and Ki 67 < 5%). Positivity for CD 10 and ER and PR, focally for Vimentin and negativity for Caldesmon and Actin.

**Conclusion**

ESN is described as a nodule composed of endometrial stromal cells located in the myometrium. It is characterized by its circumscribed and non-invasive nature. From the histological point of view, cell tabs simulating infiltration of less than 3 mm and without vascular invasion are typical. Generally, the definitive diagnosis is made in a hysterectomy sample, because an evaluation of the tumor edges is required to eliminate an LGESS and immunohistochemical criteria are required.

### Abstract 2022-RA-744-ESGO Figure 1

Proportion of patients with complete hepatitis B triple-screening at initiation of gynecologic chemotherapy (P-Chart)

### Abstract 2022-RA-744-ESGO Figure 2

Proportion of gynecologic chemotherapy prescribers ordering HBV screening with chemotherapy initiation (Run Chart)

**Conclusion**

Implementation of four interventions to increase HBV screening in gynecologic oncology chemotherapy patients significantly improved screening rates, achieving our target at 9 months with sustained improvement. Risk factor-based screening lacks sensitivity compared to universal screening which impacts management.

### Abstract 2022-RA-778-ESGO

**RARE GYNECOLOGICAL CANCERS IN A GYNECOLOGIC CANCER CENTER: 11-YEAR EXPERIENCE OF KEM**

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**Introduction/Background**

Many gynecologic cancers fulfill the criteria of a rare tumor with an annual incidence of <6 per 100,000 women. As these tumor entities are difficult to treat, specialized knowledge and skills are necessary. We analyzed the 11-year experience with rare tumors in a tertiary gynecologic oncology center.

**Methodology**

All consecutive patients with rare gynecological cancers treated at our department between 2011 and 2021 were included.
Results 1,460 patients with a rare gynecologic cancer entered our department. 1,092 patients received any kind of therapy and 368 ‘only’ had a consultation. The most common histologic types were: borderline tumors of the ovary (424 pts, 29%), low grade serous ovarian cancer (179 pts, 12.3%), ovarian sex cord-stromal tumors (164 pts, 11.2%), uterine sarcoma (142 pts, 10%), clear cell ovarian cancer (115 pts, 8%), and ovarian germ cell tumors (73 pts, 5%). There was a continuous increase of pts over the years: 237 (2011–2013), 215 (2014–2015), 276 (2016–2017), 347 (2018–2019), 385 (2020–2021) cases. In total, 905 pts came from NRW and 555 patients from other states. Conclusion We observe a centralization of patients from whole Germany with rare gynecological cancers in our center with a rising number of patients during the observation period. This provides a unique chance for further research in rare gynecologic tumors and also allows to offer prospective trials.

Methodology This is a case report of a 47-year-old patient with huge tumor of perineum localised on the left side of perineum, along the vaginal and rectal wall stretching out from the left obturator fossa till the left buttock. The tumor was previously partially debulked in other hospital giving the histopathological diagnosis of aggressive angiomyxoma with high estrogen receptors expression. The adjuvant hormonotherapy of GNRH analogues was then introduced with initial good clinical outcome. After almost one year of clinical remission the flabby tumor grew back to the vast dimensions causing many ailments. After thorough imaging diagnostics excluding local pelvic muscles infiltration and any distant metastases as well the patient was qualified for radical debulking surgery from the perineal access. The resection was completely performed when the proper dissecting plane of this locally aggressive tumor had been found with the anatomic respect of vital vessels and nerves. The perineal plastic surgery followed then the surgical complete excision simultaneously. The postoperative course was uneventful. The patologic report confirmed the previous diagnosis of aggressive angiomyxoma with clear surgical margins.

Results The patient is being strictly followed-up for one year so far with no signs of the relapse both in clinical and imaging examinations.

Conclusion Perineal aggressive angiomyxoma is a very rare tumor with unequivocal tendency of local recurrence. The scant publications suggest that complete surgical resection is the best option to prevent of the disease relapse. Systematic treatment or radiotherapy have not been proved to be effective, however antiestrogenic hormonotherapy is recommended as this kind of perineal tumor usually has pronounced hormonal receptors expression.

Abstract 2022-RA-786-ESGO Figure 1