Supplementary file 1 Search Strategy to identify randomised salt reduction trials

Ovid MEDLINE(R) from 1946 to January 2019

- 1 sodium chloride, dietary/
- 2 exp sodium, dietary/
- 3 diet, sodium-restricted/
- 4 ((sodium or salt) adj3 (restrict\$ or curb\$ or limit\$ or minim\$ or low\$ or reduc\$ or intake or diet\$ or free)).tw.
- 5 or/1-4
- 6 randomized controlled trial.pt.
- 7 controlled clinical trial.pt.
- 8 randomized.tw.
- 9 placebo.tw.
- 10 drug therapy/
- 11 randomly.tw.
- 12 trial.tw.
- 13 groups.tw.
- 14 or/6-13
- animals/ not (humans/ and animals/)
- 16 14 not 15
- 17 5 and 16
- 18 remove duplicates from 17

EMBASE from 1946 to January 2019

- 1 sodium chloride, dietary/
- 2 sodium intake/
- 3 sodium restriction/
- 4 ((sodium or salt) adj3 (restrict\$ or curb\$ or limit\$ or minimi\$ or low\$ or reduc\$ or intake or diet\$ or free)).tw.
- 5 or/1-4
- 6 randomized controlled trial/
- 7 crossover procedure/
- 8 double-blind procedure/
- 9 random\$.tw.
- 10 (crossover\$ or cross-over\$).tw.
- 11 placebo\$.tw.
- 12 (doubl\$ adj blind\$).tw.
- 13 assign\$.tw.
- 14 allocat\$.tw.
- 15 or/6-14
- 16 (animal\$ not (human\$ and animal\$)).mp.
- 17 15 not 16
- 18 5 and 17
- 19 remove duplicates from 18

Cochrane Central Register of Controlled Trials (Wiley)

- 1 MeSH descriptor Sodium, Chloride, Dietary
- 2 MeSH descriptor Sodium, Dietary explode all trees
- 3 MeSH descriptor Diet, Sodium-Restricted
- 4 sodium near3 (restrict* or curb* or limit* or minimi* or low* or reduc* or intake or diet* or free):ti,ab
- 5 salt near3 (restrict* or curb* or limit* or minimi* or low* or reduc* or intake or diet* or free):ti,ab
- 6 #1 or #2 or #3 or #4 or #5

Supplementary file 2 Study Characteristics

Gr. 1	G. I	No. of	Mean (range)	Female	White	Baseline	ъ :	Study	Change in UNa		change of BP nHg)
Study	Study country	participants	age (years)	(%)	(%)	SBP	Design	duration (days)	(mmol/24-hour)	Systolic (SE)	Diastolic (SE)
Parijs 1973 ¹	Belgium	15	41	55	NR	175	X	28	-98	-6.7 (3.48)	3.2 (4.24)
Mark 1975 ²	USA	6	27.8 (24-41)	0	NR	133	X	10	-305	-13.1 (2.77)	-7.0 (2.50)
Morgan 1981 ³	Australia	24	39	50	100	-	P	56	-88	-	-6.0 (2.41)
Skrabal 1981 ⁴	Austria	21	NR	0	NR	125	X	14	-170	-2.7 (2.36)	-3.0 (2.03)
MacGregor 1982 ⁵	UK	19	49 (30-66)	26	63	154	X	28	-76	-10.0 (2.40)	-5.0 (1.47)
Puska 1983 ⁶	Russia	72	NR	NR	100	153	P	42	-117	0.1 (3.23)	-0.7 (2.29)
Silman 1983 ⁷	UK	25	NR	NR	NR	155	P	365	-53	-8.7 (10.22)	-6.3 (4.42)
Watt 19838	UK	18	52 (31-64)	67	100	137	X	28	-56	-0.5 (1.50)	-0.3 (0.80)
Erwteman 19849	Netherlands	94	46	38	76	157	P	168	-58	-2.7 (2.20)	-3.4 (1.70)
Gillies 1984 ¹⁰	Australia	24	57	42	NR	147	X	42	-77	-2.4 (3.63)	-2.6 (2.46)
Koolen 1984 ¹¹	Netherlands	25	41.3 (22-61)	40	100	148	X	14	-208	-6.2 (3.03)	-4.6 (1.87)
Koolen 1984 ¹²	Netherlands	20	40.8 (22-61)	45	100	148	X	14	-213	-6.5 (3.11)	-4.9 (1.71)
Maxwell 1984 ¹³	US	30	47	50	NR	149	P	7	-190	-6.3 (6.77)	-4.0 (4.24)
Myers 1984 ¹⁴	Australia	125	39	NR	NR	123	X	14	-147	-6.0 (1.00)	-4.0 (1.00)
Richards 1984 ¹⁵	New Zealand	12	NR	33	100	150	X	28	-105	-5.2 (4.10)	-1.8 (3.55)
Skrabal 1984 ¹⁶	Austria	52	NR	0	NR	120	X	14	-149	-2.4 (1.22)	-1.5 (0.99)
Resnick 1985 ¹⁷	US	12	NR	NR	NR	159	X	5	-200	-3.0 (1.89)	-1.0 (1.61)
Skrabal 1985 ¹⁸	Australia	62	NR	NR	NR	120	X	14	-153	-3.0 (0.65)	-1.5 (0.54)
Ashry 1987 ¹⁹	UK	26	26	38	100	129	X	14	-227	0.0 (2.28)	-2.5 (2.57)
Grobbee 1987 ²⁰	Netherlands	40	24	15	NR	137	X	42	-72	-0.9 (1.80)	0.2 (1.67)
MacGregor 1987 ²¹	UK	15	52 (33-71)	27	67	150	X	30	-100	-13.0 (3.14)	-9.0 (3.02)
Morgan 1987 ²²	Australia	20	60.5 (50-65)	0	NR	143	P	60	-62	-6.0 (5.00)	-4.2 (2.95)
Lawton 1988 ²³	US	22	24.5 (20-31)	0	100	115	X	6	-319	-1.6 (2.04)	0.0 (1.47)
Morgan 1988 ²⁴	Australia	16	63 (48-69)	0	NR	173	X	14	-50	-3.0 (1.41)	-4.0 (1.88)
Morgan 1988 ²⁵	Australia	8	NR	NR	NR	156	X	14	-67	-7.0 (1.06)	-6.0 (1.06)
Nowson 1988 ²⁶	Australia	107	52	15	100	150	P	84	-47	-5.1 (1.42)	-4.2 (0.85)
Nowson 1988 ²⁶	Australia	105	52	15	100	149	P	84	-51	-0.2 (1.42)	0.5 (0.99)
Staessen 1988 ²⁷	Belgium	1510	41	48	NR	131	P	1825	-12	-0.2 (1.73)	-1.0 (1.09)

C4 J	C4 J	No. of	Mean (range)	Female	White	Baseline	Decign	Study duration	Change in UNa		change of BP nHg)
Study	Study country	participants	age (years)	(%)	(%)	SBP	Design	(days)	(mmol/24-hour)	Systolic (SE)	Diastolic (SE)
Chalmers 1989 ²⁸	Australia	88	59	17	100	152	X	56	-67	-3.6 (0.70)	-2.1 (0.40)
Chalmers 1989 ²⁹	Australia	108	59	17	100	144	P	56	-71	-5.5 (1.48)	-2.8 (0.85)
Dodson 1989 ³⁰	UK	9	62	33	NR	171	X	30	-76	-9.7 (5.80)	-5.1 (2.94)
Hargreaves 1989 ³¹	Australia	8	23	0	NR	129	X	14	-106	-6.0 (4.00)	-3.0 (3.00)
MacGregor 1989 ³²	UK	20	57 (42-72)	45	75	163	X	30	-141	-16.0 (4.12)	-9.0 (2.32)
Bruun 1990 ³³	Denmark	22	46.5 (29-67)	36	NR	135	X	4	-336	-6.6 (4.08)	-2.6 (3.36)
Parker 1990 ³⁴	Australia	59	52	0	NR	138	P	28	-93	1.0 (1.90)	0.4 (1.10)
Río 1990 ³⁵	Spain	15	49.2 (36-65)	47	100	149	X	14	-100	-3.4 (2.76)	-1.1 (2.03)
Sharma 1990 ³⁶	Germany	15	24 (20-31)	0	NR	107	X	7	-192	-0.9 (3.20)	-3.7 (2.69)
Sharma 1990 ³⁷	Germany	40	25 (20-31)	0	NR	113	X	7	-214	-2.1 (1.43)	-3.1 (1.22)
Carney 1991 ³⁸	Australia	11	54 (30-65)	55	NR	144	X	42	-102	-1.0 (5.57)	1.0 (3.61)
Creager 1991 ³⁹	US	17	30	0	NR	122	X	5	-168	2.0 (3.00)	2.0 (2.00)
Sharma 1991 ⁴⁰	Australia	23	24.9 (23-29)	0	NR	120	X	6	-246	-4.5 (1.43)	-2.3 (1.16)
Singer 1991 ⁴¹	UK	21	54	38	71	147	X	30	-91	-9.0 (2.34)	-3.0 (1.05)
Alli 1992 ⁴²	Italy	56	48	57	NR	149	P	365	8	-6.3 (3.06)	-3.8 (1.32)
Arroll 1992 ⁴³	New Zealand	181	55	48	NR	145	P	180	-1	-0.5 (2.50)	1.5 (1.50)
Benetos 1992 ⁴⁴	France	20	42	55	100	149	X	28	-78	-6.5 (1.67)	-3.7 (1.29)
Cobiac 1992 ⁴⁵	Australia	106	67	34	100	132	P	28	-71	-2.8 (1.41)	-1.3 (0.86)
Gow 1992 ⁴⁶	UK	9	NR	0	NR	120	X	7	-94	-8.0 (3.47)	-3.0 (3.61)
Huggins 1992 ⁴⁷	Australia	9	NR	22	NR	112	X	14	-97	-1.0 (2.00)	-2.0 (3.00)
Cutler 1992 ⁴⁸	US	744	43	29	82	125	P	540	-44	-1.7 (0.59)	-0.9 (0.42)
Fotherby 1993 ⁴⁹	UK	17	73 (66-79)	78	100	179	X	35	-79	-8.0 (3.77)	0.0 (2.39)
Nestel 1993 ⁵⁰	Australia	66	66	45	100	146	P	42	-84	-3.9 (2.94)	-1.5 (2.25)
Redon-Mas 1993 ⁵¹	Spain	418	55	46	NR	163	P	28	-109	0.9 (1.41)	1.8 (0.92)
Río 1993 ⁵²	Spain	30	49.2 (30-65)	43	NR	156	X	14	-151	-1.4 (2.26)	-0.5 (1.36)
Ruilope 1993 ⁵³	Spain	19	NR	NR	NR	157	X	21	-67	-5.9 (3.35)	-5.3 (1.62)
Ruppert 1993 ⁵⁴	Germany	163	38	40	NR	113	X	7	-274	-2.2 (0.84)	1.0 (0.51)
Sharma 1993 ⁵⁵	Germany	16	NR	0	NR	111	X	7	-224	-1.4 (5.70)	-0.5 (1.67)
Sharma 1993 ⁵⁶	Germany	15	25.3 (20-31)	0	NR	106	X	7	-174	-0.1 (2.13)	-2.1 (1.92)
Sharma 1993 ⁵⁷	Germany	18	25.1 (21-28)	0	NR	111	X	7	-218	-1.8 (1.71)	-0.4 (1.42)
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C4 I	C414	No. of	Mean (range)	Female	White	Baseline	Design	Study duration	Change in UNa		change of BP nHg)
Study	Study country	participants	age (years)	(%)	(%)	SBP	Design	(days)	(mmol/24-hour)	Systolic (SE)	Diastolic (SE)
Zoccali 1993 ⁵⁸	Italy	14	47 (30-65)	0	NR	150	X	7	-169	-13.0 (2.71)	-6.0 (1.60)
Howe 1994 ⁵⁹	Australia	56	55	45	NR	145	P	42	-78	-4.2 (2.09)	-1.5 (2.14)
Iwaoka 1994 ⁶⁰	Japan	31	46	45	0	149	X	7	-266	-14.3 (3.92)	-4.6 (1.67)
MacFadyen 1994 ⁶¹	UK	12	NR	0	NR	114	X	3	-50	7.0 (2.71)	10.0 (2.16)
Zoccali 1994 ⁶²	Italy	15	45 (30-65)	13	100	144	X	7	-163	-14.0 (4.70)	-8.0 (2.69)
Doig 1995 ⁶³	US	8	25	0	NR	-	X	4	-112	-2.3 (1.73)	0.0 (2.01)
Draaijer 1995 ⁶⁴	Netherlands	10	41	0	NR	159	X	7	-259	-7.5 (2.31)	-0.5 (0.22)
Stein 1995 ⁶⁵	US	7	34	0	100	123	X	5	-183	1.4 (3.83)	-1.2 (3.10)
Weir 1995 ⁶⁶	US	22	60	36	41	160	X	14	-136	-2.2 (2.75)	-1.4 (1.72)
Bellini 1996 ⁶⁷	Italy	43	46	0	100	167	X	14	-217	-10.3 (2.67)	-9.8 (1.35)
Ferri 1996 ⁶⁸	Italy	61	47	0	100	169	X	14	-265	-7.4 (1.78)	-3.5 (0.84)
Grey 1996 ⁶⁹	New Zealand	34	23	0	100	116	X	7	-133	1.0 (1.74)	1.0 (1.07)
Inoue 1996 ⁷⁰	UK	14	46 (21-59)	57	50	153	X	7	-293	-15.2 (2.62)	-3.7 (1.9)
Ishimitsu 1996 ⁷¹	Japan	30	54	53	0	147	X	7	-193	-11.8 (3.23)	-4.2 (1.78)
Schorr 1996 ⁷²	Germany	16	64	56	100	134	X	28	-71	-7.2 (4.90)	-2.9 (2.61)
Zoccali 1996 ⁷³	Italy	14	47 (37-59)	14	NR	140	X	7	-145	-11.0 (5.09)	-6.0 (2.78)
Cappuccio 1997 ⁷⁴	UK	47	66.8 (60-78)	49	89	163	X	30	-83	-7.2 (3.02)	-3.2 (1.42)
Cutler 1997 ⁷⁵	US	1190	44	33	80	127	P	1095	-40	-1.2 (0.50)	-0.7 (0.40)
McCarron 1997 ⁷⁶	US	99	52	42	73	139	X	28	-55	-4.9 (1.87)	-2.9 (1.10)
Meland 1997 ⁷⁷	Norway	16	50 (20-69)	19	100	145	X	56	-66	-4.0 (1.97)	-2.0 (1.25)
Schorr 1997 ⁷⁸	Germany	90	25	0	NR	110	X	7	-196	0.1 (0.83)	0.8 (0.65)
Yamamoto 1997 ⁷⁹	Japan	36	53.3 (40-69)	81	0	148	P	49	-32	-5.5 (6.40)	-3.3 (4.71)
Foo 1998 ⁸⁰	UK	18	51	56	NR	127	X	6	-149	-5.8 (3.35)	2.0 (2.06)
Gomi 1998 ⁸¹	Japan	12	51.8 (38-65)	33	0	139	X	7	-70	-1.1 (2.22)	0.3 (1.16)
Herlitz 1998 ⁸²	Sweden	6	NR	0	NR	129	X	4	-98	-5.0 (1.62)	-3.0 (0.97)
Wing 1998 ⁸³	Australia	17	61 (37-74)	18	NR	160	X	42	-59	-7.0 (2.40)	-4.0 (1.37)
Davrath 1999 ⁸⁴	US	8	25	0	NR	124	X	5	-96	8.0 (5.52)	5.0 (6.40)
Schorr 1999 ⁸⁵	Germany	187	25.1 (20-30)	0	100	111	X	7	-207	-0.2 (0.63)	0.3 (0.65)
Uzu 1999 ⁸⁶	Japan	70	50 (27-69)	33	0	153	X	7	-173	-15.4 (2.61)	-5.5 (1.37)
Boero 2000 ⁸⁷	Italy	13	51 (21-64)	23	NR	132	X	14	-209	-4.0 (1.26)	-3.0 (0.94)

C4 1	C414	No. of	Mean (range)	Female	White	Baseline	Davis	Study	Change in UNa		change of BP nHg)
Study	Study country	participants	age (years)	(%)	(%)	SBP	Design	duration (days)	(mmol/24-hour)	Systolic (SE)	Diastolic (SE)
Ames 200188	US	21	60	52	62	154	X	28	-126	-6.0 (3.33)	-2.3 (1.71)
Appel 200189	US	681	66	47	76	128	P	90	-40	-4.3 (0.89)	-2.0 (0.61)
Johnson 2001 ⁹⁰	Australia	46	69	NR	NR	-	X	14	-237	-10.8 (2.51)	-5.9 (1.61)
Akita 2003 ⁹¹	US	375	48	57	39	129	X	30	-79	-5.0 (1.27)	-2.0 (0.51)
Dishy 200392	US	25	34	40	84	112	X	6	-300	2.0 (7.81)	1.0 (1.88)
Nowson 2003 ⁹³	Australia	92	45	63	NR	118	X	28	-88	0.4 (1.20)	0.0 (1.00)
Pechere-Bertschi 2003 ⁹⁴	Switzerland	27	26 (20-40)	100	100	102	X	7	-296	-1.4 (2.34)	0.8 (1.48)
Perry 2003 ⁹⁵	UK	15	26	0	NR	115	X	5	-105	0.0 (2.89)	-2.0 (2.50)
Beeks 2004 ⁹⁶	Netherlands	117	53.6 (43-63)	43	NR	166	X	7	-99	1.2 (2.45)	-1.8 (1.06)
Berge-Landry 2004 ⁹⁷	US	48	51	21	71	144	X	28	-285	-16.0 (4.56)	-8.0 (2.28)
Gates 200498	US	12	64	50	100	144	X	28	-89	-7.0 (2.90)	-1.0 (1.83)
Forrester 2005 ⁹⁹	Nigeria	58	47	41	0	122	X	21	-72	-4.8 (1.45)	-3.2 (1.00)
Forrester 2005 ⁹⁹	Jamaica	56	41	39	0	114	X	21	-79	-5.1 (1.42)	-2.2 (1.45)
Swift 2005 ¹⁰⁰	UK	40	50	58	0	159	X	28	-78	-8.0 (2.06)	-3.0 (1.11)
Cappuccio 2006 ¹⁰¹	Ghana	1013	55	62	0	128	P	180	6	-2.5 (2.04)	-4.0 (1.61)
Ho 2007 ¹⁰²	Australia	25	49	68	NR	130	X	14	-210	-5.7 (1.50)	-2.5 (1.00)
Melander 2007 ¹⁰³	Sweden	39	53	49	100	132	X	28	-89	-6.5 (1.50)	-3.3 (1.20)
Townsend 2007 ¹⁰⁴	US	20	30	35	40	117	X	7	-171	-6.0 (4.10)	-4.0 (4.00)
Jessani 2008 ¹⁰⁵	Pakistan	184	50	53	0	122	X	7	-81	-1.0 (0.77)	0.0 (0.77)
Tzemos 2008 ¹⁰⁶	UK	16	27	0	100	121	X	5	-149	-4.0 (1.22)	-1.0 (1.45)
Visser 2008 ¹⁰⁷	Netherlands	34	27	0	100	122	X	7	-181	-5.0 (2.46)	-1.0 (1.29)
Dickinson 2009 ¹⁰⁸	Australia	29	53	76	NR	117	X	14	-92	-5.0 (2.03)	-1.0 (1.40)
He 2009 ¹⁰⁹	UK	169	50	33	42	146	X	42	-55	-4.8 (0.82)	-2.2 (0.43)
Meland 2009 ¹¹⁰	Norway	46	56 (20-75)	26	NR	128	P	56	-38	-5.0 (2.73)	-5.0 (1.49)
Paulsen 2009 ¹¹¹	UK	22	24 (22-30)	45	NR	111	X	4	-78	-1.0 (3.52)	1.0 (2.07)
Pimenta 2009 ¹¹²	US	12	56	67	50	146	X	7	-207	-22.7 (4.93)	-9.1 (2.73)
Weir 2010 ¹¹³	USA	115	52	45	86	134	X	28	-123	-9.4 (0.99)	-5.7 (0.64)
Zanchi 2010 ¹¹⁴	Switzerland	9	NR	0	NR	117	X	7	-250	-3.0 (7.94)	0.0 (4.58)
Starmans-Kool 2011 ¹¹⁵	UK	10	32 (22-40)	0	NR	114	X	14	-97	-2.0 (1.09)	0.0 (2.39)

Study	Study country	No. of	Mean (range)	Female	White	Baseline	Design	Study duration	Change in UNa		change of BP nHg)
Study	Study Country	participants	age (years)	(%)	(%)	SBP	Design	(days)	(mmol/24-hour)	Systolic (SE)	Diastolic (SE)
Carey 2012 ¹¹⁶	US	185	47.2 (18-70)	61	100	123	X	7	-203	-3.5 (1.05)	0.3 (0.65)
Carey 2012 ¹¹⁶	US, France	211	49	39	100	147	X	6	-212	-15.9 (1.25)	-9.2 (0.81)
Graffe 2012 ¹¹⁷	Denmark	21	26	52	100	110	X	4	-172	1.0 (3.36)	1.0 (1.92)
Bonfils 2013 ¹¹⁸	Denmark	36	40	58	NR	124	X	5	-140	-1.7 (2.57)	0.0 (1.30)
Mallamaci 2013 ¹¹⁹	Italy	32	48	28	NR	136	X	14	-165	-8.0 (2.20)	-3.0 (0.96)
Allen 2014 ¹²⁰	US	70	24	63	NR	116	X	5	-306	0.0 (1.40)	3.0 (1.45)
Cavka 2015 ¹²¹	Croatia	54	20	100	NR	105	P	7	-141	-5.0 (2.91)	-3.0 (2.01)
Gijsbers 2015 ¹²²	Netherlands	36	66	33	NR	137	X	28	-98	-7.5 (1.50)	-2.7 (0.76)
He 2015 ¹²³	China	553	44	52	0	126	P	105	-50	-2.3 (1.16)	-0.9 (0.94)
Markota 2015 ¹²⁴	Bosnia and Herzegovina	150	59	51	NR	175	P	60	-28	-4.9 (1.97)	-2.0 (0.60)
Matthews 2015 ¹²⁵	US	20	41	50	70	120	X	7	-205	-5.0 (1.58)	-2.5 (1.21)
Riphagen 2016 ¹²⁶	Netherlands	35	66	34	100	137	X	28	-99	-8.0 (1.50)	-2.9 (0.79)
Suckling 2016 ¹²⁷	UK	46	58	48	70	136	X	42	-49	-4.3 (1.95)	-1.6 (1.15)
Brian 2017 ¹²⁸	US	80	38	51	80	116	X	7	-287	-1.0 (0.78)	0.5 (0.71)
Gefke 2017 ¹²⁹	Sweden	10	25 (22-30)	50	100	114	X	3	-169	-1.0 (2.00)	0.0 (0.88)
Babcock 2018 ¹³⁰	US	21	38 (20-59)	50	NR	113	X	7	-177	0.0 (2.00)	3.0 (1.73)
Parvanova 2018 ¹³¹	Italy	115	64	11	NR	146	P	90	-44	-4.7 (2.02)	-3.0 (1.07)
Rorije 2018 ¹³²	Italy	12	23 (18-31)	0	NR	118	X	8	-322	-1.0 (0.82)	0.0 (2.02)
Wang 2018 ¹³³	China	90	51	64	0	122	X	7	-176	-9.6 (4.83)	-3.1 (1.56)

Abbreviations and symbols:

SBP: systolic blood pressure; Una: urinary sodium excretion; BP: blood pressure; SE: standard error; X: crossover design; P: parallel design

Supplementary file 3 Risk of bias assessment of included studies

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Parijs 1973 ¹	High risk (each patient received a number)	High risk (Those with uneven numbers were instructed to take a low-sodium diet during the first period and a high-sodium diet during the second period and vice versa for those with even numbers)	High risk (open study)	High risk (7 out of 22 participants were lost in the analysis)	Low risk
Mark 1975 ²	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Morgan 1981 ³	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	High risk (SBP not reported)
Skrabal 1981 ⁴	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
MacGregor 1982 ⁵	Unclear	Unclear	Low risk (double blinding)	Low risk	Low risk
Puska 1983 ⁶	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (4 out of 76 lost to follow up)	Low risk
Silman 1983 ⁷	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Watt 1983 ⁸	Unclear	Unclear	Low risk (double blinding)	Low risk (2 out of 20 lost)	Low risk
Erwteman 1984 ⁹	Low risk (block randomization)	Unclear	High risk (Open study, with outcome observer blinded only)	Low risk (13 out of 107 lost to follow up)	Low risk
Gillies 1984 ¹⁰	Unclear	Unclear	High risk (open study)	Low risk (4 out of 28 lost to follow up)	Low risk
Koolen 1984 ¹¹	Unclear	Unclear	High risk (Open study, with outcome observer blinded only)	Low risk (no loss to follow up)	Low risk
Koolen 1984 ¹²	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Maxwell 1984 ¹³	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Myers 1984 ¹⁴	Unclear	Unclear	Unclear	Low risk (ITT analysis)	Low risk
Richards 1984 ¹⁵	Unclear	Unclear	High risk (open study with BP measured by minicomputer)	High risk (4 out of 16 lost)	Low risk
Skrabal 1984 ¹⁶	Unclear	Unclear	High risk (open study)	Unclear	Low risk
Resnick 1985 ¹⁷	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Skrabal 1985 ¹⁸	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Ashry 1987 ¹⁹	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Grobbee 1987 ²⁰	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
MacGregor 1987 ²¹	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Morgan 1987 ²²	Low risk (block randomization)	Unclear	High risk (open study with BP observer blinded only)	Low risk	Low risk
Lawton 1988 ²³	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Morgan 1988 ²⁴	Low risk (latin square)	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Morgan 1988 ²⁵	Unclear	Unclear	Low risk (double blinding)	Unclear	Unclear
Nowson 1988 ²⁶	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk
Nowson 1988 ²⁶	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk
Staessen 1988 ²⁷	Unclear	Unclear	High risk (Neither patients nor observers were blinded)	Low risk (<0.2 loss to follow up)	High risk
Chalmers 1989 ²⁸	Unclear	Unclear	Low risk (double blinding)	Low risk (ITT analysis)	Low risk
Chalmers 1989 ²⁹	Unclear	Unclear	Low risk (double blinding)	Low risk	Low risk
Dodson 1989 ³⁰	Unclear	Unclear	Low risk (double blinding)	High risk (4 out of 13 lost to follow up)	High risk
Hargreaves 1989 ³¹	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
MacGregor 1989 ³²	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Bruun 1990 ³³	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Parker 1990 ³⁴	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Río 1990 ³⁵	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Sharma 1990 ³⁶	Unclear	Unclear	Low risk (participants blinded by placebo and BP was measured by automatic device)	Low risk (no loss to follow up)	Low risk
Sharma 1990 ³⁷	Low risk (latin square design)	Unclear	High risk (open study with BP observer blinded only)	Low risk (5 out of total 45 was excluded from the analysis due to poor compliance)	High risk
Carney 1991 ³⁸	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Creager 1991 ³⁹	Unclear	Unclear	Unclear	Low risk (no loss to follow up)	Low risk
Sharma 1991 ⁴⁰	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (2 out of 25 lost to follow up)	Low risk
Singer 1991 ⁴¹	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Alli 1992 ⁴²	Unclear	Unclear	High risk (Neither patients nor observers were blinded)	High risk (21 out of 77 lost to follow up)	High risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Arroll 1992 ⁴³	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (27 out of 208 lost to follow up)	Low risk
Benetos 1992 ⁴⁴	Low risk (computer randomization)	Unclear	Low risk (double blinding)	Low risk (2 out of 22 lost to follow up)	Low risk
Cobiac 1992 ⁴⁵	Unclear	Unclear	Low risk (double blinding)	Low risk (1 out of 107 randomized was excluded from the analysis because the patient took antihypertensive medication, which was an exclution criteria of the study)	Low risk
Gow 1992 ⁴⁶	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Huggins 1992 ⁴⁷	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Cutler 1992 ⁴⁸	Low risk	Low risk (Randomization assignments were received from the coordinating center by telephone or sealed opaque envelopes were used to convey the treatment assignment)	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk
Fotherby 1993 ⁴⁹	Unclear	Unclear	Low risk (double blinding)	Low risk (1 out of 18 lost to follow up)	Low risk
Nestel 1993 ⁵⁰	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Redon-Mas 1993 ⁵¹	Unclear	Unclear	High risk (open study)	High risk (156 out of 574 not included in the analysis because sodium excretion did not meet requirements of the study)	Low risk
Río 1993 ⁵²	Unclear	Unclear	Low risk (double blinding)	High risk (17 out of 47 lost to follow up)	Low risk
Ruilope 1993 ⁵³	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Ruppert 1993 ⁵⁴	Unclear	Unclear	Low risk (patients blinded by placebo and blood pressure measured by automated device)	Low risk (no loss to follow up)	Low risk
Sharma 1993 ⁵⁵	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Sharma 1993 ⁵⁶	Low risk (randomized according to Latin square design)	Unclear	High risk (open study)	High risk (5 out of 20 lost)	Low risk
Sharma 1993 ⁵⁷	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Zoccali 1993 ⁵⁸	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Howe 1994 ⁵⁹	Unclear	Unclear	Low risk (double blinding)	Low risk (5 out of 61 lost to follow up)	Low risk
Iwaoka 1994 ⁶⁰	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
MacFadyen 1994 ⁶¹	Unclear	Low risk (Treatments were administered in a randomised, double-blind, crossover design according to a pre-prepared schedule administered independently of the investigators by the Department of Pharmacy of the hospital)	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Zoccali 1994 ⁶²	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Doig 1995 ⁶³	Unclear	Low risk (randomization code prepared separately of the investigator)	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Draaijer 1995 ⁶⁴	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Stein 1995 ⁶⁵	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Weir 1995 ⁶⁶	Unclear	Unclear	High risk (Patients blinded only, by using placebo)	Low risk (no loss to follow up)	Low risk
Bellini 1996 ⁶⁷	Unclear	Unclear	Low risk (double blinding)	High risk (12 out of 55 lost to follow up)	High risk
Ferri 1996 ⁶⁸	Unclear	Unclear	Low risk (double blinding)	Low risk (4 out of 65 lost to follow up)	Low risk
Grey 1996 ⁶⁹	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Inoue 1996 ⁷⁰	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Ishimitsu 1996 ⁷¹	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Schorr 1996 ⁷²	Unclear	Unclear	Low risk (double blinding)	High risk (5 out of 16 lost to follow up)	Low risk
Zoccali 1996 ⁷³	Unclear	High risk (study staff were aware of the allocation)	High risk (open study)	High risk (4 out of 18 lost to follow up)	Low risk
Cappuccio 1997 ⁷⁴	Low risk (random- generated numbers handled by one not involved in the clinical assessments) in the clinical assessments)	Low risk (neither nurses nor participants were aware of the treatment allocation)	Low risk (double blinding)	Low risk (1 out of 48 lost to follow up)	Low risk
Cutler 1997 ⁷⁵	Unclear	Low risk (Randomization was performed by telephone contact with the TOHP coordinating center or by opening a sealed opaque envelope)	High risk (open study with BP observer blinded only)	Unclear	Low risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
McCarron 1997 ⁷⁶	Low risk (computer randomization)	Low risk (Placebo-controlled and study personnel blinded to the subjects' NaCl intake)	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Meland 1997 ⁷⁷	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Schorr 1997 ⁷⁸	Unclear	Unclear	High risk (Patients blinded only by using placebo)	Low risk (no loss to follow up)	Low risk
Yamamoto 1997 ⁷⁹	Unclear	Unclear	Unclear	Low risk	High risk
Foo 1998 ⁸⁰	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Gomi 1998 ⁸¹	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Herlitz 1998 ⁸²	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Wing 1998 ⁸³	Low risk (latin square)	Unclear	Low risk (double blinding)	Low risk (2 out 19 lost)	Low risk
Davrath 1999 ⁸⁴	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Schorr 1999 ⁸⁵	Unclear	Unclear	High risk (Open study, with outcome observer blinded only)	Low risk (no loss to follow up)	Low risk
Uzu 1999 ⁸⁶	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Low risk
Boero 2000 ⁸⁷	Unclear	Unclear	High risk (Intervention conducted by physicians and both the physicians and participants were aware of the allocation. Unclear if outcome observers were blinded or not)	Low risk (15 randomized, 1 withdrew consent, 1 didn't comply the protocol and 13 analyzed.)	Low risk
Ames 2001 ⁸⁸	Unclear	Unclear	High risk (Patients blinded only, by using placebo)	High risk (13 out of 30 lost to follow up)	Low risk
Appel 2001 ⁸⁹	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk
Johnson 2001 ⁹⁰	Low risk (latin square)	Unclear	Low risk (double blinding)	Low risk (6 out of 46 lost)	Low risk
Akita 2003 ⁹¹	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (9% loss to follow up)	Low risk
Dishy 2003 ⁹²	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Nowson 2003 ⁹³	Low risk (random number in excel)	Low risk (Placebo-controlled and research personnel were unaware of the randomization status of subjects and tablet allocation)	Low risk (double blinding)	High risk (20 out of 128 lost to follow up and 16 hypertensive not reported)	High risk (results for hypertensive participants not reported)
Pechere-Bertschi 2003 ⁹⁴	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (no loss to follow up)	Unclear

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Perry 2003 ⁹⁵	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Beeks 2004 ⁹⁶	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Berge-Landry 2004 ⁹⁷	Unclear	Unclear	High risk (open study)	Low risk (no loss to follow up)	Low risk
Gates 2004 ⁹⁸	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Forrester 2005 ⁹⁹	Low risk (block randomization)	Unclear	High risk (open study)	Low risk (No loss to follow up)	Low risk
Forrester 2005 ⁹⁹	Low risk (block randomization)	Unclear	Unclear	Low risk (No loss to follow up)	Low risk
Swift 2005 ¹⁰⁰	Unclear	Unclear	Low risk (double blinding)	Low risk (6 out of 46 lost to follow up)	Low risk
Cappuccio 2006 ¹⁰¹	Low risk (Villages were randomised in blocks of two, and stratified for locality (semi-urban or rural) by an independent statistician)	Low risk	High risk (Community health workers conducted the intervention and was not possible to be blinded. Only participants were blinded. Unclear if outcome assessors were blinded or not)	Low risk (ITT analysis)	Low risk
Ho 2007 ¹⁰²	Unclear	Unclear	High risk (open study with BP observer blinded only)	High risk (19 out of 44 lost to follwow up)	Low risk
Melander 2007 ¹⁰³	Unclear	Unclear	Low risk (double blinding)	Low risk (7 out of 46 lost to follow up)	Low risk
Γownsend 2007 ¹⁰⁴	Low risk (using a pre- specified randomized blocked (block=ten subjects) table generated by the GCRC biostatistician and kept by the dietician)	Low risk (using a pre-specified randomized blocked (block=ten subjects) table generated by the GCRC biostatistician and kept by the dietician)	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
essani 2008 ¹⁰⁵	Low risk (computer randomization)	Low risk	High risk (open study with BP observer blinded only)	Low risk (16 out of 200 lost to follow up)	Low risk
Γzemos 2008 ¹⁰⁶	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Visser 2008 ¹⁰⁷	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Unclear
Dickinson 2009 ¹⁰⁸	Low risk (computer randomization)	Unclear	High risk (open study with BP observer blinded only)	Low risk (2 out of 31 lost to follow up)	Low risk
He 2009 ¹⁰⁹	Low risk (computer randomization)	Low risk	Low risk (double blinding)	Low risk (16 out of 185 lost to follow up)	Low risk
Meland 2009 ¹¹⁰	Unclear (simple randomization to ensure equal number in both groups, but didn't	Low risk (The randomization list was concealed and kept inaccessible to the investigators during the trial, and was	Low risk (double blinding)	Low risk (4 out of 50 lost to follow up)	Low risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
	mention how it was done)	disclosed after all the statistical analyses had been conducted)			
Paulsen 2009 ¹¹¹	Unclear	Low risk (randomization and blinding of sodium chloride tablets was conducted by the hospital pharmacy)	Low risk (double blinding)	Low risk (5 out of 27 lost to follow up)	Low risk
Pimenta 2009 ¹¹²	Unclear	Unclear	High risk (open study)	Low risk (1 out of 13 lost to follow up)	Low risk
Weir 2010 ¹¹³	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk (17 out of 132 lost to follow up)	Low risk
Zanchi 2010 ¹¹⁴	Unclear	Unclear	Low risk (double blinding)	Low risk (1 out of 10 lost to follow up)	Low risk
Starmans-Kool 2011 ¹¹⁵	Low risk (computer randomization)	Low risk (A copy of the list was given to the hospital kitchen, and the original was kept in a sealed envelope at the department. The code was revealed when the study was finished)	Low risk (double blinding)	Low risk (4 out of 25 lost to follow up)	Low risk
Carey 2012 ¹¹⁶	Unclear	Unclear	Low risk (double blinding)	Low risk (no loss to follow up)	Low risk
Carey 2012 ¹¹⁶	Unclear	Unclear	High risk (open study)	Unclear	Low risk
Graffe 2012 ¹¹⁷	Unclear	Unclear	High risk (open study)	Unclear	Low risk
Bonfils 2013 ¹¹⁸	Low risk (tossing coin)	High risk (The primary investigator assigned participants to the interventions according to the randomization sequence. The patients were not blinded for treatment assignment)	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk
Mallamaci 2013 ¹¹⁹	Unclear	Unclear	Low risk (intervention was placebo controlled and 24-hour ambulatory BP was measured with a device)	Low risk (no loss to follow up)	Low risk
Allen 2014 ¹²⁰	Unclear	Unclear	High risk (open study with BP observer blinded only)	Unclear (No. randomized not reported and no loss to follow up was reported)	Unclear
Cavka 2015 ¹²¹	Unclear	Unclear	High risk (Patients blinded only, by using placebo)	Low risk (no loss to follow up)	Low risk
Gijsbers 2015 ¹²²	Low risk (computer- generated table)	Low risk (independent person)	Low risk (double blinding)	Low risk (1 out of 37 lost to follow up)	Low risk
He 2015 ¹²³	Low risk (random number list generated by a researcher who was blind to the identity of the participants)	Low risk	High risk (open study with BP observer blinded only)	Low risk (ITT analysis)	Low risk

Study	Sequence generation	Allocation concealment	Blinding of participants, personnel and outcome assessors	Incomplete outcome data	Selective reporting
Markota 2015 ¹²⁴	Unclear	Low risk (sealed envelope)	High risk (open study)	Low risk (no loss to follow up)	Low risk
Matthews 2015 ¹²⁵	Unclear	Unclear	Unclear	Low risk (no loss to follow up)	High risk (results only reported for 20 out of 41)
Riphagen 2016 ¹²⁶	Unclear	Unclear	Low risk (double blinding)	Low risk (1 out of 36 lost to follow up)	Low risk
Suckling 2016 ¹²⁷	Low risk (computer randomization)	Low risk (independent company)	Low risk (double blinding)	Low risk (3 out of 49 lost to follow up)	Low risk
Brian 2017 ¹²⁸	Unclear	Unclear	High risk (open study with patients not blinded to assignment)	High risk (21 out of 101 lost to follow up)	High risk
Gefke 2017 ¹²⁹	Unclear	Unclear	High risk (open study)	Low risk	Low risk
Babcock 2018 ¹³⁰	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk	Low risk
Parvanova 2018 ¹³¹	Low risk	Low risk	High risk (open study with BP observer blinded only)	Low risk	Low risk
Rorije 2018 ¹³²	Low risk (block randomization)	Unclear	High risk (open study)	Low risk	Low risk
Wang 2018 ¹³³	Unclear	Unclear	High risk (open study with BP observer blinded only)	Low risk	Low risk

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