

Presentations

[Alle Vortragsslides](#)

[\[GS5-01\] A randomized community-based trial of an angiotensin-converting enzyme inhibitor, lisino-pril or a beta blocker, carvedi-lol for the prevention of cardiotoxicity in patients with early-stage HER2-positive breast cancer receiving adjuvant trastuzumab](#)

S Munster P, Krischer J, Tamura R, Fink A, et al.

The authors conclude that: *In patients with HER2-positive breast cancer receiving trastuzumab and an anthracycline, both lisino-pril and carvedi-lol during treatment reduced cardiotoxicity in patients, but not in those with non-anthracycline containing regimens. The use of lisino-pril or carvedi-lol may allow the use of an anthracycline without compromising trastuzumab treatment in those who might benefit from an anthracycline.*

[\[GS5-02\] Cardiovascular function and the effect of exercise training during adjuvant breast cancer treatment. Results from The EBBA-II trial](#)

Thune I, Husøy A, Frydenberg H, Flote VG, et al.

The authors conclude that: *Our findings strongly support that tailored exercise training during adjuvant breast cancer treatment may counteract a decline in cardiovascular function, and in particular among those receiving chemotherapy. Our study supports incorporation of supervised clinical exercise programs into breast cancer treatment guidelines.*

Final results of the trial at SABCS 2018 (the trial closes October 15, 2018) total included N=539 (NCT02240836)

[\[GS5-03\] Lifestyle Intervention and Effect on Disease-free Survival in Early Breast Cancer Pts: Interim Analysis from the Randomized SUCCESS C Study](#)

Janni W, Rack BK, Friedl TW, Müller V, et al.

The authors conclude that: *This explorative and non-planned interim analysis indicates that the completion of a systematic telephone life style intervention program may positively impact patient outcome in early breast cancer.*

[\[GS5-04\] GS5-04: discussant Ligibel
811, 786 & 761](#)

[\[GS5-05\] Resistance to neoadjuvant
chemotherapy in triple negative breast
cancer mediated by a reversible drug-
tolerant state](#)

Echeverria GV, Ge Z, Seth S, Jeter-Jones
SL, et al.

The authors conclude that: *Collectively,
these studies reveal that a reversible
phenotypic state can confer chemoresistance
in the absence of genomic selection and that
the residual tumor state is a novel therapeutic
window for chemo-refractory TNBC.*

[\[GS5-06\] No survival benefit of
chemotherapy escalation in patients
with pCR and “high-immune” triple-
negative early breast cancer in the
neoadjuvant WSG-ADAPT-TN trial](#)

Gluz O, Nitz U, Liedtke C, Prat A, et al.

The authors conclude that: *Our
exploratory results suggest independent*

*prognostic impact of mRNA markers and
TIL's in early TNBC. Patients with both pCR
(after 12 weeks) and "high-immune"
signature (defined here by PD1) had
excellent 3y-EFS and may be candidates for
treatment de-escalation (e.g. omission of
anthracyclines), whereas "low-immune" pCR
patients may benefit from standard adjuvant
poly-chemotherapy.*

[\[GS5-07\] International pooled analysis
of the prognostic impact of disseminated
tumor cells from the bone marrow in
early breast cancer: Results from the
PADDY study](#)

Hartkopf AD, Brucker SY, Taran F-A,
Harbeck N, et al.

The authors conclude that: *Detection of
DTC in the bone marrow is an independent
prognostic marker in patients with non-
metastatic breast cancer. Further studies
should investigate the impact of DTC on
metastatic cancer progression and their role
for clinical decision making.*

San Antonio - Mosaic on a pillar of streetcar station

