



如何搜尋

- 診斷
- 檢查
- 藥物名稱
- 不須特別連接詞
- 直覺式

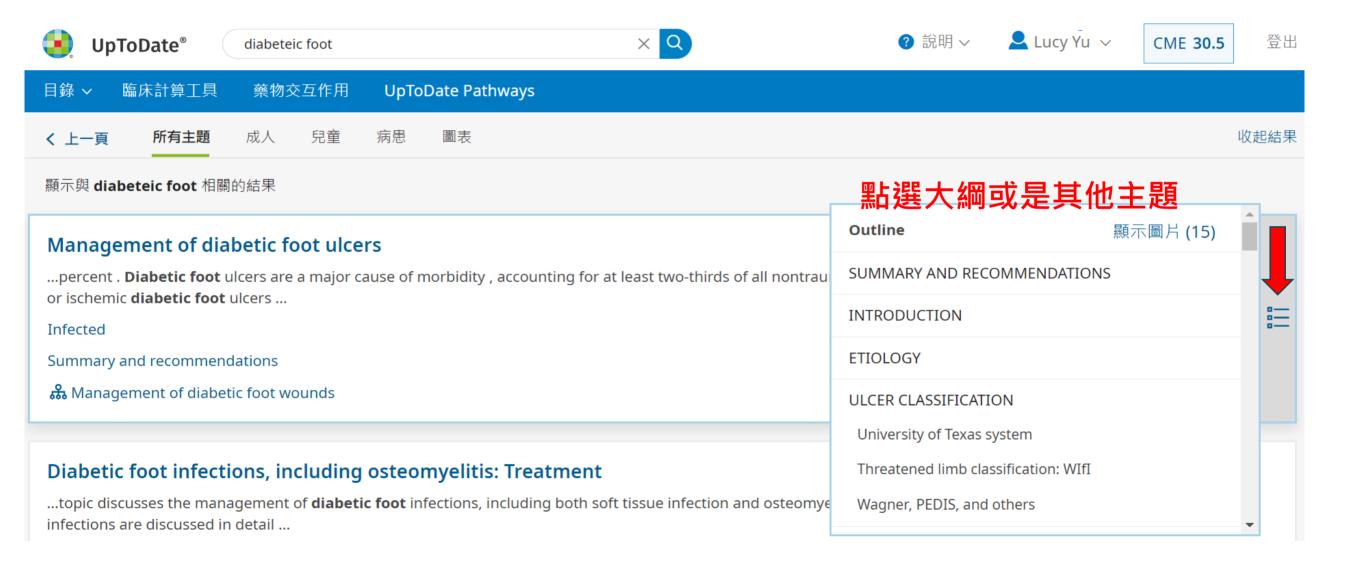
案例:

diabeteic foot

糖尿病足是糖尿病和其併發症累積而成的足部傷害,最常見的足部皮膚潰瘍、

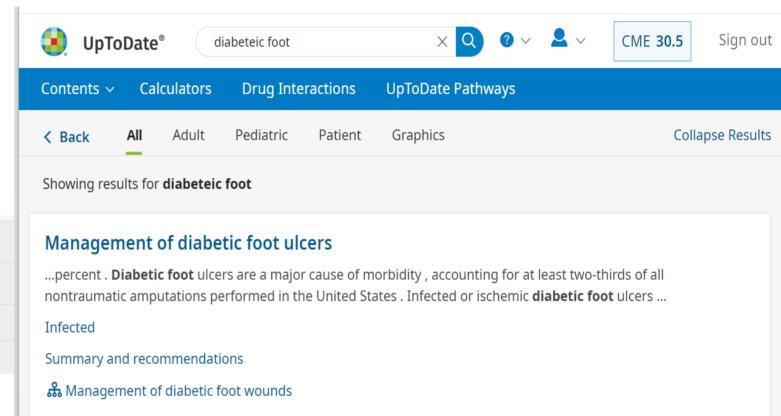
傷口不易癒合。

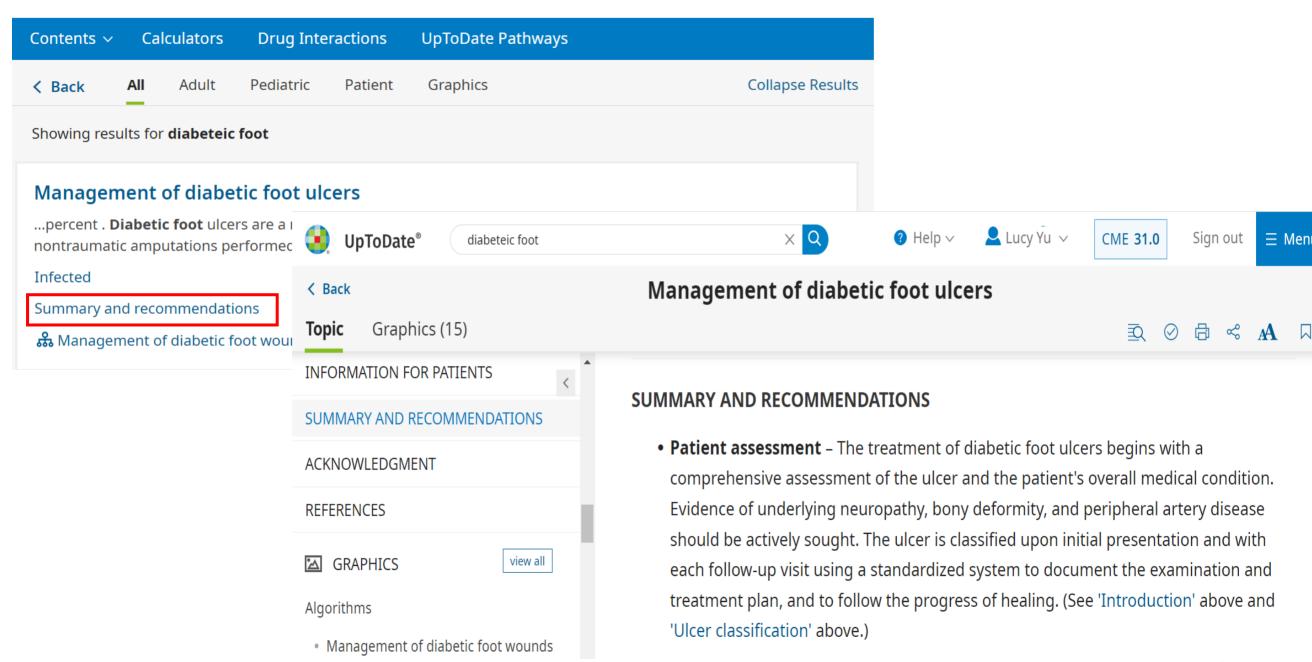




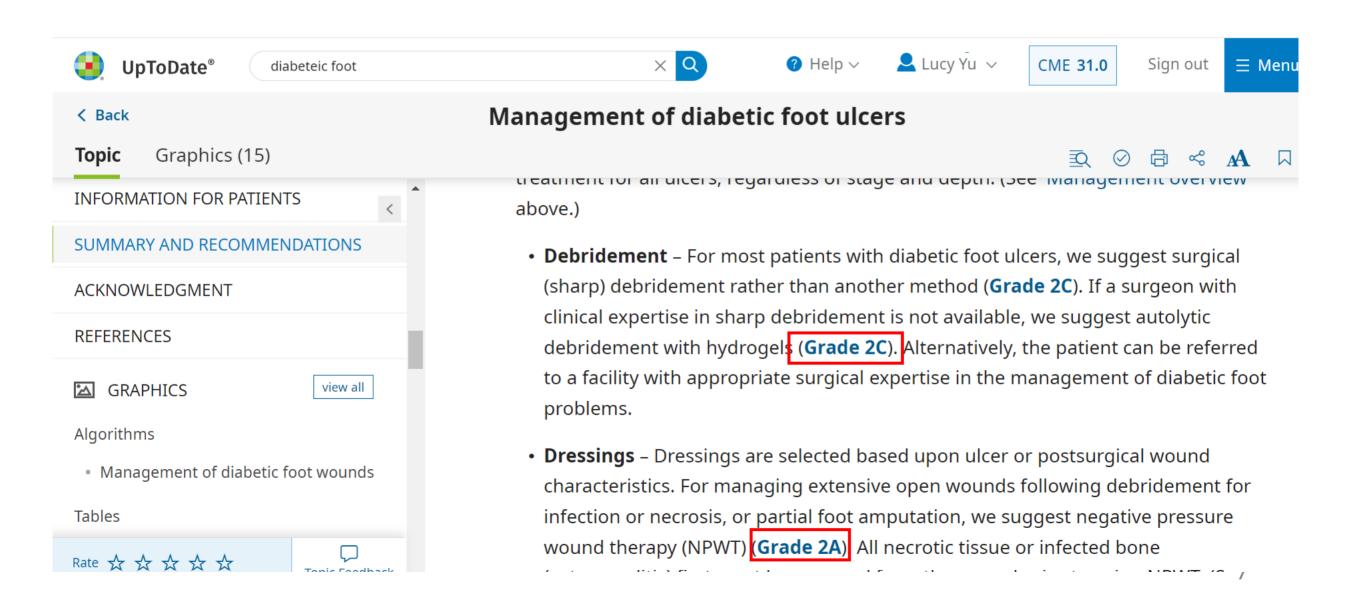
語言切換方式:請將滑鼠移到最下方







一定要在SUMMARY AND RECOMMENDATIONS才能看到證據等級





diabeteic foot







CME **31.0**

Sign (

Contents ∨

Calculators

Drug Interactions

UpToDate Pathways

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Grade 2C recommendation

A Grade 2C recommendation is a very weak recommendation; other alternatives may be equally reasonable.

Explanation:

A Grade 2 recommendation is a weak recommendation. It means "this is our suggestion, but you may want to think about it." It is unlikely that you should follow the suggested approach in all your patients, and you might reasonably choose an alternative approach. For Grade 2 recommendations, benefits and risks may be finely balanced, or the benefits and risks may be uncertain. In deciding whether to follow a Grade 2 recommendation in an individual patient, you may want to think about your patient's values and preferences or about your patient's risk aversion.

Grade C means the evidence comes from observational studies, unsystematic clinical experience, or from randomized, controlled trials with serious flaws. Any estimate of effect is uncertain.

Recommendation grades

- 1. Strong recommendation: Benefits clearly outweigh the risks and burdens (or vice versa) for most, if not all, patients
- 2. Weak recommendation: Benefits and risks closely balanced and/or uncertain

Evidence grades

- A. High-quality evidence: Consistent evidence from randomized trials, or overwhelming evidence of some other form
- B. Moderate-quality evidence: Evidence from randomized trials with important limitations, or very strong evidence of some other form
- C. Low-quality evidence: Evidence from observational studies, unsystematic clinical observations, or from randomized trials with serious flaws

For a complete description of our grading system, please see the UpToDate editorial policy.

推薦等級

- 1.強烈建議:對大多數(如果不是全部)患者而言,益處明顯大於風險和負擔(反之亦然)
- 2.弱推薦:收益和風險密切平衡和/或不確定

證據等級

- 1.高質量證據:來自隨機試驗的一致證據,或其他形式的壓倒性證據
- 2.中等質量的證據:來自具有重要局限性的隨機試驗的證據,或其他形式的非常有力的證據
- 3. 低質量證據:來自觀察性研究、非系統性臨床觀察或存在嚴重缺陷的隨機試驗的證據

藥物資訊與藥物交互作用查詢



UpToDate®

aspirin





目錄 ~ 臨床計算工具 藥物交互作用 **UpToDate Pathways**

く上一頁

所有主題

兒童 成人

病患

圖表

顯示與 aspirin 相關的結果

替代搜尋: Antiplatelet agents, Nonselective nonsteroidal antiinflammatory drugs

Aspirin for the secondary prevention of atherosclerotic cardiovascular disease

...safety between doses of 75 to 150 mg/day (called low-dose aspirin) and 160 to 325 mg/day (called medium-dose aspirin). The optimal daily dose of aspirin for long-term, secondary prevention of CVD events ...

Aspirin sensitivity

Alternatives to aspirin

Summary and recommendations

NSAIDs (including aspirin): Secondary prevention of gastroduodenal toxicity

...a PPI, patients were assigned to continue taking aspirin (80 mg per day) or to stop the aspirin and quitch to a placeho for eight weeks. Stanning agains therapy was associated with a pensionificant

藥物資訊

⊘ Aspirin

病患 兒童

查看完整主題

Aspirin: Drug information

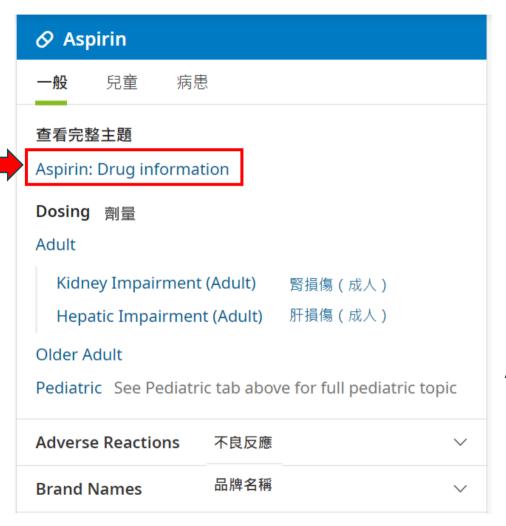
Dosing

Adult

Kidney Impairment (Adult)

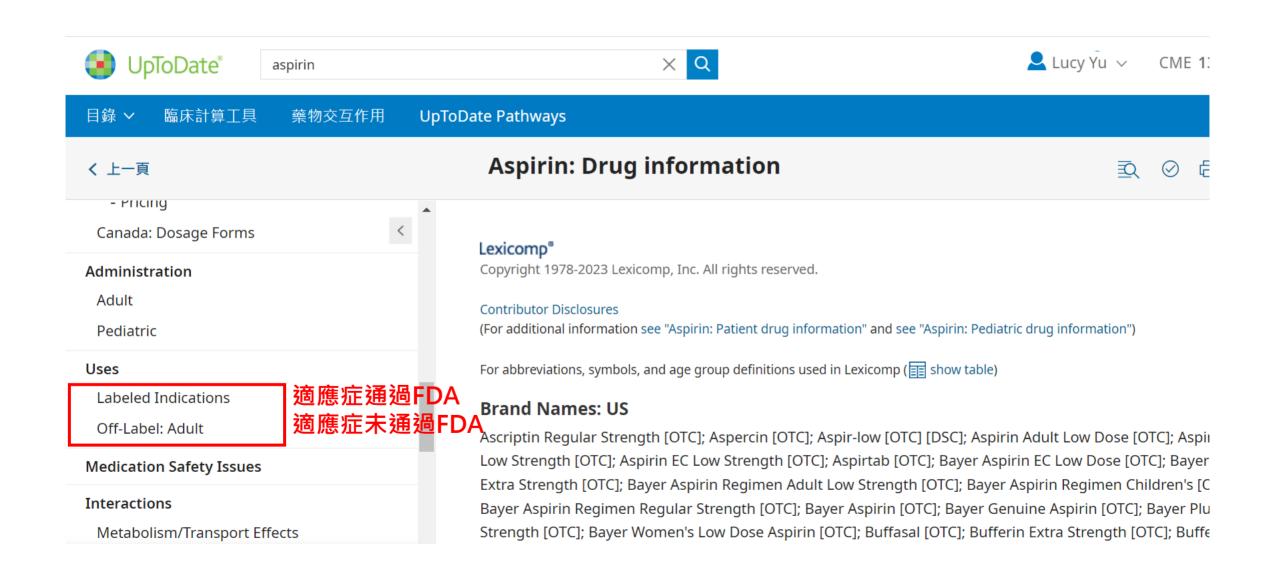
Hepatic Impairment (Adult)

Older Adult





Administration:口服或是其他方式,注意事項



不良反應



交互作用:藥物、食物



Adverse Reactions (Significant): Considerations

不良反應(嚴重):注意事項

GI effects

Blance

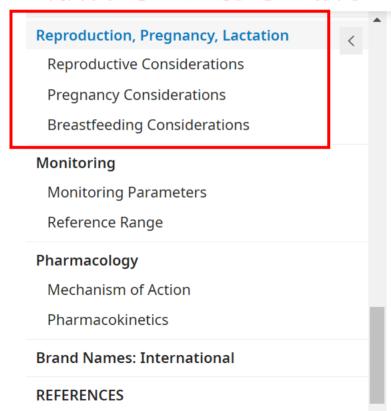
B

-. ug ucu.

(For additional information: Launch drug interactions program) Lexicomp®

Note: Interacting drugs may **not be individually listed below** if they are part of drugs within "CYP3A4 Inducers [Strong]" are NOT listed). For a complete list of on name and detailed management recommendations, use the Lexicomp drug into

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salicylate. Management: Administer with food or large volume of v powder, paprika, licorice.

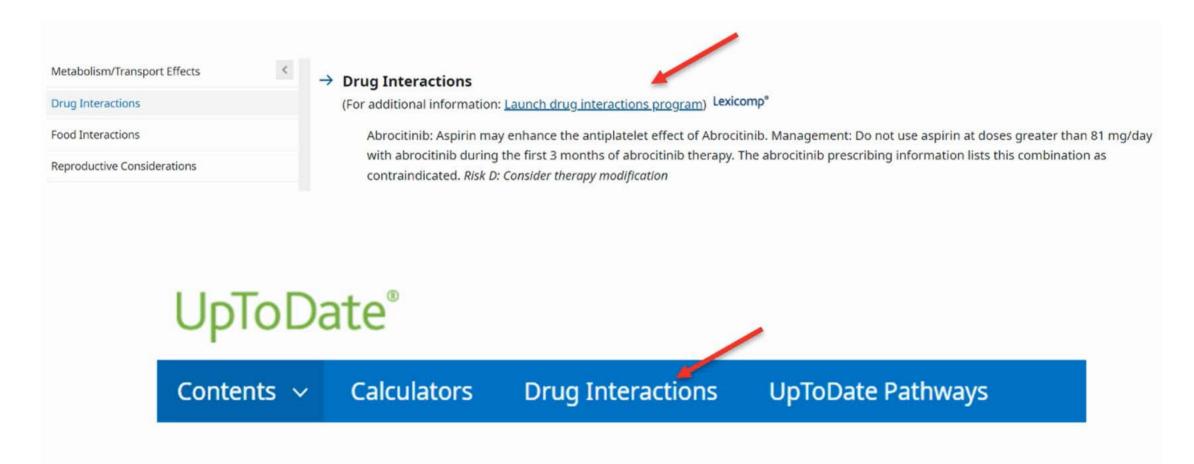
→ Reproductive Considerations

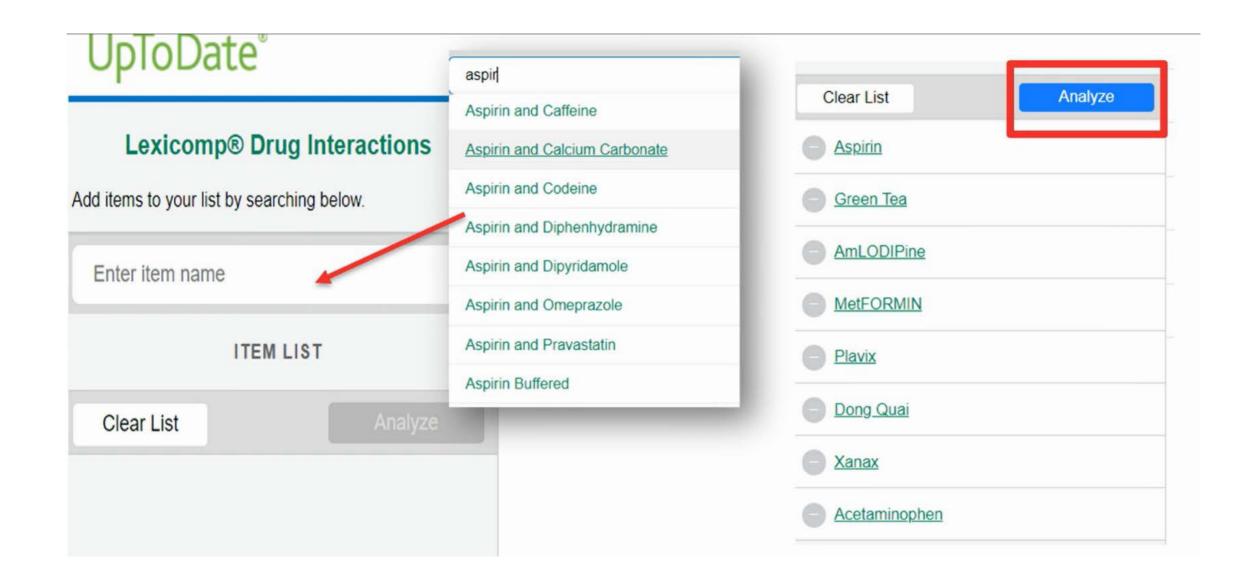
Low-dose aspirin has been evaluated to improve live birth rates in diagnosed with recurrent pregnancy loss (Hamulyák 2020; Hamuly treatment differ between guidelines. Some guidelines initiate aspi other agents (heparin or low-molecular-weight heparin) in patient [Sammaritano 2020]); ESHRE [Bender Atik 2018]; EULAR [Andreoli : in patients with a history of recurrent pregnancy loss but who are (Naimi 2021).

Pregnancy Considerations

Salicylate is present in umbilical cord and newborn serum followin (Garrettson 1975; Levy 1975; Palmisano 1969); salicylic acid and ot newborn urine following in utero exposure (Garrettson 1975).

藥物交互作用進入方式





Title Agents with Blood Glucose Lowering Effects / Salicylates

Dependencies

• Dose: This interaction is likely only of concern when using salicylate doses of 3 grams per day or more.



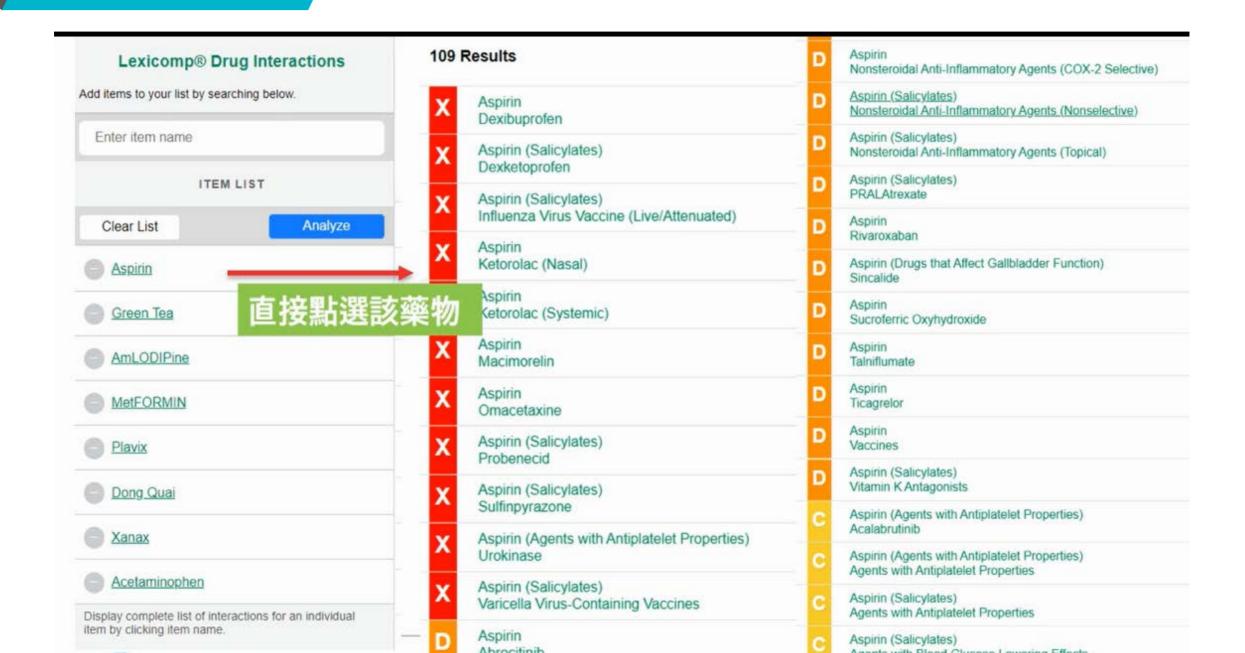
Risk Rating C: Monitor therapy

Summary Salicylates may enhance the hypoglycemic effect of Agents with Blood Glucose Lowering Effects. Severity Moderat

Patient Management Monitor for excessive pharmacological effect (e.g., hypoglycemia) in patients receiving a salicylate concurrence This is likely more of a concern in patients receiving salicylates at a dose of 3 grams or greater per day.

Salicylates Interacting Members Aminosalicylic Acid, Aspirin, Bismuth Subsalicylate, Choline Magnesium Trisalicylate, Cholin Salicylate, Triflusal

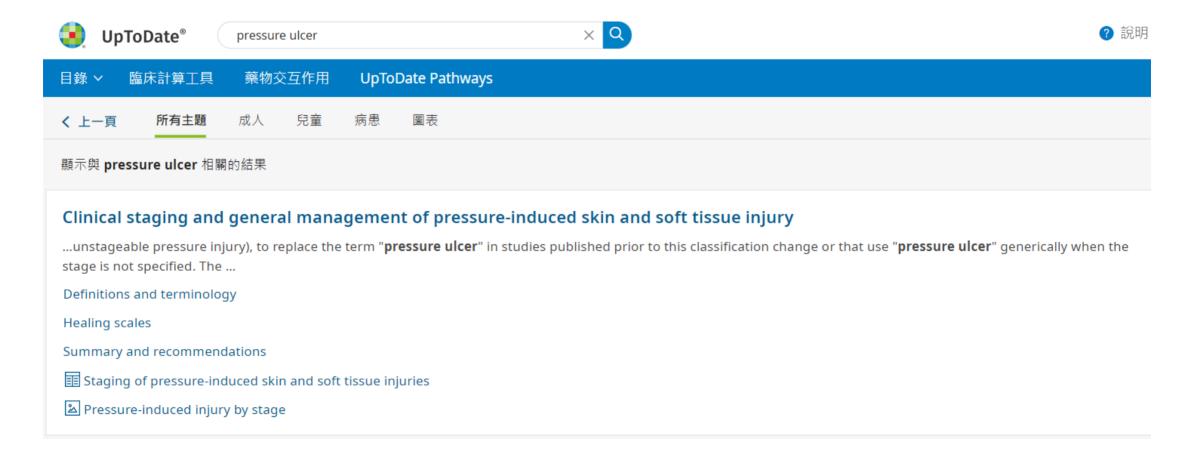
Agents with Blood Glucose Lowering Effects Interacting Members Acarbose, Albiglutide, Alogliptin, Anagliptin, Bromocripti Dapagliflozin, Disopyramide, Dulaglutide, Empagliflozin, Ertugliflozin, Evogliptin, Exenatide, Gemigliptin, Gliclazide, Glimepiride Inhalation), Insulin Aspart, Insulin Degludec, Insulin Detemir, Insulin Glargine, Insulin Glulisine, Insulin Lispro, Insulin NPH, Insulixisenatide, Lobeglitazone, Mecasermin, MetFORMIN, MiFEPRIStone, Miglitol, Mitiglinide, Nateglinide, Octreotide, Pasireotide Pramlintide, QuiNINE, Repaglinide, Rosiglitazone, Saroglitazar [INT], SAXagliptin, Semaglutide, SITagliptin, Somatostatin Acet SulfiSOXAZOLE, SUNItinib, Teneligliptin, TOLAZamide, TOLBUTamide, TraMADol, Vildagliptin, Voglibose





圖表使用

以pressure ulcer 壓瘡、褥瘡為案例





Graphics in this topic



Staging of pressure-induced skin and soft tissue injuries[1]









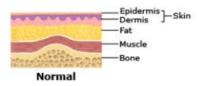


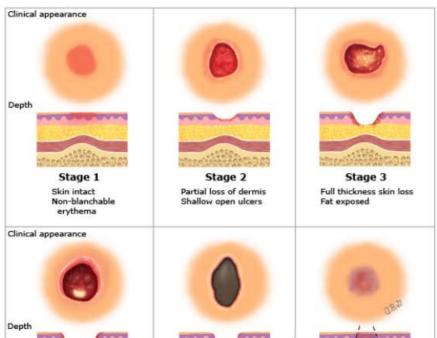


Deep tissue pressure injury

Purplish skin discoloration

Characteristics of pressure-induced injury by stage





Unstageable

pressure injury

Covered with slough

Stage 4

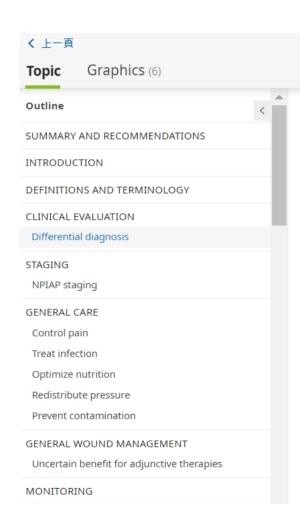
Full thickness skin loss Exposed bone, muscle These illustrations highlight the characteristics of the stages of pressure-induced injuries as defined by the National Pressure Ulcer Advisory Panel. Each panel demonstrates the clinical appearance of the involved skin as well as the cross section of the depth of injury. Most pressure-induced injuries are associated with an underlying bony prominence. There is always the potential for deeper tissue damage than appreciated on clinical examination, as illustrated by the panel "Deep tissue pressure injury."

Adapted from definitions by the National Pressure Ulcer Advisory Panel.

Graphic 50125 Version 7.0

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文章內容與圖片點選切換



Clinical staging and general management of pressure-induced skin and soft tissue injury

assessment of chronic wounds", section on 'Differentiation of chronic ulcers'.)

Pressure-related skin and soft tissue injury may also be confused with other conditions that cause skin erythema, such as cellulitis. (See "Celluliti microbiology, clinical manifestations, and diagnosis", section on 'Cellulitis and erysipelas' and "Cellulitis and skin abscess: Epidemiology, microbi diagnosis".)

STAGING

A number of staging systems have been developed to describe the extent of pressure-induced injury to the skin [6-8]. The most commonly used Pressure Injury Advisory Panel (NPIAP) (table 1 and figure 2) [6]. NPIAP suggests using the terminology "pressure injury" to describe thes degrees of skin damage due to pressure may not be associated with skin ulceration. Our usage of these terms is described above. (See 'Definitic

NPIAP staging — The NPIAP staging system is described below (table 1 and figure 2) [6]. The NPIAP stage is used to describe the initial damage. An unintended consequence of numerical staging is misinterpretation that a 'stage' implies progression or regression (forward or back not imply a step-wise sequence (either direction). In addition, the practice of changing the stage as healing occurs, known as reverse staging, is

- Stage 1 is characterized by intact skin with a localized area of nonblanchable erythema (picture 1), which may appear differently in dark of blanchable erythema or changes in sensation, temperature, or firmness may precede visual changes. Color changes do not include purport may indicate deep tissue pressure injury.
- Stage 2 is characterized by partial-thickness loss of skin with exposed dermis (picture 1). The wound bed is viable, pink or red, moist, an ruptured serum-filled blister. Adipose (fat) is not visible and deeper tissues are not visible. Granulation tissue, slough, and eschar are not presult from adverse microclimate and shear in the skin over the pelvis and shear in the heel. This stage should **not** be used to describe moi (MASD) including incontinence-associated dermatitis (IAD), intertriginous dermatitis (ITD), medical adhesive-related skin injury (MARSI), or burns, abrasions).

Charge 2 is sharesterized by full thickness loss of skip, in which adjaces (fat) is visible in the ulser and granulation tissue and enibale (ralled)



What's New



以專科的方式呈現

What's New

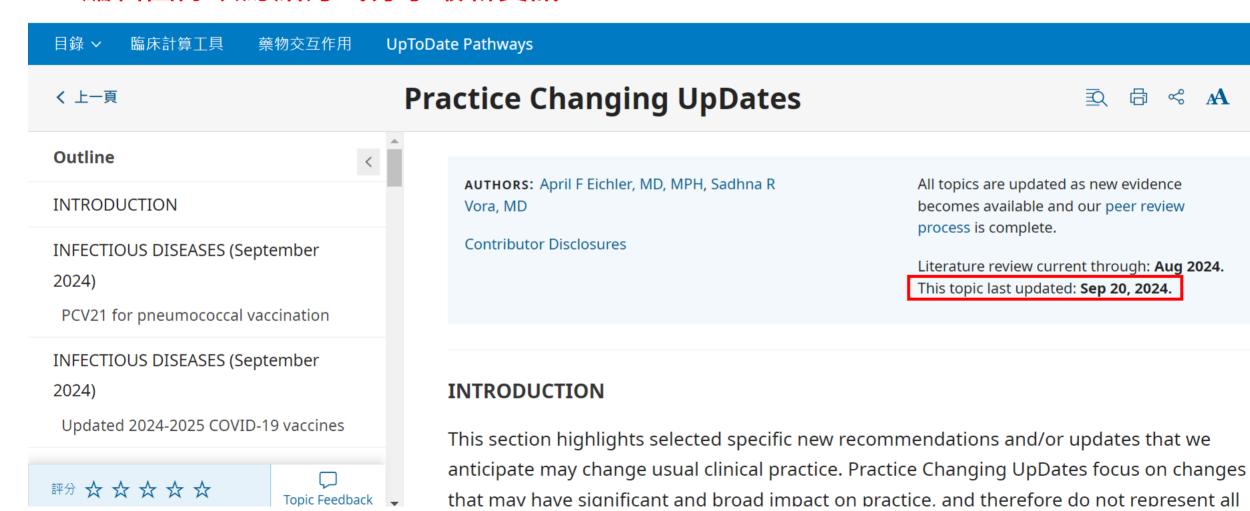
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Drug therapy	Nephrology and hypertension	Rheumatology
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Family medicine	Oncology	Surgery

編輯團隊以總結方式分享最新資訊

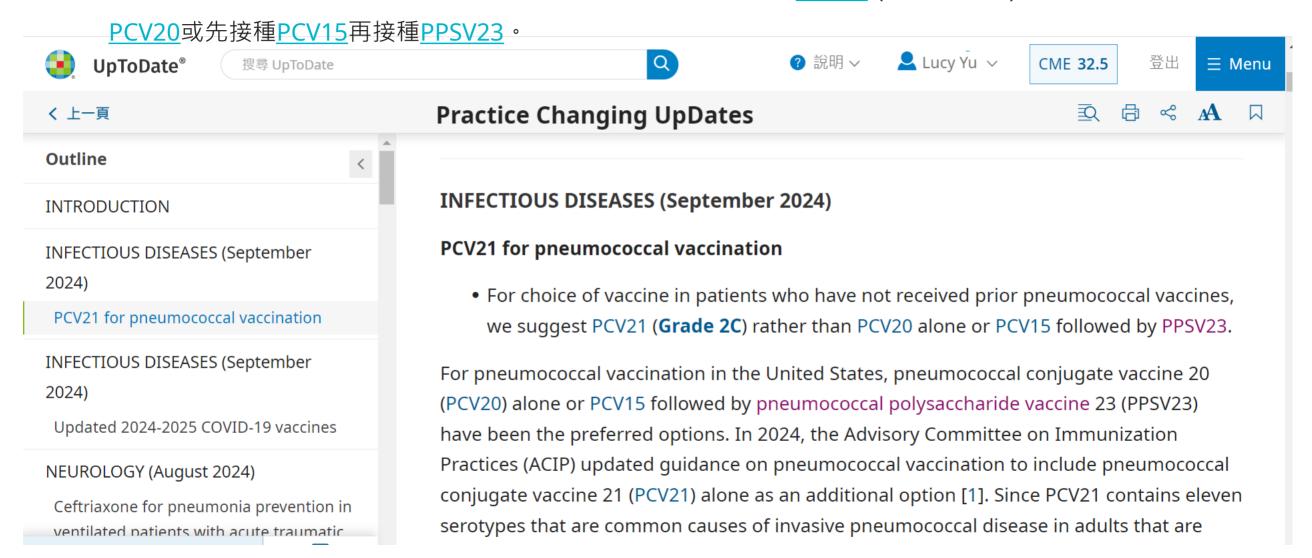


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傳染病(2024年9月)

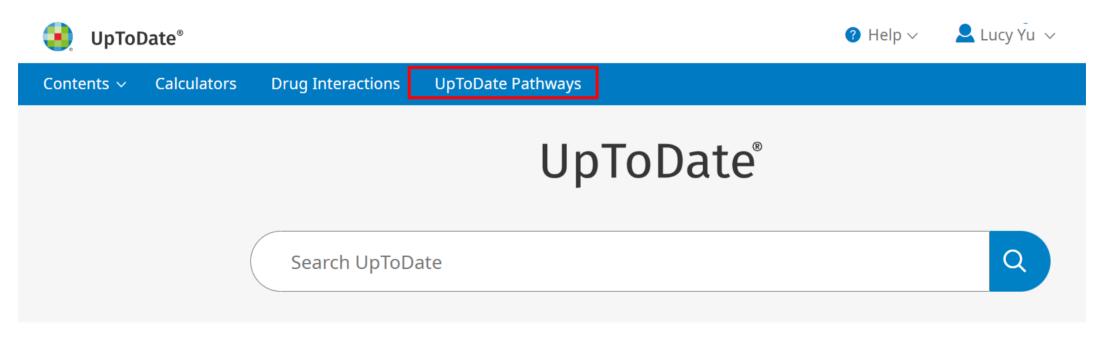
用於肺炎鏈球菌疫苗接種的 PCV21

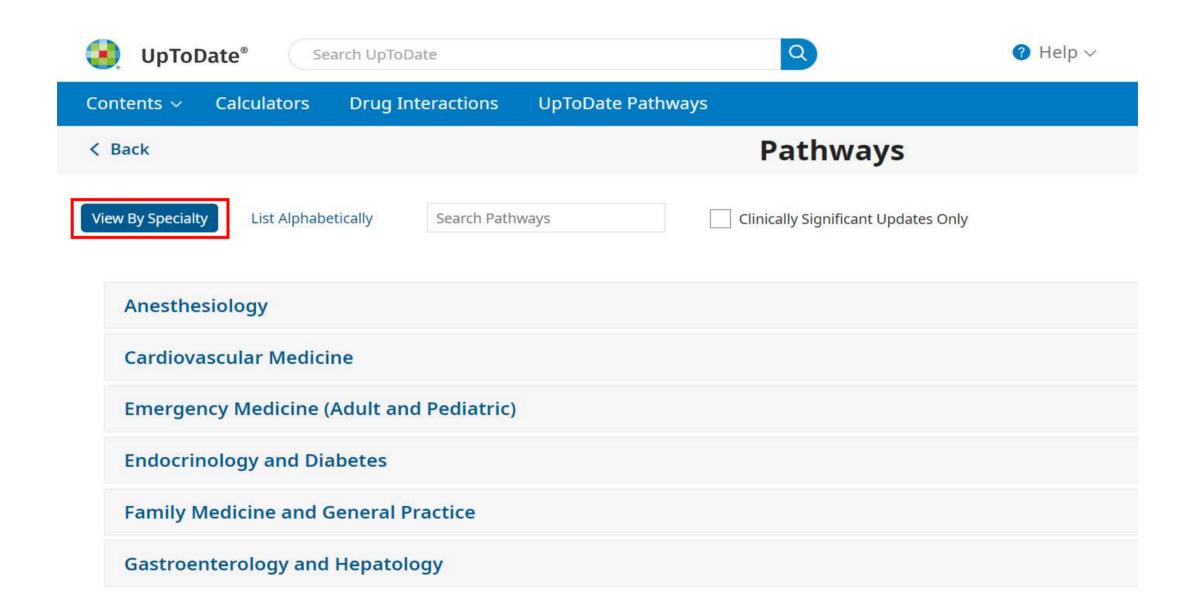
●對於先前未接種過肺炎鏈球菌疫苗的患者的疫苗選擇,我們建議PCV21 (Grade 2C),而不是單獨使用



UpToDate Pathways

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Abnormal liver tests: Initial evaluation

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Acute decompensated heart failure: Discharge checklist

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NEJM AI



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