

Patient empowerment and adherence

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Disclosure

- Nothing to disclose

Questions – True or False



True



False

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Questions – True or False

- 1. Patient empowerment is another term for adherence
- 2. Poor adherence cannot be cured, even not by patient empowerment

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Understanding adherence is not that difficult...



'Drugs don't work in patients who don't take them'
CE Koop, MD



'If we have the ball they can't score'
J Cruiff, former soccer player and coach

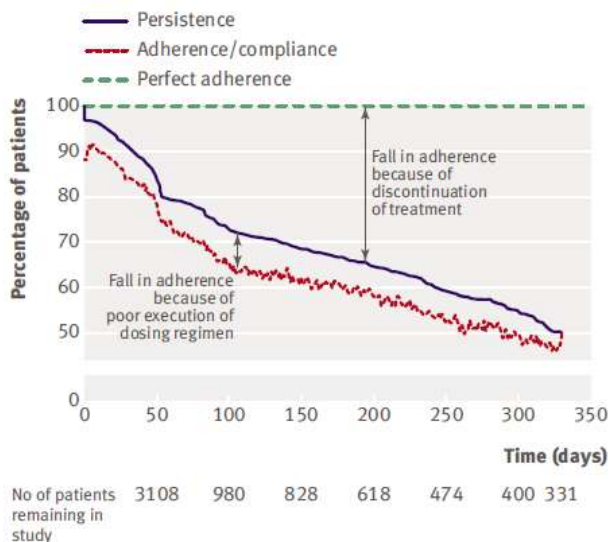
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Hypertension: what's all about

	1976-1980	1988-1991	1991-1994	1999-2000	2011-2012
Awareness (%)	51	73	68	70	82
Treatment (%)	31	55	54	59	75
Control (%)	10	29	27	34	52

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Chobanian AV. 7th report of the JNC 2003, Yoon SS. Hypertension 2015



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Vrijens B, *Bmj* 2008;1114-7.

Table 2. Association between non-compliance and discontinuation with treatment for patients who used monotherapy

	Cases*	Controls*	OR crude	OR adjusted [†]
Non-compliant (n = 9111)	1277 (14.0%)	532 (5.8%)	2.86 (2.55–3.20)	2.86 (2.52–3.24)
Stratified on gender				
Males (n = 3764)	485 (12.9%)	218 (5.8%)	2.61 (2.18–3.12)	2.52 (2.05–3.10)
Females (n = 5347)	792 (14.8%)	314 (5.9%)	3.03 (2.62–3.52)	3.06 (2.61–3.59)
Stratified on type of antihypertensive				
Diuretics (n = 2700)	494 (18.3%)	208 (7.7%)	2.96 (2.45–3.57)	3.13 (2.54–3.85)
Beta-blockers (n = 4252)	507 (11.9%)	228 (5.4%)	2.53 (2.13–2.97)	2.54 (2.11–3.06)
Calcium antagonists (n = 739)	100 (13.5%)	32 (4.3%)	3.96 (2.50–6.25)	3.65 (2.16–6.17)
ACE-inhibitors (n = 906)	116 (12.8%)	41 (4.5%)	3.27 (2.22–4.83)	3.33 (2.05–5.46)
Angiotensin II receptor antagonists (n = 435)	45 (10.3%)	14 (3.2%)	4.10 (2.05–8.19)	3.60 (1.71–7.60)
Miscellaneous (n = 79)	15 (19.0%)	9 (11.4%)	2.00 (0.75–5.33)	1.51 (0.51–4.51)
Stratified on age group				
0–19 years (n = 85)	19 (22.4%)	12 (14.1%)	1.78 (0.79–4.02)	1.76 (0.74–4.17)
20–39 years (n = 935)	163 (17.4%)	65 (7.0%)	3.13 (2.25–4.36)	3.30 (2.32–4.69)
40–59 years (n = 3810)	538 (14.1%)	230 (6.0%)	2.74 (2.31–3.26)	2.70 (2.24–3.26)
60–79 years (n = 3467)	455 (13.1%)	168 (4.8%)	3.32 (2.71–4.05)	3.46 (2.73–4.38)
≥80 years (n = 814)	102 (12.5%)	57 (7.0%)	2.00 (1.34–2.86)	1.93 (1.30–2.87)
Stratified on duration of use				
<90 days (n = 7273)	995 (13.7%)	401 (5.5%)	3.02 (2.65–3.45)	3.10 (2.67–3.59)
≥90 days (n = 1838)	282 (15.3%)	130 (7.1%)	2.41 (1.92–3.01)	2.28 (1.79–2.92)

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Van Wijk BL. *Pharmacoepidemiol Drug Saf* 2006

Adherence Within 6 mo After Diagnosis	HR* (95% CI)	P
Model 1†		
Low (PDC <40%)	1.00	<0.001§
Intermediate (PDC, 40% to 79%)	0.87 (0.73–1.03)	0.117
High (PDC ≥80%)	0.50 (0.35–0.69)	<0.001
Model 2‡		
Low (PDC <40%)	1.00	<0.001§
Intermediate (PDC, 40% to 79%)	0.86 (0.71–1.03)	0.109
High (PDC ≥80%)	0.62 (0.40–0.96)	0.032

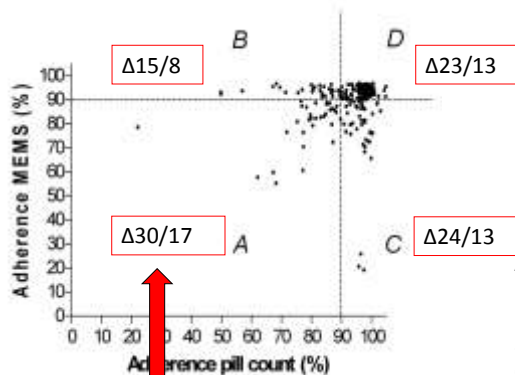
Mazzaglia G. *Circulation* 2009

Adherence†	All-cause death		Stroke		Acute myocardial infarction	
	HR (95% CI)	P value	HR (95% CI)	P value	HR (95% CI)	P value
Poor	1.00	<0.001*	1.00	0.381*	1.00	0.877*
Moderate	0.81 (0.66–0.99)	0.038	0.96 (0.69–1.33)	0.799	1.14 (0.71–1.83)	0.592
Good	0.59 (0.48–0.72)	<0.001	0.81 (0.59–1.11)	0.197	0.98 (0.62–1.55)	0.939
Excellent	0.37 (0.31–0.45)	<0.001	0.82 (0.63–1.07)	0.138	0.96 (0.65–1.40)	0.825

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Esposti LD. *Clinicoecon Outcomes Res* 2011

While knowing that...



Adherence: 82%

- A= not adherent according to MEMS and pill count (14%)
- B= adherent according MEMS, not according to pill count (14%)
- C= adherent according pill count, not according to MEMS (25%)
- D= adherent according MEMS and pill count (47%)

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Van Onzenoort HAW. *Am J Hypertens* 2010

Table 2. Effect of a clinical trial on adherence to treatment

Adherence measure	Before trial	During trial	After trial	p-value
Mean adherence, overall (% [SD])	90.6 (11)	95.6 (7)	91.8 (10)	<0.001
Mean adherence, per ATC-code* (% [SD])				
A	88.5 (17)	90.8 (16)	86.8 (20)	0.59
B	92.8 (9)	89.8 (13)	96.3 (8)	0.025
C	95.1 (9)	97.9 (7)	95.6 (9)	<0.001
G	82.6 (28)	89.7 (20)	93.3 (15)	0.20
H	90.9 (12)	95.5 (7)	87.1 (27)	0.67
L	99.0 (-)	-	84.2 (22)	0.55
M	65.1 (33)	69.0 (36)	69.9 (33)	0.61
N	74.2 (31)	74.9 (32)	76.4 (28)	0.93
R	26.9 (-)	46.6 (-)	84.0 (24)	0.079

*: Anatomical Therapeutic Code; A=Alimentary tract and metabolism; B=Blood and blood forming organs; C=Cardiovascular system; G=Genito-urinary system and sex hormones; H=Systemic hormonal preparations, excluding sex hormones and insulins; L=Antineoplastic and immunomodulating agents; M=Musculo-skeletal system; N=Nervous system; R=Respiratory system

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Van Onzenoort HAW. Hypertension 2011

Table 1. Extracted data from included studies reporting taking compliance and/or correct dosing

Study reference	Year	Design	No. of monitored patients	Monitoring period (weeks)	Aware of MEMS	Taking compliance		Correct dosing	
						Once-daily regimen	Twice-daily regimen	Once-daily regimen	Twice-daily regimen
Andrzejak et al. [10]	2000	RCT	133	24	Yes	99.9	97.5	94	78.1
Bloed et al. [12]	2002	Observational	50	56	-	-	-	83	-
Burns et al. [13]	2000	Observational	41	8	Yes	-	-	93	-
Choo et al. [14]	1999	Observational	280	12	Yes	86	-	-	-
Duffy et al. [15]	1994	RCT cross-over	320	6	Yes	94.9	81.6	84.6	64
Eisen et al. [16]	1990	RCT	88	20	Yes	96	93	-	-
Gavin et al. [17]	1999	RCT cross-over	25	4	-	101.2	90.1	92.2	72.6
Kruse et al. [19]	1994	Observational	24	30.6 ^a	Yes	88.8	87.9	84.8	79.8
Loeman et al. [20]	1997	RCT	177	20	Yes	94	91	90.2	82.2
Malton et al. [24]	1996	Observational	501	4 ^a	Yes	90.5	-	-	-
Malton et al. [23]	1992	RCT cross-over	26	4	-	94.4	-	-	-
Mengden et al. [26]	1993	RCT cross-over	18	4 ^a	No	95.7	-	81.4	-
Maurice-Vohar et al. [27]	1998	RCT	102	12	-	98.3	87.2	92.5	74.8
Neusch et al. [28]	2001	Observational	103	4	-	93	77	-	-
Rudolf et al. [30]	1990	RCT	21	10	Yes	-	89.5	-	62.6
Vaur et al. [22]	1998	Observational	212	7 ^a	Yes	-	-	85.4	-
Waeber et al. [32]	1998	Observational	38	13	Yes	-	-	80.8	-
Waeber et al. [37] ^b	1999	RCT	501	58	Yes	-	-	70.4	-
Studies with no separate compliance data for once- and twice-daily regimens.									
Bertholet et al. [11]	2000	Observational	89	4-8 ^a	Yes	-	-	91.8	-
McKinney et al. [25]	1992	RCT	34	12	No	82	-	-	-

^aAverage monitoring period. ^bCompliance with aspirin or placebo was measured in a subset of patients and was assumed to reflect compliance with anti-hypertensive treatment [42]. RCT, Randomized controlled trial.

WHO: 50%

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Wetzels GE. J Hypertension 2004

Patient empowerment

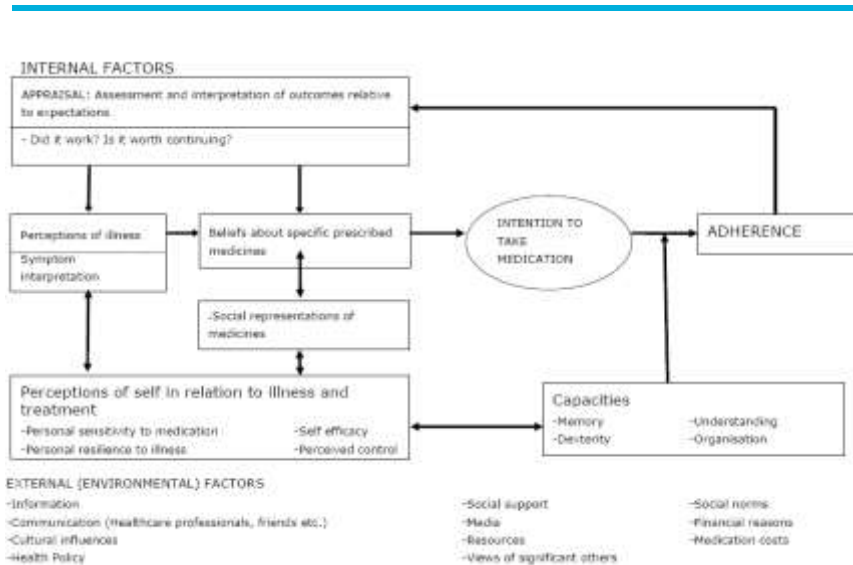


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Patient empowerment - adherence

- Patient empowerment
 - Patients are self-determining
 - Some control over their own health(care)
 - Not passive recipients of healthcare
- Adherence
 - Patients voluntarily agree with a healthcare plan
 - Not submitting to their healthcare provider (=compliance)
- Patient empowerment \approx Adherence

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Horne R. NCCSDO 2005

Interventions

- Of 17 high quality RCT, only 5 improved adherence and clinical outcome¹
 - Interventions are complex
 - Frequent interactions with patients
 - Limited effective → 'no evidence that low adherence can be cured'
- Studied interventions
 - Education, training, special pill containers, counseling, reminders, self monitoring of adherence and blood pressure, support groups, feedback and reinforcement, bi-weekly contacts: ↑adherence, ↓SBP (at 6 months)²
 - Counseling by hospital pharmacist: ↑adherence (questionnaire), ↑ controlled BP³
 - SMS service: ↑adherence (questionnaire), ↑viral suppression⁴

¹ Nieuwlaat R. Cochrane 2014

² Haynes RB, Lancet 1976

³ Morgado M. International Journal of Clinical Pharmacy 2011

⁴ Lester RT. Lancet 2010

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What about e-health interventions?

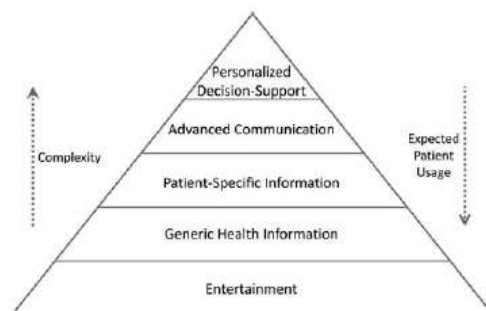
- 16 Studies since 2012: 14 websites, 1 app¹
 - 10 increased adherence
 - Combination of interventions targeting intentional and non-intentional adherence → creating patient empowerment
 - 7 applied personalized goals, including feed-back
 - 2 studied accessing patient's file
 - 1 studied personalized SMS

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¹Sieben A. Ned Tijdschr Geneeskd 2014

Other ways to empower patients?

- Patient empowerment has been conducted in the outpatient setting
- Historically limited to inpatient settings
- Systematic review (17 articles)¹
 - 3 design requirements for inpatient patient engagement
 - Remaining 14, grouped into five categories



¹Prey JE, JAMIA 2014

Self-administration of medication

- Self-administration of medication (SAM)
 - 'A 'transfer of responsibility' which should be dependent on a patient's ability to manage the tasks involved, as well as giving their consent to do so'¹
 - May increase
 - Patient knowledge
 - Patient's adherence
 - Patient satisfaction
 - Role for the hospital pharmacist?!

¹Royal Pharmaceutical Society, 2005

Why should we?

- Medication reconciliation errors at discharge¹
 - 69% no understanding of re-dosed medication
 - 82% no understanding of stopped medication
 - 62% no understanding of new medication
- Adherence²
 - 2-4 weeks after discharge: 55% non-adherent
 - 3 months after discharge: 70% non-adherent
 - Approx. 25% understood reasons for medication

¹Ziaeiian B, J Gen Intern Med 2005

²Pacina L, Drugs Aging 2014

Self-administration of medication

- Knowledge on drug name, purpose, appearance, dosage, frequency, and side-effects
 - Limited effect of SAM on patient knowledge
 - Knowledge on side effects was least known
- Adherence (pill count and questionnaire)
 - Limited effect of SAM on adherence
- Patient satisfaction (questionnaire or interview)
 - Positive responses
 - SAM should be continued following its evaluation
- Success
 - Pt who were successful shorter length of stay and fewer re-admissions

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Richardson SJ, PLoS One 2014

Conclusion

- Effectiveness of patient empowerment on adherence is limited by
 - Complexity of interventions applied
 - One-size does not fit all
- Self-administration of medication may be a tool to engage patients during hospitalization
 - Effects on adherence after discharge are limited



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Answers to the questions

- 1. Patient empowerment is another term for adherence
- 2. Poor adherence cannot be cured, even not by patient empowerment

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