</li> </ul>					
	Advanced cancer	<ul style="list-style-type: none"> <li>• Large oral doses (up to 10 g/d) of no benefit in treatment of advanced cancer</li> <li>• IV administration: can reach much higher blood levels but studies don't show benefit <ul style="list-style-type: none"> <li>○ Many CAM practitioners treat cancer pts with ≥ 25 g IV several times per wk</li> </ul> </li> </ul>					
	Common cold	<ul style="list-style-type: none"> <li>• <u>Prophylactic use:</u> <ul style="list-style-type: none"> <li>○ No evidence for reduction of <i>incidence</i> of colds in general population; may reduce in those exposed to severe physical exercise</li> <li>○ Slight reduction in <i>duration</i> of colds</li> </ul> </li> <li>• <u>Therapeutic use:</u> no evidence of benefit</li> </ul>					
Anti-oxidants	AMD (age-related macular degeneration)	<ul style="list-style-type: none"> <li>• Combo of antioxidants (B-carotene, vit C, vit E) + zinc oxide → delay progression of AMD and loss of visual acuity in pts with <i>moderate-severe</i> disease <ul style="list-style-type: none"> <li>○ NOTE: combo w/o B-carotene as effective</li> </ul> </li> <li>• No evidence that supplementation will prevent AMD</li> </ul>					
Vit D	Osteoporosis	<ul style="list-style-type: none"> <li>• Supplementation with 800 IU/day vit D (+ 1000 mg/day Ca) reduces risk of hip and non-vertebral fractures in older patients</li> </ul>					

	Reduced risk of disease	<ul style="list-style-type: none"> <li>Insufficient evidence to support increased vit D to prevent cancer or other chronic diseases <i>unrelated to skeletal health</i> <ul style="list-style-type: none"> <li>Low vit D levels may be a <u>marker</u> for poor health status, rather than a cause of it</li> </ul> </li> </ul>
Vit E	Cancer & CV disease	<ul style="list-style-type: none"> <li>Results largely negative</li> <li>Vitamin E supplements may be harmful (evidence of small increased risk of HF, MI, stroke, prostate cancer)</li> </ul>
	Alzheimer's?	<ul style="list-style-type: none"> <li>One RCT showed delay in deterioration in pts with mild-moderate AD (2000 IU/day)</li> <li>Another RCT found no effect of 2000 IU/day on rate of progression from MCI → AD</li> </ul>
	Topical	<ul style="list-style-type: none"> <li>Suggested to improve wound healing and decrease scarring of skin <ul style="list-style-type: none"> <li>Not supported by clinical trials</li> </ul> </li> </ul>
Vit K	Antagonist	<ul style="list-style-type: none"> <li>To overcome excess effects of coumarin oral anticoagulants</li> </ul>
	Newborns	<ul style="list-style-type: none"> <li>To prevent hemorrhagic disease of newborn</li> </ul>
	Topical	<ul style="list-style-type: none"> <li>No evidence that it is effective in preventing or treating bruising</li> </ul>

9. State the known physiological functions of calcium, zinc, and fluoride, and describe the consequences of their deficiency and excess intake and the use of supplements.

Calcium	Required for	<ul style="list-style-type: none"> <li>Current flow across excitable membranes</li> <li>Muscle contraction, vesicle fusion and release</li> <li>Blood coagulation</li> <li>Formation, structure &amp; remodeling of skeleton</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Mainly from milk and dairy products; also green leafy veggies and salmon, sardines</li> </ul>
	ADME	<ul style="list-style-type: none"> <li>Absorbed in intestine by active vitamin-D dependent transport &amp; facilitated diffusion <ul style="list-style-type: none"> <li>Efficiency increases as intake decreases, in part due to activation of vit D</li> </ul> </li> <li>Stored in bones (&gt;99% of body stores)</li> <li>Excreted in urine - regulated by parathyroid hormone (PTH)</li> </ul>
	Deficiency	<ul style="list-style-type: none"> <li>Common across all age groups</li> <li>If chronic, may prevent attainment of peak bone mass and may contribute to accelerated bone loss &amp; development of osteoporosis</li> </ul>
	Toxicity	<ul style="list-style-type: none"> <li>UL ranges from 1000 mg/d (infants) to 3000 mg/d (adolescents)</li> <li>Some studies suggested intake from supplements may contribute to small increase in risk of atherosclerosis and MI</li> </ul>
	Requirements	<ul style="list-style-type: none"> <li>Range from 200-1300 mg/day (depending on age, gender and status)</li> </ul>
Zinc	Function	<ul style="list-style-type: none"> <li>Essential co-factor for large number of enzymes <ul style="list-style-type: none"> <li>Required for cell growth, nucleic acid, carbohydrate and protein synthesis</li> </ul> </li> <li>Required for normal use of vitamin A</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Widely available in food; more bioavailable from meat (esp. red meat) than plants</li> </ul>
	Requirements	<ul style="list-style-type: none"> <li>8 - 14 mg/day in adults</li> </ul>
	Deficiency	<ul style="list-style-type: none"> <li>Dietary deficiency unusual but is more likely in: <ul style="list-style-type: none"> <li>Vegetarians</li> <li>Conditions when absorption is impaired, such as in Crohn's disease</li> <li>In areas where unleavened whole wheat bread is a major dietary staple</li> </ul> </li> <li>Symptoms: growth retardation, hypogonadism, delayed wound healing, disturbances in taste and smell</li> </ul>
	Excess	<ul style="list-style-type: none"> <li><u>Acute toxicity</u>: NVD, gastric bleeding at intakes &gt; 500 mg/day</li> <li><u>Chronic excess</u>: copper deficiency (&gt;150 mg/day for &gt; 6 weeks)</li> <li><u>UL</u>: 40 mg/day in adults</li> </ul>
	Use in common cold	<ul style="list-style-type: none"> <li>Zinc supplementation (≥ 75 mg/day) may reduce <i>duration</i> but not severity of colds</li> <li>Short-term use of zinc (&lt; 5 days) not associated with severe adverse effects</li> <li>Lozenges associated with unpleasant taste and nausea</li> <li>Intranasal zinc associated with loss of sense of smell</li> </ul>

Fluoride	Function	<ul style="list-style-type: none"> <li>Prevents dental caries (cavities) by:             <ol style="list-style-type: none"> <li>Inhibiting plaque                 <ul style="list-style-type: none"> <li>May kill or inhibit bacteria, inhibits their ability to produce acid from carbs</li> </ul> </li> <li>Inhibiting demineralization and enhancing remineralization of tooth enamel                 <ul style="list-style-type: none"> <li>Incorporated into tooth surface making it more resistant to acid, and speeding remineralization</li> </ul> </li> </ol> </li> </ul>
	Requirement	<ul style="list-style-type: none"> <li>AI: from 0.01 mg/day (infants) to 4 mg/day (adult men)</li> </ul>
	Deficiency	<ul style="list-style-type: none"> <li>Increased rate of tooth decay</li> <li>Dental cavities – initiated by demineralization of tooth surface by acids produced by sugar-fermenting bacteria</li> </ul>
	Sources	<ul style="list-style-type: none"> <li>Major dietary source of fluoride is drinking water</li> <li>Toothpaste, mouthwash often fluoridated</li> <li>Also contained in many prepared drinks &amp; food</li> </ul>
	Excess	<ul style="list-style-type: none"> <li>Chronic excess during tooth formation: dental fluorosis (mottled enamel of teeth)</li> <li>At higher doses: osteosclerosis (increase in bone density &amp; calcification)</li> <li>Acute toxicity: death from respiratory paralysis or cardiac failure</li> </ul>
	Supplementation	<ul style="list-style-type: none"> <li>Not recommended for children living in areas with fluoridated water</li> <li>Supplements before eruption of first permanent tooth generally not recommended             <ul style="list-style-type: none"> <li>If used, total fluoride intake from all sources should be <math>\leq 0.05 - 0.07</math> mg/kg/d to minimize risk of dental fluorosis</li> </ul> </li> <li>If used, lozenges or chewable tablets are preferred forms of supplementation</li> <li>Children 3-6 years of age should brush twice a day with “pea-sized” amount of fluoridated toothpaste</li> </ul>

## LECTURE 21 – POPULAR DIETS

### 1. List 3 reasons why people might consider dieting.

- To improve athletic performance
- To lose weight/improve appearance (concerns with body image due to social pressures)
- To become healthier (physical health) - >60% of Cdns are “overweight” and obesity is a risk factor for many diseases

### 2. Describe:

#### a. Why some diets are so popular

- Obesity is an issue all over the world – by restricting intake, people lose weight
- Huge industry – lots of media attention/marketing to get people interested
- Some are very easy to follow – “avoid carbs”
- Offer a plan/structure – don’t have to think about/plan meals
- Come in all shapes & sizes (something for everyone) – tailored to losing wt, gaining muscle, diseases, etc
- May ignore traditional dietary recommendations – “eat only steak, no potatoes”

#### b. How you can identify a “fad diet”.

DOES IT:

- Sound too good to be true? – promising extreme and rapid weight loss
- Recommend only using a single food & unlimited quantities? – ex// the Cabbage Soup Diet
- Eliminate entire food groups? – can potentially lead to nutritional deficiencies
- Recommend specific food combinations?
- Provide only anecdotal cases to support its effectiveness?
- Require special products or specific supplements?
- Lack a plan for long-term maintenance of weight loss?



- Recommends a very strict/rigid menu?
- Lack recommendations regarding exercise?
- Forget to mention:
  - Any additional costs for supplements, coaching, recipes, etc?
  - Risks associated with the diet plan?

3. Explain why it is important for a pharmacist to be aware of “popular” diets.

- Are one of the most trusted and accessible HCPs and are uniquely positioned in the public eye
- Can help contribute by being a resource to the public for general nutritional advice
- Can work collaboratively with dieticians and understand when to refer patients
- Provide access to and information on various nutritional supplements
- Play a significant role as a HCP to pts with chronic disease where self-management is key (non-drug measures)

4. Identify & describe 5 popular diets (diet characteristics, theory, claims, potential benefits and risks).

a. The 5:2 diet (Mosley Diet)

Theory	<ul style="list-style-type: none"> <li>• Intermittent fasting and/or energy restriction           <ul style="list-style-type: none"> <li>○ Lower energy intake accelerates fat/weight loss</li> <li>○ Forces the body to use up blood glucose, then stored glucose (glycogen), then fat stores</li> </ul> </li> </ul>
Claims	<ul style="list-style-type: none"> <li>• Weight loss: 1 lb/week</li> <li>• Improves brain function</li> <li>• Reduces risk of heart disease, stroke, certain cancers,</li> <li>• Promotes longevity</li> </ul>
Pros & health benefits	<ul style="list-style-type: none"> <li>• Only need to count calories 2 days of the week</li> <li>• Can be flexible with meal plan on fasting days</li> <li>• Not restrictive in foods you can eat</li> <li>• Some studies do suggest health-related benefits with intermittent fasting</li> </ul>
Cons & health risks	<ul style="list-style-type: none"> <li>• Fasting SEs: dizziness, irritability, headaches, insomnia, bad breath, dehydration</li> <li>• If fasting for prolonged periods of time:           <ul style="list-style-type: none"> <li>○ Loss of lean body mass</li> <li>○ Potential for development of nutrient deficiencies</li> </ul> </li> <li>• Not recommended in: pregnancy, diabetes, children/teens, eating disorders</li> <li>• Hard to follow – people feel hungry and may overeat on non-fasting days</li> </ul>
Bottom Line	<ul style="list-style-type: none"> <li>• Fasting days may be challenging and may not be sustainable           <ul style="list-style-type: none"> <li>○ Don't fast 2 days in a row</li> <li>○ Keep hydrated</li> <li>○ Eat nutritiously dense food on fasting days</li> </ul> </li> <li>• Limited evidence for benefits (weight loss) and information on long-term effects is lacking</li> </ul>

b. The Atkins Diet

The Diet	<ul style="list-style-type: none"> <li>• Low carb (high protein) diet that has 4 phases</li> <li>• Refers to carb intake as “net carbs” = total carb content of food – fibre &amp; sugar alcohols           <ul style="list-style-type: none"> <li>○ # grams (net) impacts blood sugar levels</li> </ul> </li> <li>• Atkins 20 (classic) and Atkins 40 (new Atkins)</li> </ul>	
	Phase I (Induction)	<ul style="list-style-type: none"> <li>• Minimum of 2 weeks duration and/or until at least 15 lbs from goal weight</li> <li>• Allowed only 20 g (net) carbs per day – must follow list           <ul style="list-style-type: none"> <li>○ Mainly fish, poultry, and other meats (devoid of net carbs)</li> <li>○ Also target 12-15 g (net) carbs from veggies (low starch)</li> </ul> </li> </ul>
	Phase II (Ongoing Weight Loss)	<ul style="list-style-type: none"> <li>• Allowed 25 g (net) carbs per day and increase by 5 net increments/week           <ul style="list-style-type: none"> <li>○ Target range: 25 – 50 g net carbs / day</li> </ul> </li> <li>• Must still follow the list           <ul style="list-style-type: none"> <li>○ Broader selection– can begin to add nuts, seeds, berries</li> </ul> </li> <li>• Continue until wt stops going down, then remove 5 g net carb from daily intake until losing wt again           <ul style="list-style-type: none"> <li>○ Duration until 10 lbs of goal weight</li> </ul> </li> </ul>

	Phase III (Maintenance)	<ul style="list-style-type: none"> <li>Gradual increase in net carb by 5-10 g increments <ul style="list-style-type: none"> <li>Target range: 50 – 80 g net carbs / day</li> </ul> </li> <li>Reintroduce new carbs as long as wt loss remains gradual &amp; is maintained <ul style="list-style-type: none"> <li>Continue this phase until reach goal weight &amp; is maintained for 1 m</li> </ul> </li> </ul>
	Phase IV (Lifetime Maintenance)	<ul style="list-style-type: none"> <li>Add more carb sources back into diet – as long as weight doesn't increase <ul style="list-style-type: none"> <li>Target range: 80 – 100 g net carbs / day</li> </ul> </li> </ul>
Theory	<ul style="list-style-type: none"> <li>Overconsumption of refined carbs are the main reason for weight gain/obesity</li> <li>Low carb diet triggers body to burn stored fat → weight loss (via ketosis) <ul style="list-style-type: none"> <li>Loss of appetite, loss of bound water</li> <li>Eat more protein which is more filling</li> </ul> </li> </ul>	
Claims	<ul style="list-style-type: none"> <li>Weight loss occurs and is maintained <ul style="list-style-type: none"> <li>Phase 1 = rapid weight loss (15 lbs in 2 wks)</li> <li>Phase 2 = 2-3 lbs loss</li> </ul> </li> <li>Improved heart health &amp; brain function</li> <li>Decreases risk factors for chronic diseases (CVD, diabetes)</li> <li>Modified Atkins type of diets might also help prevent seizures in adults</li> </ul>	
Pros	<ul style="list-style-type: none"> <li>Rapid weight loss in beginning</li> <li>Feel less hungry so eat less overall</li> <li>Might be easier to stick with compared to low fat diets</li> <li>Easy to follow – can purchase premade bars, shakes, meals</li> </ul>	
Cons / health risks	<ul style="list-style-type: none"> <li>SEs of limiting carbs: bad breath, nausea, tiredness, dizziness, insomnia, constipation</li> <li>Carbs may be replaced with high fat, processed meat – may lead to other concerns (CVD)</li> <li>Increased intake of protein may cause issues with kidney stones, osteoporosis?</li> <li>Not recommended in pregnancy</li> </ul>	
Bottom line	<ul style="list-style-type: none"> <li>Ok for short term use if weight loss occurs</li> <li>Long-term efficacy effects on overall health &amp; disease prevention are unknown</li> <li>No clear evidence that it is better than any other weight loss plan</li> <li>Hard to stick to long-term – can be complicated and time consuming to follow</li> </ul>	

c. The Dukan Diet

The diet	<ul style="list-style-type: none"> <li>Overall low carb and high protein → can eat as much as you want of the approved foods</li> <li>Oat bran supplementation and LOTS of water is recommended</li> <li>Exercise is part of the diet routine</li> <li>Composed of 4 phases <ul style="list-style-type: none"> <li>Alcohol only after reached phase 3 or 4</li> </ul> </li> </ul>	
	Phase I (Attack Phase)	<ul style="list-style-type: none"> <li>Usually lasts for 10 days</li> <li>Lean protein (chicken, turkey, eggs, fish) is allowed – must be on approved list (68 listed) <ul style="list-style-type: none"> <li>Unlimited portions</li> </ul> </li> <li>NO CARBS, including no veggies <ul style="list-style-type: none"> <li>Only 1.5 tbsp oat bran is allowed</li> </ul> </li> <li>6 cups of water daily</li> <li>Walk 20 mins per day</li> </ul>
	Phase II (Cruise Phase)	<ul style="list-style-type: none"> <li>Usually lasts for several months (3 days for each lb you want to lose)</li> <li>Add non-starchy veggies (greens) every other day (unlimited) – list of 32</li> <li>2 tbsp oat bran</li> <li>Maintain water intake (6 cups/day)</li> <li>Walk 30 mins per day</li> </ul>
	Phase III (consolidation / transition)	<ul style="list-style-type: none"> <li>Duration depends on weight loss (lasts 5 days for every lb lost)</li> <li>Add fruit, whole grain bread, hard cheese, starchy foods &amp; “celebration meals” – all very limited and prescriptive in quantity</li> <li>2 tbsp oat bran; maintain water (6 cups/day)</li> <li>Walk 25 mins per day</li> </ul>
	Phase IV (stabilization/ maintenance)	<ul style="list-style-type: none"> <li>Can eat what you want for 1 day/week; must follow rules re: protein consumption outlined in Phase I</li> <li>3 tbsp oat bran</li> <li>Maintain water (6 cups/day)</li> <li>Walk 20 mins per day</li> </ul>
Theory	<ul style="list-style-type: none"> <li>High protein – protein takes more time to digest, makes you feel fuller longer, has less calories per gram compared to carbs</li> <li>Low carb – body will burn off fat stores if carb intake is limited</li> </ul>	

Claims	<ul style="list-style-type: none"> <li>• Weight loss: 4-7 lbs in attack phase, 2-4 lbs in cruise phase, then 2lbs/week</li> </ul>
Pros/benefits	<ul style="list-style-type: none"> <li>• Weight loss at beginning is fast</li> <li>• No need to count calories</li> <li>• Very prescriptive and thus easy to follow</li> <li>• Online coaching, books, recipes, supports</li> <li>• Relies on natural foods</li> <li>• Eat as much as you like of the allowed foods</li> <li>• Emphasis on lean protein</li> </ul>
Cons/health risks	<ul style="list-style-type: none"> <li>• SEs low carbs: tiredness, dizziness, bad breath, insomnia, dry mouth, constipation, lethargy</li> <li>• Potential for nutrient deficiencies <ul style="list-style-type: none"> <li>○ Fiber supplement is required</li> <li>○ Recommendation to stay in early phases for too long</li> <li>○ Limit fruits and vegetables</li> </ul> </li> <li>• Not a lot of variety</li> <li>• Can be an issue for diabetes (esp. first 2 phases due to low carbs), renal disease?, heart disease?, vegetarians (no beans, nuts, lentils), pregnancy, depression</li> </ul>
Bottom line	<ul style="list-style-type: none"> <li>• May see quick weight loss in beginning stages but unlikely unsustainable – doesn't change overall eating habits for long term</li> <li>• May regain weight lost when go back to normal eating</li> <li>• Could be in a phase for a long time that could lead to nutritional deficiencies</li> </ul>

**d. The Paleo Diet**

The diet	<ul style="list-style-type: none"> <li>• “Caveman” diet, based on Paleolithic era → food that can be hunted/gathered</li> <li>• In general: lean protein, fish, nuts, eggs, veggies, fruits <ul style="list-style-type: none"> <li>○ Generally no upper/lower limits included – more about quality of food</li> </ul> </li> <li>• Lots of variations <ul style="list-style-type: none"> <li>○ Strict paleo – avoid dairy, grains, processed food, sugar, legumes starches, alcohol</li> <li>○ Low carb / high protein</li> </ul> </li> </ul>
Theory	<ul style="list-style-type: none"> <li>• Today's health issues &amp; diseases are a result of eating highly processed dense foods <ul style="list-style-type: none"> <li>○ Obesity, diabetes, heart disease, cancer, Parkinson's, Alzheimer's, depression</li> </ul> </li> <li>• Exclude food that humans have not genetically “adapted” to</li> <li>• Increase lean protein = satiating and helps control appetite</li> <li>• Increasing healthy fats = decreases gastric emptying, full longer</li> <li>• Encouraging veggies and fruits = antioxidants, vits, minerals (good for chronic disease)</li> </ul>
Claims	<ul style="list-style-type: none"> <li>• Better athletic performance?</li> <li>• Weight loss quick in beginning and maintained (5-10 lbs in first wks, then 1-2 lb per week)</li> <li>• Reduces risk of diseases (diabetes, heart disease, cancer)</li> <li>• Anti-inflammatory?</li> </ul>
Pros	<ul style="list-style-type: none"> <li>• Wide range of food acceptable</li> <li>• Encourages eating of nutrient dense foods and less processed food</li> <li>• Rapid weight loss in short term (and is maintained)</li> <li>• Diet also recommends exercise</li> <li>• Fairly easy to follow/understand – don't need to count calories</li> </ul>
Cons/ health risks	<ul style="list-style-type: none"> <li>• Can be costly</li> <li>• Strict paleo excludes food groups (dairy &amp; grains) and can be difficult to follow 100%</li> <li>• Can be an issue for: vegetarians</li> </ul>
Bottom Line	<ul style="list-style-type: none"> <li>• Antidote for Western Diet?</li> <li>• Can lose weight</li> <li>• High in protein and fibre – you shouldn't feel hungry and overeat</li> <li>• Exercise is highly recommended as an adjunct to diet</li> <li>• Recommends lifestyle changes and changes to eating habits – more likely to be maintained</li> </ul>

e. The South Beach Diet

The diet	<ul style="list-style-type: none"> <li>• Low carb, low sugar (high protein), high fibre, lean protein and unsaturated fats <ul style="list-style-type: none"> <li>○ Allows certain carbs based on their GI</li> </ul> </li> <li>• Recommends 3 meals a day + 2 snacks</li> <li>• 3 phases – gradual increase in carbs, while decreasing proportion of fat &amp; protein <ul style="list-style-type: none"> <li>○ No restrictions on calorie intake</li> </ul> </li> <li>• Recommends exercise</li> </ul>
Phase I (Body Reset)	<ul style="list-style-type: none"> <li>• Designed to try to eliminate cravings for sugary food/refined starches and jump start weight loss</li> <li>• Structured – 3 meals a day, 2 snacks, dessert <ul style="list-style-type: none"> <li>○ Cut out almost ALL carbs (avoid bread, rice, potatoes, pasta, sugary foods)</li> <li>○ Lean meats, nuts, beans, veggies</li> </ul> </li> <li>• Usually lasts 2 weeks</li> </ul>
Phase II (Steady Weight Loss)	<ul style="list-style-type: none"> <li>• Aim is to be the longer term weight loss phase <ul style="list-style-type: none"> <li>○ Maintain this phase until reach goal weight</li> </ul> </li> <li>• Some carbs are reintroduced in moderation (fruit, whole grains, veggies)</li> <li>• Approach is structured – one single carb reintroduced for 1 meal/day/week &amp; response is monitored <ul style="list-style-type: none"> <li>○ Continue if tolerated &amp; until 2-3 servings of right carbs introduced to daily meal plan</li> <li>○ If revert back to old unhealthy eating or weight regained, go back to phase I</li> </ul> </li> </ul>
Phase III (You Got This)	<ul style="list-style-type: none"> <li>• Start when target weight reached – aim to be maintenance phase</li> <li>• Adopting the lifestyle of making good food choices &amp; maintaining weight loss, and eating foods in moderation <ul style="list-style-type: none"> <li>○ Choose right carbs – up to 140 g carbs / day</li> <li>○ If cravings return/weight regained, go back to an earlier phase</li> </ul> </li> </ul>
Theory	<ul style="list-style-type: none"> <li>• Eating carbs with a higher GI increases blood sugars faster and for longer periods of time <ul style="list-style-type: none"> <li>○ Stimulates appetite and causes people to overeat</li> </ul> </li> <li>• Key to weight loss is eating the right carbs/fats <ul style="list-style-type: none"> <li>○ Eating “good” carbs can keep metabolism and sugar levels steady (not constantly spiking/fluctuating)</li> </ul> </li> </ul>
Claims	<ul style="list-style-type: none"> <li>• Weight loss (and maintenance): 8-13 lbs in phase I; 1-2 lbs/wk in phase II</li> <li>• Prevents some chronic diseases (diabetes, heart disease)</li> </ul>
Pros	<ul style="list-style-type: none"> <li>• Promotes consumption of veggies, lean meats and restricts fatty meats, cheeses &amp; sweets</li> <li>• No need to count calories so easy to follow for some</li> </ul>
Cons / health risks	<ul style="list-style-type: none"> <li>• SEs of low carbs: bad breath, tiredness, dizziness, insomnia, nausea, constipation</li> <li>• First phase has lots of restrictions and can be hard for some</li> <li>• Weight loss may be regained when eat normally</li> <li>• Lacks dietary fibre</li> </ul>
Bottom line	<ul style="list-style-type: none"> <li>• May be challenging to get through first phase</li> <li>• May be ok for short-term weight loss</li> <li>• No data on long-term safety and efficacy for weight loss maintenance and health related outcomes/benefits</li> </ul>

5. List some common components (principles) of a “good” diet plan.

- Easy to follow/maintain
- Focuses on healthy eating and exercise as regular habit
- Recommends nutrient-dense, high-fiber foods more often
- Limits alcohol, sugar, fat
- Has reasonable goals: decrease overall calorie intake, fat intake, exercise regularly; 0.5 – 2 lbs/wk (10-20 lbs/yr)
- Promotes long-term weight loss & maintenance and describes ways to achieve this
- Emphasizes improvements in quality of life as a success too

6. Identify reasonable approaches to achieve and maintain a healthy body weight.

- Make well balanced health food choices – choose nutrient-dense and fibre-rich foods more often
- Avoid over-processed foods
- Be realistic about overall energy intake – energy out > energy in by 500 kcal/d for weight loss
- Be mindful of portion size
- Limit fat intake to 20-35% of total calories & choose healthy fats more often
- Use little added sugar or additional salt
- Alcohol use in moderation
- Exercise regularly
- Slow down – faster eating >> higher energy intake >> increase body weight
- Drinks LOTS water
- Set goals
- Plan meals ahead of time

**LECTURE 23 – DRUG-NUTRIENT INTERACTIONS & ENTERAL FEEDING**

1. Identify and suggest appropriate resolution of the select drug-nutrient and drug-enteral food interactions presented on the lecture slides.

Types	<ul style="list-style-type: none"> <li>• <u>Chemical</u>: binding the drug and decreasing its absorption</li> <li>• <u>Physical</u>: adsorption of drug and nutrient (or feed formulation) → blockage <ul style="list-style-type: none"> <li>○ Ex// sucralfate and enteral feed</li> </ul> </li> <li>• Interaction between drug and a specific nutrient involved in the drug's metabolism</li> </ul>
Pantoprazole	<ul style="list-style-type: none"> <li>• Possible to use via NG tube (crush and dissolve tablet)</li> <li>• Consider parenteral route or changing therapy to lansoprazole or omeprazole</li> </ul>
Ciprofloxacin	<ul style="list-style-type: none"> <li>• Ciprofloxacin binds to divalent ions in the feed (reduced absorption) <ul style="list-style-type: none"> <li>○ Try administering dose during a break in feeding where possible</li> <li>○ Alternatively consider alternative quinolone (interaction is less)</li> </ul> </li> <li>• Ciprofloxacin 750 mg tablets can be crushed and dissolved (delivered via gastric tube)</li> <li>• Suspension (granules) = very thick, non-aqueous so high risk of tube blockage with fine tubes</li> </ul>

2. Discuss indications and contraindications for enteral feeding.

Indications	<ul style="list-style-type: none"> <li>• Patient with a functional GIT but where oral intake is unsafe, insufficient, or impossible <ul style="list-style-type: none"> <li>○ Support maintenance of functional integrity of gut</li> <li>○ Nutrients undergo first-pass metabolism</li> </ul> </li> </ul>	
Contraindications	Absolute	Possible
	<ul style="list-style-type: none"> <li>• Necrotizing enterocolitis</li> <li>• Bowel obstruction</li> <li>• Hemodynamic instability</li> </ul>	<ul style="list-style-type: none"> <li>• Persistent vomiting or diarrhea</li> <li>• Acute abdominal distension</li> <li>• Fistula (gastric, small or large bowel)</li> <li>• Upper GI bleeding</li> </ul>

3. Compare and contrast the different types of enteral formula.

Polymeric	<ul style="list-style-type: none"> <li>• Most common type</li> <li>• Least expensive</li> <li>• 1.5 – 2 L/day provides 100% RDA for vitamins/minerals</li> <li>• Require full digestive function for proteins, carbs and fats</li> </ul>			
	<b>Caloric density</b>	<b>Nutrient source</b>	<b>Fibre-content</b>	<b>Protein content</b>
	<ul style="list-style-type: none"> <li>• Standard: 1-1.2 kcal/mL</li> <li>• Mod: 1.5 kcal/mL</li> <li>• Dense: 2kcal/mL</li> </ul>	<ul style="list-style-type: none"> <li>• Milk-based</li> <li>• Lactose-free</li> <li>• Blenderized</li> </ul>	<ul style="list-style-type: none"> <li>• Fibre-free</li> <li>• Low-mod (1-8 g/L)</li> <li>• High (&gt;8 g/L)</li> </ul>	<ul style="list-style-type: none"> <li>• Low: nitrogen 6% of kcal</li> <li>• Standard: 11-15%</li> <li>• High: 16-25%</li> </ul>

Oligomeric	<ul style="list-style-type: none"> <li>Require minimal digestive function for proteins and medium-chain triglycerides <ul style="list-style-type: none"> <li>Often contain free amino acids</li> <li>Some pancreatic enzyme activity required for digestion of fats and oligosaccharides</li> <li>Some brush border disaccharidase activity required</li> </ul> </li> <li>Leave little residue in the colon</li> <li>Often used for patients with severe pancreatic insufficiency or short-bowel syndrome</li> </ul>	
Specialized	Renal failure	<ul style="list-style-type: none"> <li>Low-protein, low electrolytes</li> <li>Enriched with essential amino acids</li> </ul>
	Hepatic failure	<ul style="list-style-type: none"> <li>Contain 45-50% of protein as BCAAs (vs. 15-20% in standard formulas)</li> </ul>
	Glucose control	<ul style="list-style-type: none"> <li>Lower carb, higher fat (ex// glucerna)</li> </ul>
	Stress/ critically ill	<ul style="list-style-type: none"> <li>Designed for hypermetabolic patients (ex// high nitrogen content)</li> </ul>
	Pulmonary	<ul style="list-style-type: none"> <li>Increased percentage of calories from fat</li> </ul>

4. Compare and contrast the different types of enteral tubes with respect to size, material type, and placement and the benefits/disadvantages of each for the patient.

Feeding Tube Size	<ul style="list-style-type: none"> <li>External diameter of the feeding tube is measured in the French (Fr) unit <ul style="list-style-type: none"> <li>1 Fr = 0.33 mm</li> </ul> </li> </ul>			
Feeding tube position	Nasogastric (NG)	<ul style="list-style-type: none"> <li>Inserted via nose &amp; exits the stomach</li> <li>Size can vary (6 - 16 Fr); length 90 - 100 cm</li> </ul>	<ul style="list-style-type: none"> <li>Inserted via nasopharynx</li> <li>Used for days to weeks</li> </ul>	
	Nasoduodenal (ND)	<ul style="list-style-type: none"> <li>Inserted via nose &amp; exits in duodenum; usually with endoscopic or radiological imaging</li> </ul>		
	Nasojejunal tube (NJ)	<ul style="list-style-type: none"> <li>Inserted via nose &amp; exits in jejunum</li> <li>Usually inserted radiologically or endoscopically to ensure correct positioning</li> <li>Length: 150 cm = prone to blockage</li> <li>Lack of evidence re: drug absorption from this site</li> </ul>		
	Percutaneous gastrostomy	<ul style="list-style-type: none"> <li>Inserted via abdominal wall into stomach (usually endoscopically)</li> <li>Held in place with internal balloon &amp; external fixator</li> </ul>	<ul style="list-style-type: none"> <li>Direct access to GIT through skin</li> <li>Used for months to years</li> </ul>	
	Percutaneous jejunostomy			
Material Type	Selection varies with	<ul style="list-style-type: none"> <li>Intended duration of use</li> <li>Part of GIT that the feed is delivered to</li> </ul>		
	Materials		<b>Advantages</b>	<b>Disadvantages</b>
		<b>Polyurethane</b>	<ul style="list-style-type: none"> <li>More rigid</li> <li>Less flexible</li> </ul>	<ul style="list-style-type: none"> <li>Larger internal diameter compared to latex/silicone tube of same Fr size</li> </ul>
		<b>Polyvinylchloride</b>		
		<b>Silicone</b>	<ul style="list-style-type: none"> <li>Softer</li> </ul>	<ul style="list-style-type: none"> <li>Requires thicker wall to prevent stretch/ collapse</li> </ul>
<b>Latex</b>	<ul style="list-style-type: none"> <li>More flexible</li> </ul>			

5. Discuss the different types of enteral feeding products and their advantages and disadvantages.

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6. Discuss the general considerations for administering medications through enteral feeding tubes.

Implications for drug administration	<ul style="list-style-type: none"> <li>Size of lumen &amp; length of tube - narrow and/or long tubes more likely to become blocked</li> <li>Site of drug delivery: <ul style="list-style-type: none"> <li>Different pH</li> <li>Delivery beyond site of absorption</li> <li>Reduction in duration of contact with GIT</li> <li>SE may be increased due to rapid delivery of drug to lumen of small bowel</li> </ul> </li> </ul>
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Tube blockage due to	<ul style="list-style-type: none"> <li>• Small internal diameter of tube</li> <li>• Poor flushing technique or regimen</li> <li>• Gastric acid, feed and medication interactions</li> <li>• Inappropriately prepared medications (ex// inadequately crushed tablets)</li> </ul>
Medication administration key points	<ul style="list-style-type: none"> <li>• Consider if patient may be able to take some medications orally</li> <li>• Avoid adding medications directly to enteral feeding formula</li> <li>• Avoid mixing crushed or liquid medications together prior to administration</li> <li>• Change to medications with longer <math>t_{1/2}</math> but avoid use of modified/slow-release medications</li> <li>• Monitor for changes in therapeutic response</li> </ul>

7. Discuss specific medication and dosage form considerations for their administration through enteral feeding tubes.

Liquid		Advantages	Disadvantages
	Solutions	<ul style="list-style-type: none"> <li>• Ready to use</li> <li>• Accurate dosing</li> </ul>	<ul style="list-style-type: none"> <li>• Co-solvents may have p'col effect if present in sufficient quantities (ex// sorbitol &gt; 15 g/d)</li> </ul>
	Suspension	<ul style="list-style-type: none"> <li>• Usually ready-to-use</li> </ul>	<ul style="list-style-type: none"> <li>• Shorter stability/shelf-life</li> <li>• Requires adequate shaking for accurate dosing</li> <li>• Further dilution usually required (viscosity)</li> </ul>
		KEY POINTS: <ul style="list-style-type: none"> <li>• Granules in suspension may be too large to pass through feeding tube (ex// ciprofloxacin, clarithromycin, lansoprazole)</li> <li>• Suspension may be too viscous to pass through feeding tube</li> </ul>	
Solid		Advantages	Disadvantages
	Soluble Tablets	<ul style="list-style-type: none"> <li>• Drug is in solution for admin</li> <li>• Long expiry date</li> <li>• Often less costly than liquid</li> </ul>	<ul style="list-style-type: none"> <li>• Time for complete dissolution required</li> </ul>
	Effervescent Tablets	<ul style="list-style-type: none"> <li>• Long shelf-life</li> <li>• Less costly than liquids</li> </ul>	<ul style="list-style-type: none"> <li>• Larger volume of water required</li> <li>• Time for dispersion</li> <li>• Sodium content may be too high</li> <li>• Production of CO<sub>2</sub> makes syringe administration difficult</li> </ul>
	Tablets	<ul style="list-style-type: none"> <li>• Cheap (often)</li> <li>• Easily obtained</li> <li>• Most disintegrate in water</li> <li>• Crushing maybe unnecessary</li> </ul>	<ul style="list-style-type: none"> <li>• Not all tablets disintegrate easily (require crushing)</li> <li>• Crushing EC tablets will likely block tube; also absorption may be decreased</li> </ul>
	Capsules	<ul style="list-style-type: none"> <li>• Cheap</li> <li>• Often convenient (hard)</li> </ul>	<ul style="list-style-type: none"> <li>• Occupational exposure</li> <li>• Small capsules difficult to open</li> </ul>
		CAPSULE TYPES: <ul style="list-style-type: none"> <li>• Hard gelatin: open and contents mixed with water</li> <li>• Soft gelatin: usually contain oil; poorly soluble in water</li> </ul>	

**LECTURE 24 - TOTAL PARENTERAL NUTRITION**

1. To understand when TPN should be initiated.

- Inability to absorb adequate nutrients enterally (massive small bowel resection, severe malabsorption, severe vomiting/diarrhea, radiation enteritis)
- Bowel rest (ex// moderate to severe pancreatitis, IBD)
- Severe malnutrition with non-functional GIT
- Time frame
  - Non-usable GIT x 5-7 days
  - Within 1-3 days if severely malnourished with non-functioning GIT (i.e. cannot feed enterally)

2. To be able to assess a TPN formula:

Basal Energy Expenditure (BEE) x stress factor	kCal/kg/day
Maintenance x 1.2 – 1.3	No stress: 20 – 25
Surgery*, minor infection x 1.5	*Mod Stress: 25-30
Major surgery, sepsis, tumor x 1.5 – 2.0	Severe stress: 30-35

BEE calculations: ht (cm); wt (kg); age (years)

- Males:  $66 + (13.7 \times wt) + (5 \times ht) - (6.8 \times age)$
- Females:  $665 + (9.6 \times wt) + (1.7 \times ht) - (4.7 \times age)$

NOTE: use IBW as basis of calculations (formula on cheat sheet provided) unless:

- ABW > 120% IBW → use 25% of difference added onto IBW ((ABW-IBW) 0.25 + IBW))
- ABW < IBW → use ABW

a. **Macronutrients (carbohydrate, protein, fat)** – requirements provided on exam!

Protein (4 kCal/g)	Estimated requirements	<ul style="list-style-type: none"> <li>• Maintenance (healthy, unstressed): 0.8 – 1 g/kg/day</li> <li>• Moderate repletion (surgery, trauma): 1.5 g/kg/day</li> <li>• Extensive repletion (burn): 2 – 2.5 g/kg/day</li> </ul>
	Complications	<ul style="list-style-type: none"> <li>• Pre-renal azotemia (increased BUN)</li> </ul>
Dextrose (3.4 kcal/g)	Max requirement	<ul style="list-style-type: none"> <li>• 7 g/kg/day (5 mg/kg/minute)</li> <li>• 50 – 60% of total calories</li> </ul>
	Complications	<ul style="list-style-type: none"> <li>• Synthesis &amp; storage of fat (abnormal LFTs - ↑ AST, ALT) ~ 2-3 wk</li> <li>• Hyperglycemia: monitor blood glucose</li> <li>• Increased CO<sub>2</sub> production</li> </ul>
Lipids/fat (2kcal/mL)	Formulation	<ul style="list-style-type: none"> <li>• Only one formulation (20%)                             <ul style="list-style-type: none"> <li>○ 250 mL contains 4 mmol PO<sub>4</sub> &amp; 133 mcg vit K</li> </ul> </li> <li>• Contains egg phospholipids</li> </ul>
	Requirement	<ul style="list-style-type: none"> <li>• 20-30% total calories (&gt; 4% ; &lt;60%)</li> <li>• Try for &lt; 1 g/kg/day (max 2 g/kg/day)</li> </ul>
	Complications	<ul style="list-style-type: none"> <li>• Pancreatitis</li> <li>• Increased triglycerides (&lt; 4.5 mmol/L)</li> <li>• Fat overload syndrome (accumulation in RES system of liver)</li> </ul>

b. **Micronutrients (electrolytes, vitamins, trace elements).** NOTE. You will NOT be responsible for determining amount of micronutrients to add to TPN.

<b>Electrolytes</b> <ul style="list-style-type: none"> <li>• Na chloride/ acetate</li> <li>• K chloride/acetate</li> <li>• Calcium gluconate</li> <li>• Magnesium chloride</li> <li>• PO<sub>4</sub> (Na or K salt)</li> <li>• Chloride &amp; acetate</li> </ul>	Special requirements	<ul style="list-style-type: none"> <li>• Renal failure</li> <li>• Diuretic use</li> <li>• Amphotericin B</li> <li>• Refeeding syndrome</li> <li>• High fistula output</li> </ul>
Multivitamins	Need	<ul style="list-style-type: none"> <li>• Necessary for normal metabolism and cellular function</li> <li>• Do not contain vit K (only add if intralipid &lt; 200 mL/d) or iron (add iron dextran if &gt; 1 mo)</li> </ul>
	Special requirements	<ul style="list-style-type: none"> <li>• Bursn/Ig wounds: vitamin C</li> <li>• Alcoholic: thiamine, folic acid</li> <li>• Refeeding: thiamine, folic acid (coenzymes for carb metabolism)</li> </ul>
<b>Trace Elements</b> <ul style="list-style-type: none"> <li>• Cr, Cu, I, Mn, Se, Zn</li> </ul>	Need	<ul style="list-style-type: none"> <li>• Essential for proper functioning of several enzyme systems</li> </ul>
	Special Requirements	<ul style="list-style-type: none"> <li>• Extra zinc:                             <ul style="list-style-type: none"> <li>○ Burn/large wounds</li> <li>○ High fistula output/excessive diarrhea</li> </ul> </li> </ul>



3. To know how to start and monitor patients on TPN.

Fluid requirements	<ul style="list-style-type: none"> <li>Increased in fever, fistulas, diarrhea, NG suction → may require separate IV for replacement</li> <li>Decreased in: renal failure, CHF, cirrhotic ascites, pulmonary disease → concentrate macronutrients (dextrose 70%)</li> </ul>	
Monitoring	<ul style="list-style-type: none"> <li><i>Daily x 5, then twice weekly:</i> electrolytes, glucose QID, phosphate, magnesium</li> <li><i>Twice weekly x 2, then weekly:</i> albumin, calcium, urea, creatinine</li> <li><i>Weekly:</i> CBC, INR, PTT, triglycerides, LFTs (alk phos, ALT, AST, GGT)</li> <li><i>Other:</i> weights twice weekly; if on insulin, glucometer QID; other meds/conditions that can affect glucose</li> </ul>	
Feeding techniques	<b>Advantages central line</b>	<b>Disadvantages peripheral line</b>
	<ul style="list-style-type: none"> <li>Hypertonic solution okay</li> <li>More balanced formulas</li> <li>Long-term TPN</li> </ul>	<ul style="list-style-type: none"> <li>Max osmolality = 600-900 mOsm/L <ul style="list-style-type: none"> <li>Dextrose 10% = 505 mOsm/L</li> </ul> </li> <li>Major calories from FAT</li> </ul>
Discontinuing TPN	Without taper	<ul style="list-style-type: none"> <li>Once patient tolerates DAT and is taking 50% calories PO or enterally</li> <li>Stop TPN by ↓ infusion rate by 50% x 2 hrs, then D/C (VGH)</li> </ul>
	With taper	<ul style="list-style-type: none"> <li>Abrupt discontinuation (ex// CV sepsis and no other carb source)</li> <li>Taper with D5-10W via peripheral line x 2-4 hours</li> </ul>

4. To understanding major complications associated with TPN and their management

a) Refeeding

Description	<ul style="list-style-type: none"> <li>Occurs when acute provision of macronutrients (particularly carbs) that promote <u>anabolism</u> in patients who are malnourished <ul style="list-style-type: none"> <li>a) With starvation, body switches from using carbs to fat &amp; protein as main energy sources</li> <li>b) With re-introduction of carbs (TPN) → sudden shift back to glucose as predominant energy source</li> <li>c) Leads to insulin surge → anabolism → intracellular shift of <b>K, PO<sub>4</sub>, Mg</b></li> </ul> </li> <li>Also need extra <b>thiamine, folic acid</b></li> </ul>
Initiating TPN	<ul style="list-style-type: none"> <li>Start with 15-20 kCal/kg/day (~ 50% of nutritional needs) <ul style="list-style-type: none"> <li>Reduce carbs and fat, okay to give more protein</li> </ul> </li> <li>Increase to goal over 3-7 days</li> </ul>
Electrolyte abnormalities	<ul style="list-style-type: none"> <li><u>Treatment of current situation:</u> give IV boluses of K, Po<sub>4</sub>, Mg</li> <li><u>Prevention:</u> add extra K, Po<sub>4</sub>, Mg to TPN bag <ul style="list-style-type: none"> <li>Monitor electrolytes daily x 5, then twice weekly</li> </ul> </li> </ul>

b) Hyperglycemia

Diabetic pts	<ul style="list-style-type: none"> <li>Reduce initial dextrose to max 100-150 g on day 1 (adjust fat &amp; AA appropriately) <ul style="list-style-type: none"> <li>Increase slowly based on tolerance</li> </ul> </li> <li>Continue basal insulin (ex// glargine, NPH) at full or 2/3 dose</li> <li>Initiate insulin sliding scale orders (shorter-acting insulin; ex// regular or lispro)</li> </ul>
Adding insulin to TPN	<ul style="list-style-type: none"> <li>Only regular insulin is compatible with TPN <ul style="list-style-type: none"> <li>Common starting point: 0.1 units/g dextrose (i.e. 30 units for 300 g dextrose)</li> <li>or</li> <li>Add up total insulin given by sliding scale over 24 hrs and add 1/2 - 2/3 to TPN</li> </ul> </li> <li>Why not add full amount of sliding scale insulin into TPN bag? <ul style="list-style-type: none"> <li>Different routes of administration (TPN is IV, not SC)</li> <li>Safer to run sugars slightly higher than lower</li> </ul> </li> </ul>
Other causes of hyperglycemia	<ul style="list-style-type: none"> <li>Infection/sepsis</li> <li>Corticosteroids</li> <li>Stress – ACTH induces gluconeogenesis</li> <li>Excessive dextrose administration</li> </ul>

c) Elevated liver enzymes

TPN & Liver Function	Liver function changes	<ul style="list-style-type: none"> <li>Mildly elevated transaminase (AST, ALT) and alkaline phosphatase levels usually occur 2 weeks (range 1-4 weeks) after TPN is started</li> </ul>
	High-risk pts	<ul style="list-style-type: none"> <li>Pre-existing liver disease</li> <li>Pre-existing malnutrition</li> <li>Sepsis</li> <li>Extent of bowel resection</li> <li><u>Excess non-protein calories</u></li> <li><u>Little or no oral intake</u></li> <li><u>Duration on TPN</u></li> </ul>
	Drugs	<ul style="list-style-type: none"> <li>Octreotide, ceftriaxone → cause biliary sludging</li> </ul>
Cholestasis	Description	<ul style="list-style-type: none"> <li>↑ GGT, alk phos, bilirubin may reflect cholestasis (gallbladder sludge)</li> <li>Major risk factor = duration of TPN               <ul style="list-style-type: none"> <li>Lack of nutrition in bowel → impaired bile flow</li> </ul> </li> </ul>
	Management	<ul style="list-style-type: none"> <li>Avoid overfeeding</li> <li>Encourage oral/enteral intake (stimulates release of CCK → secretion of bile from gallbladder)</li> <li>If prolonged therapy necessary:               <ul style="list-style-type: none"> <li>Cycle TPN (ex// 12 hrs on; 12 hrs off)</li> <li>Ursodiol 15 mg/kg/day in 2 divided doses (bile acid)</li> </ul> </li> <li>Uncomplicated TPN-related cholestasis usually resolves within 1-4 months after stopping TPN</li> </ul>
Fatty liver (hepatic steatosis)	Description	<ul style="list-style-type: none"> <li>Elevated transaminases (AST, ALT) within 2-4 weeks after TPN initiated</li> <li>Most commonly caused by excessive carbs (&gt;7 g/kg/d) or overfeeding</li> </ul>
	Management	<ul style="list-style-type: none"> <li>Avoid overfeeding; reduce carbs</li> <li>Switch to enteral feeding ASAP</li> <li>Cycle TPN</li> </ul>