

Prevention of Cardiac Dysfunction During Adjuvant Breast Cancer Therapy (PRADA): Long-Term Follow-Up of a 2 x 2 Factorial, Randomized, Placebo-Controlled, Double-Blind Clinical Trial of Candesartan and Metoprolol

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Disclosures

Siri Lagethon Heck: None

Torbjørn Omland (Principal Investigator)

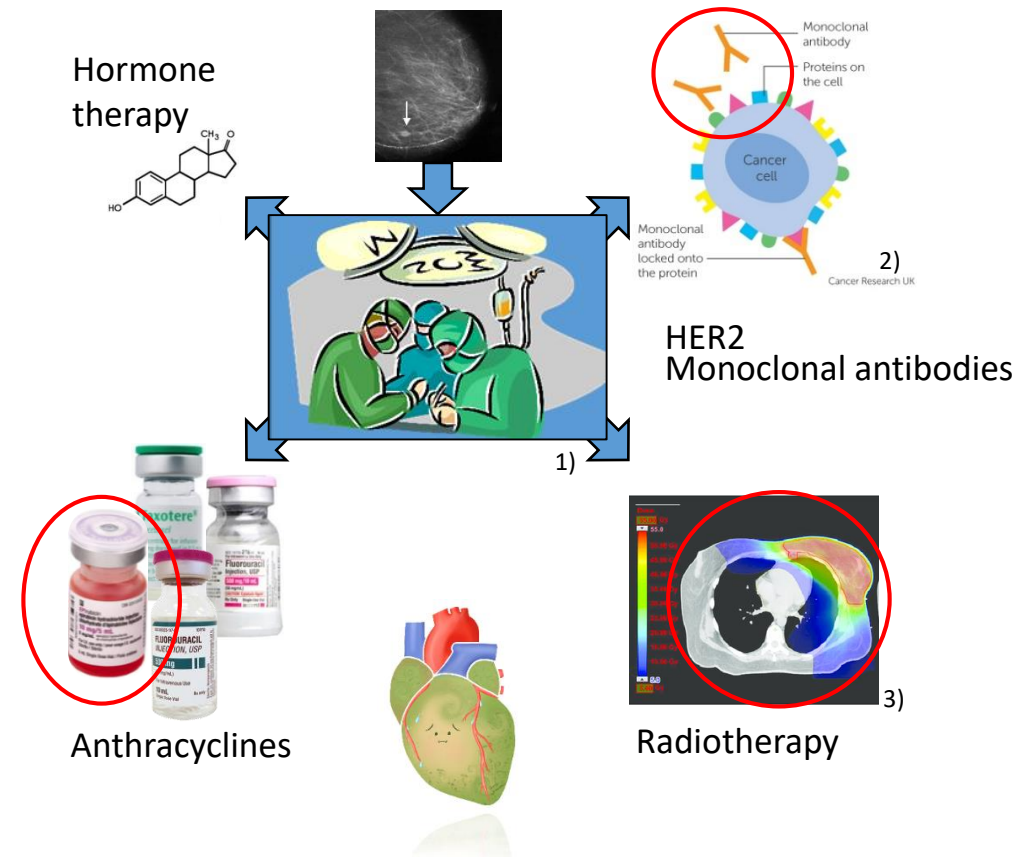
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Adjuvant breast cancer therapy prolongs survival but may harm the heart

Anthracyclines, trastuzumab and radiotherapy have cardiotoxic effects

Neurohormonal blockade may attenuate the myocardial damage during cancer treatment

Long-term effects are unknown



Trials on neurohormonal blockade during breast cancer treatment have shown modest and inconsistent beneficial effects

Study	Year	Intervention	Cancer therapy	n	Results
PRADA Gulati et al	2016	Candesartan Metoprolol	Anthracyclines Trastuzumab Radiotherapy	120	Candesartan attenuated a decline in LVEF Metoprolol attenuated troponin increase
Boekhout et al	2016	Candesartan	Trastuzumab	210	No difference in cardiotoxic events No difference in change in LVEF
MANTICORE Pituskin et al	2016	Perindopril Bisoprolol	Trastuzumab	94	No difference in change in LVEDV Attenuated decline in LVEF
CECCY Avila et al	2018	Carvedilol	Anthracyclines Taxanes	200	No difference in cardiotoxic events No difference in change in LVEF Attenuated troponin increase
Guglin et al	2019	Lisinopril Carvedilol	Trastuzumab	468	No difference in cardiotoxic events Effect in the subgroup who had received anthracyclines



PRADA: a 2 x 2 factorial, randomized, placebo-controlled, double-blind trial

Women 18-70 years with early breast cancer
Anthracycline containing adjuvant treatment

No serious concomitant illness
No significant cardiovascular disease
No prior chemo/radiotherapy
No indication/contraindications for the study drugs

Hypothesis

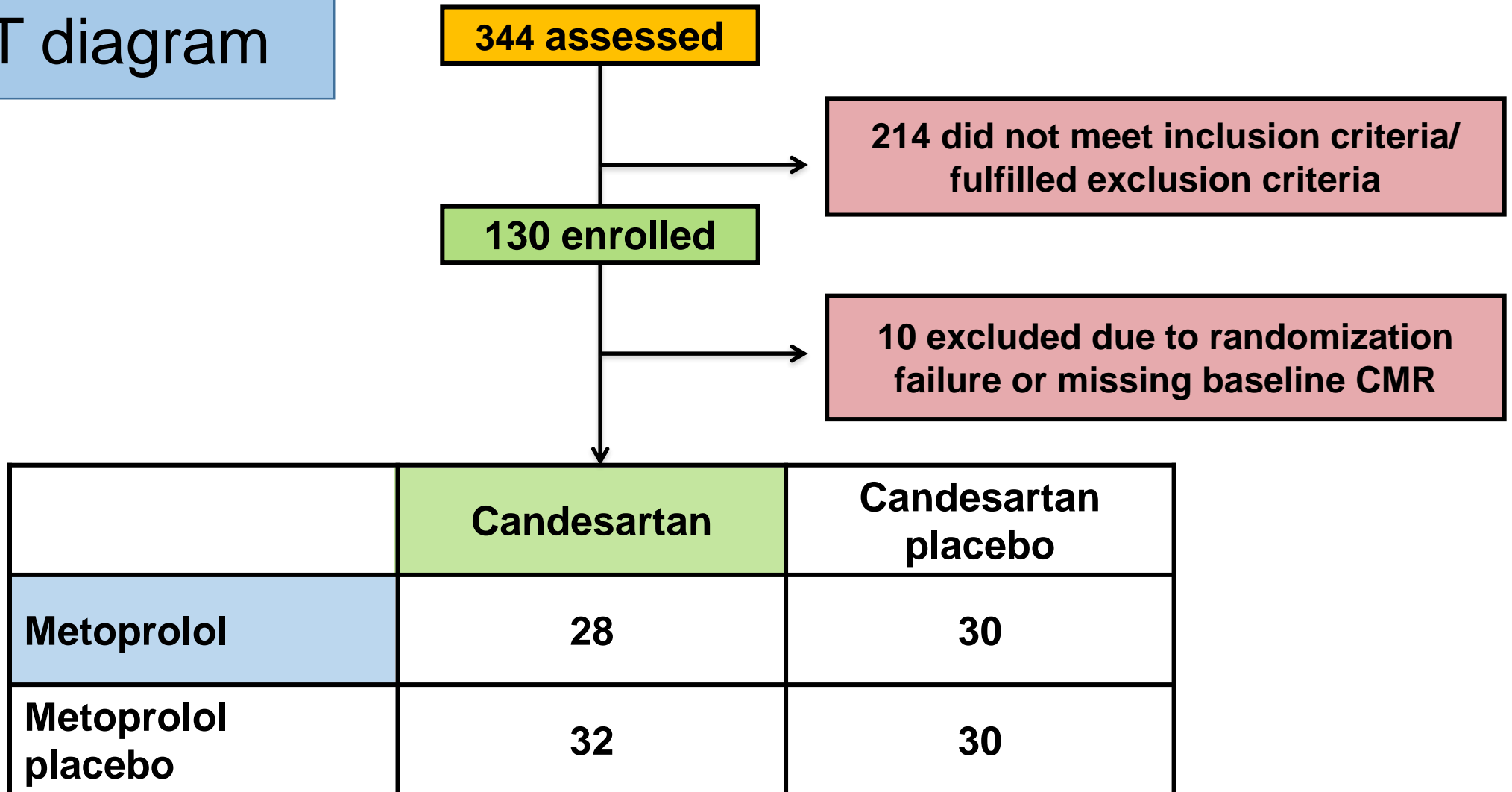
Long-term decline in cardiac function can be prevented by treatment with metoprolol and candesartan during adjuvant treatment for early breast cancer

Power calculations

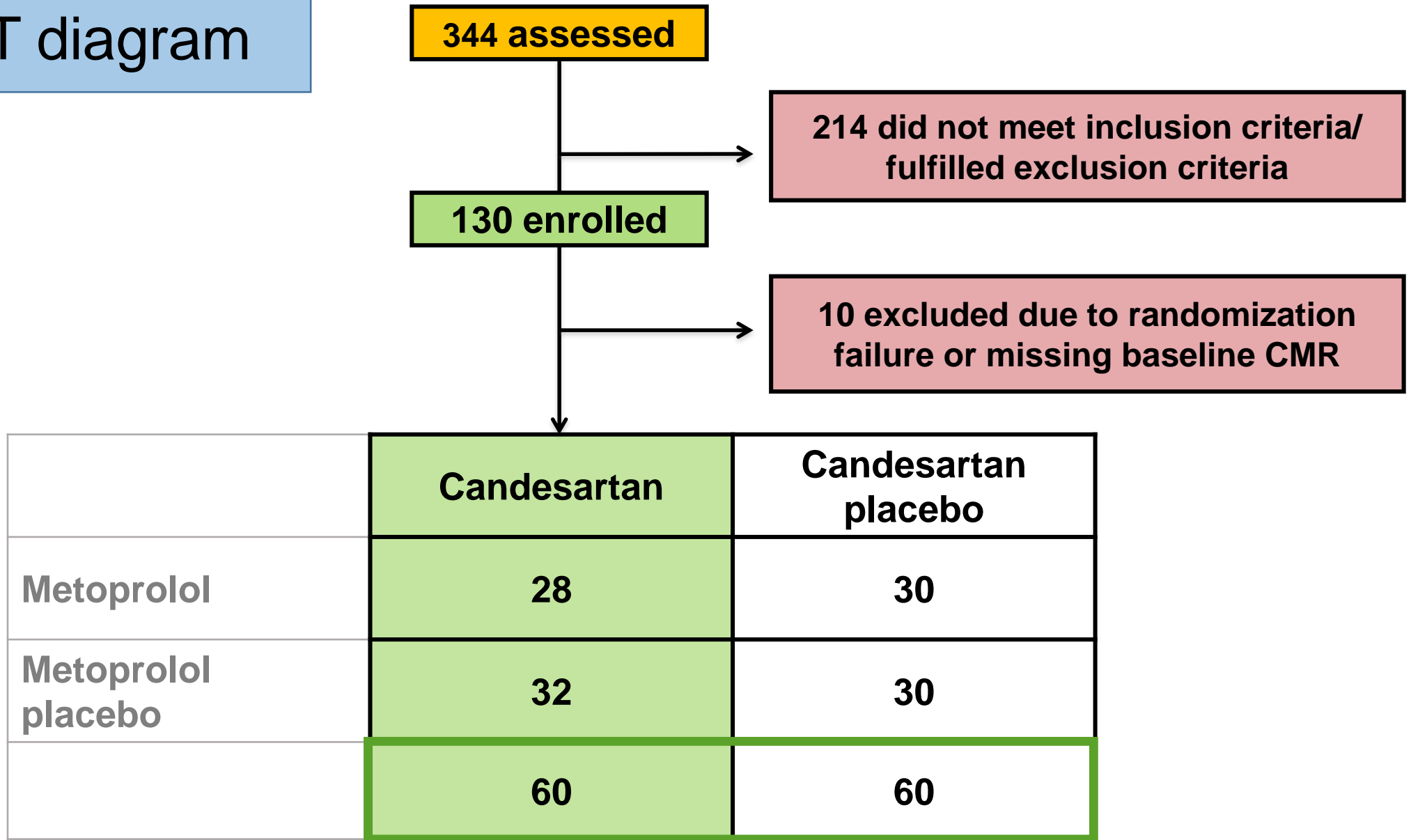
Change in LVEF	Sample size	Power
5±5%	120	95%
3±5%	120	90%
2±5%	120	60%

$\alpha = 0.05$

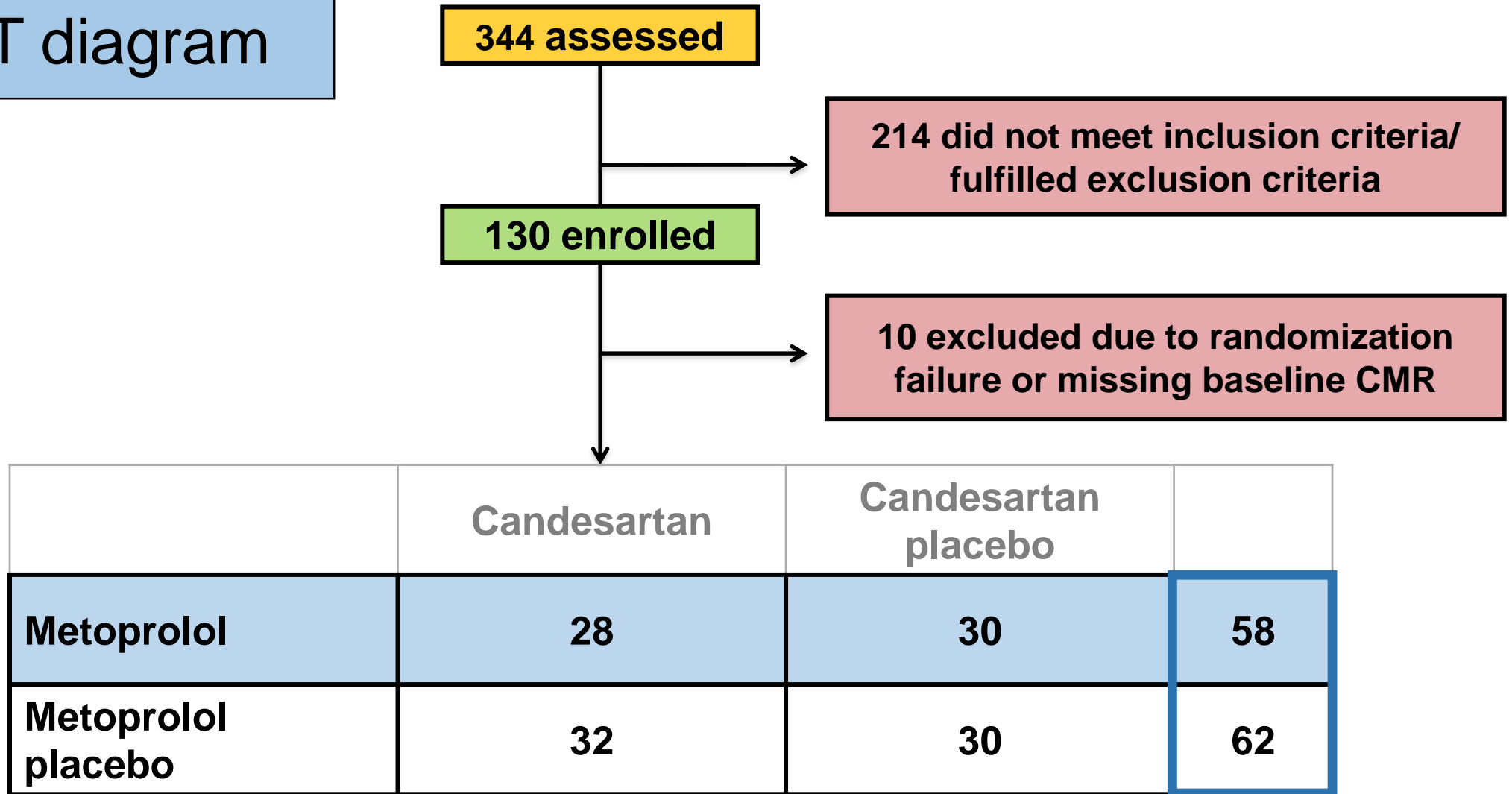
CONSORT diagram



CONSORT diagram

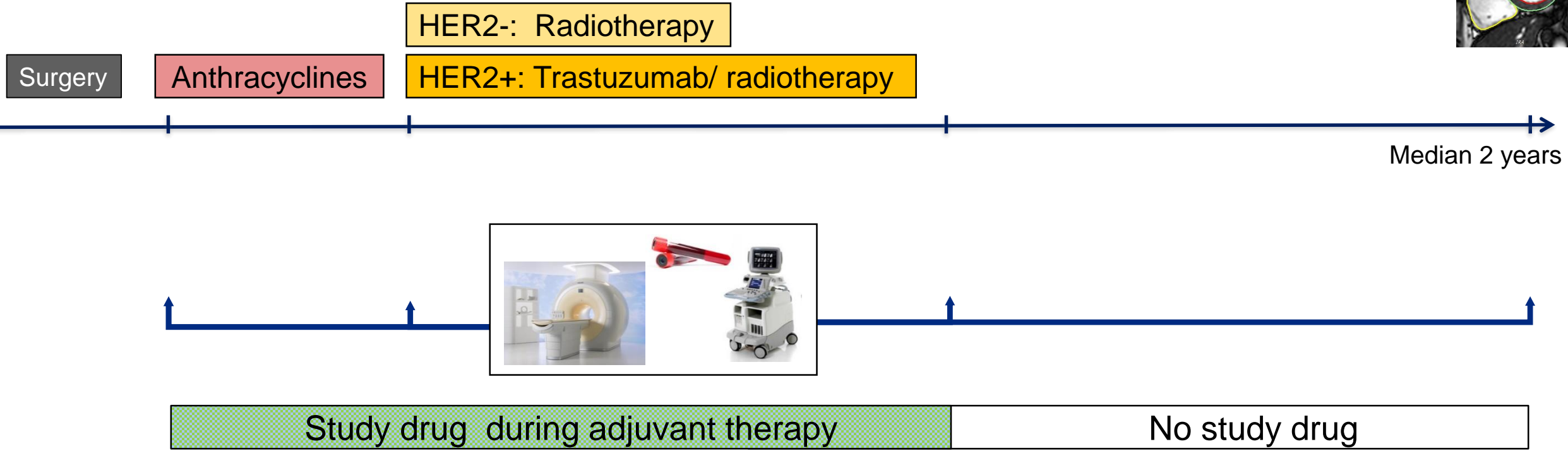
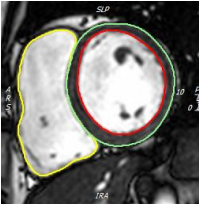


CONSORT diagram



Study flowchart

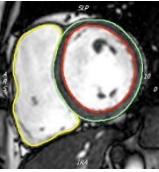
Primary endpoint: Change in LVEF



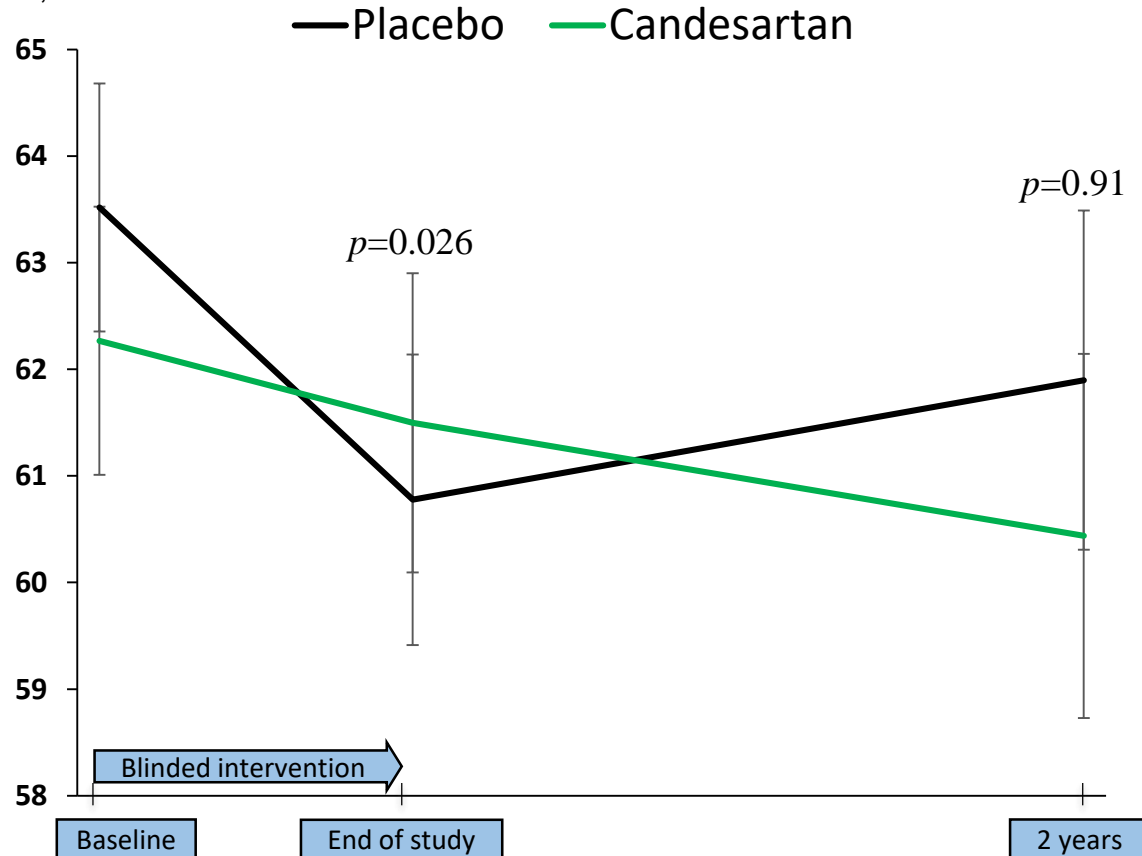
Baseline and cancer treatment characteristics

	Candesartan-Metoprolol	Candesartan-Placebo	Placebo-Metoprolol	Placebo-Placebo
N	28	32	30	30
Age at recruitment (years)	50 ± 9	52 ± 11	51 ± 9	51 ± 9
Systolic blood pressure (mmHg)	125 ± 13	132 ± 14	133 ± 12	130 ± 13
Heart rate (beats/min)	68 ± 11	68 ± 10	70 ± 12	65 ± 11
BMI	24 ± 3	26 ± 4	27 ± 6	26 ± 4
Current smokers	5/28 (18 %)	7/32 (22%)	4/30 (13%)	5/30 (17%)
Hypertension	1/28 (4%)	5/32 (16%)	2/30 (7%)	0/30 (0%)
Diabetes	0/28 (0%)	1/32 (3%)	1/30 (3%)	0/30 (0%)
Epirubicin, median dose mg/m ²	240 (240, 400)	240 (240, 400)	240 (240, 400)	240 (240, 400)
Trastuzumab	7/28 (25%)	7/32 (22%)	6/30 (20%)	7/30 (23%)
Radiation	16/28 (57%)	19/32 (59%)	20/30 (67%)	21/30 (70%)
Taxanes	24/28 (86%)	25/32 (78%)	25/30 (83%)	22/30 (73%)

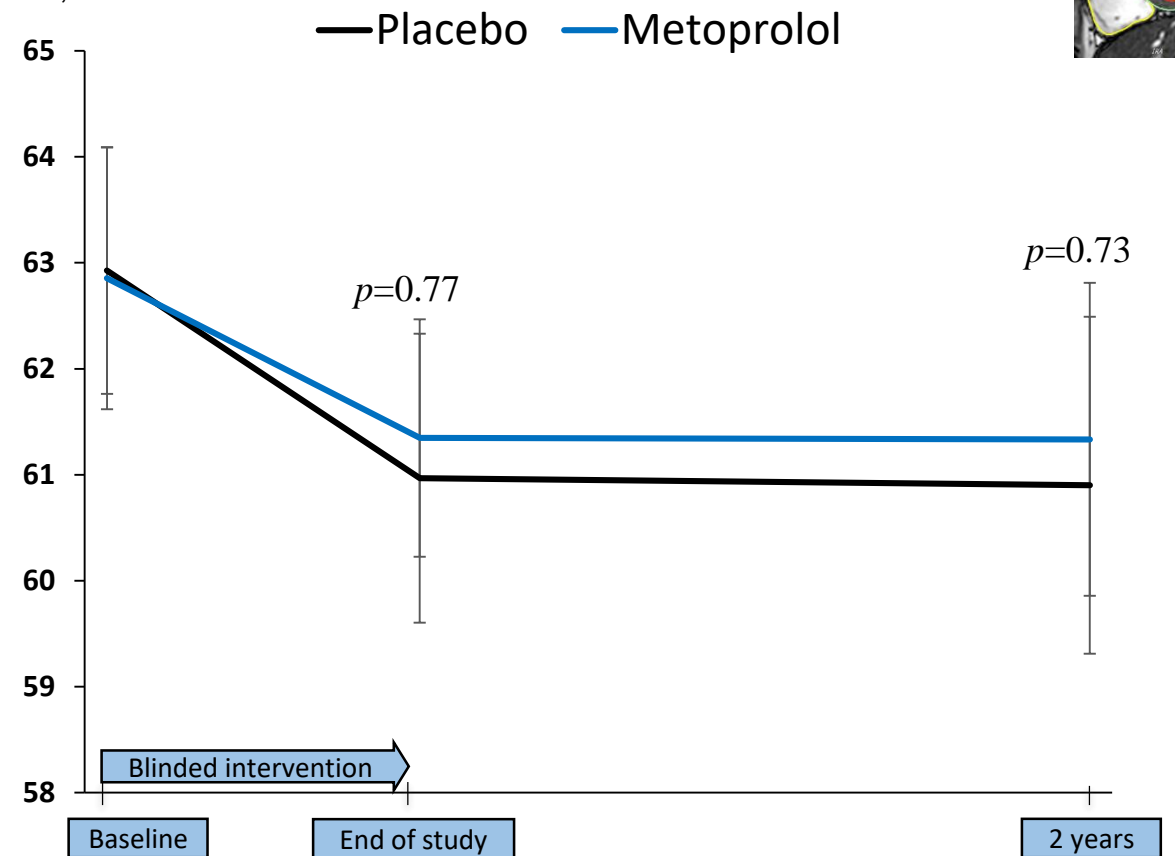
At 2 years, there were no between-group differences in change in LVEF



Observed values,
mean, 95% CI



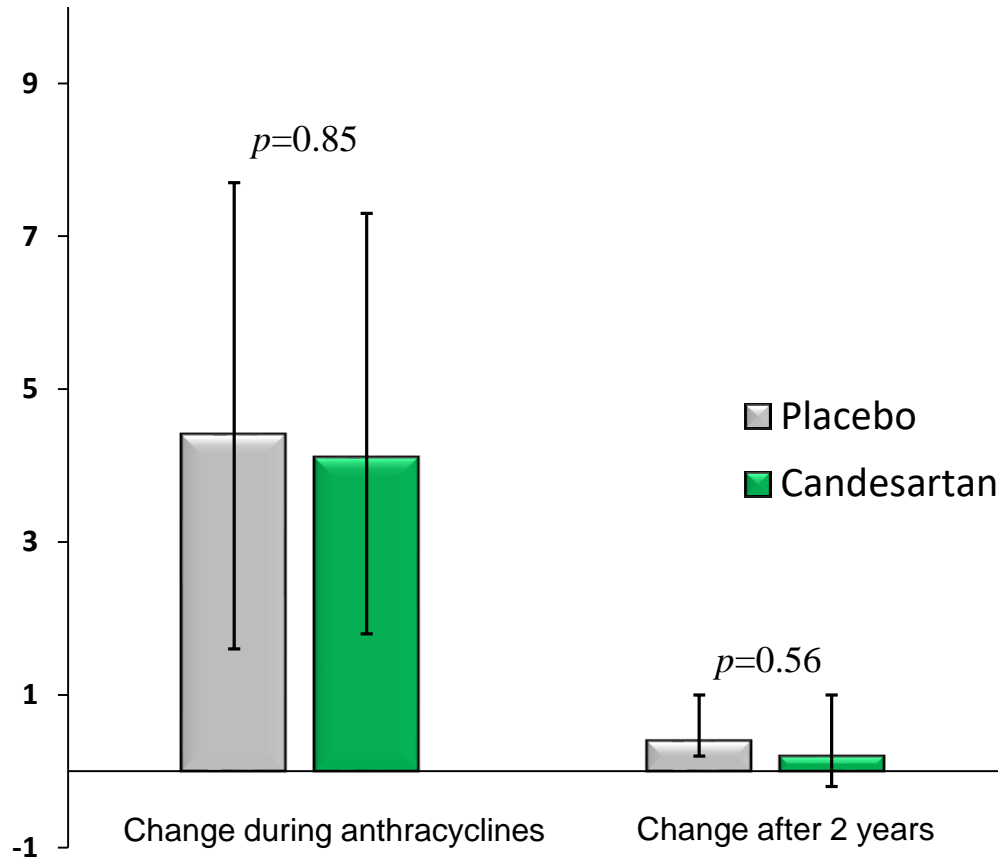
Observed values,
mean, 95% CI



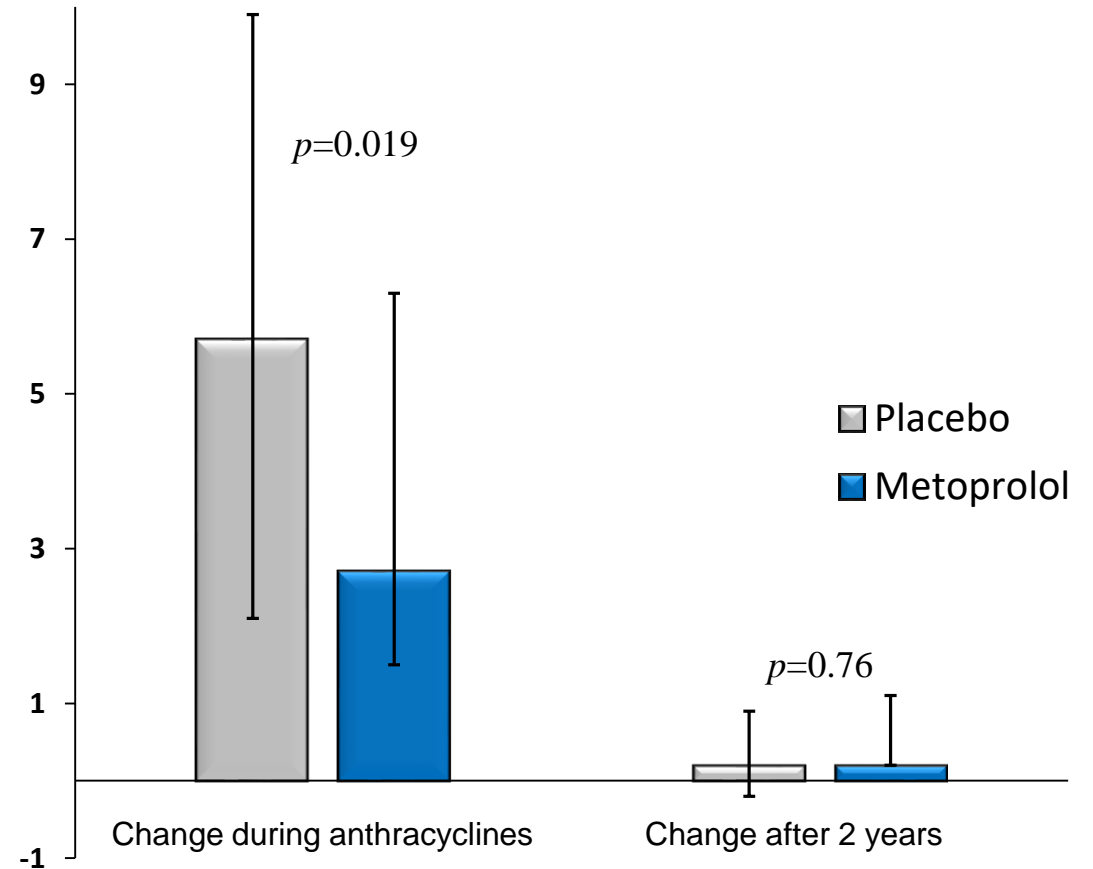
p-values are for between-group differences in the linear mixed model in the intention to treat analysis

At 2 years, there were no between-group differences in change in troponin I

Change from baseline, ng/L median, IQR

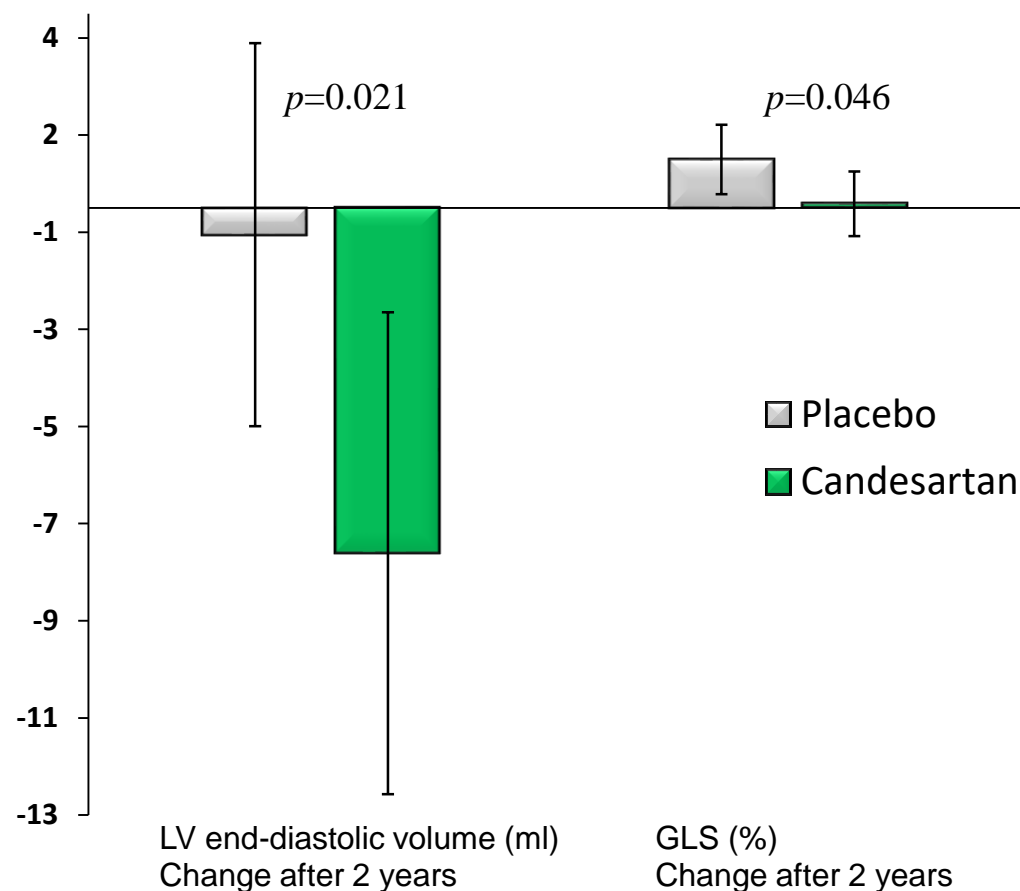


Change from baseline, ng/L median, IQR

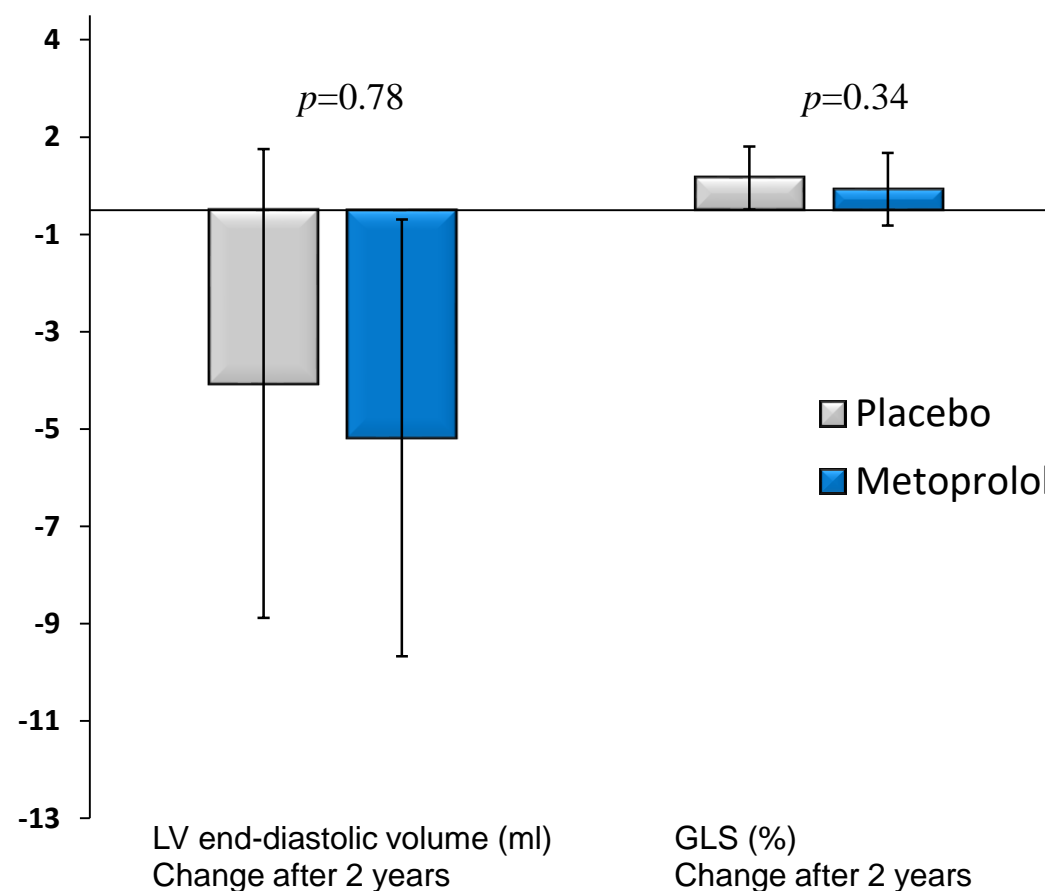


At 2 years, LV end-diastolic volume was reduced and GLS decline attenuated in the candesartan arm

Change from baseline,
Mean, 95% CI



Change from baseline,
Mean, 95% CI



Strengths

Trial design

LVEF assessed by serial CMR

2 year follow-up

Limitations

Single center

18% of participants did not have 2-year CMR

Less decline in LVEF than anticipated



Conclusion

Candesartan and metoprolol during adjuvant therapy for early breast cancer did not protect against long-term decline in LVEF



Clinical implications

Broadly administered cardioprotective therapy may not be required during adjuvant breast cancer therapy, as decline in systolic function was minor and not prevented by neurohormonal blockade

Study organization

Study steering committee

T Omland (Chair and Primary Investigator)
J Geisler (Clinical oncology)
AH Ree (Radiation oncology)
P Hoffmann (Cardiac MRI)
H Røsjø (Biobank)
K Steine (Echocardiography)

Investigators

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J Schulz-Menger (Cardiac MRI)
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Study statistician

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Data safety and monitoring board

P Smith (Chair)
O Engebråten (Clinical oncology)
FA Dahl (Biostatistician)

Akershus University Hospital

Clinical Research Unit
Division of Radiology, Surgery, Oncology and Cardiology

Oslo University Hospital

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