



# The Therapeutic Impact of Lymphadenectomy in the Management of Epithelial Ovarian Cancer: A Single Institution Experience

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## 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, complete SLND was done in 43 (51.8%) cases and the median number of removed lymph nodes was 44 (IQR 25– 75%: 38–52). Among the women who underwent a complete SLND, the median OS was 55.7 months vs 49.0 months among those where the lymph node harvest count was < 30 (*P* value – 0.16). The median PFS in the complete SLND group was 49.0 months and 43.46 months for the other group with no significant difference (*P* value – 0.18). Though there was no significant difference in OS and PFS, there was a trend towards improved survival with complete SLND group beyond 500 days.

**Conclusion:** Complete SLND group showed a trend towards improved OS and PFS, though statistically not significant. Further investigation is warranted.

*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 an advanced stage and the long-term survival is 30%-40%. The common routes of spread of

ovarian cancer are by peritoneal implantation and lymphatic dissemination [2,3].

The standard of care in early epithelial ovarian cancer (EOC) requires lymph node dissection to accurately stage the disease and to decide on the requirement of adjuvant chemotherapy.

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Cytoreduction to no visible disease followed by platinum-based chemotherapy is the standard treatment in advanced disease. Lymphadenectomy is thus an integral part of the treatment of EOC in both staging and retroperitoneal debulking.

But the clinical benefit of lymphadenectomy in women with early disease apart from providing more accurate staging is unclear [4]. In patients with advanced ovarian cancer, the effect of lymph node dissection on progression-free survival (PFS) and overall survival (OS) is also unclear but the benefit of complete cytoreduction, on the OS is well established [5-7]. The practices are variable in different centres across the globe. Removal of bulky nodes, to lymph node sampling to systematic lymph node dissection, is being advocated in various centres. Also, there is a great variation in the yield of lymph nodes dissected. Various survival outcomes have been documented with systematic lymphadenectomy and with resection of only bulky nodes [8-11]. There are very few Indian studies addressing the role of systematic lymph node dissection (SLND) in Epithelial ovarian cancer (EOC).

This study is an attempt to assess the impact of systematic pelvic and para-aortic lymphadenectomy on survival in EOC.

## **2. MATERIALS AND METHODS**

This is an observational study involving the record review of patients with a diagnosis of epithelial ovarian, fallopian tube or primary peritoneal cancer that has undergone primary or interval cytoreductive surgery in the Department of Gynecologic Oncology from January 2012 onwards. The Institutional ethics committee approval was taken prior to beginning the study.

Follow up data of the patients was prospectively updated till 30 April 2017. Patients who underwent surgery elsewhere or had received less than three courses of chemotherapy in the adjuvant setting, when indicated, were excluded from the study.

An informed written consent was obtained from patients who had completed treatment and were available for follow up. Telephonic verbal consent was obtained in women who had not visited the hospital for recent follow-up. The relevant information retrieved consisted of demographic data, clinical details, investigations, intraoperative findings, details of surgery, histopathology reports, postoperative period, adjuvant/neoadjuvant chemotherapy (NACT),

follow up, recurrence and death. All the cases were staged as per the latest FIGO staging (2014) for ovarian cancers. The FIGO stage for the cases operated prior to 2014 was reclassified to the latest FIGO staging (2014).

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

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

Descriptive statistics were reported using mean and standard deviation for continuous variables and number and percentages for the categorical variables. Kaplan -Meier curve, log-rank test, univariate and multivariate Cox regression analysis were used to explore the impact of different covariates on OS and PFS. A probability value < 0.05 was considered statistically significant. Statistical analysis was performed using SPSS statistical package 23.

## **3. RESULTS**

A total of 83 cases met the inclusion criteria of our study. Patient and tumour characteristics are shown in Tables 1 and 2 respectively. Mean age of the study cases was 51.4 years +/- 12.8. The median follows up of the patients was 30.5 months. Sixty-nine (83.1%) patients were less than 65 years. Fifty-two (62.7%) women were postmenopausal. Only eight were nulliparous (9.6%). Majority of the patients (89.2%) did not have any family history of malignancy. Thirty-seven (44.6%) women were in good performance status (Eastern Cooperative Oncology group;

ECOG 0). Forty-one women (49.4%) belonged to FIGO stage IIIC. Most of the patients had a high-grade carcinoma (n = 70; 84.3 %) and a serious histology (n =68; 81.9 %). The other histologies noted were mucinous (n = 7), endometrioid (n = 4), clear cell (n = 3) and carcinosarcoma (n =1). Lymphadenectomy was performed in 74 (89.2%) patients. Lymphadenectomy with a node count  $\geq$  30 was considered as complete systematic LND group and was found in 43 (51.8%) cases. The median number of lymph nodes extirpated was 44(IQR 25– 75%, 38–52). The median blood loss in this group was 600ml (IQR 25-75%: 500-1000ml). Among rest of the patients with the lymph node harvest of < 30, the median number of dissected nodes was 23 (IQR 25–75%: 17-27). The median blood loss in this group was 700ml (IQR 25-75%: 400-1000ml). There was no significant difference in blood transfusion rate or ICU requirement in either group. The postoperative complications were seen in 14/43 cases (32.5%) in the complete SLND group and in 12/40 cases (30%) in the other group. Lymph node metastases were present in 27 of 74 (36.5%) patients, nine patients did not undergo lymphadenectomy. Among the node-positive cases, 17 / 27 (62.9%) had normal sized non-suspicious nodes on intra-operative assessment. Complete gross cytoreduction (defined as no macroscopic tumour) was achieved in 52 (62.7%) cases, cytoreduction with the gross residual disease of 1–10 mm in 22 (26.5%) and gross residual disease > 10 mm in nine (10.8%) patients.

The mean OS was 52.4 months and PFS for all the patients was 48.76 months. Among the women who underwent a complete systematic lymphadenectomy, the median OS was 55.7 months vs 49.0 months for patients in whom the lymph node harvest count was < 30 (*P* value – 0.16). Although there was 6.7 months benefit in OS for patients with complete systematic

lymphadenectomy, especially on longer follow up, it was not statistically significant (Fig. 1).

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 (*P* value – 0.18) (Fig 2). However, a trend towards improved PFS was noted with the complete SLND group on longer follow up.

The OS was significantly different with the performance status, nodal involvement (N stage), systematic lymphadenectomy and residual disease using Kaplan Meier analysis. Subjects with better performance status, negative nodes, who underwent complete systematic lymphadenectomy with 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 was significant predictors of mortality. However, subjects with positive nodes had higher hazards of mortality as compared to negative nodes (*P* value=0.06).

Similarly, when PFS was considered, performance status, N stage and residual disease were significant predictors of recurrence. Although none of the variables was significant in the multivariate analysis, N stage and residual disease had higher hazards of recurrence (Table 3).

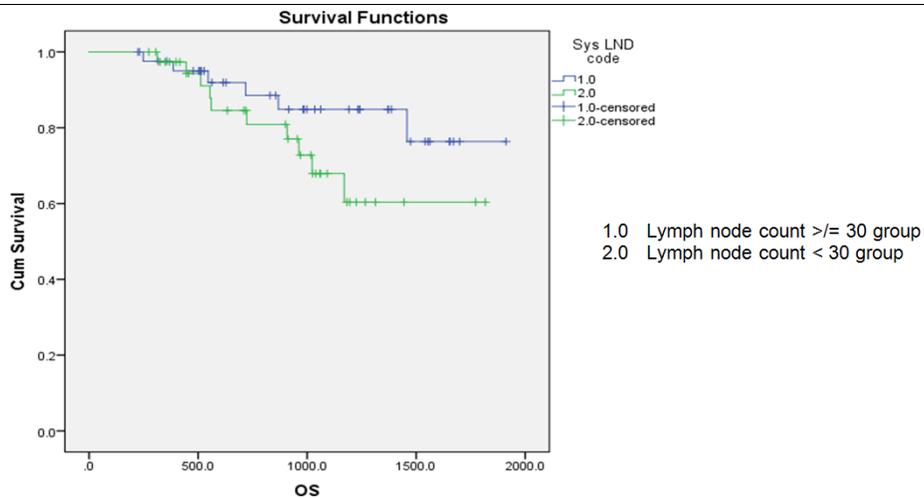
**Table 1. Patient characteristics**

<b>Variable</b>	<b>Number of patients</b>	<b>Percentage (%)</b>
Total	83	100
<b>Age</b>		
< / = 64 yrs.	69	83.1
> 64 yrs.	14	16.9
<b>Menopausal status</b>		
Postmenopausal	52	62.7
Premenopausal	31	27.3
<b>Treatment</b>		
Primary surgery	58	69.9
Neoadjuvant chemotherapy	25	30.1

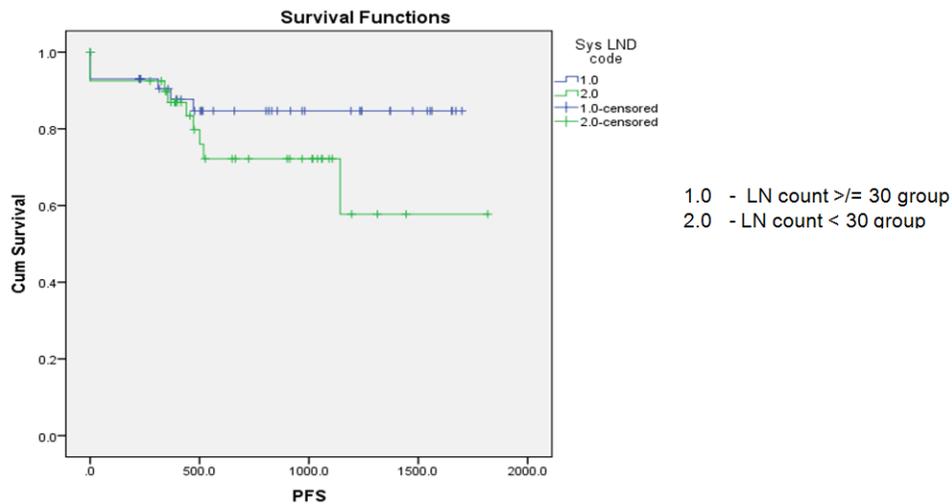
<b>Performance status</b>		
ECOG 0	37	44.6
ECOG 1	40	48.2
ECOG 2	06	7.2
<b>FIGO stage</b>		
I	21	25.2
II	3	3.6
III	51	61.5
IV	8	9.7

**Table 2. Tumour characteristics**

<b>Variable</b>	<b>Number of patients</b>	<b>Percentage (%)</b>
Total	83	100
<b>Grade</b>		
Low	13	15.7
High	70	84.3
<b>Histology</b>		
Serous	68	81.9
Others	15	18.1
<b>pN stage</b>		
Nx	9	10.8
N0	47	56.7
N1	27	32.5
<b>Lymph node resected</b>		
< 30	40	48.2
≥ 30	43	51.8
<b>Residual disease</b>		
Nil	52	62.7
< / =10mm	22	26.5
> 10mm	9	10.8
<b>Surgical complexity score</b>		
Low	9	10.8
Intermediate	68	81.9
High	6	7.3



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



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

**4. DISCUSSION**

Lymphatic spread is common in advanced EOC. 27 (36.5%) patients had positive nodes in our cohort, which is slightly less compared to other studies. This could probably be because of a high number of early-stage EOCs. In the present study 17 / 27 (62.9%) node-positive patients had normal sized non-suspicious nodes on intraoperative assessment, emphasising the inaccuracy of clinical assessment as noted in earlier studies [14].

The role of lymphadenectomy for staging procedure is well established through the therapeutic role is debated [15,16]. The impact of

lymphadenectomy in advanced ovarian cancer is less clear and guidelines are lacking whether systematic pelvic and para-aortic lymphadenectomy be performed with debulking surgery [17,18,19,20]. The decision to perform systematic lymphadenectomy is by the surgeon's discretion or the policy of the hospital.

In advanced ovarian cancer, there is a high rate of involved nodes and survival benefits are documented with systematic aortic and pelvic lymphadenectomy [9]. Studies also have shown that removal of bulky nodes improved OS in optimally debulked ovarian carcinoma and there are studies showing no advantage in resection of clinically negative nodes [17,18,19,20,21]. No

**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
<b>Age (years)</b>							
>64	69	1.56	0.50-0.856	0.43	1.07	0.29-3.93	0.91
<64	14	1	-	-	-	-	-
<b>Performance status</b>							
ECOG 0	37	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	-	-	-
<b>pN stage</b>							
N0	47	1	-	-	-	-	-
N1	27	7.54	2.06-27.51	0.002	2.07	0.24-17.57	0.50
Nx	9	4.72	0.95-23.47	0.06	5.47	0.93-31.78	0.06
<b>Lymph node count</b>							
<30	40	2.13	0.76-5.97	0.14	-	-	-
≥30	43	-	--	-	-	-	-
<b>Residual disease</b>							
1. Nil	52	1	-	-	-	-	-
1-10 mm	22	3.7	1.04-13.1	0.04	1.52	0.35-6069	0.57

> 10mm	9	7.56	2.12-26.83	0.002	3.55	0.77-16.35	0.10
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standardised techniques were mentioned in most of the studies and the number of nodes dissected were also variable.

Kim et al. in 2010, conducted a meta-analysis comparing the impact of systematic lymphadenectomy and non-systematic lymphadenectomy (random removal or less removal or no removal of pelvic and para-aortic nodes). They found an increased OS in all stage disease with systematic lymphadenectomy. But sub-analysis of the two RCTs included in their study showed no difference in OS between systematic and non-systematic lymphadenectomy [9]. In a study by Gao et al., 5-year OS in systematic lymphadenectomy group was higher than the non-systematic lymphadenectomy group. In their analysis of 14 studies, the difference was seen in observational studies and advanced stage disease and no difference was seen in the RCT, early-stage disease and residual disease  $\leq 2$ cm. The definition of unsystematic lymphadenectomy was inconsistent even in this study [21].

In the present study, in 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 was  $< 30$  but was not statistically significant. OS was significantly different between performance status, N stage, systematic lymphadenectomy and residual disease using Kaplan Meier analysis. Subjects with better performance status, negative nodes, who underwent complete systematic lymphadenectomy and nil residual disease, had a higher survival as compared to other groups.

In our study there is not much of a difference between overall and progression-free survival as the number of events has still not happened at the time of analysis and the analysis was time bound.

The lymphadenectomy in ovarian neoplasms (LION) is the only prospective randomised trial which demonstrated no significant benefit of either PFS or OS and has endorsed omitting routine lymphadenectomy in clinically node-negative advanced ovarian cancer patients with macroscopic complete tumour resection [22].

The limitation of the present study is that it is a retrospective study with fewer numbers and hence no subgroup analysis could be done.

We have included cases with varying follow-up periods after completion of treatment. A study with larger numbers and longer follow up would throw better light in this matter.

## 5. CONCLUSION

Systematic lymphadenectomy does not have a significant impact on improving overall or progression-free survival in epithelial ovarian cancers. However, N stage and residual disease had higher hazards of mortality. The complete SLND group showed a trend towards improved OS and PFS, though statistically not significant.

## CONSENT

An informed written consent was obtained from patients who had completed treatment and were available for follow up. Telephonic verbal consent was obtained in women who had not visited the hospital for recent follow-up.

## ETHICAL APPROVAL

The study was approved by the institutional ethical committee.

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## COMPETING INTERESTS

Authors have declared that no competing interests exist.

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