Effectiveness of a smoking cessation intervention for people with severe mental illness treated in ambulatory mental healthcare in the Netherlands: the KISMET trial

Müge H. Küçükaksu MSc¹, Trynke Hoekstra PhD¹, Lola Jansen MSc¹, Sanne Helmig MSc², Marcel C. Adriaanse PhD¹, Berno van Meijel PhD^{3,4,5}

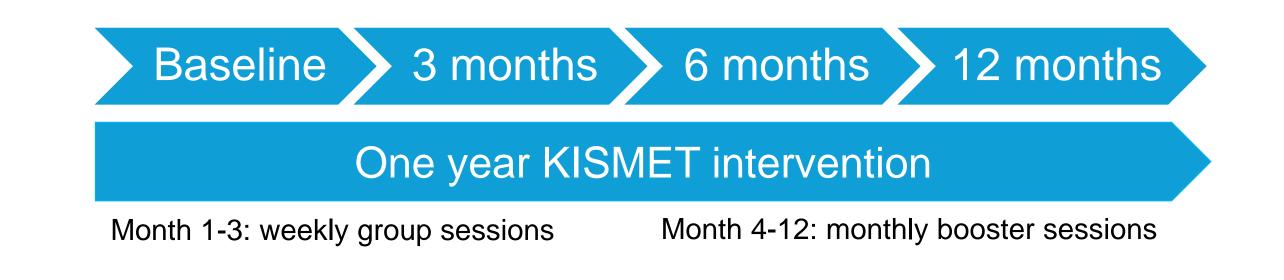
- ¹Department of Health Sciences and Amsterdam Public Health research institute, Vrije Universiteit Amsterdam, Netherlands
- ²Center for Prevention, Lifestyle and Health, Department Behavior and Health, National Institute for Public Health and the Environment, Bilthoven, Netherlands
- ³Department of Psychiatry, Amsterdam UMC and Amsterdam Public Health research institute, Amsterdam, Netherlands
- ⁴Inholland University of Applied Sciences, Centre of Expertise Prevention in Health and Social Care, Faculty of Health, Sports and Social Work, Amsterdam, The Netherlands ⁵Parnassia Psychiatric Institute, The Hague, Netherlands

Background

- Smoking is highly prevalent among people with severe mental illness (SMI): around 80% of people with SMI
- Smoking rates 2-3 times higher than in general population
- Up to 25 life years lost due to associated diseases
- Tobacco smoking is an important lifestyle factor contributing to poorer physical and mental health
- Evidence for effective smoking cessation (SC) interventions is scarce

Methods

- Research design: cluster-randomised controlled trial carried out in 21 ambulatory mental healthcare teams
- **Inclusion:** diagnosis SMI, current smoker, >18 years old, motivation to quit
- Primary outcomes: biochemically verified smoking status (through carbon monoxide) at every visit
- **Secondary outcomes include:**
- * Depression and anxiety with the Hospital Anxiety and Depression Scale (HADS) at baseline, 6and 12-months
- * Psychotic symptoms with the Positive and Negative Syndrome Scale (PANSS-6) at every visit
- Data collection and intervention timeline



Research Objective

To examine the **effectiveness** of a one-year smoking cessation intervention in ambulatory mental healthcare facilities for people with severe mental illness

The KISMET intervention

- Developed in collaboration with an expert panel applying the Delphi method
- Combined treatment consisting of three components:
- Cognitive behavioural group counselling led by two mental healthcare professionals (e.g. clinical nurse specialist, psychologist)
- Pharmacological treatment supervised by a psychiatrist (options: nicotine replacement therapy, Cytisine, Bupropion, Nortriptyline)
- III. Peer support group led by an expert-by-experience

Baseline demographics and clinical characteristics

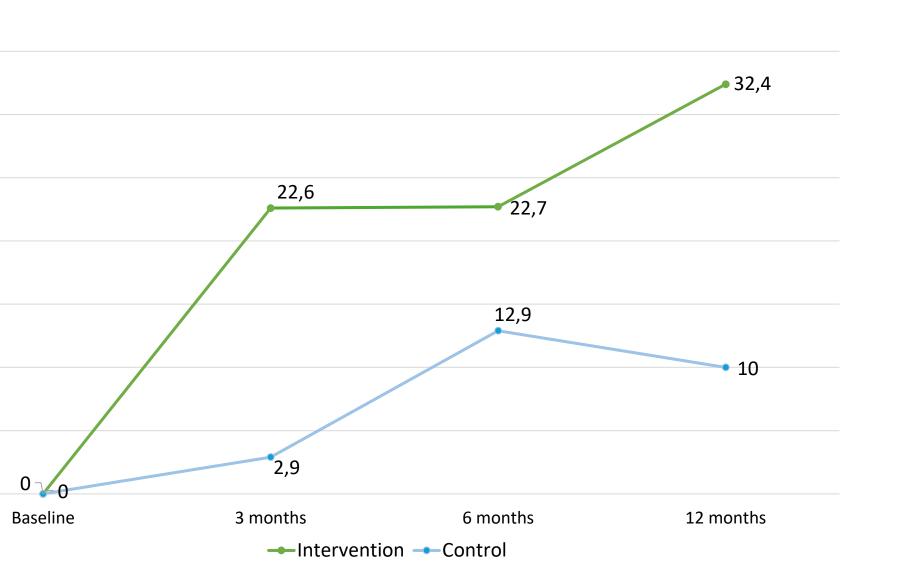
Table 1. Baseline demographics and clinical characteristics

	Intervention group	TAU
	(N=74)	(N=42)
Demographic characteristics		
Age in years (mean (SD))	47.2 (11.4)	46.3 (12.1)
Gender, no. (%)		
Woman	39 (52.7)	15 (35.7)
Man	35 (47.3)	27 (64.3)
Partner, no. (%)	12 (16.2)	11 (26.2)
No partner, no (%)	62 (83.8)	31 (73.8)
Clinical characteristics		
Primary DSM-5 diagnosis, no (%)		
Schizophrenia spectrum	45 (59.2)	16 (39.0)
Bipolar disorder	8 (10.5)	1 (2.4)
Depressive or anxiety disorder	12 (15.8)	11 (26.8)
Personality disorder	5 (6.6)	7 (17.1)
Autism spectrum	2 (2.6)	2 (4.9)
Other psychiatric disorders*	4 (5.3)	4 (9.8)
Body mass index (mean, (SD))	28.9 (7.5)	27.9 (5.4)
Number of cigarettes per day	19 (10.7)	17 (9.4)
(mean, (SD))		
HADS anxiety scale	9.1 (4.9)	8.2 (5.5)
HADS depression scale	7.6 (4.5)	7.2 (5.1)
PANSS-6 psychotic symptoms	13.2 (6.6)	10.8 (5.0)
Use of antipsychotic medication, no (%)	46 (51.7)	26 (59.1)
Use of antidepressive medication, no (%)	51 (57.3)	32 (72.7)

Key preliminary findings 1: Positive effect of KISMET intervention on smoking behaviour

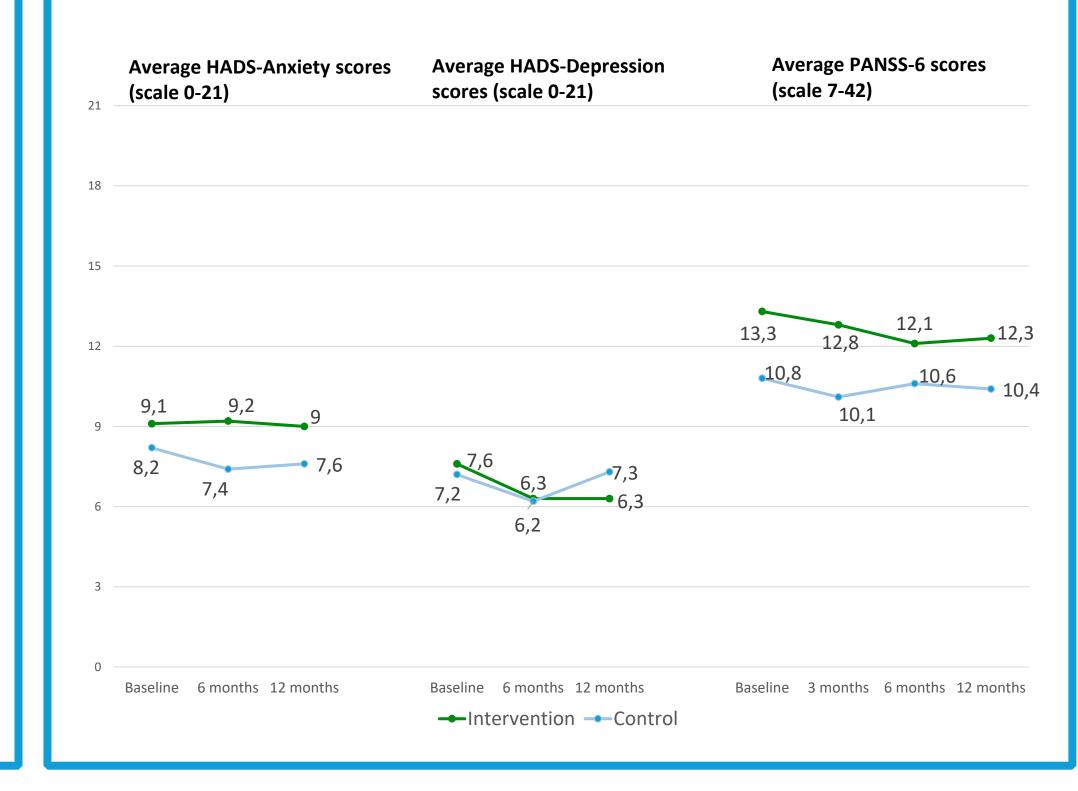
- **32.4% of patients** in the intervention group (n=37) quit smoking at 12 months compared to 10.0 % in the TAU group (n=30)
- Differences between groups not significant
- **Drop-out:** 52 patients (58.4%) in intervention group and 14 patients (31.8%) in the TAU group

Smoking cessation %



Key preliminary findings 2: No differences in mental health outcomes

No significant differences in depression, anxiety and psychotic symptoms between baseline and 12 months between groups



Conclusions and clinical implications

- Percentage of smoking cessation at 12 months favours the KISMET intervention
- No detrimental effects on depression, anxiety and psychotic symptoms of the KISMET intervention or smoking cessation within 12 months
- The KISMET intervention can be offered safely to people with severe mental illness
- Strategies to improve inclusion and reduce study attrition are needed









