

A Statical Analysis of the Role Various of Co-variables in Detemining, Number of Nodes to be Dissected in Endometrial Cancer by Cart Model, Ahpgic

Jita Parija¹, Bhagyalaxmi Nayak¹, Ashok Padhi¹, Smruti Sudha Pattnaik¹, Manoranjan Mohapatra¹, Janmejaya Mohapatra¹, Rekha Das², Padmalaya Devi³, Sushil Kumar Giri¹, Sagarika Samantray⁴, Lucy Pattnaik⁵, Sumita Mohanty⁶, Swadeep Mohanty³, Sasmita Panda⁵, Niharika Panda¹, Surendra Nath Senapathi⁵

¹Department of Gynaecoconology Oncology, AHPGIC Cuttack Utkal University, India. ²Department of Anesthesiology, AHPGIC Cuttack Utkal University, India. ³Department of Surgical Oncology, AHPGIC Cuttack Utkal University, India. ⁴Department of Pathology, AHPGIC Cuttack Utkal University, India. ⁵Department of Radiation Oncology, AHPGIC Cuttack Utkal University, India. ⁶Department of Oncopathology, AHPGIC Cuttack Utkal University, India.

Abstract

Aim and Objective: The main objective of this paper is to predict the role of covariables in determining the number of nodes to be dissected in endometrial cancer, using the best regression model. Also, to compare the accuracy of the cart model with the traditional regression model, to accurately find and predict the co-variable in determining the number of nodes to be dissected. **Material Method:** Data on 170 endometrial cancer patients with their covariates were collected from the institute ahpgic and used for the said objectives. The data for this paper was collected wherein dependent variable is total number of lymph node involved and 10 co-variates (independent variables) age, Postmenopausal Bleeding, Obstetrics History, Nodal Status, Tumor Size, Histology, Grade, Myometrial Invasion, Lymphovascular Space Invasion and Cervical Extension. **Methods:** multiple regression and cart model. **Results:** Average number of lymph node dissection among patients having tumor size less than 1.9 cm is 3.73 (approx4) and the patients having a tumor size 1.9 cm is 12.4 (approx13). Average nos of nodes dissection among the patients having prior dissected nodal staus as b/l pelvic lymphdenectomy 10.9 (approx 11) and patients having prior dissected b/l paraortic + b/l pelvic lymhadenectomy is 14.1 (approx14). Thus CART MODEL can predict with more accuracy of 95.9% than the multiple regression model which is of 88.3%, based on the selected covariates and validated by receiver operating characteristics (ROC) curve. **Conclusion:** Thus we conclude that if tumor size >1.9cm (approx> 2cm) the 12 nodes and if less than 1.9cm (approx< 2cm) then 3.73 i.e approx 4 nodes should be dissected We found that classification and regression tree (CART) model is able to predict the role of the co variate i.e tumor size in deciding the number of lymph node dissection for the EC patients with an accuracy of 95.9% based on the selected covariates and validated by receiver operating characteristics (ROC) curve. Thus CART MODEL can predict with more accuracy of 95.9% than the multiple regression model which is of 88.3%, based on the selected covariates and validated by receiver operating characteristics (ROC) curve. **Purpose:** The above study will be of help in deciding the nos of nodes to be dissected in endometrial cancer so, that unnecessary morbidities like bleeding and lymphoedem, and waste of resources i.e money and time.

Keywords: CART- Endometrial Cancer- ROC- R-square- Lymph nodes- Accuracy

Corresponding Author:

Dr. Smruti Sudha Pattnaik

Department of Gynaecoconology Oncology, AHPGIC Cuttack Utkal University, India.

Email: Drsmrutisudhapattnaik@Gmail.Com

Introduction

Lymphadenectomy remains controversial for endometrial cancer. Studies favoring omission of para-aortic lymphadenectomy are GOG 33 [1] and a Memorial Sloan Kettering series that reported 1% to 2% rates of isolated positive para-aortic nodes in clinically uterine-confined disease [2, 3]. The prognostic implications of lymph node metastasis merit upstaging from IA-B to stage IIIC. The Benedetti Panici et al [4] and Medical Research Council ASTEC (Efficacy of Systematic Pelvic Lymphadenectomy in Endometrial Cancer) 5 trials of pelvic lymphadenectomy are often quoted as evidence against a therapeutic benefit of lymphadenectomy in endometrial cancer. Both trials evaluated pelvic lymphadenectomy, and 25.2% to 33% of women in the no-lymphadenectomy arms received pelvic radiation [4, 5]. Because the Aalders et al, [6] PORTEC, [7] GOG 99, [8] ASTEC-EN.5,9 and PORTEC-210 trials showed no survival improvement from radiotherapy for early-stage endometrial cancers, using pelvic radiation to sterilize nondissected nodes has decreased [6-10]. Endometrial cancer is now managed with less lymphadenectomy and less pelvic radiation. The critical oncologic question now is: If lymph nodes are less evaluated and treated, how do we know that occult nodal metastasis is not missed and undertreated, and if this leads to decreased survival? A retrospective cohort study of women with node negative, stage I to IIIB endometrial cancer (n = 152,702) identified from the 1998-2011 National Cancer Database. Multivariable Cox proportional hazards regression tested for an association of lymph node count with survival. Restricted mean survival and relative hazard curves were plotted for survival as a function of number of removed lymph nodes. Results Among women with node-negative endometrioid endometrial cancer, for each additional five lymph nodes removed, the hazard for death decreased: stage I, the hazard ratio (HR) was 0.95 (95% CI, 0.93 to 0.97; P < .001). When grouped by grade, each additional five lymph nodes removed was also associated with decreased hazard for death: grade 1, HR was 0.96 (95% CI, 0.93 to 0.99; P = .009); grade 2, HR was 0.91 (95% CI, 0.89 to 0.94; P < .001). Conclusion was that increased lymph node count is associated with a 1% to 14% decreased hazard of death per each additional five lymph nodes removed and a 5% to 20% increased 5-year survival among women with pathologically node-negative endometrioid and serous endometrial cancers [6, 10]. Of 11,443 patients, the median age was 64 years (range, 22-74 years). In all, 78.7% had stage I disease, 10.3% had stage II disease, and 11.0% had stage III disease; 31.5% had grade 1 histology, 40.6% had grade 2 histology, and 24.3% had grade 3 histology. The median number of lymph nodes reported was 9 (range, 1-90 lymph nodes). The median number of lymph nodes and the percent of patients with positive lymph nodes have increased from 1988 to 2001.

An increasing number of lymph nodes removed was associated with a higher likelihood of identifying those with lymph node metastases. Based on the logistic regression model, the largest increase in probability of detecting at least a single positive lymph node was observed when 21 to 25 lymph nodes were resected (odds ratio [OR] of 1.45; 95% confidence interval [95% CI], 1.08-1.94 [P < .01]). Removing greater than 25 lymph nodes did not improve the statistical probability (OR of 1.23; 95% CI, 0.94-1.61 [P = .13]). The current study data suggest that the removal of 21 to 25 lymph nodes significantly increases the probability of detecting at least 1 positive lymph node in endometrioid uterine cancer. The definition of an adequate lymphadenectomy deserves further investigation [7].

The identification of the number of positive lymph node in an early stage may prevent the progress of advanced stage cancer among the cancer patients.

The effective number of lymph node dissection may vary based on the characteristics of different cancers. Early detection and dissection of affected lymph node significantly improves the survival of the patients.

At the same time, it is also important that, unnecessary dissection of the lymph nodes may cause vaginal bleeding, nerve or vessel damage, wound infection, blood clots and damage to nearby tissues. So, it is very important to identify the patients at high risk and to prevent their critical conditions it is also need to extract the required number of lymph nodes.

Therefore, it is very important to identify the patients at high risk and need to extract the required number of lymph nodes. To identify the patients at risk and to prevent their critical conditions, here we proposed a CART model to predict the required number of lymph node dissection using the information on selected co-variables. For this study, we use CART approach to predict the required number of lymph node dissection using the available information on selected co-variables. CART is a non-parametric statistical modelling technique and free from any distributional assumption, which can be used to analyze the data suffering from abnormal distribution or distribution not known. For its simplicity in modelling and interpretation, it has been widely used in Statistics, Health Science, Computer Science and Metrological Science [8]. The method was pioneered by Morgan and Sonquist (1963), later developed by [9] Breiman et al. (1984). CART can be used as an alternative technique as it has several advantages over traditional statistical techniques [10].

Materials and Methods

The data for this paper was collected on The data for this paper was collected wherein dependent variable is total number of lymph node involved and 10 co-variables (were independent variables) age, Postmenopausal Bleeding, Obstetrics History, Nodal Status, Tumor Size, Histology.

Grade, Myometrial Invasion, Lymphovascular Space Invasion and Cervical Extension. 3.2 Validation of models

The validation of a predictive model can be performed by the help of different measures viz. sensitivity, specificity, positive predictive value, negative predictive value and area under ROC curve. Some measures of validating the model are described as follows; (Table 1)

The coefficient of determination, usually denoted as R^2 or r^2 , is the proportion of the variation in the dependent variable that is predictable from the independent variable and can be calculated by using the following expression;

Where, RSS is the residual sum of squares, TSS is the total sum of squares and can be calculated by; (Figure 1).

Inclusion Criteria

Cases who underwent comprehensive surgical staging including lymphdenectomy.

Exclusion Criteria

Cervical cancers and other gynaecological malignancies like ovary.

Methods

Multiple regression and cart model. (regression tree model).

22.A total of 170 endometrial cancer patients has been taken for the said study. The descriptive statistics of the patients under study has shown in Table 2. The mean age of the patients was registered as 56.86 ± 9.01 years with the range 55 (25, 80) years. Majority patients are from the age group ≥ 57 years (54.7%). More patents (nearly 53%) are having tumor size more than or equal to 3 c.m. with the range 7.9 (0.5, 8.4) c.m. The mean tumor size observed as 02.87 ± 1.49 c.m with the range 7.9 (0.5, 8.4). Most of the patients (68.2%) suffered with postmenopausal bleeding less than, 1-year. (approx) with average 381.34 ± 622.18 days and range 2920 (0, 2920) days. The nodal dissection of the patients ranges between (0, 20) with mean number of nodes 09.56 ± 5.68 . Less

than 13 lymph node dissection was done among most of the patients (i.e., 85.9%). Considering obstetrics status, almost 70% of the patients are having one or more children (multipara) and nearly 30% are not having even a single child (nullipara). Most of the patients are with grade-2 (62.4%) and there are 15.3% and 22.3% patients with grade-1 and grade-3 respectively. The histology status shows that the endometrial glands present in nearly 85% and absent in nearly 15% of the patients. More than 80% of the patients are having myometrial invasion less than 50%. Only 13.5% and 10.6% of the patients are having cervical extension and lymphovascular space invasion positive status respectively. Major of the patients are with pelvic nodal status i.e., 57.1%. patients are with grade-2 (62.4%) and there are 15.3% and 22.3% patients with grade-1 and grade-3 respectively. The histology status shows that the endometrial glands present in nearly 85% and absent in nearly 15% of the patients. More than 80% of the patients are having myometrial invasion less than 50%. Only 13.5% and 10.6% of the patients are having cervical extension and lymphovascular space invasion positive status respectively. Major of the patients are with pelvic nodal status i.e., 57.1%. Here, Table 2 shows the results of traditional multiple regression modelling on endometrial cancer patient data to predict the number of lymph nodes of different endometrial cancer patients using different covariates. Two co-variables namely nodal status and tumor size are the significant predictors to predict the number of lymph nodes with $p < 0.05$. The multiple regression model carries a lot of loads so, the assumptions of the model for normality and homoscedasticity can be verified by Shapiro-Wilk test. The Shapiro-Wilk test statistics gives $W = 0.991$ with $p\text{-value} = 0.901$, therefore null hypothesis accepted and the normality assumptions for the multiple regression model is verified (Figure 2, 3 and 4).

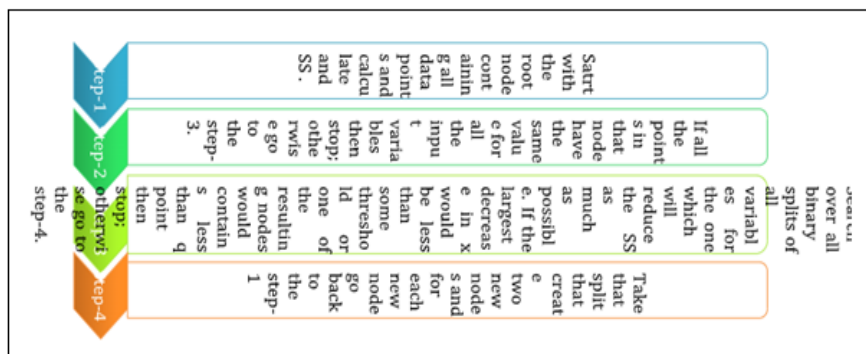


Figure 1. Residual Sum Square (RSS) Total Sum Square (TSS) Regression Tree Splitting Steps

Table 1. Predicted Value

		Predicted value	
		Category-1	Category-2
Actual value	Category-1	a (true positive)	b (false positive)
	Category-2	c (false positive)	d (true negative)

Coefficient of determination (R^2)

Table 2. Descriptive Statistics of 170 (n) Endometrial Cancer patients

Patient's Characteristics	Mean \pm SD	Range (min, max)	
Age (in years)	56.86 \pm 9.01	55 (25, 80)	
Tumor size (in c.m.)	02.87 \pm 1.49	7.9 (0.5, 8.4)	
Postmenopausal Bleeding (in days)	381.34 \pm 622.18	2920 (0, 2920)	
Node dissection (in numbers)	09.56 \pm 5.68	20 (0, 20)	
Co-variate (code)	n_i (%)	Co-variate n_i (%)	
Age		Grade	
< 57 years (0)	77 (45.3)	Grade-1 (1)	26 (15.3)
\geq 57 years (1)	93 (54.7)	Grade-2 (2)	106 (62.4)
Tumor size		Grade-3 (3)	38 (22.3)
< 3 c.m. (0)	90 (52.9)	Lymphovascular Space Invasion	
\geq 3 c.m. (1)	80 (47.1)	Negative (0)	152 (89.4)
Postmenopausal Bleeding		Positive (1)	18 (10.6)
< 381 days (0)	116 (68.2)	Cervical Extension	
\geq 381 days (1)	54 (31.8)	Negative (0)	147 (86.5)
Obstetrics History		Positive (1)	23 (13.5)
Nullipara (0)	52 (30.6)	Nodal Status	
Multipara (1)	118 (69.4)	Bplnd (pelvic) (0)	97 (57.1)
Histology		Bpand (paraaortic)(1)	73 (42.9)
Nonendometrioid (0)	26 (15.3)	Number of nodes dissect	
Endometrioid (1)	144 (84.7)	< 13 (0)	146 (85.9)
Myometrial Invasion		\geq 13 (1)	24 (14.1)
< 50% (0)	138 (81.2)		
\geq 50% (1)	18.8		



Figure 2. Systemic b/l Pelvic and Para Aortic Lymphdenectomy, with Omental Biopsy

Analysis and Result of Regression tree Algorithm

The basic regression tree growing algorithm is as follows:

Table 2 shows a total of 170 endometrial cancer patients has been taken for the said study. The descriptive statistics of the patients under study has shown in Table 2. The mean age of the patients was registered as 56.86 \pm 9.01 years with the range 55 (25, 80) years. Majority patients are from the age group \geq 57 years (54.7%). More patents (nearly 53%) are having tumor size more than or equal to 3 c.m. with the range 7.9 (0.5, 8.4) c.m . The mean tumor size observed as 02.87 \pm 1.49 c.m with the range 7.9 (0.5, 8.4). Most of the patients (68.2%) suffered with postmenopausal bleeding less than,1-year.(approx.) with average 381.34 \pm 622.18 days and range 2920 (0, 2920) days. The nodal dissection of the patients ranges between (0, 20) with mean number of nodes 09.56 \pm 5. 68.Less than 13lymph node dissection was done among most of

the patients (i.e., 85.9%). Considering obstetrics status, almost 70% of the patients are having one or more children (multipara) and nearly 30% are not having even a single child (nullipara). Most of the patients are with grade-2 (62.4%) and there are 15.3% and 22.3% patients with grade-1 and grade-3 respectively. The histology status shows that the endometrial glands present in nearly 85% and absent in nearly 15% of the patients. More than 80% of the patients are having myometrial invasion less than 50%. Only 13.5% and 10.6% of the patients are having cervical extension and lymphovascular space invasion positive status respectively.Major of the patients are with pelvic nodal status i.e., 57.1%. patients are with grade-2 (62.4%) and there are 15.3% and 22.3% patients with grade-1 and grade-3 respectively. The histology status shows that the endometrial glands present in nearly 85% and absent in nearly 15% of the patients. More than 80% of the patients are having myometrial invasion less than 50%. Only 13.5% and 10.6% of the patients are having

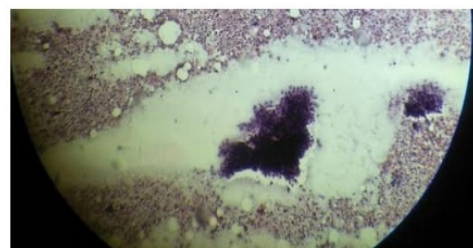


Figure 3. Pelvic Node Positive Adenocarcinima

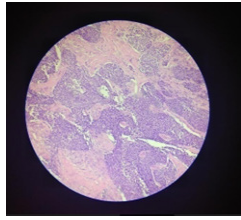


Figure 4. High Grade (3) Carcinoma Endometrium

cervical extension and lymphovascular space invasion positive status respectively. Major of the patients are with pelvic nodal status i.e., 57.1%.

Table 3 shows the results of traditional multiple regression modelling on endometrial cancer patient data to predict the number of lymph nodes of different endometrial cancer patients using different covariates. Two co-variables namely nodal status and tumor size are the significant predictors to predict the number of lymph nodes with $p < 0.05$.

At the beginning, a regression tree holds all the data points in its root node. Further, we need a splitting criterion to split the root node. Starting with covariate tumour size and the cut-point 1.9 cm, all the women with tumour size < 1.9 cm are splitted into the left daughter node and the rest into the right daughter node. A total of 56 women went to the left node and 114 women to the right node.

The choices, tumour size and 1.9 cm are the best cut-point to split the root node.

Figure 5 shows the cart model spiltting and analysis of the tumor size. Average number of lymph node dissection among patients having tumor size less than 1.9 cm is 3.73 (approx 4) and the patients having a tumor size 1.9 cm is 12.4 (approx 13). average nos of nodes dissection among the patients having prior dissected nodal staus as b/l pelvic lymphdenectomy 10.9 (approx 11) and patients having prior dissected b/l paraortic + b/l pelvic lymhadenectomy is 14.1 (approx 14) i.e the covariate tumor size i.e highlighted in green color is significant . The other co vriates are in faint blue and sky color are less significant. if tumor size > 1.9 cm approx 2cm the 13 nodes and if less than < 1.9 cm approx 2cm then 3.73 i.e approx 4 nodes (< 13) should be dissected. The second split is done on the daughter node grade as < 3 and ≥ 3 (terminal

node). Next split is done at the tumour size ≥ 1.9 cm, which divides the information on nodal status as pelvic (terminal node) and paraaortic (terminal node).

Further, patients with grade < 3 (i.e., 1 and 2) are divided into two parts as tumour size ≥ 1.45 cm (terminal node) cm and < 1.45 cm. Patients having tumour size < 1.45 cm are partitioned again by grade ≥ 2 and < 2 (terminal node). Subsequently, the patients with grade ≥ 2 are further split by obstetric status as: nullipara (0: terminal node) and multipara (1). Finally, multipara patients are divided by the last determi- nant age as < 64 (terminal node) and ≥ 64 years (terminal node).

Finally, the splitting of nodes stops after seventh split. At each node the mean number of lymph node dissection and size (%) of the root or leaf node are reported.

This tree has eight terminal nodes. The covariates tumour size, nodal status, grade, and age play an important role to construct the regression tree.

However, the variables such as postmenopausal bleeding, obstetrics history, histology, myometrial invasion, lymph vascular space invasion, cervical extension have no role in the regression tree. The output of CART model demonstrates that there are two significant predic-tors for predicting lymph node dissection and they

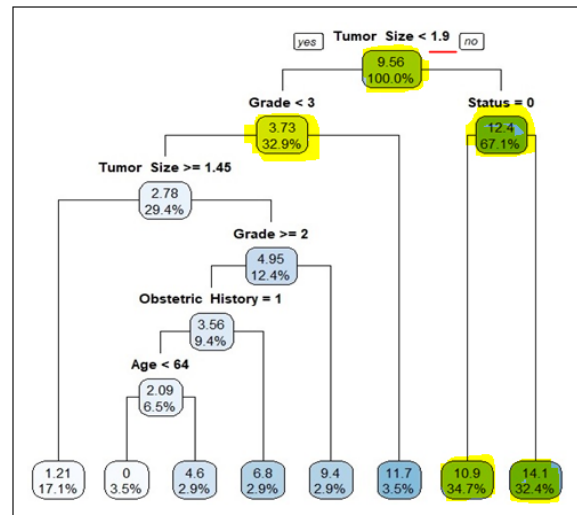


Figure 5. The Cart Model Spiltting and Analysis of the Tumor Size.

Table 3. Results of Multiple Regression Modelling

	Estimate	Std. Error	t-value	Pr(> t)
(Intercept)	1.806	3.091	0.584	0.559
Age	0.033	0.042	0.796	0.427
Postmenopausal Bleeding	-0.001	0.001	-0.039	0.969
Obstetric History	-0.435	0.861	-0.506	0.6139
Nodal status	2.312	0.752	3.074	0.002
Tumor Size	1.519	0.263	5.769	<0.001
Histology	-1.18	1.225	-0.963	0.337
Grade	0.729	0.676	1.078	0.282
Myometrial Invasion	1.202	1.009	1.191	0.235
Lymph vascular Space Invasion	-0.717	1.397	-0.513	0.608
Cervical Extension	0.912	1.192	0.766	0.445

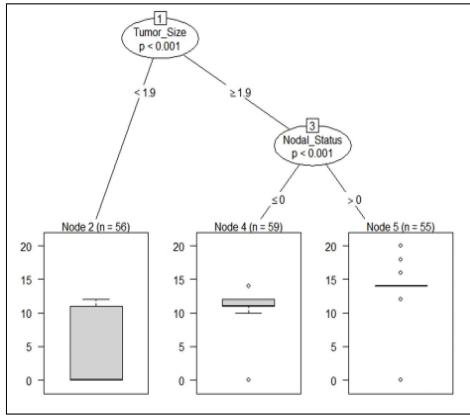


Figure 6. Modified CART Splitting with their Determinant i.e tumor size and nodal staus (i.e pelvic lymphadenectomy/ para-aortic lymphadenectomy)

are tumour size and nodal status.

Modified CART splitting with their determinants. The outcomes of the model states that the patients with tumour size ≥ 1.9 cm are at higher risk of extracting additional lymph node compared to the patients having tumour size < 1.9 cm. The patients with paraaortic +PELVIC lymhadenectomyare at High risk of additional lymph node compared with the patients having Pelvic lymphadenectomy. Thus, it is more important to dissect an additional lymph node of the patients having tumour size ≥ 1.9 cm and paraaortic nodal status (Figure 6).

We have checked the model adequacy by plotting the residuals of both multiple and CART model (not shown here). It was also verified that the observed data coin-cide with reference line passing through origin, hence validating the assumption of normality of the error term in the model (Figure 7).

It is clear that, the age predictor of the patients is

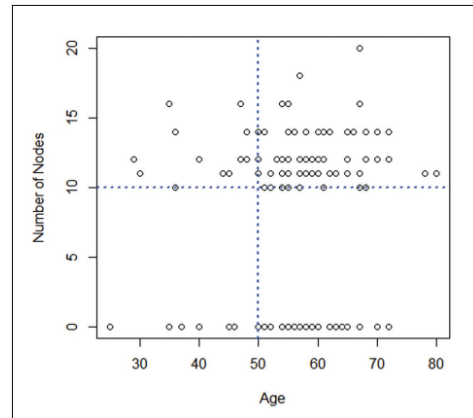


Figure 7. Scatter Plot of Age vs node

explaining the high and low risk of extraction of lymph node. Hence, the patients under study can be classified into two major groups based on their ages i.e., < 50 and ≥ 50 .

The outcomes of the model states that the patients with tumour size ≥ 1.9 cm are at higher risk of extracting additional lymph node compared to the patients having tumour size < 1.9 cm. Similarly, the patients with nodal status as paraaortic are at high risk of extracting additional lymph node compared with the patients having pelvic nodal status. Thus, it is more important to dissect an additional lymph node of the patients having tumour size ≥ 1.9 cm and paraaortic nodal status.

We have checked the model adequacy by plotting the residuals of both multiple and CART model (not shown here) shows number of lymph nodes is mostly scattered in the upper part of the blue horizontal line, indicating that the majority of patients have ≥ 10 nodes. Further, the upper data is more scattered towards the right-hand side of the blue vertical line, indicating the major patients those

Table 4 a. The Maximum Index Youden Value was 13 which was Associated Withbtst Variable i.e Age

Positive if Greater than or Equal To	Sensitivity	1-Specificity	Sensitivity + Specificity-1
-1	1	1	0
5	0.75	0.759	-0.009
10.5	0.75	0.684	0.066
11.5	0.5	0.481	0.019
13	0.417	0.335	0.082
15	0	0.044	-0.044
17	0	0.013	-0.013
19	0	0.006	-0.006
21	0	0	0

Measurement of Youden Index Test Variable, Number of Nodes Dissect State Variable, Age and Value of State Variable, 50

Table 4 b. Cross Tabulation of Predicted vs Actual Value for Both Multipleregression & CART Model for Two Groups i.e < 13 Nodes and ≥ 13 nodes

Actual	Multiple regression		Actual	Cart	
	Predictions			Predictions	
	< 13	≥ 13		< 13	≥ 13
< 13	106	6	< 13	110	2
≥ 13	40	18	≥ 13	5	53

≥ 10 lymph nodes dissected are belongs to age ≥ 50 . It is clear that, the age predictor of the patients is explaining the high and low risk of extraction of lymph node. Hence, the patients under study can be classified into two major groups based on their ages i.e., < 50 and ≥ 50 . Nos of nodes as 13 as an optimal threshold value for the dependent variable, and it can be stratified as < 13 and ≥ 13 two groups, which can be used further for validating and comparing the predictive models.

Cross tabulation of predicted vs actual value for both multiple regression & CART model for two groups i.e. < 13 nodes and ≥ 13 nodes (Table 4 a and b).

Comparison of ROC Curve of Both the Models

Therefore, we can take 13 as an optimal threshold value for the dependent variable, and it can be stratified as in order to compare the predictive power of the CART model with the multiple regression model, we draw two ROC curves shown in Figure 8. The area under the ROC curve (AUC) in 0.965 in the CART model and 0.883 in the multiple regression model. This, shows, the predictive power of the co variable i.e tumor size 1.9 cm (approx 2cm), is expected to be more sensitive and specific in the CART model than the traditional multiple regression model. Table 5 shows the value of different parameters for comparing multiple regression model and CART model. Sensitivity, specificity, PPV, NPV and accuracy are 0.726, 0.750, 0.946, 0.310 and 0.729, respectively for multiple regression model and 0.956, 0.963, 0.982, 0.913 and 0.959, respectively for the CART model. The accuracy of CART model over multiple regression model shows that CART can be considered as a better alternative to multiple regression model particularly for analysing EC data.

Discussion

From the data average number nodes becomes 9.56 (approx.10). There are two significant predictors that can be used to decide to the nos of nodes to be dissected by multiple regression model. However we found that classification and regression tree (CART) model is able to predict the role of the co variate i.e tumor size in deciding the number of lymph node dissection. Average number of lymph node dissection among patients having tumor size less than 1.9 cm is 3.73 (approx 4) and the patients having a tumor size 1.9 cm is 12.4 (approx 13). Average nos of nodes dissection among the patients having prior dissected nodal status as b/l pelvic lymphadenectomy 10.9 (approx 11) and patients having prior dissected b/l paraortic + b/l pelvic lymphadenectomy is 14.1 (approx 14). Thus CART MODEL can predict with more accuracy of 95.9% than the multiple regression model which is of 88.3%, based on the selected covariates and validated by receiver operating characteristics (ROC) curve.

Results in Context with Published Literature

A retrospective cohort study of women with node-negative, stage I to IIIB endometrial cancer (n = 152,702) identified from the 1998-2011 National Cancer Database. Multivariable Cox proportional hazards regression tested for an association of lymph node count with survival. Conclusion was that increased lymph node count is associated with a 1% to 14% decreased hazard of death per each additional five lymph nodes removed and a 5% to 20% increased 5-year survival among women with pathologically node-negative endometrioid and serous endometrial cancers [6]. CART is a non-parametric statistical modelling technique and free from any distributional assumption, which can be used to analyze

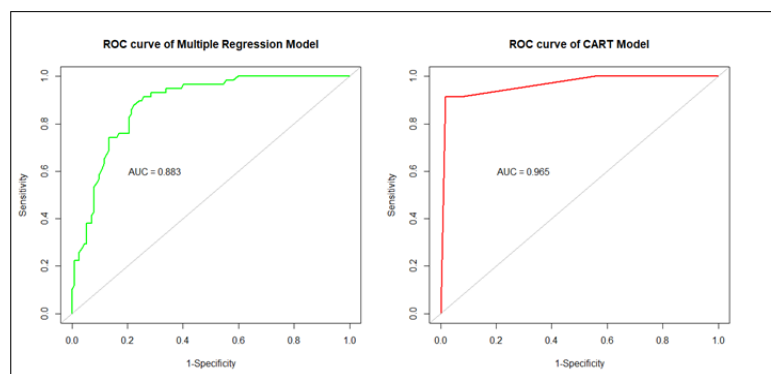


Figure 8. Comparison of ROC of Multiple Regression and Cart Model

Table 5. The Value of Different Parameters for Comparing Multiple Regression Model and CART Model

Parameters	Multiple Regression Model	Cart Model
Area under the curve (auc)	0.883	0.965
Sensitivity	0.726	0.956
Specificity	0.75	0.963
Positive predictive value	0.946	0.982
Negative predictive value	0.31	0.913
Accuracy	0.729	0.959
R ²	0.334	0.754

the data suffering from abnormal distribution or distribution not known. For its simplicity in modelling and interpretation, it has been widely used in Statistics, Health Science, Computer Science and Metrological Science [8]. The method was pioneered by Morgan and Sonquist (1963), later developed by [9] Breiman et al. (1984). CART can be used as an alternative technique as it has several advantages over traditional statistical techniques [10].

Strength and Weakness

It is found that, CART model is able to predict the number of lymph node dissection of endometrial cancer patients with an accuracy of 95.9% based on selected variables and validated using ROC curve with the area 0.965. It is also found that, CART model has the potential advancement over traditional regression model and can be used as its alternative method. The limitation of this method is that, it only considers a covariate or multiple covariates at a time and ignore the level of the co-variables. the limitation of this method is that, it only considers a covariate or multiple covariates at a time and ignore the level of the co-variables. To overcome these limitations, here we develop a regression tree.

Implication of Application in Future Research

Researchers are looking ways to estimate the number of lymph nodes for different cancer patients using some measurements such as age, sex, co-morbidities etc. The covariates tumor size and early detected nodal status from lymph node sampling are found as two significant predictors to decide the number of lymph node need to be dissected. This cart model will help us in predicting with accuracy the number of nodes to be considered dissection, so that unnecessary morbidities and mortalities can be avoided in endometrial cancer.

In conclusion, the covariates tumor size and early detected nodal status from lymph node sampling are found as two significant predictors to decide the number of lymph node need to dissecting order to avoid the critical conditions and to take appropriate remedies or treatment at an early stage to optimize their loss in terms of time and money and life by the multiple regression model.

We found that classification and regression tree (CART) model is able to predict the role of the co variate i.e tumor size in deciding the number of lymph node dissection for the EC patients with an accuracy of 95.9% based on the selected covariates and validated by receiver operating characteristics (ROC) curve. Thus CART MODEL can predict with more accuracy of 95.9% than the multiple regression model which is of 88.3%, based on the selected covariates and validated by receiver operating characteristics (ROC) curve. Thus we conclude that if tumor size >1.9cm (approx > 2cm) the 13 nodes and if less than 1.9cm (approx < 2cm) then 3.73 i.e approx 4 nodes i.e <13 should be dissected.

We demonstrated the demographic, clinical and pathological covariates that has facilitated insight into the prediction of lymph node dissection using CART m.

Purpose

The above study will be of help in deciding the nos of nodes to be dissected in endometrial cancer so, that unnecessary morbidities like bleeding and lymphoedem, and waste of resources i.e money and time.

Software used for data analysis and plotting

All the statistical analysis and plotting were done by using R (version 3.6.2) and SPSS (version 20).

References

1. Rungruang B, Olawaiye AB. Comprehensive surgical staging for endometrial cancer. *Reviews in Obstetrics & Gynecology*. 2012;5(1):28-34.
2. Creasman WT, Morrow CP, Bundy BN, Homesley HD, Graham JE, Heller PB. Surgical pathologic spread patterns of endometrial cancer. A Gynecologic Oncology Group Study. *Cancer*. 1987 Oct 15;60(8 Suppl):2035-2041. [https://doi.org/10.1002/1097-0142\(19901015\)60:8+<2035::aid-cncr2820601515>3.0.co;2-8](https://doi.org/10.1002/1097-0142(19901015)60:8+<2035::aid-cncr2820601515>3.0.co;2-8)
3. Abu-Rustum NR, Gomez JD, Alektiar KM, Soslow RA, Hensley ML, Leitao MM, Gardner GJ, et al. The incidence of isolated paraaortic nodal metastasis in surgically staged endometrial cancer patients with negative pelvic lymph nodes. *Gynecologic Oncology*. 2009 Nov;115(2):236-238. <https://doi.org/10.1016/j.ygyno.2009.07.016>
4. Benedetti Panici P, Basile S, Maneschi F, Alberto Lissoni A, Signorelli M, Scambia G, Angioli R, et al. Systematic pelvic lymphadenectomy vs. no lymphadenectomy in early-stage endometrial carcinoma: randomized clinical trial. *Journal of the National Cancer Institute*. 2008 Dec 03;100(23):1707-1716. <https://doi.org/10.1093/jnci/djn397>
5. Kitchener H, Swart AM, Qian Q, Amos C, Parmar MK. Efficacy of systematic pelvic lymphadenectomy in endometrial cancer (MRC ASTEC trial): a randomised study. *Lancet (London, England)*. 2009 Oct 01;373(9658). [https://doi.org/10.1016/S0140-6736\(08\)61766-3](https://doi.org/10.1016/S0140-6736(08)61766-3)
6. Seagle BL, Gilchrist-Scott D, Graves S, Strohl AE, Nieves-Neira W, Shahabi S. Association of Lymph Node Count and Overall Survival in Node-Negative Endometrial Cancers. *JCO clinical cancer informatics*. 2017;1. <https://doi.org/10.1200/CCI.16.00064>
7. Chan JK, Urban R, Cheung MK, Shin JY, Husain A, Teng NN, Berek JS, et al. Lymphadenectomy in endometrioid uterine cancer staging: how many lymph nodes are enough? A study of 11,443 patients. *Cancer*. 2007 06 15;109(12). <https://doi.org/10.1002/cncr.22727>
8. Faraway JJ. *Extending the linear model with R: generalized linear, mixed effects and nonparametric regression models*. Chapman and Hall/CRC. 2016.
9. Breiman L, Friedman JH, Olshen RA, Stone CJ. *Classification and regression trees*. Boca Raton, FL: Chapman & Hall. 1984;.
10. Clarke AE, Bloch DA, Danoff DS, Esdaile JM. Decreasing costs and improving outcomes in systemic lupus erythematosus: using regression trees to develop health policy. *The Journal of Rheumatology*. 1994 Dec;21(12):2246-2253.



This work is licensed under a Creative Commons Attribution-Non Commercial 4.0 International License.