REVIEW ARTICLE

Can we predict suicide and non-fatal self-harm with the Beck Hopelessness Scale? A meta-analysis

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ABSTRACT

Background. Hopelessness is considered a pre-eminent risk factor for suicide and non-fatal selfharm. We aimed to quantify the ability of the Beck Hopelessness Scale (BHS) to predict these two outcomes.

Method. Medline, Embase, PsycINFO and Cinahl were searched to January 2006. We included cohort studies in which the BHS was applied and patients were followed-up to establish subsequent suicide or non-fatal self-harm. Four studies provided usable data on suicide, and six studies provided data on non-fatal self-harm. Summary sensitivity, specificity, likelihood ratios and diagnostic odds ratios (DORs) were calculated for each study. Random effects meta-analytic pooling across studies at the standard cut-off point (\geq 9) was undertaken and summary receiver operating characteristic (ROC) curves constructed.

Results. For suicide, pooled sensitivity was 0.80 [95% confidence interval (CI) 0.68-0.90], pooled specificity was 0.42 (95% CI 0.41-0.44), and the pooled DOR was 3.39 (95% CI 1.29-8.88). For non-fatal self-harm, pooled sensitivity was 0.78 (95% CI 0.74-0.82), pooled specificity was 0.42 (95% CI 0.38-0.45), and the pooled DOR was 2.27 (95% CI 1.53-3.37).

Conclusion. The standard cut-off point on the BHS identifies a high-risk group for potential suicide, but the magnitude of the risk is lower than previously reported estimates. The standard cut-off point is also capable of identifying those who are at risk of future self-harm, but the low specificity rate means it is unlikely to be of use in targeting treatment designed to lower the rate of repetition.

INTRODUCTION

Hopelessness, defined as generalized negative expectations of the future (Beck *et al.* 1974), has been described as one of the most important long-term risk factors for suicide in clinical populations (Joiner *et al.* 2005). In a study of clinical decision making, psychiatrists gave hopelessness the greatest weighting of any single risk factor, greater even than a previous suicide attempt or depressed mood (Truant *et al.* 1991). Current practice guidelines suggest that hopelessness should be examined when assessing suicide risk and, if present, should be targeted as part of a comprehensive treatment plan (American Psychiatric Association, 2003).

The Beck Hopelessness Scale (BHS; Beck *et al.* 1974) is the only widely used standardized measure of hopelessness (Glanz *et al.* 1995). It is a 20-item scale with a true–false response format (nine items are keyed false, 11 true); each response is summed to give a severity rating ranging from 0 to 20, with high scores indicating the presence of hopelessness. Its originators,

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Beck and colleagues, report that it successfully predicts suicide in a sample of out-patients (Brown *et al.* 2000) and in-patients (Beck *et al.* 1985). The authors acknowledge that the low base rate of suicide means that any scale is likely to identify a large number of false positives, but they argue that the BHS may be useful in identifying a group who are at an increased risk of that outcome (Beck *et al.* 1990). In support of this they found that people scoring above the standard cut-off point on the scale (a score of 9 or more) were 11 times more likely to die by suicide than those scoring below that point.

Ten per cent of people who self-harm will eventually die by suicide, and about a quarter of suicides will have been preceded by an episode of non-fatal self-harm in the previous year (Owens & House, 1994). Although non-fatal self-harm and suicide can be distinguished on some risk factors, such as the patterning of psychiatric disorders, many of the risk factors appear to be common to both (Beautrais, 2001). The link between non-fatal self-harm and suicide may indicate that the BHS is also a predictor of non-fatal self-harm, a suggestion supported by a number of studies (Evans *et al.* 2004).

Although earlier reviews found only limited evidence for the effectiveness of interventions designed to reduce the rate of self-harm repetition (Hawton et al. 1998), recent reviews are more optimistic (Hepp et al. 2004; Mann et al. 2005). There are now a number of treatment approaches that appear to reduce repetition, including cognitive behaviour therapy (CBT; Brown et al. 2005), brief psychodynamic-interpersonal therapy (Guthrie et al. 2001), and individual and group psychoanalytic psychotherapy (Bateman & Fonagy, 1999). The base rate of self-harm is substantially higher than suicide, particularly among some clinical groups such as those who have had a previous episode of non-fatal selfharm. This higher base rate and the emergence of effective interventions raises the possibility that standardized scales, such as the BHS, may have a role in targeting interventions at those who are most at risk of repeating self-harm.

Some doubts, however, have been raised about the utility of the BHS. Not all studies have found that the scale successfully predicts suicide or self-harm. Beck *et al.* (1989), for example, found that the scale did not predict suicide in an alcohol-abuse group. Hawton *et al.* (1999) used the BHS in a group of adolescents who had selfharmed and found that it did not predict repetition. In those studies that have found it to predict suicide or self-harm, the information needed to comprehensively assess the performance of the test is often not reported. Although a number of studies report statistical significance, few report details of sensitivity and specificity analyses, the recommended method for assessing test performance (Sackett et al. 1991). Sensitivity is the proportion of the group with the outcome (e.g. suicide, non-fatal self-harm) that the predictive instrument correctly classifies as at risk. Specificity is the proportion of the group without the outcome correctly classified as not at risk. Moreover, studies of suicide that report these data have found the BHS to have moderate to low specificity at high levels of sensitivity (Glanz et al. 1995), a particular problem when the base rate of the outcome is likely to be low, as it is with suicide.

Over the past decade the medical literature has seen an increased use of meta-analytic strategies to assess the diagnostic and predictive accuracy of tests (Deville et al. 2002), but as yet examples in the psychiatric literature are rare. This is the first systematic review to use these techniques to assess the accuracy of the BHS as a predictor of non-fatal self-harm and suicide. The review aimed to answer three questions. First, to what extent does the standard cut-off point on the BHS identify a high-risk group for suicide? Second, does this standard cut-off point identify a high-risk group for non-fatal selfharm? Third, is the BHS a useful method of identifying those people who are most at risk for self-harm so that they can be targeted for intensive treatments designed to lower that risk?

METHOD

Data sources

The search for primary studies used four electronic databases: Medline, Cinahl, Embase and PsycINFO. Each database was searched from its earliest date to January 2006. MESH terms, individualized for each database, and a list of keywords were developed for suicide and non-fatal self-harm (e.g. overdose, parasuicide, self-injury, self-poisoning, suicide) and cohort design (e.g. longitudinal, follow-up, prospective). An adapted version of an optimal search strategy for diagnostic tests (Deville *et al.* 2002) was also used. 'Hopelessness' was not included as a search term because during piloting we identified a number of studies that met inclusion criteria but made no reference to hopelessness or synonym in the title or abstract. Instead, the initial set of retrieved abstracts was examined to assess whether the study used a cohort design and had suicide or self-harm as one of the outcomes.

Study selection

Of studies that met this initial inclusion criteria, we obtained copies of the full article if either the title or abstract referred to hopelessness or the reference list for the study, accessed through Web of Knowledge, included one of the two standard methods of referencing the BHS (BHS manual or original published description of the measure; Beck *et al.* 1974). The reference list of included studies was also examined to search for additional papers that met inclusion criteria.

In cases where more than one paper referred to the same sample, we selected the one reporting data on the largest number of people. Data from studies with smaller samples were used only if it was not possible to obtain the information necessary to quantify predictive accuracy on a larger dataset.

The full articles were examined independently by two reviewers and were included if the study used a cohort design, the BHS was administered at time 1, data were available on suicide or selfharm at time 2, and the number of people with the outcome (suicide or self-harm) was at least 10. The definition of non-fatal self-harm used in the selection of studies was based on the definition of parasuicide developed for the World Health Organization/European Study on Parasuicide (Platt *et al.* 1992). No inclusion criteria were imposed in terms of patient population or length of follow-up. The weighted κ for the decision to include a study was 1.0.

Data extraction

Two reviewers independently undertook data extraction and quality assessment using a standardized form. Information extracted included details of the referral source, sampling procedure, sample size, sample characteristics, length of follow-up, and the assessment of predictive accuracy of the BHS. Intra-class correlation coefficients (ICCs) were used to assess the reliability. ICCs ranged between 0.79 and 1.0, and the majority (73%) were above 0.90.

We constructed 2×2 tables for a cut-off point of 9 and used these to calculate sensitivity, specificity and likelihood ratios (positive and negative) (Sackett et al. 1991). We summarized likelihood ratios in preference to predictive value because these are fundamental attributes of an instrument that do not vary according to baseline risk of an outcome. We also calculated the diagnostic odds ratio (DOR), that is the ratio of the odds of a positive test among those with subsequent self-harm/suicide to the odds of a positive result among those without subsequent self-harm/suicide (Deville et al. 2002). If the study met inclusion criteria but did not provide sufficient information to quantify predictive accuracy, we contacted authors to request this additional information.

Quantitative data synthesis

We undertook a random effects meta-analysis (DerSimonian & Laird, 1986) to obtain pooled estimates of sensitivity, specificity, positive and negative likelihood ratios, and a summary DOR. Between-study heterogeneity was assessed using the I^2 statistic (Higgins *et al.* 2003), which describes the percentage of total variation across studies that is due to heterogeneity rather than chance. The I^2 statistic has several advantages over other measures of heterogeneity, including greater statistical power to detect clinical heterogeneity when fewer studies are available. As a guide, I^2 values of 25% may be considered 'low', 50% 'moderate' and 75% 'high'.

Publication and small study bias was examined by constructing Begg funnel plots (Begg, 1994) of the logit DOR, and by testing for funnel plot asymmetry using Egger's weighted regression test (Egger *et al.* 1997).

Receiver operating characteristic (ROC) curves represent the most informative way of describing the inherent trade-offs between sensitivity and specificity for a diagnostic test or predictive instrument (Knottnerus, 2002). Single plots of sensitivity and specificity were therefore calculated in ROC space for all studies. We then constructed summary ROC (sROC) curves (Deeks, 2000) within the range of data, using the method of Moses *et al.* (1993). Inverse variance

Study	Country, population and sample size	No. (%) with outcome	Maximum follow-up (years)	Sensitivity	Specificity	Likelihood ratio +ve Likelihood ratio -ve	Likelihood ratio –v
Outcome suicide							
Beck et al. (1985)	USA						
	In-patient suicidal ideation $(n = 165)$	11 (6·7)	S,	0.91 (0.59 - 1.0)	0.51(0.43 - 0.59)	1.84(1.44-2.36)	0.18(0.03 - 1.17)
Beck et al. (1990)	USA						
	Out-patient $(n = 1958)$	17 (0-9)	7-5	0.94 (0.71 - 1.0)	0.41(0.39-0.43)	1.60(1.41 - 1.81)	0.14 (0.02 - 0.96)
Nimeus et al. (1997)	Sweden						
	Previous self-harm $(n=212)$	13 (6-1)	8	0.77 (0.46 - 0.95)	0.42(0.35-0.49)	1.32 (0.96–1.82)	0.55(0.20 - 1.51)
Suominen et al. (2004) ^a	Finland						
	Previous self-harm $(n = 224)$	$17 (7 \cdot 6)$	12	0.60(0.32 - 0.84)	$0.60 \ (0.32 - 0.84) \ 0.52 \ (0.44 - 0.59)$	1.24(0.80 - 1.92)	0.78 (0.41 - 1.46)
Outcome self-harm							
Colman et al. (2004) ^a	Canada						
	Previous self-harm $(n = 369)$	92 (24·9)	1	0.71 (0.62-0.80)	$0.71 \ (0.62 - 0.80) 0.54 \ (0.49 - 0.60)$	1.56 (1.31–1.86)	0.53 (0.39-0.72)
Goldston et al. (2001)	USA						
	In-patient adolescent $(n = 180)$	45 (25.0)	4	0.84 (0.71 - 0.94)	0.49(0.40-0.58)	1.65 (1.34–2.03)	0.31 (0.16 - 0.64)
Hawton <i>et al.</i> $(2003)^{a}$	UK	~					~
	Previous self-harm $(n = 111)$	34 (30-6)	1.7	0.77(0.59 - 0.90)	0.30(0.20-0.41)	$1 \cdot 10 (0 \cdot 87 - 1 \cdot 40)$	0.76(0.36 - 1.58)
Keller & Wolfersdorf (1993)	Germany						
	In-patient depression $(n = 61)$	10 (16-4)	1	0.80(0.44 - 0.98)	0.29(0.18-0.44)	1.13 (0.79–1.62)	0.68 (0.18-2.52)
Sidley et al. (1999) ^a	UK						
•	Previous self-harm $(n = 65)$	25 (38.5)	1	0.75 (0.55-0.89)	$0.75 \ (0.55-0.89) 0.36 \ (0.19-0.56)$	1.17 (0.83–1.65)	$0.70 \ (0.31 - 1.58)$
Tyrer et al. $(2003)^a$	UK						
	Previous self-harm $(n = 430)$	183 (42.6)	1	0.81 (0.74-0.86) 0.28 (0.22-0.34)	0.28 (0.22-0.34)	1.12 (1.01–1.25)	0.69 (0.48 - 0.99)

of logits of the DOR were used to assign weights to each individual study. We also calculated a pooled area under the curve (AUC) with 95% confidence intervals (CIs) from our modelled sROC curve. An AUC value approaching 1.0 represents a perfect test and 0.5 is an uninformative test (Hanley & McNeil, 1982).

Where there was significant between-study heterogeneity, we sought to explore its causes. We first visually inspected forest plots to look for obvious outliers and we constructed Galbraith plots of the logit of the DOR to identify studies lying outside of ± 2 pooled standard deviations (Galbraith, 1988). We also attempted to undertake a meta-regression analysis of our logit sROC model using potential predictive covariates (Thompson & Higgins, 2002), and examined the effect of these in reducing between-study heterogeneity, evidenced by reductions in the I^2 statistic. A priori causes of heterogeneity examined in these analyses included the study population, length of followup and baseline risk of self-harm/suicide. All meta-regression analyses were conducted using a permutation test to minimize type 1 errors. with 1000 Monte Carlo replications to generate p values (Higgins & Thompson, 2004).

We conducted meta-analysis using STATA version 8 (StataCorp, College Station, TX, USA) and the user-written commands: metan, metareg, metabias and galbr.

RESULTS

Description of included studies

The initial broad search of abstracts identified over 28 000 references, of which 1600 met initial inclusion criteria (cohort design, suicide or self-harm as an outcome). Sixty-eight studies consisting of 51 independent samples either mentioned hopelessness in the abstract or included a reference to the BHS in the reference list. Of these, 19 met full inclusion criteria (cohort study, BHS measured at time 1, suicide or self-harm measured at time 2, *n* with outcome ≥ 10). Five of the 19 provided sufficient information to quantify predictive accuracy; for a further five we were able to obtain these data by contacting the authors.

Table 1 summarizes the characteristics of the 10 studies. Four studies (n=2559) provided data on suicide. Three had a similar rate of

suicide in the range 6-8% (Beck et al. 1985; Nimeus et al. 1997: Suominen et al. 2004). Participants in these studies had been admitted to hospital because of severe suicidal ideation or an episode of self-harm and so represent a particularly high-risk group for suicide. The one study with a substantially lower rate (0.9%)used a sample recruited from an out-patient setting (Beck et al. 1990). The length of followup varied substantially within three of the four studies depending on when a person had been recruited. Each of the four samples had more females than males; the proportion of females in the samples ranged from 54% (Beck et al. 1985) to 58% (Beck et al. 1990). All of the studies used adult samples.

The remaining six studies (n = 1216) provided data on the prediction of self-harm. Four of the samples were recruited from attendances at an emergency department after an episode of self-harm (Sidley et al. 1999; Hawton et al. 2003; Tyrer et al. 2003; Colman et al. 2004); in these studies the BHS was used to predict repetition. The two remaining studies (Keller & Wolfersdorf, 1993: Goldston et al. 2001) used in-patient psychiatric samples not recruited on the basis of previous self-harm. Most studies used a follow-up period of 1 year; only one study used a substantially longer follow-up (Goldston et al. 2001). The proportion of females in the studies ranged from 45% (Sidley et al. 1999) to 68% (Tyrer et al. 2003). With the exception of one study, which used an adolescent sample (Goldston et al. 2001), all examined self-harm in adults. Three of the six studies also provided information on people who had died by suicide during the follow-up period. In all of the studies the number was small (range 2-7), and these were included as part of the outcome group in the analyses reported here.

Quantitative analysis of suicide prediction

When we combined psychometric attributes across the four suicide studies, we found low to moderate between-study heterogeneity, except in the case of specificity, and no evidence of publication or small study bias (p = 0.12). Pooled sensitivity was 0.80 (95% CI 0.68–0.90, $I^2 = 57\%$) and specificity was 0.42 (95% CI 0.41–0.44, $I^2 = 76\%$). The likelihood ratio for a positive test was 1.55 (95% CI 1.31–1.83, $I^2 = 44\%$) and the likelihood ratio for a negative

test was $0.45 (95\% \text{ CI } 0.20-1.03, I^2 = 49\%)$. The pooled DOR was $3.39 (95\% \text{ CI } 1.29-8.88, I^2 = 37\%)$ (see Fig. 1 and Table 2).

When we summarized individual studies within ROC space, we found tests were gathered around the top-middle section of the graph, and a summary ROC curve provides a line of best fit, with 95% confidence limits, between studies for the cut-off point of 9 or over (see online Appendix). The pooled AUC was 0.70 (95% CI 0.59-0.85). The small number of studies meant that we were unable to explore sources of between-study heterogeneity using meta-regression.

Quantitative analysis of non-fatal self-harm prediction

For the self-harm studies, we found low to moderate between-study heterogeneity, again except for specificity, and no evidence of publication or small study bias (p=0.93). Pooled sensitivity was 0.78 (95 % CI 0.74–0.82, $I^2 = 0$ %) and specificity was 0.42 (95% CI 0.38-0.45, $I^2 = 90\%$). The pooled likelihood ratio for a positive test was $1.29 (95\% \text{ CI } 1.09 - 1.52, I^2 = 74\%)$ and the likelihood ratio for a negative test was $0.58 (95\% \text{ CI } 0.47-0.71, I^2 = 0\%)$. The pooled DOR was 2.27 (95% CI 1.53–3.37, $I^2 = 35\%$) (see Fig. 2 and Table 2). We conducted a sensitivity analysis by removing the results of Goldston et al. (2001), which used an adolescent population; the results were not substantially altered by the removal of this study (sensitivity = 0.77, 95% CI 0.72-0.81; specificity = 0.40, 95% CI 0.37-0.44; positive likelihood ratio = 1.22, 95% CI 1.04-1.44; negative likelihood ratio = 0.61, 95% CI 0.49-0.76). We also conducted a further analysis in which we removed data from both Goldston et al. (2001) and Keller & Wolfersdorf (1993), which, unlike the other four studies, were not predicting the repetition of non-fatal self-harm. The removal of these two studies also produced similar results to the full analysis (sensitivity = 0.77; 95% CI 0.72-0.81; specificity = 0.41, 95% CI 0.37-0.45; positive likelihood ratio = 1.23, 95% CI 1.02–1.49; negative likelihood ratio = 0.61, 95% CI 0.49-0.76).

When we summarized individual studies within ROC space, we found tests were gathered within the top-left corner of the graph, and a summary ROC curve provides a line of best fit,

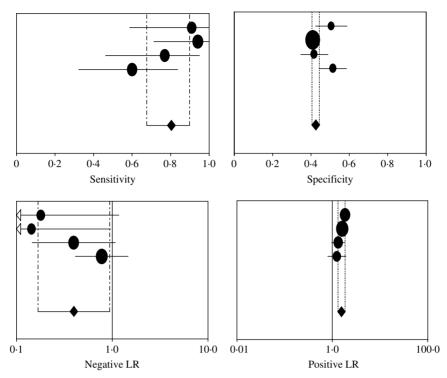


FIG. 1. Random effects pooled sensitivity, specificity and likelihood ratios (LR) for suicide studies. (Data from the following studies: Beck et al. 1985, 1990; Nimeus et al. 1997; Suominen et al. 2004.)

with 95% confidence limits, between studies for the cut-off point of 9 or over (see online Appendix). The pooled AUC was 0.63 (95% CI 0.57-0.70).

When we conducted a meta-regression, we found that study setting (emergency department versus in-patient ($\beta = 0.74$, 95% CI -0.51 to 0.99, p = 0.18, $I^2 = 93$ %), length of follow-up ($\beta = 0.31$, 95% CI -0.16 to 0.79, p = 0.14, $I^2 = 93$ %) and baseline risk ($\beta = -0.04$, 95% CI -0.10 to 0.03, p = 0.19, $I^2 = 71$ %) were not significantly related to the DOR.

DISCUSSION

This is the first systematic review to provide a quantitative summary of the performance of the BHS in predicting suicide and self-harm. We identified four cohort studies that examined suicide (total n=2559) and six that examined non-fatal self-harm (n=1216).

The accurate prediction of suicide is inevitably compromised by the low base rate of the event, which leads to substantial over-inclusion even for scales with high sensitivity and specificity. However, as Beck et al. (1990) argue, the BHS is designed to identify a *potential* for suicide, rather than behaviour itself. In support of this, they found that people scoring 9 or above on the measure were 11 times more likely to kill themselves than those scoring less than 9, which corresponds to the DOR of 11.12 (95% CI 1.45-84.04) calculated in our analysis for that study. The other paper by Beck et al. (1985) reported a similarly high DOR (10.26; 95%) CI 1.28-82.13). However, our main finding for the suicide studies is that the capacity of the BHS to identify this suicide potential is less than that reported in the original validation studies. Two subsequent studies (Nimeus et al. 1997; Suominen et al. 2004) found a notably lower DOR and the pooled DOR (3.39; 95% CI 1.29-8.88) was also substantially lower than that quoted in the research by Beck and colleagues. Although a score of 9 or above on the BHS does confer an increased risk of suicide, and therefore

Outcome	Psychometric attribute	Random effects pooled estimate (95% CI)	I ² between study variance (%)
Suicide (four studies)	Sensitivity	0.80 (0.68-0.90)	57
	Specificity	0.42(0.41 - 0.44)	76
	+ ve likelihood ratio	1.55 (1.31–1.83)	44
	-ve likelihood ratio	0.45 (0.20-1.03)	49
	Diagnostic odds ratio	3.39 (1.29-8.88)	37
Self-harm (six studies)	Sensitivity	0.78 (0.74-0.82)	0
	Specificity	0.42(0.38-0.45)	90
	+ ve likelihood ratio	1.29 (1.09–1.52)	74
	- ve likelihood ratio	0.58 (0.47-0.71)	0
	Diagnostic odds ratio	2.27 (1.53-3.37)	35

 Table 2.
 Random-effects meta-analyses of predictive accuracy

CI, Confidence interval.

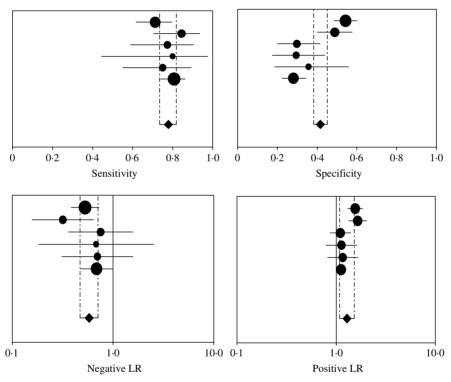


FIG. 2. Random effects pooled sensitivity, specificity and likelihood ratios (LR) for non-fatal self-harm studies. (Data from the following studies: Keller & Wolfersdorf, 1993; Sidley *et al.* 1999; Goldston *et al.* 2001; Hawton *et al.* 2003; Tyrer *et al.* 2003; Colman *et al.* 2004.)

identifies a suicide-potential group, the magnitude of that risk appears to be less than previously reported. There were too few studies to explore how methodological differences between the studies could account for this difference. However, the two later studies were conducted in Scandinavia and used translated versions of the BHS. It is possible that the psychometric properties of the translations are substantially different from the English version.

The pooled DOR for the self-harm studies is 2.27 (95% CI 1.53-3.37), which suggests that the standard cut-off point on the BHS does identify a group that is at increased risk of future

self-harm. This does not necessarily suggest. however, that the scale would be of use in identifying a group at high risk of repetition so that it can be targeted for treatment. Previous systematic reviews of treatments for self-harm have concluded that psychological and pharmacological interventions have limited effects on the likelihood of future self-harm (Hawton et al. 1998), but a number of more recent studies and reviews suggest that psychological interventions can significantly reduce the repetition rate (Bateman & Fonagy, 1999: Guthrie et al. 2001: Hepp et al. 2004; Brown et al. 2005; Mann et al. 2005). These treatments are costly and intensive, so it is likely that only a small proportion of a group who may be at risk for self-harm, such as those attending an emergency setting after an initial episode of self-harm, could be provided with these interventions. Effective methods for identifying those at highest risk of repetition may be required to target treatment, and the data from our analysis can be used to assess whether the BHS would be a useful aid to decision making. The six self-harm studies and the four that examine repetition of self-harm in accident and emergency samples produce very similar sensitivity (0.78 and 0.77) and specificity estimates (0.42 and 0.41), but we use the values for the four repetition studies, because these more closely match the prediction situation under discussion. The pooled sensitivity of 0.77 would identify the majority of people who would repeat self-harm in the absence of intensive treatment, but the low pooled specificity rate (0.41)means that the BHS would identify more people as at risk than could be feasibly offered treatment. This can be illustrated by using data on the rate of repetition for people seen in an emergency setting after an episode of self-harm. A systematic review of these data found a repetition rate of 16% over the course of a year (Owens *et al.* 2002). If we combine this rate with the pooled sensitivity and specificity values, then over half (62%) of people would be identified as at risk of repetition according to a cut-off point of 9 on the BHS, a higher figure than could be offered an intensive treatment such as CBT or psychodynamic-interpersonal therapy. It remains possible that the BHS may be of use if a higher cut-off point were to be selected, but this cannot be assessed from the data reported here.

It is also unclear whether the BHS at the standard cut-off point would be of use as a screening tool to identify people for a more intensive risk assessment. Screening tests with high sensitivity may be acceptable even when accompanied by low or moderate specificity, because the subsequent stage can be used to more clearly differentiate those at risk from those who are not. However, this may not apply in this decision-making context. The screening test typically needs to produce a higher level of sensitivity than that reported for the BHS at a cut-off point of 9. At this cut-off point, about one-quarter of people who would subsequently self-harm would be classified as not at risk, a miss rate that may be considered unacceptably high. Furthermore, in the typical multi-stage screening process, the subsequent stages of assessment are expected to deliver a much more accurate assessment of risk than the initial screen, but it is unclear whether such a test exists for non-fatal self-harm. No other psychometric measure has markedly outperformed the BHS and the limited evidence on clinical prediction for this type of risk suggests that a clinical interview may not improve accuracy (Janofsky et al. 1988).

We note that nine of the 19 studies identified as containing potentially useful data could not be included in the analysis because, despite our best efforts in corresponding with authors, we were unable to obtain the necessary information required to construct 2×2 tables for the metaanalysis. As may be expected, we found it difficult to obtain data from studies conducted some time ago, either because we were unable to contact the authors or because the authors stated that the original data were no longer available. Although the absence of these nine studies may have introduced a selection bias, it is difficult to speculate on whether this would have improved or worsened the pooled sensitivity and specificity values. Specific guidelines on the publication of diagnostic and predictive studies have been published to improve the quality of data that are made available (Bossuyt et al. 2003). Adherence to these guidelines will make it easier to carry out future reviews of the performance of the BHS.

The small number of studies included in our analyses meant that we were unable to conduct a meta-regression for suicide and had to be conservative in our use of predictive covariates in our meta-regression for non-fatal self-harm. We probably lacked sufficient statistical power to detect a substantial effect of population and study setting characteristics in predicting the psychometric performance of the BHS instrument. For example, we did not find a relationship between length of follow-up and psychometric properties, when one might logically have been expected to exist. A useful future exercise will be to repeat the analytic methods outlined in this paper as and when further research emerges relating to the predictive properties of the BHS. A greater number of individual studies with a greater range of settings, populations and periods of follow-up will allow the causes of underlying heterogeneity of the performance characteristics to be explored in more detail using techniques such as meta-regression.

This systematic review confirms that hopelessness, as measured by the BHS, is a risk factor for both suicide and non-fatal self-harm. However, our findings suggest two important caveats to this general conclusion. First, the extent to which a score of 9 or above on the BHS indicates an increased risk of suicide is notably lower than that reported in the original validation studies. Second, the standard cut-off point on the BHS is unlikely to be of use in identifying a group for treatment designed to reduce the rate of repetition of non-fatal selfharm, because that cut-off point would identify a higher proportion of people as at risk than could be feasibly offered treatment.

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DECLARATION OF INTEREST

None.

NOTE

Supplementary information accompanies this paper on the Journal's website (http://journals. cambridge.org).

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