

A survival predicting model for patients with papillary renal cell carcinoma

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We commend the work of Yan *et al.* (1) for their retrospective study developing nomograms to predict the survival outcomes of patients with papillary renal cell carcinoma (pRCC), which may aid in clinical decision-making. As such, there are a few points that we would like to bring up.

The study included patients whose pRCC was the first tumor but did not exclude multiple primary cancers. However, another SEER-based study showed renal cell carcinoma (RCC) patients were at a significantly higher risk of developing a second malignancy, which may affect patients' survival (2).

RCC surgery in the SEER database includes cryosurgery, partial nephrectomy, and radical nephrectomy, etc. Previous studies have also proved that the prognosis of RCC patients with early T stage receiving different types of surgical treatment is different (3,4). It would be more meaningful to stratify patients according to the type of surgery that they received; in this way, readers can easily see which surgical treatment improves pRCC patient's survival better.

Because there is lack of granularity in SEER about type of chemotherapy and only a small number of patients had received chemotherapy (3.25%), the authors' finding that chemotherapy was a negative prognostic factor should be treated with reservation. Previous case reports showed pRCC patients with long-term cancer control by chemotherapy (5,6).

The number of patients younger than 40 years old and older than or equal to 80 years old in their study is too

small. Continuous variables, such as age, can be divided based on the optimal cut-off value generated by X-tile software, making grouping more reasonable.

SEER database provides information of four metastatic sites (lung, liver, bone and brain), which could be considered for inclusion in their study. This is because considering whether patients had the above four sites of metastases rather than whether they were in M1 stage will improve predictive accuracy of their model to some extent.

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Guo and Liu. Nomograms for predicting survival in pRCC

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