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Imminent death: clinician certainty and accuracy of prognostic predictions

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ABSTRACT

Objectives To determine the accuracy of predictions of dying at different cut-off thresholds and to acknowledge the extent of clinical uncertainty.

Design Secondary analysis of data from a prospective cohort study.

Setting An online prognostic test, accessible by eligible participants across the UK.

Participants Eligible participants were members of the Association of Palliative Medicine. 99/166 completed the test (60%), resulting in 1980 estimates (99 participants × 20 summaries).

Main outcome measures The probability of death occurring within 72 hours (0% certain survival–100% certain death) for 20 patient summaries. The estimates were analysed using five different thresholds: 50/50%, 40/60%, 30/70%, 20/80% and 10/90%, with percentage values between these extremes being regarded as ‘indeterminate’. The positive predictive value (PPV), negative predictive value (NPV) and the number of indeterminate cases were calculated for each cut-off.

Results Using a <50% versus >50% threshold produced a PPV of 62%, an NPV of 74% and 5% indeterminate cases. When the threshold was changed to ≤10% vs ≥90%, the PPV and NPV increased to 75% and 88%, respectively, at the expense of an increase of indeterminate cases up to 62%.

Conclusion When doctors assign a very high (≥90%) or very low (≤10%) probability of imminent death, their prognostic accuracy is improved; however, this increases the number of ‘indeterminate’ cases. This suggests that clinical predictions may continue to have a role for routine prognostication but that other approaches (such as the use of prognostic scores) may be required for those cases where doctors’ estimates are indeterminate.

INTRODUCTION

Palliative care doctors are frequently asked to make predictions about the future health outcomes for their patients; this is what they base treatment decisions on. One such prediction is about whether a patient is imminently dying. This has been shown to be a difficult task.^{1–3} Towards the end of life, accurate recognition of the dying phase can enable a ‘good death’ to occur, with the patient’s final wishes being achieved, surrounded by loved ones and harmful interventions ceased.^{4–6} These wishes cannot be fulfilled if the clinical team are not able to recognise accurately that the person is dying. Although clinician predictions of survival are frequently inaccurate and overoptimistic,^{7–9} they are still quite well correlated with actual survival and so are still recommended for use in routine clinical practice.¹⁰

The ‘More Care: Less Pathway’ report into the limitations of the Liverpool Care Pathway (LCP) described, among other things, the dissatisfaction of relatives with the information they were given about how long their relative was expected to live. A particular issue arose when patients, who were not in fact imminently dying, were cared for using the LCP. The report included a recommendation that more research was needed into the best way to communicate prognostic uncertainty. A recent report from the Royal College of General Practitioners highlighted how difficult doctors find it to initiate a discussion about dying with people who have a terminal disease.¹¹ The report describes many barriers to this discussion including the fact that some doctors feel uncomfortable discussing uncertainty. For these reasons, it is vital to improve doctors’ ability to recognise when patients are imminently dying in order to initiate conversations about



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prognosis, including the appropriate communication of uncertainty.

There is a lack of evidence on the final days of life on which to base clinical practice^{1,2} and little is understood about how doctors recognise dying.¹² Furthermore, it has been recognised that uncertainty might be an inherent part of predicting dying.^{13,14} For these reasons, we wanted to examine how certain a doctor needs to be that a death is likely to be imminent in order to have the highest diagnostic accuracy.

In a previously reported study, we asked palliative care doctors across the UK to participate in an online prognostic task.³ The task presented 20 patient summaries of real patients who had been referred to specialist palliative care, of whom half died within 72 hours and half survived beyond 72 hours. As part of this task, the doctors were asked to read the summaries and provide a probability estimate of that patient dying within the next 72 hours.

For the purpose of this paper, we analysed the results of the prognostic task to explore the following:

1. Do palliative care doctors use the full range of probability estimates from 0% (certain survival) to 100% (certain death) when asked if a patient is imminently dying?
2. What is the effect on prediction accuracy of using different probability thresholds to predict which patients will or will not die imminently.

METHODS

Study design

This is a secondary analysis of data collected as part of a prospective study to investigate how palliative care doctors recognise imminently dying patients.³ This paper follows Standards for Reporting Diagnostic accuracy studies (STARD) reporting guidelines (see online supplementary file 1).

Participants

All medical doctors who were members of the Association of Palliative Medicine (APM) were invited to participate. Recruitment took place between 22 April 2016 and 1 July 2016.

Patient and public involvement

Members of the South West London Cancer Research Network, consisting of patients and bereaved relatives, reviewed the study documentation for the wider prospective observational study.³

Procedure

An email was distributed via the association secretariat to all APM members containing a web link to an online prognostic test. Consent was sought via a tick box prior to participation. Participants were asked to provide some demographic information about themselves before completing the study.

The prognostic test contained 20 patient summaries, presented as 'vignettes'. Each summary was about a patient who had been referred to specialist palliative

care services, in either a hospice or hospital setting. In addition to a narrative summary of the patient's condition, supplementary information was presented: blood test results, medications and observations such as temperature or oxygen saturation level (where available). All patients were monitored to record if death occurred within 72 hours so that this actual outcome could be compared with the doctors' probability estimates. An example of a patient summary can be seen in online supplementary file 2 to show the level of detail presented.

Participants were asked to read through each patient summary and provide an estimated percentage probability (any number between 0% and 100%) that the patient would die within the next 72 hours.

Test methods

The prognostic test of interest was the percentage estimate of the likelihood of death in the next 72 hours provided by the participants. Previous literature has suggested that 50% is an estimate often used in place of 'I don't know'.¹⁵ Therefore, we started by assuming that every prediction about the probability of imminent death made by a clinician indicated either that the patient was likely to die (if the estimate was greater than 50%) or that the patient was likely to survive (if the estimate was less than 50%). Estimates of exactly 50% were regarded as being 'indeterminate'.

Then, we investigated the effect of classifying patients with scores in between two more extreme values as having an 'indeterminate' prognosis. For instance, we investigated how the accuracy of predictions would be affected if only estimates of imminent death of $\geq 60\%$ were taken to indicate that a patient would be likely to die; an estimate of imminent death of $\leq 40\%$ indicated that a patient would be likely to survive; and an estimate between 41% and 59% was regarded as being 'indeterminate'. This process was repeated for different threshold values. Thus, we also looked at using threshold values of $\leq 30\%$, 31%–69% and $\geq 70\%$; values of $\leq 20\%$, 21%–79% and $\geq 80\%$; and finally at values of $\leq 10\%$, 11%–89% and $\geq 90\%$. In total, we looked at five different ways of grouping the clinicians' survival probabilities. The 30%/70% cut-off range has previously been suggested as the most clinically meaningful grouping.^{16,17}

Analysis

The demographics of the respondents and of the patients described in the vignettes were summarised using mean (SD) and number (%). For each of the 20 patient vignettes, the clinicians' predictions of the probability of imminent death were summarised across all participating clinicians using median and IQR. The clinicians' predictions were then compared with the known patient survival outcomes. The accuracy of their predictions under these different threshold values was summarised in a series of 2×2 tables, showing

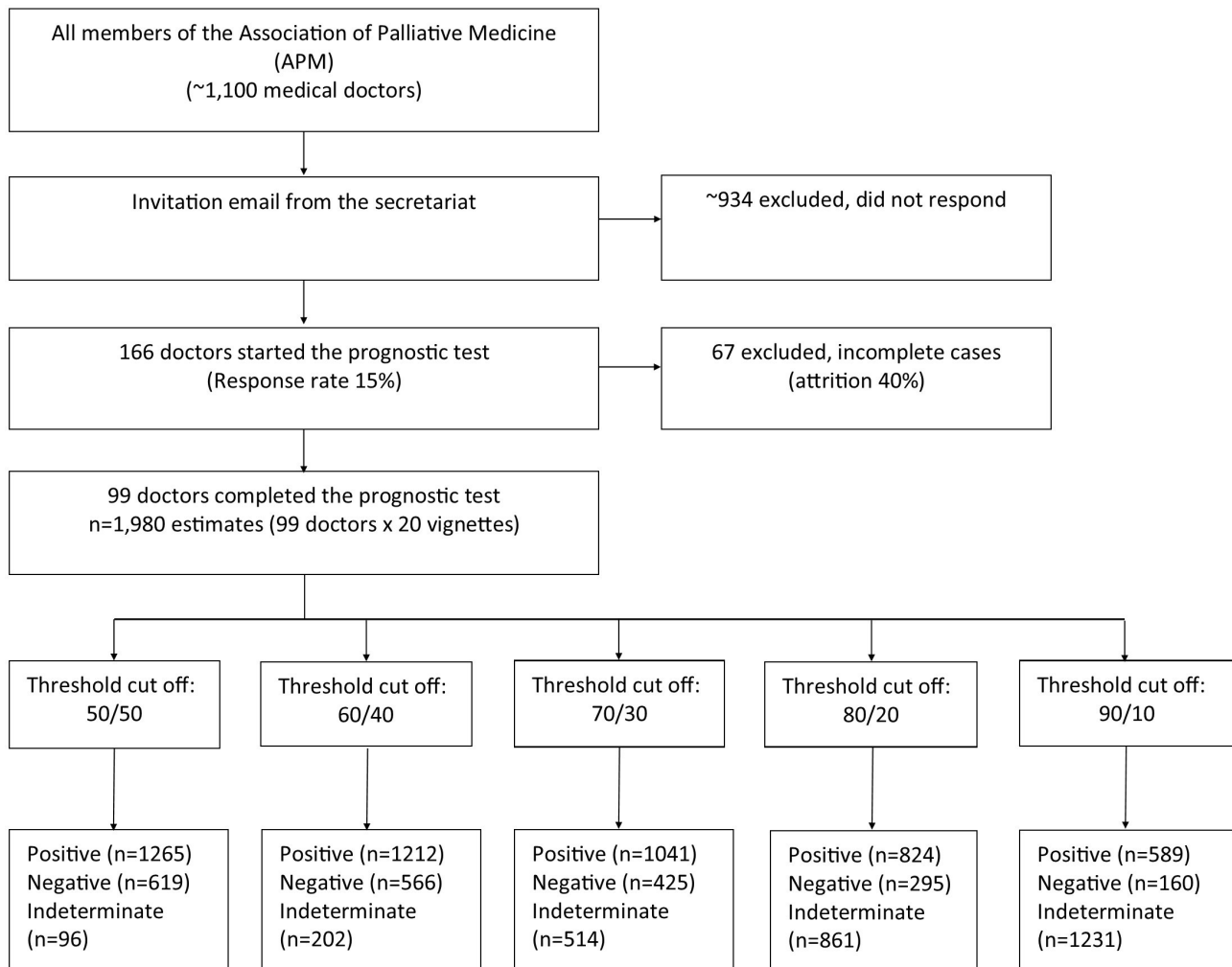


Figure 1 Study flow diagram.

true positives, false positives, true negatives and false negatives (where ‘positive’=death within 72 hours, and ‘negative’=survived 72 hours). The indeterminate predictions were omitted from this analysis. The positive predictive value (the proportion of patients who died when the clinician predicted dying) and the negative predictive value (the proportion of patients who survived when the clinician predicted survival) were calculated.

RESULTS

A total of 166 participants started the test, of whom 99 (60%) provided survival probabilities for all 20 case histories. Data from participants who did not complete all of the cases were excluded from the analysis because they were judged to have lost interest in the task and/or not treated it sufficiently seriously. The majority of non-completers (41/67) stopped the test after reading the first case history. All but two other participants completed less than half of the assessments. Thus, there were 1980 estimates provided by the participants (99 doctors each provided 20 estimates) included in the

analysis. **Figure 1** shows the flow of participants in the study, and the number of predictions of death/survival within 72 hours according to each of the five threshold value groupings. In this figure, ‘positive’ refers to the number of estimates predicting death within 72 hours, and ‘negative’ refers to the number of estimates predicting survival. **Table 1** shows the demographics of respondents. **Table 2** presents a brief demographic background of the

Table 1 Demographics of participants

Demographic	N=99
Gender (n, %)	
Male	21 (21)
Female	78 (79)
Age (mean, SD)	41 (9.5)
Ethnicity (n, %)	
White British	87 (88)
Other	12 (12)
Years since qualified (mean, SD)	17 (10.3)
Years in specialist palliative care (mean, SD)	12 (8.4)

Table 2 Characteristics of the patients presented within the prognostic test

Demographics	n (%)
Total number of participants	20
Gender	
Male	13 (65)
Female	7 (35)
Age (mean, SD)	71.45 (18.88)
Ethnicity	
White British	13 (65)
Other	7 (35)
Cancer diagnosis?	
Yes	14 (70)
No	6 (30)
Charlson score (mean, SD)	5.00 (1.89)
Length of survival	
Within 72 hours	10 (50)
Outside 72 hours	10 (50)

patients presented to the participants within the case summaries.

Use of percentage probability estimates

Figure 2 shows the median and IQR of the probability estimates given by the 99 respondents for each of the 20 vignettes included in the prognostic test. This does not represent the order in which they were presented to the participants. Vignettes 1–10 are those where the patient died within 72 hours. Vignettes 11–20 are those where the patient survived for over 72 hours. For 6 patients, the median predicted probability of dying was between 90% and 100%, 2 patients had a median estimate between 0% and 20% of dying and 12 patients had a median estimate between 20% and 80%.

Participants used the full range of the scale (0%–100%) in their survival predictions. There was one vignette in which the median response by the participants was greater than 90% probability of dying and the patient survived beyond 72 hours.

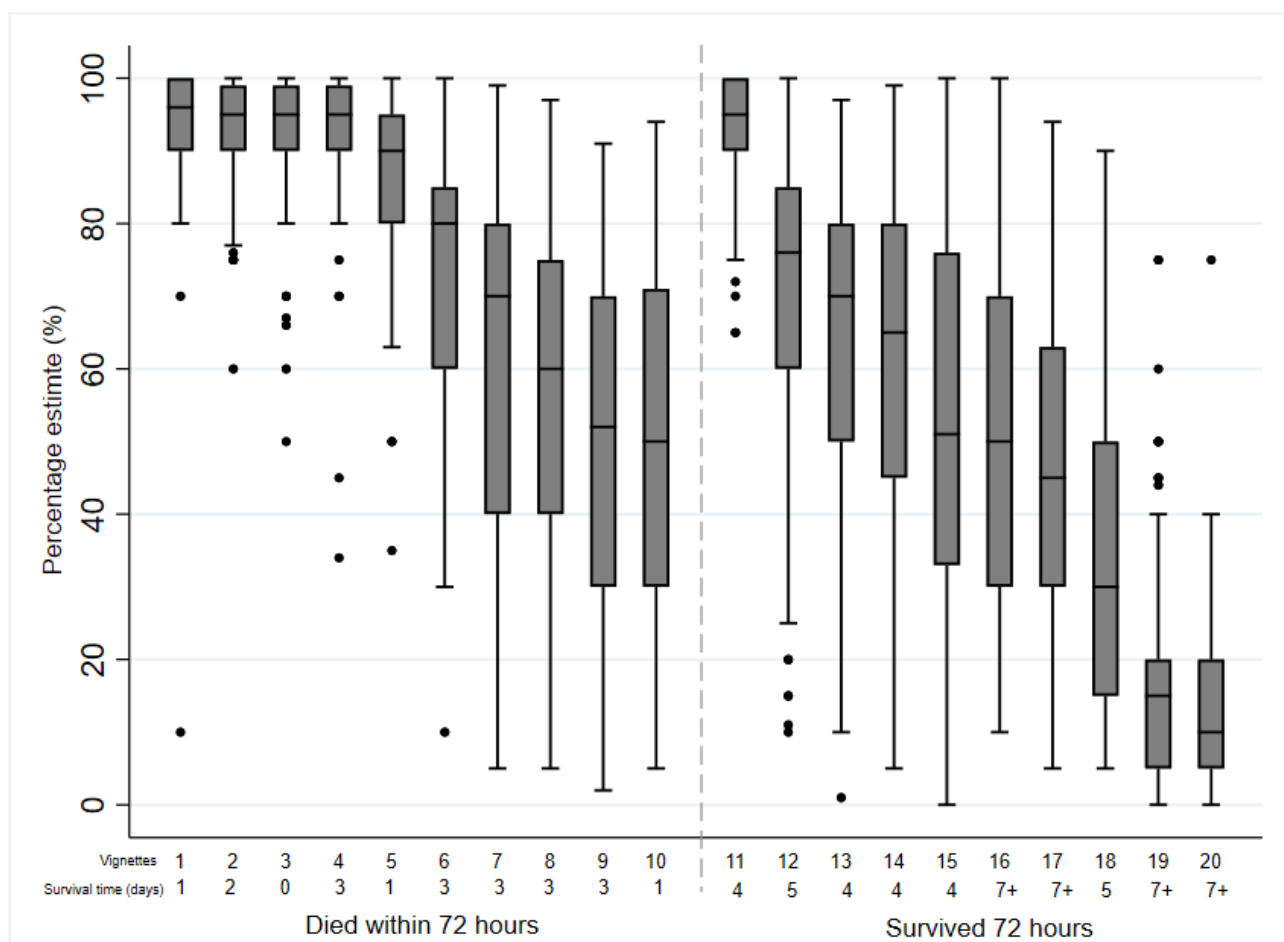
**Figure 2** Boxplot of responses to each vignette.

Table 3 Accuracy of clinician probability estimates using different double thresholds (including indeterminate values)

Estimate cut-off	Prediction*	Outcome (within 72 hours)			Outcomes		Proportion of indeterminate cases
		Died N	Survived	Total	PPV	NPV	
>50%	Die	787	478	1265	62%		
<50%	Survive	163	456	619		74%	
50%–50%	<i>Indeterminate</i>	40	56	96			5%
≥60%	Die	766	446	1212	63%		
≤40%	Survive	144	422	566		75%	
41%–59%	<i>Indeterminate</i>	80	122	202			10%
≥70%	Die	685	356	1041	66%		
≤30%	Survive	87	338	425		80%	
31%–69%	<i>Indeterminate</i>	218	296	514			26%
≥80%	Die	577	247	824	70%		
≤20%	Survive	39	256	295		87%	
21%–79%	<i>Indeterminate</i>	374	487	861			43%
≥90%	Die	440	149	589	75%		
≤10%	Survive	19	141	160		88%	
11%–89%	<i>Indeterminate</i>	531	700	1231			62%

*Each estimate was categorised in to a group if the estimate was above or below the cut-off. Scores that were *indeterminate* were omitted from the analysis. PPV (positive predictive value; the proportion of patients who died when the clinician predicted dying, eg, 787/1265 for the 50% cut-off), NPV (negative predictive value; the proportion of patients who survived when the clinician predicted survival, eg, 456/619 for the 50% cut-off).

The assessment of accuracy

Table 3 shows the accuracy of clinicians' predictions about the probability of imminent death at each of the five thresholds. The resulting number of 'indeterminate' scores is also presented. The positive predictive value of the clinical prediction (ie, the proportion of estimates of imminent death that correctly predicted that the patient would die imminently) improved from 62% (at the 50% threshold) to 75% at the 10%/90% threshold. The negative predictive value of the clinical prediction (ie, the proportion of estimates of survival beyond 72 hours that correctly predicted the outcome) improved from 74% (at the 50% threshold) to 88% at the 10%/90% threshold.

Indeterminate cases

Applying the simplest threshold of <50% or >50%, 96 estimates (5%) were ascribed a probability of exactly 50% and were regarded as being indeterminate. The number of indeterminate cases increased steadily with each change in the thresholds applied until at the ≤10%/≥90% threshold, 1231 estimates were deemed to be indeterminate (1231/1980; 62% of all estimates). In these 1231 indeterminate estimates, 700 (57%) patients survived and 531 (43%) died.

DISCUSSION

The principal finding from this study is that when using a <50% or >50% threshold, and ignoring predictions of exactly 50%, clinical predictions of imminent death are not very accurate (accuracy 66%). However, accuracy could be improved by adjusting the threshold values that are taken to indicate 'likely

death' or 'likely survival'. The best way to maximise accuracy was to only regard those predictions with a probability of ≥90% as indicating imminent death, and only those predictions with a probability of ≤10% as indicating likely survival beyond 72 hours, with all other values in between being regarded as 'indeterminate'. The drawback with this strategy was that a relatively large proportion of clinician estimates would have been regarded as indeterminate. In our study 14/20 vignettes were ascribed a median probability estimate in the range between 10% and 90%, suggesting a majority of predictions would be regarded as uninformative using this approach.

It makes logical sense that the higher the probability estimate provided by the doctor, the more likely it is that the outcome will occur. However, these results illustrate how challenging it is to predict the imminent future, even by specialist palliative care doctors. The results suggest it is important to acknowledge and discuss uncertainty whatever the level of the clinician's estimate, as is illustrated by one of the patient summaries used in the prognostic task. This patient survived for longer than 72 hours despite the median probability ascribed to his or her death being greater than 90%. Perhaps, however, clinical estimates in the range of 11%–89% should be regarded with greater circumspection.

Ultimately, the decision about which is the 'best' threshold to use is a qualitative and subjective one. What this analysis does is to quantitatively illustrate the effect of using different thresholds on the accuracy of clinicians' predictions. Even when using the

highest threshold level ($\geq 90\%$), clinicians' predictions of imminent death were only correct on 75% of occasions. Accuracy gradually increases for every rise in threshold value but at the extreme values the number of indeterminate cases escalates rapidly.

The European Association for Palliative Care (EAPC) has recommended that clinical predictions of survival should still hold an important role in prognostication, while accepting that, taken in isolation, clinical predictions of survival are not very accurate.¹⁰ Other means to estimate prognosis in palliative care patients have been developed^{16–18–20} and are still being evaluated.^{21–23} One question that therefore arises is when and how to combine clinical predictions of survival with prognostic scores, multidisciplinary team (MDT) decisions or algorithms. Our study suggests that clinical predictions of survival are at the most accurate when the thresholds of $\leq 10\%$ and $\geq 90\%$ are employed; however, there are important implications for the number of 'indeterminate' estimates that arise as a result of this.

Strengths and limitations

This is the first study to measure the accuracy of multiple doctors' predictions about the imminent death of the same group of patients, thus allowing a direct comparison of their prognostic accuracy.

The prognostic test could be criticised for lacking some face validity as the doctors did not get to review the patients at a face-to-face assessment. Nonetheless, the vignettes all concerned real cases and included a detailed descriptive summary of the patient including routinely available blood results and other clinical findings. A further limitation of this study is that the findings are based on these specific 20 case histories, which may not be 'representative' of all the patients who will be observed in palliative care. Half of the vignettes in this sample died, and this might not be representative of survival times in palliative care. Further research is required to see if a different set of 20 summaries would produce similar results.

CONCLUSION

Our results show that even when patients are very ($>90\%$) confident that a patient will die within the next 72 hours, these estimates are only correct on 75% of occasions (and at the expense of a high number of indeterminate cases). It is therefore important for clinicians to convey this level of uncertainty in their communications with patients and relatives and not to forget that even their most confident predictions are inaccurate 25% of the time. This suggests that while clinical predictions will continue to have a role for routine prognostication, other approaches (such as the use of prognostic scores) may be required for those cases where doctors' estimates are indeterminate.

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Contributors NW developed the research concept, designed the data collection tools, monitored and collected the data for the study, wrote the statistical analysis plan, cleaned and analysed the data and drafted and revised the paper. She is the guarantor. FR and VV assisted in the study design and the statistical analysis of the data. PH and PS developed the research concept and oversaw the data collection and analysis. CT developed the study website to collect the data and monitored the data collection. All authors reviewed and provided comments on the drafted paper. All authors approve the final version.

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REFERENCES

- 1 Neuberger J, Guthrie C, Aaronovitch D. *More care, less pathway: a review of the Liverpool care pathway*. Department of Health CC, 2013.
- 2 Parry R, Seymour J, Whittaker B, *et al*. Rapid evidence review: pathways focused on the dying phase in end of life care and their key components 2013.
- 3 White N, Harries P, Harris AJL, *et al*. How do palliative care doctors recognise imminently dying patients? A judgement analysis. *BMJ Open* 2018;8.
- 4 National Institute for Health and Care Excellence (NICE). Care of dying adults in the last days of life 2017.
- 5 The End of Life Care Strategy. *Department of Health*, 2008.

- 6 Ellershaw J, Ward C. Care of the dying patient: the last hours or days of life. *Bmj* 2003;326.
- 7 White N, Reid F, Harris A, *et al.* A systematic review of predictions of survival in palliative care: how accurate are clinicians and who are the experts? *Plos One* 2016;11:e0161407.
- 8 Glare *Pet al.* A systematic review of physicians' survival predictions in terminally ill cancer patients. *BMJ* 2003;327.
- 9 Parkes CM. Accuracy of predictions of survival in later stages of cancer. *BMJ* 1972;2:29–31.
- 10 Maltoni M, Caraceni A, Brunelli C, *et al.* Prognostic factors in advanced cancer patients: evidence-based clinical recommendations—a study by the Steering Committee of the European Association for Palliative Care. *J Clin Oncol* 2005;23:6240–8.
- 11 Royal College of Physicians. Talking about dying: how to begin honest conversations about what lies ahead 2018.
- 12 Taylor P, Dowding D, Johnson M. Clinical decision making in the recognition of dying: a qualitative interview study. *BMC Palliative Care* 2017;16.
- 13 Schwartz A, Bergus G. *Medical decision making*. A Physician's Guide Cambridge, Cambridge Medicine, 2008.
- 14 Ruegger J, Hodgkinson S, Field-Smith A, *et al.* Care of adults in the last days of life: summary of NICE guidance. *BMJ: British Medical Journal* 2015;351.
- 15 Bruine de Bruin W, Carman KG. Measuring risk perceptions: what does the excessive use of 50% mean? *Medical Decision Making* 2012;32:232–6.
- 16 Hui D, Kilgore K, Nguyen L, *et al.* The accuracy of probabilistic versus temporal clinician prediction of survival for patients with advanced cancer: a preliminary report 2011;16:1642–8.
- 17 Casarett DJ, Farrington S, Craig T, *et al.* The art versus science of predicting prognosis: can a prognostic index predict short-term mortality better than experienced nurses do? 2012;15:703–8.
- 18 Gwilliam B, Keeley V, Todd C, *et al.* Development of prognosis in palliative care study (PIPs) predictor models to improve prognostication in advanced cancer: prospective cohort study. *BMJ* 2011;343:d4920.
- 19 Morita T, Tsunoda J, Inoue S, *et al.* The palliative prognostic index: a scoring system for survival prediction of terminally ill cancer patients. *Support Care Cancer* 1999;7:128–33.
- 20 Pirovano M, Maltoni M, Nanni O, *et al.* A new palliative prognostic score: a first step for the staging of terminally ill cancer patients. *J Pain Symptom Manag* 1999;17:231–9.
- 21 Kalpakidou AK, Todd C, Keeley V, *et al.* The prognosis in palliative care Study II (PiPS2): study protocol for a multi-centre, prospective, observational, cohort study. *BMC Palliative Care* 2018;17.
- 22 Kim E-S, Lee J-K, Kim M-H, *et al.* Validation of the prognosis in palliative care study predictor models in terminal cancer patients. *Korean J Fam Med* 2014;35:283–94.
- 23 Baba M, Maeda I, Morita T, *et al.* Survival prediction for advanced cancer patients in the real world: a comparison of the palliative prognostic score, Delirium-Palliative prognostic score, palliative prognostic index and modified prognosis in palliative care study predictor model. *Eur J Cancer* 2015;51:1618–29.