內科部住院醫師醫學研究教學--以健保資料庫研究為例

內科部研究中心 研究員 許秋婷

聯絡方式

分機:7388、code: 179297

位 置:三期13F血液透析室旁的討論室

from Bedsides to Bed from Bed to Benches

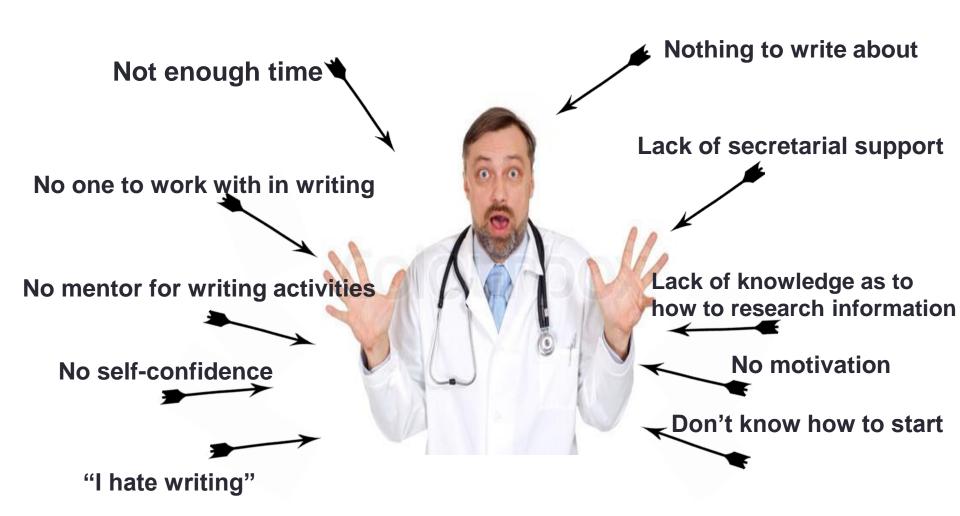








Reasons Given for not Writing



Reasons Why We Write

- 1. Gain intellectual stimulation
- 2. Share ideas
- 3. Report research
- 4. Express an opinion
- 5. Generate discussion
- 6. Advance one's discipline
- 7. Assert "ownership" of a topic
- 8. Attain promotion/tenure
- 9. Report a case
- 10. Enhance one's personal reputation
- 11. Achieve some small measure of immortality by publishing our ideas
- 12. Earn income







醫師為何要執行醫學研究?

老年人可以 吃重鹹嗎? 中風與慢性 腎臟病的關係?

可以幫我開哪個柯P的神奇藥嗎?

自己連呼 吸也會胖?

An apple a day, keep doctor away?



吃A藥比B 藥好嗎?

長期吃安眠藥會不會得癌症?

我都是靠這個乙型阻斷劑…才撐過議員的質詢

地表最強的神藥????

手術使用乙型阻斷劑增加死亡率?

根據發表在JAMA的分析結果,非心臟手術使用乙型阻斷劑增加死亡風險達27%





手術使用乙型阻斷劑增加死亡率?

根據發表在JAMA的分析結果,非心臟手術使用乙型阻斷劑增加死亡風險達27%



JAMA Clinical Evidence Synopsis | May 26, 2015

Perioperative Use of β-Blockers in Cardiac and Noncardiac Surgery

Hermann Blessberger, MD1; Juergen Kammler, MD1; Clemens Steinwender, MD1

[+] Author Affiliations

我者

JAMA. 2015;313(20):2070-2071. doi:10.1001/jama.2015.1883.

Text Size: A A A

ABSTRACT

Article Figures References

ABSTRACT | SUMMARY OF FINDINGS | DISCUSSION | ARTICLE INFORMATION | REFERENCES

Clinical Question Are β -blockers associated with lower rates of mortality and morbidity after cardiac or noncardiac surgery?

Bottom Line In cardiac surgery, β -blockers are associated with a lower incidence of supraventricular tachycardias (SVTs) and ventricular arrhythmias. In noncardiac surgery, β -blockers are associated with a possible increase in mortality and strokes, a lower incidence of acute myocardial infarctions (AMIs) and SVTs, and an increase in bradycardia and hypotension. If tolerated, long-term β -blocker treatment should be continued perioperatively, whereas the decision to start a β -blocker should be individualized, weighing risks and benefits.

非心臟手術使用乙型阻斷劑可能增加死亡率~~~ 研究結果顯示:

- (1) 死亡率顯著增加27% (RR 1.27, 1.01-1.59);
- (2) 腦血管事件顯著增加109% (RR 2.09, 1.14-3.82);



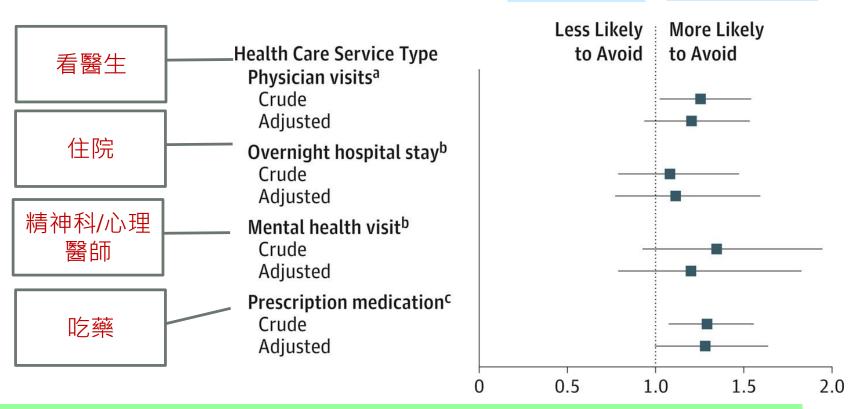
From: Association Between Apple Consumption and Physician Visits: Appealing the Conventional Wisdom That an Apple a Day Keeps the Doctor Away

JAMA Intern Med. 2015;175(5):777-783. doi:10.1001/jamainternmed.2014.5466

一天一顆蘋果,真的就不用看醫生了嗎?

吃了變差

吃了比較好



研究結果顯示: 每天吃蘋果,沒有辦法降低看醫生、住院或吃

藥的風險

pple

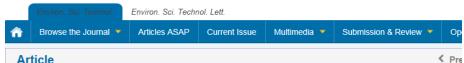
include covariates for age, sex, race or ethnicity educational attainment, body mass index, smoking status, and health insurance type.

Date of download: 8/9/2015

胖子的悲哀--連呼吸都會胖

Environmental Science & Technology





Activation of Human Peroxisome Proliferator-Activated Nuclear Receptors (PPARγ1) by Semi-Volatile Compounds (SVOCs) and Chemical Mixtures in Indoor Dust

Mingliang Fang †, Thomas F. Webster ‡, and Heather M. Stapleton *†

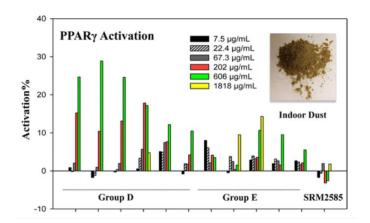
- † Nicholas School of the Environment, Duke University, Durham, North Carolina 27708, United States
- Department of Environmental Health, Boston University School of Public Health, Boston, Massachusetts 02118, United States

Environ. Sci. Technol., Article ASAP DOI: 10.1021/acs.est.5b01523

Publication Date (Web): July 14, 2015

Copyright © 2015 American Chemical Society

*Phone: 919-613-8717, Fax: 919-684-8741, E-mail: heather.stapleton@duke.edu.



吸空氣會胖是真的!美研究:灰塵中的化學物質使人發胖

NOWnews - 2015年8月2日 下午1:51



國際中心/綜合報導

不少長年飽受肥胖所苦的人會宣稱「自己 連呼吸都會胖」,不過這可不是他們在誇 大其詞,近日國外有項研究發現,這是真 的!

美國北卡羅萊納州的杜克大學研究團隊發 現,灰塵中含有會使人發胖的化學物質,

影響人體的脂肪代謝率、細胞增長與死亡等功能。孩童若在發育期間吸入過量的灰塵,未 來將導致肥胖。

灰塵中會使人發胖的化學物質,也存在於塑膠、潤滑油和阻燃劑之中,它與促使人發胖的 PPARgamma受體相同,有左右肥胖細胞的作用。

研究人員根據孩童平時會接觸到的灰塵量,在家中、辦公室和健身房收集了25個灰塵樣本,結果他們發現,一半以上都有PPARgamma活躍的跡象。據美國環保署統計,孩童每天大約會接觸到家中50毫克的灰塵。

From: Dietary Sodium Content, Mortality, and Risk for Cardiovascular Events in Older Adults: The Health, Aging, and Body Composition (Health ABC) Study

JAMA Intern Med. 2015;175(3):410-419. doi:10.1001/jamainternmed.2014.6278

攝取過多鹽類,會增加高血壓和心血管疾病風險?

- 一項前瞻性世代研究結果,這項研究針對高齡長者 (年齡71~80歲) (n=2,642),以飲食問卷的方式調查飲食鈉含量,結果發現:
- (1) 飲食鈉含量這檔事,與死亡風險並沒有顯著 關係 (HR 1.03 per 1 g, 95% CI 0.98-1.09, p=0.27)
- (2) 校正可能的影響因子後,攝取過少與過多的 鈉,並沒有顯著增加死亡風險

研究顯示吃重鹹並未增加死亡率

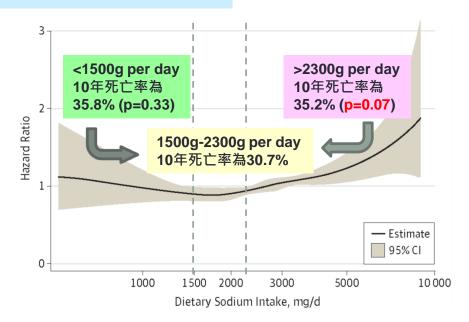


Figure Legend:

Restricted Cubic Spline Model of Dietary Sodium Intake as a Univariate Predictor of Mortality. The cubic spline model improved the likelihood ratio χ^2 over the linear model (from χ^2_1 = 10.71 to χ^2_4 = 12.33), but the gain in fit did not justify the increased model complexity. The Bayesian information criterion, which penalizes for unnecessary complexity, indicated that the linear model is preferable.

Hypertension **Hypertension**

PP.11.23: ASSOCIATION BETWEEN SALT INTAKE AND INDICES OF MICRO- AND MACROVASCULAR FUNCTION.

Triantafyllou, A.; Anyfanti, P.; Zabulis, X.; Triantafyllou, G.; Karamaounas, P.; Gkolias, V.; Gkaliagkousi, E.; Douma, S.

Abstract

Objective: Multiple studies have linked sodium intake with office blood pressure (BP) levels and increased cardiovascular mortality and morbidity. However, the association between increased salt consumption and 24-hour ambulatory BP monitoring (ABPM) parameters remains underinvestigated. Moreover, it still remains ambiguous whether it promotes subclinical micro- and macrovascular damage independent of BP levels and simultaneously in different target organs.

Design and method: Consecutive newly diagnosed, otherwise healthy, never-treated hypertensive patients and healthy volunteers underwent ABPM and blood sampling. Sodium intake was estimated in 24-hour urine samples. Structural alterations of dermal capillaries (capillary density per visual field) were evaluated using special software analysis of nailfold capillaroscopy images. Functional microvascular alterations of the kidney were assessed by estimation of microalbuminuria. The Sphygmocor device was used to assess arterial stiffness by measurement of pulse wave velocity (PWV), and Aortic augmentation Index (Alx).

Results: The study included 193 participants. Sodium excretion was significantly associated with ABPM (r = 0.216, p < 0.01), day-time systolic blood pressure (SBP) (r = 0.207, p < 0.05), and night-time SBP (r = 0.213, p < 0.01). The association between sodium excretion and 24-hour SBP (r = 0.170, p < 0.05), as well as night-time SBP (r = 0.173, p < 0.05) remained significant even after adjustment for other parameters. Although sodium excretion was not associated with PWV or capillary density, a significant correlation was found between sodium excretion and Alx, as well as microalbuminuria. In the multiple linear regression model, 24-hour SBP (p = 0.008), albuminuria (p = 0.019) and Alx (p = 0.042) remained significant predictors of sodium excretion, even after adjustment for age, BMI, office BP, smoking, glomerular filtration rate (GFR), capillary rarefaction and aldosterone levels.

Conclusions: This is the first study demonstrating a significant association between salt intake and indices of functional microvascular (microalbuminuria) and macrovascular (Alx) involvement in a population free from the long-standing effects of essential hypertension and after accounting for several factors including aldosterone. In addition, it was shown that of all 24-hour ABPM parameters, 24-hour and night-time SBP exhibit the most powerful association with salt consumption. Our findings highlight the detrimental effects of excessive dietary salt intake and should serve as a reminder to promote lifestyle changes in hypertensive patients.



Table of Contents

Vol. 40, No. 1, 2015

Issue release date: August 2015

Section title: Original Paper

Their Relationship to Overhydration in Chronic Kidney Disease Patients

Hallvass A.E.C.^a · Claro L.M.^a · Gonçalves S.^a · Olandoski M.^a · Nerbass F.B.^{a, b} · Aita C.A.M.^a · de Moraes T.P.^a · Pecoits-Filho R.^a

^aSchool of Medicine, Pontifícia Universidade Católica do Paraná, Curitiba, and ^bNutrition Department, Pro-Rim Foundation, Joinvile, Santa Catarina, Brazil

Abstract Purchase FullText PDF Login / Register

Abstract

The purpose of this study was to estimate sodium intake in a group of patients with chronic kidney disease (CKD) and to correlate the results with the urinary excretion values of sodium and signs of fluid overload. We included patients with CKD in different stages. Urinary sodium was measured in 24 h urine samples. Body composition monitor (BCM) was used to estimate the hydration status. Sixty patients $(38 \pm 15 \text{ ml/min of GFR})$ presented $4.14 \pm 1.71 \text{ g/24}$ h of urinary sodium excretion. Overhydration was detected in 50% of the patients by the BCM. There was a positive correlation between the measured sodium excretion values and BCM, ICW, ECW and TBW. In conclusion, markers of overhydration evaluated by BCM were positively correlated with urinary sodium excretion.

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From Bed to Benches From Wet-lab to Dry-lab

健保資料庫之合作平台

内科部研究中心 提出申請 彰化師範大學 中國醫藥大學

健保研究資料庫的好, JAMA都知道:

JAMA Internal Medicine

Formerly Archives of Internal Medicine

Invited Commentary

Invited Commentary

Nationwide Population Science Lessons From the Taiwan National Health Insurance Research Database

Ann W. Hsing, PhD; John P. A. Ioannidis, MD, DSc

Nationwide Population Science: Lessons From Taiwan

非資料庫研究: Find data bases to fit a research question

資料庫研究: Find research questions to fit existing data sets

事起頭難 Turning idea into a research Question (Clear research aim, rationale, hypothesis)

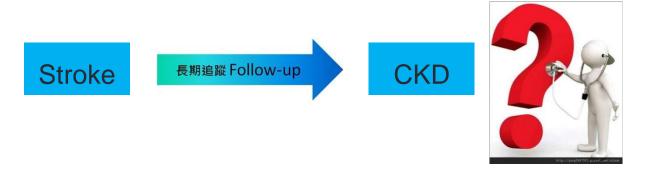
分享第一篇健保資料庫文章的構思過程

- 探討 Stroke 和 Chronic kidney disease 相關性研究



定論: CKD as a Risk Factor for Stroke

Stroke and CKD share similar cardiometabolic (心臟代謝) risk factors.





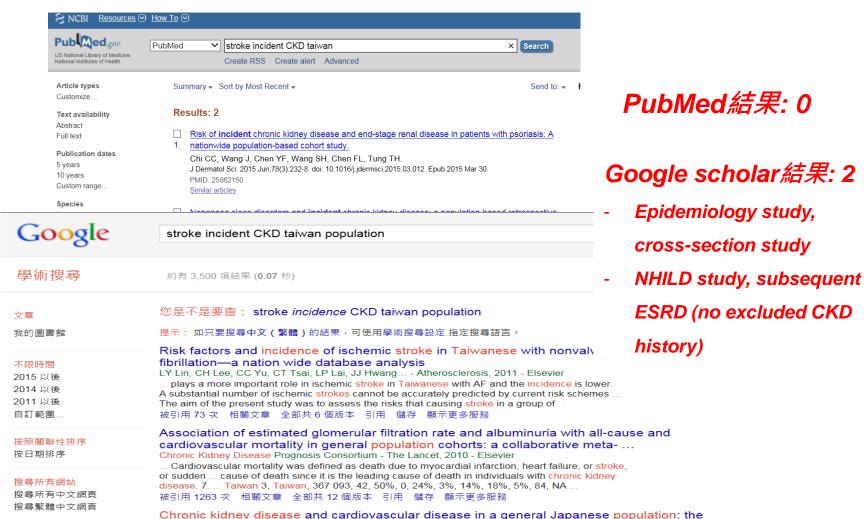
出發總是要有方向一文獻查詢

Start-Up

Systematic review of existing evidence

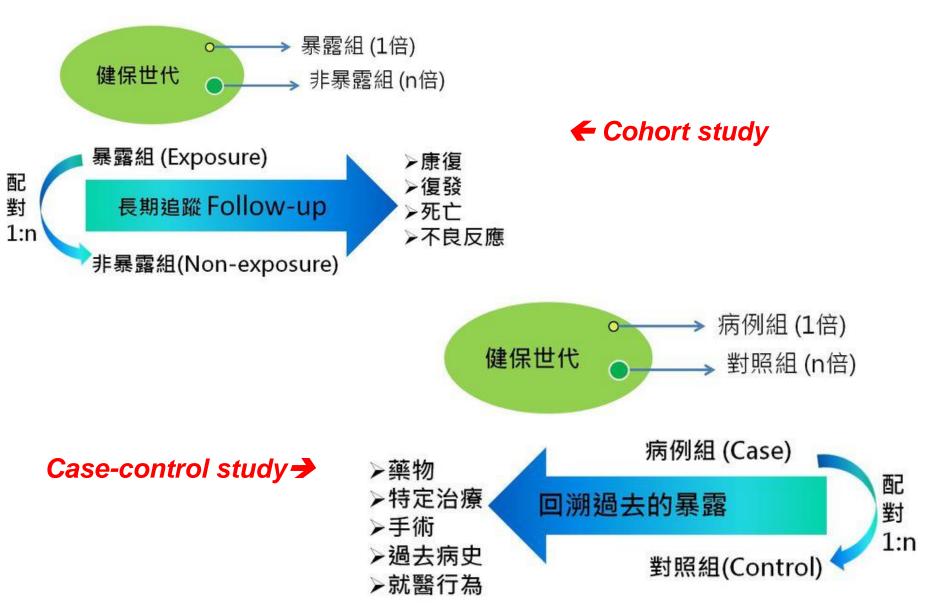
Searched by **PubMed** & **Google scholar** ...

- 先確認 是否有stroke 和 CKD 相關流行病學文章或其他健保資料庫文章



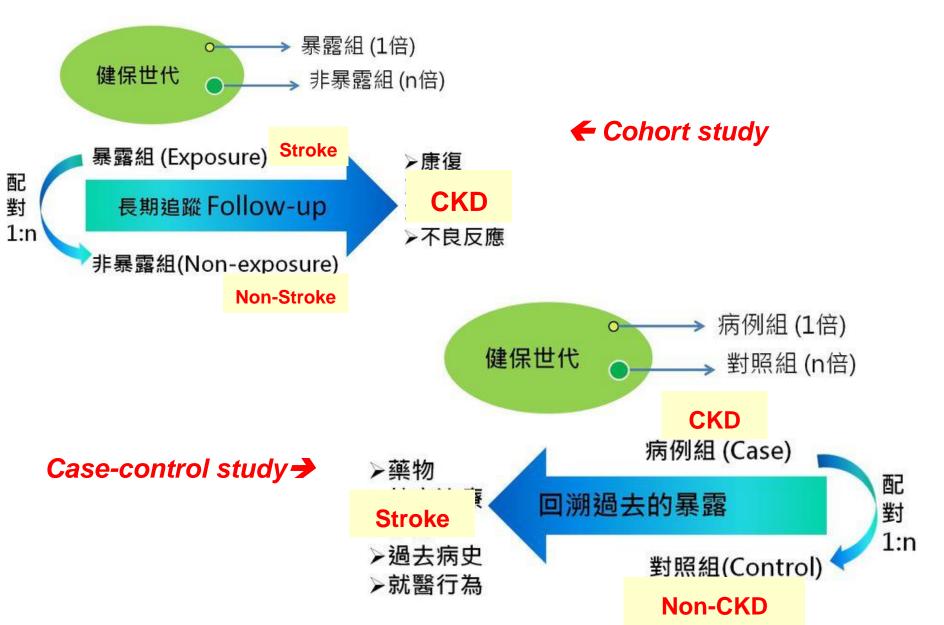
How to do?

Develop appropriate design to answer question



How to do?

Develop appropriate design to answer question





Qualitative Data

研究對象: Study population

- (1) 資料庫:
- (2) 納入條件: (疾病需註明ICD-9 or藥物 ATC碼or 處置碼) (頻率定義)
- (3) 排除條件 Exclusion:(疾病需註明 ICD-9 or藥物ATC碼or 處置碼) (頻率定義)
- (4) 自變項或Exposure (X):
 - I. Exposure:

II. Non-Exposure:

備註:

配對控制變數 Matched case-control: (1:m→ m=) (Ex: Age, gender, year of diagnosis....)

備註:

始____年~__年止

統計控制變數Confounder:

Primary Outcome (Y₁):

Secondary Outcome (Y₂):

Third Outcome (Y₃):

(疾病需註明ICD-9 or藥物ATC 碼or處置碼) (頻率定義)

備註:

End-point criteria:



Qualitative Data





Qualitative Data

研究對象: Study population

- (1) 資料庫:健保資料庫
- (2) 納入條件: (疾病需註明ICD-9 or藥物 ATC碼or 處置碼) (頻率定義) All population
- (3) 排除條件 Exclusion:(疾病需註明 ICD-9 or藥物ATC碼or 處置碼) (頻率定義)
- (4) 自變項或Exposure (X):
 - I. Exposure:

Stoke:

ischemic (ICD-9-CM 433-

438) hemorrhagic (ICD-9-

CM 430-432)

II. Non-Exposure:

Non-stoke population

備註:

配對控制變數 Matched case-control: (1:m→ m=) (Ex: Age, gender, year of diagnosis....) 備註:

始____年_年止

統計控制變數Confounder:

Primary Outcome (Y₁):

CKD (ICD-9-CM 580-589)

Secondary Outcome (Y₂):

ESRD (ICD-9-CM 585 585.1)

Third Outcome (Y₃):

Death (ID檔有註明死亡,低

估)

Censored: (退保日期)

(疾病需註明ICD-9 or藥物ATC 碼or 處置碼) (頻率定義)

備註:

End-point criteria:



Qualitative Data

研究對象: Study population

- (1) 資料庫:健保資料庫
- (2) 納入條件: (疾病需註明ICD-9 or藥物 ATC碼or 處置碼) (頻率定義) All population
- (3) 排除條件 Exclusion:(疾病需註明 ICD-9 or藥物ATC碼or 處置碼) (頻率定義)
 - history of CKD (ICD-9-CM codes 580-589) and ESRD (ICD-9-CM code 585) before the index date
 - aged less than 18 years
- (4) 自變項或Exposure (X):
 - I. Exposure:

Stoke:

ischemic (ICD-9-CM 433-438) hemorrhagic (ICD-9-CM 430-432)

II. Non-Exposure:

Non-stoke population

備註:

配對控制變數 Matched case-control: (1:m→ m=) (Ex: Age, gender, year of diagnosis....) 備註:

始____年_年止

統計控制變數Confounder:

Primary Outcome (Y₁):

CKD (ICD-9-CM 580-589)

Secondary Outcome (Y₂):

ESRD (ICD-9-CM 585 585.1+EPO) EPO(ATC:)

Third Outcome (Y₃):

Death (ID檔有註明死亡,低估)

Censored: (退保日期)

(疾病需註明ICD-9 or藥物ATC 碼or處置碼) (頻率定義)

備註:

健保局首頁 > 「藥材專區」 > 「藥品」 > 「健保用藥品項」 > 「健保用藥品項104年08月壓縮總檔」

連結以下網址

(http://www.nhi.gov.tw/webdata/webdata.aspx? menu=21&menu_id=713&WD_ID=849&webda ta_id=1139) °



11. 健保用藥且適用「罕見疾病防治及藥物法」之品項檔(104.06.22更新) 🖾 🙆



Qualitative Data

研究對象: Study population

- (1) 資料庫:健保資料庫
- (2) 納入條件: (疾病需註明ICD-9 or藥物 ATC碼or 處置碼) (頻率定義) All population
- (3) 排除條件 Exclusion:(疾病需註明 ICD-9 or藥物ATC碼or 處置碼) (頻率定義)
 - history of CKD (ICD-9-CM codes 580-589) before the index date
 - aged less than 18 years
- (4) 自變項或Exposure (X):
 - I. Exposure:

Stoke:

ischemic (ICD-9-CM 433-438) hemorrhagic (ICD-9-CM 432-432)

CM 430-432)

II. Non-Exposure:Non-stoke population

備註:

配對控制變數 Matched case-control: (1:m→ m=4) (Ex: Age, gender, year of diagnosis....)
Age, gender, year of index

備註:

始____年_年止

統計控制變數Confounder:

Primary Outcome (Y₁):

CKD (ICD-9-CM 580-589)

Secondary Outcome (Y₂): advanced CKD (ICD-9-CM

585 585.1)+EPO (ATC:) Third Outcome (Y₃):

Death (ID檔有註明死亡,低估)

Censored: (退保日期)

(疾病需註明ICD-9 or藥物ATC 碼or處置碼) (頻率定義)

備註:

End-point criteria:



Qualitative Data

研究對象: Study population

- (1) 資料庫:健保資料庫
- (2) 納入條件: (疾病需註明ICD-9 or藥物 ATC碼or 處置碼) (頻率定義) All population
- (3) 排除條件 Exclusion:(疾病需註明 ICD-9 or藥物ATC碼or 處置碼) (頻率定義)
 - history of CKD (ICD-9-CM codes 580-589) before the index date
 - aged less than 18 years
- (4) 自變項或Exposure (X):
 - I. Exposure:

Stoke:

ischemic (ICD-9-CM 433-438) hemorrhagic (ICD-9-CM 430-432)

II. Non-Exposure:Non-stoke population

備註:

配對控制變數 Matched case-control: (1:m→ m=4) (Ex: Age, gender, year of diagnosis....)
Age, gender, year of index
備註:

始_1996__年~_2012_年止

統計控制變數Confounder:

- 1. Income
- 2. Urbanization
- 3. Hypertension (ICD-9-CM code 401 to 405)
- 4. diabetes mellitus (ICD-9-CM code 250)
- 5. hyperlipidemia (ICD-9-CM code 272)
- 6. endocarditis (ICD-9-CM codes 063-42, 074-22, 039-20-039-24, 098-84, 112-81, 115-04, 115-14, 115-94, 421-0-421-9, 424-9)
- 7. atrial fibrillation (AF, ICD-9-CM code 427-31)
- 8. ischemic heart disease (CAD)(ICD-9-CM codes 410-414)
- 9. congestive heart failure (CHF; ICD- 9-CM code 428)
- 10. peripheral artery occlusive disease (PAOD; ICD-9-CM codes 443–444).

Primary Outcome (Y₁):

CKD (ICD-9-CM 580-589)

Secondary Outcome (Y₂): advanced CKD (ICD-9-CM 585 585.1)+EPO (ATC:)

Third Outcome (Y₃):

Death (ID檔有註明死亡,低估)

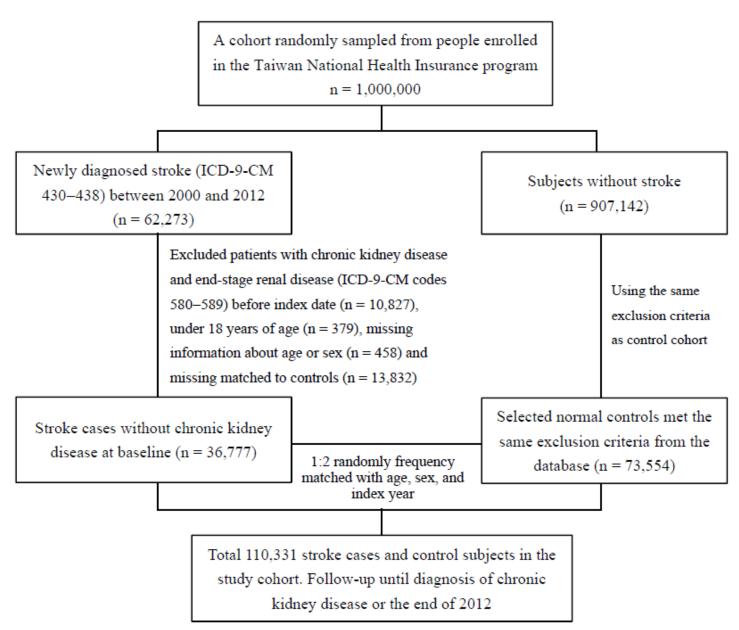
Censored: (退保日期)

(疾病需註明ICD-9 or藥物ATC 碼or處置碼)(頻率定義) 借註:

 Death is a competing risk of Y₁, Y₂

End-point criteria:

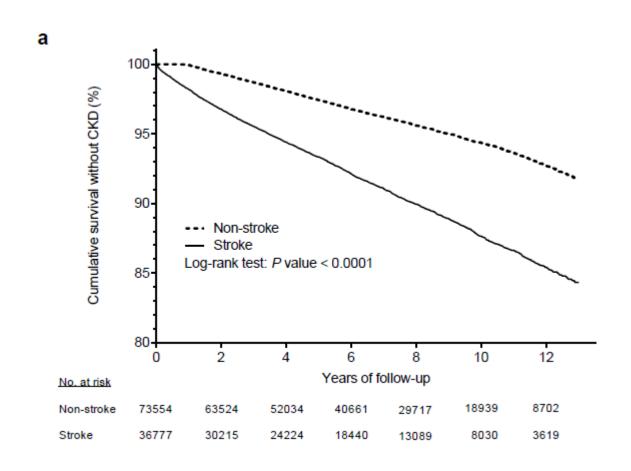
Study flow-chart



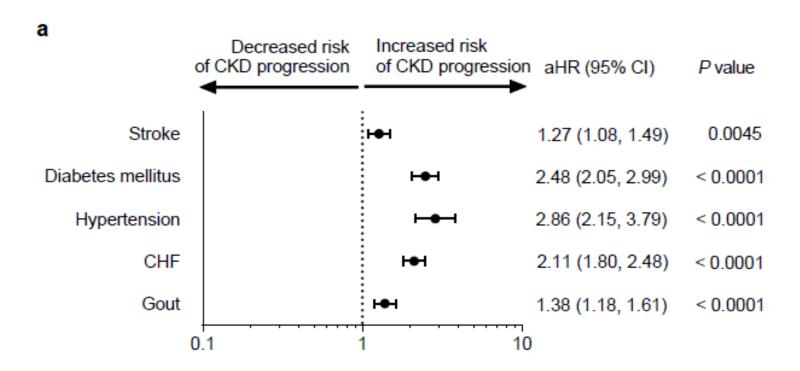
Frequency match

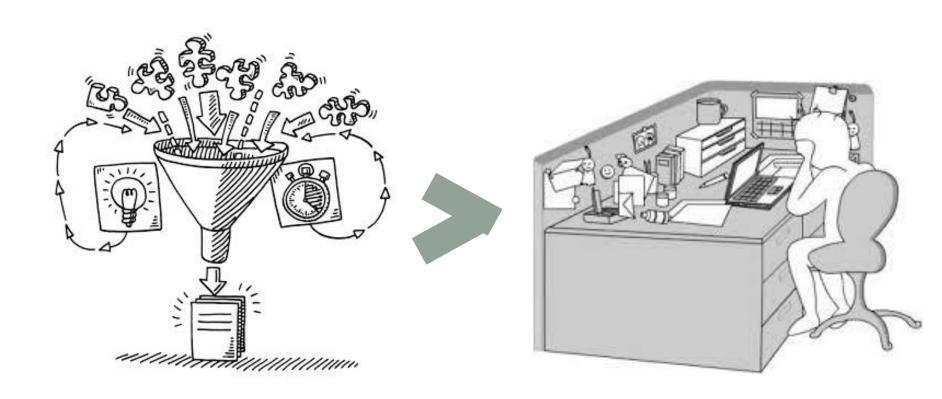
	Non-stroke	Stroke	Total cohort	P value*
Number of patients	73,554	36,777	110,331	
Gender, n (%)				
Female	36,060 (49.03%)	18,030 (49.03%)	54,090 (49.03%)	1.000
Male	37,494 (50.97%)	18,747 (50.97%)	56,241 (50.97%)	
Age, mean \pm SD, years Age stratified, n (%)	58.96 ± 12.32	59.16 ± 12.28	59.03 ± 12.31	0.011
< 50	15,312 (20.82%)	7,656 (20.82%)	22,968 (20.82%)	1.000
50 - 64	34,062 (46.31%)	17,031 (46.31%)	51,093 (46.31%)	
≥ 65	24,180 (32.87%)	12,090 (32.87%)	36,270 (32.87%)	

Kaplan-Meier curve



Multivariate analysis





Thanks for your attention!!!!