Short Research Article

2 Title: A Randomized, Double-Blind, Placebo-Controlled Study of a Blend of Herbal Extracts 3 Taken Once Per Day for Weight Loss in Healthy Volunteers 4 5 **ABSTRACT** 6 Aims: We previously demonstrated that a blend of herbal extracts (Weighlevel®; a mixture of 7 extracts from the leaves of Alchemilla vulgaris, Olea europaea, Mentha longifolia and from the 8 seeds of Cuminum cyminum) taken 3 times per day produces weight loss in preclinical and 9 clinical studies. The aim of the present study was to test the efficacy of a new slow-release 10 formulation (Weighlevel® One) taken once per day on change in body weight and related 11 measures. 12 **Study design:** Randomized, double-blind, placebo-controlled study. 13 Place and Duration of Study: Health Clinics in Copenhagen, Denmark between 7 January 14 2016 and 5 March 2016. **Methodology:** Thirty-six adult subjects were randomized to consume the herbal blend (n = 20)15 16 or placebo (n = 16) once per day for 8 weeks. Weight and waist circumference were assessed 17 weekly. The primary endpoint was the change from baseline in body weight for the herbal blend 18 group compared with placebo. Secondary assessments included waist circumference, appetite. 19 craving, bowel health, and safety and tolerability. 20 **Results:** After 8 weeks, the herbal blend group lost an average of 3.7 kg (95% CI of 3.0 to 4.5 21 kg); whereas the placebo group lost 0.1 kg (95% CI of -0.7 to 1.0 kg). This difference in mean 22 weight loss between the herbal blend and placebo groups was statistically significant (P < .001).

A statistically significant reduction in waist circumference was also observed for the herbal blend

compared with placebo (P < .001). The herbal blend was well tolerated; no adverse events were

1

23

24

25

reported.

26 Conclusion: Daily administration of this blend of herbal extracts, administered once daily, may 27 produce weight loss. 28 29 Keywords: Alchemilla vulgaris, Olea europea, Mentha longifolia, Cuminum cyminum, overweight, obesity, slow release 30 31 32 1. INTRODUCTION 33 Obesity rates worldwide have nearly tripled over the last four decades [1]. The World Health 34 Organization estimates that nearly 2 billion adults are currently overweight or obese [1]. 35 Sustained weight loss of as little as 3% to 5% can produce clinically meaningful reductions in 36 cardiometabolic risk factors such as blood glucose and lipids, with larger weight losses producing greater benefits [2]. However, many people struggle to lose weight through diet and 37 38 exercise alone. 39 Some plants used in traditional Greco-Arab and Islamic medicine have properties which may aid 40 41 in weight loss. Alchemilla vulgaris (Lady's mantle) is used in traditional Arabic medicine for 42 weight loss and to treat stomach and intestinal pain [3]. It also has anti-inflammatory properties 43 [4]. Olea europea (olive) improves insulin sensitivity and may reduce blood pressure and 44 plasma lipids [5-7]. Mentha longifolia (wild mint) is traditionally used to treat gastrointestinal 45 disorders and also has antimicrobial properties [8]. Supplementation with Cuminum cyminum 46 (cumin) has been reported to produce weight loss in overweight subjects [9]. 47 48 We previously demonstrated that a blend of herbal extracts (Weighlevel®; a mixture of extracts 49 from the leaves of Alchemilla vulgaris, Olea europaea, and Mentha longifolia and from the 50 seeds of Cuminum cyminum) taken 3 times per day is effective in producing weight loss in both 51 preclinical and clinical studies [10, 11]. However, the rate of medication compliance tends to

decrease as the number of daily doses increases. A systematic review found that medication compliance dropped to 65% for medications taken 3 times per day, and compliance was significantly higher for once daily regimens [12]. This suggests that patients would be more likely to consume the herbal blend as instructed if the dosing regimen were once daily rather than 3 times per day.

The aim of the present study was to test the efficacy of a new slow-release formulation (Weighlevel® One) taken once per day on the change in body weight and other weight-related measures in a randomized, double-blind, placebo-controlled study. We hypothesized that the new herbal blend formulation would produce incremental and sustained weight loss over the course of the 8-week study.

2. MATERIAL AND METHODS

2.1 Participants

Participants were recruited by a specially trained qualified nurse (Erla Øregaard, who is a recognized as a specialist in patient safety by the Danish Health Authority) from Health Clinics in Copenhagen, Denmark between 7 January 2016 and 5 March 2016. Eligible participants were generally healthy, not pregnant, were unsatisfied with their current weight, interested in losing weight, and agreed to follow the study protocol.

2.2 Study Design

73

74

75

76

77

78

79

80

81

82

83

84

85

86

87

88

This study was a randomized, double-blind, placebo-controlled trial. Participants were randomized to receive either 1 tablet of herbal blend or 1 tablet of placebo per day for 8 weeks. Herbal blend and placebo tablets were provided free of charge. The herbal blend (Weighlevel®One) is a mixture of extracts from the leaves of *Alchemilla vulgaris*, *Olea* europaea, Mentha longifolia and from the seeds of Cuminum cyminum and a patented slow release component (Propol®). These ingredients are Generally Regarded as Safe (GRAS). The matching placebo and herbal blend each had a net weight of 0.88 grams and were manufactured by ProPharma (Copenhagen). Treatment allocation was concealed, and blinding was maintained by administering the herbal blend or placebo in coded containers. The participants, recruiting nurse, and investigators remained blinded during the study. The primary endpoint was the change in body weight from baseline to endpoint for the herbal blend compared with placebo. Secondary endpoints were the change in waist circumference and Visual Analog Scale (VAS) assessments from baseline to endpoint for the herbal blend and placebo. Safety and tolerability were also evaluated through reporting of adverse events.

89

90

91

92

93

94

Participants were instructed to take 1 tablet in the morning and to maintain their daily eating or exercise routine. Body weight and waist circumference measurements were obtained at the same time and week-day throughout the study. Weekly follow-up reports were conducted online. In person visits occurred at baseline and at Weeks 4 and 8. Adverse events were assessed throughout the study.

95

96

97

98

2.3 Visual Analog Scale Assessments

Appetite, craving for sweets, and bowel health were assessed using VAS which have been reported to be reliable in appetite research [13]. Each visual analog scale consisted of a clear

unmarked plastic strip. Participants were asked to place their finger on the strip in response to the following questions: How hungry are you today? (appetite), How much have you been craving sugar/sweets today? (craving), and Are you having bowel movements daily/how does your bowel feel? (bowel health). Each plastic strip was then given a numerical rating from 1 to 5 by the investigator in which higher numbers indicated an improvement: Appetite (1=hungry, 5=low appetite), Craving (1=craving sugar, 5= no craving), Bowel Health (1=infrequent bowel movements, uncomfortable bowel, 5=bowel ok).

2.4 Statistical Analysis

All results reported are for the Intent-to-Treat Population. Missing values for the 4 subjects that dropped out of the study early are accounted for by maximum likelihood, using a missing at random (MAR) assumption. A repeated-measures mixed-effects model was used to compare changes from baseline in treatment and control groups. A random subject effect and fixed treatment, baseline value and time effects were included in the model.

3. RESULTS

3.1 Participants

A flow diagram of participant disposition is shown in Figure 1. A total of 50 volunteers were assessed for eligibility. Reasons for exclusion from the study were: did not meet inclusion criteria (n=14) and declined to participate (n=4). Thirty-six participants were randomized to receive either 1 tablet of herbal blend (n=20) or 1 tablet of placebo (n=16) per day for 8 weeks. Four participants dropped out after randomization and before the first weekly online follow-up visit. Demographics and baseline characteristics were similar for the 2 treatment groups (see Table 1).

Figure 1. Study Flow Diagram

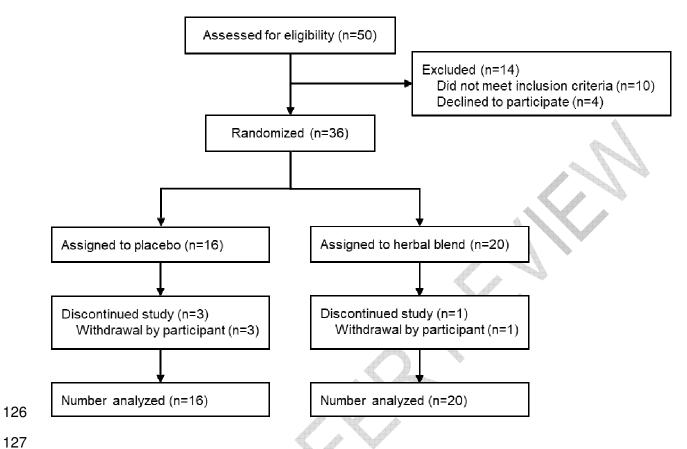


Table 1. Demographics and baseline characteristics

	Placebo	Herbal Blend
	(N=16)	(N=20)
Age (years)	47.1 (9.8)	45.3 (9.1)
Sex, Female	14 (87.5%)	19 (95.0%)
Weight (kg)	79.2 (10.0)	79.1 (15.2)
Height (cm)	171.9 (6.2)	170.8 (7.9)
BMI (kg/m²)	26.8 (3.5)	27.1 (4.4)
Waist circumference (cm)	91.4 (8.6)	93.4 (12.1)

^{129 *} Data are mean (SD) or n (%)

3.2 Body Weight and Waist Circumference

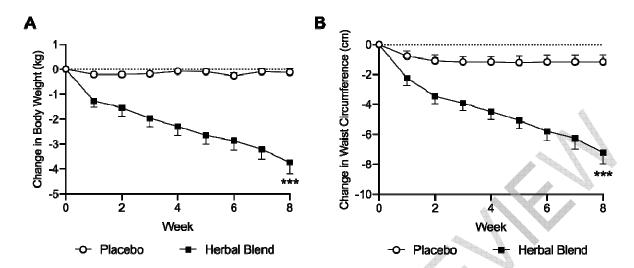
After 8 weeks, the herbal blend group lost an average of 3.7 kg (95% CI: 3.0 to 4.5 kg; P<.001) compared with 0.1 kg lost for the placebo group (95% CI: -0.7 to 1.0 kg) (Table 2). This was equivalent to a 4.7% reduction in weight in the group that received the herbal blend, compared to a 0.2% reduction in the placebo group (Table 2, P<.001). The group that was treated with the herbal blend demonstrated continued weight loss that was sustained for the duration of the 8-week study (Figure 2A), with mean weight loss of 0.46 kg per week. In contrast, mean weight loss in the placebo group was generally unchanged from baseline. Waist circumference was also reduced by 7.2 cm in the herbal blend group compared with a reduction of 1.2 cm in the placebo group (Figure 2B and Table 2, P<.001).

Table 2. Change from baseline to Week 8 in weight, waist circumference, and visual
 analog scale ratings

Change	Placebo	Herbal Blend	LS Mean	P value
	(N=16)	(N=20)	Difference	
Body Weight (kg)	-0.1 (0.4)	-3.7 (0.4)	-3.6 (0.6)	P <.001
95% CI	[-1.0, 0.7]	[-4.5, -3.0]	[-4.7, -2.5]	
Body Weight (%)	-0.2 (0.4)	-4.7 (0.3)	-4.5 (0.5)	P <.001
95% CI	[-1.0, 0.7]	[-5.7, -4.0]	[-5.7, 3.4]	
Waist Circumference (cm)	-1.2 (0.8)	-7.2 (0.6)	-6.1 (1.0)	<i>P</i> <.001
95% CI	[-2.7, 0.4]	[-8.5, -5.9]	[-8.1, -4.0]	
VAS Appetite	-0.4 (0.2)	0.6 (0.1)	1.0 (0.2)	P <.001
95% CI	[-0.7, 0.0]	[0.3, 0.9]	[0.5, 1.4]	
VAS Cravings	-0.2 (0.3)	0.9 (0.2)	1.2 (0.3)	P<.001
95% CI	[-0.8, 0.3]	[0.5, 1.4]	[0.5, 1.9]	
VAS Bowel Health	-0.5 (0.3)	0.3 (0.3)	0.9 (0.4)	P=.002
95% CI	[-1.2, 0.1]	[-0.2, 0.9]	[0.0, 1.7]	

^{*} Data are least-squares mean (SE) and 95% confidence interval. VAS = visual analog scale.

Figure 2. Herbal blend reduced body weight and waist circumference

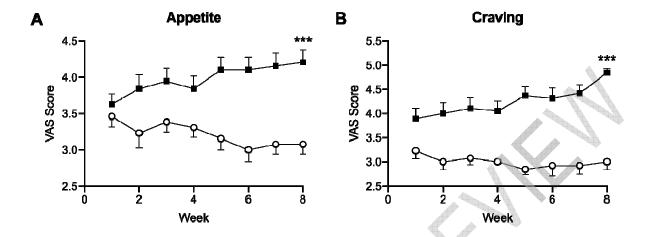


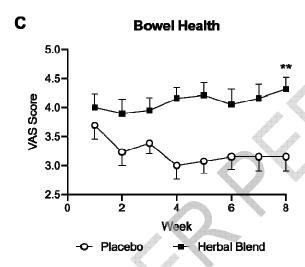
A) Change in body weight. Mean baseline body weight was 77.9 kg for the placebo group and 78.1 kg for the herbal blend group. B) Change in waist circumference. Mean baseline waist circumference was 91.2 cm for the placebo group and 93.0 cm for the herbal blend group. Data are mean ±SE for available data (placebo n=13, herbal blend n=19). ***P <.001 compared to placebo.

3.3 Visual Analog Scale Assessments

Statistically significant improvements in patient-reported appetite, craving for sweets, and bowel health were observed compared to placebo at Week 8 (all P < .01, Table 2 and Figure 3). These improvements in patient-reported appetite and craving for sweets in the herbal blend group continued for the duration of the study whereas there was a slight worsening in the placebo group (Table 2 and Figure 3A and B). Bowel health also appeared to worsen in the placebo group, compared to an improvement in the herbal blend group (Table 2 and Figure 3C).

Figure 3. Herbal blend improves subjective ratings of appetite, craving and bowel health





Increases in score demonstrate improvements. A) Visual analog scale rating of appetite (1=hungry, 5=low appetite). B) Visual analog scale rating of craving for sweets (1=craving sugar, 5= no craving). C) Visual analog scale rating of bowel health (1=infrequent bowel movements, uncomfortable bowel, 5=bowel ok). Data are mean \pm SE for available data (placebo n=13, herbal blend n=19). **P<.01, ***P<.001 compared to placebo. VAS = visual analog scale.

3.4 Safety

The herbal blend was well tolerated. No adverse events or changes in wellbeing were reported.

4. DISCUSSION

The results presented here demonstrate that the slow-release herbal blend taken once per day produced statistically significant weight loss in healthy adults. After 8 weeks of treatment, participants in the herbal blend group lost an average of 4.7% of their baseline body weight compared with 0.2% weight loss in the placebo group. The herbal blend was also well tolerated and there were no safety concerns. This is in agreement with previous studies where the herbal blend was administered 3 times per day and produced weight loss of approximately 7% after 2 months of treatment and 10-13% weight loss after 3 months [10, 11]. The weight loss observed with the once per day formulation was slightly lower than previously observed with the 3 times per day formulation. This difference may be related to the higher mean baseline body weight of participants in previous studies. It may also be related to differences between populations where previous studies were performed (Galilee, Israel) and where the current study was performed (Copenhagen, Denmark). However, weight loss continued for the duration of the 8-week study and there was no evidence of a plateau, suggesting that additional weight loss may be possible with continued administration of the herbal blend.

In addition to weight loss, there was a statistically significant reduction in waist circumference that corresponded with the reduction in body weight in participants who received the herbal blend compared with placebo. Whether the reductions in body weight and waist circumference (both measures of the metabolic syndrome) reflect an improvement in other weight-related comorbidities, such as lipids, blood pressure, and blood glucose remains to be determined. However, the reduction in body weight approaches a 5% reduction from baseline, which has been determined to represent clinically meaningful weight loss that reduces the incidence of diabetes, reduces blood pressure, and improves lipids [14]. Thus, improvements in cardiometabolic markers may be due to weight loss as well as to independent effects of

components of the herbal blend. For example, olive leaf extracts have been shown to inhibit intestinal glucose absorption and improve blood pressure, lipids, and markers of inflammation [5-7] while cumin has been demonstrated to reduce elevated blood glucose by improving glucose utilization [15].

The mechanism for weight loss with the herbal blend is hypothesized to be attributed to multiple effects including increased thermogenesis resulting in fat depletion, reduced blood glucose, and beneficial change in digestion. Increased thermogenesis with the herbal blend has been demonstrated in Sprague-Dawley rats [11]. This is consistent with the reports of metabolic stimulation with extracts of *Alchemilla vulgaris* L. [16] and olive leaf [17]. Olive leaf and cumin have been shown to improve blood glucose by inhibiting intestinal glucose absorption [5, 6] and improving glucose utilization [15]. *Alchemilla vulgaris* L. and cumin have been shown to regulate digestive enzymes [18, 19] while mint has been demonstrated to increase gastric emptying and passage of food through the gastrointestinal tract [20].

This study demonstrates that weight loss with the herbal blend is also likely attributed to a reduction in caloric intake. Participants who received the herbal blend reported improvements in appetite, craving, and bowel health, measured by VAS, which has demonstrated efficacy in assessing appetite [13]. The reductions in appetite and craving with the herbal supplement were observed in the context of weight loss and continued for the duration of the study and may reflect reduced caloric intake. The improvement in bowel health is consistent with the improvements in digestion demonstrated by *Alchemilla vulgaris* and mint [18-20] and is notable given that fecal incontinence is common in individuals with obesity [21].

There were several limitations to this study. Because this was an 8 week study with a relatively small sample size, the extent of weight loss with the herbal blend is unclear, although results

228 are consistent with previous studies [10, 11]. In addition, improvements in appetite and craving 229 indicate that reduced caloric intake may contribute to weight loss with the herbal blend. 230 However, caloric intake was not assessed so the contribution of reduced caloric intake to weight 231 loss has yet to be determined. Finally, the effects of the herbal blend of cardiometabolic 232 markers, such as lipids, blood pressure, and blood glucose were not assessed. The effects of the herbal blend on markers of cardiometabolic risk requires further study. 233 234 235 5. CONCLUSION 236 In summary, the 4.7% weight loss in participants treated with the herbal blend was statistically 237 significant and well within the range that would be expected to produce beneficial effects on 238 markers of cardiometabolic risk [2]. The ease of use of a once per day formulation is expected 239 to improve adherence and provide meaningful improvements in weight-related health. 240 241 242 Consent 243 Informed and written consent was obtained from all participants prior to participation in the 244 study. 245 246 **Ethical approval** 247 All experiments were examined and approved by the appropriate ethics committee represented 248 by professor, dr. med Steen Lindkær-Jensen (Aarhus University, Aarhus, Denmark and Imperial 249 College, London, UK), and were examined and performed in accordance with the ethical 250 standards laid down in the 1964 Declaration of Helsinki. 251 252

253

COMPETING INTERESTS DISCLAIMER:

Authors have declared that no competing interests exist. The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

260

261

254

255

256

257

258

259

262 References

- World Health Organization. *Obesity and overweight*. 2018 Feb 16, 2018 [cited 2019 Feb
 12]; Available from: https://www.who.int/news-room/fact-sheets/detail/obesity-and-overweight.
- Jensen, M.D., et al., 2013 AHA/ACC/TOS guideline for the management of overweight
 and obesity in adults: a report of the American College of Cardiology/American Heart
 Association Task Force on Practice Guidelines and The Obesity Society. J Am Coll
 Cardiol, 2014. 63(25 Pt B): p. 2985-3023.
- Said, O., et al., Ethnopharmacological survey of medicinal herbs in Israel, the Golan
 Heights and the West Bank region. J Ethnopharmacol, 2002. 83(3): p. 251-65.
- 272 4. Schink, A., et al., *Screening of herbal extracts for TLR2- and TLR4-dependent anti-*273 *inflammatory effects.* PLoS One, 2018. **13**(10): p. e0203907.
- Wainstein, J., et al., Olive leaf extract as a hypoglycemic agent in both human diabetic
 subjects and in rats. J Med Food, 2012. 15(7): p. 605-10.
- de Bock, M., et al., Olive (Olea europaea L.) leaf polyphenols improve insulin sensitivity
 in middle-aged overweight men: a randomized, placebo-controlled, crossover trial. PLoS
 One, 2013. **8**(3): p. e57622.

- 279 7. Lockyer, S., et al., Impact of phenolic-rich olive leaf extract on blood pressure, plasma
- 280 *lipids and inflammatory markers: a randomised controlled trial.* Eur J Nutr, 2017. **56**(4):
- 281 p. 1421-1432.
- 8. Mikaili, P., et al., *Pharmacological and therapeutic effects of Mentha Longifolia L. and its*
- 283 *main constituent, menthol.* Anc Sci Life, 2013. **33**(2): p. 131-8.
- 284 9. Taghizadeh, M., et al., Effect of the cumin cyminum L. Intake on Weight Loss, Metabolic
- 285 Profiles and Biomarkers of Oxidative Stress in Overweight Subjects: A Randomized
- 286 Double-Blind Placebo-Controlled Clinical Trial. Ann Nutr Metab, 2015. 66(2-3): p. 117-
- 287 24.
- 288 10. Said, O., et al., A double blinded- randomized clinical study with "weighlevel", a
- combination of four medicinal plants used in traditional arabic and islamic medicine. The
- 290 Open Complementary Medicine Journal, 2010. 2: p. 1-6.
- 291 11. Said, O., et al., Weight loss in animals and humans treated with "weighlevel", a
- combination of four medicinal plants used in traditional arabic and islamic medicine. Evid
- 293 Based Complement Alternat Med, 2011. **2011**: p. 874538.
- 294 12. Claxton, A.J., J. Cramer, and C. Pierce, A systematic review of the associations between
- 295 dose regimens and medication compliance. Clin Ther, 2001. **23**(8): p. 1296-310.
- 296 13. Flint, A., et al., Reproducibility, power and validity of visual analogue scales in
- 297 assessment of appetite sensations in single test meal studies. Int J Obes Relat Metab
- 298 Disord, 2000. **24**(1): p. 38-48.
- 299 14. Williamson, D.A., G.A. Bray, and D.H. Ryan, *Is 5% weight loss a satisfactory criterion to*
- 300 define clinically significant weight loss? Obesity (Silver Spring), 2015. 23(12): p. 2319-
- 301 20.
- 302 15. Roman-Ramos, R., J.L. Flores-Saenz, and F.J. Alarcon-Aguilar, *Anti-hyperglycemic*
- effect of some edible plants. J Ethnopharmacol, 1995. **48**(1): p. 25-32.

304 16. Borodin Iu, I., et al., [Effect of polyphenol fraction from Alchemilla vulgaris on the 305 morphofunctional state of the thyroid in rats exposed to cold]. Biull Eksp Biol Med, 1999. 306 **127**(6): p. 697-9. 307 17. Al-Qarawi, A.A., M.A. Al-Damegh, and S.A. ElMougy, Effect of freeze dried extract of 308 Olea europaea on the pituitary-thyroid axis in rats. Phytother Res, 2002. 16(3): p. 286-7. 309 18. Jonadet, M., et al., [Flavonoids extracted from Ribes nigrum L. and Alchemilla vulgaris 310 L.: 1. In vitro inhibitory activities on elastase, trypsin and chymotrypsin. 2. 311 Angioprotective activities compared in vivol. J Pharmacol, 1986. 17(1): p. 21-7. 312 19. Platel, K. and K. Srinivasan, Influence of dietary spices and their active principles on 313 pancreatic digestive enzymes in albino rats. Nahrung, 2000. 44(1): p. 42-6. 314 20. Spirling, L.I. and I.R. Daniels, *Botanical perspectives on health peppermint: more than* 315 just an after-dinner mint. J R Soc Promot Health, 2001. 121(1): p. 62-3. 316 21. Pares, D., et al., Bowel habits and fecal incontinence in patients with obesity undergoing 317 evaluation for weight loss: the importance of stool consistency. Dis Colon Rectum, 2012. 318 **55**(5): p. 599-604.