實證醫學簡介-文獻搜尋及資料庫介紹

實證醫學的五大進行步驟

- 1.Assess the clinical condition Assess 分析病患的情況了解病患的臨床需求
- 2.Formulate an answerable question. Ask 提問: 由個案的臨床資料形成可回答的臨床問題
- 3. Track down the best evidence. Acquire 尋找最佳的實證(各種文獻及資料庫,包括發表及未發表的資料)
- 4. Critically appraise the evidence for validity, impact, and applicability.

 Appraisal

評估各種醫學報告的可信度、臨床重要性,以及可應用性

■ 5.Integrate with our clinical expertise and patient values.

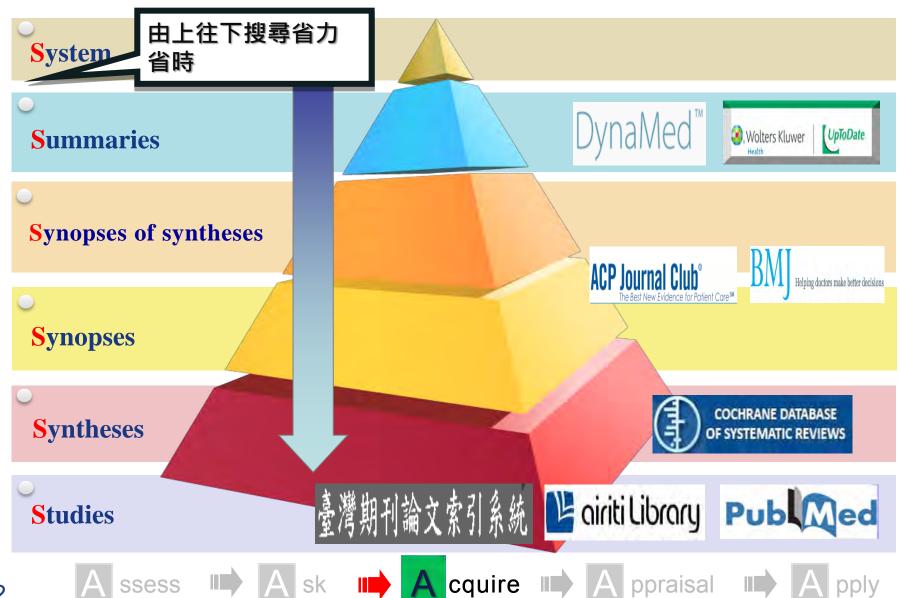
整合並應用於實際患者的治療決策 (臨床應用) Apply



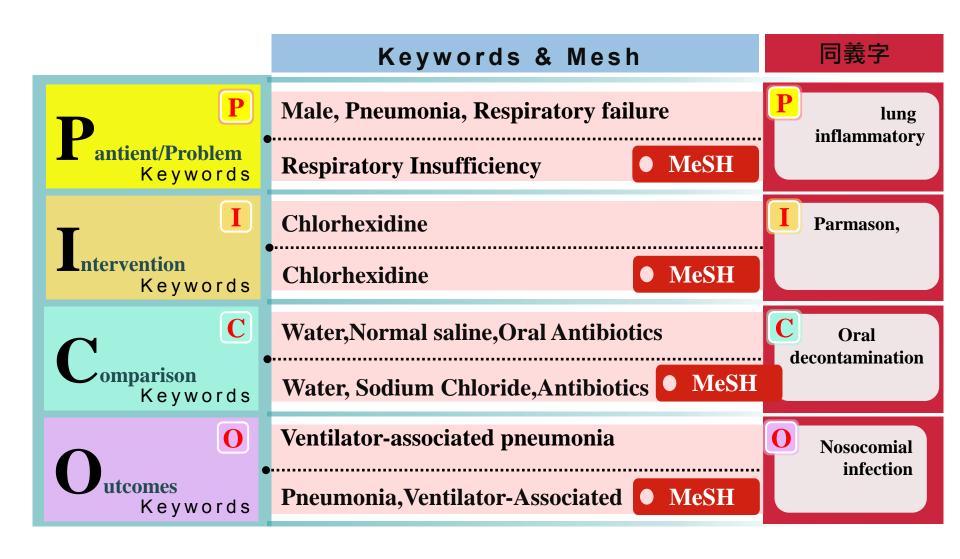
資料庫介紹,文獻搜尋

- 布林邏輯
- Summary型資料庫
- Synopses型資料庫
- Syntheses型資料庫
- · Study型資料庫
- 檢索策略
- 檢索技巧初階

資料庫金字塔65



PICO AND 布林邏輯





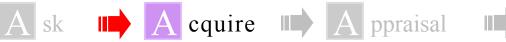
















Summary型資料庫

• 為何用Summaries型資料庫

DynaMed™

搜尋: 簡單、可搜尋全文內容

呈現: 評讀精簡資訊、推薦與證據

應用: 快速支援臨床決策



• 怎麼用Summaries型資料庫 將拿手主題來搜尋所有資料庫一輪 挑出一兩個滿意的資料庫拿來應急 對呈現的證據永遠保持合理懷疑

DynaMed

The Evidence Based Clinical Database

每日更新主題評論實證醫學資料庫



DynaMed 實證醫學主題評論資料庫

- 市面唯一「每日更新」的EBM 主題評論資料庫
- 完整的實證等級資訊與實證文獻參考來源
- 條列式、架構化的主題內容呈現方式
- 可依照足以改變臨床決策的更新文獻進行篩選
- Journal of Clinical Epidemiology
- 1. Timeliness of content updating: 內容的更新 (DynaMed 獲評為 No.1)
- 2. Breadth of coverage: 內容的廣度(DynaMed 獲評為No.3)
- 3. Quality of evidence reporting: 實證證據的品質(DynaMed 獲評為No.2

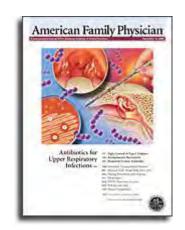
資訊來源

每日審視逾500種以上的頂尖醫學期刊、重要醫學二次文獻實證醫學文獻資源、藥物資訊資源、臨床診療指引









Cochrane Database of Systematic Reviews

Annals of Family Medicine

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Горіс	Update Reason	Date	
Medication and drug exposure in pregnancy	interferon beta use during pregnancy associated with increased risk of preterm birth in women with multiple sclerosis (Neurology 2012 Sep 11)	03/13/2013 05:04:00 PM	
Medication and drug exposure in pregnancy	exposure to NSAIDs during first trimester of pregnancy may not increase risk of major congenital malformations, but COX2 inhibitors may increase risk of musculoskeletal malformations (J Rheumatol 2012 Nov)	03/13/2013 04:57:00 PM	
HER2 inhibitors for breast cancer	trastuzumab does not appear to increase risk of central nervous system relapse (Lancet Oncol 2013 Mar)	03/13/2013 04:27:00 PM	

證據等級和推薦等級顯示方式

提供容易判讀的實證等級 (summary by FACT)

Level of Evidence:

Level 1 (likely reliable) Evidence

Level 2 (mid-level) Evidence

Level 3 (lacking direct) Evidence



提供容易判讀的建議等級(Summary by Guidelines)

Recommendations:

Grade A recommendation (consistent high-quality evidence)

Grade B recommendation (inconsistent or limited evidence)

Grade C recommendation (lacking direct evidence)



http://www.dynamicmedical.com/levels.php

Achilles tendinopathy

Treatment overview:

- rest and ice considered first-line therapy for acute Achilles tendinopathy (grade C recommendation [lacking direct evidence])
- if complete or partial runture non-weight hearing, immediate orthogodic consult, see Achilles tendon rupture
- · conservati 實證等級答訊
 - · res 貝亞寸巛貝미
 - analgesics may reduce symptoms in the state term but do ... have long-term benefit
 - topical NSAID (niflumic acid) might reduct pain and hasten return a jour level of activity (level 2 [mid-level] evidence)
 - oral NSAIDs may be no better an placebo for improving pain and function (level 2 [mid-level] evidence)
 - also consider ice and elevation after ctivity
- calf stretching and strengthening exerci
 - necessary for recovery of function after conservative treatment
 - eccentric calf muscle training pears more effective than concentric calf muscle training (and watchful waiting) for recovery of function tendinosis (level 2 [mid-level] evidence)
- additional treatment considerations
 - topical glyceryl trinitrate plus physical therapy may reduce symptoms compared to physical therapy alone for chronic noninsertional achievidence)
 - steroid injection not recommended due to limited inconsistent evidence (grade B recommendation [inconsistent or limited evidence])
 - shock wave therapy has inconsistent evidence for effect on pain in chronic Achilles tendinopathy
 - insufficient evidence to clearly define optimal treatment for acute or chronic Achilles tendonitis, based on Cochrane review of 9 trials

Activity:

Rest, ice and activity modification:

- rest and ice considered first-line therapy for acute Achilles tendinopathy (grade C recommendation [lacking direct evidence])
 - based on expert opinion
 - Reference J Fam Pract 2008 Apr;57(4):261 EBSCO host Full Text
- rest

Achilles tendinopathy

Treatment overview:

- rest and ice considered first-line therapy for acute Achilles tendinonathy (grade C recommendation [lacking direct evidence])
- if complete or partial rupture non-weight bearing, immediate

conconcitive treatment for 4.6 weeks if acute symptom adema

建議等級資訊

t do not have long-term benefit

- topical NSAID ifflumic as imight reduct pain and hasten return to previous level of activity (level 2 [mid-level] evidence)
- oral NSAIDs may no better an placebo for improving pain and function (level 2 [mid-level] evidence)

hes

- also consider ice and elevath after activity
- calf stretching and strengthening exerces
 - necessary for recovery of function fer conservation treatment
 - eccentric calf muscle training appears three effective the concentric calf muscle training (and watchful waiting) for recovery of function tendinosis (level 2 [mid-level] evidence)
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Top General Information (including ICD-9/-10 Codes) Causes and Risk Factors Complications and Associated Conditions History Physical Diagnosis Prognosis □ Treatment Treatment overview Activity Medications Surgery Consultation and referral Other management

Send Comment to Editor

Achilles tendinopathy

- 54 patients aged 18-70 years with chronic tendinopathy 2-7 cm above Achilles tendon insertion randomized to (saline injection plus usual care)
- pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity level score at 24 weeks improved significantly in both groups (not significantly different between the pain and activity different between the pain activity different between the pain and activity different between the pain activity different betwe
- Reference JAMA 2010 Jan 13;303(2):144, commentary can be found in JAMA 2010 May 5;303(17):1696

Surgery:

- last resort for patients with chronic Achilles tendon pain unrelieved by nonoperative treatment
- surgeries include
 - · resection of prominent tuberosity
 - debridement of bursa
 - · excision of thickened, scarred paratenon
 - · removal of accessible calcific deposits within tendon
- surgery reported to be beneficial (level 3 [lacking direct] evidence)
 - based on uncontrolled case series
 - retrospective study of 35 patients with painful Achilles tendon syndrome unrelieved by nonoperative treatmen years
 - mean pain scores improved from 4.7 preoperatively to 1.5 postoperatively, 90% patients had significant improved participation
 - 14 patients had altered sensation at surgical scar
 - Reference Am J Sports Med 2002 May-Jun;30(3):318 in J Musculoskel Med 2002 Dec;19(12):516

Consultation and referral:

- physical therapy in selected cases for muscle rehabilitation
 - · ultrasound modality, commonly used prior to manually-assisted calf and Achilles stretching
 - · transverse friction massage may convert tendinosis to tendonitis and initiate acute-phase immune response
- · if complete or partial rupture non-weight bearing, immediate orthopedic consult

Other management:

insufficient evidence to clearly define optimal treatment for acute or chronic Achilles tendonitis



alopecia

Browse: A B C D E F G H I J K L M N O P Q R S T U V W X Y Z





Browse Categories

利用關鍵字查找

Alopecia areata	3	Alopecia areata
Androgenetic alopecia - work in progress		⊞ General Information (including ICD-9/-10 Codes)
Minoxidil (Topical)		Causes and Risk Factors
Finasteride		Complications and Associated Conditions
Toxicities of chemotherapeutic agents		History
Action Commission Comm		Physical
Minoxidil (Systemic)		
Discoid lupus erythematosus		⊞ Prognosis
Tinea capitis		☐ Treatment
Temozolomide		Treatment overview
Chemotherapy for advanced or recurrent non-small cell lung cancer (NSCLC)		Medications
		Consultation and referral
Trichotillomania		Other management
Irinotecan		Prevention and Screening
Pentosan		① References including Reviews and Guidelines
Gastric lymphoma		Patient Information
Polycystic ovary syndrome		Acknowledgements
Chronic mucocutaneous candidiasis		
Acne		
Hepatitis B		

Alopecia areata

Treatment overview:

- usually no treatment indicated or necessary
- efalizumab (Raptiva) 1-2 mg/kg subcutaneously weekly (withdrawn from market in United States and European Union)

Medications:

- inconclusive evidence on topical and systemic interventions for alopecia
 - based on Cochrane review of trials with methodologic limitations
 - systematic review of 17 randomized trials evaluating topical and systemic interventions for alopecia areata, alopecia

治療方式說明

- most trials were small, only 1 trial had adequate allocation concealment, 2 trials reported use of intention-to-treat-a
- interventions included topical and oral corticosteroids, topical cyclosporine, photodynamic therapy and topical minox
- no intervention showed significant treatment benefit in terms of hair growth compared to placebo
- no trial evaluated self-assessed hair growth or quality of life
- no randomized trials identified that evaluated diphencyprone, dinitrochlorobenzene, intralesional corticosteroids, or of
- Reference Cochrane Database Syst Rev 2008 Apr 16;(2):CD004413 EBSCO host Full Text
- intralesional steroids
- · avoid systemic steroids
- PUVA and anthralin (irritation) have also been used
- minoxidil not generally recommended
- topical diphenylcyclopropenone reported to have about 45% to 79% response rates in case series (level 3 [lacking direction])
 - topical diphenylcyclopropenone reported to have 45% response rate at 6 months (25 patients had complete regro patients with chronic extensive alopecia areata (J Am Acad Dermatol 2001 Jan;44(1):73)
 - topical diphenylcyclopropenone reported to have 79% response rate (6 complete and 16 partial responders) in operate areata, 13 of 22 responders (51%) had partial recurrence over 6-12 months; side effects included eczematous reaction, and hyperpigmentation (BMC Dermatology 2005 May 26;5:6)
 - diphenylcyclopropenone (diphencyprone) associated with 78% cumulative response rate at 32 months in series of 3 JAMA 2001 Nov 21;286(19):2384)
- efalizumab may not be effective for alopecia areata (level 2 [mid-level] evidence)
 - based on small randomized trial

alopecia

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- Allergic Disorders
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 - ☐ Cardiovascular medications
 - ★ ACE inhibitors
 - Angiotensin II receptor blockers
 - Antianginal agents
 - **±** Antiarrhythmic agents

 - Antihypertensive agents

 - Antiplatelet agents
 - ☐ Beta blockers
 - Acebutolol Rx
 - Atenolol Rx
 - Beta blockers during acute ST-elevation myocardial infarction (STEMI)
 - Beta blockers for heart failure
 - Betaxolol (Systemic) Rx
 - Bisoprolol Rx
 - Carvedilol Rx
 - Esmolol Rx
 - Laborated Dv

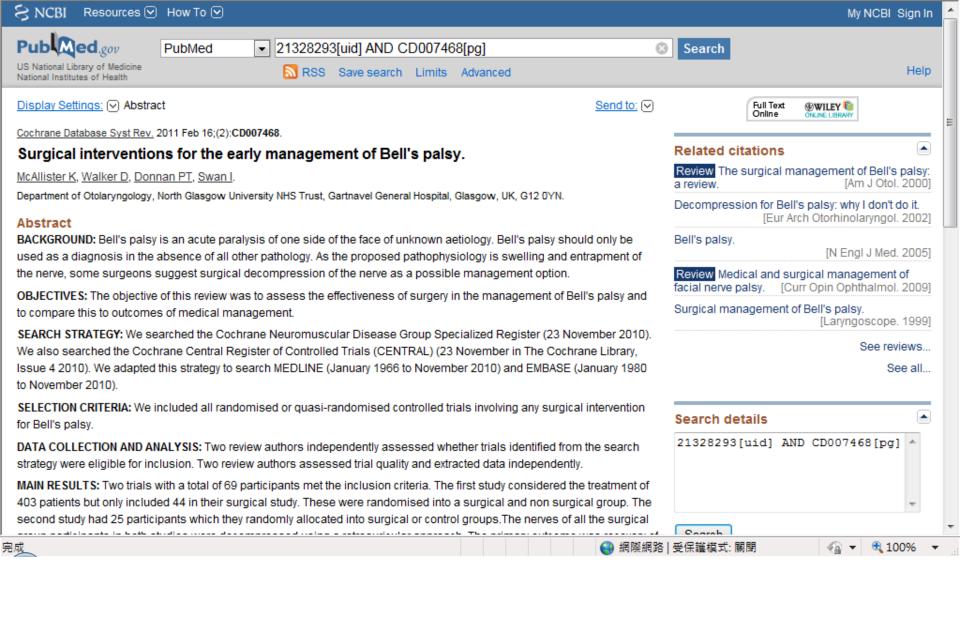


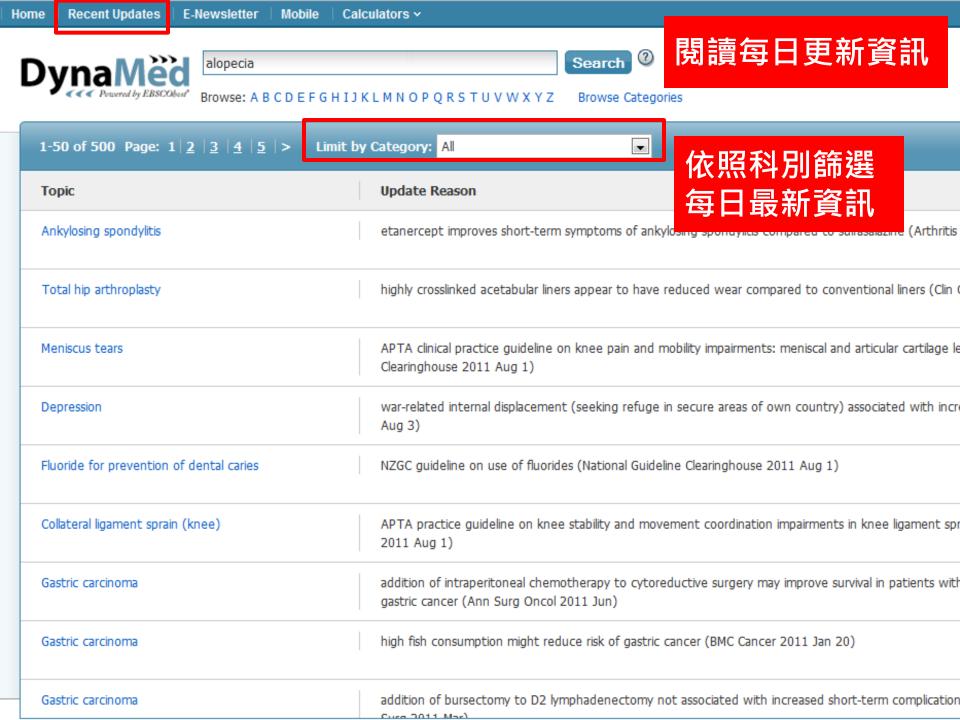
Bells palsy

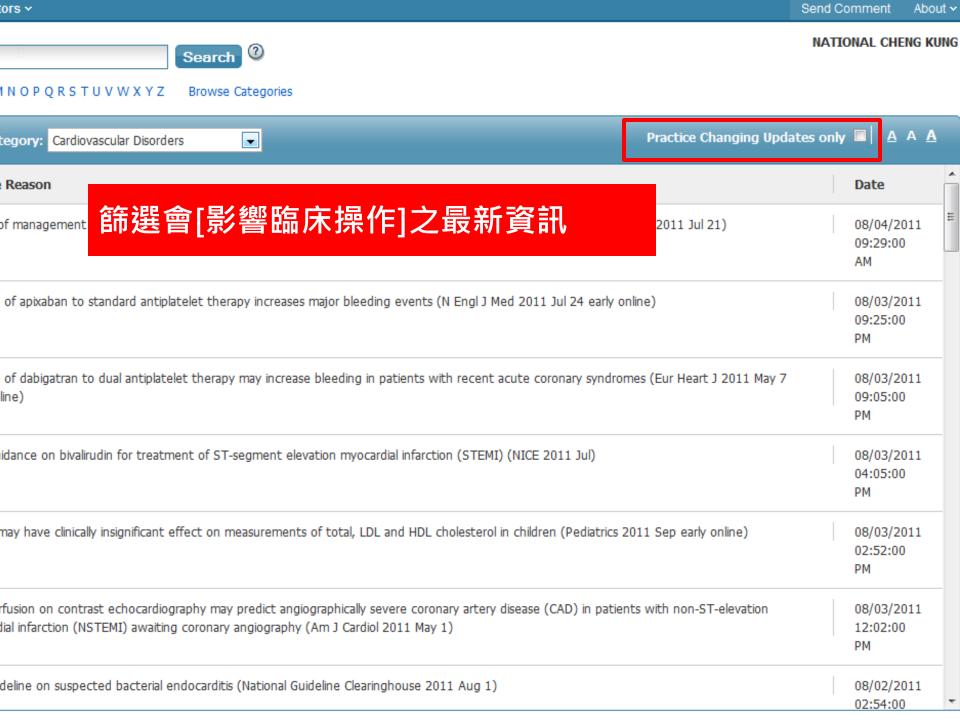
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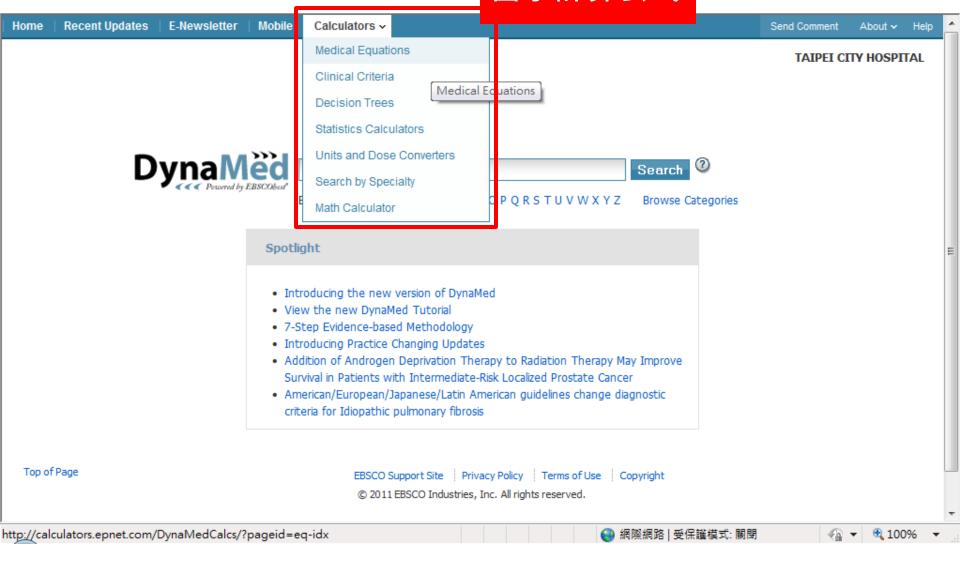








醫學計算公式





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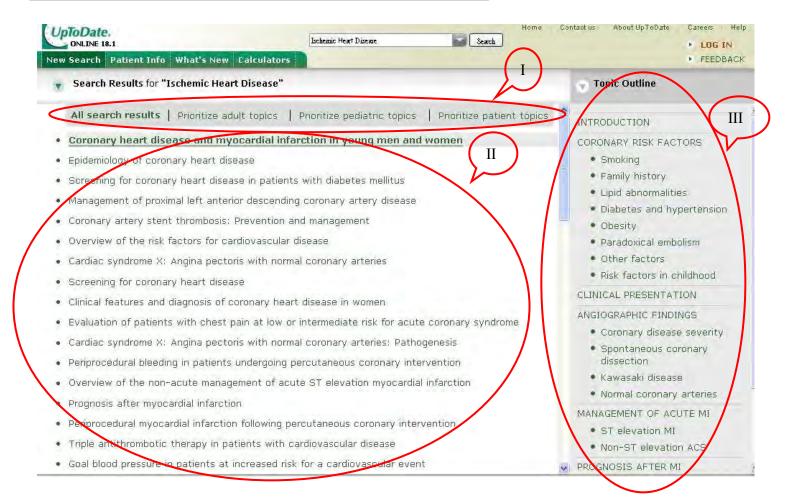
- UpToDate 提供即時實證醫學及臨床醫療資訊,
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- 文獻中附有圖片,包括圖表、X光片、相片、影像檔等,及MEDLINE的引用文獻摘要。

Evidence Grading

RECOMMENDATION GRADES					
Grade 1	Strong Recommendation	Benefits clearly outweigh the risks and burdens (or vice versa) for most, if not all, patients			
	"We recommend"				
Grade 2	Weaker Recommendation	Benefits and risks closely balanced and/or			
	"We suggest"	uncertain			
EVIDENCE GRADES					
Grade A	High Quality Evidence	Consistent evidence from randomized trials, or overwhelming evidence of some other form			
Grade B	Moderate Quality Evidence	Evidence from randomized trials with important limitations, or very strong evidence of some other form			
Grade C	Low Quality Evidence	Evidence from observational studies, unsystematic clinical observations, or from randomized trials with serious flaws			

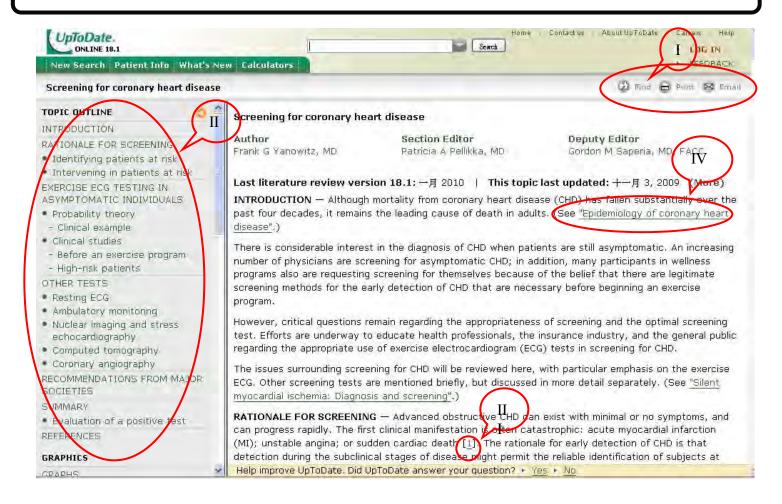
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I.不同的篩選策略可來進行文章排列 II.所檢索出來的文章結果 III.文章所顯示出的文章內容標準



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- 1. 提供搜尋、列印及email功能
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Uptodate

New Search:



Uptodate

- Side effects Combination therapy - Short-term efficacy - Long-term efficacy Antimuscarinics Herbal therapies Other MODIFICATION OF THERAPY OTHER STRATEGIES ECONOMIC CONSIDERATIONS SUMMARY AND RECOMMENDATIONS REFERENCES GRAPHICS FIGURES · Prevalence of BPH with age · Einsctorida in BALL

Herbal therapies — Herbal therapies for BPH are commonly used in Europe; these remedies comprised 70 percent of spending for drug treatment of prostatism in Germany in 1997 [48]. Saw palmetto is approved by the German Commission E for stage I and II (mild to moderate) BPH. Two herbal extracts have officially been approved for the treatment of prostatism in France. No herbal therapies have been approved by the United States Food and Drug Administration for this purpose, although many men probably try these treatments. There is a substantial placebo effect associated with herbal therapy, as there is for most drugs used to treat BPH.

The data concerning efficacy of these therapies are conflicting. In systematic reviews of controlled trials, saw palmetto plant extract was as effective as finasteride in relieving the symptoms of prostate obstruction, although it did not decrease prostate volume [49]. A subsequent placebocontrolled trial found no evidence that saw palmetto was superior to Help improve UpToDate. Did this topic answer your question? • Yes • No

Synopses

- 針對單一文獻進行嚴格評讀後寫成一至兩 頁的評論,評論內容從可信度,重要性, 臨床推薦度皆有提及。
- 可提供臨床人員對於有興趣文獻之專家評讀看法。
- 針對已經有共識的文章加以評讀,通常更新慢,文章數量少。





ACP J Club: 濃縮10頁原始研究為1頁EBM評論

Review: On-demand maintenance therapy with proton pump inhibitors is as effective as continuous therapy for nonerosive GERD

ACP Journal Club. 2007 Nov-Dec;147:69.

Pace F, Tonini M, Pallotta S, Molteni P, Porro GB. Systematic review: maintenance treatment of gastro-oesophageal reflux disease with proton pump inhibitors taken 'on-demand'. Aliment Pharmacol

Ther. 2007; 26:195-204. [PubMed ID: 17593065] 原始研究

Question 臨床問題 The state of th

In patients with gastroesophageal reflux disease (GERD), is on-demand maintenance therapy with proton pump inhibitors (PPIs) effective for controlling symptoms and preventing relapse?

Methods 研究方法

treatment groups.

Data sources: MEDLINE and Cochrane Controlled Trials Register (to December 2008), and reference lists.

Study selection and assessment; Studies published in English as **fu** articles that compared on-demand maintenance therapy, using PPIs **o** receptor antagonists, with placebo or continuous maintenance therapy patients with GERD, with or without erosive esophagitis. 17 articles reporting on 18 studies (n = 14 142) met the selection criteria. All studies used PPIs (esomeprazole, 20 or 40 mg, lansoprazole, 15 or 30 mg;

omeprazole, 10 or 20 mg; pantoprazole, 20 or 40 mg; or rabeprazole, 1L 20 mg); 1 study also included an H_2 -receptor antagonist (ranitidine, 300 mg). In the 14 randomized controlled trials (RCTs), symptomatic patients were first given a short course (4 to 8 wk) of continuous treatment, and

then responding patients were randomized to the different maintenance

Main results 結果

Because of differences in outcome measures among studies, metaanalysis was not done. On-demand PPI therapy was more effective than placebo for all levels of GERD severity (Table). It was more effective than continuous PPI therapy in patients with nonerosive GERD but not in patients with erosive or uninvestigated GERD (Table). 1 RCT (n = 6017) showed that continuous PPI therapy provided better quality of life than on-

Ciniclusions 結論

In patients with gastroesophageal reflux disease (GERD), on-demand maintenance therapy with proton pump inhibitors (PPIs) is more effective than placebo for controlling symptoms. On-demand PPI therapy is as effective as continuous PPI therapy in patients with nonerosive GERD but not in those with more severe disease.

Commentary 評論

patients, with continuous PPI use being the most common prescription pattern (1). Because of cost and safety issues, several alternatives to this approach have been suggested, including intermittent therapy (sustained periods of continuous therapy followed by discontinuation until symptoms recur) and on-demand therapy (patient-driven use based on day-to-day symptoms). Many experts recommend the use of on-demand therapy for nonerosive GERD (2), a view that is supported by the review by Pace and colleagues. The designs, outcomes, and even patient populations (as evidenced by varying placebo response rates) of the included studies are heterodeneous: thus, the authors have appropriately refrained from doing a

GERD is a chronic condition necessitating long-term therapy in most

Galveston, Texas, USA References 參考書日

Rajasekhara R. Mummadi, MD

University of Texas Medical Branch

Pankaj J. Pasricha, MD

1. Nocon M, Labenz J, Jaspersen D, et al. Long-term treatment of patients with gastro-oesophageal reflux disease in routine care-results from the ProGERD study. Aliment Pharmacol Ther. 2007;25:715-22. [PubMed ID: 17311605]