

## *Clinical Study*

# **Development of a Score Predicting Survival after Palliative Reirradiation**

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Received 5 May 2014; Accepted 2 September 2014; Published 21 September 2014

Academic Editor: Thomas E. Adrian

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*Purpose.* To develop a prognostic model for predicting survival after palliative reirradiation (PR). *Methods and Materials.* We analyzed all 87 PR courses administered at a dedicated palliative radiotherapy facility between 20.06.2007 (opening) and 31.12.2009. Uni- and multivariate survival analyses were performed, the previously published survival prediction score (SPS) was evaluated, and a PR-specific prognostic score was calculated. *Results.* In multivariate analysis, four parameters significantly influenced survival: performance status, use of steroids, presence of liver metastases, and pleural effusion. Based on these parameters, a 4-tiered score was developed. Median survival was 24.5 months for the favorable group, 9.7 and 2.8 months for the two intermediate groups, and 1.1 months for the unfavorable group (P = 0.019 for comparison between the two favorable groups and  $P \le 0.002$  for all other pairwise comparisons). All patients in the unfavorable group died within 2 months. *Conclusion.* The performance of PR-specific score was promising and might facilitate identification of patients who survive long enough to benefit from PR. It should be validated in independent patient groups, ideally from several institutions and countries.

## 1. Introduction

Palliative reirradiation is currently used in a variety of clinical settings, including but not limited to bone and brain metastases or lung and gynecological cancers [1–4]. The number of scientific publications on this topic has increased in recent years [5]. In a well-defined geographical part of Norway, palliative reirradiation contributed 10% to all palliative radiotherapy series administered during a 12-month period [6]. Randomized trials comparing single versus multiple fractions for painful bone metastases reported retreatment rates of 11-42% after a single fraction and 0–24% after multiple fractions, as summarized by Chow et al. [1]. Comparable to palliative radiotherapy in general, clinicians attempt to tailor treatment regimens to patients' prognosis, thereby minimizing undesirable over- and undertreatment. Decision aids such as prognostic scores and nomograms might facilitate rapid and reproducible assessment of patients' survival expectation by

transforming the complex set of patient- and disease-related prognostic factors into a standardized tool. Ideally, prognostic scores are easy to administer and valid across different institutions and countries [7]. The Survival Prediction Score (SPS), developed and validated by Chow et al. in patient cohorts treated with palliative radiotherapy, is among the tools that might be widely applicable, because it is based on three readily available parameters: primary cancer type, site of metastases, and performance status [8]. Its performance has never been tested specifically in patients undergoing palliative reirradiation. Together with a large number of other baseline factors potentially impacting survival, we analyzed SPS in a single-institution cohort study.

## 2. Methods

We retrospectively reviewed the records of all consecutive patients who received palliative reirradiation at a single hospital with dedicated palliative radiotherapy unit. The patients started their treatment in the time period from June 20, 2007 (date of opening of the dedicated palliative radiotherapy unit), to December 31, 2009. Reirradiation was defined as partial or complete field overlap (examples of partial overlap: initial course included thoracic vertebrae Th4-6 and reirradiation Th6-8; initial course of radical prostate radiotherapy followed by pelvic bone metastasis irradiation). A total of 87 reirradiation courses were studied. Stereotactic radiotherapy was unavailable and thus not included in the present series. All medical records, treatment details, and information on date of death were available in the hospital's electronic patient record (EPR) system. The survival status and date of death or last follow-up of the patients were obtained from the EPR. Patients who were lost to followup were censored on the date of last documented contact (personal appointment, telephone conversation, and blood test). Median follow-up for all surviving/censored patients was 5.4 months. Survival time was measured from the start of reirradiation. Actuarial survival curves were generated by Kaplan-Meier method and compared by log-rank test (analyses performed with IBM SPSS Statistics 20). Multivariate analyses were performed by Cox regression (backward conditional method). We assigned SPS as described by Chow et al. [8], that is, based on three variables (nonbreast cancer, metastases other than bone, and Karnofsky performance status (KPS)  $\leq$  60): poor prognosis group when all three were present, intermediate prognosis group when two were present, and good prognosis group when 0-1 were present. Our own prognostic scores were developed as previously described by Rades et al. [9, 10]. In brief, the score for each predictive factor was determined by dividing the actuarial death rate at prespecified time points (given as the percentage) by 10. For example, patients with good KPS were assigned 0 points and those with poor KPS 1.5 points (rate of death at 1 month (15%) divided by 10). The total score represented the sum of the scores for each predictive factor. Two time points reflecting poor prognosis or short survival were chosen, 1 month and 2 months, because there is no generally agreed definition of sufficient survival expectation, justifying initiation of palliative radiotherapy. Given that recent research and discussions focused on overtreatment, for example, use of radiation therapy in the last 30 days of life, we felt that predicting short survival might be more important [11–14].

## 3. Results

Median age at the time of reirradiation was 67 years (range 38–90 years). Prostate (29%) and non-small cell lung cancer (NSCLC, 11%) were the most common primary tumors. Additional baseline information is shown in Table 1. Bone metastases were the prevailing target for reirradiation. The most common regime consisted of 10 fractions of 3 Gy (43%). Other common regimes included 8 Gy single fraction (uncomplicated bone metastases) and 5 fractions of 4 Gy (various sites and indications). Five courses (6%) remained incomplete, typically because of earlier than expected clinical

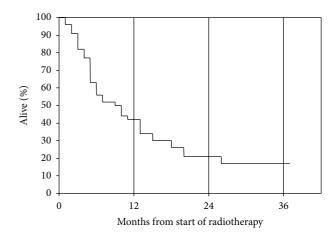


FIGURE 1: Actuarial overall survival after palliative reirradiation (Kaplan-Meier estimate).

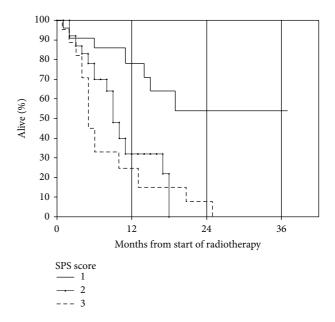


FIGURE 2: Actuarial overall survival after palliative reirradiation stratified by SPS score (Kaplan-Meier estimate): group 1 (n = 23), median not reached; group 2 (n = 26), median 6.7 months; group 3 (n = 38), median 4.1 months; P = 0.26 for group 2 versus 3 and P < 0.05 for other comparisons.

deterioration. Median survival of this small group of patients was 2.8 months. Overall median survival from reirradiation was 8.6 months and 1-year survival rate 42% (Figure 1). Six percent of patients received radiotherapy during the final month of life. Seventeen percent of patients died within 2 months.

We analyzed the potential prognostic impact of all baseline parameters shown in Table 1 and assigned SPS score. However, the performance of this score was unsatisfactory because two of the three patient groups had similar survival (Figure 2). As shown in Table 2, two components of the SPS score (metastases location and performance status) significantly influenced survival, while primary tumor type

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Characteristic	No.	%	Characteristic
Entire cohort	87		Number of meta
Gender			0
Male	65	75	1 (e.g., lungs o
Female	22	25	2 (e.g., lungs
Family status <sup>1</sup>			3
Single	20	23	4
Married	55	63	Progressive dise
Partner	5	6	No
Missing information	7	8	Yes
Karnofsky performance status			Missing infor
90–100	31	36	Systemic cancer
70-80	30	34	No
≤60	26	30	Within 4 wee
Primary tumor site			Within 3 mor Earlier
Prostate	25	29	Missing infor
Breast	9	10	Use of opioid an
Lung (non-small cell)	10	11	No
Colorectal	8	9	Yes
Bladder	5	6	Missing infor
Kidney	6	7	Use of steroids a
Skin (malignant melanoma)	3	3	No
Other	21	24	Yes
Dose/fractionation (intention-to-treat)			Missing infor
10 fractions of 3 Gy	24	28	Serum hemoglo
Single fraction of 8 Gy	19	22	Low <sup>2</sup>
5 fractions of 4 Gy	15	17	Normal
12–15 fractions of 2.5 Gy	4	5	Missing infor
Other	25	29	Serum albumin <sup>1</sup>
Reirradiation target types			Low <sup>2</sup>
Bone metastases	69	79	Normal
Brain metastases	5	6	Missing infor
Lung metastases or primary tumor	6	7	Serum lactate de
Other	7	8	Normal <sup>2</sup>
Known brain metastases			Elevated
No	80	92	Missing infor
One or more	7	8	Serum alkaline j
Known liver metastases			Normal <sup>2</sup>
No	68	78	Elevated
One or more	19	22	Missing infor
Known lung metastases			Serum creatinin
No	65	75	Low <sup>2</sup>
One or more	22	25	Normal Elevated
Known adrenal gland metastases			Missing infor
No	76	87	Serum C-reactiv
One or more	11	13	Normal <sup>2</sup>
Known bone metastases			Elevated but l
No	14	16	Elevated 30–6
One or more	73	84	Elevated >60
Metastatic spinal cord compression			Missing infor
No	80	92	Thrombocyte co
Yes (radiologic or symptomatic)	7	8	Low <sup>2</sup>
Pleural effusion			Normal
No	81	93	High
Yes (radiologic or symptomatic)	6	7	Missing infor

TABLE 1: Continued.			
Characteristic	No.	%	
Number of metastatic sites			
0	10	11	
1 (e.g., lungs only)	37	4	
2 (e.g., lungs and brain)	27	3	
3	11	13	
4	2	2	
Progressive disease outside RT target volume <sup>1</sup>			
No	27	3	
Yes	55	6	
Missing information	5	6	
Systemic cancer treatment <sup>1</sup>			
No	23	2	
Within 4 weeks before RT	21	2	
Within 3 months before RT	14	10	
Earlier	17	2	
Missing information	12	14	
Use of opioid analgetics at start of RT <sup>1</sup>			
No	21	2	
Yes	54	6	
Missing information	12	14	
Use of steroids at start of RT <sup>1</sup>			
No	32	3	
Yes	38	4	
Missing information	17	2	
Serum hemoglobin <sup>1</sup>			
Low <sup>2</sup>	66	7	
Normal	16	18	
Missing information	5	6	
Serum albumin <sup>1</sup>			
Low <sup>2</sup>	17	2	
Normal	42	4	
Missing information	28	3	
Serum lactate dehydrogenase <sup>1</sup>			
Normal <sup>2</sup>	14	10	
Elevated	35	4	
Missing information	38	4	
Serum alkaline phosphatase <sup>1</sup>			
Normal <sup>2</sup>	25	2	
Elevated	29	3	
Missing information	33	3	
Serum creatinine <sup>1</sup>			
Low <sup>2</sup>	13	1	
Normal	48	5	
Elevated	15	12	
Missing information	11	1.	
Serum C-reactive protein <sup>1</sup>			
Normal <sup>2</sup>	20	2	
Elevated but less than 30 mg/L	27	3	
Elevated 30-60 mg/L	14	10	
Elevated >60 mg/L	17	2	
Missing information	9	10	
Thrombocyte count <sup>1</sup>			
Low <sup>2</sup>	11	13	
Normal	45	5	
High	19	2	
Missing information	12	14	

TABLE 1: Continued.

Characteristic	No.	%
Charlson comorbidity index <sup>1</sup>		
0	7	8
1-2	44	51
3 or more	28	32
Missing information	8	9
Smoking status <sup>1</sup>		
Current smoker	34	39
No	34	39
Missing information	19	22

RT: radiotherapy.

<sup>1</sup>Missing information in some cases.

<sup>2</sup>Hematology and blood chemistry results refer to institutional limits of normal; only test results obtained within one week before RT were considered.

did not. In multivariate analysis, a total of four parameters significantly influenced survival: KPS, use of steroids, presence of liver metastases, and pleural effusion. Based on these parameters, a new 4-tiered prognostic score was developed. As described in Section 2, we compared two different variants, which are shown in Table 3. When applying a short-survival-definition of 1 month (variant 1), the resulting survival curves separated clearly (Figure 3). Median survival was 24.5 months for the favorable group, 9.7 and 2.8 months for the intermediate groups, and 1.1 months for the unfavorable group (P = 0.024 for comparison between)the two favorable groups and  $P \leq 0.003$  for all other pair-wise comparisons). Thirty-three percent of patients in the unfavorable group died within 1 month and all within 2 months. When applying a short-survival-definition of 2 months (variant 2), the resulting survival curves separated equally clear (Figure 4). Median survival was exactly the same as in variant 1 (P = 0.019 for comparison between the two favorable groups and  $P \leq 0.002$  for all other pair-wise comparisons). Since the unfavorable group included exactly the same patients, 33% died within 1 month and all within 2 months. Because of its superior significance level, variant 2 might be the preferred assignment method.

#### 4. Discussion

Palliative reirradiation is an important treatment option, providing symptom improvement in many patients with bone metastases [1] and other conditions [15]. While most previous studies were small and often retrospective, the randomized bone metastases study by Chow et al. comparing different fractionation regimens included 850 patients [1]. Median survival in the two arms was 9.3 and 9.7 months, respectively. This result is comparable to the 8.6 months reported in our own, bone metastases-dominated study. However, survival of individual patients might be as short as few days or as long as several years (Figure 1). Therefore, prognostic scores might be valuable decision aids when prescribing palliative reirradiation. Chow et al. have previously published several reports on a score for patients receiving palliative radiotherapy in general, the SPS. Development of this prediction model started in

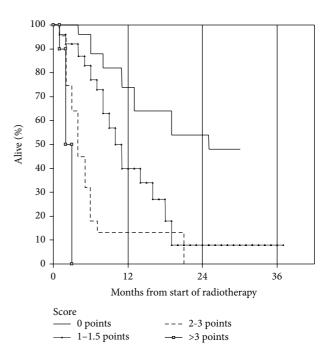


FIGURE 3: Actuarial overall survival after palliative reirradiation stratified by new score, variant 1 (Kaplan-Meier estimate): median 24.5 (0 points) versus 9.7 (1–1.5 points) versus 2.8 (2-3 points) versus 1.1 months (>3 points), P = 0.024 for comparison between group 1 and 2,  $P \leq 0.003$  for all other pair-wise comparisons. Number of patients in each group: 20, 24, 20, and 6. Missing information to assign score in 17 patients.

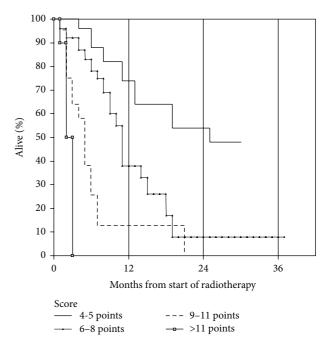


FIGURE 4: Actuarial overall survival after palliative reirradiation stratified by new score, variant 2 (Kaplan-Meier estimate): median 24.5 (4-5 points) versus 9.7 (6–8 points) versus 2.8 (9–11 points) versus 1.1 months (>11 points), P = 0.019 for comparison between group 1 and 2,  $P \leq 0.002$  for all other pair-wise comparisons. Number of patients in each group: 20, 26, 18, and 6. Missing information to assign score in 17 patients.

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TABLE 2: Prognostic factors for survival. All baseline variables shown in Table 1 were analyzed (univariate, log-rank test). Those with *P* value <0.1 were carried forward to multivariate Cox regression analysis and are shown here.

Characteristic	Median survival (months)		alue	
		Univariate <sup>1</sup>	Multivariate	
Karnofsky PS				
90–100	18.3		0.0001	
70-80	9.4	0.0001	0.0001	
≤60	2.1			
Known brain metastases				
No	9.7	0.008	n.s.	
Yes	3.6			
Known liver metastases				
No	9.7	0.037	0.039	
Yes	2.8			
Pleural effusion				
No	9.4	0.007	0.039	
Yes	1.3			
Number of metastatic sites				
Max. 2	9.7	0.054	n.s.	
3 or more	2.8			
Progressive disease outside RT target volume				
No	12.6	0.033	n.s.	
Yes	5.5			
Use of opioid analgetics				
No	24.5	0.02	n.s.	
Yes	5.2			
Use of steroids				
No	12.2	0.002	0.015	
Yes	3.6			
Serum albumin				
Low	9.7	0.001	n.s.	
Normal	2.8			
Serum alkaline phosphatase				
Normal	15.1	0.027	n.s.	
Elevated	4.1	/		
Serum creatinine				
Low	1.6			
Normal	9.7	0.0001	n.s.	
Elevated	15.1			
Serum C-reactive protein				
Normal	18.3			
Elevated but less than 30 mg/L	12.6	0.0001	n.s.	
Elevated 30–60 mg/L	5.3		11.5.	
Elevated >60 mg/L	2.6			
Thrombocyte count				
Low	12.7			
Normal	9.7	0.038	n.s.	
High	4.0			
Number of abnormal blood tests <sup>2</sup>				
Max. 1	12.7			
2	5.8	0.008	n.s.	
3 or more	3.0			

TABLE 2: Continued.

Characteristic	Median survival (months)	Px	<i>P</i> value		
	median survivar (months)	Univariate <sup>1</sup>	Multivariate		
Smoking status					
Current smoker	4.3	0.063	<b>n</b> 6		
No	9.7		n.s.		
Time from first cancer diagnosis					
Shorter than median (47 months)	5.3	0.089	20		
Longer than median	9.7	0.089	n.s.		

RT: radiotherapy; PS: performance status.

<sup>1</sup>If more than 2 groups, *P* value from log-rank test pooled over all strata.

<sup>2</sup>All tests shown in Table 1 were considered. Significance levels were not corrected for multiple tests.

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TABLE 3: Prognostic scores based on four parameters predicting survival in multivariate analysis. Endpoints: death within 1 month (variant 1) and death within 2 months (variant 2).

Parameter	Died within 1 month	Points <sup>1</sup>	Died within 2 months	Points <sup>1</sup>
Karnofsky PS				
70–100	2%	0	7%	1
≤60	15%	1.5	39%	4
Known liver metastases				
No	4%	0	8%	1
Yes	11%	1	49%	5
Pleural effusion				
No	4%	0	14%	1
Yes	33%	3	50%	5
Use of steroids				
No	3%	0	10%	1
Yes	11%	1	28%	3
Minimum sum score		0		4
Maximum sum score		6.5		17

PS: performance status.

<sup>1</sup>Death rate divided by 10.

395 patients referred to their palliative radiotherapy program [16]. Later, they refined their original six-parameter-model by reducing the number of variables to three (primary cancer type, site of metastases, and performance status), arriving at the SPS [8, 17]. We hypothesized that this score might also predict survival of patients receiving reirradiation but discovered that further studies, which also include other models, are needed. The performance of the SPS score (Figure 2) can be explained by the fact that not all adverse SPS features (nonbreast cancer, metastases other than bone, and poor performance status) influenced prognosis of reirradiated patients. In the present study, metastases location and performance status significantly influenced survival, while primary tumor type did not.

Disadvantages of our study include its retrospective design and limited number of patients, especially regarding subgroups. Not all patients had complete information on all baseline parameters recorded in the EPR system. The majority of reirradiation courses consisted of hypofractionated regimens, mostly 1–15 fractions, with dose/fractionation parameters reflecting a patient's expected prognosis (clinical estimate). We did not use any particular prognostic models or scores when assigning treatment regime during the time period covered in our study. Nevertheless, more than 90% of patients who were offered reirradiation also completed their treatment. Only 6% were treated during the final month of life, suggesting that our clinical decision making was largely successful, even if further improvement should be attempted.

Our score based on KPS, use of steroids, presence of liver metastases, and pleural effusion performed promisingly. To the best of our knowledge, no other scores related specifically to palliative reirradiation exist. One of the clinical aims of applying prognostic scores might be avoidance of overtreatment in patients with very short survival [18]. Recently, Tanvetyanon et al. have reported on use of radiotherapy in the last 30 days of life in the United States [19]. They used a SEER-Medicare linked database to obtain a large study cohort of 202,299 patients who died as a result of lung, breast, prostate, colorectal, and pancreas cancers (top five cancer causes of death) between January 1, 2000, and December 31, 2007. The rate of radiotherapy in the last 30 days of life, by many regarded as inappropriate overtreatment, though this point of view is controversial, was 7.6%. No data on reirradiation were reported in this study, and no attempt was made to develop predictive models. Before our new score can be widely implemented, external validation is necessary. In the future, it might become possible to study narrowly defined patient groups, if sufficiently large databases can be created. For example, Tanvetyanon et al. have published prognostic factors for survival after salvage reirradiation in patients with head and neck cancer [19]. Rades et al. have developed scores specific to metastatic spinal cord compression [20, 21], and Sperduto et al. to brain metastases [22], both related to first line treatment rather than reirradiation.

#### 5. Conclusions

Prognostic factors for survival might change during the course of disease, for example, from first line to subsequent treatments. The performance of the newly developed score was promising and might facilitate identification of patients who survive long enough to benefit from palliative reirradiation. It should be validated in independent patient groups, ideally from several institutions and countries.

### **Conflict of Interests**

The authors declare that there is no conflict of interests regarding the publication of this paper.

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