

#### **ESMO** Preceptorship Programme

Breast Cancer – Lisboa, Postugal – 16 - 17 Setember 2016

European Society for Medical Oncology

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# Long Response in Cutaneous Metastasis of Breast Cancer



- 70 y.o. Woman
- ECOG PS 0
- Past medical history: Dyslipidemia; Peripheral angiopathy
- Drug history: warfine, sinvastatin, gabapentin, tramadol, metoclopramide
- No history of drug / alcohol intake
- Gynecological history:

Menarche at age 14, menopause at age 50, G3P2, 1 expontaneous abortion, Hormonal contraception for 20 years, hormonal therapeutic substitution for 5 y.

• Palpable breast Lump  $\rightarrow$  Mammography:

**Right Breast Nodule 5.5cm Bi-Rads 5** 

11.2005



	Right Modified Radical Mastectomy	11.2005
$\odot$	Ductal Invasive Carcinoma pT3N3M0.	
	ER-pos, PR- pos, HER2-pos – Intrinsic subtype Luminal B-like	
	Adjuvante ChT + RT + HT	4. 2006
	<ul> <li>FAC 6 cycles with good tolerance. Doxorrubicin cumulative dose 200mg/m2</li> </ul>	
	<ul> <li>RT right chest wall and lymph node areas, total 50 Gy</li> </ul>	10.2006
	<ul> <li>HT with Anastrozole</li> </ul>	
~	Cutono au rolance	DFS 20 m
۲	Cutaneous relapse	
•	Cutaneous relapse 1 <sup>st</sup> Line Paliative ChT with Capecitabina + Trastuzumab	DFS 20 m 07.2007
•	-	
•	1 <sup>st</sup> Line Paliative ChT with <b>Capecitabina + Trastuzumab</b>	
	1 <sup>st</sup> Line Paliative ChT with <b>Capecitabina + Trastuzumab</b> 16 cycles; Best response: Partial Response	07.2007
	1 <sup>st</sup> Line Paliative ChT with <b>Capecitabina + Trastuzumab</b> 16 cycles; Best response: Partial Response	07.2007 05.2008
•	1 <sup>st</sup> Line Paliative ChT with <b>Capecitabina + Trastuzumab</b> 16 cycles; Best response: Partial Response For exhaust ChT suspends capecitabine, keeps <b>Trastuzumab and starts Letrozole</b> .	07.2007 05.2008



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#### Long Response in Cutaneous Metastasis of Breast Cancer

#### **Case Overview**

۲	Cutaneous Progression		
	3 <sup>rd</sup> Line Paliative ChT with Vinorelbin + Trastuzumab		06.2010
	<ul> <li>Vascular leg ulcer – contraindication to keep vinorelbine</li> </ul>		
	<ul> <li>Best response: Parcial Response</li> </ul>		
	Started Letrozol + Trastuzumab	PFS 6 m	03.2011
	Slight cutaneous progression $\rightarrow$ Changes AI for <b>Exemestan + Trastuzum</b>	nab	09.2011
۲	Cutaneous Progression	PFS 5 m	
	4rd Line Paliative ChT with Paclitaxel (weekly) + Trastuzumab		02.2012
	<ul> <li>Best Response: Parcial Response</li> </ul>		
	<ul> <li>Peripheral neuropathy Grade 2 + Exhaust Treatment</li> </ul>		
	Fulvestran + Trastuzumab	PFS 12 m	12.2012
۲	Cutaneous Progression		
	<ul> <li>Inclusion in TDM-1 clinical trial (TDM1 arm)</li> </ul>		12.2013
	<ul> <li>Abandoned Clinical Trial for failure to timely treatment, in management for</li> </ul>	ollowing	
	excessive anticoagulation with ear bleeding.		01.2015
	<ul> <li>Best Response: complete response</li> </ul>		



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ST PRACTICE

- Inclusion in TDM-1 clinical trial (TDM1 arm)
  - Best Response: complete response



Figure 1. End of TDM1 Treatment



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03.2015

**PFS 12 m** 

Tamoxifen + Trastuzumab

• Cutaneous Recorrence - Ulcerative lesion with satellite skin nodules 03.2016



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• Cutaneous Recorrence

03.2016





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۲	Cutaneous Recorrence - ulverative lesion with satellite skin nodules	03.2016
	Brachitherapy	
	Restarted TDM1	07.2016
	Last medical visit – ECOG 1. Complete woud healing.	

→ Clear benefit of adding trastuzumab even after progression in HER2-positive breast cancer.

→ After multiple chemotherapy lines TDM1 represents addicional benefit for this patient, who has already achieved an OS of more than 10 years.

→ Particularly in the case of cutaneous metastasis, only with prompt recognition of progression, with a close surveillance, comes the opportunity to treat the progression of disease, to improve survival rates.



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