

Using the Palliative Performance Scale to Estimate Survival for Patients at the End of Life: A Systematic Review of the Literature

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Abstract

Background: The Palliative Performance Scale (PPS) has been widely used for survival prediction among patients with cancer; however, few studies have reviewed PPS scores in heterogeneous palliative care populations across multiple care settings.

Objective: The aim of this systematic review was to determine how the PPS tool has been used to estimate survival at the end of life.

Methods: This review followed the Preferred Reporting Items for Systematic Reviews and Meta-Analyses guidelines. PubMed, Embase, and the Cochrane Library were searched for the existing literature published from 2008 to 2017. We synthesized study characteristics, the PPS scores at baseline, and primary outcomes, and explored differences in survival estimates by diagnosis. The quality of the studies was assessed using the Good ReseArch for Comparative Effectiveness (GRACE) checklist.

Results: Seventeen studies were included in this review (nine with cancer and eight with mixed diagnoses). All included studies reported that the PPS exhibited a significant association with survival. Survival estimates ranged from 1 to 3 days for patients with PPS scores of 10% compared with 5 to 36 days for those with scores of 30%. The categorical cut-points for the PPS scores were not consistently reported across studies.

Conclusion: This review provides a broad overview on the prognostic value of the PPS tool for survival among multiple patient populations across care settings. Consistent reporting of PPS scores would facilitate the comparison of survival estimates across end-of-life diagnoses.

Keywords: hospices; palliative care; Palliative Performance Scale; prognosis; review; survival

Introduction

KNOWING HOW MUCH TIME is left is a critical issue for terminally ill palliative care patients. Accurate survival prognostication enables patients and their families to make better decisions about goals of care and to prepare for the end of life.^{1,2} It is also important for healthcare providers to proactively refer patients for hospice care at the appropriate time.¹ A clear prognosis helps hospice organizations better allocate the intensity of palliative care services, including end-of-life treatments.³ Given the importance of accurate survival estimates, valid measures are needed.

The Palliative Performance Scale (PPS)⁴ is a reliable and valid tool that has been used to measure functional perfor-

mance of,^{2,7} and to predict survival among, palliative care cancer patients.^{6,10–13} The PPS is a modification of the Karnofsky Performance Scale,⁸ which was originally developed for cancer patients and later adapted to be more generalizable to other end-of-life diagnosis. The PPS scores measure five functional domains: ambulation, activity level and evidence of disease, self-care, oral intake, and level of consciousness (Appendix A). Each of the five domains is divided into 11 levels ranging from 0% to 100% in 10%-point intervals, with 0% indicating death and 100% being fully ambulatory and healthy. The PPS is simple to use in various healthcare settings,^{1,9} and the assessment of the PPS can be conducted by multiple healthcare professionals, including physicians and registered nurses.¹

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This systematic literature review critically evaluates the relationship between PPS scores and length of survival in patients with end-of-life diagnoses. This is particularly relevant because palliative and hospice services have expanded over the last decade to include multiple end-of-life diagnoses after initially focusing on patients with cancer. The aim of this systematic literature review was to determine how the PPS tool has been used to estimate survival at the end of life in both patients with cancer and patients with other end-of-life diagnoses.

Methods

This systematic review was conducted based on the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) guidelines.¹⁶

Search strategy and selection criteria

A search of the literature was conducted between September and October 2017 using the following electronic databases: PubMed, Embase, and the Cochrane Library. The search strategy was developed through a literature review and consultation with an informationist in the health sciences library at Columbia University. The searches used MeSH terms for PubMed, Emtree terms for Embase, and free text terms. Boolean searching techniques using “AND” and “OR” included the following search terms: “palliative performance scale,” “prognosis,” “mortality,” “survival,” “rehospitalization,” and “length of services.” More detailed search terms used for each database are outlined in Appendix B.

Studies were included if they met the following criteria: (1) published between January 2008 and October 2017, (2) written in English, (3) original research study with data, (4) studies that investigated survival times or survival/mortality proportions after the PPS scores were administered, and (5) studies that examined either the prognostic value of the PPS tool for survival or association between the PPS and survival/mortality. Studies were excluded if they (1) provided only a discussion, opinion, commentary, review, or editorial; (2) were a published conference abstract only or presentation slides; (3) examined healthcare providers’ outcomes (e.g., level of knowledge, perspectives) without patient outcomes; or (4) involved patients who were <18 years old.

Data extraction and analysis

Titles were screened (D.B.), and abstracts were independently reviewed by two reviewers (D.B. and R.M.C.) to identify eligible studies. During the abstract review, the two reviewers met and compared opinions on included and excluded articles, and consensus was reached through discussion when there were different opinions. Full-text articles were reviewed by two reviewers (D.B. and R.M.C.). The following data were extracted in the final review and presented in a table of an Excel spreadsheet: (1) author, (2) year of publication, (3) country, (4) study design, (5) study setting, (6) patient characteristics (number of participants at baseline, mean age, gender, race/ethnicity, diagnosis type, and PPS scores at baseline), and (7) patient outcomes measured. After the coding process, the extracted data were analyzed and synthesized using descriptive statistics.

Quality assessment

The Good ReseArch for Comparative Effectiveness (GRACE) checklist¹⁷ was used to assess the quality of the included studies. This tool was chosen because it is validated and designed to systematically assess the rigor of observational research studies.^{17,18} The GRACE checklist consists of 11 items, including 6 related to data and 5 related to methods. The 11 items are scored by dichotomized responses of sufficient (“+”) or insufficient (“-”). Four reviewers (D.B., D.R., L.J., and R.M.C.) independently assessed the quality of the studies. Consensus was reached through discussion when there were discrepancies.

Results

Study selection

A flow diagram illustrating our review process is shown in Figure 1. The initial search yielded 219 articles from three electronic databases. Of the 219 articles, 72 articles were duplicates and removed. Titles of the remaining 147 articles were screened based on the inclusion/exclusion criteria of this review for eligibility. The titles for 53 of these articles passed an initial screening, and the abstracts were more closely reviewed. During the abstract review, 22 articles were excluded because they did not meet the inclusion criteria of this review. The remaining 31 articles were included for the full-text review, and then 14 articles were excluded for several reasons (e.g., not in English, presentation slides). In total, 17 articles were included for the final review.

Study information

All 17 reviewed studies were published between 2009 and 2017 in seven countries, with the majority in Canada ($n=6$; 35%) and the United States ($n=5$; 29%) (Fig. 2A). The most common study design was a prospective study design ($n=9$; 53%), followed by retrospective ($n=8$; 47%). Studies included in this review were conducted in inpatient palliative care units ($n=4$; 23.5%), outpatient/ambulatory care/community-dwelling settings ($n=4$; 23.5%), and mixed palliative care settings ($n=4$; 23.5%), including skilled nursing facilities, inpatient palliative care units, home hospice settings, and outpatient palliative care settings (Fig. 2B).

Overall, nine studies (53%) included patients diagnosed with cancer, and eight studies (47%) included palliative care patients with mixed diagnoses such as cancer, cardiovascular disease, and stroke. The majority of the studies ($n=11$; 65%) recorded patients’ PPS scores by clinicians (physician or registered nurse) who cared for patients in the study settings. Of the remaining six studies, four did not clearly report who measured and recorded the PPS scores, while two studies stated that patients’ PPS scores were recorded by research team members (e.g., physician, registered nurse) who did not involve in caring for patients in the study settings (Tables 1 and 2).

PPS scores at baseline

Of the 17 studies, 13 (76%) reported the detailed PPS scores on admission or during a baseline interview. Of the four remaining studies, most either did not specify the initial PPS scores or reported only the PPS range, mean, or median. Among the 13 studies that reported the details of initial PPS

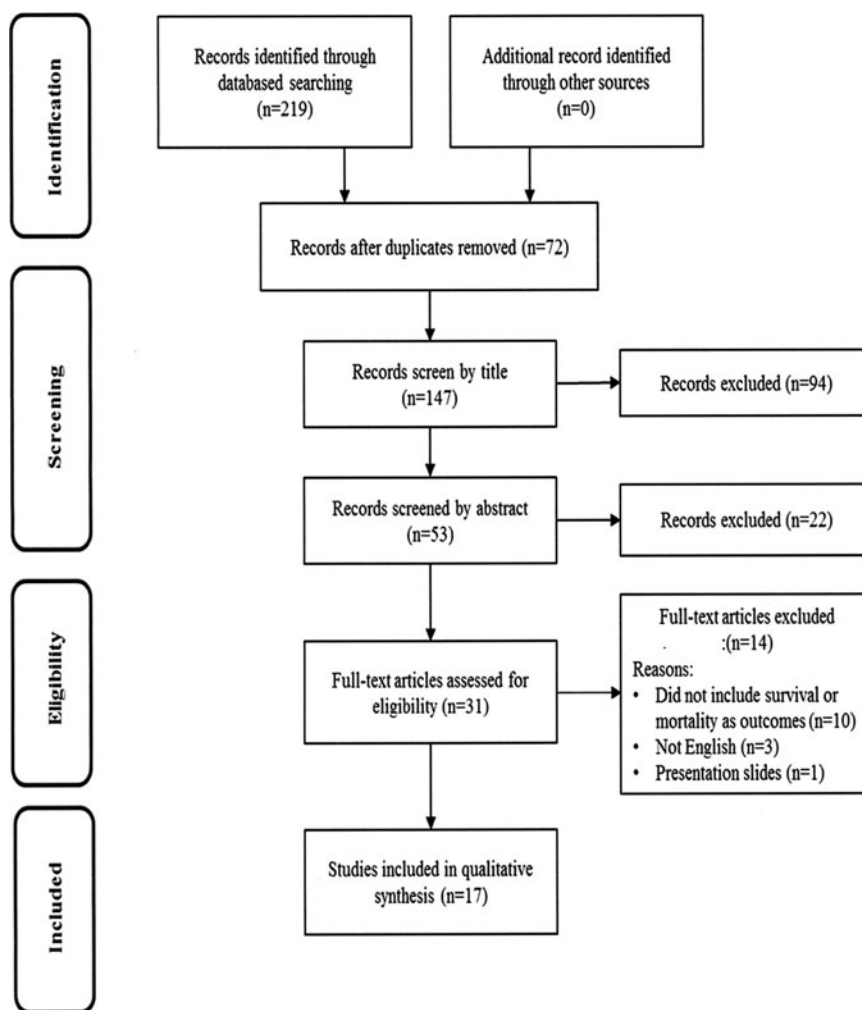


FIG. 1. PRISMA flow diagram. PRISMA, Preferred Reporting Items for Systematic Reviews and Meta-Analyses.

scores, seven^{1,10,14,19–22} reported the initial PPS as a categorical variable that ranged from three to nine cut-points. Each of these seven studies chose different categorical cut-points. For example, Maltoni et al.²⁰ reported three categories (PPS 10–20%, PPS 30–50%, PPS ≥60%). Of the 13 studies, the 6 remaining^{2,5,12,13,23,24} reported the initial PPS scores as discrete variables, which had different ranges of PPS scores (e.g., PPS 10–70%, PPS 40–90%). Of the six studies, two studies^{2,5} reported PPS ranging from 10% to 70%, while the four remaining studies reported various ranges. For example, O'Mahony et al.²³ reported PPS scores ranging from 40% to 90%, while Lau et al.¹¹ reported PPS scores ranging from 10% to 80%.

Patient outcomes measured

The included studies measured patient outcomes as either survival times ($n=14$; 82%) or survival/mortality proportions ($n=3$; 18%). PPS scores were represented as categorical or discrete variables within different cohorts. In this review, survival time was defined as the period between the date of admission or a baseline interview and the date of death. Thirteen studies (76%) included covariates in their prediction models when using the PPS to estimate survival times. The

most common covariate was age ($n=10$), followed by gender ($n=9$) and diagnosis or cancer site ($n=8$).

Survival estimates by PPS scores

As described above, most studies ($n=15$; 88%) reported survival time or proportions predicted by PPS scores. Specifically, survival estimates among five studies with PPS scores of 10% ranged from one to three days.^{1,11–13,25} The survival ranges for patients who scored 30% on the initial PPS were substantially wider (5–36 days). To examine the accuracy of PPS scores in predicting survival, three studies (18%)^{10,20,26} measured the predictive validity of the PPS by using sensitivity, specificity, or the area under the curve (AUC). Mei et al.¹⁰ showed that the study model increased predictive accuracy of survival in patients within 90 days, with a positive predictive value (PPV) of 79% and an AUC of 0.70. Kim et al.²⁶ reported that when the PPS scores are <30%, the predictive accuracy of the PPS for three- and four-week survival was acceptable (AUC = 0.729 and 0.771, respectively). The findings from these two studies differed from Maltoni et al.²⁰ who found that the predictive accuracy of the PPS at 30 days was less than a PPV of 50%.

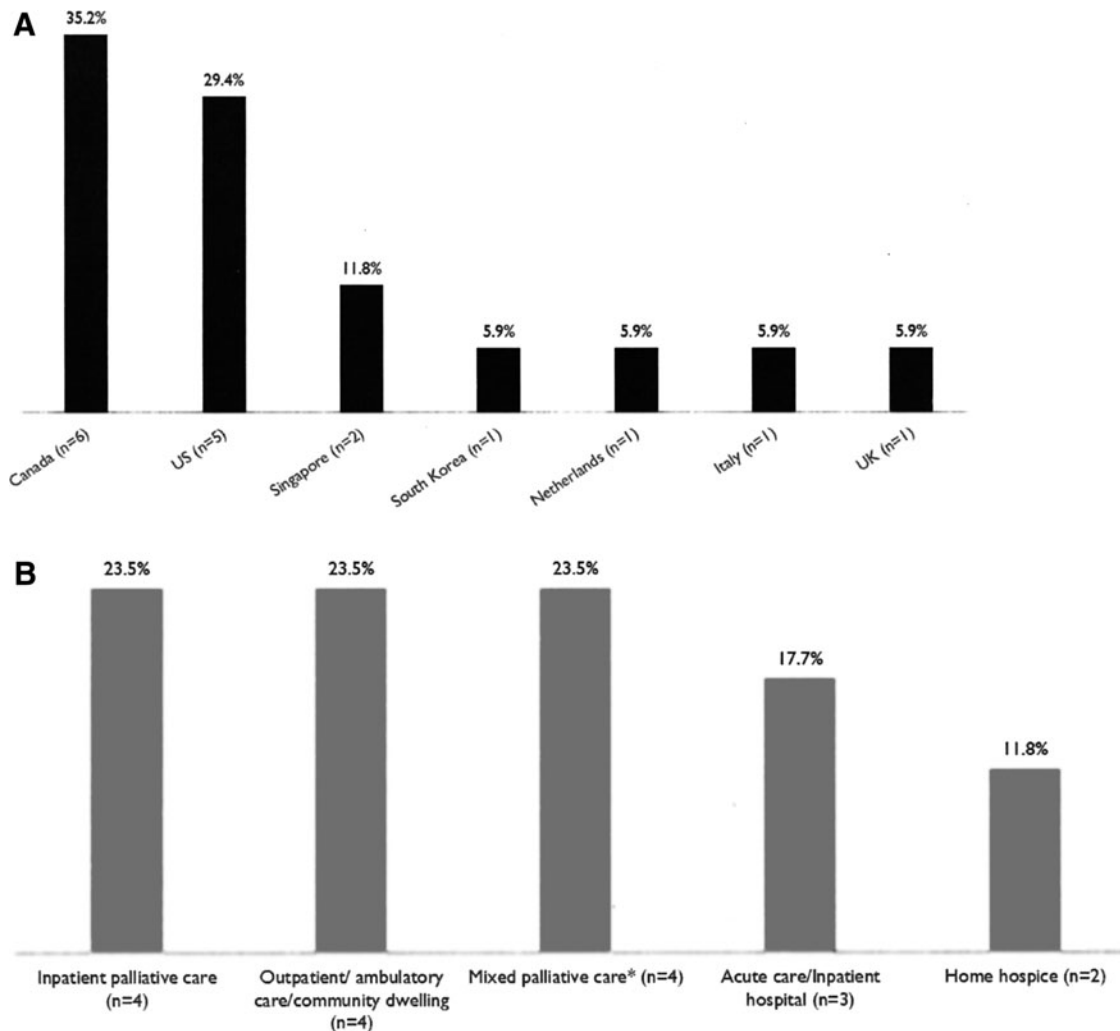


FIG. 2. Characteristics of included studies. **(A)** Countries. **(B)** Study settings. *Mixed palliative care settings included skilled nursing facilities, inpatient palliative care units, home hospice settings, and outpatient palliative care settings.

To better understand the use of the PPS for survival prediction based on patient diagnoses, we analyzed the prognostic value of the PPS tool on survival by dividing all included studies into two diagnosis groups (cancer vs. mixed diagnosis).

Survival estimates for cancer patients

The PPS survival among studies that only included cancer patients ($n=9$) is summarized in Table 1. The studies were most commonly conducted in Canada ($n=3$) and the United States ($n=2$). The most common study settings were outpatient/ambulatory care/community-dwelling settings ($n=4$), followed by inpatient palliative care units ($n=3$). The number of participants at baseline ranged from 62 to 11,342. Of the nine studies, seven reported mean age of patients with cancer, which ranged from 61 to 71 years. Eight studies reported the gender of patients; four studies included a higher percentage of females, and the remaining four studies consisted of a higher percentage of males. Most studies ($n=8$) did not examine the race or ethnicity of study participants; however, one study¹³ conducted in the United States reported that the largest racial or ethnic group was

African American (46.3%), followed by Caucasian (30.9%). All but one of the studies² reported specific types of participants' cancer diagnoses. Of the nine studies that included cancer patients, eight reported initial PPS scores, and each study chose to report different ranges of the PPS scores. Five of these studies reported categorical scoring of the PPS.^{10,14,19–21} However, these scoring categories differed considerably across studies, ranging from three to nine categories. Three studies^{2,13,23} reported the initial PPS scores as a discrete variable and had different ranges of PPS scores in their patient populations.

All nine studies reported detailed survival time stratified by initial PPS scores recorded on admission or during a baseline interview, and the survival time was reported by categorical or discrete variables of the PPS scores within different cohorts. All nine studies reported a significant association between PPS scores and survival among palliative care patients with advanced cancer.

Survival estimates for mixed diagnosis patients

The characteristics among eight of the studies with a mixed diagnosis palliative care sample are summarized in Table 2.

TABLE 1. SUMMARY OF PALLIATIVE PERFORMANCE SCALE SURVIVAL AMONG PATIENTS WITH CANCER (N=9)

Author, country, year	n	Study settings, years of data collection	Age, gender	Diagnoses	PPS recorded by	PPS at baseline (n, %)	Primary outcomes
Jang, Canada, 2014	1655	Oncology Palliative Care Clinic at Princess Margaret Cancer Centre 2007–2010	Median: 65 Male: 809 (49%) Female: 846 (51%)	GI, lung, genitourinary, breast, gynecological, head and neck, CNS, hematological, other	Physician	PPS 10–30%: 24 (1%) PPS 40–50%: 315 (19%) PPS 60–70%: 784 (48%) PPS 80–100%: 508 (31%)	Survival time (median days, 95% CI) PPS 10–30%: 22 (12–102) PPS 40–50%: 51 (44–60) PPS 60–70%: 115 (105–131) PPS 80–100%: 221 (197–244)
Jansen, Netherland, 2015	78	Residential hospice 2010	Mean: 70.6 Male: 47 (60%) Female: 31 (40%)	Terminal primary cancer diagnosis patients	Physician	PPS 10%: 1 (1%) PPS 20%: 14 (18%) PPS 30%: 8 (10%) PPS 40%: 21 (27%) PPS 50%: 21 (27%) PPS 60%: 10 (13%) PPS 70%: 3 (4%)	Survival time (median days, IQR) PPS 10–20%: 6 (4–11) PPS 30–50%: 17.5 (11.75–41.5) PPS >50%: 41 (22.5–75)
Kim, South Korea, 2014	415	Kyungpook National University Hospital and Medical Center 2009–2011	Mean: 60.7 Male: 240 (57.8%) Female: 175 (42.2%)	Terminal cancer patients (stomach, lung, colon, biliary track, pancreatic, liver, uterine cervical, breast, ovarian, prostate)	Not reported	Not recorded	Survival time PPS <30%: 3–4 weeks PPS cutoff point for survival <3 and <4 weeks was 30 or below AUC 3 weeks: 0.729 (95% CI 0.684–0.772) AUC 4 weeks: 0.771 (95% CI 0.727–0.810)
Maltoni, Italy, 2012	549	Three Italian hospices 2009–2010	Median: 71 Male: 269 (49%) Female: 280 (51%)	Gastrointestinal, respiratory, genitourinary, other	Physician	PPS 10–20%: 227 (41.3%) PPS 30–50%: 314 (57.2%) PPS ≥60%: 8 (1.5%)	Survival time (median days, 95% CI) PPS 10–20%: 11 (9–13) PPS 30–50%: 32 (28–41) PPS ≥60%: 139 (67, n.r.)
Mei, Singapore, 2013	296	Palliative Care Unit of Tan Tock Seng Hospital 2009–2011	Mean: 68.9 Male: 159 (53.72%) Female: 137 (46.28%)	Breast, CUP, GI, GU, head and neck, MSK pulmonary, skin	Not reported	PPSV2 10–30%: 140 (39.32%) PPSV2 40–60%: 199 (55.90%) PPSV2 70–100%: 17 (4.78%)	Survival time (median days) PPS 10–30%: 19 PPS 40–60%: 43 PPS 70–100%: 59
Myers, Canada, 2015	368	Outpatient Palliative Care Clinic within the Odette Cancer Centre at Sunnybrook Health Sciences Centre 2011–2013	Mean: 64.8 Male: 160 (44%) Female: 208 (56%)	Gastrointestinal, breast, lung, gynecological, genitourinary, head and neck; melanoma, hematology, unknown prim, other	Physician	PPS ≤40%: 18 (5%) PPS 50%: 68 (18%) PPS 60%: 113 (31%) PPS 70%: 99 (27%) PPS 80%: 59 (16%) PPS 90%: 11 (3%) Median PPS: 60	Survival time (median days) PPS 40%: 28 PPS 50%: 71 PPS 60%: 104 PPS 70%: 173 PPS 80%: 197 PPS 90%: 226

(continued)

TABLE 1. (CONTINUED)

<i>Author, country, year</i>	<i>n</i>	<i>Study settings, years of data collection</i>	<i>Age, gender</i>	<i>Diagnoses</i>	<i>PPS recorded by</i>	<i>PPS at baseline (n, %)</i>	<i>Primary outcomes</i>
O'Mahony, United States, 2016	62	Outpatient oncology clinic at University Medical Center (dates not recorded)	Mean: 63.7 Gender: unspecified	Lung	Physician	(range 20–90) PPS 40%: 1 (1.61%) PPS 50%: 10 (16.31%) PPS 60%: 11 (17.74%) PPS 70%: 31 (50%) PPS 80%: 5 (8.06%) PPS 90%: 4 (6.45%)	Survival time Changes in PPS scores between baseline and 1 month were not significantly associated with shorter survival time PPS was associated with survival time when stratified by median score at baseline
Seow, Canada, 2013	11,342	Ambulatory setting 2007–2009	Mean: 65 Male: 4144 (52.58%) Female: 3738 (47.42%)	Breast, CNS, gastrointestinal, genitourinary, gynecologic, hematology, head and neck, other, primary unknown sarcoma, skin, lung	Physician or nurse	PPS ≤20%: 40 (0.3%) PPS 30%: 109 (1%) PPS 40%: 263 (2.3%) PPS 50%: 800 (7.1%) PPS 60%: 1222 (10.8%) PPS 70%: 1794 (15.8%) PPS 80%: 2105 (18.6%) PPS 90%: 2431 (21.4%) PPS 100%: 2578 (22.7%)	Survival time (median days) PPS 40%: 154 PPS 50%: 231 PPS 60%: 315 PPS 70%: 441 PPS 80%: 686 PPS 90%: >800
Weng, United States, 2009	492	Home hospice program 2005–2006	Mean: 67 Male: 219 (44.5%); Female: 273 (55.5%)	Lung, GI, GU, breast, hematological, neurological head and neck, muscle, other	Nurse	PPS 10%: 32 (6.5%) PPS 20%: 35 (7.1%) PPS 30%: 57 (11.6%) PPS 40%: 181 (36.8%) PPS 50%: 122 (24.8%) PPS 60%: 44 (8.9%) PPS 70%: 18 (3.7%) PPS 80%: 2 (0.4%) PPS 90%: 0 (0%) PPS 100%: 1 (0.2%)	Survival time (median days, SD) PPS 10%: 3 (6.2) PPS 20%: 5 (33.5) PPS 30%: 10 (19.6) PPS 40%: 19 (54.4) PPS 50%: 29.5 (59.6) PPS 60%: 35 (50.8) PPS 70%: 48 (60.7) PPS 80%: 17.5 (4.9)

AUC, area under the curve; PBS, Palliative Performance Scale.

TABLE 2. SUMMARY OF PALLIATIVE PERFORMANCE SCALE SURVIVAL AMONG PATIENTS WITH MIXED DIAGNOSES (N=8)

Author, country, year	n	Study settings, years of data collection	Age, gender	Diagnoses	PPS recorded by	PPS at baseline, (n, %)	Primary outcomes
Babcock, United States, 2016	123	Emergency department in John Dempsey Hospital 2013	Mean: 78.7 Male: 48 (39%) Female: 75 (61%)	Cardiac disease: 33 (27%) Infectious disease: 25 (21%) Abdominal pain: 20 (16%) Fracture: 6 (5%) Dehydration: 6 (5%) Cancer: 5 (4%) COPD: 3 (2%) Anemia: 3 (2%) Renal failure: 3 (2%) Miscellaneous: 19 (15%)	Research team member (physician)	PPS mean: 70.7% PPS range: 30–100%	Three-month survival proportions PPS 0–30%: 42.9% PPS 40–60%: 85.1% PPS 70–100%: 97.1% Six-month survival proportions (mean days) PPS 0–30%: 14.3% (107) PPS 40–60%: 74.5% (151) PPS 70–100%: 95.7% (175)
Chan, Singapore, 2013	400	Tan Tock Seng Hospital (acute care hospital) 2009–2010	<45:10 (2.5%) 45–64: 99 (24.8) 65–74: 104 (26%) 75–84: 126 (31.5%) ≥85: 61 (15.3%) Male: 176 (44%) Female: 224 (56%)	Cancer: 340 (85%) Noncancer: 60 (15%)	Research team member (registered nurse)	PPSv2 10%: 15 (3.8%) PPSv2 20%: 9 (2.3%) PPSv2 30%: 104 (26%) PPSv2 40%: 104 (26%) PPSv2 50%: 87 (21.8%) PPSv2 60%: 44 (11%) PPSv2 ≥70%: 37 (9.3%)	Survival time (median days, 95% CI) PPSv2 10%: 2 (1.33, 2.67) PPSv2 20%: 8 (0.0, 16.77) PPSv2 30%: 23 (10.51, 35.49) PPSv2 40%: 25 (17.89, 32.14) PPSv2 50%: 43 (28.04, 57.96) PPSv2 60%: 41 (23.36, 58.64) PPSv2 ≥70%: 116 (45.49, 186.51)
Downing, Canada, 2010	3328	Victoria Hospice Society Palliative care unit 1993–2005	Mean: 70.2 Male: 1595 (48.1%) Female: 1723 (51.9%)	Cancer: 2939 (88.3%) Noncancer: 389 (11.7%)	Physician or nurse	PPSv2 10%: 221 (6.6%) PPSv2 20%: 335 (10.1%) PPSv2 30%: 1028 (30.9%) PPSv2 40%: 1098 (33%) PPSv2 50%: 508 (15.3%) PPSv2 60%: 128 (3.9%) PPSv2 70%: 10 (0.3%)	Survival time (median days) ^a PPS 30%: ~17 PPS 40%: ~30 PPS 50%: ~50 PPS 60%: ~65
Harris, United States, 2014	118,532	Home, nursing home or hospice, hospital/inpatient unit in New Mexico, California, Florida, Pennsylvania, Wisconsin, Michigan, Ohio, Texas, and Kansas/Missouri 2008–2012	≥65: 97,068 (81.9%) <65: 21,464 (18.1%) Male: 53,944 (45.5%) Female: 64,588 (54.5%)	Cancer: 45,601 (35.8%) Debility: 13,466 (11.4%) Cardiovascular disease: 16,522 (13.9%) Dementia: 11,931 (10.1%) Pulmonary: 8037 (6.8%) Stroke: 5559 (4.7%) Other: 17,416 (14.7%)	Not reported	PPS <30%: 26,005 (21.9%) PPS 30%: 22,227 (18.8%) PPS 40%: 24,792 (20.9%) PPS 50%: 19,900 (16.8%) PPS ≥60%: 5731 (4.8%) Missing: 19,877 (16.8%)	Six-month mortality Six-month mortality probability ranged from 32.6% to 99.5%. PPS scores, OR (95% CI) <30: reference 30: 0.22 40: 0.10 50: 0.07 >60: 0.04

(continued)

TABLE 2. (CONTINUED)

<i>Author, country, year</i>	<i>n</i>	<i>Study settings, years of data collection</i>	<i>Age, gender</i>	<i>Diagnoses</i>	<i>PPS recorded by</i>	<i>PPS at baseline, (n, %)</i>	<i>Primary outcomes</i>
Lau, Canada, 2009	6066	Victoria Hospice palliative care, home or palliative inpatient unit 1993–2005	Mean: 71.5 Male: 2892 (47.7%) Female: 3156 (52.0%) Unknown: 18 (0.3%)	Cancer: 5097 (84.0%) Noncancer: 756 (12.5%) AIDS: 75 (1.2%); Cardiovascular: 328 (5.4%) Neurological: 69 (1.1%) Respiratory: 157 (2.6%) Others: 127 (2.1%) Not recorded: 213 (3.5%)	Physician or nurse	Not recorded	Survival time (median days, 95% CI) PPS 10%: 1 (1–1) PPS 20%: 2 (2–2) PPS 30%: 5 (5–5) PPS 40%: 13 (12–14) PPS 50%: 28 (25–31) PPS 60%: 43 (38–48) PPS 70%: 63 (48–78)
Lau, Canada, 2009	513	William Osler Health Centre in Toronto, Canada (inpatient and outpatient units) 2005–2006	Mean: 74.91 Male: 256 (49.9%) Female: 257 (50.1%)	Cancer: 347 (67.6%) Noncancer: 166 (32.4%)	Physician or nurse	PPSv2 10%: 27 (5.3%) PPSv2 20%: 71 (13.8%) PPSv2 30%: 118 (23%) PPSv2 40%: 56 (10.9%) PPSv2 50%: 88 (17.2%) PPSv2 60%: 56 (10.9%) PPSv2 70%: 81 (15.8%) PPSv2 80%: 16 (3.1%)	Survival time (median, days, 95% CI) PPSv2 10%: 2 (1–3) PPSv2 20%: 6 (4–8) PPSv2 30%: 12 (9–15) PPSv2 40%: 31 (15–47) PPSv2 50%: 35 (29–41) PPSv2 60%: 50 (33–67) PPSv2 70%: 110 (77–143) PPSv2 80%: 71 (0–196)
Linklater, United Kingdom, 2012	666	Specialist palliative care unit, Primary, Secondary, nursing home (dates not recorded)	Unspecified Male: 281 (42.2%) Female: 385 (57.8%)	Malignant: 397 (59.6%) Nonmalignant: 163 (24.5%) Dementia/frailty: 106 (15.9%)	Physician or nurse	Not recorded	Survival time (median days, IQR) days PPS 10%: 3 (1–11) PPS 30%: 35.5 (17.25–123.5) PPS 50%: 52 (18–141) PPS 100%: 138 (115.5–335.5)
McGreevy, United States, 2017	153	Surgical ICU 2012	Mean: 70.8 Male: 93 (60.8%) Female: 60 (39.2%)	Traumatic brain injury	Not reported	PPS 50%: 6 (3.9%) PPS 60%: 9 (5.9%) PPS 70%: 20 (13.1%) PPS 80%: 23 (15%) PPS 90%: 41 (26.8%) PPS 100%: 54 (35.3%)	In-hospital mortality In-hospital mortality: 28 (18.3%) In a model adjusting for age, injury severity score, and traumatic brain injury, PPS was a significant predictor of in-hospital mortality

^aEstimated survival time in days based on Kaplan–Meier survival curves in the text.

The included studies were conducted in Canada ($n=3$), the United States ($n=3$), Singapore ($n=1$), and the United Kingdom ($n=1$). The most common study settings were mixed palliative care settings ($n=4$), followed by acute care units/inpatient hospitals ($n=3$). The number of participants in these studies ranged from 123 to 118,532. Of the eight studies, five^{5,9,11,12,24} reported the mean age of study participants ranging from 70 to 79 years. Seven^{1,5,9,11,12,22,25} of the eight studies included a higher percentage of females. Three studies^{9,22,24} conducted in the United States examined the race or ethnicity of the study participants, and revealed that the majority of the study participants were Caucasian.

Of the eight studies, five examined detailed initial PPS scores, and reported them with different ranges of the scores across the studies. Two studies^{1,22} reported categorical initial PPS scores using five and seven categories, respectively. Three studies^{5,24,27} reported the initial PPS scores as a discrete variable, and each chose different ranges of PPS scores (e.g., PPS 10% ~ PPS 70%, PPS 10% ~ PPS 80%, and PPS 50% ~ PPS 100%).

Regarding patient outcomes estimated by categorical or discrete variables of the PPS scores, most studies ($n=6$, 75%) measured survival time or proportions, and the remaining two studies^{22,24} analyzed mortality rates. All eight studies revealed that the PPS was a significant predictor for survival time or mortality rate.

Discussion

In this systematic literature review, we critically evaluated the relationship between PPS scores and survival times or proportions by synthesizing research across diverse care settings and analyzing differences in the characteristics of studies between two diagnosis groups (cancer vs. mixed diagnosis). The PPS was a significant predictor of survival for patients with both cancer and other end-of-life diagnoses. As such, this review confirms previous study findings among patients with cancer⁷ and supports the use of the PPS tool for estimating the length of survival among patients with mixed end-of-life diagnoses such as heart disease, pulmonary disease, and dementia.

Most studies included in this review were conducted in Canada and in the United States. This was consistent with findings from a previous review,⁷ and is most likely explained by the fact that the PPS was developed at the Victoria Hospice Society in Canada.²⁵ Specifically, because there were already established research teams in Canada, the evidence base establishing the use of PPS as a tool for estimating survival is larger compared with other countries.

Both prospective and retrospective designs were used in the studies included in this review. Prospective designs were defined as studies in which researchers recruited participants to assess the PPS at baseline and collected follow-up data about survival length or proportions. Retrospective designs were defined as studies in which the researchers used medical record data to define PPS scores and survival/mortality after the point at which the patient had received treatment. The retrospective studies may be limited by less accurate measurements of PPS scores compared with prospective studies because trained researchers could not interview palliative care patients.²⁸

Across studies, there was wide variation in how the PPS scores and subsequent survival were reported. Some studies examined the PPS scores using categorical variables, while others used discrete measures of the PPS. Specifically, studies reporting the categorical PPS scores included various numbers of categories and cut-points. Studies reporting the discrete measure of the PPS also described different ranges of the PPS and subsequent survival estimates. Moreover, these studies did not present the rationale regarding how or why they chose ways to categorize PPS scores or decided to use certain ranges of PPS scores. Downing et al.¹⁵ also reported the need for the consistency in reporting survival probabilities and length of survival based on the PPS scores. The lack of consistency in categorization of PPS scores hampers our ability to draw comparisons between studies utilizing survival analysis among different palliative care populations. Downing et al.¹⁵ suggested the following recommendations for consistently reporting survival based on the PPS scores: (1) reporting median survival times according to PPS levels; (2) using PPS quartiles with confidence intervals to convey information about early, mid, and late survivors; or (3) establishing a life table from PPS survival curves.

A majority of the studies focused on examining the association between the PPS and survival, while only three studies investigated the predictive accuracy of the PPS by using sensitivity, specificity, or AUC. Examining the optimal cutoff points of the PPS for the prediction of survival is essential to provide helpful information about when and which diagnoses the PPS tool is effectively predictive of survival. More studies investigating the predictive accuracy of the PPS in estimating survival across heterogeneous palliative care populations are needed to help healthcare providers predict accurate survival times using the PPS tool.

Limitations

This review included only studies written in English; thus, it may have missed studies that report the relationship between the PPS and survival in non-English-speaking countries. The PPS scores were not reported consistently across studies; thus, the heterogeneity in how PPS scores were reported in relation to survival times/proportions limited our ability to conduct a meta-analysis. Inconsistencies may also be introduced in survival estimation through the recent reorientation of palliative care away from specialized clinicians trained in palliative medicine toward a “palliative approach” in which palliative knowledge and expertise is integrated into models of care that do not formally specialize in palliative care.²⁹ This issue may be an important topic for future reviews investigating the association of PPS scores with survival among patients nearing the end of life.

Conclusion

The findings from this review confirm that the PPS tool has been used more frequently to estimate survival in patients with cancer, and that there is a lot of opportunity to apply the tool to palliative care patients with different end-of-life diagnoses. In studies that include the PPS as a measure, there should be more consistency in how it is reported to support

generalizability of the measure across study settings and disease diagnoses. Additional research is needed to determine evidence-based cut-points for different patient populations. Moreover, further research is needed in countries where the PPS is less widely used, such as countries in South America, Africa, Europe, and Asia, to obtain more reliable and validated measures for survival prediction by considering different races, ethnicities, and cultures. Specifically, Europe mainly uses the Karnofsky Performance Scale, and Japan incorporates the PPS into the Palliative Prognostic Index.³⁰ More research is also needed on how the PPS tool can be used in other underrepresented palliative care patient populations such as those with heart disease. Finally, we recommend that the PPS is used in acute care or home care settings to identify those who should receive palliative care services earlier.

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APPENDIX A. PALLIATIVE PERFORMANCE SCALE

PPS level (%)	Ambulation	Activity and evidence of disease	Self-care	Intake	Conscious level
100	Full	Normal activity & work No evidence of disease	Full	Normal	Full
90	Full	Normal activity & work No evidence of disease	Full	Normal	Full
80	Full	Normal activity & work No evidence of disease	Full	Normal or Reduced	Full
70	Reduced	Unable Normal Job/Work Significant disease	Full	Normal or Reduced	Full
60	Reduced	Unable hobby/house work Significant disease	Occasional assistance necessary	Normal or Reduced	Full or Confusion
50	Mainly Sit/Lie	Unable to do any work Extensive disease	Considerable assistance required	Normal or Reduced	Full or Confusion
40	Mainly in Bed	Unable to do most activity Extensive disease	Mainly assistance	Minimal to sips	Full or Drowsy +/- Confusion
30	Totally Bed Bound	Unable to do any activity Extensive disease	Total Care	Mouth care only	Full or Drowsy +/- Confusion
20	Totally Bed Bound	Unable to do any activity Extensive disease	Total Care		Full or Drowsy +/- Confusion
10	Totally Bed Bound	Unable to do any activity Extensive disease	Total Care		Drowsy or Coma +/- Confusion
0	Death	-	-	-	-

PPS = Palliative Performance Scale.

APPENDIX B. SEARCH TERMS

	PubMed	Embase	Cochrane library
Phase 1.	“palliative performance scale”[tiab] OR ppsv2[tiab]	‘palliative performance scale’/exp OR ‘palliative performance scale’:ti,ab OR ppsv2:ti,ab OR ppsv2	“palliative performance scale”:ti,ab OR ppsv2:ti,ab
Phase 2.	“prognosis”[mesh] OR prognosis[tiab] OR prognos*[tiab] OR predict*[tiab] OR “mortality”[mesh] OR mortality[tiab] OR “survival” [mesh] OR survival[tiab] OR rehospitization[tiab] OR rehospitization[tiab] OR length of service*[tiab]	‘prognosis’/exp OR ‘prognosis’:ti,ab OR ‘prognos*’:ti,ab OR ‘predict*’:ti,ab OR ‘mortality’/exp OR ‘mortality’:ti,ab OR ‘survival’/exp OR ‘survival’:ti,ab OR ‘hospital readmission’/exp OR ‘rehospitalization’:ti,ab OR ‘re-hospitalization’:ti,ab	prognosis (mesh) OR prognos*:ti,ab OR predict*:ti,ab OR mortality (mesh) OR mortality:ti,ab OR survival (mesh) OR survival:ti,ab OR rehospitization:ti,ab OR rehospitization: ti,ab OR length of service*:ti,ab
		Phase 1 AND Phase 2	