



## Computer-based personalized feedback intervention for cigarette smoking and prescription analgesic misuse among persons living with HIV (PLWH)



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### ABSTRACT

Pain, tobacco cigarette smoking, and prescription opioid misuse are all highly prevalent among persons living with HIV (PLWH). Smoking and pain medication misuse can lead to deleterious outcomes, including more severe pain and physical impairment. However, we are not aware of any interventions that have attempted to address these issues in an integrated manner. Participants ( $N = 68$ ) were recruited from an outpatient infectious disease clinic and randomized to either a computer-based personalized feedback intervention (Integrated PFI) that aimed to increase motivation, confidence, and intention to quit smoking, and decrease intentions to misuse prescription analgesic medications, or a Control PFI. Results indicated that PLWH who received the Integrated PFI (vs. Control PFI) evinced greater post-treatment knowledge of interrelations between pain and tobacco smoking. Moreover, participants who received the Integrated PFI and smoked at least 10 cigarettes per day (but not  $< 10$  CPD) reported greater confidence and readiness/intention to quit smoking. Effects of the Integrated PFI on knowledge of pain and opioid misuse, and attitudes/intentions regarding prescription pain medication misuse were not statistically-significant. Taken together, these results indicate that this novel intervention strategy may offer promise for addressing a critical public health need in a population that is generally underrepresented in clinical research.

Rates of cigarette smoking among individuals with chronic pain (~24–68%; Michna et al., 2004; Orhurhu, Pittelkow, & Hooten, 2015) and persons living with HIV (PLWH; 45–74%; Vidrine, 2009) are substantially higher than those observed in the general population (14%; Norris, Schiller, & Clarke, 2018). Pain is the second most common symptom reported among PLWH (Carr, 2004), and the point prevalence of pain among PLWH ranges from 54% to 83% using a three-month recall period (Parker, Stein, & Jelsma, 2014). Pain among PLWH is often frequent and persistent (Breitbart et al., 1996), and can be attributed to various causes, including HIV/AIDS, side effects of medication, and living conditions (Miaskowski et al., 2011). Notably, there is evidence that pain is more intense among PLWH who smoke cigarettes (Patel et al., 2006; Turner et al., 2001), possibly via smoking-related dysregulation of the endogenous opioid system (Shi, Weingarten, Mantilla, Hooten, & Warner, 2010).

Consistent with a reciprocal model of pain and tobacco smoking (Ditre, Brandon, Zale, & Meagher, 2011; Zale, Maisto, & Ditre, 2016), pain has been shown to motivate smoking (Ditre & Brandon, 2008), pain patients endorse the use of cigarettes to cope with pain (Hooten et al., 2011b; Patterson et al., 2012), and cigarette smoking has been implicated in the onset and exacerbation of painful conditions (Shiri, Karppinen, Leino-Arjas, Solovieva, & Viikari-Juntura, 2010; Sugiyama et al., 2010). Furthermore, cross-sectional evidence suggests that perceived interrelations between pain and tobacco smoking (e.g., that

smoking can help one cope with pain) are associated with more severe pain and functional impairment (i.e., interference with daily activities, work, and recreational/social activities) among individuals with chronic pain and PLWH (Ditre, Zale, Heckman, & Hendricks, 2017; Weinberger, Seng, Ditre, Willoughby, & Shuter, 2018).

Given the ubiquity of pain among PLWH, it is not surprising that they tend to be prescribed more analgesic medications than persons in the general population (Silverberg et al., 2012). There is also evidence that opioid misuse behaviors (e.g., taking more medication than prescribed, seeking multiple prescriptions) are especially prevalent among PLWH (Hansen et al., 2011; Tsao, Dobalian, & Stein, 2005), with rates of misuse as high as 62% (Robinson-Papp, Elliott, Simpson, Morgello, & Bank, 2012). Cigarette smoking has been associated with an elevated risk for engaging in aberrant drug-related behaviors (Michna et al., 2004), and chronic pain patients using opioids have reported that opioid consumption stimulates smoking (Hooten et al., 2011a). Moreover, those who endorse smoking for pain-cope have demonstrated greater reliance on analgesic medications (Jamison, Stetson, & Parris, 1991), and there is reason to suspect that complex nicotine-opioid interactions may lead to cross-tolerance (Shi et al., 2010) and diminished pain relief (Zevin & Benowitz, 1999).

Previous research has demonstrated that interventions for PLWH can enhance motivation/success in quitting smoking (e.g., Moadel et al., 2012; Vidrine, Arduino, Lazev, & Gritz, 2006; Wewers, Neidig, &

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Kihm, 2000), and improve pain outcomes (e.g., Trafton et al., 2012). However, despite the high prevalence and deleterious effects of co-occurring pain, tobacco smoking, and prescription analgesic misuse among PLWH, we are not aware of any interventions that have attempted to address these issues in an integrated manner. Integrated treatments are often preferable to traditional approaches for treating co-occurring disorders (e.g., sequential and parallel treatment), because they are more efficient, cost-effective, and do not require that one condition takes precedent over another (e.g., Mueser, 2003). One promising therapeutic format is the computer-based personalized feedback intervention (PFI; e.g., Noar, 2011; Riper et al., 2009). Drawing on motivational and social psychology perspectives (Bandura, 1994, pp. 25–59; Miller & Rollnick, 2002), PFIs rely on the presentation of discrepant information, including personalized profiles (e.g., regarding smoking rate and chronic pain status), risk factors (e.g., cigarette smoking as a risk factor for chronic pain), and normative comparisons (e.g., perceptions regarding others' misuse of prescription pain medications). Computer-based PFIs are portable, adaptable, easy to implement, and can be delivered to a large number of patients by non-specialized care providers, thereby reducing patient burden and making intervention participation more feasible (Cunningham, 2007, pp. 399–416; Hasin et al., 2013). Computer-based PFIs have been recommended for HIV care settings (Noar, 2011), and they enhance standardization of treatment in terms of format, contact time, and focus of content.

The goal of this study was to pilot test a newly developed computer-based PFI for PLWH who are prescribed analgesic medication aimed at: (1) increasing motivation, confidence, and intention to quit smoking, and (2) decreasing positive attitudes and intentions toward the misuse of analgesic medications (e.g., prescription opioids). The integrated intervention included a novel psychoeducation component that was informed by several theories of health behavior change, including the transtheoretical model (Prochaska, Redding, & Evers, 2008), the health belief model (Champion & Skinner, 2008), and the theory of planned behavior (Ajzen, 1991). In accordance with these models, it is our contention that PLWH may become more motivated to quit smoking and less motivated to misuse prescription pain medication as they develop discrepancy between the continuation of these behaviors and their desired pain outcomes. Indeed, developing discrepancy between continued substance use and desired outcomes is a core component of evidence-based motivational substance use interventions, including PFIs (e.g., Abrams & Niaura, 2003; Larimer et al., 2007). Moreover, psychoeducation regarding interrelations between pain, smoking, and analgesic misuse may increase motivation for behavior change because individuals may begin to perceive themselves as immediately susceptible to the negative pain-related effects of smoking and medication misuse and may come to understand the pain-related benefits of quitting smoking and taking pain medication as prescribed. Finally, in accordance with the theory of planned behavior, providing personalized feedback regarding interrelations between pain, smoking, and analgesic misuse may change attitudes towards these health behaviors, perceived norms regarding the consequences of these behaviors, and perceived behavioral control, which may, in turn, alter behavioral intentions.

Participants were randomized to receive either the Integrated PFI that addressed pain, smoking, and analgesic misuse, or a Control PFI, which addressed health behaviors that were not related to pain or substance use (e.g., nutrition, physical activity). We hypothesized that participants randomized to the Integrated PFI intervention would evince greater post-treatment knowledge regarding pain-smoking interrelations, as well as greater motivation, confidence, and intention to quit smoking. We also hypothesized that participants in the PFI condition would evince greater knowledge regarding pain and the misuse of analgesic medications, as well as fewer maladaptive attitudes regarding prescription opioid misuse, and reduced intention to misuse prescription pain medications in the future.

## 1. Method

### 1.1. Participants

Participants were recruited from a university hospital-based outpatient infectious disease clinic in central New York. Medical records were screened for the following inclusion criteria: (1) current cigarette smoking, and (2) current use of prescription analgesic medication. Participants aged  $\leq 30$  years and those who self-reported a current attempt to quit smoking were excluded. A total of 68 participants met these criteria and completed the single study session.

### 1.2. Procedure

Prospective participants were identified via medical chart review of patients presenting to a university hospital-based outpatient infectious disease clinic in central New York. During routine clinic visits, a designated health care provider informed patients about the opportunity to participate in the study and obtained verbal consent regarding their willingness to be introduced to a trained research assistant who worked in collaboration with clinic staff. Patients who provided permission were then introduced to the research assistant, who described the study, assessed additional inclusion/exclusion criteria, and scheduled a study appointment. Of 140 patients who completed the initial screening, 68 met all inclusion criteria. Upon arrival to the study appointment, informed consent was obtained, and participants were assured that all information collected for the study would be kept confidential and would not be shared with their health care providers. Participants were then administered a brief computer-based baseline questionnaire that would be used to collect sociodemographic data and generate personalized content for the PFI. Participants were then randomized to either PFI or Control computer-based PFI conditions. Responses to baseline questionnaires were immediately transmitted into the personalized feedback interventions, so there was no delay between completion of the baseline measures and commencement of the intervention. Primary outcome measures were administered on the same computer immediately following the intervention. All participants completed the same pre- and post-intervention questionnaires, regardless of condition assignment. Study sessions typically lasted approximately 1.5 h, and participants were compensated \$45 for the completion of the visit.

### 1.3. Intervention conditions

All participants received a brief computer-based intervention. Both interventions included visual and audio components (presented at a 5th grade reading level) and took approximately 20–30 min to complete. All feedback was presented in a non-confrontational tone, and incorporated personalized information collected during the baseline assessment.

#### 1.3.1. Integrated PFI

The Integrated PFI included content relevant to interrelations between pain, tobacco smoking, and analgesic medication use/misuse in the context of HIV and aging. Components of this intervention included personalized profiles for chronic pain (e.g., intensity, duration, goals for future functioning), smoking (e.g., cigarettes smoked per day, past quit attempts, motivation to quit), analgesic medication use/misuse (e.g., prescription compliance), and perceptions of pain-smoking interrelations. The Integrated PFI also incorporated pain-smoking psychoeducation (e.g., reciprocal relations between pain and smoking; risks of continued smoking vs. benefits of quitting in the context of chronic pain, HIV, and aging), and feedback regarding analgesic medication use/misuse (e.g., use vs. misuse; relations between pain, smoking, and the efficacy of prescription opioid medications in the context of HIV and aging). In addition, participants were given normative feedback

regarding the percentage of pain patients that increase their dose of pain medication on their own, and the percentage of pain patients that seek more than one doctor for pain medications. Participants were presented with responses regarding perceived prevalence of these behaviors, and then received feedback regarding the perceptions of others, as well as the actual percentage of pain patients who engage in these maladaptive health behaviors. Reference group data was derived from the empirical literature (e.g., McDonald & Carlson, 2013). Finally, the Integrated PFI addressed the utility of employing more adaptive (i.e., non-substance related) strategies for coping with pain (e.g., distraction, deep breathing), and prompted participants to consider alternatives to cigarette smoking and the misuse of prescription opioids.

### 1.3.2. Control PFI

The Control PFI included content relevant to the importance of nutrition, exercise, and medication adherence in the context of HIV and aging. The Control PFI content was selected to clearly distinguish it from the PFI (i.e., there was no mention of smoking or opioid misuse), while also providing benefit to PLWH. Components of this intervention included psychoeducation regarding nutrition and exercise (e.g., health consequences of poor nutrition and physical inactivity), and personalized feedback regarding perceptions of interrelations between nutrition, exercise, and HIV and aging (e.g., using nutrition as part of HIV symptom management, using exercise to reduce the negative effects of HIV and aging). Control PFI components were approximately equivalent to the PFI in terms of personalization, duration, graphic design, and focus on health-related behaviors.

## 1.4. Primary outcome measures

### 1.4.1. Knowledge of pain-smoking interrelations

The 8-item Pain and Smoking Questionnaire (PSQ) was developed for this study to assess knowledge of pain-smoking interrelations, including whether cigarette smoking can cause chronic pain, contribute to pain-related functional impairment, reduce the effectiveness of prescription analgesics, or aid in efforts to cope with pain. Items also assessed knowledge of whether pain can motivate smoking behavior and whether quitting smoking may be associated with improved pain and physical functioning. Response options included *yes*, *no*, or *not sure/don't know*. The PSQ was administered before and after intervention delivery, and total scores reflect the number of items that were answered correctly (range 0–8). This measure demonstrated good internal consistency in the current sample (Cronbach's  $\alpha = 0.82$ ).

### 1.4.2. Motivation, intention, and confidence to quit smoking

The Rulers for Smoking Cessation (RSC; Boudreaux et al., 2012) is a widely-used measure that includes three items to assess: perceived importance of quitting (“How important is stopping smoking to you?” 0 = *Not important at all*; 10 = *Most important goal of my life*); readiness/intention to quit smoking (“How ready are you to quit smoking within the next month?” 0 = *Not at all*; 10 = *100% ready*); and confidence in quitting (“How confident are you that you will quit smoking within the next month?” 0 = *Not at all*; 10 = *100% confident*).

### 1.4.3. Knowledge of pain-prescription analgesic interrelations

The 6-item Pain and Analgesic Medication Questionnaire (PAMQ) was developed for this study to assess knowledge of interrelations between pain and use/misuse of analgesic medication, including whether misusing opioids can make pain worse over time, reduce their effectiveness, or lead to tolerance. Items also assessed knowledge of what constitutes misuse (e.g., taking more than prescribed, using another person's medication), and whether taking opioids as prescribed can improve pain/functioning and reduce the risk of dependence. Response options included *yes*, *no*, or *not sure/don't know*. The PAMQ was administered before and after intervention delivery, and total scores reflect the number of items that were answered correctly (range 0–6).

This measure demonstrated acceptable internal consistency (Cronbach's  $\alpha = 0.79$ ).

### 1.4.4. Attitudes and beliefs regarding prescription analgesic use

Attitudes and beliefs regarding prescription analgesic use were assessed pre- and post-treatment using an adapted version of the Attitudes and Beliefs Questionnaire (ABQ; Passik et al., 2000). The ABQ includes items that query a range of aberrant drug-related attitudes and behaviors in the context of cancer and HIV, and we adapted this questionnaire for a more general population (i.e., so items were not specific to cancer or HIV). Participants indicated their agreement with 15 statements (e.g., “People should consider taking any prescription pain medication that relieves their pain, even if their primary doctor would not approve”), using a scale ranging from 0 (*strongly disagree*) to 4 (*strongly agree*). Two additional items assessed the percentage of patients they believe either increase their dose of pain medicine on their own or seek out more than one physician for prescription pain medications (1 = 0–25%, 2 = 26–50%, 3 = 51–75%, 4 = 76–100%). Items were summed to generate a total ABQ score, with higher scores representing a greater number of maladaptive attitudes and beliefs regarding the use/misuse of prescription pain medications. The ABQ demonstrated excellent internal consistency ( $\alpha = 0.91$ ).

### 1.4.5. Intention to misuse prescription analgesics

Intention to misuse opioids and other prescription pain medications in the next 30 days was assessed using an adapted version of the Current Opioid Misuse Measure (COMM; Butler et al., 2007), which consisted of 10 items using a 5-point Likert scale ranging from 0 (*extremely unlikely*) to 4 (*extremely likely*). Seven items from the 17-item COMM were excluded from the COMM-Intention measure because they could not be adapted for a future orientation (e.g., “In the past 30 days, how often have you had trouble with thinking clearly or had memory problems?”). The COMM-Intention items were administered before and after intervention delivery, and higher scores reflect greater intentions to misuse prescription analgesic medications in future (range: 10–50). The COMM-Intention demonstrated excellent internal consistency ( $\alpha = 0.94$ ).

## 1.5. Other measures

### 1.5.1. Smoking characteristics

Smoking characteristics (e.g., number of cigarettes smoked per day, number of years smoking) were assessed via self-report.

### 1.5.2. Pain characteristics

Pain characteristics were all assessed via self-report. First, participants were asked to indicate their primary pain location (i.e., location that hurt the most over the past three months). Response options consisted of back, head, face, neck, shoulders, arms, hands, chest, breast, stomach, abdomen, hips, legs, and feet. Second, a single item was used to assess pain duration (“How long have you been experiencing this type of pain?”; CDC, 2007). Third, pain intensity was assessed using the characteristic pain intensity subscale of the Graded Chronic Pain Scale (GCPS; Von Korff, Ormel, Keefe, & Dworkin, 1992). The GCPS provides a reliable and valid method of assessing global pain severity across a range of chronically painful conditions. Participants rated the intensity of their pain *right now*, their *worst* pain in the past 24 h, and their pain *on average* during the past 24 h using separate 0–10 numerical rating scales. Consistent with GCPS scoring instructions (Von Korff, 2011), these items were then averaged and multiplied by 10 to yield a continuous composite score (range 0–100) of characteristic pain intensity/severity.

### 1.5.3. HIV/AIDS symptom count

The AIDS Clinical Trial Group Symptom Distress Module (SDM) is a 20-item measure used to assess HIV/AIDS symptoms (Justice et al.,

2001). Participants indicated whether they have experienced any of 20 different symptoms (e.g., fatigue or loss of energy, feeling dizzy or lightheaded) over the past two weeks, and subsequently rated the degree to which they have been bothered by each symptom on a scale ranging from 1 (*doesn't bother me*) to 4 (*bothers me a lot*). The SDM has been shown to have excellent construct validity (Justice et al., 2001), and demonstrated good internal consistency in the current sample ( $\alpha = 0.89$ ).

1.6. Data analytic plan

Analyses were conducted using SPSS Version 22 (IBM Corp, 2013). First, group differences in baseline variables (e.g., demographics, smoking characteristics, pain characteristics) were examined using t-tests and chi-square analyses to verify that randomization was successful. No differences in baseline characteristics were observed between intervention conditions (all  $ps > .05$ ). Second, the distributions of all outcome variables were examined for normality, and skewness and kurtosis fell within acceptable ranges (George & Mallery, 2003). Third, we tested the effects of the intervention on each outcome measure (PSQ, RSC, PAMQ, ABQ, and COMM-Intention) using analysis of covariance (ANCOVA), with the baseline score entered as a covariate for each outcome. This approach to analyzing pretest-posttest outcomes has been recommended by researchers, and has shown to be a more powerful technique compared to repeated measures ANOVAs and gain scores (e.g., Huck & McLean, 1975). Given the observed variability in number of cigarettes smoked per day ( $SD = 10.97$ , with approximately 40% of participants smoking fewer than 10 CPD), we conducted exploratory analyses to test the effects of the interaction between treatment condition assignment and CPD (dichotomized using a median split as  $<$  or  $\geq$  10 CPD) on all smoking outcomes. Finally, the magnitude of group differences was examined using partial eta squared ( $\eta_p^2$ ), with values of 0.01, 0.09, and 0.25 characterizing effects as small, medium, or large.

2. Results

2.1. Participant characteristics

Participants included 68 PLWH (39.7% female;  $M_{age} = 51.35$  years,  $SD = 7.94$ , range: 32–70), who reported smoking an average of approximately 13 CPD ( $SD = 10.97$ ). Nearly half of the sample (47%) identified as black or African American. All participants reported an annual income below \$50,000, more than half reported earning less than \$10,000 per year, and only 3 participants completed four years of college. Participants reported experiencing an average of 12.78 HIV/AIDS symptoms over the past two weeks ( $SD = 6.63$ ), with “fatigue or loss of energy”, “pain, numbness, or tingling in the hands or feet”, “difficulty falling or staying asleep”, and “muscle aches or joint pain” representing the most frequently endorsed symptoms. Over 95% of the sample endorsed pain lasting longer than 3 months, which is a commonly used cutoff for indexing chronic pain (e.g., Merskey, 1986). Half of all participants reported neck/back as primary source of pain, and over one-third reported lower extremity pain. The mean characteristic pain intensity score was high ( $M = 54.41$ ,  $SD = 31.61$ ; Von Korff, Dworkin, & Le Resche, 1990). Finally, participants reported using a variety of prescription analgesics, including both opioid (e.g., Oxycodone, Hydrocodone, Percocet) and non-opioid (e.g., Gabapentin, Lyrica, Naproxen) pain medications. Complete sociodemographic and clinical data are presented in Table 1.

2.2. Smoking-relevant outcomes

2.2.1. Pain-smoking knowledge

Participants randomized to the Integrated PFI demonstrated greater knowledge regarding pain-smoking interrelations at post-treatment,

**Table 1**  
Sociodemographic, smoking, and pain characteristics.

	PFI n = 32	Control n = 36	Total N = 68
	n (%)	n (%)	n (%)
Gender			
Female	14 (43.8%)	13 (36.1%)	27 (39.7%)
Race			
White	12 (37.5%)	16 (44.4%)	28 (41.2%)
Black or African American	16 (50.0%)	16 (44.4%)	32 (47.1%)
American Indian/Alaska Native	4 (12.5%)	4 (11.1%)	8 (11.8%)
Ethnicity			
Hispanic	2 (6.3%)	6 (16.7%)	8 (11.8%)
Income			
< \$10,000	16 (50.0%)	19 (52.8%)	35 (51.5%)
\$10,000 - \$19,999	11 (34.4%)	8 (22.2%)	19 (27.9%)
\$20,000 - \$29,999	3 (9.4%)	3 (8.3%)	6 (8.8%)
\$30,000 - \$39,999	1 (3.1%)	5 (13.9%)	6 (8.8%)
\$40,000 - \$49,999	1 (3.1%)	1 (2.8%)	2 (2.9%)
Education			
Did not graduate high school	15 (46.9%)	10 (27.8%)	25 (36.8%)
High school graduate	7 (21.9%)	9 (25.0%)	16 (23.5%)
Some college	4 (12.5%)	9 (25.0%)	13 (19.1%)
Technical school/Associate's degree	6 (18.8%)	5 (13.9%)	11 (16.2%)
4-year college degree	0 (0.0%)	2 (5.6%)	2 (2.9%)
School beyond 4-year college degree	0 (0.0%)	1 (2.8%)	1 (1.5%)
Primary Pain Location			
Neck/Back	14 (43.8%)	19 (52.8%)	33 (48.6%)
Lower Extremity	14 (43.8%)	11 (30.6%)	25 (36.8%)
Upper Extremity	2 (6.3%)	3 (8.3%)	5 (7.4%)
Chest	1 (3.1%)	0 (0.0%)	1 (1.5%)
Stomach	0 (0.0%)	1 (2.8%)	1 (1.5%)
Head	1 (3.1%)	2 (5.6%)	3 (4.4%)
Pain Duration			
< 3 months	3 (9.4%)	1 (2.8%)	3 (5.9%)
3–12 months	7 (21.9%)	5 (13.9%)	12 (17.6%)
> 12 months	22 (68.8%)	30 (83.3%)	52 (76.5%)
Pain Medication			
Opioid	15 (46.9%)	23 (63.9%)	38 (55.9%)
Non-opioid	17 (53.1%)	13 (36.1%)	30 (44.1%)
	M (SD)	M (SD)	M (SD)
Age	51.91 (8.23)	50.86 (7.74)	51.35 (7.94)
Characteristic pain intensity <sup>a</sup>	57.60 (32.41)	51.57 (31.07)	54.41 (31.61)
Cigarettes per day	10.75 (7.55)	15.06 (13.07)	13.03 (10.97)
Years smoking	28.13 (10.69)	29.00 (10.17)	28.59 (10.35)
ACTG Symptom Count <sup>b</sup>	13.91 (6.82)	12.00 (6.48)	12.78 (6.63)

Note.

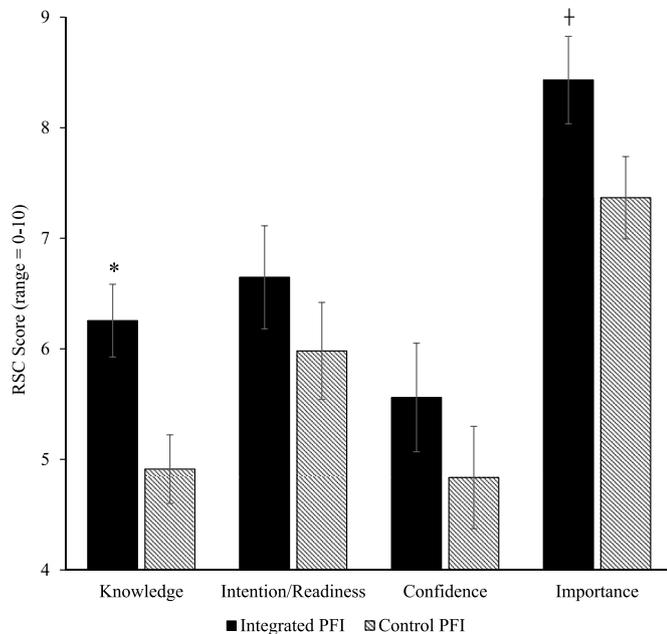
<sup>a</sup> Graded Chronic Pain Scale - Characteristic Pain Intensity Subscale.

<sup>b</sup> AIDS Clinical Trial Group Symptom Distress Module – Symptom Count.

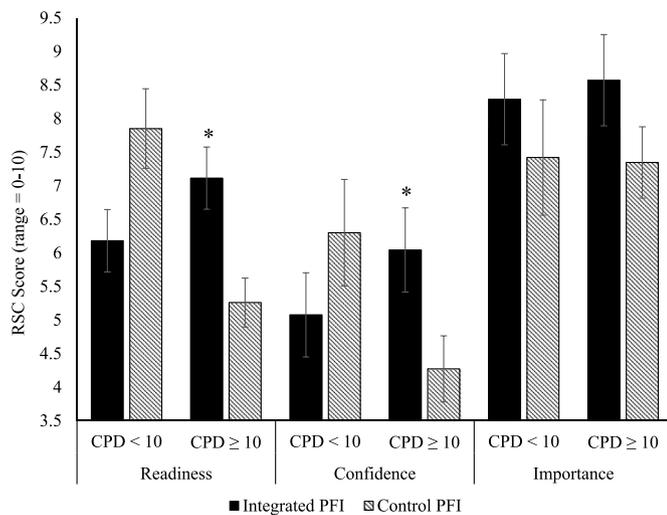
when compared to those randomized to the Control PFI ( $M = 6.26$ ,  $SE = 0.33$  vs.  $M = 4.91$ ,  $SE = 0.31$ ;  $F[1, 65] = 8.84$ ,  $p < .01$ ,  $\eta_p^2 = 0.12$ ; see Fig. 1), with an observed effect that may be characterized as medium-to-large in magnitude.

2.2.2. Readiness/intention to quit smoking

Although we did not observe a main effect for post-treatment readiness/intention to quit smoking ( $F[1, 65] = 1.08$ ,  $p = .30$ ,  $\eta_p^2 = 0.02$ ), adjusted means were in the expected direction (PFI  $M = 6.65$ ,  $SE = 0.47$  vs. Control PFI  $M = 5.98$ ,  $SE = 0.44$ ; see Fig. 1). The intervention condition  $\times$  CPD interaction term, however, was statistically-significant ( $F[1, 63] = 7.46$ ,  $p < .01$ ,  $\eta_p^2 = 0.11$ ; see Fig. 2), indicating that the Integrated PFI resulted in greater readiness/intention to engage a quit attempt among participants who smoked at least



**Fig. 1.** Smoking-Relevant Outcomes as a Function of Intervention Condition Assignment. *Note.* \* $p < .05$ , † $p < .10$ . RSC = Rulers for Smoking Cessation. Knowledge = knowledge regarding pain-smoking interrelations; Importance = importance of stopping smoking; Readiness = Intention/readiness to quit smoking within the next month; Confidence = Confidence in quitting smoking within the next month.



**Fig. 2.** Readiness, Confidence, and Importance of Quitting Smoking as a Function of Interaction between Intervention Condition and Cigarettes Smoked Per Day. *Note.* \* $p < .05$ . RSC = Rulers for Smoking Cessation. CPD = Cigarettes smoked per day; Importance = importance of stopping smoking; Readiness = Intention/readiness to quit smoking within the next month; Confidence = Confidence in quitting smoking within the next month.

10 CPD (but not among those who smoked less than 10 CPD), with partial eta squared indicating a medium-to-large effect size.

**2.2.3. Confidence in quitting smoking**

Although we did not observe a main effect for post-treatment confidence in quitting over the next month ( $F[1, 65] = 1.15, p = .29, \eta_p^2 = 0.02$ ), adjusted means were in the expected direction (PFI  $M = 5.56, SE = 0.49$  vs. Control PFI  $M = 4.84, SE = 0.46$ ; see Fig. 1). Again, however, the condition  $\times$  CPD interaction term was statistically-significant ( $F[1, 63] = 4.62, p = .04, \eta_p^2 = 0.07$ ), indicating that the

Integrated PFI resulted in greater confidence to engage a quit attempt only among participants who smoked at least 10 CPD, with an effect that may be characterized as medium in magnitude.

**2.2.4. Perceived importance of quitting**

A trend-level main-effect was observed for perceived importance of quitting smoking ( $F[1, 65] = 3.73, p = .06, \eta_p^2 = 0.05$ ), such that post-treatment ratings of quitting importance were higher among participants randomized to the ( $M = 8.43, SE = 0.40$ ) vs. Control ( $M = 7.37, SE = 0.37$ ) PFI. Partial eta squared indicates a medium sized main effect, and the intervention condition  $\times$  CPD interaction did not reach statistical significance ( $F[1, 63] = 0.10, p = .76, \eta_p^2 = 0.00$ ).

**2.3. Prescription opioid/pain medication-relevant outcomes**

**2.3.1. Pain-prescription analgesic knowledge**

Although adjusted means were in the expected direction (Integrated PFI  $M = 5.00, SE = 0.29$  vs. Control PFI  $M = 4.42, SE = 0.28$ ), group differences in post-treatment knowledge regarding interrelations between pain and the use/misuse of analgesic medication did not reach statistical significance ( $F[1, 65] = 2.05, p = .16, \eta_p^2 = 0.03$ ).

**2.3.2. Attitudes and beliefs regarding prescription opioid/analgesic misuse**

Although participants randomized to the Integrated PFI (vs. Control PFI) endorsed fewer post-treatment maladaptive attitudes and beliefs regarding the misuse of prescription pain medications ( $M = 15.97, SE = 1.90$  vs.  $M = 18.41, SE = 1.79$ ), this difference was not statistically-significant ( $F[1, 65] = 0.87, p = .35, \eta_p^2 = 0.01$ ).

**2.3.3. Intention to misuse prescription pain medications in the future**

No main effect was observed regarding intention to misuse opioids and other prescription pain medications in the next 30 days as a function of condition assignment (Integrated PFI  $M = 16.57, SE = 1.27$  vs. Control PFI  $M = 16.42, SE = 1.20$ ;  $F[1, 65] = 0.01, p = .93, \eta_p^2 < 0.001$ ).

**3. Discussion**

This is the first study to examine the effects of a brief integrated computer-based PFI that was developed to increase motivation to quit smoking and decrease intention to misuse prescription opioid medications among people living with HIV. In terms of smoking-related variables, results indicated that participants randomized to the Integrated PFI intervention (vs. Control PFI) evinced greater post-treatment knowledge of interrelations between pain and tobacco smoking (e.g., that smoking is a unique risk factor in the onset/progression of chronic pain, and that quitting smoking may improve pain and physical impairment). Two significant interactions further indicated that the Integrated PFI resulted in greater confidence in quitting and readiness/intention to quit only among heavier smoking PLWH (i.e., those who reported smoking at least 10 cigarettes per day). Finally, we observed a trend-level main effect, such that Integrated PFI participants reported greater perceived importance of quitting smoking. Effect sizes ranged from medium to medium-large in magnitude, and these findings are generally consistent with evidence that providing individuals with explicit links between their smoking and health may increase motivation to quit smoking (McCaul et al., 2006).

In terms of pain medication misuse outcomes, results were less robust. Specifically, although adjusted means were generally consistent with hypotheses, we observed no statistically-significant differences as a function of treatment condition assignment regarding pain-opioid misuse knowledge, prescription medication misuse beliefs/attitudes, or intention to misuse opioids in the next 30 days. Collectively, effects ranged from small to less than small in magnitude. One possible explanation for these null findings is that participants were not selected based on current misuse or risk for future misuse of prescription pain

medications. Nearly 1/3 of participants in this study endorsed zero intention for misuse at baseline, which may have limited our ability to detect change in related outcomes. Future research may benefit from recruiting smokers with PLWH who are determined to be at risk for aberrant use of prescription opioids.

Tobacco smoking, chronic pain, and prescription analgesic misuse are all highly prevalent among PLWH, and unique interrelations between pain, smoking, and the misuse of prescription analgesia represent a prime target for novel intervention in this population. Consistent with this perspective, smokers who are not yet ready to quit should receive brief interventions to increase motivation to quit and promote future quit attempts (Fiore et al., 2008). In addition, the authors of a seminal review of smoking cessation among PLWH observed that approximately 80% of HIV + smokers had not considered quitting and suggested that new interventions for this population should be more accessible and specifically target motivation to quit (Niaura et al., 2000). Single-session PFIs (without therapeutic guidance) have been shown to be both efficacious and cost-effective in reducing problematic substance use (Riper et al., 2009), and these data suggest that a brief computer-based PFI may function as a “first line of defense” in addressing confidence and readiness to quit smoking in HIV care settings.

Despite novel and encouraging aspects of this approach, several important limitations should be considered. First, the current sample was relatively small, and although these data provide preliminary support for a larger efficacy study (perhaps focusing on smoking-related outcomes), indices of statistical significance and effect size should be interpreted with caution. Future work should recruit a larger sample of PLWH, which would result in greater power to detect statistically significant effects and would allow for the inclusion of additional theoretically relevant covariates (e.g., pain intensity). Second, because primary outcomes were assessed immediately following treatment delivery, there is some potential for demand, and the extent to which these effects are maintained beyond the initial assessment period remains unclear. Similarly, although these data provide support for intervention-related change in motivation to quit smoking and intention to misuse prescription pain medication, the effects of the intervention on behavior are unknown. Future work would benefit from employing longer-follow up periods, with an emphasis on rates of engagement and subsequent outcomes of post-treatment smoking cessation attempts. Third, given that the Integrated PFI was compared to a control intervention that did not address pain-smoking relations or motivation to quit smoking, additional research is needed to determine the incremental utility of incorporating tailored/integrated pain-smoking content into brief motivational smoking treatments. Finally, participants were not selected based on current misuse or risk for future misuse of prescription pain medications at baseline, and future work should recruit a sample of patients who engage in aberrant medication-related behaviors.

In summary, results of this pilot study indicate that a brief (20–30 min) computer-based personalized feedback intervention may result in greater knowledge regarding pain-smoking interrelations and enhance confidence/intention to quit smoking, particularly among heavier cigarette smokers living with HIV and chronic pain. There is an emerging global emphasis on the adaptation of behavioral interventions for PLWH that can be easily disseminated (Collins et al., 2011), and by developing a portable/integrated smoking intervention for PLWH with chronic pain, our novel intervention strategy addresses a critical public health need in a population that is generally underrepresented in clinical research.

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## Conflicts of interest

We have no conflicts of interest to declare.

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