

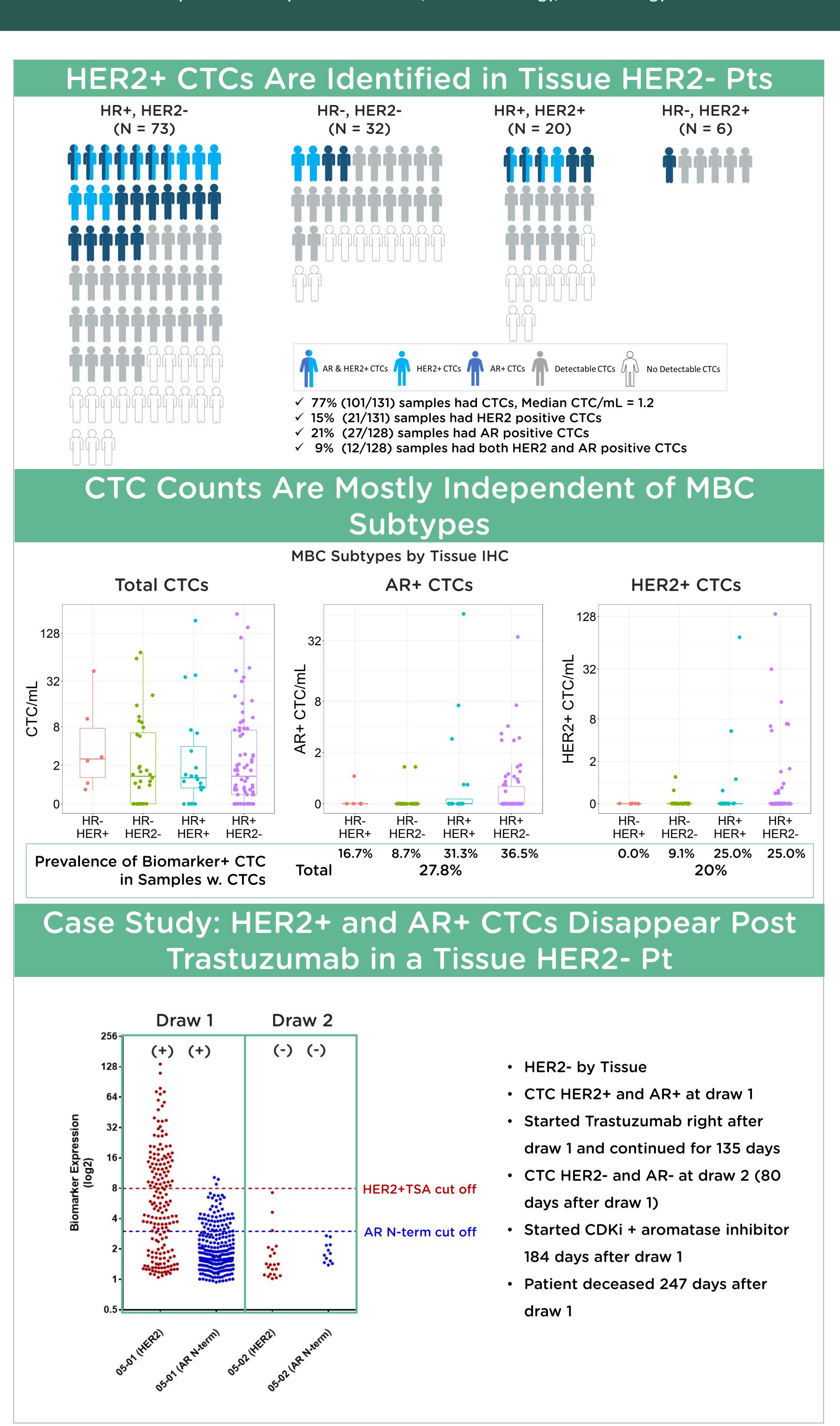
# HER2, AR protein expression and chromosomal instability in circulating tumor cells (CTCs) of metastatic breast cancer (MBC) patients (pts)



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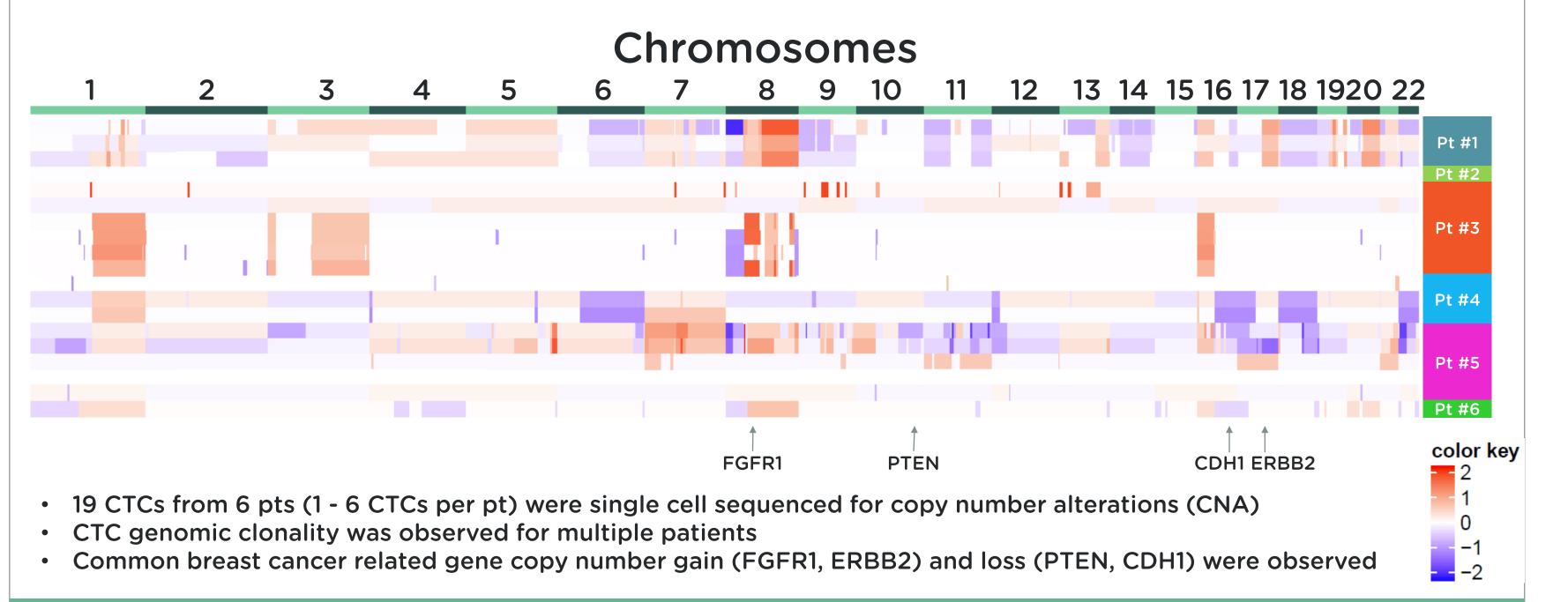
### Upregulation of HER2 and androgen receptor (AR) are mechanisms of acquired resistance to endocrine therapy Measurement of these proteins and their localization requires metastatic biopsies, which are costly, invasive, and prone to under-sampling A CTC-based test could expand the clinical utility of these biomarkers MBC blood samples were characterized for CTC prevalence, HER2 and AR expression on treatment and at time of disease progression using the Epic Sciences platform Methods A. The Epic Sciences CTC Platform 1) Epic Sciences Platform 2) Single Cell Capture and Sequencing 4) SINGLE CELL WGA 5) LIBRARY PREPARATION 6) WHOLE GENOME SEQUENCING & BIOINFORMATICS Single cell dual index/barcodec The state of the s B. Example Cell Images HER2+ CTC All Channels **CD45** AR+ CTC CD45 All Channels DAPI C. Consort 131 blood samples from 82 MBC Pts: • 73 ER/PR+, HER2-•131 Samples tested for HER2 expression • 32 ER/PR-, HER2-•128 samples tested for AR expression • 20 ER/PR+, HER2+ • 6 ER/PR-, HER2+

Background



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### Heterogeneous Genome Profiles Are Observed Across Multiple Patients



#### Conclusions

- The majority (77.1%) of metastatic breast cancer patients had detectable CTCs
- Diverse expression of HER2 and AR were observed and these endocrine therapy resistance markers could potentially guide subsequent therapy selection
- Prospective evaluation of HER2 and AR on MBC pts' CTCs as predictive biomarkers of benefit from inhibitors of these proteins is needed