Objectives Women with cervical cancer who undergo radical hysterectomy are often treated postoperatively with chemoradiation. The patient selection that minimizes adjuvant treatment is valuable. We compared two methods for predicting postoperative adjuvant treatment of patients with stage IB2 cervical cancer.

Methods This multicenter retrospective study included 272 women with IB2 tumors. A receiver operating characteristic curve (ROC) analysis was used to determine the optimal tumor cutoff size to predict adjuvant treatment. A second analysis compared the rate of adjuvant treatment between women with and without lymph vascular space involvement (LVSI).

Results According to the ROC, the optimal cutoff value of tumor size for predicting adjuvant treatment was 2.95 cm (sensitivity 0.70, specificity 0.67). Tumors were \( \geq 3.0 \text{ cm} \) in 166 (61.0%) women. The rate of adjuvant treatment was higher in women with larger tumor diameter (73.8% vs. 47.9%, \( p<0.001 \)). Among women with LVSI, rates were higher of positive lymph nodes (41.0% vs. 14.5%, \( p<0.0001 \)) and postoperative adjuvant treatment (83.3% vs. 53.7%, \( p<0.0001 \)). Among women with tumor size \( \geq 3.0 \text{ cm} \) and LVSI, the rate of adjuvant treatment was 90.0%. In the multivariate analysis, both tumor size \( \geq 3.0 \text{ cm} \) and LVSI were independently associated with adjuvant treatment (OR 3.9, 95% CI 2.1–7.1; \( p<0.0001 \) and OR 4.9, 95% CI 2.4–10.0; \( p<0.0001 \), respectively).

Conclusions These data should be weighed in multidisciplinary consultation with radiation oncologists when deciding treatment strategy.

**EPV035/#138** RADICAL TRACHELECTOMY. EXPERIENCE IN KAZIOR

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**Objectives** To investigate pregnancy outcomes in women after radical trachelectomy (RT) in Kazior for early-stage cervical cancer

**Methods** Systematic analysis of the data of the cancer register of the Republic of Kazakhstan

**Results** Since 2013, radical trachelectomy has been performed at Kazior. From 2013 to 2021, 8 operations were performed, 7 of them by abdominal access, 3 by laparoscopic approach. 6 (75%) of the patients had stage 1B1 from 2 to 4 cm; 2 (25%) had a 1A1 stage. The average age of patients was 28 years (from 26 to 37 years). 5 (62.5%) patients were nulliparous, 2 patients had 2 children, 1 patient had 1 child.

LVSI was negative in preoperative histological examination, and resection margins were also negative. The histological form of the tumor in all cases was squamous cell carcinoma. On average, 11 lymph nodes were removed. In 1 patient (12.5%) after histological examination LVSI was positive, in 7 it was negative. None of the patients had metastases to the pelvic lymph nodes. During express history, the resection margins were negative in all patients. Patients in the postoperative period were not prescribed chemoradiation therapy, of the 8 patients who retained fertility, there were 5 pregnancies, 2 miscarriages at 9–10 weeks, and 3 deliveries at 36–37 weeks of gestation.

**Conclusions** Thus, in 2013–2021, 8 radical trachelectomy operations were successfully performed. The data presented in this publication demonstrate that patients with stage IB1 tumors ranging in size from 2 to 4 cm and with favorable histology are acceptable candidates for attempted radical trachelectomy.

**EPV036/#142** EUROPEAN NETWORK FOR GYNAECOLOGICAL ONCOLOGICAL TRIAL (ENGOT)-CX11/ GYNECOLOGIC ONCOLOGY GROUP (GOG) 3047/ KEYNOTE-A18: PHASE 3 TRIAL OF PEMBROLIZUMAB PLUS CHEMORADIOTHERAPY IN HIGH-RISK LOCALLY ADVANCED CERVICAL CANCER

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**Objectives** High-risk locally advanced cervical cancer has a poor prognosis. External beam radiotherapy (EBRT) with concurrent chemotherapy followed by brachytherapy is the standard of care. The immunostimulatory activity of pembrolizumab may be enhanced by concurrent chemoradiotherapy (CCRT). Pembrolizumab monotherapy is approved for patients with PD-L1–positive recurrent or metastatic cervical cancer that progressed during or after chemotherapy. The phase 3 ENGOT-cx11/GOG 3047/KEYNOTE-A18 (NCT04221945) study is evaluating pembrolizumab with CCRT in patients with locally advanced cervical cancer.

**Methods** 980 patients with high-risk (FIGO 2014 stage IB2-IIb with node-positive disease or stage III-IVA), locally
advanced, previously untreated cervical cancer will be randomized 1:1 to receive either 5 cycles of pembrolizumab 200 mg Q3W plus CCRT followed by 15 cycles of pembrolizumab 400 mg Q6W or 5 cycles of placebo Q3W plus CCRT followed by 15 cycles of placebo Q6W. CCRT includes 5 cycles (optional 6th dose) of cisplatin 40 mg/m² Q1W plus EBRT followed by brachytherapy. Randomization is stratified by planned EBRT type (intensity-modulated radiotherapy [IMRT] or volumetric-modulated arc therapy [VMAT] vs non-IMRT or non-VMAT), screening stage cancer (IB2-IIb vs III-IVA), and planned total radiotherapy dose. Treatment will continue for 20 cycles or disease progression, unacceptable toxicity, or withdrawal. Primary endpoints are PFS per RECIST v1.1 by investigator and OS. Secondary endpoints include PFS at 2 years, OS at 3 years, CR at 12 weeks, ORR, and safety. Enrollment began May 2020 and is planned for 193 sites in 30 countries.

**Results** Not applicable

**Conclusions** Not applicable

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**EPV037/#157**

**INCIDENCE OF CERVICAL CANCER AND THE HPV VACCINE IN THE UNITED STATES: ARE WE SEEING RESULTS OF VACCINATION EFFORTS?**

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**Objectives** To determine the incidence and trends of cervical cancer in the United States in relation to the HPV vaccine.

**Methods** Data were obtained from the U.S. Cancer Statistics program from 2001–2017. SEER*Stat 8.3.8 and Joinpoint regression program 4.8.0.1 were used to calculate incidence trends.

**Results** Over the last 17 years, cervical cancer incidence is decreasing at an average annual percent change (AAPC) of -1.03% (p<0.001). We performed a subset analysis of women who were 9–13 years old in 2006 when the HPV vaccine was approved, now 20–24 years old in 2017. In the pre-vaccine era (2001–2011), the incidence of cancer decreased 2.3% annually (p=0.038), of note, after the introduction of the vaccine (2011–2017), it decreased at 9.6% per year (p=0.002). In the pre-vaccine era (2001–2012), the incidence of new diagnoses of squamous cell carcinoma observed a decrease of 3.1% annually (p=0.004). However, in the post-vaccine era (2012–2017), there was an 11.8% decline in new cases per year (p=0.007). Although there is a decrease in older age groups, there is no difference in the trends pre and post vaccine era, particularly in the age groups who were not eligible for vaccination at that time.

**Conclusions** In our population analysis, our data suggest that the HPV vaccination may have decreased in incidence of cervical cancer in the younger cohort after its approval.

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**EPV039/#175**

**RELATIVE IMPORTANCE OF INDIVIDUAL INSURANCE STATUS AND HOSPITAL PAYER MIX ON SURVIVAL FOR WOMEN WITH CERVICAL CANCER**

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**Objectives** Safety-net hospitals (SNH) are important sites of care especially for vulnerable groups (e.g., uninsured/Medic-aid). We examined the relative contributions of individual insurance status and hospital payer mix on quality of care and survival for women with cervical cancer.