

Factors Affecting Survival in Egyptian Patients with Advanced Vulval Cancer

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Abstract: INTRODUCTION: Vulvar carcinoma accounts for about 4% of the female reproductive tract cancers and a total of 0.6% of all malignant cancers in females. PATIENTS AND METHODS: This retrospective study aimed to identify the possible risk factors which can impact survival in Egyptian patients with vulval cancer who received treatment at Ain Shams University department of clinical oncology and nuclear medicine in the period from 1-1-2007 till 1-1-2012. RESULTS: Our study included 64 patients [median age 63 years, median overall survival (OS) was 16.5 months]. The factors associated with favourable impact on OS were: Rural residence (17 versus 10 months for patients from urban residence, $p=0.002$), premenopausal status (19 versus 16 months in post menopausal patients, $p<0.001$), low grade histology (19 versus 10 months in patients with high grade, $p<0.001$) and negative resection margins (19 versus 10 months in case of positive margin, $p<0.001$). The factors associated with poor OS were: higher number of offspring (≥ 5 offspring, OS 10 versus 19 months if less than 5 offspring, $p<0.001$), patients presenting with ulcer rather than mass (11 versus 19 months, $p<0.001$ and those with bilateral disease (10 versus 17 months in unilateral disease, $p=0.04$), presence of ≥ 4 positive groin lymph nodes metastases (OS =16 months versus 17 months, $p=0.047$) tumor size ≥ 4 cm (10 versus 17 months in case of <4 cm, $p=0.001$) and depth of stromal invasion (17 versus 16 months in case of <1 cm and ≥ 1 cm respectively, $p=0.015$) CONCLUSION: Urbanization has been linked with more aggressive disease leading to decrease of OS. The age of onset of intercourse did not affect survival as was expected however multiple offspring which could be related to more frequent intercourse had an effect on survival. The offered treatment modalities did not show a superiority probably because all the patients presented in an advanced stage. The anatomical nature of this cancer might be the cause in delayed diagnosis which warrants health education projects for early detection. Further studies are required to assess the risk factors for development of vulval cancer in Egyptian patients.

Keywords: Vulva, Cancer, Retrospective

1. Introduction

Invasive vulvar carcinoma is considered to be one of the rare cancers of female reproductive tract accounting for about 4% of the female genital cancers and 0.6% of all malignant cancers reported in females. Vulval cancers are reported in about 2.5 per 100,000 women years in the developed countries, unfortunately, this cancer is 2-3 times more frequent in the developing world. [1] According to the American Cancer Society, in 2018, there were about 6000 recorded new cases of vulvar cancer and about 1,200 estimated mortalities. The five-year survival rates for vulvar cancer is expected to be 70%. [2] On the other hand, in Egypt, the crude incidence

rates per 100,000 is about 0.3%. [3]

Vulvar carcinoma usually occurs in postmenopausal females with a notable surge in the age-specific incidence with age (50-70 years of age). The labia majora were found to be the most common anatomical site for origin of vulval tumors. The most common histology seen in vulvar cancers is squamous cell carcinoma (SCC) accounting for 90-95% of the cases followed by melanoma, adenocarcinoma, sarcoma as well as basal cell carcinoma. [4, 5]

The most common known etiology for development of vulvar tumors involves infection with the Human Papilloma Virus (HPV). [6] In this group of females there is an association with high-risk sexual activity in addition to

smoking. In the remaining cases (most commonly elderly females with no apparent association to HPV infection) vulval cancer is commonly found on top of chronic dystrophic or inflammatory changes. [4]

The presenting symptoms are commonly local and include itching and pruritus, contact bleeding, offensive discharge, localized pain, mass or ulcer. Biopsy is required to establish the diagnosis. Other diagnostic measures include pelvic ultrasound, CT or MRI scans. In some cases further metastatic work up might be required. [4, 7]

Initial stage is considered the most important factor affecting the future treatment options and the following expected response to therapy. The overall 5-year survival rate of patients with stage I epidermoid invasive cancer might reach 85-90%. Unfortunately, the expected survival rate falls with increasing stage; however, an approximate 5-year survival rate of 40% can be reached, even in patients with positive lymph node metastasis. [8, 9]

Other factors include histological grade and stromal invasion. In a review of the National Cancer Data Base, patients with positive inguinal lymph nodes for malignancy were found to have 5-year variable survival rates (About 64% with 2-cm lesions and 43% with lesions measuring >2 cm in size). In patients with primary tumors of any size, the survival rate was similar whether 1 lymph node was positive or 2-3 lymph nodes were positive (55% vs 59%). However, the survival rate in patients with 4 or more positive nodes was only 33% at 5 years.[8]

Adequate groin treatment is the single most important factor in reducing mortality from vulvar tumors. It is recommended that both inguinal and femoral lymph nodes should be removed, and appropriate dissection should involve the removal of at least 8–10 nodes. [9]

Treatment options depend largely on the presenting stage and they include surgical resection, radiation (Brachytherapy and External beam therapy) as well as chemotherapy (CT). Early-stage vulvar cancer is primarily treated by surgical resection (radical vulvectomy and inguinofemoral lymphadenectomy). The surgical margins ideally should be ≥ 1 cm. Re-excision is considered if margins were found to be positive or less than 8 mm. As an alternative, adjuvant radiation therapy may be considered rather than re-excision. [10]

Adjuvant radiation is determined based on pathological risk factors such as positive lympho-vascular invasion, bilateral malignant disease and positive lymph nodes (LN). Stage II disease with tumor extension up to the distal third of the urethra or vaginal distal third or with malignant anal involvement can be treated with radical local excision in addition to bilateral inguinofemoral lymphadenectomy. Radiation therapy to these regions can also be considered. A National Cancer Data Base (NCDB) analysed the data of 1797 patients with malignant vulval tumors who underwent radical surgery with confirmed positive inguinal nodal involvement and then treated with adjuvant radiation therapy, concluded that the addition of adjuvant chemotherapy to the management plan resulted in a 38% decline in mortality risk.[11]

In case of locally advanced vulvar cancers (namely: bulky stage III and stage IV), treatment options include radical surgery versus concurrent chemotherapy with radiation therapy. It was found that there is no statistically significant difference in overall survival (OS) or in treatment-related adverse events when chemoradiation (primary or neoadjuvant) was compared with primary radical surgical management.[12]

In case of metastatic vulvar cancer (stage IVB), chemotherapy treatment protocols used for metastatic vulvar cancer are similar to those used for metastatic cervical cancers, and combinations of radiotherapy with concurrent chemotherapy can be considered in some cases. [13]

2. Patients and Methods

2.1. Study Design

A retrospective analysis was done for the hospital records of patients diagnosed with vulvar cancer and received treatment at Ain Shams University department of clinical oncology and nuclear medicine in the period from 1-1-2007 till 1-1-2012.

Sixty four patients with a histo-pathological proof of vulvar carcinoma were included in the analysis. The various risk factors were identified and correlated with treatment outcome and survival. Patient risk factors included: age, residence, smoking status, parity, offspring, menstrual status, family history of cancer, past history of cancer and age of onset of sexual activity. Pathological risk factors included: histology, grade, site, depth of invasion, post-operative margins, tumor size, positive lymph nodes, total dissected lymph nodes and stage of disease (TNM). Treatment factors studied included surgery (biopsy versus radical surgery), groin lymph node dissection (LND), concurrent chemotherapy with radiotherapy versus radiotherapy alone. The data collected were analyzed and compared with the published literature. Clinical response was assessed using Response Evaluation Criteria in Solid Tumors criteria version 1.1. (RECIST 1.1)

Overall survival (OS) period was defined as the interval between the pathological diagnosis of the disease (surgery, or core biopsy) and death from any cause.

2.2. Statistical Analysis

All the collected Data was converted into codes and entered to the Statistical Package for Social Science (IBM SPSS) version 20. The qualitative data identified were given as numbers and percentages while quantitative data were presented as mean, median, standard deviations and ranges. Comparisons between two groups (with qualitative data) were done with *Chi-square test* and/or *Fisher exact test* (used instead of Chi-square test when the expected count was found less than 5). *Kaplan Mayer analysis* was used to assess the relation of overall survival (OS) with the other studied parameters. The confidence interval was set to 95% and the margin of error accepted was set to 5%. So, the p-value was considered significant as the following: $P > 0.05$: Non significant, $P < 0.05$: Significant, $P < 0.01$: Highly significant.

3. Results

The median age of the 64 patients included in the study was 63 years (Range 43-68 years). Most of the patients had a good performance status; ECOG PS 1 (81.3%). About 80% of the patients lived in rural areas. Most of the patients (75%) were postmenopausal. All of the patients in the study were multipara and had at least 2 offspring. Regarding the age of onset of sexual activity, half the patients started at age of 18 years. Considering the first presentation, vulval mass was the most common (48.4%), followed by vulval pain (28.1%) and bleeding (23.4%). All the reported cases showed squamous histology (56.3% of cases were low grade) with bilateral disease seen in 37.5%. Unfortunately, all patients were advanced; stage III A (31.3%) and III B (68.8%). Twenty five patients had 4 positive lymph nodes (about 40%), 11 patients with 5 +ve L.N (17.2%), 14 patients with 6 +ve L.N (21.9%), and 14 patients with 12 +ve L.N (21.9%). Thirty one specimens were with negative margins (48.4%) and 33 patients with positive margins (51.6%).

Forty patients underwent surgery with radical intent (62.5%) and the other 24 were palliative surgery (37.5%). Twenty eight patients had radical vulvectomy (43.8%), 12 patients hemivulvectomy (18.8%), 16 patients underwent excision (25%), and the other 8 only did a biopsy (12.5%). Thirty eight patients had lymph node dissection (59.4%), in the other 26 patients, no dissection of lymph nodes was done (40.6%).

As regards the radiation given, 42 patients had the radiation dose of 60 Gy (65.6%), and the other 22 patients had 66Gy (34.4%). Radiation was delivered by multi-energy linear accelerator using 3-D conformal technology with PERCISE planning system. Forty eight patients had concurrent chemotherapy with the radiotherapy (75%), the other 16 had radiation therapy only (25%).

The median overall survival (OS) was 16.5 months (Range

7-20 months). Rural residence of the patient was associated with a favourable statistically significant OS of 17 months versus 10 months in patients from urban residence ($p=0.002$). Pre menopausal patients had a statistically significant longer OS of 19 months versus 16 months in post menopausal patients ($p<0.001$). Higher number of offspring (≥ 5 offspring) was associated with lower OS of 10 months as compared to 19 months in case of <5 offspring ($p<0.001$). As regard findings at initial clinical examination, patients presenting with ulcer had a lower OS (11 months) as compared to patients presenting with a mass (19 months, $p=0.001$) and patients presenting with bilateral vulval disease were associated with the lower OS of 10 months versus 17 months in case of unilateral disease ($p=0.04$). While patients with low grade disease experienced a better OS (19 months versus 10 months in patients with high grade, $p=0.001$). Patients with ≥ 4 positive lymph nodes were associated with unfavourable survival outcome (16 months) in comparison to those with less than 4 positive groin LN (17 months, $p=0.047$). Patients with negative resection margins were associated with the highest OS (19 versus 10 months in case of positive margin, $p<0.001$). The depth of stromal invasion had an impact on survival, where an invasion ≥ 1 cm was associated with worse OS (17 versus 16 months in case of depth <1 cm, $p=0.015$). The tumor size also affected OS where tumors ≥ 4 cm were associated with worse OS (10 months in comparison to 17 months if tumor size < 4 cm, $p=0.001$).

Other studied factors including radical versus palliative surgery, groin lymph node dissection, dose of radiation therapy, concurrent radiation and chemotherapy versus radiation only did not have a statistically significant impact on overall survival.

COX regression multivariate analysis showed that menstrual status, pathological grade and resection margins had an impact on survival.

Table 1. Descriptive data of the studied sample.

Factor Assessed		No.	%
Age Groups (years)	41-50	20	31.25
	51-60	0	0
	61-70	44	68.75
Performance (PS)	1	52	81.3
	2	12	18.8
Residence	Rural	51	79.7
	Urban	13	20.3
Family History	No	56	87.5
	yes	8	12.5
Menstrual Status	Post-menopause (Pre)	48	75
	Pre-menopause (Post)	16	25
	2	8	12.5
Number of offspring	3	16	25
	4	8	12.5
	5	32	50
Age of Onset of Sexual Activity	16	10	15.6
	18	32	50.0
	19	14	21.9
	21	8	12.5
Presentation	Bleeding (B)	15	23.4
	Mass (M)	31	48.4
	Pain (P)	18	28.1

Factor Assessed		No.	%
Examination	Mass (M)	31	48.4
	Ulcer (U)	33	51.6
Site	Bilateral Labia Major	24	37.5
	Clitorius	10	15.6
	Unilateral Labium Major	30	46.9
Grade	Low Grade (1)	28	43.8
	High Grade (2)	36	56.3
Stage	IIIA	20	31.3
	IIIB	44	68.8
Tumor size (cm)	3	21	32.8
	4	12	18.8
	5	10	15.6
	8	1	1.6
	9	18	28.1
	10	2	3.1
	4	25	39
Positive LN	5	11	17.2
	6	14	21.9
	12	14	21.9
Depth (cm)	0.7	3	4.7
	0.8	7	10.9
	0.9	7	10.9
	1	33	51.6
Margin	1.5	14	21.9
	Negative (n)	31	48.4
	Positive (y)	33	51.6
Surgery Aim (Radical verus Not)	n	24	37.5
	R	40	62.5
Surgery	Biopsy	8	12.5
	Excision	16	25
	Hemivulvectomy	12	18.8
	Radical Vulvectomy	28	43.8
Groin LN Dissection	n	26	40.6
	y	38	59.4
Radiation Dose	60 Gy	42	65.6
	66 Gy	22	34.4
Concurrent Chemotherapy and Radiation	n	16	25
	y	48	75

Table 2. Descriptive data of String Variables in the studied population.

	Age (years)	Offspring	Age of Onset of Sexual Activity (Years)	Tumor size (cm)	Positive LN	Depth (cm)	OS (months)
Median	63	3	18	4	5	1	16.5
Std. Deviation	8.141	1.153	1.397	2.606	3.094	0.2478	4.9618
Minimum	43	2	5	3	4	0.7	7
Maximum	68	5	16	10	12	1.5	20

Table 3. Factors associated with an impact on overall survival.

Factor		Median OS (months)	95% CI (months)	p -value
Residance	Rural (R)	17	16.385-17.615	0.002
	Urban (U)	10	8.777-11.223	
Menstrual Status	Postmenopause (post)	16	12.103-19.897	<0.001
	Premenopause (pre)	19	17.693-20.307	
Offspring	<5	19	18.040-19.960	<0.001
	≥5	10	8.073-11.927	
Clinical Presentation	Mass (M)	19	16.853-21.147	0.001
	Ulcer (U)	11	7.483-14.517	
Disease Lateralization	Bilateral (bi)	10	7.952-12.048	0.04
	Unilateral (uni)	17	16.604-17.396	
Pathological Grade	Low grade (1)	19	17.829-20.171	<0.001
	High grade (2)	10	7.850-12.150	
Tumor Size (cm)	≥ 4 (n)	10	6.619-13.381	0.001
	<4 (y)	17	13.636-20.364	
Depth of Stromal invasion (cm)	<1	17	14.983-19.017	0.015
	≥1	16	7.984-24.016	
Resection Margins	Negative (n)	19	18.523-19.477	<0.001

Factor		Median OS (months)	95% CI (months)	p -value
Positive Groin Lymph nodes	Positive (y)	10	8.010-11.990	0.047
	≥4	16	12.399-19.601	
	<4	17	13.996-20.004	

Table 4. COX regression analysis for the studied variables in univariate analysis.

Factor	B	SE	Wald	df	Sig.	Exp (B)
Residence	.478	.547	.764	1	.382	1.614
Menstrual Status	1.129	.487	5.379	1	.020	3.091
Offspring	-.461	.389	1.404	1	.236	.631
Clinical Presentation	-.295	.482	.375	1	.540	.744
Disease Lateralization	1.127	.602	3.496	1	.062	3.085
Grade	-2.320	.648	12.822	1	.000	.098
Tumor size	-.123	.514	.057	1	.811	.884
Positive Groin Lymph nodes	.422	.492	.737	1	.391	1.526
Depth of Stromal invasion	-.135	.405	.111	1	.739	.874
Resection Margins	-1.414	.477	8.797	1	.003	.243

4. Discussion

Primary vulvar tumors are the 20th most common female cancers, mainly seen in age groups 50-70 years. [14] Okolo *et al.* examined 78 cases of vulvar cancer from 1981 to 2008. The mean age in this clinical study was 49.7 years with a peak incidence found in the fifth decade. [15] In a study by Singh *et al.* which comprised 41 cases of vulvar cancer, the mean age was 52 years, but the peak incidence was observed in the sixth and seventh decades. Incidence is significantly high in multiparous and postmenopausal women, however, not related to survival. [16]

These findings were confirmed by our study where all the studied females were multipara and 75% were postmenopausal. In addition, the median age was 63 years, however the sample included patients less than 50 years (15.2%). However, in our study, Pre menopausal patients had a statistically significant longer OS of 19 months versus 16 months in post menopausal patients ($p < 0.001$). Higher number of offspring (≥ 5 offspring) was associated with lower OS of 10 months as compared to 19 months in case of < 5 offspring ($p < 0.001$).

The most common type of vulvar cancer is squamous cell carcinoma (SCC) accounting for more than 90% of the cases; the remaining types are rare and include: melanoma, adenocarcinoma, sarcoma and basal cell carcinoma. Okolo *et al.* showed that 73.61% of vulvar cancers included in their study were squamous cell type (SCC). [15] Similarly, in a study by Singh *et al.*, SCC accounted for 97.56% of the cases. [16]

Furthermore, a study from Tunisia by Kehila *et al.* showed similar findings, which is very similar to the results of our study where all patients were SCC. [17]

Kosary *et al.* studied the prognostic impact of FIGO stage, histology, histologic grade, age and race in survival for cancers of the female gynecological (cervix, endometrium, ovary, vulva, vagina) using cases obtained from the National Cancer Institute's Surveillance, Epidemiology, and End Results (SEER) program that were diagnosed between 1973 and 1987. [18]

Paladini *et al.* published a study of 75 patients with inguinal and/or pelvic lymph node malignant metastases from vulvar cancer of squamous histology. Among the parameters examined in this study were: size as well as location of the primary cancer (T), depth of invasion of the stromal tissues, histological grade, positive cancerous lympho-vascular space involvement (LVSI), local immune response, FIGO stage, number of involved lymph nodes (LN). Among the variables related to the primary tumors, only size of the primary tumor and LVSI had an impact on with survival ($P < 0.003$ and $P < 0.02$, respectively). [19] These findings are similar to our results where we found that patients with primary tumors ≥ 4 cm were associated with worse OS (10 months in comparison to 17 months if tumor size < 4 cm, $p = 0.001$).

While some clinical studies did not observe any association between the number of metastatic lymph nodes and the risk of recurrence, other trials have identified that two or more positive lymph nodes, extracapsular spread as well as metastatic lesions were poor prognostic factors. [20]

More recent clinical studies and reviews show that even one positive lymph-node metastasis significantly affects prognosis negatively compared with lymph node-negative patients and in a further analysis by Woelber *et al.*, the number of affected nodes was highly statistically significant as a prognostic factor in the patients who did not receive adjuvant treatment. [21] What is interesting in our analysis is that we evaluated the significance of the number of lymph node metastases in a group of patients who were lymph node positive while most previous studies compared lymph node positive with negative patients in general.

5. Conclusion

We have found that patients with urban residence have been linked with more aggressive disease leading to decrease of OS. The age of onset of intercourse did not affect survival as we were expecting, however, multiple offspring which could be related to more frequent intercourse had an effect on survival. The offered treatment modalities did not show a superiority probably because all the patients presented in an advanced stage. The anatomical nature of this cancer might

be the cause in delayed diagnosis which warrants health education projects for early detection. Further studies are required to assess the risk factors for development of vulval cancer in Egyptian patients.

References

- [1] Howlader N, Noone AM, Krapcho M, et al. SEER Cancer Statistics Review, 1975-2012, National Cancer Institute. Bethesda, MD: April 2015.
- [2] American Cancer Society. *Cancer Facts and Figures 2018*. Atlanta, Ga: American Cancer Society; 2018.
- [3] Ibrahim AS, Khaled HM, Mikhail N, Baraka H and Kamel H. Cancer Incidence in Egypt: Results of the National Population-Based Cancer Registry Program. *J Cancer Epidemiol*. 2014; 2014: 437971.
- [4] Eifel PJ, Berek JS, Markman MA. Cancer of the cervix, vagina, and vulva. In: DeVita VT Jr., Lawrence TS, Rosenberg SA, editors. *Cancer: Principles and Practice of Oncology*. 9th ed. Philadelphia, PA: Lippincott Williams & Wilkins; 2011. p. 1311-44.
- [5] Hacker NF, Eifel PJ, van der Veldenc J. FIGO cancer report 2012. Cancer of the vulva. *Int J Gynecol Obstet* 2012; 119 Suppl 2: S90-6.
- [6] Rakislova N, Saco A, Sierra A, Del Pino M, Ordi J. Role of Human Papillomavirus in Vulvar Cancer. *Adv Anat Pathol*. 2017 Jul; 24 (4): 201-214.
- [7] Committee Opinion No. 675 Summary: Management of Vulvar Intraepithelial Neoplasia. *Obstet Gynecol*. 2016 Oct. 128 (4): 937-8.
- [8] Creasman WT, Phillips JL, Menck HR. The National Cancer Data Base report on early stage invasive vulvar carcinoma. The American College of Surgeons Commission on Cancer and the American Cancer Society. *Cancer*. 1997 Aug 1. 80 (3): 505-13.
- [9] Hacker NF. Vulvar cancer. In: Berek JS, Hacker NF, editors. *Berek and Hacker's Gynecologic Oncology*. 5th ed. Philadelphia: Lippincott Williams & Wilkins; 2009. p. 536-75.
- [10] Khanna N, Rauh LA, Lachiewicz MP, Horowitz IR. Margins for cervical and vulvar cancer. *J Surg Oncol*. 2016 Mar. 113 (3): 304-9.
- [11] Gill BS, Bernard ME, Lin JF, Balasubramani GK, Rajagopalan MS, Sukumyanich P, et al. Impact of adjuvant chemotherapy with radiation for node-positive vulvar cancer: A National Cancer Data Base (NCDB) analysis. *Gynecol Oncol*. 2015 Jun. 137 (3): 365-72.
- [12] Shylasree TS, Bryant A, Howells RE. Chemoradiation for advanced primary vulvar cancer. *Cochrane Database Syst Rev*. 2011 Apr 13.
- [13] Gadducci A, Cionini L, Romanini A, Fanucchi A, Genazzani AR. Old and new perspectives in the management of high-risk, locally advanced or recurrent, and metastatic vulvar cancer. *Crit Rev Oncol Hematol*. Ireland: 2006. 227-41.
- [14] Nicoletto MO, Parenti A, Del Bianco P, Lombardi G, Pedrini L, Pizzi S, Carli P, Della Palma M, Pastorelli D, Corti L, Becagli L. Vulvar cancer prognostic factors. *Anticancer Res*. 2010 Jun; 30 (6): 2311-7.
- [15] Okolo CA, Odubanjo MO, Awolude OA, Akang EE. A review of vulvar and vaginal cancers in Ibadan, Nigeria. *N Am J Med Sci* 2013; 6: 76-81.
- [16] Singh N, Negi N, Srivastava K, Agarwal G. A cohort study of vulvar cancer over a period of 10 years and review of literature. *Indian J Cancer* 2016; 53: 412-5.
- [17] Kehila M, Chanoufi M. Vulvar cancer in Tunisia: Epidemiological and clinicopathological features multicentric study. *Journal of the Egyptian National Cancer Institute* Volume 29, Issue 2, June 2017, Pages 95-98.
- [18] Kosary CL. FIGO stage, histology, histologic grade, age and race as prognostic factors in determining survival for cancers of the female gynecological system: an analysis of 1973-87 SEER CASES of cancers of the endometrium, cervix, ovary, vulva and vagina. *Seminars in surgical oncology* 10 (1), 31-46, 1994.
- [19] Paladini D, Cross P, Lopes A, Monaghan J. Prognostic significance of lymph node variables in squamous cell carcinoma of the vulva. *Cancer* 74 (9), 2491-2496, 1994.
- [20] Lataifeh I, Carraro Nascimento M., Nicklin J., Perrin L., Crandon A., Obermair A. (2004) Patterns of recurrence and disease-free survival in advanced squamous cell carcinoma of the vulva. *Gynecol Oncol* 95: 701-705.
- [21] Oonk M., van Hemel B., Hollema H., de Hullu J., Ansink A., Vergote I., et al. (2010) Size of sentinel-node metastasis and chances of non-sentinel-node involvement and survival in early stage vulvar cancer: results from GROINSS-V, a multicentre observational study. *Lancet Oncol* 11: 646-652.