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## Phase II Study of Pembrolizumab Efficacy and Safety in Women with Recurrent Small Cell Neuroendocrine Carcinoma of the Lower Genital Tract

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### Author Contribution Statement

Drs. Aung Naing and Michael Frumovitz were involved in all aspects of the study including study design, patient recruitment and study administration, collection and analysis of data, and writing and approval of the final manuscript. Drs. Shannon Westin, Timothy Yap, Ahnert J. Rodon, Daniel Karp, and Amir Jazaeri were involved in patient recruitment and study administration, collection of data, and writing and approval of the final manuscript. Gloria Salvo, Abdulrazzak Zarifa, Mingxuan Xu, and Abdulrahman Abonofal were involved in study administration, collection of data, and writing and approval of the final manuscript.

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Drs. Abonofal, Salvo, Xu and Zarifa have nothing to disclose.

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## Abstract

**Objective**—To investigate the efficacy and safety of pembrolizumab in women with recurrent small cell neuroendocrine tumors of the lower genital tract.

**Methods**—We conducted an open-label, investigator-initiated phase II basket trial of pembrolizumab 200 mg intravenously every 3 weeks in patients with rare tumors ([ClinicalTrials.gov: NCT02721732](https://clinicaltrials.gov/ct2/show/study/NCT02721732)). The trial had prespecified cohorts, including small cell malignancies of extrapulmonary origin. Eligibility criteria included disease progression during standard treatment in the 6 months before study enrollment. Patients were enrolled from February 2017 to February 2019. The primary endpoint was the proportion of patients alive without progression at 27 weeks. Response to pembrolizumab was evaluated every 9 weeks (3 cycles) with radiographic imaging.

**Results**—Seven women with gynecologic extrapulmonary small cell carcinoma were enrolled, 6 with cervical and 1 with vulvar carcinoma. No patient was progression free at 27 weeks. At first radiologic assessment, 1 patient had stable disease, while 6 had progression. The single patient with stable disease at 6 weeks had disease progression at 14 weeks. The median progression-free interval was 2.1 months (range 0.8–3.3 months). Severe treatment-related adverse events (grade 3) were seen in 2 of 7 patients (29%); 1 patient had grade 3 asymptomatic elevation of serum alkaline phosphatase, and 1 had grade 3 asymptomatic elevation of serum alanine aminotransferase.

**Conclusions**—Pembrolizumab alone showed minimal activity in women with recurrent small cell neuroendocrine tumors of the lower genital tract. Treatment was well tolerated in the majority of study participants, and the rate of severe adverse events was low.

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## Introduction

High-grade neuroendocrine carcinomas of the cervix (small cell, large cell, and undifferentiated) account for < 2% of all newly diagnosed cervical cancers. These tumors are highly aggressive and have high rates of recurrence. Even though > 70% of women with high-grade neuroendocrine carcinoma of the cervix are diagnosed with early-stage disease, the 5-year survival rate for all patients with this disease is < 30% [1]. Consensus guidelines detailing recommended therapies for newly diagnosed patients have been published, but none of these guidelines offer options for recurrent disease [2, 3]. The National Comprehensive Cancer Network guidelines for treating cervical cancer specifically exclude high-grade neuroendocrine carcinoma [4]. Combination chemotherapy with topotecan, paclitaxel, and bevacizumab has emerged as a common regimen for recurrent small cell neuroendocrine carcinoma of the cervix but even with these drugs, median overall survival after first recurrence is < 10 months [5]. There are very few active regimens for women with recurrent disease, and new treatment options are desperately needed.

Many therapeutic approaches for treating women with high-grade neuroendocrine carcinomas of the cervix have been extrapolated from studies in small cell lung cancer as the diseases appear histologically alike with similar clinical behavior. Studies have demonstrated the activity of single-agent checkpoint inhibitors in treating recurrent small cell lung cancer. The KEYNOTE-028 study reported an objective response rate of 33% (1 complete response,

7 partial responses) for the anti-PD-1 antibody pembrolizumab in 24 patients with recurrent small cell lung cancer [6]. The CheckMate-032 study also showed good activity for PD-1 inhibitors in recurrent small cell lung cancer: 10 (10%) of 98 patients had a partial response to single-agent nivolumab, and an additional 22 (22%) had stable disease [7].

Immune checkpoint inhibitors have also shown promise in the most common types of cervical cancer. Over 98% of cervical cancers are of squamous, adenocarcinoma, or adenosquamous histologies. In 98 patients with such tumors, pembrolizumab demonstrated an overall response rate of 12% (3 complete responses and 9 partial responses) [8]. Nivolumab as a single agent has been explored in 2 different studies in cervical cancer. In a study of 19 patients, the objective response rate was 26% (3 complete responses and 2 partial responses), and another 8 patients (42%) had stable disease [9]. Results of a second study, however, were less impressive: only 1 (4%) of 25 evaluable patients achieved a partial response (duration of response, 3.8 months), and another 9 (36%) had stable disease [10]. The median duration of response for those with stable disease was only 5.7 months.

Although there is a single case report of nivolumab as an active agent in a woman with recurrent high-grade neuroendocrine carcinoma of the cervix [11], we identified no prospective studies evaluating the activity of PD-1/PD-L1 inhibitors in high-grade neuroendocrine carcinomas of the cervix in a search of PubMed. As part of a multi-arm basket trial for patients with rare tumors, we evaluated the safety and clinical efficacy of pembrolizumab in a cohort of women with small cell neuroendocrine carcinomas of the lower genital tract.

## Methods

This phase II, open-label study of single-agent pembrolizumab ([ClinicalTrials.gov: NCT02721732](https://clinicaltrials.gov/ct2/show/study/NCT02721732)) was approved by both the US Food and Drug Administration and the Institutional Review Board at The University of Texas MD Anderson Cancer Center. All patients were enrolled at MD Anderson Cancer Center. Patients with recurrent or advanced rare tumors were enrolled into one of 10 cohorts: 1) squamous cell carcinoma of the skin, 2) small cell malignancies of extrapulmonary origin, 3) adrenocortical carcinoma, 4) medullary renal cell carcinoma, 5) carcinoma of unknown primary, 6) penile carcinoma, 7) vascular sarcoma, 8) testicular cancer, later relabeled as germ cell tumor, 9) paraganglioma-pheochromocytoma, and 10) other rare tumor histologies. Results of the entire study have been reported elsewhere [12]. Here, we report on the patients with gynecologic cancers enrolled in the prespecified extrapulmonary small cell carcinoma cohort.

To be eligible for cohort 2 of the umbrella trial, patients had to be ≥ 18 years old with histologically confirmed recurrent or metastatic extrapulmonary small cell malignancy. Patients had to have experienced disease progression during standard therapies within the previous 6 months. Patients treated with prior immunotherapy were excluded. Patients were eligible regardless of PD-L1 expression. In this report, we included only patients with extrapulmonary small cell carcinoma of gynecologic origin.

Pembrolizumab was administered at a fixed dose of 200 mg intravenously every 3 weeks. Patients remained on treatment until documented disease progression, completion of 24 months of treatment, withdrawal of consent or investigator's decision to stop treatment, or unacceptable adverse event. Patients could be removed from the study for either clinical or radiologic disease progression.

The primary endpoint of the study was the proportion of patients who were alive and progression free at 27 weeks (9 cycles). Secondary endpoints included objective response rate (partial or complete response), clinical benefit rate 4 months (complete response, partial response, or stable disease), and safety and tolerability.

Tumor response was assessed every 9 weeks (3 cycles) using Response Evaluation Criteria in Solid Tumors (RECIST) version 1.1 and/or immune-related RECIST (irRECIST). Patients who had radiographic progression but clinically stable disease were allowed to continue pembrolizumab until confirmation of disease progression by a follow-up scan 4 weeks after initial documentation of progression. Adverse events were assessed using the National Cancer Institute Common Terminology Criteria for Adverse Events version 4.03.

When fresh or archived tissue was available, PD-L1 biomarker analysis was performed by Qualtek assay using Merck 22C3 antibody. An H-score (range, 0–300) was assigned on the basis of the percentage and intensity of membrane staining. Through recursive partitioning analysis, a score of 42.5 was identified as the optimal cut-off point for PD-L1 H-score. Tumorinfiltrating lymphocytes (TILs) within tumor nests were scored on a scale of 0 to 3, with 0 denoting absence and 3 representing intense intratumoral lymphocytic infiltration. Boardcertified pathologists performed all biomarker evaluations.

The study utilized a Simon's optimal 2-stage design for each of the 10 cohorts. The first stage accrued 12 patients, and 3 patients had to meet the primary endpoint (nonprogression at 27 weeks) in order for the second stage, with accrual of an additional 13 patients, to be opened. In this report, we present data for the patients with gynecologic malignancies in cohort 2 (extrapulmonary small cell cancers). Descriptive statistics are used to summarize patient characteristics. Patients were included in the outcome analysis if they received 1 adequate onstudy tumor assessment and were included in the safety analysis if they received 1 dose of pembrolizumab. The best overall response was defined as the best response observed from the start of the treatment until disease progression or discontinuation of treatment for any reason. Waterfall plots were constructed to demonstrate best overall response based on irRECIST.

## Results

Twelve patients with extrapulmonary small cell carcinoma were enrolled in this cohort between February 2017 and February 2019. Six patients had cervical cancer, 4 had prostate cancer, 1 had vulvar cancer, and 1 had small cell carcinoma of unknown primary. No patients in the cohort met the primary endpoint of nonprogression at 27 weeks, and therefore no further patients with extrapulmonary small cell carcinoma were enrolled.

Seven patients had a small cell carcinoma of gynecologic origin. Demographics for this group are summarized in Table 1. The median age was 41 years (range, 31–76), and the median number of prior therapies was 2 (range, 1–3).

Two patients (28%), 1 with vulvar cancer and 1 with cervical cancer, had clinical progression prior to first imaging assessment. At first imaging assessment, 4 patients (57%) had immune-related progression by irRECIST criteria and 1 patient (14%) had stable disease. The patient with stable disease had imaging early at 6 weeks due to a suspicion of clinical progression but was noted to have 10% decrease in tumor size. However, at her next assessment, 8 weeks later, she was noted to have 40% increase in tumor size. The best overall responses are summarized in Figure 1. The median progression-free survival was 2.1 months (range, 0.8–3.3 months, 95% confidence interval, 0.8–3.2 months). Durations of response are shown in Figure 2. At the time of data analysis, none of the 7 patients were alive.

Treatment-related adverse events are summarized in Table 2. Fatigue and elevation of liver transaminases were the most commonly reported treatment-related adverse events. The majority of treatment-related adverse events were grade 1 or 2. Two patients had grade 3 adverse events: 1 had asymptomatic elevation of alkaline phosphatase, and the other had asymptomatic elevation of alanine aminotransferase. There were no grade 4 adverse events or treatment related deaths in this cohort.

Six patients (86%) had tissue available for analysis of PD-L1 membrane staining and assessment of the presence of TILs within tumor nests. On the basis of percentage and intensity of membrane staining, 2 patients had an H-score of 0, 3 patients had an H-score of 1 (indicating PD-L1 staining of 1+ in 1% of the tumor cells), and 1 patient had an H-score of 3 (indicating PD-L1 staining of 1+ in 3% of tumor cells). No patient had 2+ staining in any cells. All 6 patients had TILs within tumor nests; 4 patients had a score of 1, 1 patient had a score of 2, and 1 patient had a score of 3. A TIL score of 2 (representing high TILs) was noted in 2 (33%) of 6 patients.

## Discussion

Our group recently reported a phase II study of pembrolizumab in patients with rare tumors. Anti-tumor activity was observed in patients with adrenal cortical carcinoma, paraganglioma, squamous cell carcinoma of skin, and cancers of unknown origin [12]. However, in this phase II clinical trial, no patients with gynecologic small cell neuroendocrine carcinoma showed response to single-agent pembrolizumab. One patient had stable disease at her first assessment, at 6 weeks, but was noted to have progression < 2 months later. The first imaging was performed early at 6 weeks as opposed to 9 weeks due to a clinical suspicion for progression. The other 6 patients had disease progression after only 2 to 3 cycles of treatment. As recurrent small cell carcinoma of the cervix is incurable and is associated with a median survival after first relapse of < 10 months, more clinical trials for women with this malignancy are desperately needed [5].

The relationship between biomarkers and response to PD-1/PD-L1 inhibitors has varied widely across disease sites, tumor histologies, and specific checkpoint inhibitors. In the study reported here, a PD-L1 score was assigned that was based on the percentage and intensity of membrane staining. No patient had more than 1+ staining in any tumor cells, and no patient had more than 3% of cells testing positive. There is no means to translate these findings into the more commonly used combined positive score (CPS) for PD-L1 protein expression, although presumably these modest findings from the Qualtek assay would be similar to a combined positive score of 0. In a separate study of 23 high-grade neuroendocrine carcinomas stained for PD-L1 expression, only 1 (4%) tested positive for the receptor (combined positive score = 1) [13].

In KEYNOTE-028, in which investigators found an overall response rate of 33% in patients with small cell lung cancer treated with single-agent pembrolizumab, PD-L1 expression was a requirement for eligibility, so tumors from all 24 patients enrolled were positive for PD-L1 expression. In contrast, in Checkmate-032, in which investigators found a response rate of 10% in patients with recurrent small cell lung cancer, PD-L1 expression was not a requirement for inclusion, and in fact 86% of patients tested had PD-L1-negative tumors [7]. In that study, there was no relationship between response and PD-L1 expression.

For the most part, treatment strategies for small cell neuroendocrine carcinomas of the gynecologic tract have been borrowed from approaches that have proven successful in patients with small cell lung cancer [2]. However, data from our previous study of whole exome sequencing of small cell neuroendocrine tumors of the cervix show that these tumors may be more similar to squamous carcinomas and adenocarcinomas of the cervix than to small cell lung cancer [14]. In KEYNOTE-158, a study of single-agent pembrolizumab in women with recurrent squamous carcinoma, adenocarcinoma, or adenosquamous carcinoma, the only responses observed occurred in patients with PD-L1-expressing tumors [8]. In contrast, in Checkmate-358, a study of single-agent nivolumab in a similar patient population, responses did not seem to be related to the presence or absence of PD-L1 expression [9].

In addition to PD-L1 expression, tumor mutation burden has emerged as a potential predictor of response to immunotherapy. Clinical trials of checkpoint inhibitors in patients with melanoma and non-small cell lung cancer have shown improved survival for those with high tumor mutation burden [15–17]. Tumors that have a mutational load above 10 somatic mutations per megabase are considered most likely to be able to produce neoantigens that can be detected by T-cells [18]. Although a small number of small cell lung cancer and cervical cancer specimens have > 10 mutations per megabase, the overwhelming majority of these tumors will not be considered to harbor high mutational burden [18]. We previously evaluated tumor mutational burden in 15 small cell cervical cancer tumor specimens through whole exome sequencing and found a median of only 1.7 somatic mutations per megabase [14]. Another marker associated with high tumor mutation burden is high microsatellite instability. In our previous evaluation of 25 high-grade neuroendocrine cervical cancer tumor specimens, none were found to have high microsatellite instability [13].

Even though traditional biomarkers of immunotherapy response such as PD-L1 expression, microsatellite instability, and tumor mutational burden might suggest a lack of response to checkpoint inhibitors for high grade neuroendocrine carcinomas of the cervix, there may still be a role for immunotherapy in these tumors. Immunohistochemistry staining for PARP expression in small cell lung cancer specimens has predicted *in vitro* responses to PARP inhibitors [19]. Similarly, immunohistochemistry studies have demonstrated high PARP expression of >90% in small cell cervical cancer specimens [13]. DNA damage repair (DDR) inhibitors such as PARP inhibitors have been shown to upregulate PD-L1 expression and enhance responses to immunotherapy in cell lines [20]. We believe the combination of DDR and checkpoint inhibitors may be an active therapeutic approach for patients with high grade neuroendocrine carcinomas and will soon open a phase II basket trial of the PARP inhibitor niraparib and the anti-PD1 agent TSR-042 for neuroendocrine small cell carcinomas for any primary site.

One of the major limitations of this study is the small sample size. Performing clinical trials in rare tumors is challenging for many reasons, among them difficulty with recruiting patients and securing funding. Research in rare tumors requires recruiting from small populations of potential patients spread across large geographical areas, which makes trial accrual and completion difficult [21]. However, social media has emerged as a very effective means to “advertise” trials to focused groups of women with rare gynecologic cancers, potentially improving accrual [22]. In addition, one might consider opening trials at a few select centers chosen for geographic location to optimize access to patients from around the world. Small, innovative single-arm clinical trials focused on “pertinence, validity, and precision” could positively change therapeutic options for patients with rare diseases with only a relatively few patients enrolled in any single trial [21]. In this study, we were able to accrue 7 patients with small cell neuroendocrine carcinoma of the gynecologic tract in 2 years, an accrual rate much higher than the rate of < 2 patients per year that our institution uses to define a low-accruing trial. [23].

A second factor that can make performing clinical trials in rare tumors challenging is that drug companies may be hesitant to fund expensive studies for a small return on their investment even if drugs are Food and Drug Administration–approved for an indication in a specific rare tumor. Currently multiple drug companies are making drugs in the same class such as PD-1/PDL1 and PARP inhibitors. They therefore may be incentivized to expand indications for their drug into rare tumors as a means to differentiate themselves from a competitor’s compound. We may be in an era where pharmaceutical companies are more willing to explore supporting trials in rare malignancies.

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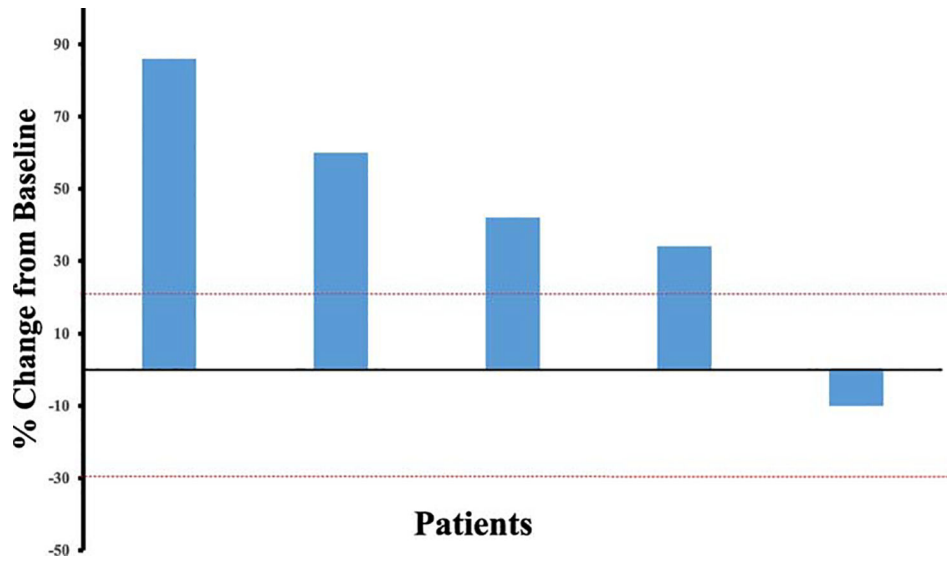
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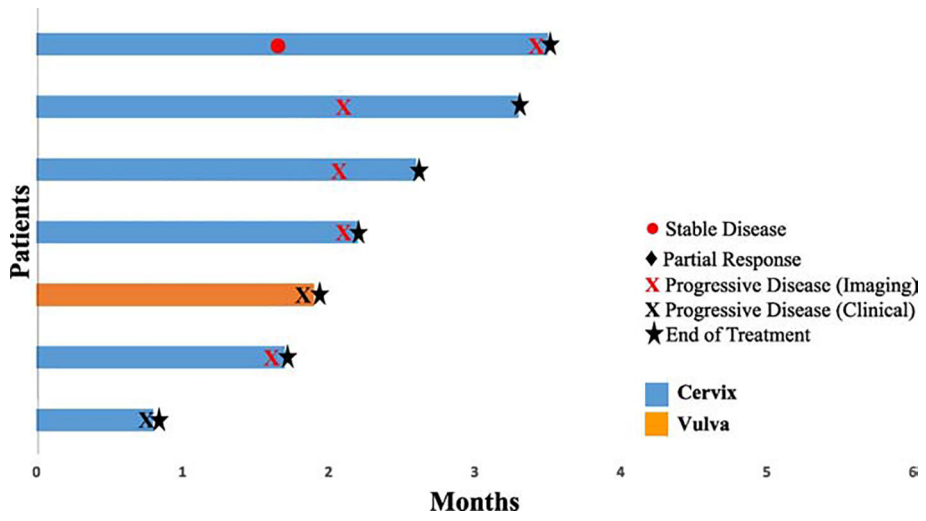
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**Highlights**

- Single-agent pembrolizumab had minimal antitumor activity against small cell neuroendocrine tumors of the gynecologic tract
- Single-agent pembrolizumab had acceptable toxicity in patients with small cell neuroendocrine tumors of the gynecologic tract
- Primary small cell neuroendocrine tumors of the gynecologic tract was associated with low PD-L1 expression



**Figure 1:** Waterfall plot illustrating response to pembrolizumab in 5 evaluable patients with cervical cancer. Two patients (1 with vulvar cancer and 1 with cervical cancer) had clinical progression before first imaging assessment and are not included in this figure.



**Figure 2:**  
Time to and duration of response to pembrolizumab in 7 evaluable patients

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**Table 1:**

## Patient demographics

Characteristic	n (%)
Primary tumor site	
Cervix	6 (86)
Vulva	1 (14)
Median age, years (range)	
	41 (31–76)
Race	
Caucasian	6 (86)
Hispanic	1 (14)
ECOG performance status	
0	2 (29)
1	5 (71)
Number of prior therapies	
1	2 (29)
2	3 (43)
3	2 (29)

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**Table 2:**

Treatment-related adverse events during pembrolizumab therapy

Adverse event	No. (%) of patients	
	Any grade	Grade 3
Fatigue	2 (29)	0
Elevated ALT/AST	2 (29)	1 (14)
Elevated alkaline phosphatase	1 (14)	1 (14)
Arthralgia	1 (14)	0
Maculopapular rash	1 (14)	0

ALT - alanine aminotransferase, AST - aspartate aminotransferase

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