Evidence based medicine

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臨床情境

影響小莊健康決策的因素



CYCH ER EBM

你是專業的醫療人員也是實證醫學專家,請分析PICO架構

圖然可以!PICO 架構是實證醫學中用來形成臨床問題並指導文獻檢索的一種有用工具。以下是 PICO的基本組成及分析方法:

請以你的介紹的架構,幫我分析下面的臨床情境,並以表格方式呈現 PICO

關於P,要包含基本資訊,相關共病,患者價值,社經地位;關於I,要包含藥物選擇,介入劑量,途徑頻率,治療時間;關於C,要包含對照選擇,安慰劑組,標準治療,輔助治療;關於O,要包含重要指標及替代指標或者類別變項及連續變項。如果在臨床情境裡面沒有,請提醒缺乏哪些元素。

###

小莊是一位30出頭的年輕醫療工作者,由於工作的性質,常常需要長時間在電腦前坐著。忙碌的醫院生活讓他幾乎沒有時間運動,下班後總是覺得身心疲憊,只想吃點鹹酥雞來犒賞自己,然後早早洗洗睡。然而,隨著年齡增長,小莊漸漸感受到體力的下降和代謝的變慢。最近的公司健檢報告顯示出讓他頗為擔心的結果:

⑤ 以下是根據您提供的臨床情境,按照 PICO 架構進行的分析,並以表格形式呈現:

PICO 元素	内容分析
P (Population)	基本資訊:30出頭男性,醫療工作者,久坐生活型態。 相關共病:肥胖、脂肪肝、高三酸甘油酯、低 IDL。 患者價值:希望有效減重且不影響生活工作,偏好可執行性高的治療方式。 社經地位:醫療工作者,壓力大,生活忙碌,時間有限。
I (Intervention)	藥物選擇:GLP-1類(liraglutide、semaglutide)、SGLT2 抑制劑。 介入劑量:需依臨床建議確定,建議參考標準劑量或逐步調整劑量。 途徑頻率:GLP-1類通常皮下注射(每日或每週);SGLT2 抑制劑為口服(每日)。 治療時間:中長期(通常超過 12 週以觀察減重和代謝改善效果)。
C (Comparison)	對照選擇:生活方式改變(飲食控制、運動)。 安慰劑組:有些研究可能包含安慰劑作為對照。 標準治療:僅透過飲食和運動改善健康指標的非藥物方案。 輔助治療:中醫療法(如針灸或中藥)。
O (Outcome)	重要指標: - 類別變項:是否達到臨床顯著的體重減少(如≥5%體重減輕)。 - 連續變項:體重變化(公斤)、BMI變化。 替代指標:三酸甘油酯降低、IDL提升、肝臟脂肪含量改善(影像學或血液指標)。 缺失指標:患者感受(如生活品質變化)、藥物依從性、長期效果(需特別關注)。

臨床情境

主要問題

各種減重方式的成效

病人期望

小莊希望達到減重效果

醫療考量

減重療效與安全性、有無副作用

背景知識

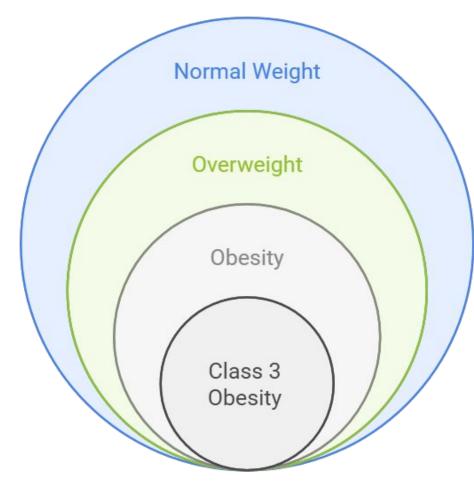
Hierarchy of Weight Classifications

Baseline health risk

Elevated health risks

Increased health risks

Severe health risks and complications



Overweight:

BMI 25~29.9 kg/m2

Obesity:

BMI \geq 30 kg/m2

Class 3 Obesity:

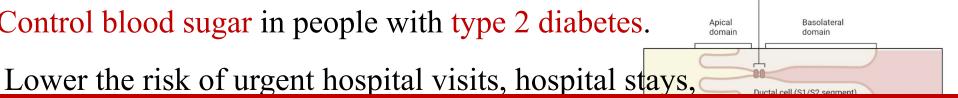
BMI \ge 40 kg/m2 or \ge 35 kg/m2 in the presence of comorbidities)

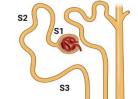


背景知識-SGLT-2 Inhibitor

What is this drug used for?

- Control blood sugar in people with type 2 diabetes.

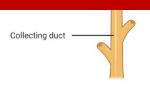




No FDA Approval for body weight loss

stays, and death, in people with certain kidney problems.











Ask Acquire Appraisal Apply Audit

根據臨床問題形成第一個 PICO

Adult male with obesity, fatty liver, hyperlipidemia, without DM

治療型問題

SGLT-2 inhibitor with standard care

□ 診斷型問題

Placebo, standard care, diet, exercise

□ 傷害型問題

預後型問題

- 0
- Body weight loss, BMI reduction, □ 篩檢型問題 side effect, waist circumference, hip circumference

根據臨床問題形成第二個 PICO

Adult male with obesity, fatty liver, hyperlipidemia, without DM

- ▶ 治療型問題
- Intermittent fasting with standard care
- 〕診斷型問題

Normal dietary

」傷害型問題

預後型問題





Ask Acquire Appraisal Apply Audit

CYCH ER EBM 檢索策略

	Keywords / Free text / Synonyms	[MeSH terms]	中文
P	(overweight OR obes*) AND (non-diabe* OR no-diabe* OR "without diabe*") AND (adult) ("fatty liver" OR "liver steatosis" OR "hepatic steatosis" OR "hepatic fat infiltration" OR "fatty liver disease") ("dyslipidemia" OR "hyperlipidemia" OR "lipid disorders")	"Obesity"[Mesh]" "Fatty Liver"[Mesh] AND "Non-alcoholic Fatty Liver Disease"[Mesh] Hyperlipidemias Dyslipidemias	肥胖、過重、超重 非糖尿病、脂肪肝 血脂異常
	sodium glucose cotransporter 2 inhibitors OR canagliflozin OR dapagliflozin OR empagliflozin OR ipragliflozin OR tofogliflozin OR luseogliflozin OR sergliflozin OR remogliflozin OR ertugliflozin OR sotagliflozin	"Sodium-Glucose Transporter 2 Inhibitors"[Mesh] AND "Sodium-Glucose Transporter 2"[Mesh]	鈉-葡萄糖共同轉運器-2 抑制劑
C	Control groups, Placebo Standard therapy Diet control, physical activity	"Control groups"[Mesh] "Placebo"[Mesh] "Diet"[Mesh]	對照組、安慰劑 標準治療、飲食控制 運動
0	weight loss, weight reduction, body weight decrease, weight management, BMI reduction side effect, waist circumference, hip circumference	"Body Mass Index"[Mesh] Body Weight"[Mesh] "Weights and Measures"[Mesh "Weight Reduction Programs"[Mesh]	減重、BMI 腰圍、臀圍 副作用
		CYCH ER EBM	I

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PICO	英文同義詞	控制詞彙 MeSH	控制詞彙 Emtree
P (Population)	obesity, obese, overweight, adiposity, high BMI, excessive weight, body weight disorders, weight gain	Obesity, Overweight	Obesity, Overweight
	non-diabetic, non diabetes, no diabetes, without diabetes	Non-Diabetic	Non-Diabetic, Non Diabetes
	adult, adults, young adult*, middle- aged, middle-aged person*, people	Adult	Adult
(Intervention)	SGLT2 inhibitor, SGLT-2 inhibitor, SGLT 2 inhibitors, sodium-glucose cotransporter 2 inhibitor, sodium glucose co-transporter 2 inhibitor	Sodium-Glucose Transporter 2 Inhibitors	Sodium-Glucose Cotransporter 2 Inhibitor
	dapagliflozin, canagliflozin, empagliflozin, ertugliflozin, remogliflozin, sergliflozin	Dapagliflozin, Canagliflozin, Empagliflozin, Ertugliflozin	Dapagliflozin, Canagliflozin, Empagliflozin
C (Comparison)	placebo, standard care, usual care, dietary intervention, diet modification, physical activity, exercise	Placebo, Diet, Exercise	Placebo, Diet, Exercise
O (Outcome)	weight loss, weight reduction, body weight decrease, weight wanagement, BMI reduction, obesity	Weight Loss, Body Weight, Body Mass Index	Weight Loss, Body Weight, Body Mass Index

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檢索策略

首選Level1: SR of RCT

Oxford Centre for Evidence-Based Medicine 2011 Levels of Evidence

Question	Step 1 (Level 1*)	Step 2 (Level 2*)	Step 3 (Level 3*)	Step 4 (Level 4*)	Step 5 (Level 5)
How common is the problem?		Systematic review of surveys that allow matching to local circumstances**	Local non-random sample**	Case-series**	n/a
Is this diagnostic or monitoring test accurate? (Diagnosis)	of cross sectional studies with consistently applied reference	Individual cross sectional studies with consistently applied reference standard and blinding	Non-consecutive studies, or studies without consistently applied reference standards**	Case-control studies, or "poor or non-independent reference standard**	Mechanism-based reasoning
	Systematic review of inception cohort studies	Inception cohort studies	Cohort study or control arm of randomized trial*	Case-series or case- control studies, or poor quality prognostic cohort study**	n/a
	of randomized trials or n-of-1 trials		Non-randomized controlled cohort/follow-up study**	Case-series, case-control studies, or historically controlled studies**	Mechanism-based reasoning
COMMON harms?		or (exceptionally) observational study with dramatic effect	Non-randomized controlled cohort/follow-up study (post-marketing surveillance) provided there are sufficient numbers to rule out a common harm. (For long-term harms the duration of follow-up must be sufficient.)**	Case-series, case-control, or historically controlled studies**	Mechanism-based reasoning
What are the RARE harms? (Treatment Harms)	trials or n-of-1 trial	Randomized trial or (exceptionally) observational study with dramatic effect			
Is this (early detection) test worthwhile? (Screening)	Systematic review of randomized trials		Non -randomized controlled cohort/follow-up study**	Case-series, case-control, or historically controlled studies**	Mechanism-based reasoning

V//VII VIL V > 1 L

檢索策略

System

Summary

Synopses of synthesis

Synthesis

Synopses of studies

Studies









華藝線上圖書館

檢索策略

臨床問題

Systematic Review Meta-Analysis RCT Cohort Study

限定5年內 Full text English 中文 台灣本土文 獻

符 合 PICO CYCH ER EBM

檢索策略

Annals of Internal Medicine®

Articles | 21 August 2007

How Quickly Do Systematic Reviews Go Out of Date? A Survival Analysis FREE

Authors: Kaveh G. Shojania, MD, Margaret Sampson, MLIS, Mohammed T. Ansari, MBBS, MMedSc, MPhil, Jun Ji, MD, MHA, Steve Doucette, MSc, and David Moher, PhD | <u>AUTHOR, ARTICLE, & DISCLOSURE INFORMATION</u>

Publication: Annals of Internal Medicine • Volume 147, Number 4

The median survival time without substantive new evidence for the meta-analyses was 5.5 years

搜尋Pubmed

TEXT AVAILABILITY

Abstract

Free full text

✓ Full text

ARTICLE TYPE **Books and Documents** Clinical Trial Meta-Analysis Randomized Controlled Trial Review Systematic Review **PUBLICATION DATE** 1 year 5 years 10 years **Custom Range**

輸入關鍵字 適當使用布林運算『AND』、『OR』

限定適當文章類型

『Meta-Analysis』文章

『Systematic Reviews』文章

『Randomized controlled Trial』文章

限定搜尋範圍

限定『5年』文章

限定『Full text』全文可供評讀

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搜尋Pubmed

Search	Actions	Details	Query	Results	Time
#6	•••	>	Search: #1 AND #2 Filters: in the last 5 years, Meta-Analysis, Systematic Review Sort by: Most Recent	2	21:59:38
#5	•••	>	Search: #1 AND #2 Filters: in the last 5 years, Meta-Analysis Sort by: Most Recent	2	21:59:24
#4	•••	>	Search: #1 AND #2 Filters: in the last 5 years Sort by: Most Recent	19	21:59:20
#3	•••	>	Search: #1 AND #2 Sort by: Most Recent	33	21:59:17
#2	•••	>	Search: sodium glucose cotransporter 2 inhibitors OR canagliflozin OR dapagliflozin OR empagliflozin OR ipragliflozin OR tofogliflozin OR luseogliflozin OR sergliflozin OR remogliflozin OR ertugliflozin OR sotagliflozin Sort by: Most Recent	13,639	21:59:05
#1	•••	~	Search: (overweight OR obes*) AND (non-diabe* OR no-diabe* OR "without diabe*") AND (adult) Sort by: Most Recent ("overweight" [MeSH Terms] OR "overweight" [All Fields] OR "overweighted" [All Fields] OR "overweightness" [All Fields] OR "overweights" [All Fields] OR "obes*" [All Fields]) AND ("non diabe*" [All Fields] OR "no diabe*" [All Fields] OR "without diabe*" [All Fields]) AND ("adult" [MeSH Terms] OR "adult" [All Fields] OR "adults" [All Fields] OR "adults" [All Fields] OR "overweight" [MeSH Terms] OR "overweight" [All Fields] OR "overweighted" [All Fields] OR "overweightness" [All Fields] OR "overweights" [All Fields] adult: "adult" [MeSH Terms] OR "adult" [All Fields] OR "adults" [All Fields] Fields] OR "adult's" [All Fields]	4,474	21:58:55

SR作為filters 篩選level 1文獻

P AND I

確認advanced search正確

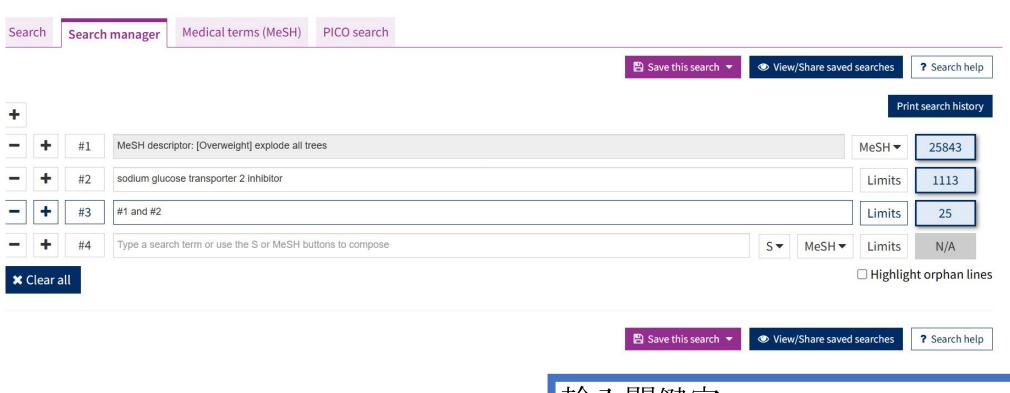
使用MeSH terms

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搜尋Cochrane

Advanced Search



輸入關鍵字 適當使用布林運算『AND』、『OR』 不限語言類型地區

搜尋Embase

Final Keywords:

('obesity' AND 'sodium glucose cotransporter 2 inhibitor' AND non-diabetic adult)/br AND ('body weight loss') AND ([Systematic Review]/lim OR [Randomized Controlled Trial]/lim OR [Meta Analysis]/lim) AND [2020-2024]/py



輸入關鍵字
『obesity』
『sodium glucose
cotransporter 2 inhibitor』
『body weight loss』
『non-diabetic adult』
適當使用布林運算
『AND』、『OR』
不限語言類型地區

搜尋結果

	輸入P、I 及適當同義 詞		選擇SR、MA 文章		選擇5年內 文章		符合臨床問 題文章
Cochrane Library	25	>	0	>	0	>	0
PublMed	33	>	19	•	2	•	1
Embase®	95	>	19	>	4	•	1
oiriti Librory 華藝線上圖書館	1	>	0	•	0	•	0

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選擇文獻

Efficacy and safety of GLP-1 receptor agonists versus SGLT-2 inhibitors in overweight/obese patients with or without diabetes mellitus: a systematic review and network meta-analysis	SR of RCTs	61 Trials	Up to 2022/1	P I C	× × ×
Sodium-Glucose Co-Transporter-2 Inhibitors in Non-Diabetic Adults With Overweight or Obesity: A Systematic Review and Meta-Analysis	SR of RCTs	6 Trials	Up to 2021/2	P I C	
Effect of Sodium-Glucose Cotransporter 2 Inhibitors on Weight Reduction in Overweight and Obese Populations without Diabetes: A Systematic Review and a Meta-Analysis	SR of RCTs	6 Trials	Up to 2021/5	P I C	X
	CYCH TK TOM -				

選擇最佳文獻



▶ Front Endocrinol (Lausanne). 2021 Aug 16;12:706914. doi: 10.3389/fendo.2021.706914 🖸

Sodium-Glucose Co-Transporter-2 Inhibitors in Non-Diabetic Adults With Overweight or Obesity: A Systematic Review and Meta-Analysis

Hanrui Zheng 1,2,†, Min Liu 3,†, Sheyu Li 4,5, Qingyang Shi 5, Shengzhao Zhang 6, Yiling Zhou 4, Na Su 1,2,*

▶ Author information ▶ Article notes ▶ Copyright and License information

PMCID: PMC8415407 PMID: <u>34484120</u>

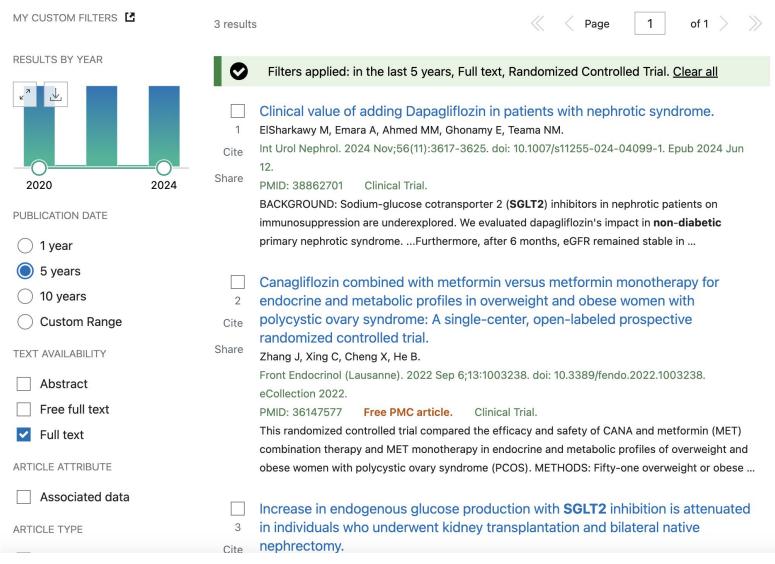


最佳研究設計 SR of RCT

✔ 較新的發表年份

✔ 最符合臨床情境 PICO

確認是否遺漏更新更好的 RCT



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Ask Acquire **Appraisal** Apply Audit

評讀工具



CASP Checklist: 10 questions to help you make sense of a Systematic Review

How to use this appraisal tool: Three broad issues need to be considered when appraising a systematic review study:

Are the results of the study valid? (Section A)

What are the results? (Section B)

Will the results help locally? (Section C)

✔ 針對效度直觀分析

✔ 針對結果直觀分析

✔ 共10個問題探討各面向

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此系統性回顧是否問了一個清楚、明確的臨床問題?



C

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Background: Sodium-glucose-cotransporter-2 (SGLT2) inhibitors have proven to be effective in improving glycemic control and lowering body weight in patients with type 2 diabetes mellitus. However, the efficacy and safety on weight loss in adults with overweight or obesity but not diabetes remain unclear. In this article, we aimed to identify the efficacy and safety of SGLT2 inhibitors in adults with overweight or obesity but not diabetes in randomized controlled studies (RCTs).

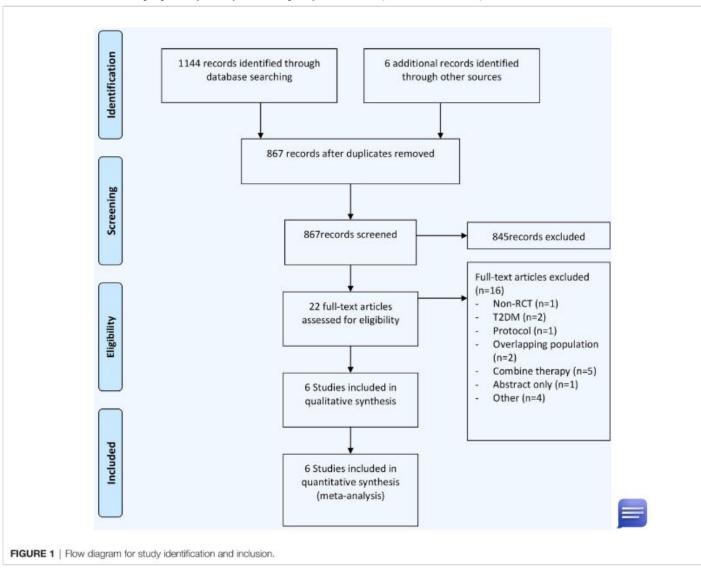
符合臨床問題

PICO清楚定義

P



作者是否尋找適當研究型態的文獻?



收錄符合問題RCT

清楚定義納入排除條件



你認為所有重要且相關的研究都被納入?

Literature Search

An extensive search for RCTs in Medline (Ovid SP), Embase (Ovid SP), Cochrane Central Register of Controlled Trials (Ovid SP), for studies published from the creation time of databases until February 20th, 2021, using the keywords: "Sodium-Glucose Transporter 2", "sodium glucose cotransporter 2 inhibitors", "canagliflozin", "dapagliflozin", "empagliflozin", "ipragliflozin", "tofogliflozin", "luseogliflozin", "sergliflozin", "remogliflozin", "ertugliflozin", "sotagliflozin", "overweight", "obesity" and "obese" (Supplementary Information 2), ClinicalTrial.gov was screened for potentially eligible studies. The reference lists of relevant published researches investigating the use of SGLT2 inhibitors in non-diabetes with overweight or obesity were also reviewed for potentially relevant studies. We contacted authors by email if the full-text was not available or if the outcomes were not enough.

搜尋各種資料庫

Protocol說明納入排除

不限年代語言國家種族

與專家聯繫

包含已/未發表文獻

搜尋英文/中文文獻

從參考資料再搜尋



你認為所有重要且相關的研究都被納入?

primary outcomes of interest, namely body weight and body mass index (BMI). Weight loss ≥ 5%, Waist circumference (WC), Hip circumference (HC), Waist/hip ratio (W/H) and adverse events were secondary outcomes. The adverse events included general adverse events and serious adverse events; (4) Study design: randomized controlled trials (RCTs) limited to the English language without restrictions of study size, follow-up length or publication year. No ethical approval and no contact with individual patients were required. The exclusion criteria were as follows: (1) including participants with pregnant; (2) animal experiments; (3) studies published in a language other than Chinese or English; (4) published as abstract only; (5) including patients with prediabetes.

搜尋各種資料庫

Protocol說明納入排除

不限年代語言國家種族

與專家聯繫

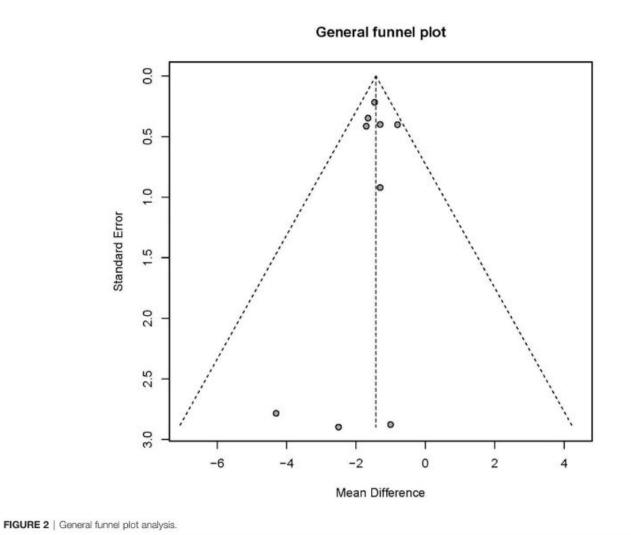
包含已/未發表文獻

搜尋英文/中文文獻

從參考資料再搜尋



你認為所有重要且相關的研究都被納入?



搜尋各種資料庫

Protocol說明納入排除

不限年代語言國家種族

與專家聯繫

包含已/未發表文獻

搜尋英文/中文文獻

從參考資料再搜尋



作者是否評估所納入研究文獻的品質?

Quality Assessment

Two independent reviewers (HZ and ML) assessed the risk of bias of the included studies according to the Cochrane Handbook for Systematic Reviews of Interventions (version 5.1), and the disagreement were resolved by consulting the 3rd reviewer (NS). We assessed the quality of the included studies concerning 7 aspects including random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, and selective reporting and other bias (12). Each of them was judged as low, high or unclear risk. Grading of Recommendations Assessment, Development and Evaluation (GRADE) (13, 14) tool was used to assess the evidence quality and provide evidence for future guidelines, concerning inconsistency, indirectness, imprecision, and other bias.

2個作者獨立評讀

使用GRADE評讀工具



作者將研究結果進行合併, 這樣的合併是否合理 ?

	SGLT	2 inhibi	tors	PI	acebo	E		Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV. Fixed, 95% Cl	IV. Fixed, 95% CI
Bays2014(CANA 100mg/d)	-2.8	2.9	85	-1.1	2.5	86	11.8%	-1.70 [-2.51, -0.89]	
Bays2014(CANA 300mg/d)	-2.4	2.9	96	-1.1	2.5	86	12.7%	-1.30 [-2.08, -0.52]	
Bays2014(CANA 50mg/d)	-1.9	2.9	95	-1.1	2.5	86	12.6%	-0.80 [-1.59, -0.01]	
Gonzalez-Ortiz2017(DAPA 10mg/d)	-3	4.88	13	1.3	8.77	13	0.3%	-4.30 [-9.76, 1.16]	·
Hollander2017(CANA 300mg/d)	-1.9	6.18	78	-0.6	5.23	76	2.4%	-1.30 [-3.11, 0.51]	
Hussey2010(SER 1500mg/d)	-1.55	0.25	6	-0.09	0.47	6	43.0%	-1.46 [-1.89, -1.03]	-
Hussey2010(SER 3000mg/d)	-1.74	0.71	6	-0.09	0.47	6	16.8%	-1.65 [-2.33, -0.97]	
Napolitano2014(REMO 750mg/d)	-7.6	3.46	8	-5.1	8.71	11	0.2%	-2.50 [-8.18, 3.18]	•
Napolitano2014(SER 3000mg/d)	-6.1	3.32	8	-5.1	8.71	11	0.2%	-1.00 [-6.64, 4.64]	· · ·
Total (95% CI)			395			381	100.0%	-1.42 [-1.70, -1.14]	•
Heterogeneity: Chi2 = 4.65, df = 8 (P =	= 0.79); l²	= 0%							
Test for overall effect: Z = 9.98 (P < 0.	.00001)		•						Favours SGLT2 inhibitors Favours Placebo

FIGURE 3 | Changes of body weight in patients receiving SGLT2 inhibitors versus placebo. CI, confidence interval; IV, inverse variance; SD, standardized deviation.

Body weight

異質性:低度(I²: 0%)

Fixed effects model

次族群分析

敏感性測試



這篇系統性文獻回顧的整體結果為何?

	SGLT	2 inhibi	tors	PI	acebo			Mean Difference	Mean Difference
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV. Fixed, 95% Cl	IV, Fixed, 95% CI
Bays2014(CANA 100mg/d)	-2.8	2.9	85	-1.1	2.5	86	11.8%	-1.70 [-2.51, -0.89]	-
Bays2014(CANA 300mg/d)	-2.4	2.9	96	-1.1	2.5	86	12.7%	-1.30 [-2.08, -0.52]	
Bays2014(CANA 50mg/d)	-1.9	2.9	95	-1.1	2.5	86	12.6%	-0.80 [-1.59, -0.01]	
Gonzalez-Ortiz2017(DAPA 10mg/d)	-3	4.88	13	1.3	8.77	13	0.3%	-4.30 [-9.76, 1.16]	•
Hollander2017(CANA 300mg/d)	-1.9	6.18	78	-0.6	5.23	76	2.4%	-1.30 [-3.11, 0.51]	
Hussey2010(SER 1500mg/d)	-1.55	0.25	6	-0.09	0.47	6	43.0%	-1.46 [-1.89, -1.03]	
Hussey2010(SER 3000mg/d)	-1.74	0.71	6	-0.09	0.47	6	16.8%	-1.65 [-2.33, -0.97]	
Napolitano2014(REMO 750mg/d)	-7.6	3.46	8	-5.1	8.71	11	0.2%	-2.50 [-8.18, 3.18]	•
Napolitano2014(SER 3000mg/d)	-6.1	3.32	8	-5.1	8.71	11	0.2%	-1.00 [-6.64, 4.64]	•
Total (95% CI)			395			381	100.0%	-1.42 [-1.70, -1.14]	•
Heterogeneity: Chi ² = 4.65, df = 8 (P =	0.79); l²	= 0%						r-cuence Track this all all	
Test for overall effect: Z = 9.98 (P < 0	.00001)	0.0966							Favours SGLT2 inhibitors Favours Placebo

Body weight

P值<0.05有顯著差異

FIGURE 3 | Changes of body weight in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.



結果精準嗎?

		Weight	T-4-1	200								
			Lotai	SD	Mean	Total	SD	Mean	Study or Subgroup			
	-1.70 [-2.51, -0.89]	11.8%	86	2.5	-1.1	85	2.9	-2.8	Bays2014(CANA 100mg/d)			
	-1.30 [-2.08, -0.52]	12.7%	86	2.5	-1.1	96	2.9	-2.4	Bays2014(CANA 300mg/d)			
	-0.80 [-1.59, -0.01]	12.6%	86	2.5	-1.1	95	2.9	-1.9	Bays2014(CANA 50mg/d)			
+	-4.30 [-9.76, 1.16]	0.3%	13	8.77	1.3	13	4.88	-3	Gonzalez-Ortiz2017(DAPA 10mg/d)			
-	-1.30 [-3.11, 0.51]	2.4%	76	5.23	-0.6	78	6.18	-1.9	Hollander2017(CANA 300mg/d)			
	-1.46 [-1.89, -1.03]	43.0%	6	0.47	-0.09	6	0.25	-1.55	Hussey2010(SER 1500mg/d)			
	-1.65 [-2.33, -0.97]	16.8%	6	0.47	-0.09	6	0.71	-1.74	Hussey2010(SER 3000mg/d)			
+	-2.50 [-8.18, 3.18]	0.2%	11	8.71	-5.1	8	3.46	-7.6	Napolitano2014(REMO 750mg/d)			
←	-1.00 [-6.64, 4.64]	0.2%	11	8.71	-5.1	8	3.32	-6.1	Napolitano2014(SER 3000mg/d)			
	-1.42 [-1.70, -1.14]	100.0%	381			395			Total (95% CI)			
_	-1.42 [-1.70, -1.14]	100.0%	381			395	= 0%		Total (95% CI) Heterogeneity: Chi ² = 4.65, df = 8 (P = Test for overall effect: Z = 9.98 (P < 0			
	←	-4.30 [-9.76, 1.16] -1.30 [-3.11, 0.51] -1.46 [-1.89, -1.03] -1.65 [-2.33, -0.97] -2.50 [-8.18, 3.18] -1.00 [-6.64, 4.64]	0.3% -4.30 [-9.76, 1.16] 2.4% -1.30 [-3.11, 0.51] 43.0% -1.46 [-1.89, -1.03] 16.8% -1.65 [-2.33, -0.97] 0.2% -2.50 [-8.18, 3.18]	13 0.3% -4.30 [-9.76, 1.16] 76 2.4% -1.30 [-3.11, 0.51] 6 43.0% -1.46 [-1.89, -1.03] 6 16.8% -1.65 [-2.33, -0.97] 11 0.2% -2.50 [-8.18, 3.18] 11 0.2% -1.00 [-6.64, 4.64]	8.77 13 0.3% -4.30 [-9.76, 1.16] 5.23 76 2.4% -1.30 [-3.11, 0.51] 0.47 6 43.0% -1.46 [-1.89, -1.03] 0.47 6 16.8% -1.65 [-2.33, -0.97] 8.71 11 0.2% -2.50 [-8.18, 3.18] 8.71 11 0.2% -1.00 [-6.64, 4.64]	1.3 8.77 13 0.3% -4.30 [-9.76, 1.16] -0.6 5.23 76 2.4% -1.30 [-3.11, 0.51] -0.09 0.47 6 43.0% -1.46 [-1.89, -1.03] -0.09 0.47 6 16.8% -1.65 [-2.33, -0.97] -5.1 8.71 11 0.2% -2.50 [-8.18, 3.18] -5.1 8.71 11 0.2% -1.00 [-6.64, 4.64]	13 1.3 8.77 13 0.3% -4.30 [-9.76, 1.16] 78 -0.6 5.23 76 2.4% -1.30 [-3.11, 0.51] 6 -0.09 0.47 6 43.0% -1.46 [-1.89, -1.03] 6 -0.09 0.47 6 16.8% -1.65 [-2.33, -0.97] 8 -5.1 8.71 11 0.2% -2.50 [-8.18, 3.18] 8 -5.1 8.71 11 0.2% -1.00 [-6.64, 4.64]	4.88 13 1.3 8.77 13 0.3% -4.30 [-9.76, 1.16] 6.18 78 -0.6 5.23 76 2.4% -1.30 [-3.11, 0.51] 0.25 6 -0.09 0.47 6 43.0% -1.46 [-1.89, -1.03] 0.71 6 -0.09 0.47 6 16.8% -1.65 [-2.33, -0.97] 3.46 8 -5.1 8.71 11 0.2% -2.50 [-8.18, 3.18] 3.32 8 -5.1 8.71 11 0.2% -1.00 [-6.64, 4.64]	-3 4.88 13 1.3 8.77 13 0.3% -4.30 [-9.76, 1.16] -1.9 6.18 78 -0.6 5.23 76 2.4% -1.30 [-3.11, 0.51] -1.55 0.25 6 -0.09 0.47 6 43.0% -1.46 [-1.89, -1.03] -1.74 0.71 6 -0.09 0.47 6 16.8% -1.65 [-2.33, -0.97] -7.6 3.46 8 -5.1 8.71 11 0.2% -2.50 [-8.18, 3.18] -6.1 3.32 8 -5.1 8.71 11 0.2% -1.00 [-6.64, 4.64] 395 381 100.0% -1.42 [-1.70, -1.14]			

FIGURE 3 | Changes of body weight in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

Body weight

Sample size: 776

95%CI: -1.42 (-1.70, -1.14) 無碰到統計無效線

95%CI:-1.42 (-1.70, -1.14) 無碰到臨床無效線



作者將研究結果進行合併, 這樣的合併是否合理 ?

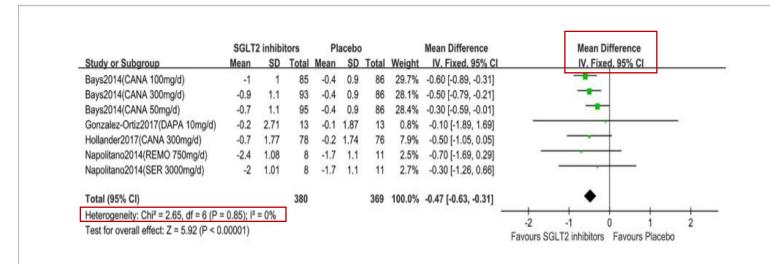


FIGURE 4 | Changes of BMI in patients receiving SGLT2 inhibitors *versus* placebo. BMI, body mass index; CI, confidence interval; IV, inverse variance; SD, standardized deviation.

BMI

異質性: 低度(I²:0%)

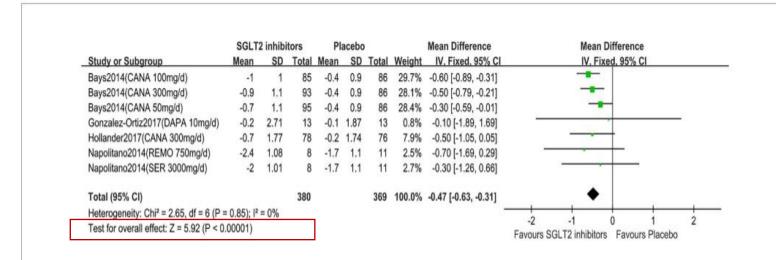
Fixed effects model

次族群分析

敏感性測試



這篇系統性文獻回顧的整體結果為何?



BMI

P值<0.05有顯著差異

FIGURE 4 | Changes of BMI in patients receiving SGLT2 inhibitors *versus* placebo. BMI, body mass index; CI, confidence interval; IV, inverse variance; SD, standardized deviation.



結果精準嗎?

Study or Subgroup Mean SD Total Mean SD Total Weight IV, Fixed, 95% CI IV, Fixed, 95% CI Bays2014(CANA 100mg/d) -1 1 85 -0.4 0.9 86 29.7% -0.60 [-0.89, -0.31] Bays2014(CANA 300mg/d) -0.9 1.1 93 -0.4 0.9 86 28.1% -0.50 [-0.79, -0.21] Bays2014(CANA 50mg/d) -0.7 1.1 95 -0.4 0.9 86 28.4% -0.30 [-0.59, -0.01] Gonzalez-Ortiz2017(DAPA 10mg/d) -0.2 2.71 13 -0.1 1.87 13 0.8% -0.10 [-1.89, 1.69] Hollander2017(CANA 300mg/d) -0.7 1.77 78 -0.2 1.74 76 7.9% -0.50 [-1.05, 0.05] Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]		20000	2 inhibi		1/2/2	acebo			Mean Difference			an Diffe	100000	
Bays2014(CANA 300mg/d) -0.9 1.1 93 -0.4 0.9 86 28.1% -0.50 [-0.79, -0.21] Bays2014(CANA 50mg/d) -0.7 1.1 95 -0.4 0.9 86 28.4% -0.30 [-0.59, -0.01] Gonzalez-Ortiz2017(DAPA 10mg/d) -0.2 2.71 13 -0.1 1.87 13 0.8% -0.10 [-1.89, 1.69] Hollander2017(CANA 300mg/d) -0.7 1.77 78 -0.2 1.74 76 7.9% -0.50 [-1.05, 0.05] Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.5% -0.70 [-1.69, 0.29] Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Fixed, 95% CI		IV.	Fixed, 9	95% CI	
Bays2014(CANA 50mg/d) -0.7 1.1 95 -0.4 0.9 86 28.4% -0.30 [-0.79, -0.21] Gonzalez-Ortiz2017(DAPA 10mg/d) -0.2 2.71 13 -0.1 1.87 13 0.8% -0.10 [-1.89, 1.69] Hollander2017(CANA 300mg/d) -0.7 1.77 78 -0.2 1.74 76 7.9% -0.50 [-1.05, 0.05] Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.5% -0.70 [-1.89, 0.29] Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Bays2014(CANA 100mg/d)	-1	1	85	-0.4	0.9	86	29.7%	-0.60 [-0.89, -0.31]					
Gonzalez-Ortiz2017(DAPA 10mg/d) -0.2 2.71 13 -0.1 1.87 13 0.8% -0.10 [-1.89, 1.69] Hollander2017(CANA 300mg/d) -0.7 1.77 78 -0.2 1.74 76 7.9% -0.50 [-1.05, 0.05] Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.5% -0.70 [-1.69, 0.29] Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Bays2014(CANA 300mg/d)	-0.9	1.1	93	-0.4	0.9	86	28.1%	-0.50 [-0.79, -0.21]		-			
Hollander2017(CANA 300mg/d) -0.7 1.77 78 -0.2 1.74 76 7.9% -0.50 [-1.05, 0.05] Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.5% -0.70 [-1.69, 0.29] Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Bays2014(CANA 50mg/d)	-0.7	1.1	95	-0.4	0.9	86	28.4%	-0.30 [-0.59, -0.01]					
Napolitano2014(REMO 750mg/d) -2.4 1.08 8 -1.7 1.1 11 2.5% -0.70 [-1.69, 0.29] Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Gonzalez-Ortiz2017(DAPA 10mg/d)	-0.2	2.71	13	-0.1	1.87	13	0.8%	-0.10 [-1.89, 1.69]	-		-		-0
Napolitano2014(SER 3000mg/d) -2 1.01 8 -1.7 1.1 11 2.7% -0.30 [-1.26, 0.66]	Hollander2017(CANA 300mg/d)	-0.7	1.77	78	-0.2	1.74	76	7.9%	-0.50 [-1.05, 0.05]		-	•		
	Napolitano2014(REMO 750mg/d)	-2.4	1.08	8	-1.7	1.1	11	2.5%	-0.70 [-1.69, 0.29]			\neg		
T-1-1/050/ CD	Napolitano2014(SER 3000mg/d)	-2	1.01	8	-1.7	1.1	11	2.7%	-0.30 [-1.26, 0.66]			*	-	
10tal (95% Cl) 380 369 100.0% -0.47 [-0.63, -0.31]	Total (95% CI)			380			369	100.0%	-0.47 [-0.63, -0.31]			•		

FIGURE 4 | Changes of BMI in patients receiving SGLT2 inhibitors *versus* placebo. BMI, body mass index; CI, confidence interval; IV, inverse variance; SD, standardized deviation.

BMI

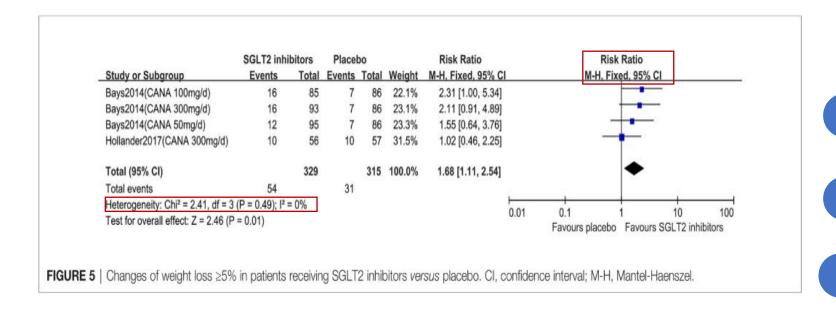
Sample size: 749

95%CI:-0.47(-0.63, -0.31) 無碰到統計無效線

95%CI: -0.47(-0.63, -0.31) 無碰到臨床無效線



作者將研究結果進行合併, 這樣的合併是否合理 ?



Weight loss ≥ 5%

異質性: 低度(I²:0%)

Fixed effects model

次族群分析

敏感性測試



這篇系統性文獻回顧的整體結果為何?

Weight loss ≥ 5%

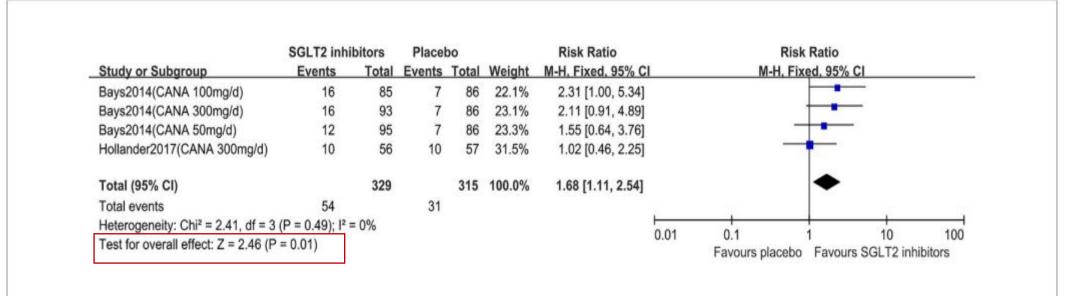


FIGURE 5 | Changes of weight loss ≥5% in patients receiving SGLT2 inhibitors versus placebo. CI, confidence interval; M-H, Mantel-Haenszel.

P值<0.05有顯著差異



結果精準嗎?

	SGLT2 inhi		Placel	18		Risk Ratio			Risk Ratio		
Study or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% C		M	-H, Fixed, 95%	CI	
Bays2014(CANA 100mg/d)	16	85	7	86	22.1%	2.31 [1.00, 5.34]			-	_	
Bays2014(CANA 300mg/d)	16	93	7	86	23.1%	2.11 [0.91, 4.89]			-	-	
Bays2014(CANA 50mg/d)	12	95	7	86	23.3%	1.55 [0.64, 3.76]			-		
Hollander2017(CANA 300mg/d)	10	56	10	57	31.5%	1.02 [0.46, 2.25]			_		
Total (95% CI)		329		315	100.0%	1.68 [1.11, 2.54]			•		
Total events	54	1.50000	31	JAN STATE	100000000000000000000000000000000000000		W	200		33	
Heterogeneity: Chi ² = 2.41, df = 3	(P = 0.49); I ²	= 0%					0.04	1		10	- 5
Test for overall effect: Z = 2.46 (P	= 0.01)						0.01	0.1 Favours pl		10 s SGLT2 inh	

FIGURE 5 | Changes of weight loss ≥5% in patients receiving SGLT2 inhibitors versus placebo. CI, confidence interval; M-H, Mantel-Haenszel.

Weight loss ≥ 5%

Sample size: 644

95%CI:1.68(1.11, 2.54)無 碰到統計無效線

95%CI: 1.68(1.11, 2.54)有 碰到臨床無效線

NNT:16



作者將研究結果進行合併, 這樣的合併是否合理 ?

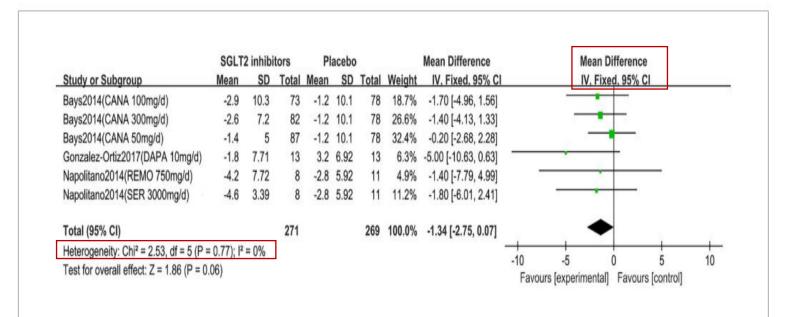


FIGURE 6 | Changes of waist circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

腰圍

異質性: 低度(I²:0%)

Fixed effects model

次族群分析

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這篇系統性文獻回顧的整體結果為何?

腰圍

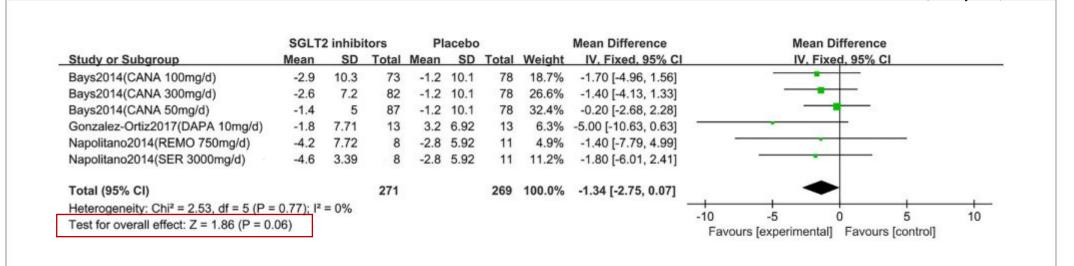


FIGURE 6 | Changes of waist circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

P值>0.05無顯著差異



結果精準嗎?

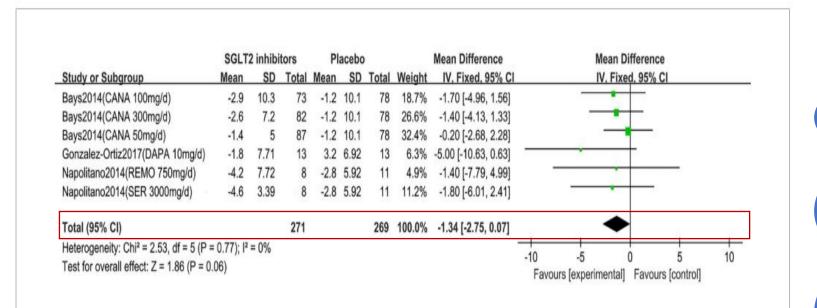


FIGURE 6 | Changes of waist circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

腰圍

Sample size: 540

95%CI:-1.34(-2.75, 0.07)有 碰到統計無效線

95%CI: -1.34(-2.75, 0.07) 有碰到臨床無效線



作者將研究結果進行合併, 這樣的合併是否合理 ?

Mean Difference Mean Difference SGLT2 inhibitors Placebo Study or Subgroup SD Total Mean SD Total Weight IV, Fixed, 95% CI IV, Fixed, 95% CI Bays2014(CANA 100mg/d) 0.1 10.4 77 19.2% -2.20 [-5.58, 1.18] Bays2014(CANA 300mg/d) 81 0.1 10.4 77 26.1% -3.10 [-5.99, -0.21] Bays2014(CANA 50mg/d) 86 0.1 10.4 77 35.7% -2.10 [-4.57, 0.37] Napolitano2014(REMO 750mg/d) 8 -3.1 3.3 9.4% 0.10 [-4.71, 4.91] Napolitano2014(SER 3000mg/d) -1.9 6.28 8 -3.1 3.3 11 1.20 [-3.57, 5.97] Total (95% CI) 253 100.0% -1.86 [-3.33, -0.38] Heterogeneity: $Chi^2 = 3.00$, df = 4 (P = 0.56); $I^2 = 0\%$ 0 Test for overall effect: Z = 2.46 (P = 0.01) Favours [experimental] Favours [control]

FIGURE 7 | Changes of hip circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

臀圍

異質性: 低度(I²:0%)

Fixed-effects model

次族群分析

敏感性測試



這篇系統性文獻回顧的整體結果為何?

臀圍

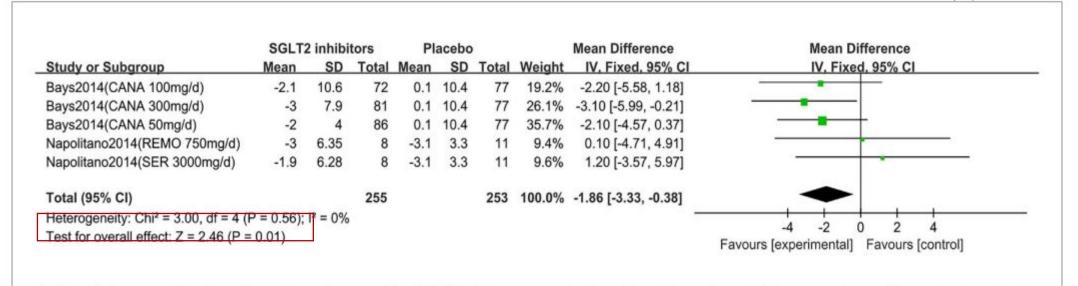


FIGURE 7 | Changes of hip circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

P值<0.05有顯著差異



結果精準嗎?

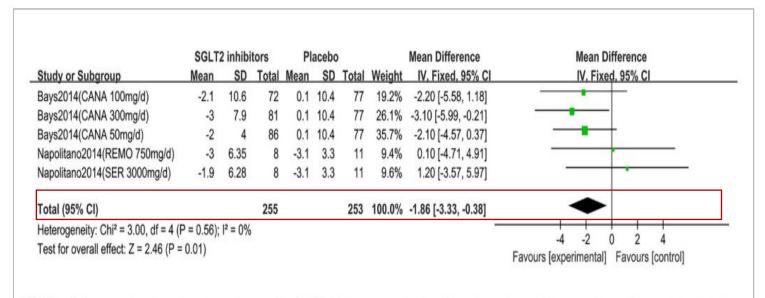


FIGURE 7 | Changes of hip circumference in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

臀圍

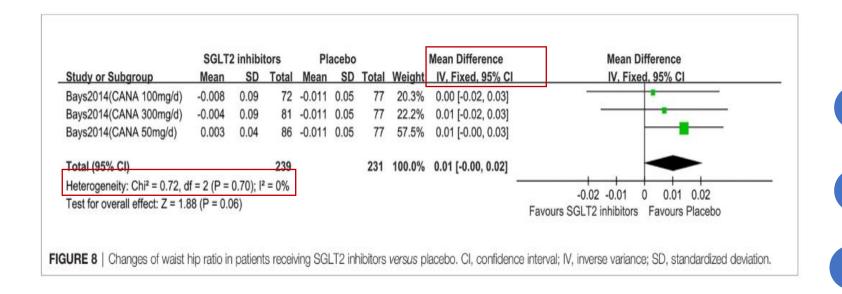
Sample size: 508

95%CI:-1.85(-3.33, -0.38) 無碰到統計無效線

95%CI: -1.85(-3.33, -0.38) 有碰到臨床無效線



作者將研究結果進行合併, 這樣的合併是否合理 ?



腰臀比

異質性: 低度(I²: 0%)

Fixed-effects model

次族群分析

敏感性測試



這篇系統性文獻回顧的整體結果為何?

腰臀比

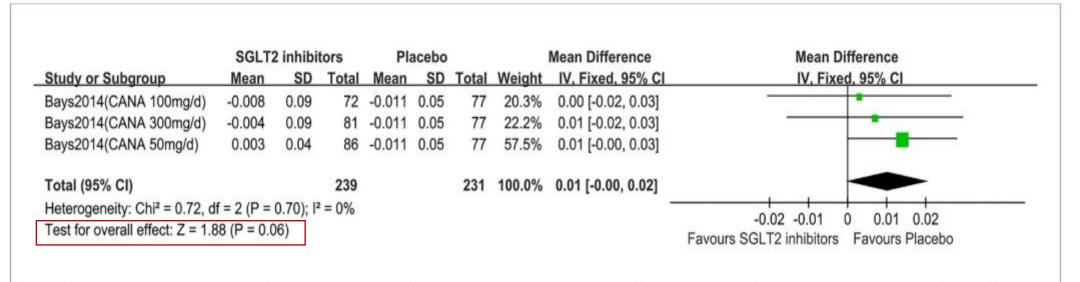
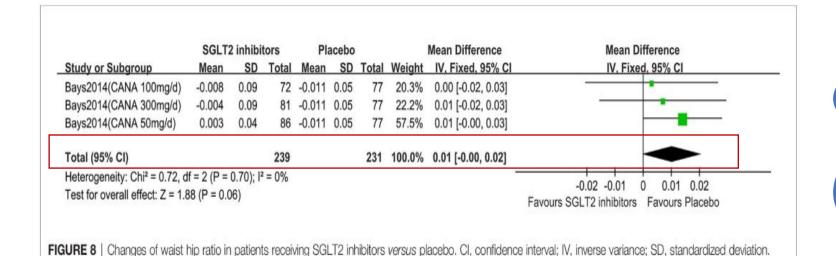


FIGURE 8 | Changes of waist hip ratio in patients receiving SGLT2 inhibitors versus placebo. Cl, confidence interval; IV, inverse variance; SD, standardized deviation.

P值>0.05無顯著差異



結果精準嗎?



腰臀比

Sample size: 470

95%CI:0.01(-0.00, 0.03)有 碰到統計無效線

95%CI: 0.01(-0.00, 0.03)有 碰到臨床無效線



評定個別研究結果的證據品質: GRADE

			降級			證據
	風險偏差	不一致性	間接性	不精確性	發表偏差	品質 GRADE
體重	1		1	•		Very low
BMI	1		•			Low
體重下降5%	o o		1			Moderate
腰圍	1		1			Low
臀圍	1		•			Low
腰臀比			1			Moderate

CYCH ER EBM

CYCH ER EBM

此研究結果是否可應用到當地的族群?

	評讀文獻	臨床情境
性別	Female 較多	Male
共病	無糖尿病	無糖尿病
種族	歐美較多	台灣
年齡	18-60 years old	30 years old
疾病	肥胖、超重	肥胖、脂肪肝
介入	SGLT2 inhibitor	SGLT2 inhibitor
	CYCH ER EBM	

此研究結果是否可應用到當地的族群?

TABLE 1 | Baseline characteristics of each included study (n=6)

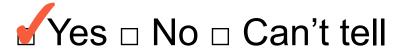
Author (year)	1	С	Nun	nber	Male	e (%)	Mear (yea	•	Mean weigh	body nt (kg)		n BMI /m²)	Patients	Country	Follow-up (weeks)	Outcomes	Funding
			Т	С	т	С	Т	С	т	С	Т	С					
Færch2021 (15)	DAPA 10mg qd	PBO	30	30	43	40	61.4 ± 8.5	57.2 ± 9.9	M:103.7 ± 17.6 F:82.5 ± 13.7	M:99.0 ± 15.1 F:92.1 ± 25.3	M:31.6 ± 3.8 F:30.4 ± 5.1	M:30.4 ± 3.2 F:34.0 ± 8.4	Overweight/ Obese with PreDM	Denmark	13	BW,BMI,WC,W/H	Novo Nordisk Foundation,AstraZeneca AB
Hollander2017 (16)	CANA 300mg qd	PBO	84	82	19	18.3	45.2 ± 11.0	44.8 ± 11.1	103.3 ± 19.1	104.3 ± 18.2	37.3 ± 4.7	38.0 ± 5.2	Overweight/ Obese Without DM	US	26	BW,BW(loss) ≥5%, BMI,WC	Janssen Research & Development, LLC
Gonzalez- Ortiz2017 (17)	DAPA 10mg qd	PBO	13	13	15.4	15.4	46.5 ± 5.2	45.0 ± 6.8	68.0 ± 4.6	73.0 ± 8.2	27.3 ± 2.0	27.3 ± 1.6	Overweight Without DM	NR	12	BW,BMI,WC	
Bays2014 (18)	CANA 50mg qd	PBO		89	12	16	44.9 ± 11.8	45.1 ± 11.9	98.1 ± 16.0	102.2 ± 19.9	36.6 ± 5.3	36.6 ± 5.5	Overweight/ Obese Without	US and Puerto Rico	12	BW,BW(loss) ≥5%, BMI, WC,HC,W/H	Janssen Global Services, LLC
	CANA 100mg ad	PBO	93	89	12	16	45.8 ± 11.0	45.1 ± 11.9	105.0 ± 16.6	102.2 ± 19.9	37.9 ± 5.1	36.6 ± 5.5	DM				
	CANA 300mg	PBO	96	89	10	16	43.5 ± 11.0	45.1 ± 11.9	100.2 ± 18.0	102.2 ± 19.9	36.9 ± 5.3	36.6 ± 5.5					
Napolitano2014 (19)	REMO 250mg tid	PBO	9	12	7	'3	42 ±	13.0	101 ±	± 14.6	33 =	± 2.4	Healthy obese	UK	8	BW,BMI,WC,HC	Glaxo Smith Kline
	SER 1000mg tid	PBO	9	12													
Hussey2010 (20)	SER 500mg tid	PBO	6	6	44	1.4	18-	-55	N	IR	25	-35	Healthy Overweight/	US	2	BW	Glaxo Smith Kline
	SER 1000mg tid	PBO	6	6									Obese				

文獻與情境類似

較少亞洲國家 種族bias:moderate

可應用到當地族群

DM, diabetes mellitus; NA, not applicable; I, Intervention C, Control; DAPA, dapagliflozin; CANA, canagliflozin; REMO, remogliflozin; SER, sergliflozin; PBO, placebo; BMI, body mass index; BW, body weight; WC, Waist circumference; HC, Hip circumference; W/H, Waist/hip ratio.



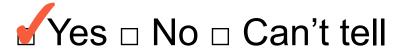
是否所有重要的臨床結果都被考量到?

TABLE 1 | Baseline characteristics of each included study (n=6).

Author (year)	1	С	Nur	nber	Male	e (%)	Mear (yea	•	Mean weigh	body nt (kg)		n BMI /m²)	Patients	Country	Follow-up (weeks)	Outcomes	Funding
			Т	С	т	С	Т	С	Т	С	Т	С					
Færch2021 (15)	DAPA 10mg qd	PBO	30	30	43	40	61.4 ± 8.5	57.2 ± 9.9	M:103.7 ± 17.6 F:82.5 ± 13.7	M:99.0 ± 15.1 F:92.1 ± 25.3	M:31.6 ± 3.8 F:30.4 ± 5.1	M:30.4 ± 3.2 F:34.0 ± 8.4	Overweight/ Obese with PreDM	Denmark	13	BW,BMI,WC,W/H	Novo Nordisk Foundation, Astra Zeneca AB
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Bays2014 (18)	CANA 50mg qd CANA 100mg	PBO PBO	0.0	89 89	12 12	16 16	44.9 ± 11.8 45.8 ± 11.0	45.1 ± 11.9 45.1 ± 11.9	98.1 ± 16.0 105.0 ± 16.6	102.2 ± 19.9 102.2 ± 19.9	36.6 ± 5.3 37.9 ± 5.1	36.6 ± 5.5 36.6 ± 5.5	Overweight/ Obese Without DM	US and Puerto Rico	12	BW,BW(loss) ≥5%, BMI, WC,HC,W/H	Janssen Global Services, LLC
	qd CANA 300mg qd	PBO	96	89	10	16	43.5 ± 11.0	45.1 ± 11.9	100.2 ± 18.0	102.2 ± 19.9	36.9 ± 5.3	36.6 ± 5.5					
Napolitano2014 (19)	REMO 250mg tid	PBO	9	12	7	'3	42 ±	13.0	101 ±	14.6	33 =	± 2.4	Healthy obese	UK	8	BW,BMI,WC,HC	Glaxo Smith Kline
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	SER 1000mg tid	PBO	6	6									Obese				

有考慮到各種臨床結果

DM, diabetes mellitus; NA, not applicable; I, Intervention C, Control; DAPA, dapagliflozin; CANA, canagliflozin; REMO, remogliflozi; SER, sergliflozin; PBO, placebo; BMI, body mass index; BW, body weight; WC, Waist circumference; HC, Hip circumference; W/H, Waist/hip ratio.



是否所有重要的臨床結果都被考量到?

有考慮到各種臨床結果

TABLE 3 | Adverse events reported in all included studies.

Adverse events	Numbers (Studies)	SGLT2 in	hibitors	Con	trol	Relative risk (95% CI)
		Events	Total	Events	Total	
AEs leading to discontinuation	4	22	401	9	385	2.25 (1.08 to 4.68)
hypoglycemia	1	3	287	6	267	0.49 (0.13 to 1.83)
Urinary tract infection	5	31	420	20	398	1.45 (0.83 to 2.54)
Genital mycotic infection	4	51	393	9	374	5.36 (2.72 to 10.59)
Vulvovaginal mycotic infection	2	31	371	3	349	9.14 (3.00 to 27.78)
Nausea	2	23	371	6	349	3.54 (1.47 to 8.53)
Sinusitis	2	11	371	3	349	2.67 (0.90 to 7.95)

Cl, confidence interval.



付出傷害和花費換得介入措施的益處是否 值得?

介入措施

針對非糖尿病,超重或肥胖 患者,使用SGLT2 inhibitor

花費

健保只給付糖尿病患者 花費約50元



優點

有減輕體重、BMI的效果



副作用

可能有低血糖、泌尿道感染、噁心等副作用



付出傷害和花費換得介入措施的益處是否 值得?

項目	有形支出	無形支出	預估總價
看診費用	一次450元	看診心理負擔	一個月需4000元
輾轉車費	一趟250元	來回舟車勞頓	一個月需1000元
請假成本	時薪190元	減少休息時間	一個月需3000元

證據等級	缺點	優點	其他選擇
Level 3	效果因人而異	清熱消腫排膿	民俗療法
Level 3	治療效果有限	便宜簡單上手	衛生保健
Level 3	需要持之以恆	調理生理功能	飲食運動



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AI輔助

成本類型	介入方式(SGLT2 抑制劑)	對照組方式(生活方式改變)
直接成本	1. 藥物費用:約 400~600 USD/月(國際市場價)	1. 飲食顧問費:約 50~100 USD/次
	2. 醫療費用:定期檢查(如血檢、肝功能檢測),約 100~200 USD/次	2. 健身房會員費:約 30~50 USD/月
		3. 健康食品額外支出:約 100~200 USD/月
間接成	1. 醫療訪問時間損耗:每月 1~2 小時	1. 運動時間損耗:每週 3~5 小時
	2. 交通費用:約 10~50 USD/月	2. 調整飲食規劃的時間損耗:約每週2
無形成本	1. 副作用壓力:輕微副作用(如泌尿感染),可能導致心理不適	1. 自律壓力:飲食限制或規律運動的心 理負擔
	2. 長期依從壓力:需定期服藥	2. 持續性壓力:無法達到減重目標的挫 敗感
總成本	約 600~800 USD/月(包括所有直接、間接及無形成本)	約 150~300 USD/月(以最低限度的方式計算)

評定證據等級

	清楚明確的臨床問題
	收納適當研究類型
V	所有研究都被納入
	研究文獻的品質
	結果合併的合理性
	整體結果
	精準性
D	臨床應用
	成本效益

研究品質差



研究不精準

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	CASP question	Y/N
1	此系統性回顧是否問了一個清楚、明確的臨床問題?	Υ
2	作者是否尋找適當研究型態的文獻?	Υ
3	你認為所有重要且相關的研究都被納入?	Υ
4	作者是否評估所納入研究文獻的品質?	Υ
5	作者將研究結果進行合併, 這樣的合併是否合理?	Υ
6	這篇系統性文獻回顧的整體結果為何?	Υ
7	結果精準嗎?	N
8	此研究結果是否可應用到當地的族群?	Υ
9	是否所有重要的臨床結果都被考量到?	Υ
10	付出傷害和花費換得介入措施的益處是否值得?	Υ





Ask Acquire Appraisal **Apply** Audit

有形無形支出及隱藏成本

000	SGLT2 inhibitor	飲食運動
醫療成本	一個月藥費約兩千 第一個月門診追蹤約兩千	門診追蹤 保健食品
生產力損失	請假成本 照護成本	飲食運動費用
社會成本	影響病人照護 同事關係	影響病人照護 同事關係
心靈成本	生活品質 副作用	自律壓力 心理壓力
總成本	一個月約一萬元	一個月約兩千元

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成本效益比

CER: Control event rate

= 16

RR: Relative risk

NNT

9.8% x 0.68

需要治療 16 名患者, 才能使 1 名患者 達到體重下降 ≥5% 的效果

ICER

COPE

= NNT x 時間 x 治療費用 = 16 x 3m x 4000/ =192k





Ask Acquire Appraisal Apply **Audit**

4E Model 從證據到應用

實證醫學 Evidence

證據&建議等級:level 2

GRADE: moderate ~ very

low

患者期待 Expectation

小莊希望達到減重效果 不常運動

專家意見 Expert

建議對於肥胖過重非糖尿病的患者,使用SGLT2 inhibitor可達到減重效果

社經考量 Environment

非健保給付條件要自費用藥和門診追蹤

CYCH ER EBM

Evidence to Decision (EtD) framework

這個問題重要嗎	重要
整體證據品質如何	證據品質中~低
價值觀與重要預後確定性	可能有不確定性
效果是否足夠大	效果不夠大
副作用是否足夠小	副作用可能小
效果是否超越副作用	效果可能大於副作用
單位好處所需花費小	單位好處所需花費可能小
是否可被使用者接受	接受度高
是否可被使用者執行	執行度高

Moberg, J., Oxman, A.D., Rosenbaum, S. et al. The GRADE Evidence to Decision (EtD) framework for health system and public health decisions. Health Res Policy Sys 16, 45 (2018)

共享決策

文獻利>弊

李宜恭醫師《醫策會SDM上課教材》





是

我們建議應該做並解釋行動方案

我們建議考慮做 提供選項輔助決策

柄人 偏好



我們建議考慮做 提供選項輔助決策

我們建議考慮做 提供選項輔助決策

回答病人問題一以去學術化術語方式



小莊先生您好,經過我們團隊縝密的實證 搜尋後,目前現有最佳證據是由**系統性回 顧文獻**支持,對於**肥胖過重非糖尿病**的患 者,使用**SGLT2** inhibitor可達到**減重**效果 ,用藥和門診一個月約**花費四千元**,血脂 異常和脂肪肝是**高危險族群**,可以同時**搭 配飲食控制和養成運動習慣。**



AI輔助

我想要您協助評估文章的下面這個題目:

###

作者是否尋找適當研究型態的文獻?

其中需要評估

Q1: 收錄符合問題RCT (Randomized Controlled Trial),若有則這題

為YES

Q2:清楚定義納入條件,若有則這題為YES Q3:清楚定義排除條件,若有則這題為YES

各項小細項問題定義清楚

###

###

顯示方法為

Q1:yes OR no {文獻參考的原始原文}

Q2:yes OR no {文獻參考的原始原文}

Q3:yes OR no {文獻參考的原始原文}

請將結果以表格呈現

以下是針對 Frontiers 文獻的「作者是否尋找適當研究型態的文獻」評估結果:

問題	回答	文獻參考的原始原文
Q1	YES	"We included studies meeting the following criteria: (1) Participants: adults with overweight or obesity but not diabetes undergoing SGLT2 inhibitors based on the study definition; (4) Study design: randomized controlled trials (RCTs)" (Sodium-Glucose Co-Trans)
Q2	YES	"We included studies meeting the following criteria: (2) Interventions/comparisons: using SGLT2 inhibitors as a monotherapy and placebo as the control. All included participants received standardized advice on diet and physical activity throughout the trial; (3) Outcomes: reporting one of the primary outcomes of interest, namely body weight and body mass index (BMI)." (Sodium-Glucose Co-Trans)
Q3	YES	"The exclusion criteria were as follows: (1) including participants with pregnant; (2) animal experiments; (3) studies published in a language other than Chinese or English; (4) published as abstract only; (5) including patients with prediabetes." (Sodium-Glucose Co-Trans)

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AI輔助

我想要您協助評估文章的下面這個題目:

###

Do you think the important, relevant studies were included? 作者有沒有可能遺漏掉重要、相關的研究?

其中需要評估

Q1:搜尋三個以上資料庫,實證資料庫包含Cochrane Library、

EMBASE、MEDLINE、pubmed等,若三種以上則這題為Yes

Q2:關鍵字明確,包含MESH TERM與其他同義字詞,若有則這題為

YES

Q3:搜尋是否限制使用英語文獻,有限制則是YES

Q4:搜尋reference list、會議壁報論文等,若有則這題為YES

Q5:與專家聯繫或是與收入文獻的原作者聯繫等,若有則這題為YES

各項小細項問題定義清楚

###

###

顯示方法為

Q1:yes OR no {文獻參考的原始原文}

Q2:yes OR no {文獻參考的原始原文}

Q3:yes OR no {文獻參考的原始原文}

Q4:yes OR no {文獻參考的原始原文}



請將結果以表格呈現

以下是針對 Frontiers 文獻的「作者是否可能遺漏重要、相關的研究」的評估結果:

問題	回答	文獻參考的原始原文
Q1	YES	"We searched for RCTs concerning SGLT2 inhibitors in adults with overweight or obesity but not diabetes in Medline (Ovid SP), Embase (Ovid SP), Cochrane Central Register of Controlled Trials (Ovid SP), and ClinicalTrials.gov up to February 2021." (Sodium-Glucose Co-Trans)
Q2	YES	"using the keywords: 'Sodium-Glucose Transporter 2', 'sodium glucose cotransporter 2 inhibitors', 'canagliflozin', 'dapagliflozin', 'empagliflozin', 'ipragliflozin', 'tofogliflozin', 'luseogliflozin', 'sergliflozin', 'remogliflozin', 'ertugliflozin', 'sotagliflozin', 'overweight', 'obesity' and 'obese'." (Sodium-Glucose Co-Trans)
Q3	YES	"We included studies meeting the following criteria limited to the English language without restrictions of study size, follow-up length or publication year." (Sodium-Glucose Co-Trans)
Q4	YES	"The reference lists of relevant published researches investigating the use of SGLT2 inhibitors in non-diabetes with overweight or obesity were also reviewed for potentially relevant studies." (Sodium-Glucose Co-Trans)
Q5	YES	"We contacted authors by email if the full-text was not available or if the outcomes were not enough." (Sodium-Glucose Co-Trans)

CYCH ER EBM

AI輔助

我想要您協助評估文章的下面這個題目:

###

作者是否評估所納入研究文獻的品質?

其中需要評估

Q1:兩個以上author或reviewer獨立評讀,若有則這題為YES

Q2:使用適當的工具來評讀風險偏差或證據等級,若有則這題為YES

各項小細項問題定義清楚

###

###

顯示方法為

Q1:yes OR no {文獻參考的原始原文}

Q2:yes OR no {文獻參考的原始原文}

請將結果以表格呈現

以下是針對 Frontiers 文獻「作者是否評估所納入研究文獻的品質」的評估結果:

問題	回答	文獻參考的原始原文
Q1	YES	"Two independent reviewers (HZ and ML) assessed the risk of bias of the included studies according to the Cochrane Handbook for Systematic Reviews of Interventions (version 5.1), and the disagreement were resolved by consulting the 3rd reviewer (NS)." (Sodium-Glucose Co-Trans)
Q2	YES	"We assessed the quality of the included studies concerning 7 aspects including random sequence generation, allocation concealment, blinding of participants and personnel, blinding of outcome assessment, incomplete outcome data, and selective reporting and other bias." (Sodium-Glucose Co-Trans)

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AI輔助

關於研究結果,你可以把數據列出,並且分析是否具有統計上意義以及臨床上意義嗎?

關於研究結果,你可以把主要結果與次要結果數據列出,包含點估計值以及信賴區間,並且分析是否具有統計上意義以及臨床上意義,證據等級與證據品質?請用表格來整理。縱軸包含主要結果、次要結果,而橫軸包含點估計值 | 信賴區間 |統計上意義 | 原始數據判斷依據|臨床上意義| 原始數據判斷依據| 證據等級 |證據品質。臨床上意義,類別變項請使用NNT去表示,連續變項請使用SMD來表示。如果結果包含判斷內容,請提供判斷邏輯,提供我們可以複查檢視。

結果 類型	點估計值	信賴區間	統計 上意 義	原始數據判斷依據	臨床上意義	原始數據判斷 依據	證據等級	證據品質
主要 結果								
體重 變化	-1.42 kg	-1.70 to -1.14	有	p < 0.00001	SMD: 0.35	根據小幅體重 減輕的效果計 算	中	低(樣本 量小)
BMI 變化	-0.47 kg/m²	-0.63 to -0.31	有	p < 0.00001	SMD: 0.33	根據小幅BMI 改善計算	中	低(樣本 量小)
次要 結果								
腰圍 變化	-1.34 cm	-2.75 to 0.07	無	p = 0.06	無臨床意 義	腰圍變化小且 不顯著	低	低
不良 事件	RR: 1.45	0.83 to 2.54	無	p > 0.05	NNT無 法計算	不良事件未顯 著升高	中	中

CYCH ER EBM

AI輔助

屬性	我們的病人	文獻中的族群	相似程度 (0~10分)
病人 特性	30出頭男性,醫療工作者,久 坐生活型態,超重/肥胖,脂肪 肝	非糖尿病,超重或肥胖患者(年齡範圍18-61歲, 多數為女性),平均基線BMI約36 kg/m²,體重分 布68~105 kg	7
國家/ 種族	台灣,亞洲地區	參與者來自美國、歐洲(丹麥、英國等)及其他 國家,亞洲代表不足	5
介入 治療	SGLT2 抑制劑(如 Dapagliflozin, Canagliflozin)	使用SGLT2抑制劑(Canagliflozin, Dapagliflozin等),劑量包括50 300 mg/day,治療期間多為 2 26週	9
對照 處置	安慰劑或生活方式改變(如飲食控制、運動)	與安慰劑對照,所有受試者均接受標準化的飲食 和運動建議	9
預期 指標	體重下降、BMI減少、脂肪肝改善、血脂改善(IDL和TG)	體重減少(-1.42 kg, MD)、BMI降低(-0.47 kg/m², MD);未直接討論脂肪肝或血脂改善	6

AI輔助

PDA 結構

1. 診斷

• 病情: 小莊, 30 出頭, 肥胖 (BMI > 30), 脂肪肝, 三酸甘油酯 (TG) 過高。

• 目標:體重控制、脂肪肝改善、降低代謝風險。

項目	SGLT2 抑制劑		
優點	1. 可輕微減重(平均體重減少 1.42 kg)		
	 路低代謝相關風險(如心血管風險、血糖控制)。 		
	3. 服用簡便,依從性高。		
缺點	 減重效果有限,需與其他方案結合(如飲食)。 		
	2. 成本較高(每年約 8,400 USD)。		
	 可能出現副作用(如泌尿感染、生殖感染)。 		
相對適合族 群	1. 減重需求不大,但需降低代謝風險的患 者。		
	2. 無法持續執行飲食和運動計劃的患者。		

偏好/價值觀	SGLT2 抑制劑
減重效果	会会会
成本承受能力	全全
副作用容忍度	***
執行可行性	***
長期健康改善重要性	***

• 對小莊的適配建議:

- 若以減重為主要目標,且能自律進行生活方式改變,建議首選生活方式改變。
- 若代謝改善是首要目標,且對減重要求不高,且願意接受藥物副作用和成本,可選擇 SGLT2抑制劑。
- 混合策略:建議將生活方式改變與 SGLT2 抑制劑結合使用,以達到更好的綜合效果。