

Original Research Article

THE THERAPEUTIC IMPACT OF LYMPHADENECTOMY IN THE MANAGEMENT OF EPITHELIAL OVARIAN CANCER: A SINGLE INSTITUTION EXPERIENCE

ABSTRACT

Aim: To assess the impact of lymphadenectomy on overall survival (OS) and progression free survival (PFS) in patients with epithelial ovarian cancer (EOC).

Study design: Retrospective observational study

Place and duration of study: All patients diagnosed with epithelial ovarian, fallopian tube or primary peritoneal cancer treated in the Department of Gynecologic Oncology in a tertiary care hospital in South India from January 2012 onwards. All patients' follow up data was prospectively updated till 30 April 2017.

Methodology: We included 83 patients who met the inclusion criteria. The patients were classified into two groups based on the number of lymph nodes (LN) harvested (< 30 lymph nodes and \geq 30 nodes). Lymphadenectomy was considered systematic (SLND) when the harvest was \geq 30 nodes on the pathologic specimen.

Results: Out of the 83 cases, 70 (84.3%) had a high-grade carcinoma and 68 (81.9%) had serous histology. Lymph node dissection was performed in 74 (89.2%) patients. Lymph node harvest of > 30 nodes was seen in 43 (51.8%) cases and the median number of removed lymph nodes was 44 (IQR 25– 75%: 38–52). Complete cytoreduction (defined as no macroscopic tumor) was achieved in 52 (62.7%) cases. Kaplan-Meier curve and log rank test were used to compare the survival curves between clinical variables. Univariate cox regression analysis revealed that performance status, node positivity and residual disease were the significant predictors of mortality considering the overall survival and progression free survival. However, multivariate cox regression analysis revealed that none of the variables were significant predictors of mortality.

Conclusion: Systematic lymphadenectomy does not have a significant impact on improving overall survival (OS) or progression free survival (PFS) in epithelial ovarian cancers. Performance status and residual disease were important predictors of mortality.

Keywords:

Keywords Systematic lymphadenectomy, epithelial ovarian cancer, survival, cytoreduction

1. INTRODUCTION

Ovarian cancer is the most fatal of gynecologic malignancies [1] The disease is often diagnosed in advanced stage and long-term survival is 30%-40%. The common routes of spread of ovarian cancer are by peritoneal implantation and lymphatic dissemination [2,3].

21 Surgical staging in early cases and primary cytoreductive surgery followed by platinum-
22 based chemotherapy is the standard treatment. The clinical benefit of lymphadenectomy in
23 women with early disease apart from providing more accurate staging is unclear [4] The
24 effect of lymph node dissection on progression-free survival (PFS) and OS in patients with
25 advanced ovarian cancer is still unknown. The benefit of complete cytoreduction, on the
26 overall survival (OS) is well established [5-7]. Whether optimum surgery for epithelial ovarian
27 cancer (EOC) should include retroperitoneal lymphadenectomy along with intraperitoneal
28 procedures as part of maximal cytoreductive is not well defined. Various survival outcomes
29 have been documented with systematic lymphadenectomy and with resection of only bulky
30 nodes [8-11]. The practices are variable in different centers across the globe. There are very
31 few Indian studies addressing the role of systematic lymph node dissection (SLND) in
32 Epithelial ovarian cancer (EOC). This study is an attempt to assess the impact of systematic
33 pelvic and para-aortic lymphadenectomy on survival in EOC.
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36 37 **2. MATERIAL AND METHODS:**

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39 This was an observational study involving the record review of patients with a diagnosis of
40 epithelial ovarian, fallopian tube or primary peritoneal cancer who underwent primary surgery
41 or interval cytoreduction in the Department of Gynecologic Oncology in a tertiary care
42 hospital from January 2012 onwards. Follow up data of the patients was prospectively
43 updated till 30 April 2017. Women who were operated elsewhere or who received less than
44 three courses of adjuvant chemotherapy, when indicated, were also excluded from the
45 study.

46 The relevant information retrieved consisted of demographic data, clinical details,
47 investigations, intraoperative findings, details of surgery, histopathology reports,
48 postoperative period, adjuvant / neoadjuvant chemotherapy (NACT), follow up, recurrence
49 and death. All the cases were staged as per the FIGO 2014 staging for ovarian cancers. The
50 FIGO stage for the cases operated prior to 2014 was reclassified according to the FIGO
51 2014 staging.

52 Systematic pelvic LND involved removal of all pelvic lymphatic tissues in front of, behind,
53 and between the iliac vessels up to the bifurcation of the aorta, and down to the obturator
54 fossa and the pelvic floor. Para-aortic lymphadenectomy extended up to the renal vessels,
55 removing all lymphatic tissues around and between the aorta and vena cava. The number of
56 LN harvested was considered as representative of the extent of dissection. Systematic
57 lymphadenectomy (SLND) was defined as a complete procedure when at least 30 lymph
58 nodes were reported in the pathologic specimen. [12] Surgery was performed by the same
59 dedicated team over the period of study. The Surgical procedures were quantified using the
60 Surgical Complexity Score (SCS) previously described by Aletti et al [13].

61 OS was calculated as the number of months from the date of diagnosis to either the date of
62 death or the date censored and PFS was calculated as the number of months from the date
63 of diagnosis to either the date of recurrence or the date censored.

64 Descriptive statistics were reported using mean and standard deviation for continuous
65 variables and number and percentages for the categorical variables. Kaplan -Meier curve,
66 log rank test, univariate and multivariate Cox regression analysis were used to explore the
67 impact of different covariates on OS and PFS. Probability value < 0.05 was considered
68 statistically significant. Statistical analysis was performed using an SPSS statistical package
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72**3. RESULTS**

Variable	Number of patients	Percentage (%)
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73 [(A total of 83 cases met the inclusion criteria for this study. Patient and tumor characteristics
74 are shown in Table 1 and 2 respectively. Mean age of the study cases was 51.4 years +/-
75 12.8 with median follow up of 30.5 months. Sixty-nine (83.1%) were younger than 65 years.
76 Fifty-two (62.7%) women were postmenopausal. Only eight were nulliparous (9.6%). Most
77 of the patients (89.2%) did not have a family history of any malignancy. Thirty-seven (44.6%)
78 women were in a good performance status (Eastern Cooperative Oncology group; ECOG 0).
79 Forty-one women (49.4%) belonged to FIGO stage IIIC. Most of the patients had a high-
80 grade carcinoma (n = 70; 84.3 %) and a serous histology (n =68; 81.9 %).
81 Lymphadenectomy was performed in 74 (89.2%) patients. Lymphadenectomy with a node
82 count \geq 30 (complete systematic LND group) was found in 43 (51.8%) cases and the
83 median number of removed lymph nodes was 44(IQR 25– 75%: 38–52). The median blood
84 loss in this group was 600ml (IQR 25-75%: 500-1000ml). Among the rest of the patients
85 (lymph node count < 30), the median number of dissected nodes was 23 (IQR 25–75%: 17-
86 27). The median blood loss in this group was 700ml (IQR 25-75%: 400-1000ml). There was
87 no significant difference in the number of cases which needed ICU care, blood transfusion
88 and postoperative morbidity in either group. Lymph node metastases was present in 27 of 74
89 (36.5%) patients, nine patients did not undergo lymphadenectomy. Among the node positive
90 cases, 17 / 27 (62.9%) had normal sized non-suspicious nodes on intra-operative
91 assessment. Complete gross cytoreduction (defined as no macroscopic tumor) was
92 achieved in 52 (62.7%) cases, cytoreduction with gross residual disease of 1–10 mm in 22
93 (26.5%) and gross residual disease > 10 mm in nine (10.8%) patients.

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Total		83	100
Variable	Number of patients		Percentage (%)
<= 64 yrs	69	83.1	
> 64 yrs	14	16.9	
Menopausal status			
Postmenopausal	52	62.7	
Premenopausal	31	27.3	
Treatment			
Primary surgery	58	69.9	
Neoadjuvant chemotherapy	25	30.1	
Performance status			
ECOG 0	37	44.2	
ECOG > 0	46	55.4	
FIGO stage			
I	21	25.2	
II	3	3.6	
III	51	61.5	
IV	8	9.7	

Table 1. Patient characteristics

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Table 2. Tumor characteristics

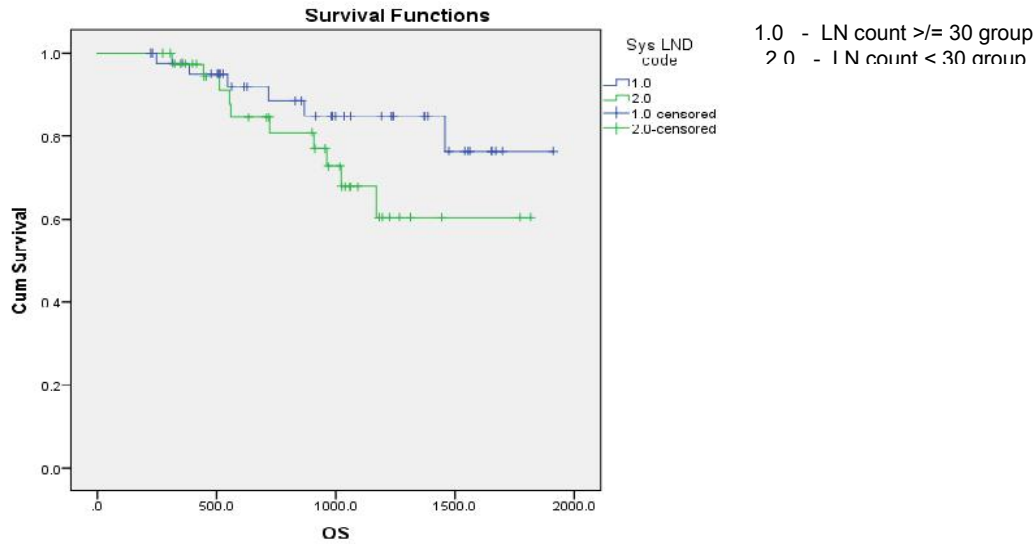
Total	83	100
Grade		
Low	13	15.7
High	70	84.3
Histology		
Serous	68	81.9
Others	15	18.1
pN stage		
Nx	9	10.8
N0	47	56.7
N1	27	32.5
Lymph node resected		
< 30	40	48.2
≥ 30	43	51.8
Residual disease		
Nil	52	62.7
< / =10mm	22	26.5
> 10mm	9	10.8
Surgical complexity score		
Low	9	10.8

Intermediate	68	81.9
High	6	7.3

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Considering the outcome as mortality, the mean OS and PFS for the all the patients was 52.4 months and 48.76 months respectively. Among the women who underwent a complete systematic lymphadenectomy, the median OS was 55.7 months vs 49.0 months among those where the lymph node harvest count was < 30. Although median survival was slightly higher among women who underwent a complete systematic lymphadenectomy, there was no significant difference as compared to women where the lymph node harvest count was < 30. (Fig 1)

Fig 1. Overall survival (OS) in days. Kaplan - Meier

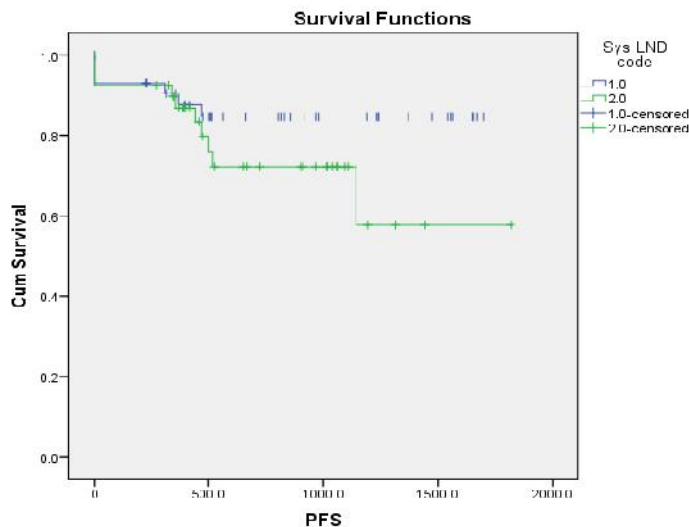


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The median PFS in the group which underwent complete systematic lymphadenectomy was 49.0 months and 43.46 months for the other group with no significant difference . (Fig 2)

Fig 2. Progression free survival (PFS) in days. Kaplan - Meier

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OS was significantly different between performance status, nodal involvement (N

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stage), systematic lymphadenectomy and residual disease using Kaplan Meier analysis. Subjects with better performance status, negative nodes, who underwent complete systematic lymphadenectomy and had nil residual disease, had a higher survival as compared to other groups.

Univariate cox regression analysis revealed that performance status, N stage and residual disease were the significant predictors of mortality considering the overall survival. Subjects with poor performance status had 12.6 times higher hazards, positive nodes had 2.2 times higher hazards and residual disease had 27 times higher hazards as compared to other groups.

Multivariate cox regression revealed that none of the above variables were significant predictors of mortality. However, subjects with positive nodes had higher hazards of mortality as compared to negative nodes (p=0.06). Similarly, when progression free survival was considered, performance status, N stage and residual disease were significant predictors of recurrence. Although none of the variables were significant in the multivariate analysis, N stage and residual disease had higher hazards of mortality (Table 3).

Table 3. Univariate and multivariate analysis of prognostic factors

Variable	N	Univariate analysis			Multivariate analysis		
		HR	95%CI	P value	HR	95%CI	P value
Age (years)							
>64	1.	1.56	0.50-0.856	0.43	1.07	0.29-3.93	0.91

Variable	N	Univariate analysis			Multivariate analysis		
<64	2.	1	-	-	-	-	-
Performance status							
ECOG 0	3.	1	-	-	-	-	-
ECOG 1	40	0.040	0.004-0.360	0.004	0.21	0.01-3.75	0.29
ECOG 2	6	0.428	0.136-1.348	0.14	-	-	-
pN stage							
N0	4.	1	-	-	-	-	-
N1	5.	7.54	2.06-27.51	0.002	2.07	0.24-17.57	0.50
Nx	6.	4.72	0.95-23.47	0.06	5.47	0.93-31.78	0.06
Lymph node count							
<30	7.	2.13	0.76-5.97	0.14	-	-	-
≥30	8.	-	--		-	-	-
Residual disease							
9. Nil	10.	1	-	-	-	-	-
1-10 mm	11.	3.7	1.04-13.1	0.04	1.52	0.35-6069	0.57
> 10mm	12.	7.56	2.12-26.83	0.002	3.55	0.77-16.35	0.10

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3.1 DISCUSSION:

Comprehensive surgical staging in early disease and optimal cytoreduction in advanced stage with platinum-based chemotherapy is the mainstay treatment of EOC [14-16]. Lymphatic spread is common in advanced EOC. Histopathology showed 27 (36.5%) positive nodes in our cohort, which is slightly less compared to other studies. This could probably be

191 because of high number of early stage EOCs. In the present study 17 / 27 (62.9%) node
192 positive patients had normal sized non-suspicious nodes on intraoperative assessment,
193 emphasizing the inaccuracy of clinical assessment as noted in earlier studies [17].
194 The role of lymphadenectomy for staging procedure is well established though the
195 therapeutic role is debated [18,19]. The impact of lymphadenectomy in advanced ovarian
196 cancer is less clear and guidelines are lacking whether systematic pelvic and para aortic
197 lymphadenectomy be performed with debulking surgery [20-24]. The decision to perform
198 systematic lymphadenectomy is by the surgeon's discretion or the policy of the hospital.
199 In advanced ovarian cancer there is a high rate of involved nodes and survival benefits are
200 documented with systematic aortic and pelvic lymphadenectomy [25]. Studies also have
201 shown that removal of bulky nodes improved OS in optimally debulked ovarian carcinoma
202 and there are studies showing no advantage in resection of clinically negative nodes [20-
203 24,26]. No standardized techniques were mentioned in most of the studies and number of
204 nodes dissected were also variable. The node count for systematic verses incomplete nodal
205 dissection was also not uniform. The lymphadenectomy in ovarian neoplasms (LION) is the
206 only prospective randomized trial which demonstrated no significant benefit of either PFS or
207 OS and endorsed omitting routine lymphadenectomy in clinically node negative advanced
208 ovarian cancer patients with macroscopic complete tumor resection [26].
209 Kim et al in 2010, conducted a meta-analysis comparing the impact of systematic
210 lymphadenectomy and non-systematic lymphadenectomy (random removal or less removal
211 or no removal of pelvic and para aortic nodes). They found an increased OS in all stage
212 disease with systematic lymphadenectomy. But sub-analysis of the two RCT included in their
213 study showed no difference in OS between systematic and non-systematic
214 lymphadenectomy [24]. In a study by Gao et al, 5-year OS in systematic lymphadenectomy
215 group was higher than the non-systematic lymphadenectomy group. In their analysis of 14
216 studies, the difference was seen in observation studies and advanced stage disease and no
217 difference was seen in the RCT, early stage disease and residual disease \leq 2cm. The
218 definition of unsystematic lymphadenectomy was inconsistent even in this study [27]
219 In the present study, in women who underwent a complete systematic lymphadenectomy,
220 the median OS was 55.7 months vs 49.0 months among those where the lymph node
221 harvest was $<$ 30, but was not statistically significant. Overall survival was significantly
222 different between performance status, N stage, systematic lymphadenectomy and residual
223 disease using Kaplan Meier analysis. Subjects with better performance status, negative
224 nodes, who underwent complete systematic lymphadenectomy and nil residual disease, had
225 a higher survival as compared to other groups.
226 In our study there is not much of a difference between overall and progression free survival
227 as the number of events have still not happened at the time of analysis and the analysis was
228 time bound.
229 The limitations of the present study is that it is a retrospective study with less numbers and
230 hence no sub group analysis could be done.
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234 4. CONCLUSION

235 Systematic lymphadenectomy does not have a significant impact on improving overall or
236 progression free survival in epithelial ovarian cancers. However, N stage and residual
237 disease had higher hazards of mortality.

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