A powered bone marrow biopsy system versus manual methods: a systematic review and meta-analysis of randomised trials

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ABSTRACT
A significant number of bone marrow biopsies are performed annually. Outcomes of patient pain and sample size of biopsy are important issues as perception of patient pain is underestimated by clinicians and sample size assists in accurate diagnosis. Manual extraction of marrow biopsy is the main method used. Recently a powered system has been introduced. The objectives of this systematic review and meta-analysis were to determine if the powered system reduces patient pain and improves sample capture. A PubMed and Cochrane search for randomised controlled trials was undertaken comparing the powered system with manual methods. Risk of bias assessment and meta-analysis of outcomes using appropriate statistical methods were performed. Five randomised controlled trials were identified. Patient pain (measured via visual analogue scale (VAS)—100 point scale) was significantly reduced using the powered system: mean difference = −6.57; 95% CI −12.93 to −0.22; p = 0.04. The relative reduction in pain was 17%–25% with the powered system. Sample biopsy size (length in mm) was also significantly increased with the powered system: mean difference = 3.65 mm; 95% CI 1.61 mm to 5.68 mm; p = 0.0005. The relative increase in sample size was 33% with the powered system. Operator ease of use (as measured via VAS) and adverse events were similar. Despite limited operator experience, patients experienced less pain and sample sizes were increased without an increase in adverse events with the powered system. The powered system may offer an option in obtaining samples from patients whose pain is compromised by disease and may reduce redos.

INTRODUCTION
While bone marrow biopsy is a relatively common procedure (600 000 US procedures) and straightforward procedure, there are issues associated with it, namely, patient morbidity (eg, haemorrhage, patient pain); adequacy of trephine length (and volume of biopsy sample); and ease of use for the clinician performing the procedure. It has been demonstrated in several trials that the greater the length (in millimetre) of the biopsy sample, the greater the likelihood of lesions being detected. Bone marrow biopsy sampling technologies have until recently mainly consisted of a manual method. This method is most commonly carried out on the posterior superior iliac spine with a patient in the right or left lateral position. Local anaesthesia is provided via infiltrating an adequate area of the periosteum. Commonly a skin excision (with a lancet) is made prior to inserting the biopsy needle. With manual methods, once the needle contacts the bone, it is advanced by slowly rotating clockwise and counterclockwise until the cortical bone is penetrated and the marrow cavity is entered. Ideally the length of the biopsy specimen from an adult should measure at least 20 mm in length.

Recently, a new battery-powered bone marrow biopsy system was developed (OnControl Vidacare Corporation, Shavano Park, Texas, USA) and cleared by the US Food and Drug Administration (510(K) clearance number K072045). The OnControl system uses a battery-powered drill to insert the marrow needle into the iliac bone of adult haematology patients.

Several randomised controlled trials (RCTs) have been undertaken comparing the use of the powered bone marrow biopsy system with manual methods. However, the results from these studies have been equivocal. Further, the RCTs performed to date have been done on a relatively small number of patients. By combining like outcomes from multiple studies, a more definitive conclusion might be made on the utility of a powered bone marrow biopsy system.

The objective of this systematic review and meta-analysis is to determine whether the powered bone marrow biopsy system does provide for significantly different/improved outcomes on the endpoints of pain and sample size.

MATERIALS AND METHODS
RCTs comparing the powered system with manual biopsy methods were included. Participants included both patients with disorders of the blood and bone marrow. Patients were to never have had a previous bone marrow biopsy or other type of biopsies.

The outcome measures of importance that were evaluated were:
- Pain as measured via a visual analogue scale (VAS), either a 10 point or 100 point scale
- Bone marrow length (in mm) or volume (in mm³)
- Complications or adverse events (as defined by the trial)
- Length of time for the procedure (which as per above has been defined as an independent predictor of pain).
**Search method**
The search method used can be found in online supplementary appendix 1.

**Data collection and analysis**
Data collection and analysis used a data collection form identified in online supplementary appendix 2. Data collection was first collected and analysed by one of the authors (JV). It was further adjudicated by the second author (MM) independently. Any disagreements were discussed between the two authors and a final agreement on including/excluding a trial was made by consensus.

**Assessment of risk of bias in included studies**
The risk of bias for each of the studies was reported and is summarised in the risk of bias figure and in online supplementary appendix 3.

Two review authors independently assessed each included study for assessing risk of bias. A risk of bias table was completed for each eligible study. An assessment of risk of bias using a ‘risk of bias summary figure’ was compiled. This display of internal validity indicates the weight the reader may give the results of each study.

**Statistics**
Statistical analysis was performed using the statistical package that accompanies Cochrane Review Manager V.5.1. For continuous data (eg, pain scores, time of procedure, length of trephine), we used the mean difference (inverse variance, random or fixed effects model) if outcomes were measured in the same way between trials. The results of binary outcomes (ie, complication or not) were presented as risk ratios (Mantel-Haenszel, random or fixed effects model) with corresponding 95% CIs.

Attempts were made to contact authors of included studies in order to obtain any missing data and/or to clarify biases. If data remained missing, we left it as missing and did not attempt to impute values. We assumed the data to be missing at random and therefore no bias would be introduced.

If trials could be combined, assessment of statistical heterogeneity was made using the $I^2$ statistic in order to determine appropriateness for meta-analysis. If the $I^2$ statistic was at or below 60%, the heterogeneity was considered moderate and meta-analysis was deemed appropriate. If the value was greater than 60%, sensitivity analyses were undertaken in an attempt to identify which studies were most likely causing the problem and are reviewed in the Discussion section. As well, in examining small size studies and heterogeneity, a comparison of fixed and random effects models was employed. If the estimates were similar, it would be concluded that any small-study effects would have little effect on the intervention effect estimate.

Funnel plots were used to assess reporting bias for those studies combined for meta-analysis purposes. In the absence of bias, the funnel plot should approximate a symmetrical (inverted) funnel. If bias exists, an asymmetrical appearance of the funnel plot would exist. Each outcome was reported separately. Furthermore, an assessment was made of publication bias (including a review of unpublished studies), location bias (types of journals) and language bias.

**RESULTS**

**Results of the search**
The PRISMA flow diagram (figure 1) shows the results of the search.

![PRISMA flow diagram](image)

**Included studies**
Eight studies were included for further evaluation. Of these eight, two were excluded as they were abstracts of follow on published manuscripts. One additional study was excluded as it was an interim analysis of a single arm validation study.

**Risk of bias in included studies**
The main risks of bias: blinding of operators was not possible between the two methods. Further, biases existed related to the experience levels of the operators between the two methods—with minimal to no experience with the powered system and significantly more expertise with the manual method (see figure 2 for risk of bias summary of trials; see online supplementary appendix 3 for additional details of each study). All funnel plots demonstrated symmetry.

**Effects of interventions**
In combining the five trials on the outcome of overall pain, as measured by the patient via the VAS, there was a significant difference in the pain score, favouring less pain with the powered bone marrow system: mean difference $=−6.57$; 95% CI $=−12.93$ to $−0.22$; $p=0.04$; $I^2=25%$ (figure 3) (inverse variance, random effects model). On a relative basis, this represents an approximate 17% reduction in overall pain over the average VAS score for manual biopsy of 39. Further, three trials could be combined on the outcome of pain upon needle insertion. A lessening of pain was observed with the powered bone marrow system: mean difference $=−7.87$; 95% CI $=−15.77$ to 0.44; $p=0.05$; $I^2=23%$ (figure 4) (inverse variance, random effects model). On a relative basis, this represents an approximate 24% reduction in needle insertion pain over the average VAS score for manual biopsy of 33.5. Last, procedure time (in seconds) was significantly less with the powered system:

![Risk of bias graph](image)
mean difference $= -85.35 \text{ s;}$ 95% CI $= -145.14 \text{ s to } -25.56 \text{ s;}$ p=0.005; $I^2= 89\%$ (figure not shown) (inverse variance, random effects model).

Sample biopsy length as measured in millimetres (mm) in all five studies was found to be significantly larger with the powered system: mean difference $= 3.65 \text{ mm;}$ 95%; CI $1.61 \text{ mm to } 5.68 \text{ mm;}$ p=0.0005; although heterogeneity was moderately high at $I^2=62\%$ (figure 5) (inverse variance, random effects model). On a relative basis, this represents an approximate 33% increase in the trephine length over the weighted average mean length for manual biopsy of 10.97 mm. As well, in the two trials$^{14, 19}$ that measured the overall volume in mm$^3$, sample size was found to be significantly larger with the powered system: mean difference $= 18.26 \text{ mm}^3;$ 95%; CI $13.26 \text{ mm}^3$ to $23.26 \text{ mm}^3;$ p<0.00001; $I^2= 0\%$ (figure not shown) (inverse variance, random effects model).

As it relates to adverse events (as defined in the trial, ie, haematoma, extended pain, skin ensnared on shaft of needle), there was no statistically significant difference: risk ratio $= 3.56;$ 95% CI $0.59$ to $21.42;$ p=0.17; $I^2=0\%$ (Mantel-Haenszel, random effects model) (figure not shown).

Last, as it relates to operator satisfaction, there was no statistically significant difference between the two methods in the two trials that measured this via a VAS$^{14, 19}$ mean difference $= 1.41;$ 95%; CI $-0.75$ to $3.56;$ p=0.20; although heterogeneity was high at $I^2=95\%$ (figure not shown).

**DISCUSSION**

Despite very limited experience (eg, <5 uses) by the majority of the operators using the powered bone marrow biopsy system involved in the randomised trials (compared with more extensive experience with manual biopsy methods), it was found that patients experienced significantly less overall pain and operators were able to obtain significantly larger sample sizes with the powered bone marrow biopsy system. Additionally, there was no difference in adverse outcomes between the two methods employed (powered vs manual). These results may speak of an ease of use issue measured in several of the trials (albeit differently) favouring the powered method.

The issue of pain reduction in patients undergoing bone marrow biopsy has been studied extensively in the literature and appears to be one of more important issues with bone marrow biopsy procedures$^{10, 20-22}$ which show that clinicians underestimate the level of pain patients experience.$^{10, 22}$ A technology such as the powered system appears to assist in lowering the levels of pain experienced by patients and may help with this perception mismatch. When examined on a relative basis, this represents a 17%–24% reduction in pain when using the powered system, which could represent the difference between no pain and mild discomfort or mild discomfort and abject pain. This pain reduction may be especially helpful in patients whose pain threshold may be compromised due to disease states such as cancer and the associated treatments they undergo. Additionally, the significantly shorter time to perform the procedure, favouring the powered system, may aid in a reduction of pain.

The amount of bone marrow obtained is also important in helping pathologists accurately assess for the presence/absence of disease. It has been found in several studies that the larger the sample size, the greater the likelihood of accurately identifying lesions.$^{4, 5}$ Larger sample sizes have improved the ‘positivity’ of diagnosis by upwards of 13%–15%.$^{5, 6}$ Further, the informative value of a bone marrow biopsy in detecting discrete marrow lesions is enhanced in a longer sample size due to the fact that it provides a large amount of marrow tissue for histological evaluation and more likely a sample from the innermost part of the bone marrow where abnormalities might be present.$^{23}$ Again the powered system demonstrates that a significantly larger sample is obtained versus manual methods. As identified in the results section, the relative increase in sample trephine length of 33% with the powered system could be the difference in not having to repeat a biopsy and/or in accurately diagnosing a disease state. Last, the improved amount of sample capture with the powered system may reduce the need for repeat procedures as noted above.

Ease of use issues is also important for those performing the bone marrow biopsy. While not mentioned in the peer reviewed literature, performing manual biopsy procedures on a regular basis can result in operator problems such as carpal tunnel syndrome and wrist sprains, especially in patients with hard bones.$^{24}$ Perhaps the powered system might alleviate this issue.

**Heterogeneity issues**

Two of the comparisons resulted in high heterogeneity between the studies: time in seconds for the procedure and sample length (in mm). Regarding the time in seconds meta-analysis,
the 2011 Miller trial was excluded from the analysis with the heterogeneity statistic reduced to 0% (with findings significant at the p<0.00001 level). Regarding the sample length size meta-analysis, the 2013 Bucher study was excluded and the heterogeneity statistic reduced to 31% (with findings significant at the p<0.00001 level). The findings from both Miller et al and Bucher et al while favouring the powered system were not as different in their magnitude as the other studies that were included. Further, additional analyses related to each, that is, pain (measured via a VAS) (with time in seconds of the procedure an independent predictor of pain) and volume of sample (measured in mm³, also related to sample length) demonstrated a significant difference, favouring the powered system. Based on these additional findings, it was assumed that the findings of increased sample length and time in seconds for the powered procedure were statistically different and the findings homogeneous.

Quality of the evidence
In evaluating the risk of bias graphs, the overall quality of the evidence might be considered average—with a similar amount of low and high risk of bias findings from all the studies. This review only included English language publications.

CONCLUSIONS
Despite limited operator experience, this systematic review and meta-analysis demonstrates a lowering of patient pain and an improved amount of sample capture with the powered system, with similar complications and adverse events. It also demonstrates, even with limited operator powered system experience, that it is easy to use.

Figure 5 Forest plot overall trephine length (mm).

and approved the final version of the review prior to submission. MM: conceived the statistical analysis protocol and reviewed all of the statistical analysis prior to submission. Both authors are guarantors of the review.

Competing interests Jeff Voigt is a reimbursement consultant for the manufacturer of the powered bone marrow biopsy system. Michael Mosier has no interests to declare.

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Correction

J Voigt, M Mosier. A powered bone marrow biopsy system versus manual methods: a systematic review and meta-analysis of randomised trials. J Clin Pathol 2013;66:9 792–796. Figure 5 in this paper was incorrect. It depicted adverse events identified and did not represent the overall trephine length analysis. The correct Figure 5 is shown below.

![Corrected Figure 5](image_url)