



Lo mejor del año en 10 minutos

Dr. Javier Sobrino Martínez
Unidad de HTA



Madrid 24 de abril de 2015



FUNDACIÓ HOSPITAL DE
l'Esperit Sant

Arterial Hypertension

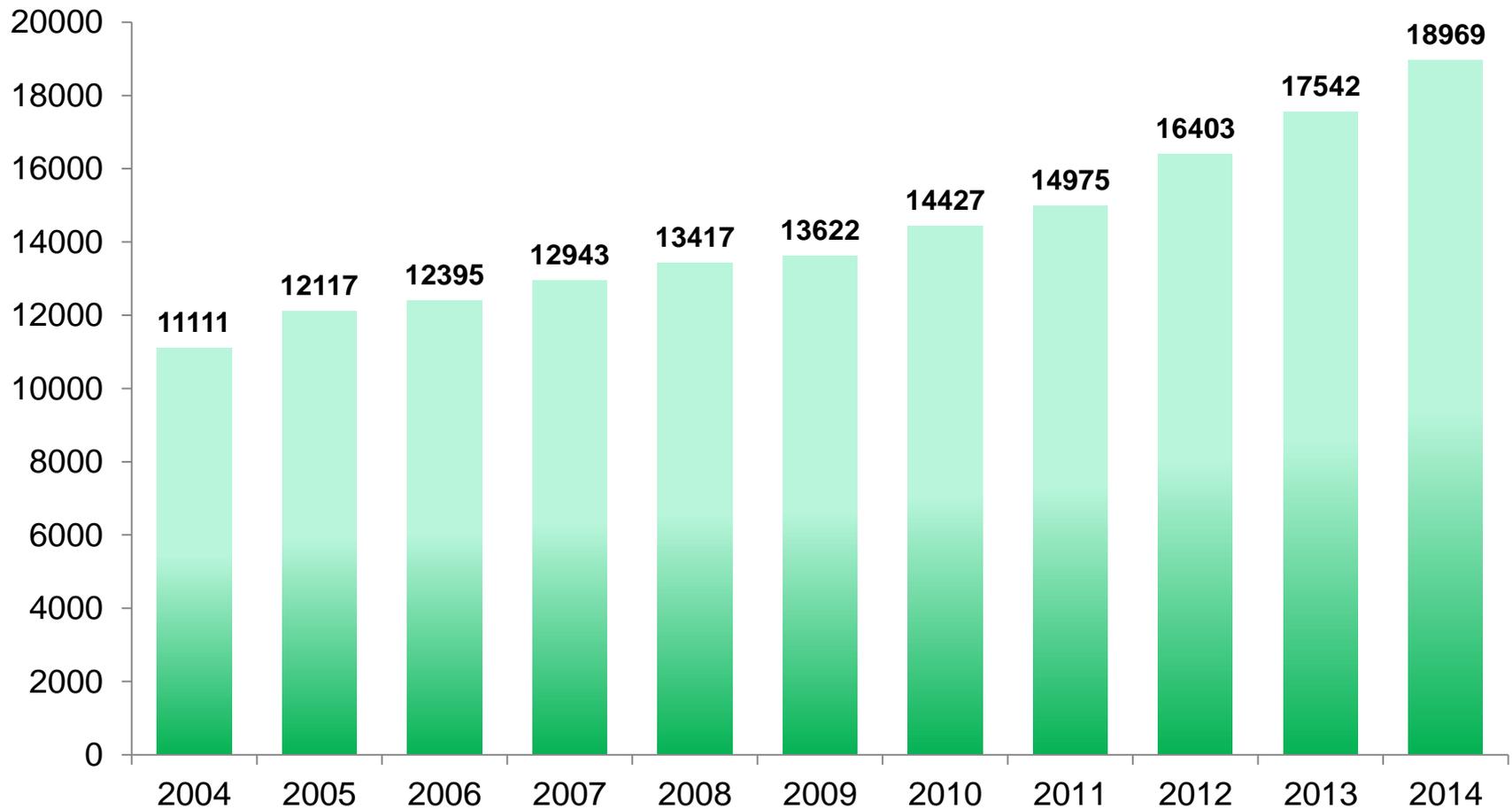




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TRENDS
Most Viewed (Last Week)

IMAGES IN CLINICAL MEDICINE
Incisional Keloid [38,880 views]
April 9, 2015 | I.V. Salas and A.F. Mirabell

ORIGINAL ARTICLE
Free Full Text | Video | Comments

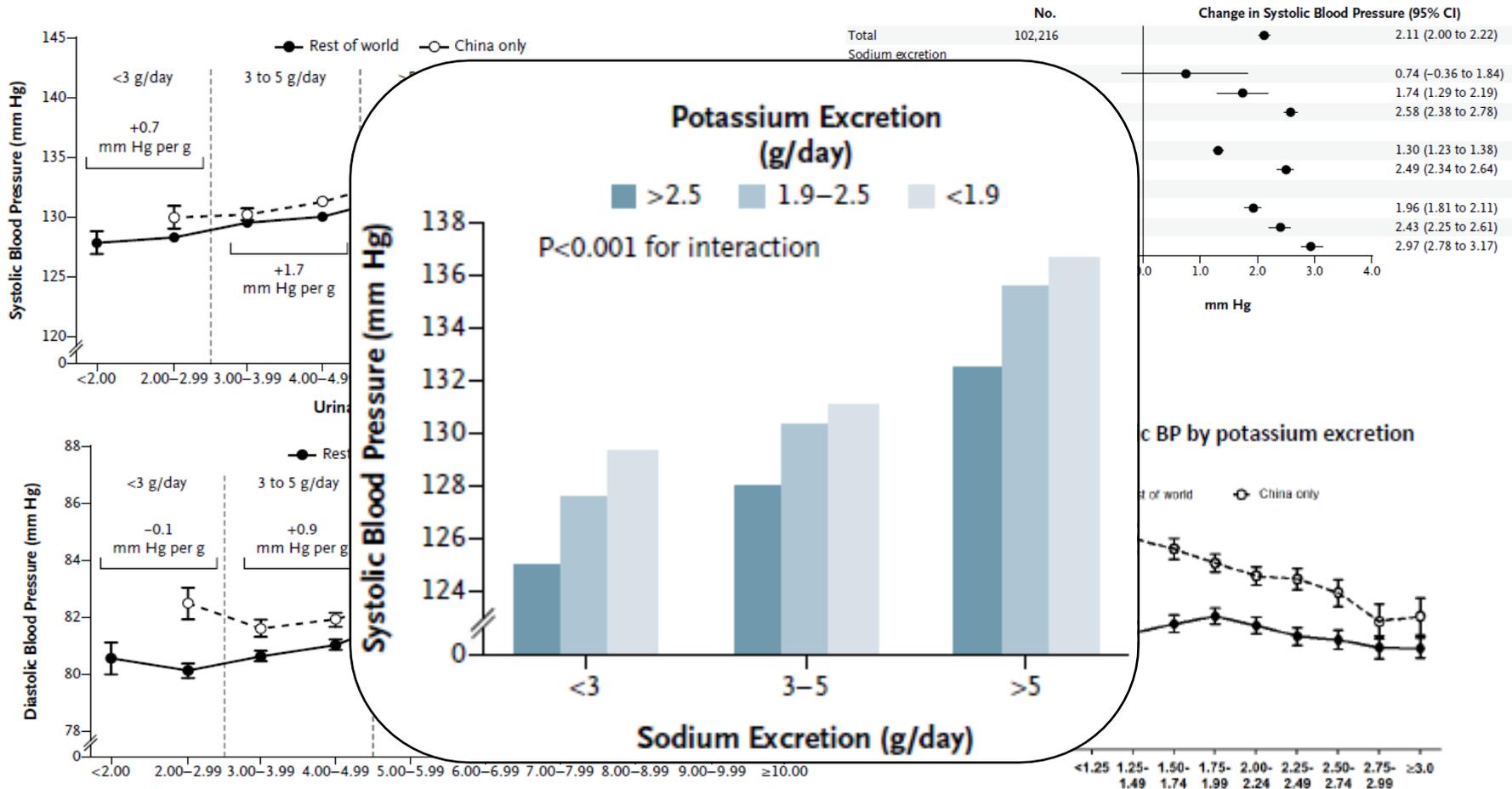
[VIEW THE DISCUSSIONS >](#)

Mente A. N Engl J Med. 2014; 371: 601-11

Relación de la ingesta de sodio y potasio con la presión arterial, la mortalidad total y la morbilidad cardiovascular

- Estudio PURE (*Prospective Urban Rural Epidemiology*)
- 102.000 individuos (18-70 años. Edad media 51 años) de 18 países de los 5 continentes abarcando todas las condiciones sociales y económicas.
- La EU de Na y K en orina matinal
- El cálculo estimado en 24 h Fórmula de Kawasaki.
- EU de 1 g de Na \approx Ingesta de 2,5 g de sal

EU media de Na 4,93 g/día; de K 2,12 g/d
44 % EU Na > 5 g /d (12,5 g de sal).
46 % EU Na entre 3-5 g/d.



	No.
Total Sodium excretion	102,216

Change in Systolic Blood Pressure (95% CI)

Change in BP by potassium excretion

Potassium Excretion (g/day)

■ >2.5 ■ 1.9-2.5 ■ <1.9

P<0.001 for interaction

Systolic Blood Pressure (mm Hg)

Sodium Excretion (g/day)

Systolic Blood Pressure (mm Hg)

Diastolic Blood Pressure (mm Hg)

Urinary

● Rest of world ○ China only

● Rest of world ○ China only

<3 g/day 3 to 5 g/day

<3 g/day 3 to 5 g/day

<2.00 2.00-2.99 3.00-3.99 4.00-4.99

<2.00 2.00-2.99 3.00-3.99 4.00-4.99 5.00-5.99 6.00-6.99 7.00-7.99 8.00-8.99 9.00-9.99 ≥10.00

<1.25 1.25-1.49 1.50-1.74 1.75-1.99 2.00-2.24 2.25-2.49 2.50-2.74 2.75-2.99 ≥3.00

Media de seguimiento 3,7 años

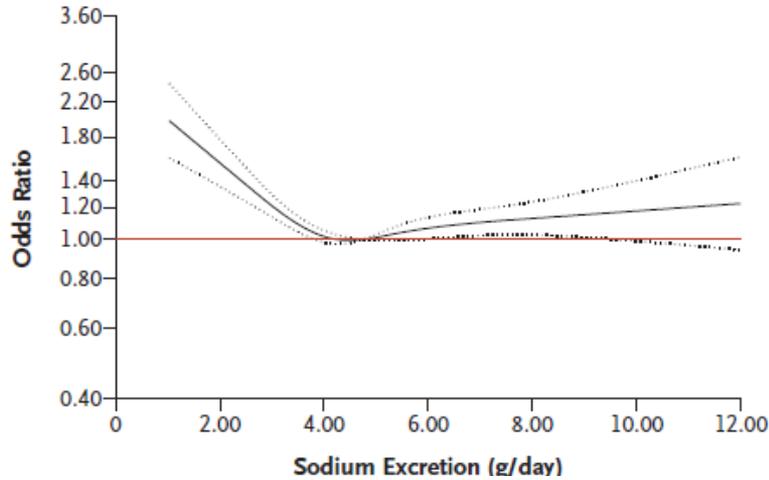
Variable compuesta:

Mortalidad + Eventos CV: 3.317 sujetos (3,3%)

Muertes: 1.976

O'Donnell M. *N Engl J Med.* 2014; 371: 612-23

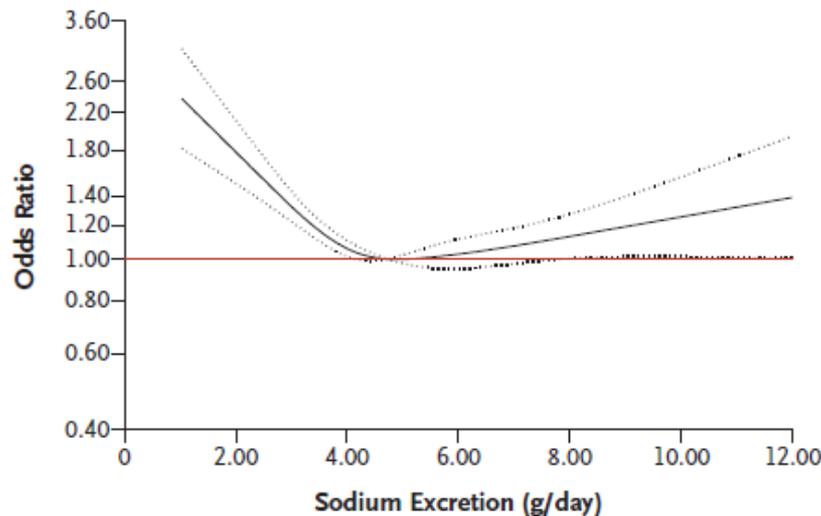
Estimated Sodium Excretion and Risk of Death or Cardiovascular Events



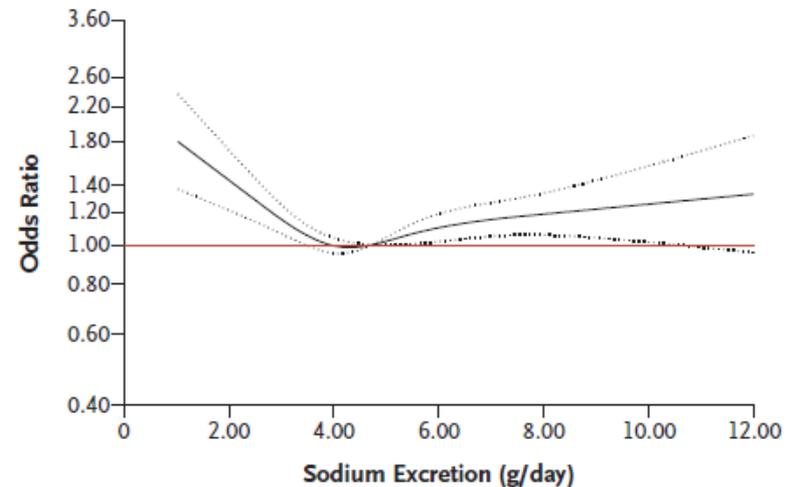
Referencia 4-6 g/dia

- > 7 gr/dia: OR 1,15 (1,02–1,30)
- < 3 gr/dia OR 1,27 (1,12–1,44)

Estimated Sodium Excretion and Risk of Death from Any Cause



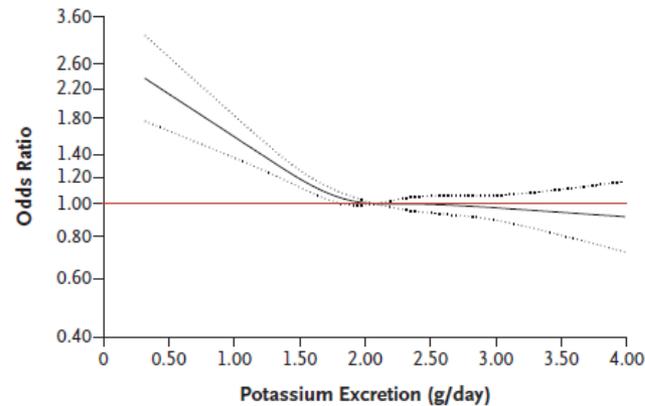
Estimated Sodium Excretion and Risk of Major Cardiovascular Events



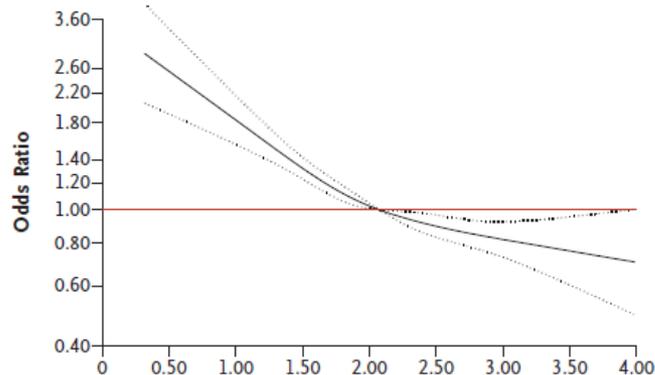
Asociación entre EU de K con Mortalidad y Eventos Cardiovasculares

	<1.50 g/d	1.50–1.99 g/d	2.00–2.49 g/d	2.50–3.00 g/d	>3.00 g/d
OR	1.00	0.86 (0.77–0.97)	0.81 (0.73–0.91)	0.86 (0.75–0.98)	0.78 (0.67–0.91)

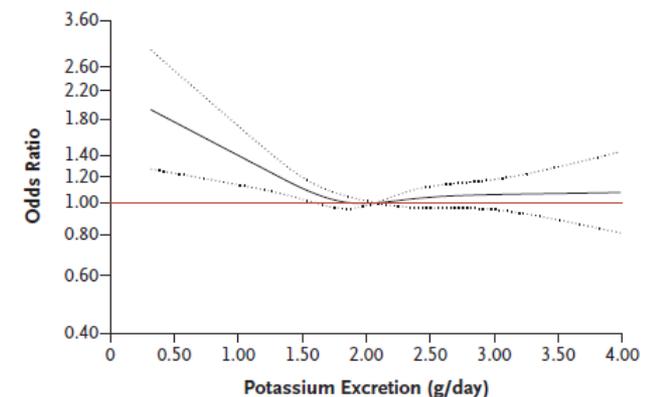
Estimated Potassium Excretion and Risk of Death or Cardiovascular Events



Estimated Potassium Excretion and Risk of Death from Any Cause

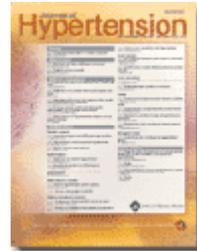


Estimated Potassium Excretion and Risk of Major Cardiovascular Events



Prognostic impact from clinic, daytime, and nighttime systolic blood pressure in nine cohorts of 13 844 patients with hypertension

The ABC-H Investigators.
Journal of Hypertension 2014,
32:2332-40



Metaanálisis de 9 estudios de cohortes de pacientes hipertensos seguidos como mínimo 1 año
13.844 pacientes de Europa, Brasil y Japon

Hazard ratio (IC 95%) para ECV por cada incremento de 10 mmHg de PAS

	Total eventos CV	Cardiopatía isquémica	Ictus
Sin ajustar			
PAS nocturna	1,25 (1,22-1,29)	1,13 (1,05-1,22)	1,29 (1,19-1,39)
PAS diurna	1,20 (1,15-1,26)	1,08 (0,99-1,18)	1,29 (1,20-1,38)
PAS clínica	1,11 (1,06-1,16)	1,13 (0,95-1,34)	1,13 (1,06-1,21)
Con ajuste			
PAS nocturna	1,26 (1,20-1,31)	1,22 (1,13-1,31)	1,26 (1,09-1,46)
PAS diurna	1,01 (0,94-1,08)	0,97 (0,88-1,07)	1,04 (0,92-1,17)
PAS clínica	1,00 (0,95-1,05)	1,01 (0,93-1,09)	1,00 (0,97-1,03)





The importance of night-time systolic blood pressure in diabetic patients: Dublin Outcome Study

Draman M.S.

J Hypertens. 2015 Apr 16. [Epub ahead of print]

Mohd S. Draman^a, Eamon Dolan^c, Lelane van der Poel^c, Tommy Kyaw Tun^b, John H. McDermott^b, Seamus Sreenan^b, and Eoin O'Brien^d

- 11.291 pacientes sin medicación antihipertensiva
- 859 diabéticos
- Seguimiento medio 5,3 años
- 74 muertes

	Diabéticos	No Diabéticos	p
PAS dia (mmHg)	146,4	145,1	n.s.
PAS noche (mmHg)	131,2	126,4	<0,0001
% Non-dipper	47,4	35,5	<0,0001

TABLE 2. Hazard ratios for a 10-mmHg increase in night-time and daytime SBP for overall and cause-specific cardiovascular mortality in diabetic patients

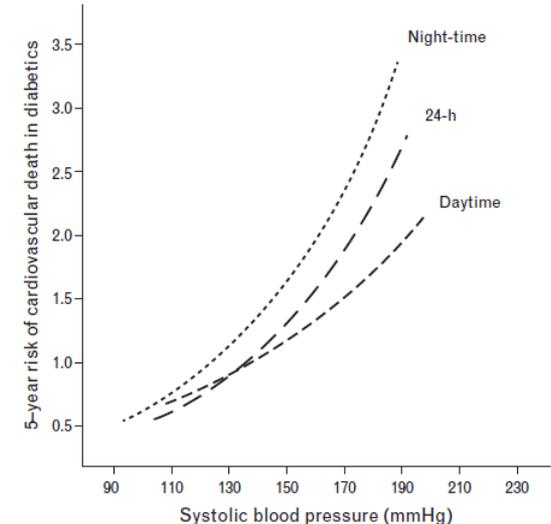
	Cardiovascular	Stroke	Cardiac
Number of events	74	27	47
Daytime	1.10 (1.03–1.17)*	1.12 (1.00–1.25)	1.10 (1.02–1.19)*
Night time	1.25 (1.11–1.39)*	1.38 (1.13–1.58)*	1.16 (1.02–1.25)*
Daytime ^a	0.99 (0.96–1.17)	0.97 (0.91–1.14)	0.99 (0.96–1.24)
Night time ^a	1.32 (1.12–1.69)*	1.95 (1.18–3.27)*	1.24 (0.99–1.56)

Hazard rates (95% confidence intervals) for each 10-mmHg increase in systolic pressure with adjustments applied for baseline characteristics including sex, age, BMI, history of cardiovascular events and smoking status.

^aDaytime further adjusted for night-time blood pressure and vice versa.

* $P < 0.05$.

Riesgo Mortalidad en diabeticos non-dipper comparados con dipper
HR 2,11, (IC 95% 1,11–4,01; $p < 0.05$).





Isolated Systolic Hypertension in Young and Middle-Aged Adults and 31-Year Risk for Cardiovascular Mortality



The Chicago Heart Association Detection Project in Industry Study

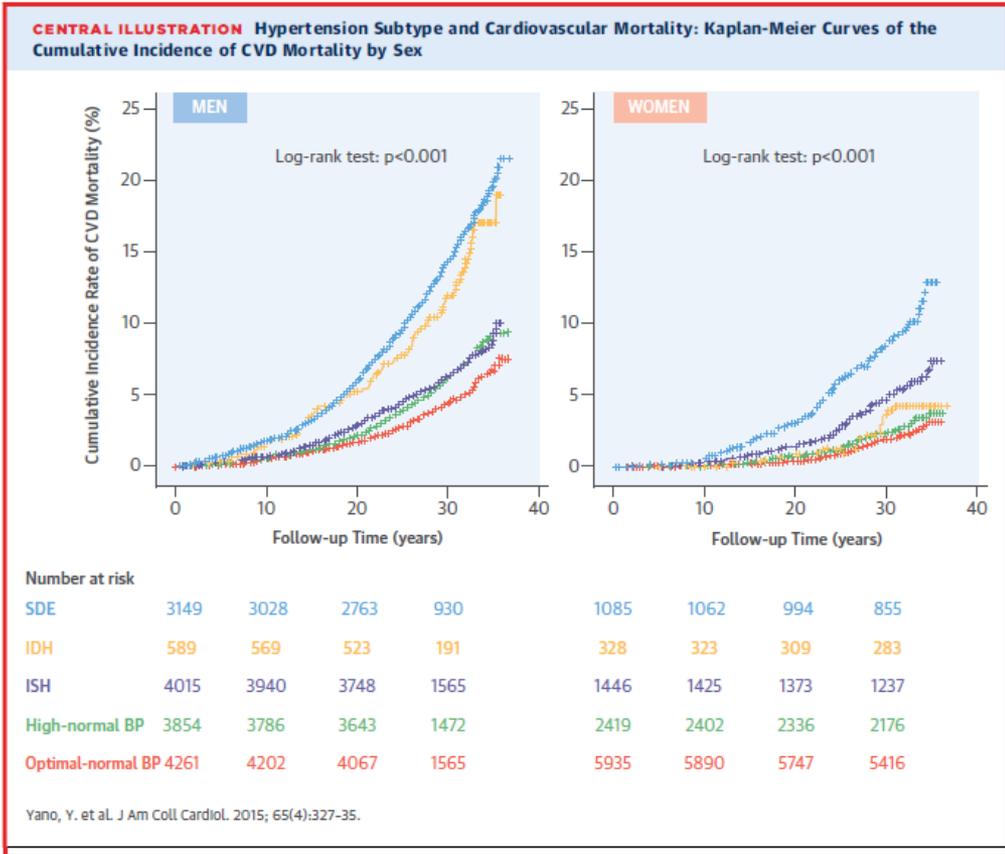
Yuichiro Yano, MD, PhD,* Jeremiah Stamler, MD,* Daniel B. Garside, MA,*† Martha L. Daviglus, MD, PhD,*† Stanley S. Franklin, MD,‡ Mercedes R. Carnethon, PhD,* Kiang Liu, PhD,* Philip Greenland, MD,* Donald M. Lloyd-Jones, MD, ScM*

27.081 sujetos entre 18 y 49 años, sin ECV previa, ni tto antihipertensivo
 Seguimiento medio 31,1 años

TABLE 4 Sex-Specific Unadjusted and Multivariate-Adjusted HR (95% CIs) for BP as a Continuous Variable for Risk for CVD Mortality, CHD Mortality, and Stroke Mortality

Model	Men (n = 15,868)		Women (n = 11,213)	
	SBP, 10 mm Hg	DBP, 5 mm Hg	SBP, 10 mm Hg	DBP, 5 mm Hg
CVD mortality				
Model 1 (unadjusted)	1.26 (1.23–1.30)‡	1.25 (1.22–1.28)‡	1.37 (1.31–1.43)‡	1.28 (1.23–1.33)‡
Model 2	1.18 (1.15–1.21)‡	1.15 (1.13–1.18)‡	1.19 (1.13–1.25)‡	1.14 (1.09–1.19)‡
Model 3	1.13 (1.10–1.17)‡	1.12 (1.09–1.15)‡	1.15 (1.09–1.21)‡	1.11 (1.07–1.17)‡
Model 4	1.06 (1.01–1.10)†	1.08 (1.05–1.12)‡	1.10 (1.02–1.19)*	1.05 (0.99–1.12)
CHD mortality				
Model 1 (unadjusted)	1.25 (1.20–1.29)‡	1.24 (1.20–1.27)‡	1.41 (1.33–1.49)‡	1.30 (1.24–1.37)‡
Model 2	1.16 (1.12–1.20)‡	1.14 (1.11–1.17)‡	1.23 (1.15–1.31)‡	1.16 (1.10–1.23)‡
Model 3	1.11 (1.07–1.16)‡	1.10 (1.07–1.14)‡	1.17 (1.10–1.25)‡	1.12 (1.06–1.19)‡
Model 4	1.05 (0.997–1.10)	1.08 (1.03–1.12)‡	1.14 (1.03–1.25)†	1.04 (0.96–1.12)
Stroke mortality				
Model 1 (unadjusted)	1.39 (1.29–1.51)‡	1.38 (1.29–1.47)‡	1.35 (1.22–1.50)‡	1.27 (1.16–1.39)‡
Model 2	1.29 (1.19–1.40)‡	1.27 (1.19–1.36)‡	1.19 (1.06–1.33)†	1.14 (1.03–1.26)†
Model 3	1.26 (1.16–1.37)‡	1.25 (1.17–1.35)‡	1.19 (1.06–1.34)†	1.14 (1.03–1.26)*
Model 4	1.08 (0.96–1.23)	1.19 (1.07–1.33)†	1.13 (0.95–1.34)	1.06 (0.92–1.22)

Sex-specific unadjusted and adjusted HRs (95% CI) of BP (continuous variable) for risk for CVD mortality, CHD mortality, and stroke mortality are shown. As adjustment factors, model 2 includes demographic variables (age at baseline, race, and education) plus SBP (or DBP), model 3 includes demographic variables plus clinical characteristics (body mass index, current smoking, total cholesterol, and diabetes) plus SBP (or DBP), and model 4 includes demographic variables plus clinical characteristics plus SBP plus DBP. Statistical significance was defined as p < 0.05. *p < 0.05; †p < 0.01; ‡p < 0.001. Abbreviations as in Tables 1 and 3.



Isolated Systolic Hypertension in Young and Middle-Aged Adults and 31-Year Risk for Cardiovascular Mortality



The Chicago Heart Association Detection Project in Industry Study

Yuichiro Yano, MD, PhD,* Jeremiah Stamler, MD,* Daniel B. Garside, MA,*† Martha L. Daviglius, MD, PhD,*† Stanley S. Franklin, MD,‡ Mercedes R. Carnethon, PhD,* Kiang Liu, PhD,* Philip Greenland, MD,* Donald M. Lloyd-Jones, MD, ScM*



TABLE 3 Sex-Specific Unadjusted and Multivariate-Adjusted HRs (95% CIs) for Risk for CVD Mortality, CHD Mortality, and Stroke Mortality by Hypertension Subtype

	Men (n = 15,868)					Women (n = 11,213)				
	Optimal-Normal BP	High-Normal BP	ISH	IDH	SDH	Optimal-Normal BP	High-Normal BP	ISH	IDH	SDH
CVD mortality										
Person-ys of follow-up	134,247	120,694	124,588	17,741	92,124	188,288	76,827	45,054	10,198	32,839
Number of events (incidence per 100,000 person-ys)	221 (165)	276 (229)	282 (226)	79 (445)	460 (499)	139 (74)	73 (95)	82 (182)	13 (12.7)	103 (314)
Relative risk: HR (95% CI)										
Model 1 (unadjusted)	1 (reference)	1.39 (1.16-1.66)†	1.39 (1.16-1.65)†	2.75 (2.13-3.56)†	3.16 (2.69-3.71)†	1.00 (reference)	1.28 (0.96-1.70)	2.49 (1.90-3.27)†	1.73 (0.98-3.06)	4.37 (3.39-5.64)†
Model 2	1 (reference)	1.31 (1.10-1.57)†	1.36 (1.14-1.63)†	1.93 (1.49-2.49)†	2.16 (1.83-2.54)†	1.00 (reference)	1.04 (0.78-1.38)	1.66 (1.26-2.19)†	1.21 (0.68-2.13)	2.13 (1.64-2.77)†
Model 3	1 (reference)	1.25 (1.05-1.50)*	1.23 (1.03-1.46)*	1.68 (1.29-2.17)†	1.77 (1.49-2.09)†	1.00 (reference)	1.00 (0.75-1.33)	1.55 (1.18-2.05)†	1.05 (0.59-1.85)	1.79 (1.36-2.37)†
CHD mortality										
Number of events (incidence per 100,000 person-ys)	152 (113)	200 (166)	203 (163)	55 (310)	315 (342)	71 (38)	39 (51)	60 (133)	9 (88)	64 (194)
Relative risk: HR (95% CI)										
Model 1 (unadjusted)	1.00 (reference)	1.46 (1.18-1.81)†	1.45 (1.18-1.79)†	2.78 (2.04-3.79)†	3.13 (2.58-3.79)†	1.00 (reference)	1.34 (0.91-1.98)	3.57 (2.53-5.04)†	2.35 (1.18-4.71)*	5.33 (3.80-7.47)†
Model 2	1.00 (reference)	1.38 (1.12-1.70)†	1.42 (1.15-1.75)†	1.94 (1.43-2.65)†	2.13 (1.75-2.59)†	1.00 (reference)	1.09 (0.73-1.61)	2.35 (1.66-3.33)†	1.62 (0.81-3.25)	2.58 (1.83-3.66)†
Model 3	1.00 (reference)	1.32 (1.07-1.63)*	1.28 (1.04-1.58)*	1.71 (1.25-2.33)†	1.75 (1.43-2.14)†	1.00 (reference)	1.03 (0.70-1.52)	2.12 (1.49-3.01)†	1.33 (0.66-2.68)	1.97 (1.36-2.85)†
Stroke mortality										
Number of events (incidence per 100,000 person-ys)	24 (18)	26 (22)	26 (21)	6 (34)	63 (68)	28 (15)	17 (22)	14 (31)	1 (10)	18 (55)
Relative risk: HR (95% CI)										
Model 1 (unadjusted)	1.00 (reference)	1.20 (0.69-2.09)	1.18 (0.68-2.05)	1.93 (0.79-4.72)	4.05 (2.53-6.49)†	1.00 (reference)	1.48 (0.81-2.70)	2.10 (1.11-3.99)*	0.66 (0.90-4.86)	3.77 (2.09-6.82)†
Model 2	1.00 (reference)	1.14 (0.66-1.99)	1.17 (0.67-2.04)	1.32 (0.54-3.24)	2.74 (1.70-4.41)†	1.00 (reference)	1.22 (0.66-2.23)	1.47 (0.77-2.81)	0.47 (0.06-3.44)	1.92 (1.04-3.53)*
Model 3	1.00 (reference)	1.12 (0.64-1.95)	1.08 (0.62-1.90)	1.15 (0.47-2.83)	2.40 (1.46-3.94)†	1.00 (reference)	1.23 (0.67-2.25)	1.46 (0.76-2.82)	0.45 (0.06-3.30)	1.91 (1.01-3.62)*

Sex-specific unadjusted and adjusted HRs (95% CI) for risk of CVD mortality, CHD mortality, and stroke mortality among each hypertension subtype are shown. As a adjustment factors, model 2 includes demographic variables (age at baseline, race, and education), and model 3 includes demographic variables plus body mass index, current smoking, total cholesterol, and diabetes. Statistical significance was defined as p < 0.05. †p < 0.05; ‡p < 0.01; §p < 0.001.

CHD = coronary heart disease; CI = confidence interval; CVD = cardiovascular disease; HR = hazard ratio; other abbreviations as in Table 1.



Effects of Blood Pressure Reduction in Mild Hypertension

A Systematic Review and Meta-analysis

Johan Sundström, MD, PhD; Hisatomi Arima, MD, PhD; Rod Jackson, MBChB, PhD; Fiona Turnbull, MBChB, MPH (Hons), PhD; Kazem Rahimi, MD; John Chalmers, MD, PhD; Mark Woodward, PhD; and Bruce Neal, MBChB, PhD, on behalf of the Blood Pressure Lowering Treatment Trialists' Collaboration*

Ann Intern Med. 2015;162:184-191.

- **Selección de estudios:** Pacientes sin enfermedad cardiovascular con HTA grado 1 (140-150/90-99 mmHg) que fueron asignados aleatoriamente a tratamiento activo (fármacos antihipertensivos: IECAs, calcioantagonistas, diuréticos (95%) o intensificación del tratamiento vs control: placebo o tratamiento menos intenso.
- **Extracción de los datos:** Se extrajeron los datos de pacientes individuales de los ensayos clínicos del BPLTCC y datos agregados de los otros ensayos clínicos

15.266 pacientes incluidos con seguimiento medio 4,4 años
 Edad media 63 años; 40% mujeres
 PA medias iniciales 146/84

Evento	Porcentaje	Riesgo a 5 años (%)
Eventos Cardiovasculares	5,1	7,4
Enfermedad coronaria	2,8	2,8
Ictus	1,8	4,6
Insuficiencia Cardíaca	2,5	2,4
Muertes Cardiovasculares	3,9	3,1
Muertes totales	4,4	6,6

Effects of Blood Pressure Reduction in Mild Hypertension

A Systematic Review and Meta-analysis

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Ann Intern Med. 2015;162:184-191.

Diferencia media final entre activo vs control
PAS 3,6 mmHg PAD 2,4 mmHg

Evento	Odds Ratio	IC 95%
Eventos CV	0,86	0,74-1,01
Enfermedad coronaria	0,91	0,74-1,12
Ictus	0,72	0,55-0,94
Insuficiencia Cardiaca	0,80	0,57-1,12
Muertes CV	0,75	0,57-0,98
Muertes totales	0,78	0,67-0,92

Original Investigation

**Effect of Self-monitoring and Medication Self-titration
on Systolic Blood Pressure in Hypertensive Patients
at High Risk of Cardiovascular Disease
The TASMINE-SR Randomized Clinical Trial**

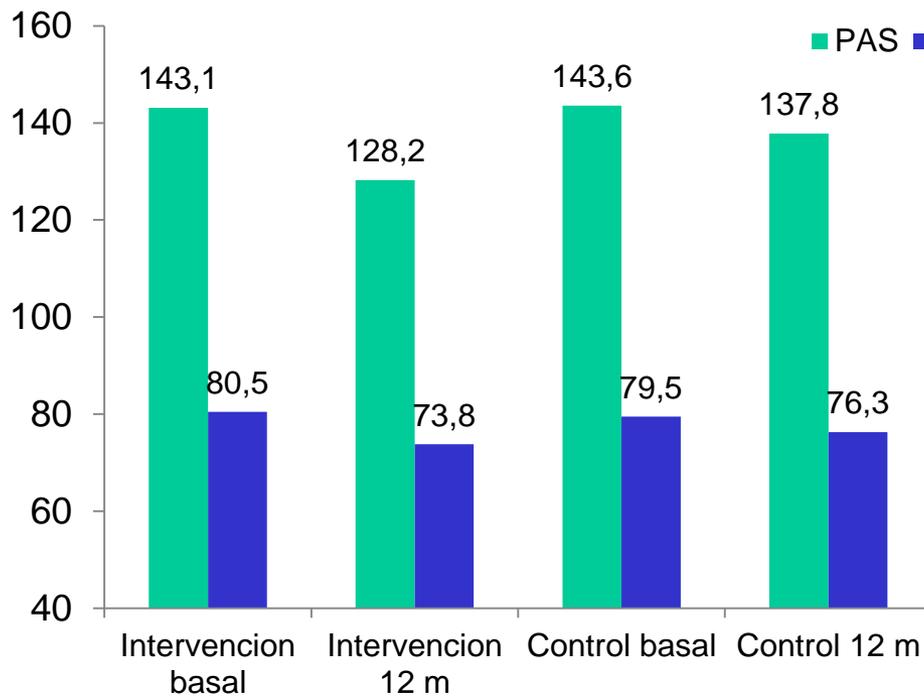
JAMA. 2014;312(8):799-808

Richard J. McManus, FRCGP; Jonathan Mant, MD; M. Sayeed Haque, PhD; Emma P. Bray, PhD;
Stirling Bryan, PhD; Sheila M. Greenfield, PhD; Miren I. Jones, PhD; Sue Jowett, PhD; Paul Little, MD;
Cristina Penaloza, MA; Claire Schwartz, PhD; Helen Shackelford, RGN; Claire Shovelton, PhD;
Jinu Varghese, RGN; Bryan Williams, MD; F.D. Richard Hobbs, FMedSci

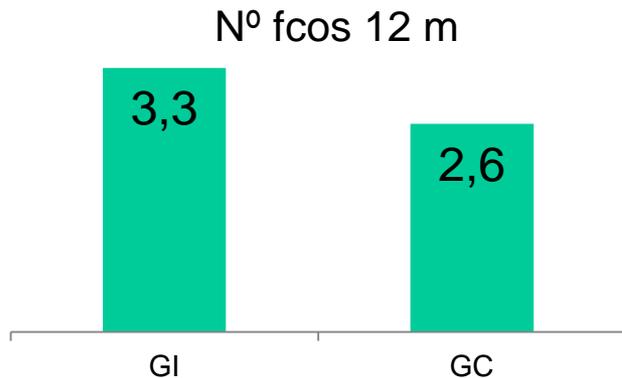
Objetivo determinar si en pacientes con ECV establecida, DM o ERC
La auto-monitorización de la PA y del tratamiento antihipertensivo, con
ajuste de la dosis de acuerdo a un algoritmo, es más eficaz que el
tratamiento habitual en el control de las cifras de PAS

Estudio aleatorizado incluidos 552 pacientes > 35 años
PA clínica objetivo < 130/80 AMPA < 120/75

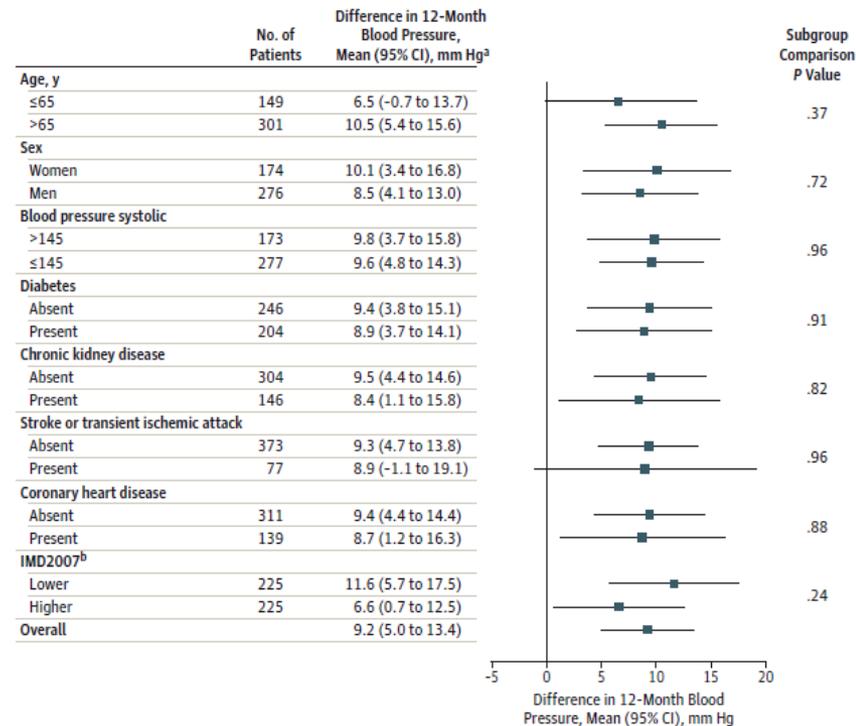
Grupo intervención: AMPA 2 medidas matinales diarias la primera
semana de cada mes si AMPA con al menos 4 lecturas elevadas 2
meses consecutivos, modificaban el tratamiento según algoritmo
previo establecido con su medico de AP



Diferencia entre Intervención vs Control a 12 meses:
PAS 9,2 mmHg PAD 3,4 mmHg



Los fármacos añadidos con mayor frecuencia fueron calcioantagonistas y diuréticos tiazídicos



Less-Tight versus Tight Control of Hypertension in Pregnancy

Laura A. Magee, M.D., Peter von Dadelszen, M.B., Ch.B., D.Phil., Evelyn Rey, M.D., Susan Ross, M.B.A., Ph.D., Elizabeth Asztalos, M.D., Kellie E. Murphy, M.D., Jennifer Menzies, M.Sc., Johanna Sanchez, M.I.P.H., Joel Singer, Ph.D., Amiram Gafni, D.Sc., André Gruslin, M.D.,* Michael Helewa, M.D., Eileen Hutton, Ph.D., Shoo K. Lee, M.D., Ph.D., Terry Lee, Ph.D., Alexander G. Logan, M.D., Wessel Ganzevoort, M.D., Ph.D., Ross Welch, M.B., B.S., D.A., M.D., Jim G. Thornton, M.B., Ch.B., M.D., and Jean-Marie Moutquin, M.D.

- Mujeres gestantes hipertensas entre la 14 y la 33 semana de gestación
- No HTA grave y no proteinuria previa
- Aleatorizadas a control estricto PAD 85 mmHg vs menos estricto PAD 100 mmHg
- Se incluyeron 987 pacientes
- Objetivo compuesto: interrupción embarazo o cuidados perinatales intensivos

Table 2. Primary and Other Perinatal Outcomes.*

Variable	Less-Tight Control (N=493)	Tight Control (N=488)	Adjusted Odds Ratio (95% CI)†
Primary outcome — no. (%)	155 (31.4)	150 (30.7)	1.02 (0.77–1.35)
Pregnancy loss — no. (%)	15 (3.0)	13 (2.7)	1.14 (0.53–2.45)
Miscarriage	0	1 (0.2)	
Ectopic pregnancy	0	0	
Elective termination‡	1 (0.2)	1 (0.2)	
Perinatal death	14 (2.8)	11 (2.3)	1.25 (0.56–2.81)
Stillbirth	12 (2.4)	7 (1.4)	
Neonatal death	2 (0.4)	4 (0.8)	
High-level neonatal care for >48 hr — no./total no. (%)§	141/480 (29.4)	139/479 (29.0)	1.00 (0.75–1.33)
Gestational age at delivery — wk	36.8±3.4	37.2±3.1	
Small-for-gestational-age newborns — no./total no. (%)¶			
Birth weight <10th percentile	79/491 (16.1)	96/488 (19.7)	0.78 (0.56–1.08)
Birth weight <3rd percentile	23/491 (4.7)	26/488 (5.3)	0.92 (0.51–1.63)
Other perinatal outcomes of liveborn infants			
Respiratory complications — no./total no. (%)			
Clinical respiratory problem	82/480 (17.1)	67/479 (14.0)	1.19 (0.83–1.71)
Administration of oxygen beyond the first 10 min of life	34/479 (7.1)	25/477 (5.2)	1.24 (0.72–2.14)
Ventilatory support (with or without intubation) beyond the first 10 min of life	35/478 (7.3)	38/479 (7.9)	0.86 (0.53–1.40)
Use of surfactant	28/480 (5.8)	26/479 (5.4)	0.97 (0.55–1.69)
At least one serious neonatal complication — no./total no. (%)	40/480 (8.3)	40/479 (8.4)	0.96 (0.60–1.52)

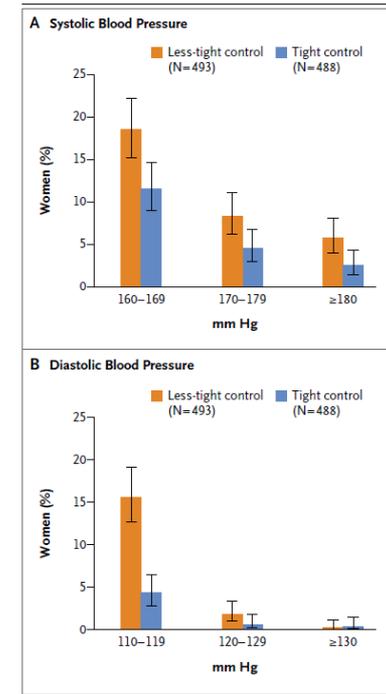


Table 3. Secondary and Other Maternal Outcomes.*

Variable	Less-Tight Control (N=493)	Tight Control (N=488)	Adjusted Odds Ratio (95% CI)†
Serious maternal complications — no. (%)‡	18 (3.7)	10 (2.0)	1.74 (0.79–3.84)
Uncontrolled hypertension	0	0	
Transient ischemic attack or stroke	0	1 (0.2)	
Pulmonary edema	2 (0.4)	1 (0.2)	
Renal failure	0	1 (0.2)	
Transfusion§	16 (3.2)	8 (1.6)	
Placental abruption — no. (%)	11 (2.2)	11 (2.3)	0.94 (0.40–2.21)
Severe hypertension — no. (%)	200 (40.6)	134 (27.5)	1.80 (1.34–2.38)
Preeclampsia — no./total no. (%)	241/493 (48.9)	223/488 (45.7)	1.14 (0.88–1.47)
Defined only by new proteinuria¶	148/493 (30.0)	132/488 (27.0)	1.08 (0.74–1.59)
At least one symptom of preeclampsia	171/493 (34.7)	156/488 (32.0)	1.11 (0.84–1.46)
Abnormal laboratory test results			
Platelet count <100×10 ⁹ /liter	21/493 (4.3)	8/488 (1.6)	2.63 (1.15–6.05)
Elevated AST or ALT level, with symptoms	21/492 (4.3)	9/488 (1.8)	2.33 (1.05–5.16)
Elevated LDH level, with symptoms	16/491 (3.3)	9/488 (1.8)	1.78 (0.77–4.11)
HELLP syndrome**	9/493 (1.8)	2/488 (0.4)	4.35 (0.93–20.35)
Serum creatinine level >2.3 mg/dl	0	1/488 (0.2)	

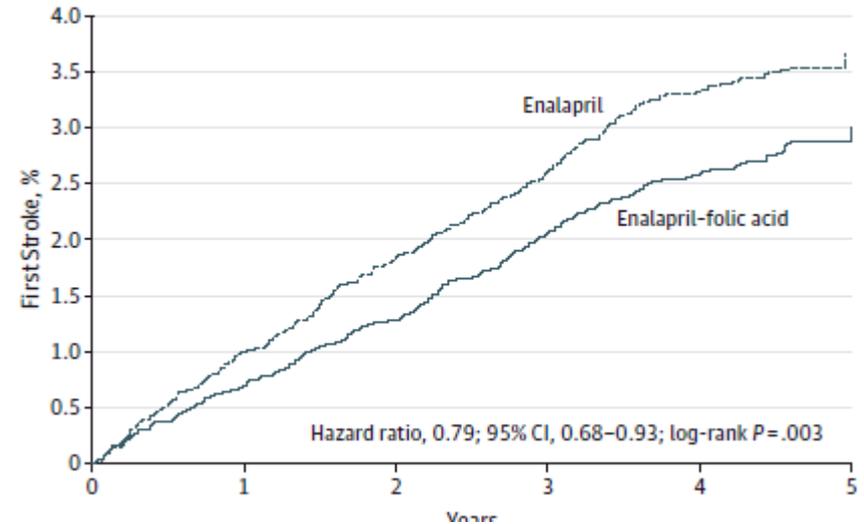


Efficacy of Folic Acid Therapy in Primary Prevention of Stroke Among Adults With Hypertension in China

The CSPPT Randomized Clinical Trial

Yong Huo, JAMA 2015 Apr 7;313(13):1325-35

Hipertensos entre 45-75 años sin ECV previa
Enalapril 10 mg + ácido fólico 0,8 mg vs enalapril 10 mg
 20.702 chinos incluidos
 Seguimiento medio 4,5 años



Outcomes	No. (%) With Outcome		Hazard Ratio (95% CI) ^a	<i>P</i> Value ^b
	Enalapril-Folic Acid (n = 10 348)	Enalapril (n = 10 354)		
First stroke (primary outcome) ^c	282 (2.7)	355 (3.4)	0.79 (0.68-0.93) ^d	.003
Secondary outcomes				
Ischemic stroke	223 (2.2)	292 (2.8)	0.76 (0.64-0.91)	.002
Hemorrhagic stroke	58 (0.56)	62 (0.60)	0.93 (0.65-1.34)	.71
Composite of stroke, myocardial infarction, or death due to cardiovascular causes	324 (3.1)	405 (3.9)	0.80 (0.69-0.92)	.002
Myocardial infarction ^e	25 (0.24)	24 (0.23)	1.04 (0.60-1.82)	.89
Death due to cardiovascular causes ^f	43 (0.4)	43 (0.4)	1.00 (0.66-1.53)	>.99
All-cause death	302 (2.9)	320 (3.1)	0.94 (0.81-1.10)	.47

Caffeine and Blood Pressure



Associations of Ambulatory Blood Pressure With Urinary Caffeine and Caffeine Metabolite Excretions

Idris Guessous.
Hypertension. 2015;65:691-696.

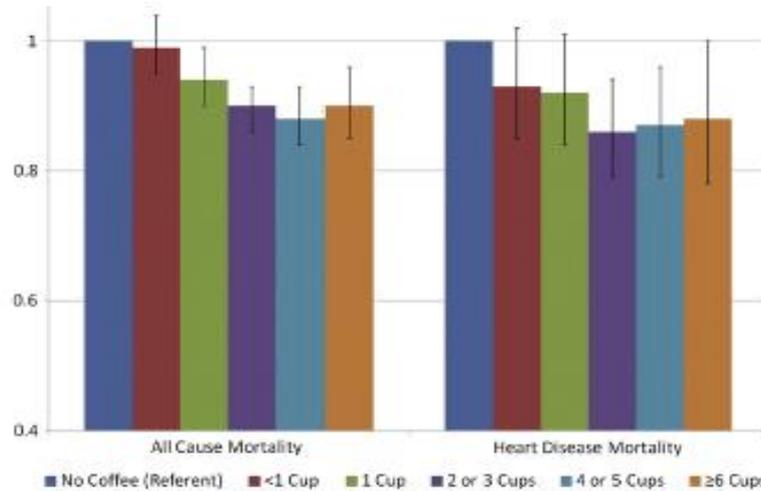
Idris Guessous, Menno Pruijm, Belén Ponte, Daniel Ackermann, Georg Ehret, Nicolas Ansermot, Philippe Vuistiner, Jan Staessen, Yumei Gu, Fred Paccoud, Markus Mohaupt, Bruno Vost, Antoinette Pechère-Berstchi, Pierre-Yves Hombres, J.H. O'Keefe. J Am Coll Cardiol 2013;62:1043-51

- 836 suizos
- MAPA de 24 horas
- Medicion de cafeina

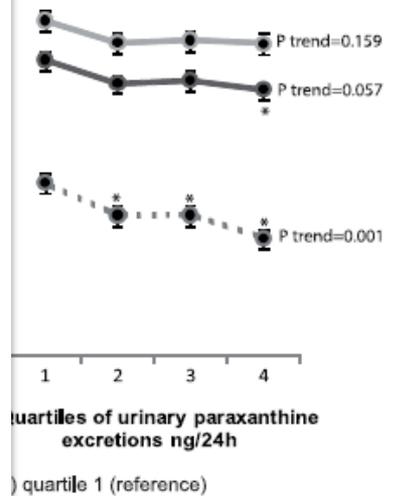
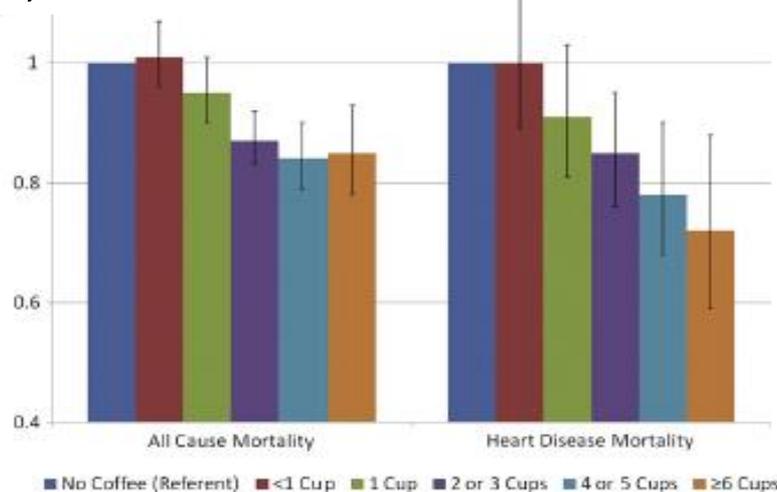
Table 2. Adjusted Associations of Ambulatory Blood Pressure With 24-h Methylxanthines Excretions

Methylxanthine	Systolic BP	
	Beta, SE	P Value
Caffeine*		
24 h	-0.642, 0.296	0.03
Day	-0.505, 0.313	0.11
Night	-1.107, 0.315	<0.001
Paraxanthine*		
24 h	-0.718, 0.343	0.04
Day	-0.545, 0.362	0.15
Night	-1.376, 0.364	<0.001
Theophylline*		
24 h	-0.633, 0.341	0.06
Day	-0.458, 0.360	0.21
Night	-1.183, 0.363	0.001
Theobromine*		
24 h	0.302, 0.338	0.36
Day	0.325, 0.357	0.36
Night	0.003, 0.361	0.98

Models are adjusted for age, sex, body mass index, diabetes mellitus, current alcohol use, antihypertensive treatment, blood Na⁺ and K⁺ values highlighted in bold are statistically significant. BP, blood pressure; and GFR, glomerular filtration rate. *Log-transformed.



Mujeres



HTA 2014 en 30 segundos



- Mayor consumo de sal mayor incremento de PA y a la inversa con K
- El riesgo mas bajo de mortalidad y morbilidad CV con consumo de 10-15 g sal/día y mayor ingesta de K.
- Los valores de PAS nocturna son los mejores predictores de mortalidad CV en hipertensos y en diabéticos
- La HSA en el joven presenta un mayor riesgo CV
- El tto antihipertensivo previene ictus y mortalidad en HTA grado 1
- La AMPA y auto ajuste tto antiHTA reduce + PAS a 12 meses que cuidado habitual.
- Un objetivo de control estricto de PA no confiere mejor pronostico materno-infantil en gestantes hipertensas.
- El acido fólico asociado a enalapril redujo la incidencia de ictus en hipertensos
- A mayor consumo de café menores valores de PAS ambulatoria



MUCHAS GRACIAS



SEMI-FINAL 1

FC BARCELONA

VS

FC BAYERN MÜNCHEN

SEMI-FINAL 2

JUVENTUS

VS

REAL MADRID CF

FINAL
BERLIN 2015



Barcelona-Bayer

AS



UEFA Club Competitions
Semi-finals draws

