

## Clinicopathologic Features and Prognostic Factors in Male Breast Cancer A Single Centre Experience



İzzet Doğan<sup>1</sup>, Esra Aydın<sup>1</sup>, Naziye Ak<sup>1</sup>, Kamuran İbiş<sup>2</sup>, Özge Şükrüoğlu<sup>3</sup>, Seda Kılıç Erciyas<sup>3</sup>, Hülya Yazıcı<sup>3</sup>, Adnan Aydıner<sup>1</sup>, Pınar Saip<sup>1</sup>

<sup>1</sup>Istanbul University Institute of Oncology, Department of Medical Oncology <sup>2</sup>Istanbul University Institute of Oncology, Department of Radiation Oncology <sup>3</sup>Istanbul University Institute of Oncology, Department of Cancer Genetics

**Objectives:** To evaluate clinicopathologic features and prognostic factors in male breast cancer (MBC).

**Methods:** The data of 80 patients were retrospectively reviewed. The clinicopathological features, receptor status (ER, PR, and Her2), BRCA mutation status, tumor stage, and treatment data were recorded. Kaplan Meier method and Cox regression analysis were used for overall survival (OS) analysis.

**Results:** The median follow-up period was 47 months (range: 2.2-214.6 months). The median age at diagnosis was 61 years (range: 25-86 years). BRCA mutations were detected in 31% of the patients (Table-1). The most common histological type of tumor was invasive ductal carcinoma (84%). The ratio of estrogen and progesterone receptor positivity was 93.6% and 74.4%, respectively. Her2 overexpression was present in 16.9%. Locoregional recurrence and distant metastases were observed in 1 (1.4%), and 21 (28.4%) patients, respectively. The median OS was  $120.9 \pm 25.8$  months (70.3-171.5) (Figure-1). The five years OS was  $74.9 \pm 6\%$ . BRCA mutation status did not affect OS in univariate analysis (p=0.501). Smoking (p=0.037) and CA15-3 levels (p=0.033) at diagnosis were significantly associated with OS in univariate analysis; however, it was not confirmed by multivariate analysis (Table-2).

**Conclusion:** In this study, we showed to BRCA mutations, smoking, regular alcohol consumption, body mass index (BMI), and family history of breast cancer were not affect on the OS.

	Number of Patients (%)		
Age at diagnosis, years	N:80		
≤ 50	13 (16.3)		
50-65	38 (47.5)		
≥ 65	29 (36.2)		
BRCA mutation status	N:42		
No mutation	29 (69)		
BRCA mutation-positive	13 (31)		
BRCA1 mutation	4 (9.5)		
BRCA2 mutation	8 (19)		
BRCA1 and 2 mutation	1(2.5)		
Family history of breast cancer	N:65		
Yes	19 (29.2)		
No	46 (70.8)		
Body Mass Index (BMI) kg/m <sup>2</sup>	N:50		
<25	12 (24)		
25-30	25 (50)		
≥30	13 (26)		
Smoking status	N:64		
Current	39 (60.9)		
Never	25 (39.1)		
Regular alcohol consumption	N:60		
Yes	16 (26.7)		
No	44 (73.3)		
Stage at diagnosis	N:75		
Stage 1	20 (26.6)		
Stage 2	26 (34.7)		
Stage 3	23 (30.7)		
Stage 4	6 (8)		
pT status	N:68		
pT1-pT2	53 (78)		
pT3-pT4	15 (22)		
pN status	N:69		
Node negative	31 (44.9)		
Node positive	38 (55.1)		
pN1	22 (31.9)		
pN2	8 (11.6)		
pN3	8 (11.6)		
Grade	N:63		
1	2 (3.2)		
2	26 (41.3)		
3	35 (55.5)		

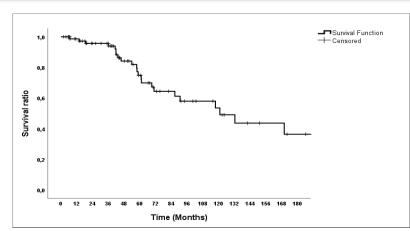


Table 2. Univariate and multivariate analysis for overall survival				
	Univariate	Multivariate analysis		
	analysis			
	P-value	P-value	HR	
Age			(95% CI)	
(<65 vs. ≥65)	0.199			
BRCA mutation status	0.501			
(yes vs. no)				
Family history of breast cancer	0.557			
(yes vs. no)				
Smoking status	0.037	0.098	2.89	
(never vs. current)			(0.82-10.18)	
Regular alcohol consumption	0.963			
(yes vs. no)				
Body Mass Index (BMI) kg/m <sup>2</sup>	0.643			
(obese vs. non-obese)				
CA15-3 levels at diagnosis	0.033			
(high vs. normal)				
Stage at diagnosis	0.002	0.004	11.13	
(stage 1-2-3 vs. stage 4)			(2.14-57.89)	