#### **Resource Highlight**

#### *UpToDate*



#### **SMHCS Medical Library**



### **UpToDate**

- Evidenced-based point of care clinical medical database providing recommendations across 26 specialties. UpToDate integrates and links to LexiDrug (formerely LexiComp)
- With UpToDate:
  - Answer Questions Quickly
  - Increase Clinical Knowledge
  - Improve Patient Care
  - Earn CME
  - Share Patient Education
- UpToDate is available via website and mobile app.





#### **Quick Overview**

UpToDate		
Evidence-based	More expert opinion,Less well referenced	
Built-in drug info	From Lexicomp	
Patient information	Yes, UTD written materials	
Breadth	Monitors over 460 journals plus Cochrane. Also monitors 10,000+ national and international guidelines.	
Peer Reviewed	Yes	



# **Style of Presentation - UpToDate**

• UpToDate favors longer, **narrative** style delivery.

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	Clinical presentation, diagnosis, and initial e	
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Outline <		
SUMMARY AND RECOMMENDATIONS	CLINICAL PRESENTATION	
INTRODUCTION	• Type 2 diabetes – Type 2 diabetes is by far the most common type of diabe	etes in adults (>90 percent) and is characterized by hyperglycemia usually due to progressive loss of insulin
CLINICAL PRESENTATION	secretion from the beta cell superimposed on a background of insulin resistance, resulting in relative insulin deficiency. The majority of patients are asymptomatic at presentation, with hyperglycemia noted on routine laboratory evaluation, prompting further testing. The frequency of symptomatic diabetes has been decreasing in parallel with improved efforts to diagnose	
DIAGNOSTIC CRITERIA	diabetes earlier through screening. (See "Screening for type 2 diabetes mel	
Diabetes	The classic symptoms of hyperalycemia (including polyuria, polydinsia, poc	turia, blurred vision, and weight loss) are often noted only in retrospect after a blood glucose value has been shown
<ul> <li>Symptomatic hyperglycemia</li> </ul>		s significantly above 180 mg/dL (10 mmol/L), exceeding the renal threshold for glucose reabsorption, which leads to
<ul> <li>Asymptomatic hyperglycemia</li> </ul>	increased urinary glucose excretion. Glycosuria causes osmotic diuresis (ie,	polyuria) and hypovolemia, which in turn can lead to polydipsia. Patients who replete their volume losses with
Prediabetes	concentrated sugar drinks, such as non-diet sodas, exacerbate their hyperg	Jlycemia and osmotic diuresis.
DIAGNOSTIC TESTS	Rarely adults with type 2 diabetes can present with a hyperosmolar hyperg	lycemic state, characterized by marked hyperglycemia, severe dehydration, and obtundation, but without
Fasting and two-hour plasma glucose		e 2 diabetes is also uncommon in adults but may occur under certain circumstances (usually severe infection or
A1C	other acute liness). (see "Diabetic ketoacidosis and hyperosmolar hypergiy mellitus".)	cemic state in adults: Clinical features, evaluation, and diagnosis" and "Syndromes of ketosis-prone diabetes
A1C, FPG, and OGTT as predictors of cardiovascular risk		
DIFFERENTIAL DIAGNOSIS	<ul> <li>Type 1 diabetes – Type 1 diabetes is characterized by autoimmune destruction</li> <li>5 to 10 percent of diabetes in adults.</li> </ul>	tion of the pancreatic beta cells, leading to absolute insulin deficiency. Type 1 diabetes accounts for approximately
Other causes of hyperglycemia	DKA may be the initial presentation in approximately 25 percent of adults w	vith newly diagnosed type 1 diabetes. Compared with children, the loss of insulin secretory capacity usually is less
Classification of diabetes		liabetes typically have a longer estimated period prior to diagnosis and are likely to have more protracted
VALUATION		. In 2 to 12 percent of adults, the clinical presentation is similar to that of type 2 diabetes (older-age onset and not
Comprehensive history	initially insulin dependent), with autoimmune-mediated insulin deficiency d (LADA). (See "Classification of diabetes mellitus and genetic diabetic syndro	leveloping later in the course of disease [1]. This is sometimes referred to as latent autoimmune diabetes of adults
Biochemical testing	(LADA). (See Classification of diabetes mellitus and genetic diabetic syndro	Thes ', section on Eatent autominiate diabetes in adults (EADA) .)
Differentiating the cause		
Diabetes-related complications and comorbid	DIAGNOSTIC CRITERIA	
Rate ☆☆☆☆☆ Topic Feedback	Fasting plasma glucose (EPG), two-bour plasma glucose during a 75 g oral gluc	cose tolerance test (OGTT), or glycated hemoglobin (A1C) may be used for diagnostic testing. OGTT is not commonly



# **Style of Presentation - Grading Labels**

**UpToDate** includes grading labels in the treatment recommendations.

#### SUMMARY AND RECOMMENDATIONS

- Epidemiology Hip fractures are common worldwide and substantially increase the risk of death and major morbidity in older adults. (See "Hip fracture in older adults: Epidemiology and medical management".)
- Clinical anatomy and fracture classification Hip fractures are classified by anatomic location and fracture type. The general categories include intracapsular (femoral neck and head) and extracapsular (intertrochanteric and subtrochanteric) fractures. Fracture classification is described further in the text. Intracapsular fractures have higher rates of nonunion, malunion, and avascular necrosis of the femoral head because the blood supply is more easily disrupted ( $\sim$  figure 2 and  $\sim$  figure 3). (See 'Fracture classification' above and 'Anatomic considerations' above.)
- Initial care Initial management of the patient with a hip fracture consists primarily of providing adequate analgesia and consulting an orthopedic surgeon. Regional nerve blocks can be effective. It is prudent to obtain blood for type and crossmatch in patients with any two of the following risk factors: age over 75 years, initial hemoglobin below 12 g/dL (SI 120 g/L), and a peritrochanteric fracture. We suggest not using skin or skeletal tractor (Grade 2B). (See 'Initial assessment and medical management' above.)

High-quality evidence: Consistent evidence from randomized trials, or overwhelming evidence of some other form

Α

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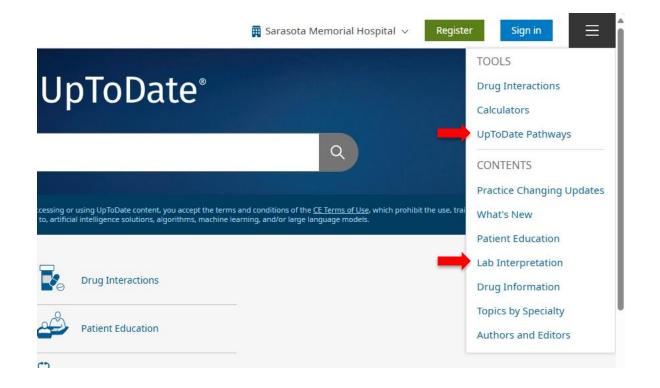
Moderate-quality evidence: Evidence from randomized trials with important limitations, or very strong evidence of some other form

Low-quality evidence: Evidence from observational studies, unsystematic clinical observations, or from randomized trials with serious flaws



# **Unique Tools – Pathways & Lab Interpretation**

- **UpToDate Pathways** provide interactive guides to help make appropriate decisions related to specific clinical questions exclusively utilizing synthesized content from UpToDate and underlying evidence in society guidelines.
- Lab Interpretations enable you to quickly and accurately interpret and decide next steps on abnormal lab results.
- Pathways and Lab Interpretations are available on both the web and mobile app versions of UpToDate.





# Contact the Library for Assistance and in-person or virtual training.

Email: Medical-Library@SMH.com

Website: https://library.smh.com

