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# Effects of Education and Experience on Primary Care Providers' Perspectives of Obesity Treatments during a Pragmatic Trial

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#### **Abstract**

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**Objective**—To examine the impact of a one-year pragmatic obesity trial on primary care providers' (PCPs) perspectives of treatment.

**Methods**—PCPs from four intervention (PCP-I) and five control clinics (PCP-C) completed preand post-intervention surveys on weight loss counseling, comfort discussing obesity treatments and perceived effectiveness of interventions; questions were rated on 0–10 Likert scales. Only PCP-I received patient updates and education about obesity management.

**Results**—Eighty PCPs completed pre-intervention surveys [pre] (71% female, 71% physicians); 82 PCPs completed post-intervention surveys [post] (66% female, 70% physicians). PCPs were most comfortable discussing exercise (median 8, interquartile range 7–9), even after the trial (P=0.71). PCPs were least comfortable discussing phentermine/topiramate ER (4, 2–6), but developed more comfort (pre 3, 1.5–6; post 5, 3–7; P<0.001). Only PCP-I became more comfortable discussing phentermine (pre 7, 4–8; post 8, 7–9; P=0.026). After the trial, PCPs rated phentermine/topiramate ER more effective (pre 5, 3–6; post 7, 5–8; P<0.001); only PCP-I rated exercise less effective (pre 7, 4–8.5; post 5, 3–7; P=0.035) and phentermine more effective (pre 5, 5–7; post 7, 6–8; P<0.001).

**Conclusions**—PCPs initially overvalued exercise and undervalued medications. PCPs exposed to education and experience gave higher comfort and effectiveness ratings to weight loss medications.

#### Keywords

Obesity; Obesity Treatment; Primary Care; Exercise; Pharmacologic Therapy

#### INTRODUCTION

Obesity affects one-third of U.S. adults and increases the risk for common chronic medical conditions.(1–3) Forecasts predict the prevalence of obesity will increase by 33% and that of severe obesity by 130% between 2010 and 2030.(4) Additionally, obesity has increased medical costs, particularly in the Medicare and Medicaid populations.(5, 6) Despite recognition of obesity as a national epidemic, less than half of patients with obesity reported receiving weight loss counseling even when they had obesity-related comorbidities.(7, 8) Specifically, weight loss counseling by primary care providers (PCPs) is not increasing and may be declining, and most PCPs provide no weight loss counseling at all.(9) Studies have identified several barriers to providing weight loss counseling including lack of time, lack of comfort in discussing treatments and a lack of reimbursement.(10, 11) Despite these challenges, stakeholders continue to encourage PCPs to take an active role in obesity management.(12, 13)

Although very little structured obesity treatment is delivered within primary care, effective tools to facilitate weight loss exist. Meta-analyses have demonstrated that meal replacements(14) and group behavioral programs(15) produce clinically significant weight loss in real-world settings. Anti-obesity medications are also effective.(16) However, prescription rates are low due to concerns over safety, modest weight loss and questions surrounding the appropriateness of pharmacological treatment for obesity.(17) Low prescribing rates may lead to PCPs' discomfort in discussing medications due to a lack of

knowledge about and experience with the relative effectiveness of medications compared to more frequently suggested modalities like dietary change and exercise.

Since obesity disproportionately effects racial and socioeconomic minorities (18–20) and non-white survey respondents more likely want weight-related discussions with their providers, (21) we assessed the obesity treatment-related perceptions of PCPs at urban clinics serving culturally diverse patients within an integrated safety net health care institution. As part of a one-year pragmatic clinical trial using a "toolbox" approach to weight loss management, (22) we provided education to PCPs and offered intervention patients the ability to choose from several non-surgical obesity treatment modalities at low out-of-pocket costs. We sought to determine the impact of our pragmatic trial on those patients' PCPs regarding their 1) views on obesity and weight loss counseling, 2) comfort discussing non-surgical obesity treatments and 3) perceived effectiveness of those tools. The findings extended our understanding of current views and barriers toward obesity management within primary care.

## **METHODS**

### Study Design

We conducted cross-sectional surveys of PCPs at nine primary care clinics affiliated with Denver Health and Hospital Authority (DH) before the initiation and after the completion of a pragmatic clinical trial: A Toolbox Approach to Obesity Treatment in Primary Care Trial (Toolbox).(22) Briefly, Toolbox was a one-year, open-label obesity treatment trial within primary care clinics at DH, with a registry-based comparator group. Patients in the intervention cohort were provided with computer-based education at the first visit and offered tools for weight loss management at monthly clinic visits. Patients at the four intervention clinics could choose a tool from a variety of evidence-based weight loss treatments at a low out-of-pocket cost (\$5–10 copay/month): partial meal replacement regimen (i.e., protein shakes and small portion-controlled meals, meant to replace two of three self-selected meals), recreation center vouchers, obesity pharmacotherapy (phentermine or phentermine/topiramate ER), commercial weight loss program vouchers, and a clinic-based group behavioral weight loss program.

PCPs from intervention clinics (PCP-I) were provided four half-hour education sessions during the trial which were conducted by the principle investigator of the trial during regularly scheduled staff meetings. During these sessions, information was presented on the relative effectiveness of each of the weight loss treatment strategies used in the trial, details of obesity treatment guidelines, the low rates of utilization of these tools reported in the literature and strategies for discussing weight management with patients during busy office visits. The modest effectiveness of exercise as a weight loss tool and the greater effectiveness of medications was highlighted. Details of the Toolbox design were presented in the initial session and details of the study's progress and challenges were highlighted in later sessions. There was an opportunity for discussion and questions. Attendance was taken only at the first session when pre-intervention surveys were circulated. Throughout the study, PCP-I were given status updates whenever their patients enrolled in the intervention, were lost to follow up, chose or switched tools (including which tool was selected) or

achieved 5% weight loss. PCPs from the five control clinics (PCP-C), were not provided any education nor did they gain any firsthand experience with patients receiving obesity treatments through the trial.

The primary outcome of the parent trial was percentage of patients participating in the intervention who achieved 5% weight loss after one year as compared to a registry-based comparator group who did not receive the intervention.

#### Participants and procedures

**Survey Development**—A short survey was developed to assess PCPs' perspectives regarding the significance of obesity as a problem for their patients, barriers to providing weight loss advice, comfort discussing various weight loss tools and the perceived effectiveness of those same weight loss interventions. The survey incorporated a number of items from a questionnaire previously developed and validated by investigators from The Johns Hopkins University School of Medicine and Harvard T.H. Chan School of Public Health (KG, SB, JC),(11, 23) Surveys were reviewed and approved by the Colorado Multiple Institutional Review Board (see Supporting Information, Surveys S1 and S2, for pre- and post-intervention questionnaires).

**Setting, Participants and Survey Implementation**—DH is an integrated safety net health care system serving a low socioeconomic, ethnically diverse and medically underserved population in Colorado. As of 2014, DH served a population consisting of roughly two-thirds racial/ethnic minorities including 43% Hispanic, 14% African American and 6% either Asian American, Native American, or multi-racial.

The participants for this study were PCPs who worked at the nine primary care clinics affiliated with DH, geographically distributed around the greater Denver metropolitan area. Clinics varied markedly by patient volume, the demographics of the populations served, the total number of providers at each site and the relative number of nurse practitioners (NP)/ physician assistants (PA-C)/family medicine physicians/internal medicine physicians. Clinics were randomized to either intervention (four clinics, PCP-I) or control (five clinics at baseline and included in the analyses, PCP-C). Pre-intervention surveys were distributed, completed and collected at standing monthly team meetings held at each clinic site between December 2013 and July 2014. During the post-intervention period between September 2016 and December 2016, providers were notified that the trial concluded and surveys were again distributed at standing team meetings. In an effort to survey all PCPs at clinic sites, PCPs who were not in attendance at meetings where surveys were distributed (n=9 pre-intervention, n=12 post-intervention) were contacted individually and given surveys to return to study staff.

All surveys were completed anonymously with the intention that PCPs would respond honestly to questions about weight loss interventions without fear of their answers being judged by study staff. To accomplish this, demographic data were collected separately from all PCPs who completed surveys. As a result of this strategy we were unable to correlate individual PCPs' responses on the surveys to demographic variables or to compare changes in individual PCPs' responses between pre-intervention and post-intervention surveys.

#### **Outcome Measure**

On a 0–10 Likert scale (0 being "Least comfortable" and 10 being "Most comfortable"), PCPs rated their comfort discussing individual weight loss tools with their patients: lifestyle modification programs, portion-controlled foods, exercise, phentermine and phentermine/ topiramate ER. This list of obesity interventions in the survey was similar to those offered in the Toolbox trial. PCPs also rated the effectiveness of these tools for weight loss on a 0–10 Likert scale (0 being "Least effective" and 10 being "Most effective"). Additionally, PCPs views on the importance of obesity as a problem and their comfort in general weight loss counseling were measured on similar 0–10 Likert scales.

#### Statistical Analyses

The median and interquartile range (IQR) were calculated for each item rating on both the pre-intervention and post-intervention surveys. Since ratings were ordinal and non-normally distributed, medians were used instead of means for group comparison. The Wilcoxon rank sum test was used to compare within-group changes among the PCP-I and PCP-C groups. The p-values reported were not adjusted for multiple comparisons. Analyses were conducted using SAS version 9.4.

#### **RESULTS**

## **Demographics**

The demographics for individuals completing surveys are depicted in Table 1 (80 of 85 preintervention participants, 82 of 82 post-intervention participants). The non-physician providers (n=23 pre-intervention, n=25 post-intervention) who completed surveys included NPs (n=5 pre-intervention, n=7 post-intervention), PA-Cs (n=15 pre-intervention, n=15 post-intervention) intervention), doctors of pharmacy (PharmD, n=1 pre-intervention, n=1 post-intervention), and registered nurses (RN, n=2 pre-intervention, n=1 post-intervention). Of all PCPs practicing at the clinic sites during the study, 85% (85 of 100) completed pre-intervention and 68% (82 of 121) completed post-intervention surveys. Eighty-nine percent (76 of 85) in the pre-intervention group and 85% (70 of 82) in the post-intervention group attended the initial team meetings at which the first surveys were distributed and education was delivered. The difference in the total number of PCPs between the pre- and post-intervention periods is related to the addition of new providers in support of the new clinic (non-intervention) opening during the later portion of the trial and an increase in the number of part-time providers who may not have been available for the post-intervention survey. Since surveys were anonymous and demographic data were collected separately from survey responses, we were unable to compare pre- to post-intervention survey results by individual providers to determine if differences in provider type (e.g., MD/DO vs. NP/PA-C/PharmD/RN, etc.) or attendance at educational sessions led to differential views on obesity treatment.

#### **Provider Views on Obesity and Weight Loss Counseling**

Distributions of the ratings for all survey items are displayed in Table 2. All providers identified obesity as a significant problem for their patients at both time points. Similarly, providers from both clinic groups rated their comfort discussing weight with their patients

an 8 out of 10 before and after the Toolbox study period. Providers were less optimistic that their advice had an impact or were comfortable in counseling patients on their own; these responses also did not change over time.

## **Perceived Comfort in Discussing Various Weight Loss Tools**

Distributions of comfort ratings for specific weight loss tools are depicted in Table 3 (and Supporting Information, Figure S1). PCPs from all clinics were most comfortable discussing exercise with no change in comfort ratings after the trial. Similarly, comfort ratings did not change significantly for all providers regarding lifestyle modification programs or portion-controlled foods, though these were both rated lower than exercise. Unlike PCP-C, PCP-I were significantly more comfortable discussing phentermine (pre-PCP-I: median 7, IQR 4–8; post-PCP-I: 8, 7–9; P=0.026) and phentermine/topiramate ER (pre-PCP-I: 4, 2–6; post-PCP-I: 6, 5–8; P<0.001) after compared to before the intervention.

## **Perceived Effectiveness of Various Weight Loss Tools**

Distributions of the effectiveness ratings for specific weight loss tools are displayed in Table 4 (and Supporting Information, Figure S2). PCPs in both types of clinics gave higher effectiveness ratings to lifestyle intervention, portion control, and exercise than they did for medications in the pre-intervention survey. Unlike PCP-C, post-intervention PCP-I felt exercise was significantly less effective than they did pre-intervention (pre-PCP-I: 7, 4–8.5; post-PCP-I: 5, 3–7; p=0.035) and reported phentermine to be more effective after the intervention (pre-PCP-I: 5, 5–7; post-PCP-I: 7, 6–8; p<0.001). Interestingly, both PCP-C and PCP-I gave higher effectiveness ratings to phentermine/topiramate ER after the trial (pre-PCP-C: 4, 2–5; post-PCP-C: 6, 4–7; p=0.005; pre-PCP-I: 6, 5–8; post-PCP-I: 7, 6–8; p=0.002).

### DISCUSSION

While little is known about the content of patient-provider discussions about obesity and weight loss, prior studies have found that when weight loss counseling does occur, discussions often fail to include guideline-recommended assessments and treatment plans. (24) After our one-year, open-label pragmatic trial in urban safety net primary care clinics, we found that PCPs, regardless of working at intervention or control clinics, overvalued exercise and undervalued obesity medications compared to what the literature shows about their respective effectiveness in weight loss. However, intervention PCPs who received provider education and patient updates were significantly more comfortable discussing and gave higher effectiveness ratings to obesity medications after the trial. Our study is one of the first to assess changes in provider attitudes in response to education and real-world obesity intervention within primary care.

Providers appeared to undervalue the effectiveness of weight loss medications since phentermine and phentermine/topiramate ER were rated as the least effective weight loss tools prior to the intervention. The post-intervention survey revealed that PCPs gave weight loss medications higher effectiveness ratings after the intervention. Perhaps PCP-I perspectives changed through the four education sessions on weight loss tools and when they

saw that their patients were losing weight with one or more of the Toolbox tools. Trials have shown that medications provide equivalent or greater weight loss compared to the other lifestyle interventions included in our survey. According to a recent review of U.S. Food and Drug Administration (FDA)-approved anti-obesity medications, the average placebo-subtracted weight loss among trials was 4.8 kg with phentermine given over 12–28 months and 9.1 kg with phentermine/topiramate ER 15mg/92mg given over one year.(25)

Since many individuals treated with weight loss interventions initially lose weight and then regain it, the duration of treatment and follow up are important in assessing the relative effectiveness of different treatments. In contrast to weight loss medications, a meta-analysis showed mean weight losses of 2.6–4.4 kg with reduced-calorie diets and 7.0–7.3 kg with partial meal replacements after one year compared to various control groups whose treatments included an isoenergetic control diet, an isoenergetic traditional low-fat diet, an isoenergetic diabetic diet, the American Diabetes Association Diet, a 1,500 kcal/day control diet and a traditional lifestyle-group.(14) A 12-week Weight Watchers (WW) program produced 2.4 kg comparator-subtracted weight loss on average at one year. (26) A systematic review of 45 studies (including 39 randomized controlled trials) of WW and/or various commercial and proprietary weight loss programs found 0.1-4.9% greater weight loss among programs at one year compared to control/education and/or counseling; kg weight losses were not reported.(27) Large behavioral weight loss trials have resulted in an average of 5.5 kg weight loss after four years and only about 2 kg weight loss after ten years compared to usual care (e.g., Diabetes Prevention Program and Diabetes Prevention Program Outcomes Study).(28, 29)

Our providers serve a socioeconomically disadvantaged population; therefore, high costs of weight loss medications could explain why PCPs were less comfortable discussing these options but this should not have affected their perceptions of medication effectiveness. One can speculate as to why PCPs undervalue medications for weight loss (i.e., physician stigma against treating obesity, lack of training in obesity management and medications, fear of prescribing anti-obesity medications given historical concerns over safety, limited patient requests), but little is known about current provider views about anti-obesity medications and how these views may relate to prescribing patterns. This is a particularly relevant given the recent approval by the FDA of several new weight loss medications and the apparent low level of uptake of these medications as compared to new glucose lowering medications.(30)

In contrast to the level of efficacy PCPs attributed to the practice, physical activity without dietary restriction has been shown to have only a modest effect on body weight, typically providing a weight loss of less than 3% of initial body weight.(31) Although exercise has numerous health benefits, a review of randomized controlled trials comparing weight loss in groups assigned to physical activity alone versus groups assigned to no intervention found that only 1–3 kg of weight was lost in most studies conducted over roughly one year (range 4–16 months).(32) National physical activity guidelines do not include an evidence statement supporting the notion that physical activity results in significant weight loss, but instead stress that the "health benefits of physical activity are generally independent of body weight."(33) PCPs seem to overvalue exercise as a weight loss intervention. Our results are unlikely to be unique to our group of PCPs as a previous study found that PCPs were more

likely to counsel on physical activity than on diet or weight control.(34) Another study found that increased physical activity and dietary advice were most commonly discussed during talks about weight loss.(35) Small qualitative studies have found that clinicians often base advice on their own experiences with weight.(36) Perhaps, PCPs are more comfortable discussing exercise because they have more personal experience with this activity than dietary restriction or weight loss medications.

Consistent with previous studies, providers gave a low rating to the impact their counseling efforts have on patients' actions to lose weight.(37) It has been suggested that inadequate weight management counseling during primary care visits may in part be due to providers' perceived futility based on how they view their patients' ability to lose weight as well as environmental factors beyond their control.(38) Such a pessimistic attitude from providers may not be warranted, however, as a recent meta-analysis found that most studies have demonstrated a positive effect of provider weight loss advice on patient attempts to change behaviors related to their weight.(39) Studies have also shown that patients report that they want their physicians to address weight during visits, to give specific individualized weight loss management plans, and to provide encouragement to foster self-motivation for weight loss.(40)

This study has a number of limitations. We surveyed a small sample of PCPs from a single health care system, which primarily serves a socioeconomically disadvantaged population. Differences in baseline ratings between control and intervention clinics (e.g., the comfort and effectiveness ratings were significantly lower for phentermine/topiramate ER and the effectiveness rating was significantly higher for exercise among PCP-C compared to PCP-I, see Supporting Information, Table S1 and S2) may represent uncontrolled variation between the control and intervention clinics such as the characteristics, experiences, prior education and training of the providers at the different clinics. Interestingly, comfort and effectiveness ratings for other types of weight loss tools were not significantly different at baseline. There were also no significant differences in any of the questions related to provider views on obesity and weight loss counseling at baseline (see Supporting Information, Table S3). Altogether, the above differences likely do not change the main conclusions of this study that exercise was overvalued, medications were undervalued and intervention clinic providers' views changed more from the pre- to post-intervention survey.

The surveys were anonymous, and therefore we were unable to determine whether the same providers completed both surveys and we noted that gender was different between the pre/post groups. Given that we were unable to link pre- and post-intervention responses, we treated the two samples of providers as independent, which could underestimate the variance and thus inflate type I error. To protect anonymity, we could not match provider demographics with responses or directly assess changes to individual PCPs' perceptions and habits regarding weight loss counseling and treatment. Determining whether our PCPs' degree or provider type led to differential outcomes in comfort or effectiveness ratings would have been interesting to investigate as there was a greater proportion of non-physician PCPs in control clinics compared to intervention clinics (Table 1). A recent web-based survey to assess beliefs, practices and knowledge regarding obesity management among PCPs (family physicians and internists), OB-GYN providers and NPs revealed that rates of

pharmacotherapy prescribing were lower and aversion to bariatric surgery was significantly higher among NPs and OB-GYN providers compared to the PCP physicians.(41)

We were also unable to determine the degree of contact with the Toolbox intervention for each PCP at an intervention clinic, though, as mentioned previously, PCP-I were given four provider education sessions during the trial and received status updates when their patients enrolled in the study, which tool they chose, when they switched or added tools or achieved 5% weight loss. We only included phentermine and phentermine/topiramate ER in our survey because those were the only two medications offered in the Toolbox intervention. Although not measured, the amount of prior training PCPs had with the obesity interventions offered in the trial, particularly phentermine and phentermine/topiramate ER, likely was related to survey ratings. Providers may have had more familiarity with other FDA-approved weight loss medications such as orlistat, lorcaserin, naltrexone/bupropion or liraglutide. Phentermine and phentermine/topiramate ER, in particular, may raise unique concerns among PCPs as phentermine was part of the "fen-phen" combination taken off the market in 1997, is FDA-approved for only 3 months duration, is a stimulant and is a controlled substance. PCPs may be reluctant to prescribe topiramate because of the cognitive dysfunction that is seen at the higher doses used to treat migraines and seizures. However, as discussed before, weight loss prescription practices are very low and the most common medication prescribed was phentermine, both at DH and nationally.(42, 43) Additionally, unmeasured beliefs toward weight loss may confound the relationship between PCP comfort discussing weight loss and perceived efficacy of the weight loss tools, thus biasing our results towards or away from the null. Lastly, given that Colorado has the lowest selfreported adult obesity rates and highest physical activity rates in the country, (44, 45) our findings may not be generalizable to other settings.

In conclusion, our results suggest that providers may be spending their limited counseling time discussing exercise at the expense of discussing more effective weight loss interventions. The Toolbox trial demonstrated that providing education to clinicians and providing clinical experience with a variety of evidence-based medical weight loss tools within the primary care setting of a pragmatic clinical trial improved PCPs' comfort in discussing and their perceived effectiveness of weight loss medications.

## Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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#### STUDY IMPORTANCE QUESTIONS

## 1. What is already known about the subject?

- Weight loss counseling is inconsistent in primary care.
- More information is needed regarding primary care providers' (PCPs) views on obesity treatment options.
- PCPs tend to overvalue exercise and undervalue weight loss medications in treating obesity.

## 2. What does your study add?

- A one-year pragmatic obesity trial that provided education about and experience with weight loss treatments demonstrated a shift in PCPs' perspectives, specifically regarding exercise and weight loss medications.
- PCPs felt more comfortable discussing and gave higher effectiveness ratings to weight loss medications after this pragmatic trial, bringing their views more in line with published effectiveness data.

Table 1

Characteristics of the Surveyed Primary Care Providers

Demographic	ıphic	Pre-intervention survey $\begin{array}{c} n{=}80^* \\ n\ (\%) \end{array}$	vention survey n=80* n (%)	Post-intervention survey n=82 n (%)	ation survey 82 %)
		PCP-C <sup>†</sup> 37 (46)	$PCP-C^{\dagger}$ 37 (46) $PCP-I^{\sharp}$ 43 (54) $PCP-C$ 41 (57) $PCP-I$ 41 (43)	PCP-C 41 (57)	PCP-I 41 (43)
Gender	Gender Female	25 (68)	32 (74)	29 (71)	25 (61)
December	Physician §	24 (63)	33 (77)	24 (59)	(08) 88
angar	Non-physician#	13 (35)	10 (23)	17 (41)	8 (20)

Demographics missing for five providers on the pre-intervention survey

 $^{\uparrow}$ PCP-C = Control Clinic PCPs

\$ Included MD or DO  $^{\prime\prime}_{\rm Included}$  NP, PA-C, Pharm D, RN

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Provider Views on Obesity and Weight Loss Counseling

Table 2

p-value 0.45 0.25 0.28 0.93  $\begin{array}{c} \text{Intervention Clinic PCPs (PCP-I)} \\ n{=}89 \end{array}$ Post-intervention n=41 Median (IQR) 8 (6–8) 8 (6-2) 6 (5-7) 7 (5–8) Pre-intervention n=48 Median (IQR) 9 (8-10) 8 (6-9) 5.5 7 (6-8) p-value‡ 0.85 0.52 0.53 0.11 Control Clinic PCPs (PCP-C) n=78 Post-intervention n=41 Median (IQR) 7 (6-8) 6 (6-8) 8 (7-9) 5 (4-6) Pre-intervention n=37 Median  $(IQR)^{\dagger}$ 8 (8-10) 8 (6-2) (8-9) 6 (5-7) Do you think your advice to a patient to take action to lose weight has an impact? How comfortable are you in counseling patients on your own for weight loss? How comfortable are you overall in discussing weight with your patients? How significant a problem do you think obesity is for your patients? Question\*

Questions rated from 0 (Least significant, comfortable, or impact) to 10 (Most significant, comfortable, or impact)

 $<sup>^{7}</sup>$ IQR = Interquartile range (Q1–Q3)

 $<sup>\</sup>vec{t}$ -values calculated using Wilcoxon rank sum test

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Table 3

Perceived Comfort in Discussing Various Weight Loss Tools\*

·I)		p-value	0.56	0.29	06:0	0.026	<0.001
Intervention Clinic PCPs (PCP-I) n=89	Post-intervention n=41	Median (IQR)	(8–9) L	9 (2–5)	(6-L) 8	8 8	9 9
Interventio	Pre-intervention n=48	Median (IQR)	7 (6–8)	5 (4–7)	8 (6– <i>L</i> )	7# ( <b>4–8</b> )	4" (2–6)
		p-value‡	0.19	0.54	0.84	0.23	0.17
Control Clinic PCPs (PCP-C) n=78	Post-intervention n=41	Median (IQR)	(6 <del>-</del> 9)	6 (4–7)	8 (7–10)	6 (4–8)	4 <i>§</i> (1–5.5)
Control	Pre-intervention n=37	Median $(\mathbf{IQR})^{\dagger}$	7 (5–8)	6 (3–7)	8 8	5 (3–7)	2 (1–5)
			Lifestyle modification programs/commercial weight loss programs	Portion-controlled foods/meal replacements	Exercise	Phentermine	Phentermine/topiramate ER

Questions rated from 0 (Least comfortable) to 10 (Most comfortable)

 $<sup>\</sup>vec{T}$ IQR = Interquartile range (Q1–Q3)

<sup>†</sup> p-values calculated using Wilcoxon rank sum test

 $<sup>\</sup>stackrel{\mathcal{S}}{=} 40$  for post-intervention PCP-C and phentermine/topiramate ER

 $_{\rm II}\!=\!47$  for pre-intervention PCP-I and phentermine and phentermine/topiramate ER

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Table 4

Perceived Effectiveness of Various Weight Loss Tools\*

Intervention in Pre-intervention in District Controlled Foots/meal replacementsPre-intervention in District Controlled Foots/meal Pre-intervention in District Controlled Foots/meal replacementsPre-intervention in District Controlled Foots/meal Pre-intervention in District Controlled Foots		Control (	Control Clinic PCPs (PCP-C) n=78		Interventio	Intervention Clinic PCPs (PCP-I) $n{=}89$	[]
Median (IQR)     Median (IQR)     p-value*     Median (IQR)       8     7     0.13     7#       (6-9)     (5-8)     0.99     7       (5-7)     (4-8)     0.99     7       (6-9)     7     0.12     7       (6-9)     (5-9)     0.27     5**       (4-7)     (4-7)     (4-7)     (5-7)       4.8     6%     0.005     677       (2-5)     (4-7)     0.005     677		Pre-intervention n=37	Post-intervention n=41		Pre-intervention n=48	Post-intervention n=41	
8 7 0.13 7#   (6-9) (5-8) 0.13 7   (5-7) (4-8) 0.99 7   (6-9) (5-9) 0.12 7   (4-7) (6-9) 0.27 5**   (4-7) (4-7) 0.005 6**   (2-5) (4-7) 0.005 6**		Median (IQR) <sup>†</sup>	Median (IQR)	p-value*	Median (IQR)	Median (IQR)	p-value
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Lifestyle modification programs/commercial weight loss programs	(6 <del>-</del> 9)	7 (5–8)	0.13	7# (5–8)	7 (6–8)	0.75
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	Portion-controlled foods/meal replacements	6 (5–7)	6 (4–8)	0.99	7 (5–8)	7 (6–8)	0.21
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$	Exercise	(6 <del>-</del> 9)	7 (5–9)	0.12	7 (4–8.5)	5 (3–7)	0.035
$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Phentermine	5 (4–7)	6 <sup>  </sup> (4–7)	0.27	\$ ** (5-7)	7 (6 <del>-8</del> )	<0.001
	Phentermine/topiramate ER	4 <i>§</i> (2–5)	6¶ (4–7)	0.005	6 <sup>77</sup> (5–8)	7 ( <del>6–8</del> )	0.002

Questions rated from 0 (Least effective) to 10 (Most effective)

 $\P=38$  for post-intervention PCP-C and phentermine/topiramate ER

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 $<sup>\</sup>vec{T}$ IQR = Interquartile range (Q1-Q3)

 $<sup>\</sup>overset{\rlap{}}{f}_{\text{p-values}}$  calculated using Wilcoxon rank sum test

 $<sup>^{\$}</sup>$  n=32 for pre-intervention PCP-C and phentermine/topiramate ER

<sup>#</sup> =40 for post-intervention PCP-C and phentermine

<sup>#</sup> n=47 for pre-intervention PCP-I and lifestyle

<sup>\*\*</sup> n=46 for pre-intervention PCP-I and phentermine

 $<sup>^{\</sup>uparrow\,\uparrow}_{\rm n=45}$  for pre-intervention PCP-I and phentermine/topiramate ER