

# **Primary Tumor Extirpation** in the Setting of Stage IV Breast Cancer

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### Background

- Surgical management of patients stage IV breast cancer is controve
- Existing studies in Stage IV breas cancer have not closely evaluated role of patient response to inducti systemic therapy (IST) in its relationship to survival outcomes.

#### Objective

Evaluate our recent experience wi patients undergoing resection of the primary tumor in the setting of metastatic breast cancer.

#### **Methods**

- Institutional database reviewed from 2008-2018 to identify patients with diagnosis of de novo stage IV brea cancer defined as metastasis diagnosed within 4 months of breat cancer diagnosis.
- Patients were included if they underwent primary tumor extirpati (lumpectomy, mastectomy).
- Patients were grouped according their response in the primary disea site to IST into progression (progressive primary disease on I or no progression (Response to IS nonprogressive primary, comprisi complete, partial and stable response).

	Mayo	o Clinic, R
	Results	Table 1
with ersial. st d the ion	<ul> <li>45 patients were included.</li> <li>Median age was 55 years, 4 patients had a diagnosis of inflammatory breast cancer.</li> <li>Operations were wide local excision (n=11), total mastectomy (n=29), skinsparing (n=4) or nipple-sparing mastectomy (n=1).</li> <li>Demographics and tumor characteristics were similar between the two groups, despite slightly higher percentage of T4 and N2-3 disease and ≥5 metastatic sites in the progressive primary disease compared to the nonprogressive primary</li> </ul>	Table 1No. of PatDistant DiResponseSystemicComplePartialStableProgresMedian SeMedian SeS-year Su5-year SuMedian DiDisease-FSurvival, r
	<ul> <li>group (p&gt;0.05). Patient characteristics and survival data are shown in Table 1, and Figures 1,2.</li> <li>Predictors of overall survival are summarized in Table 2.</li> </ul>	2-year Dis Disease-F Survival 5-year Dis Disease-F Survival
om h a east	Figure 1.	Alive
ast	Lopapility 1.0 – 1.0 – 1.0	1.0 <u>.</u> 같
ion	filipting $1.0 - $	Overall Survival Probability 8.0 8.0 8.0 8.0 8.0 8.0 8.0 8.0
to ase	0.4 – 0.2 –	4.0 arkiv erall Surviv 0.2
ST) ST/ ng	Bood trepting       0.0       Response to IST         Progression on IST       0         0       5       10       15       20       25       30       35	ð 0.0

Months Since Surgery

## Malke Asaad, MD<sup>1</sup>, Jennifer A. Yonkus, MD<sup>2</sup>, Tanya L. Hoskin, MS<sup>3</sup>, Tina J. Hieken, MD<sup>2</sup>, James W. Jakub, MD<sup>2</sup>, Jennifer Racz, MD<sup>2</sup>, Judy C. Boughey, MD<sup>2</sup>, Amy C. Degnim, MD<sup>2</sup>

	Overall N(%)	Progression N(%)	No Progression N(%)	p-value		
Patients	45	13(29)	29(64)			
nt Disease onse to mic Therapy						
nplete	16(38)	3(23)	13(45)	0.013*		
tial	14(33)	4(31)	10(34)			
ble	6(14)	1(8)	5(17)			
gression	4(10)	4(31)	0(0)			
n Survival, ns	117	117	NA	0.378		
<sup>·</sup> Survival	90%	83%	92%			
<sup>·</sup> Survival	76%	71%	81%			
n Distant se-Free /al, months	31	11	50	0.028*		
<sup>·</sup> Distant se-Free /al	53%	22%	62%			
<sup>·</sup> Distant se-Free /al	39%	NA	45%			
	33(73)	4(31)	7(24)	0.713		

#### . Patient Characteristics and Survival Analysis

Figure 2.

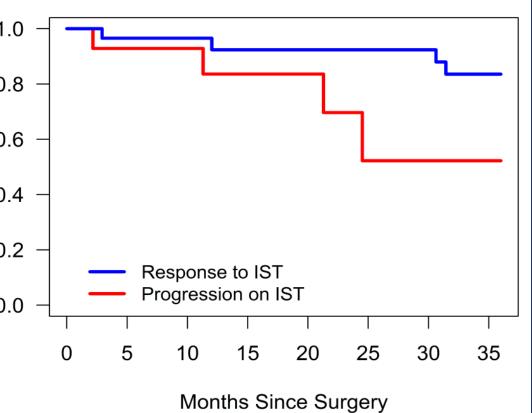


Table 2. Predictors of Decreased Overall Survival					
Predictor	Hazard Ratio (95% CI)	p-value			
Age ≥55 vs. <55	3.9 (1.1, 20.7)	0.03			
Hormone Receptor: positive vs. negative	0.4 (0.1, 1.3)	0.10			
Her-2 Receptor Status: positive vs. negative	0.3 (0.03, 1.1)	0.08			
No. of metastasis: 1 vs. >1	2.0 (0.6, 7.0)	0.23			
Metastatic site: bone vs. other	0.8 (0.3, 2.6)	0.76			
Clinical T3/T4 vs. T1/T2	2.5 (0.8, 8.0)	0.12			
Clinical N2/N3 vs. N1	7.5 (2.0, 26.9)	0.005			
Primary Disease Response to Systemic Therapy	0.3 (0.1, 1.2)	0.08			
Distant Disease Response to Systemic Therapy	0.4 (0.1, 1.6)	0.21			
Closest Margin, per 1 cm	1.3 (0.6, 2.6)	0.45			
Pathologic Tumor Size, per 1 cm increase	1.4 (1.2, 1.6)	<0.001			
Number of Positive Nodes, per 1 node increase	1.1 (1.05, 1.16)	<0.001			

#### **Conclusions**

Surgery is reasonable to consider in stage IV breast cancer patients with good performance status, low disease burden and good response to systemic therapy.

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