Managing Acute Insomnia in Prison: Evaluation of a ‘one-shot’ Cognitive Behavioural Therapy for Insomnia (CBT-I) intervention

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ABSTRACT

Insomnia is a serious condition that affects over 60% of the prison population and has been associated with aggression, anger, impulsivity, suicidality, and increased prison health care use. Nonpharmacological interventions for prison inmates are scarce despite the high prevalence and significant consequences of insomnia among those incarcerated. The aim of the present study was to examine the preliminary efficacy and effectiveness of a one-shot of Cognitive Behavioural Therapy for Insomnia (CBT-I) for prison inmates with acute insomnia in an open trial. The intervention consisted of one 60-70 minute session of CBT-I and a self-management pamphlet. A consecutive series of 30 adult male offenders with acute insomnia from a UK prison completed measures of prospective sleep (daily sleep diary), insomnia symptoms severity (Insomnia Severity Index), and mood symptoms (Patient Health Questionnaire and General Anxiety Disorder) one week before and four weeks after receiving the intervention. Pairwise t-tests revealed that a single-shot of CBT-I was effective in reducing the severity of insomnia in adult male offenders (t=(29), 12.65,p<0.001). Further, the results demonstrated high effect sizes for reductions in depressive (dz=1.07) and anxious (dz=1.06) symptoms, as well as insomnia severity (dz=2.25). A single shot session of CBT-I is effective in managing acute insomnia and mood (depression, anxiety) symptoms in adult male prison inmates. Future research should focus on testing if the single shot CBT-I intervention can be implemented and disseminated in other settings and populations (e.g., female and juvenile/youth offenders).

Key Words: Acute Insomnia, Prison, Depression, Anxiety, Cognitive Behavioural Therapy for Insomnia

INTRODUCTION

An estimated 11 to 81% of prison inmates report insomnia symptoms (Dewa, Kyle, Hassan, Shaw & Senior, 2015). The wide variation among studies may be due to different criteria used to define insomnia as well as different measures used to assess insomnia symptoms. For example, requests for hypnotics were used as an indicator of insomnia in one study (10.9%- Kjelsberg & Hartvig, 2005) whereas other studies, with much higher prevalence rates, used clinical screening instruments or general health questionnaires with one or two sleep-related items embedded within (e.g. 74%- Singleton, Meltzer & Gatward, 1998; 71.5%- O’Brien, 2001). The only study to date which has used an insomnia-specific measure, the Sleep Condition Indicator (need reference), suggests an overall prevalence rate for Insomnia Disorder of 61.6%, with women significantly more likely to report insomnia, in prison, compared to men(Dewa, Hassan, Shaw & Senior, 2017).

The consequences of insomnia in prison inmates include aggression, anger, impulsivity and increased prison health care use (Barker, Ireland, Cu & Ireland, 2016; Ireland & Culpin, 2006; Feron, Paulus, Tonglet, Lorant & Pestiaux, 2005). Importantly, as insomnia has been associated with suicide ideation, suicide attempts and completed suicides both in the general population and in prison populations, the additive impact of insomnia and disrupted mood is likely to increase the vulnerability to suicide in this population(Carli, Roy, Bevilacqua, Maggi, Cesaro, Sarchiapone, 2011). As such there is a critical need to address the development and treatment of insomnia in prison inmates.

Despite a significant amount of research focused on the development and treatment of other mental and physical conditions in the prison population, there is a dearth of literature examining the development and treatment of insomnia within the prison environment(Dewa et al, 2015). Spielman’s three-factor model of insomnia may be beneficial in attempting to understand the development, and maintenance of insomnia in this population. The model posits that insomnia usually begins with the combination of a predisposition towards insomnia (e.g., high emotional reactivity) paired with a precipitating event (e.g., a stressful event). The transition to chronic insomnia is usually perpetuated by several factors, which may include an increased effort to induce sleep in response to distress about poor sleep and conditioned arousal (whereby the bed becomes a cue for arousal rather than sleep), which serve to maintain the sleep problem. Within this context, imprisonment may act as the initial precipitating factor contributing to the onset of insomnia (Elger & Sekera, 2009). In fact, in one study, 60% of non-substance misusing prisoners reported symptoms of insomnia starting within their first few weeks of imprisonment(Elger, 2004). The high prevalence rate of mental health disorders and substance abuse in prison inmates in conjunction with the stress of sentencing and incarceration may also serve to precipitate insomnia. One study demonstrated that 83% of prison inmates with insomnia reported anxiety and/or depression as the initial cause(Elger, 2004).

In addition, the constraints of the prison environment may also serve as a perpetuating factor that continues to exacerbate insomnia, including the separation from family/loved ones, strictly enforced routines and sleep/wake schedules, limited exercise/physical activity, shared living/sleeping space, lack of privacy, safety concerns, limited access to sunlight, and environmental factors (e.g., noise, light, temperature, mattress/bedding; Singleton, Meltzer & Gatward, 1998; Elger, 2009; Eytan, Haller, Wolff, Cerutti, Sebo &Bertrand, 2011; Toller, 1978; Hassan, Edge, Senior & Shaw, 2013). Further, spending a significant amount of time in a prison cell on a cot, that is not only used for sleep but also activities of daily living (e.g., sitting, socializing, watching television, reading, writing, eating, napping) may contribute to conditioned arousal serving to perpetuate insomnia (Ireland & Culpin, 2006). Therefore, it is not surprising that hypnotic and anxiolytic use is ten times higher for prison inmates (even when limiting to patients with no known history of substance misuse) compared to community dwelling outpatients. (Elger, Goehring, Reyaz & Morabia, 2002).

Cognitive behavioural therapy (CBT), a psychotherapeutic approach that targets cognitions and behaviours that cause and maintain a problem, has been adapted to many psychiatric disorders including insomnia. Cognitive Behavioral Therapy for insomnia (CBT-I) is an effective non-pharmacological treatment for insomnia***.*** CBT-I is a structured, short-term, skill-focused psychotherapy aimed at changing maladaptive cognitions (i.e. thoughts and beliefs) and behaviours contributing to insomnia. The weight of evidence supporting CBTI, summarized in several meta-analyses, led to its recognition as a first-line treatment for insomnia by the British Association of Psychopharmacology, European Sleep Research Society and American College of Physicians (Wilson, Nutt, Alford, Argyropoulos, Baldwin, Bateson, ... & Gringras, P, 2010; Riemann, Baglioni, Bassetti, Bjorvatn, Dolenc Groselj, Ellis…& Hertenstein, 2017; Qaseem, Kansagara, Forciea, Cooke & Denberg, 2016). Improvements following CBTI are equivalent to those achieved during acute treatment with hypnotic medications and its effects are more durable after treatment discontinuation (Edinger, Wohlgemuth, Radtke, Marsh & Quillian, 2001; Morin, Colecchi, Stone, Sood & Brink, 1999). Further, studies have shown that the treatment of insomnia using CBT-I is associated with reductions in depressive and anxious symptomology in addition to suicide ideation(Taylor & Pruiksma, 2014; Manber, Bernert, Suh, Nowakowski, Siebern & Ong, 2014; Trockel, Karlin, Taylor, Brown & Manber, 2015).

Although it would appear that a full CBT-I delivery model would be the treatment of choice in the prison population, there may be a more timely and cost effective method for managing insomnia in this context. Considering the increased vulnerability to develop insomnia within the prison environment, it may be prudent to address insomnia during its acute phase, as it is likely to be easier to treat and have better prognostic outcomes as opposed to when it is in its chronic phase (Ellis, Gehrman, Espie, Riemann & Perlis, 2012). To that end, Ellis and colleagues developed a ‘one-shot’ CBT-I intervention specifically designed to circumvent the transition from acute to chronic insomnia(Ellis, Cushing & Germain, 2015). The one-shot involves a self-help pamphlet and a single 60-70 minute therapy session. In their first study, using a community sample from the general population, Ellis and colleagues showed an initial remission rate of 60% in those who received the intervention compared to 15% in the Wait List Control group one-month post therapy. Furthermore, at three months post therapy 73% of those in the treatment group had remitted. In a subsequent study it was demonstrated that in addition to being equally effective delivered in groups as it was in an individual face-to-face format; there were significant reductions in depression and anxiety symptomology one-month post treatment(Boullin, Ellwood & Ellis, 2016). As such it appears that the brief ‘one-shot’ treatment may be a beneficial intervention and adaptable to the prison setting.

Therefore, the aim of the present study was to examine the preliminary efficacy and effectiveness of a one-shot CBT-I intervention for acute insomnia. It was hypothesised that the intervention would be an effective treatment, as evidenced by significant reductions in insomnia-related symptoms and improvements in sleep in adult male prisoners post treatment. A secondary hypothesis was that the one-shot CBT-I intervention would also be associated with significant reductions in depressive and anxious symptomology.

METHOD

Design

An open trial of a one shot CBT-I intervention in a prison setting to inmates with acute insomnia was tested. It was decided that there would be no control group as the intervention aimed to help individuals who suffer from acute insomnia, which has a short duration (2 weeks-3 months– Ellis et al, 2012). As such, withholding treatment from a control group could exacerbate their symptoms and lead to them developing chronic insomnia for which the specific intervention under examination has yet to be tested and alternative treatments within this particular prison are limited. Ethical approval was obtained from the Ethics Committee at Northumbria University, the National Health Service Health Research Authority and the North East Forensic Psychology Services Ethics Committee.

Participants

A consecutive series of thirty Category C male prisoners (a closed prison for individuals who cannot be trusted in open conditions, however the likelihood of them escaping is low) aged 21-60 were recruited over a period of four months. All participants (n = 30) identified themselves as white British. Participants were selected through self-referral or staff referral to a ‘Mental Health in Reach Team’ in a prison in the North East of the United Kingdom. To be included in the study participants had to have a principle complaint of acute insomnia. Acute insomnia was defined based on meeting all criteria for DSM-5 Insomnia Disorder but with a duration of between two weeks and three months (see procedure). Participants were excluded if, following clinical interview, it was determined that they did not have insomnia or it was already chronic (defined as greater than three months). Moreover, if participants presented with any co-morbidity, physical or psychological, or substance abuse they were excluded from taking part. In these cases participants were referred to specialist medical/psychological services for treatment as usual.

Materials

*Insomnia Severity Index.*

The Insomnia Severity Index (ISI) is a 7-item questionnaire used to assess the nature, severity and impact of insomnia(Morin, 1993). A 5-point Likert scale (0=not at all – 4=very severe) is used to rate each item. The total questionnaire yields a score between 0-28. Reliability and validity of the ISI is well documented(Bastien, Vallieres & Morin, 2001), showing a high level of internal consistency (Cronbach’s Alpha = 0.74). The ISI has also been shown to be a clinically useful outcome tool in insomnia treatment research (Morin, Belleville, Belanger & Ivers, 2011).

*Patient Health Questionnaire.*

The Patient Health Questionnaire (PHQ) is a 9-item questionnaire used to screen, diagnose and monitor the severity of depression(Kroenke, Spitzer & Williams, 2001). It uses a 4-point Likert scale (0=not at all – 3=everyday), yielding a score between 0-27. The PHQ has high levels of internal consistency, with a Cronbach’s Alpha of 0.89(Kroenke, Spitzer & Williams, 2001) identifying it as a valid and reliable instrument. For the present study, Question 3 (‘Trouble Falling asleep or staying asleep or sleeping too much) was omitted from all analyses (as such the range for the PHQ was between 0-24).

*Generalised Anxiety Disorder.*

The Generalised Anxiety Disorder (GAD) is a 7-item questionnaire used to measure anxiety symptoms(Spitzer, Kroenke, Williams & Lowe, 2006). The GAD uses a 4-point Likert scale (0=not at all – 3=everyday) totalling a score between 0-21. It shows high levels of internal consistency (Cronbach Alpha = 0.91) suggesting it is a valid and reliable psychometric instrumentfor assessing the symptoms of anxiety, and has been used with a wide variety of populations (Lowe, Decker, Muller, Brahler, Schellberg, Herzog & Herzberg, 2008).

*Daily Sleep Diary.*

The Consensus Sleep Diary(Carney, Buysse, Ancoli-Israel, Edinger, Krystal, Lichstein & Morin, 2012) was used throughout the present study. The sleep diary asks the participant to report: i) what time the participant went to bed, ii) what time they intended to sleep, iii) how long they were awake during the night, iv) what time they woke for the final time, v) what time they got out of bed and vi) how much sleep the participant felt they got that night. Participants were instructed to estimate these times, not relying on a clock, and complete the diary every morning within a 20-40 minute window of waking. From this data the following variables were derived by averaging the previous week’s data: Time in Bed (TIB), Sleep Latency (SL), Wake After Sleep Onset (WASO), Total Sleep Time (TST) and Sleep Efficiency (SE). SE was calculated using the following formula (TST/TIB x 100) to derive a percentage.

*‘One-Shot’ CBT-I Intervention.*

Full details of the one-shot intervention have been described previously(see Ellis, Cushing & Germain, 2015 for more details). The single CBT-I session remained the same as it had in the two previous studies (Ellis, Cushing & Germain, 2015; Boullin, Ellwood & Ellis, 2016)except two diagrams were used to facilitate discussion on the body’s natural sleep cycle and how predisposing, precipitating and perpetuating factors can affect sleep: Borberly’s two-process model of sleep(Borbely, 1982) and Spielman’s 3P model(Spielman, Saskin & Thorpy, 1987) respectively. As with the previous studies, Sleep Restriction, including the rules for weekly titration, was the main focus of the single session and the previous weeks sleep diaries were used to set the initial prescribed sleep schedule (i.e., time to bed and time out of bed). The initial prescription was based upon the average total sleep time from the previous week becoming the time in bed for the following week. Finally, participants were told to continue titration at weekly intervals until they had reached a stable SE between 85-90% and were satisfied with their sleep. The self-help pamphlet outlined the principles of Stimulus Control, Cognitive Control, and the use of Imagery Distraction Techniques (a copy of the pamphlet is available from the corresponding author). There were two modifications made to the pamphlet in order to accommodate the prison environment – (1) where previously Stimulus Control instructions suggested the bedroom should only be used for sleep and sex, sex was omitted from these instructions and (2) within the Stimulus Control instructions, participants were not instructed to leave the bedroom but rather identify a ‘non-sleep’ space in their cell and go there if they were unable to sleep. The single session remained the same as it had in the two previous studies (Ellis, Cushing & Germain, 2015; Boullin, Ellwood & Ellis, 2016)except two diagrams were used to facilitate discussion on the body’s natural sleep cycle and how predisposing, precipitating and perpetuating factors can affect sleep: Borberly’s two-process model of sleep(Borbely, 1982) and Spielman’s 3P model(Spielman, Saskin & Thorpy, 1987) respectively. As with the previous studies, Sleep Restriction, including the rules for weekly titration, was the main focus of the single session and the previous weeks sleep diaries were used to set the initial prescribed sleep schedule (i.e., time to bed and time out of bed). The initial prescription was based upon the average total sleep time from the previous week becoming the time in bed for the following week. Finally, participants were told to continue titration at weekly intervals until they had reached a stable SE between 85-90% and were satisfied with their sleep. The same therapist (CR) delivered the intervention in an individual face-to-face setting. One other modification to the study protocol from that of the previous studies was that participants were seen at weekly intervals over the following four weeks (i.e. until study completion), if they requested, to check their sleep diary calculations and titration schedules were correct. No additional guidance was given at these sessions. Of the 30 prisoners who took part, 29 (96.67%) attended an additional support session.

Procedure

Participants that had self-referred or been referred to the Mental Health in Reach Team on the basis of a principle complaint of acute insomnia were specifically referred to the first author for assessment and treatment (CR). CR is a Senior Assistant Psychologist with eight years of experience working with a closed forensic setting and was trained and supervised by the corresponding author with respect to the ‘one-shot’ CBT-I (JGE – A qualified Somnologist with 8 years’ experience delivering CBT-I). As is standard in this environment, full clinical interviews are undertaken on prisoners when they either self-refer or a referred to the Mental Health in Reach Team for any issue. For the present study, however, an additional sleep disorder clinical interview was undertaken by the clinician (CR) to determine whether participants met criteria for acute insomnia and whether they had an occult sleep disorder that may have been masking as acute insomnia (e.g. Obstructive Sleep Apnoea, Periodic Limb Movement Disorder, a Circadian Rhythm Disorder). If the participant met study criteria they were informed about the nature, duration and level of support available for the study and asked whether they would like to take part. If participants agreed, informed consent was gained and the participant enrolled. At this meeting the participant was given the sleep diary, instructed on its use and completed the PHQ, GAD and ISI for baseline assessment. Appointments were also made for the following week for the intervention session and for the final follow-up assessment (four weeks following the intervention). At the follow-up appointment sleep diaries were collected and participants completed the same measures from baseline. Finally, participants were debriefed and, if requested, were referred on to another service.

Prison Routine/Environment

Prisoners would be locked in their cells between 19:00 and 19:30 in the evening and remain there until between 07:00 and 07:30 in the morning. On waking, and if employed, prisoners would leave their wing at 08:00, return to their cells for lunch between 11.30 and 13.30, and then return to work from 14:00 until 16:00. The evening meal, in their cell, would be at approximately 17:00 and then there would be an opportunity to leave the cell for recreation for approximately 1-2 hours prior to being locked in for the night. During recreation periods all prisoners had access to the gym and an outside courtyard. All cells contained a single bed, television, small window, curtains and bathroom. Although prisoners had control over what time they turned out the lights at night they had no control over ambient temperature or noise levels.

Analytic Strategy

Paired t-tests were used to determine significant differences between pre and post scores on the ISI, GAD and PHQ. Additionally, paired t-tests were undertaken on sleep diary derived measures of SL, WASO, TST and SE. Within group Cohen’s dz’s were then calculated to examine the effect size of changes on the scales (ISI, GAD-7 and PHQ) and sleep variables (SL, WASO, TST and SE). Finally, remission status was defined as a reduction >7 points on the ISI between baseline and follow up(Morin, Belleville, Belanger & Ivers, 2011). Missing data of less than 5% on measures was treated by mean substitution and over 5% by casewise deletion.

RESULTS

During the study duration 30 referrals (both self and by others) were made. No participants were excluded based on inclusion/exclusion criteria and no participant refused to enrol and complete the intervention and assessments. Data was collected from 30 males – mean age 33.13 + 8.85 years. Descriptive data on the sample can be seen in Table 1.

INSERT TABLE 1 HERE

Paired t-tests showed that participants experienced a significant reduction in insomnia-related symptoms (ISI) one month post intervention (t(29)=12.65, p<0.001). Significant reductions were also observed for both anxious (t(29)= 5.03, p<0.001) and depressive symptomology (t(29)= 4.88, p<0.001). In terms of changes in sleep continuity, there were significant increases in TST (t(29)= 4.93, p<0.001) and SE (t(29)= 5.64, p<0.001) at follow up. Moreover, there were significant reductions in SL (t(29)= 5.92, p<0.001) and WASO (t(29)= 5.57, p<0.001). The effect sizes, calculated using between group mean change scores (pre-treatment to follow up) on each variable, were dz=2.25 (ISI), 1.07 (PHQ), 1.06 (GAD), 0.86 (TST), 1.13 (SL), 0.90 (WASO) and 1.04 (SE).

Remission rates based upon pre-post changes in ISI score were 73.33% and compliance (as defined by the number of nights participants stayed within their prescribed time to bed and/or prescribed time out of bed in the first week post therapy within a margin of 15 minutes) was 90%.

DISCUSSION

The aim of the present study was to determine the preliminary effectiveness and efficacy of a ‘one-shot’ CBT-I intervention in male prisoners with acute insomnia. A secondary aim was to determine whether the treatment also reduced symptoms of depression and anxiety. With respect to the primary aim, the intervention was found to be efficacious, with 73% of prisoners remitting at the one-month follow-up. Significant reductions in insomnia symptoms, and improvements in subjective sleep from the sleep diary, were also observed. Furthermore, the effect sizes on these variables were also strong, ranging from moderate to large. Whilst these findings are in line with the previous literature on the impact of a single shot CBT-I intervention for acute insomnia (Ellis, Cushing & Germain, 2015; Boullin, Ellwood & Ellis, 2016),they also suggest that the intervention can be successfully delivered in a prison environment with minor modifications. Considering the relatively high prevalence of chronic insomnia in prison inmates (+60%) and the existing literature suggesting the process of incarceration may act as a precipitant to the development of insomnia itself, the results should be viewed as an important first-step in managing insomnia in the prison inmates with the potential to reduce or prevent violence, suicide attempts, and health care utilization of prison inmates. Interestingly all those who took part in the study were asked, on completion, what they felt were the most beneficial aspects. Anecdotally, most of the comments revolved around the pamphlet and specifically the stimulus control instructions. It is, however, speculative to suggest that stimulus control was the most powerful component of the intervention in this context, rather that this is what the population felt was the most helpful.

The results also demonstrated the intervention was effective in reducing anxiety and depression symptoms (i.e. Aim 2). Again, these findings support a prior study testing the effectiveness of reducing mood symptoms using a single shot CBT-I intervention (Boullin, Ellwood & Ellis, 2016). As rates of anxiety and depression are high in prison populations, and in conjunction with insomnia, have been associated with increased suicide risk(Hassan et al, 2013), these findings are particularly relevant in this context. Prison suicide has been a long-standing issue across the globe (Webb, 2017) and to have another potential intervention to help manage suicide risk in this environment is important. Future studies should examine the effectiveness of the present intervention on reducing suicidal thoughts, intentions and actions/attempts as well as other risky behaviours (self-mutilation/cutting, substance use).

This study had several limitations. First, there was no comparator control condition. The decision not to have a control group was made on ethical grounds when studying prisoners (a population deemed “vulnerable” in terms of manipulating for research) with acute insomnia. Whilst this significantly limits our capacity to discuss efficacy, beyond preliminary, this study is an important first step to demonstrate the feasibility of the intervention in this context. Data from Ellis and colleagues (2015) RCT demonstrated a one-month remission rate of 15% in the control group compared to a 60% remission rate in participants who were treated using a similar ‘one-shot’ CBT-I intervention. Further, the second study using the one-shot intervention demonstrated a one-month remission rate, following treatment, of 76.28% which is broadly comparable to the results from the present study. Considering the remission rate in the present study (i.e. 73%), it appears that the intervention is helpful when used in this context, although therapist factors should not be ignored. However, as we do not know the natural remission rate for acute insomnia in a prison population, this suggestion remains speculative. Future research should examine the efficacy of the intervention in a fully powered randomised control trial, perhaps with a full CBT-I arm embedded for those with chronic insomnia. Second, is the follow-up period in the present study. A one-month follow-up is a relatively short time to track both the durability and indeed the potential for relapse following delivery of the intervention. Whilst this was chosen on a largely theoretical basis (i.e. the intervention was designed to prevent from the transition from acute to chronic insomnia) and feasibility given the environmental setting, future studies are needed to determine the long-term impact of the intervention. Other potential factors that could have resulted in the superior outcomes, at least compared to the earlier RCT (Ellis, Cushing and Germain, 2015) include the fact that: i) the sample only included those who were relatively healthy and ii) the availability of the additional support sessions. With regard to the latter issue however although four support sessions were offered, the uptake was limited with 29 prisoners attending an additional session. Unfortunately an analysis as to whether the additional support sessions impacted on treatment cannot be undertaken due to the disparity in who did and who did not utilize those sessions.

A degree of caution should be taken when generalising the findings from the current study to other prison environments. The forensic setting utilised in this study was unique, as it will probably be in most prison settings. Prisons are likely to have differing routines, environmental configurations and different classes of prisoners, severity of crimes/offenses, each of which could impact on both the practicality and acceptability of integrating this intervention. Specifically in the present study the locations where prisoners resided were single cells (i.e. each participant had a cell to themselves, one single bed and a chair). This allowed them to use their cell to complete the behavioural components of the intervention (i.e. Stimulus Control and Sleep Restriction) without disturbing other prisoners. Whilst not entirely comparable, it is known that partners who share a sleep environment can influence the process of full CBT-I(Ellis, Deary & Troxel, 2015) and so it would be interesting to see how the results would fare if incorporated into a shared cell environment. Furthermore, as the present study only used a sample of male prisoners it is unknown whether these effects would be comparable in a women’s prison or juvenile offenders. Considering the prevalence of insomnia is higher in incarcerated women compared to men(Dewa et al, 2017) a comparable study in a women’s prison is needed.

In summary, the identification and management of insomnia is fundamental in the prison environment due to the forensic population being more vulnerable to developing insomnia(Elger, 2004) and the established link between insomnia, poor mood and suicide risk (Carli et al, 2011). The findings from the present study suggest a single-shot of CBT-I, with an accompanying self-help pamphlet, has some promise in this setting for individuals with acute insomnia. Further research into this relatively unchartered area of insomnia treatment, however, is warranted to determine whether these effects are comparable in other closed forensic settings and populations.

REFERENCES

Barker LF, Ireland JL, Cu S, Ireland CA (2016). Sleep and its association with aggression among prisoners: Quantity or quality? IJLP, 47, 115-121.

Bastien, C. H., Vallières, A., & Morin, C. M. (2001). Validation of the Insomnia Severity Index as an outcome measure for insomnia research. *Sleep medicine*, *2*(4), 297-307.

Borbély, A. A. (1982). A two process model of sleep regulation. *Hum neurobiol*, *1*(3), 195-204.

Boullin, P., Ellwood, C., & Ellis, J. G. (2016). Group vs. Individual Treatment for Acute Insomnia: A Pilot Study Evaluating a “One-Shot” Treatment Strategy. *Brain sciences*, *7*(1), 1-11.

Carli, V., Roy, A., Bevilacqua, L., Maggi, S., Cesaro, C., Sarchiapone. M. (2011). Insomnia and suicidal behaviour in prisoners. *Psychiatry Research.* 185, 141-144.

Carney, C. E., Buysse, D. J., Ancoli-Israel, S., Edinger, J. D., Krystal, A. D., Lichstein, K. L., & Morin, C. M. (2012). The consensus sleep diary: standardizing prospective sleep self-monitoring. *Sleep*, *35*(2), 287-302.

Christensen, H., Batterham, P. J., Gosling, J. A., Ritterband, L. M., Griffiths, K. M., Thorndike, F. P., ... & Mackinnon, A. J. (2016). Effectiveness of an online insomnia program (SHUTi) for prevention of depressive episodes (the GoodNight Study): a randomised controlled trial. *The Lancet Psychiatry*, *3*(4), 333-341.

Dewa, L. H., Hassan, L., Shaw, J. J., & Senior, J. (2017). Trouble sleeping inside: a cross-sectional study of the prevalence and associated risk factors of insomnia in adult prison populations in England. *Sleep medicine*, *32*, 129-136.

Dewa, L. H., Kyle, S. D., Hassan, L., Shaw, J., Senior, J. (2015). Prevalence, associated factors and management of insomnia in prison populations: An integrative review. *Sleep Medicine Reviews.* 24, 13-27.

Edinger JD, Wohlgemuth WK, Radtke RA, Marsh GR, Quillian RE. Cognitive behavioral therapy for treatment of chronic primary insomnia: a randomized controlled trial. Jama. 2001;285(14):1856-64.

Elger, B. S. (2004). Prevalence, Types and Possible Causes of Insomnia in a Swiss Remand Prison. *European Journal of Epidemiology.* 19, 665-677.

Elger, B. S. (2009). Prison life: television, sports, work, stress and insomnia in remand prison. *International Journal of Law and Psychiatry* 32, 74-83.

Elger, B. S., & Sekera, E. (2009). Prospective Evaluation of Insomnia in Prison using the Pittsberg Sleep Quality /index: Which are the Factors of Predicting Insomnia? *International Journal of Psychiatry in Clinical Practice.* 13, 206-217.

Elger BS, Goehring C, Revaz SA, Morabia A. 2002. Presciption hypnotics and tranquilisers at the Geneva prison’s outpatient service in comparison to an urban outpatient medical service. SP, 47, 39-43.

Ellis, J. G., Cushing, T., & Germain, A. (2015). Treating acute insomnia: a randomized controlled trial of a “single-shot” of cognitive behavioral therapy for insomnia. *Sleep*, *38*(6), 971-978.

Ellis, J. G., Deary, V., & Troxel, W. M. (2015). The role of perceived partner alliance on the efficacy of CBT-I: preliminary findings from the Partner Alliance in Insomnia Research Study (PAIRS). *Behavioral sleep medicine*, *13*(1), 64-72.

Ellis, J. G., Gehrman, P., Espie, C. A., Riemann, D., & Perlis, M. L. (2012). Acute insomnia: current conceptualizations and future directions. *Sleep medicine reviews*, *16*(1), 5-14.

Eytan, A., Haller, D. M., Wolff, H., Cerutti, B., Sebo, P., Bertrand, D. (2011). Psychiatric symptoms, psychological distress and somatic comorbidity among remand prisoners in Switzerland. *International Journal of Law and Psychiatry.* 34, 13-19.

Feron, J. M., Paulus, D., Tonglet, R., Lorant, V., & Pestiaux, D. (2005). Substantial use of primary health care by prisoners: epidemiological description and possible explanations. *Journal of Epidemiology & Community Health*, *59*(8), 651-655.

Hassan, L., Edge, D., Senior, J., Shaw, J., (2013). Staff and patient perspectives on the purpose of psychotropic prescribing in prisons: care or control? *General Hospital Psychiatry.* 35, 433-438.

Hassan, L., Senior, J., Frisher, M., Edge, D., Shaw, J. (2014). A comparison of psychotropic medication prescribing patterns in East of England Prisons and the general population. *Journal of Psychopharmacology.* 28, 357-362.

Ireland, J., L., Culpin, V. (2006). The Relationship between Sleeping Problems and aggression, Anger and Impulsivity in a Population of Juvenile and Young Offenders. *Journal of Adolescent Health.* 38, 649-655.

Katic, B., Heywood, J., Turek, F., Chiauzzi, E., Vaughan, T. E., Simacek, K., ... & Renger, J. J. (2015). New approach for analyzing self-reporting of insomnia symptoms reveals a high rate of comorbid insomnia across a wide spectrum of chronic diseases. *Sleep medicine*, *16*(11), 1332-1341.

Kjelsberg, E., & Hartvig, P. (2005). Can morbidity be inferred from prescription drug use? Results from a nation-wide prison population study. *European journal of epidemiology*, *20*(7), 587-592.

Kroenke, K., Spitzer, R. L., & Williams, J. B. (2001). The phq‐9. *Journal of general internal medicine*, *16*(9), 606-613.

Löwe, B., Decker, O., Müller, S., Brähler, E., Schellberg, D., Herzog, W., & Herzberg, P. Y. (2008). Validation and standardization of the Generalized Anxiety Disorder Screener (GAD-7) in the general population. *Medical care*, *46*(3), 266-274.

Manber, R., Bernert, R. A., Suh, S., Nowakowski, S., Siebern, A. T., & Ong, J. C. (2014). CBT for insomnia in patients with high and low depressive symptom severity: adherence and clinical outcomes. *FOCUS*, *12*(1), 90-98.

Mitchell, M. D., Gehrman, P., Perlis, M., Umschied, C. A. (2012). Comparative effectiveness of cognitive behavioural therapy for insomnia: a systematic review. *BMC Family Practise.* 13, 40.

Morin, C. M. (2015). Cognitive behavioral therapy for chronic insomnia: state of the science versus current clinical practices. *Annals of internal medicine*, *163*(3), 236-237.

Morin, C. M. (1993). *Insomnia: Psychological assessment and management*. Guilford Press.

Morin, C. M., Belleville, G., Bélanger, L., & Ivers, H. (2011). The Insomnia Severity Index: psychometric indicators to detect insomnia cases and evaluate treatment response. *Sleep*, *34*(5), 601-608.

Morin CM, Colecchi C, Stone J, Sood R, Brink D. Behavioral and pharmacological therapies for late-life insomnia: a randomized controlled trial. Jama. 1999 Mar 17;281(11):991-9.

O'Brien, M. (2001). *Psychiatric morbidity among women prisoners in England and Wales: further analysis of data from the 1997 ONS survey of psychiatric morbidity among prisoners in England and Wales*. Stationery Office.

Ohayon, M. M. (2002). Epidemiology of insomnia: what we know and what we still need to learn. *Sleep medicine reviews*, *6*(2), 97-111.

Ohayon, M. M., & Roth, T. (2003). Place of chronic insomnia in the course of depressive and anxiety disorders. *Journal of psychiatric research*, *37*(1), 9-15.

Okajima, I., Komada, Y., Inoue, Y. (2011). A meta-analysis on the treatment effectiveness of cognitive behavioural therapy for primary insomnia. *Sleep and Biological Rhythms.* 9, 24-34.

Qaseem, A., Kansagara, D., Forciea, M. A., Cooke, M.and Denberg, T. D. (2016). Management of chronic insomnia disorder in adults: a clinical practice guideline from the American College of Physicians. *Ann. Intern. Med.*, **165**: 125–133.

Riemann, D., Baglioni, C., Bassetti, C., Bjorvatn, B., Dolenc Groselj, L., Ellis, J. G., ... & Hertenstein, E. (2017). European guideline for the diagnosis and treatment of insomnia. *Journal of sleep research*, *26*(6), 675-700.

Schutte-Rodin, S., Broch, L., Buysse, D., Dorsey, C., & Sateia, M. (2008). Clinical guideline for the evaluation and management of chronic insomnia in adults. *Journal of clinical sleep medicine: JCSM: official publication of the American Academy of Sleep Medicine*, *4*(5), 487.

Singleton, N., Meltzer, H., Gatward, R. (1998*). Psychiatric morbidity among prisoners in England and Wales.* London: UK.

Spielman, A. J., Saskin, P., & Thorpy, M. J. (1987). Treatment of chronic insomnia by restriction of time in bed. *Sleep*, *10*(1), 45-56.

Smith, M. T., & Perlis, M. L. (2006). Who is a candidate for cognitive-behavioral therapy for insomnia? *Health psychology*, *25*(1), 15.

Spitzer, R. L., Kroenke, K., Williams, J. B., & Löwe, B. (2006). A brief measure for assessing generalized anxiety disorder: the GAD-7. *Archives of internal medicine*, *166*(10), 1092-1097.

Taylor, D. J., Mallory, L. J., Lichstein, K. L., Durrence, H. H., Riedel, B. W., & Bush, A. J. (2007). Comorbidity of chronic insomnia with medical problems. *Sleep*, *30*(2), 213-218.

Taylor, D. J., & Pruiksma, K. E. (2014). Cognitive and behavioural therapy for insomnia (CBT-I) in psychiatric populations: a systematic review. *International review of psychiatry*, *26*(2), 205-213.

Toler, H. C. (1978). The treatment of insomnia with relaxation and stimulus-control instructions among incarcerated males. *Criminal Justice Review.* 5, 117-130.

Trockel, M., Karlin, B. E., Taylor, C. B., Brown, G. K., & Manber, R. (2015). Effects of cognitive behavioral therapy for insomnia on suicidal ideation in veterans. *Sleep*, *38*(2), 259-265.

Webb, M. 2017. Suicide in Prison. The Encyclopedia of Corrections. 1–2.

Wilson, S. J., Nutt, D. J., Alford, C., Argyropoulos, S. V., Baldwin, D. S., Bateson, A. N., ... & Gringras, P. (2010). British Association for Psychopharmacology consensus statement on evidence-based treatment of insomnia, parasomnias and circadian rhythm disorders. *Journal of Psychopharmacology*, *24*(11), 1577-1601.