

Rare tumours of the breast

Emanuela Esposito^{1,2}, Michelino De Laurentiis¹

¹Department of Breast and Thoracic Oncology, Istituto Nazionale Tumori, IRCCS-Fondazione G. Pascale, Naples, Italy; ²Department of Clinical Medicine and Surgery, University of Naples Federico II, Naples, Italy

Correspondence to: Emanuela Esposito, MD, FEBS. Department of Breast and Thoracic Oncology, Istituto Nazionale Tumori, IRCCS-Fondazione G. Pascale, Via Mariano Semmola, 53 Naples, Italy. Email: emanuelaexpo@hotmail.it.

Submitted Sep 29, 2018. Accepted for publication Oct 08, 2019. doi: 10.21037/tcr.2019.10.05 View this article at: http://dx.doi.org/10.21037/tcr.2019.10.05

Breast cancer is the most commonly occurring cancer in women. One in eight woman is diagnosed with breast cancer and over two million new cases have been registered in 2018 (1). About 90% of breast cancers are classified into classic type and certain therapies are very well described and straightened by randomised controlled trials (2). Namely, classic infiltrating ductal carcinoma (IDC) is the most common type of breast cancer, accounting for 80% of breast cancer diagnoses (2). Second in incidence is infiltrating lobular carcinoma (ILC) and accounts for 10% to 15% of breast cancers. Less than 10% of breast cancers are defined as "rare" and there is a paucity of literature on this topic. Neither randomised trials nor consensus on the optimal treatment of rare breast cancers is available. Data derive from case reports, national registries and few case series.

Rare tumours of the breast can be divided into epithelial and mesenchymal (3). Rare epithelial breast cancers are a heterogeneous group of malignancies with different behaviours and prognoses. Since the early 2000s, the discovery that breast cancers can be classified into specific molecular subtypes based on their global gene expression profiles has deeply shaped the current understanding of inter-tumor heterogeneity (4). Intrinsic classification of breast cancer into five subtypes, which influences the choice of adjuvant treatment, is defined based on genetic research of common ductal breast cancer, without the inclusion of rare histological types without knowledge of rare histological types, some prognostically favorable rare breast cancers may be erroneously classified among the group of prognostically unfavorable tumors due to their immunohistochemical characteristics, which may result in unnecessary aggressive adjuvant treatment (5,6).

Management of "rare" breast cancer is multidisciplinary; it includes locoregional (surgery and radiation therapy) and systemic therapy approaches.

Very little is known about specific conditions such as male breast cancer, breast cancer in pregnancy, inflammatory breast cancer or even about rare site of breast cancer metastases. The manuscripts included in this "Rare Tumours of the Breast" focussed issue of *Translational Cancer Research* evaluate the evidence for rare breast cancers, is aimed to clarify difficult aspects on the management of these disease and describe how to switch from bench to bedside through translational research to achieve the best treatment for patient diagnosed with a "rare" tumour of the breast.

Acknowledgments

Funding: None.

Footnote

Provenance and Peer Review: This article was commissioned by the editorial office for the focused issue "Rare Tumors of the Breast" published in *Translational Cancer Research*. This article did not undergo external peer review.

Conflicts of Interest: Both authors have completed the ICMJE uniform disclosure form (available at http://dx.doi. org/10.21037/tcr.2019.10.05). The focused issue "Rare Tumors of the Breast" was commissioned by the editorial office without any funding or sponsorship. EE and MDL served as the unpaid Guest Editors for the focused issue.

Esposito and Laurentiis. Rare tumours of the breast

EE also serves as the unpaid editorial board member of *Translational Cancer Research* from May 2018 to Apr 2020.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

Open Access Statement: This is an Open Access article distributed in accordance with the Creative Commons Attribution-NonCommercial-NoDerivs 4.0 International License (CC BY-NC-ND 4.0), which permits the non-commercial replication and distribution of the article with the strict proviso that no changes or edits are made and the original work is properly cited (including links to both the formal publication through the relevant DOI and the license). See: https://creativecommons.org/licenses/by-nc-nd/4.0/.

Cite this article as: Esposito E, De Laurentiis M. Rare tumours of the breast. Transl Cancer Res 2019;8(Suppl 5):S443-S444. doi: 10.21037/tcr.2019.10.05

References

- 1. Available online: https://www.wcrf.org/dietandcancer/ cancer-trends/breast-cancer-statistics
- 2. Sharma GN, Dave R, Sanadya J, et al. Various types and management of breast cancer: an overview. J Adv Pharm Technol Res 2010;1:109-26.
- Acevedo C, Amaya C, López-Guerra JL. Rare breast tumors: Review of the literature. Rep Pract Oncol Radiother 2013;19:267-74.
- 4. Skibinski A, Kuperwasser C. The origin of breast tumor heterogeneity. Oncogene 2015;34:5309-16.
- 5. Perou CM, Sørile T, Eisen MB, et al. Molecular portraits of human breast tumours. Nature 2000;406:747-52.
- 6. Guteša I, Roth A, Milas I, et al. Mirko Gulan and Mladen Stanec Rare Breast Tumors. Libri Oncol 2014;42:75-81.

S444