

AKC 13. Milli Konqresi, 13-14 Dekabr 2024, Bakı



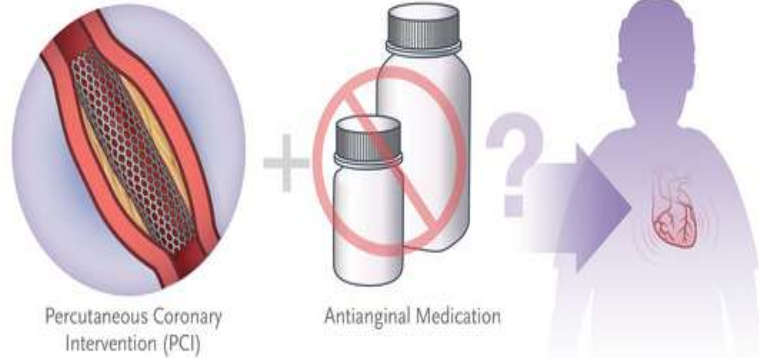
**Dərman müalicələri çağ atlayıb,
hər kəsin damarına müdaxilə şərt deyil.**

Fuad Səmədov

ORBITA 2

A Placebo-Controlled Trial of Percutaneous Coronary Intervention for Stable Angina

Rajkumar CA et al. DOI: 10.1056/NEJMoa2310610



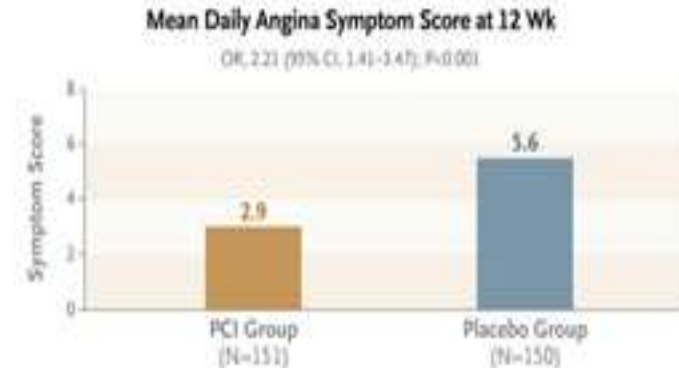
301
patients

Inclusion criteria: Patients suitable for PCI, with angina or equivalent symptoms, anatomical evidence of significant coronary stenosis in ≥ 1 vessel, either: a. Invasive angiogram indicating $\geq 70\%$ stenosis b. CT coronary angiography indicating $\geq 90\%$ stenosis, and evidence of ischemia.

PCI group
(n=151)

VS.

placebo procedure
(n=150)



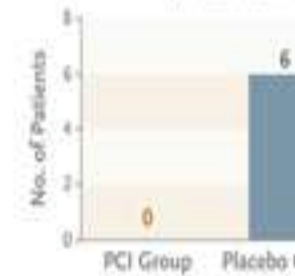
701

Mean treadmill exercise time (sec)

641

Spontaneous MI

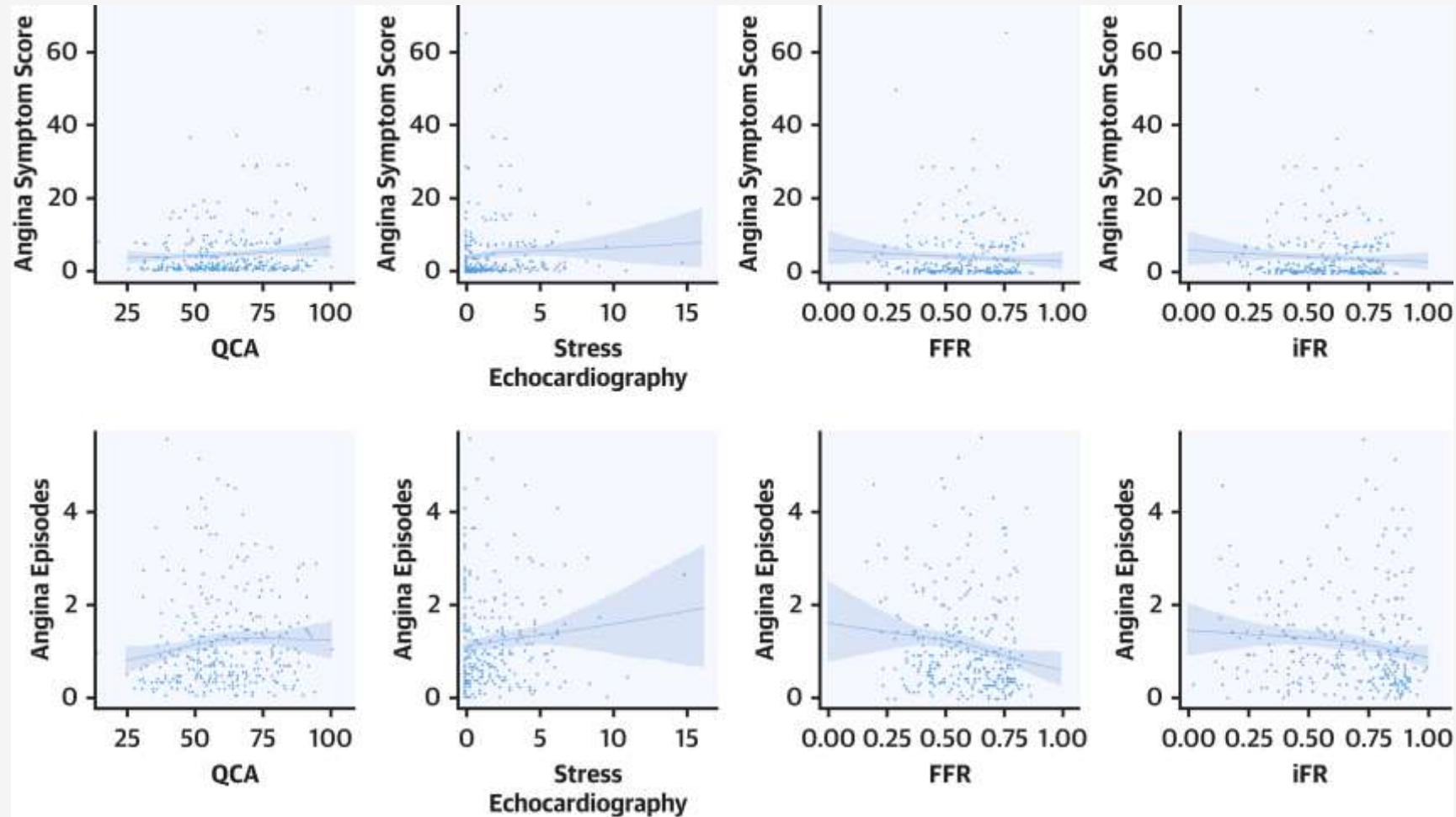
Stroke



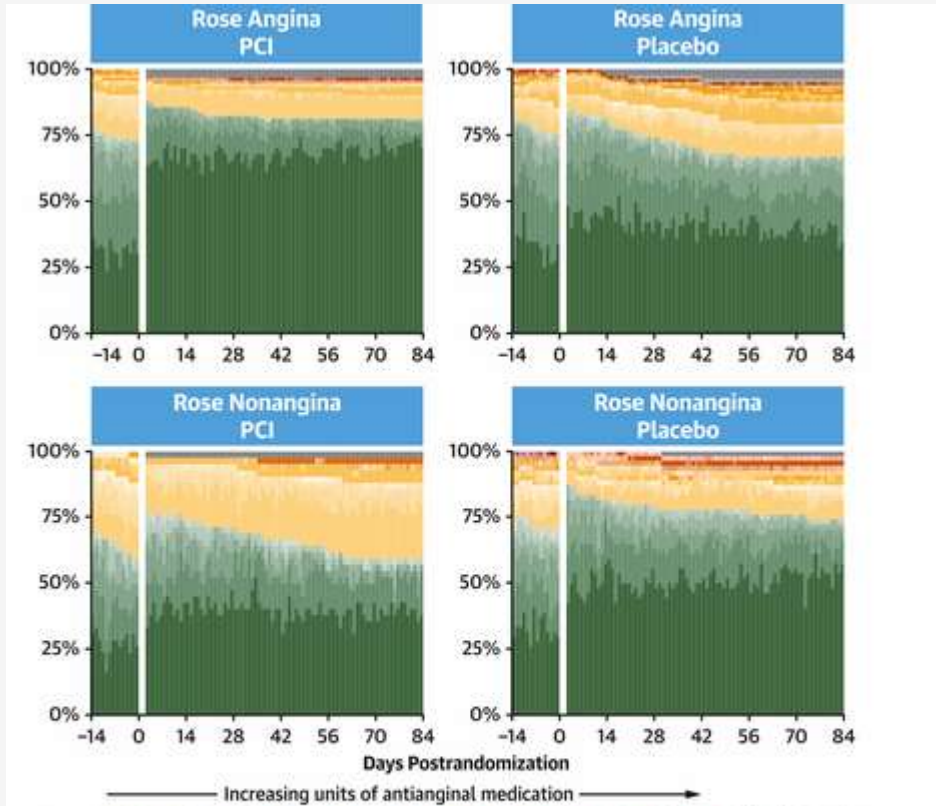
CONCLUSIONS

Among patients with stable angina who were receiving little or no antianginal medication and who had objective evidence of ischemia, PCI resulted in a better health status with respect to angina than a placebo procedure at 12 weeks.

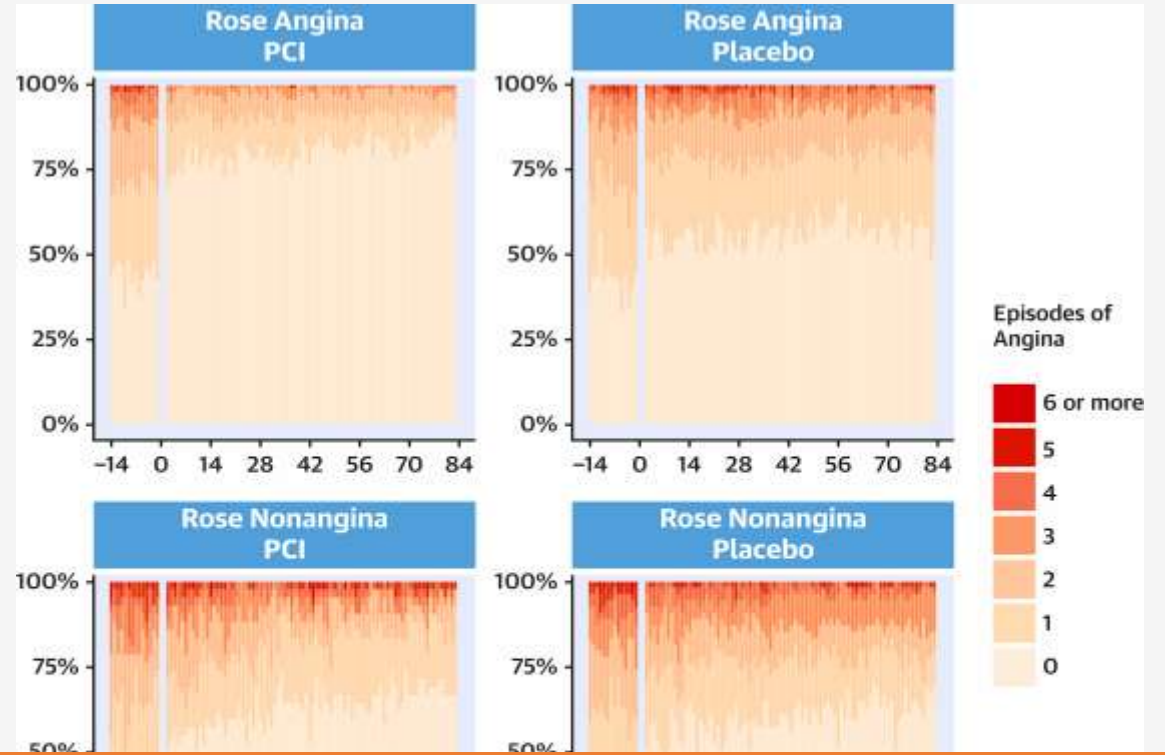
ORBITA 2 substudy – simptom/lezyon asılılığı



ORBITA 2 substudy – Rose Angina score

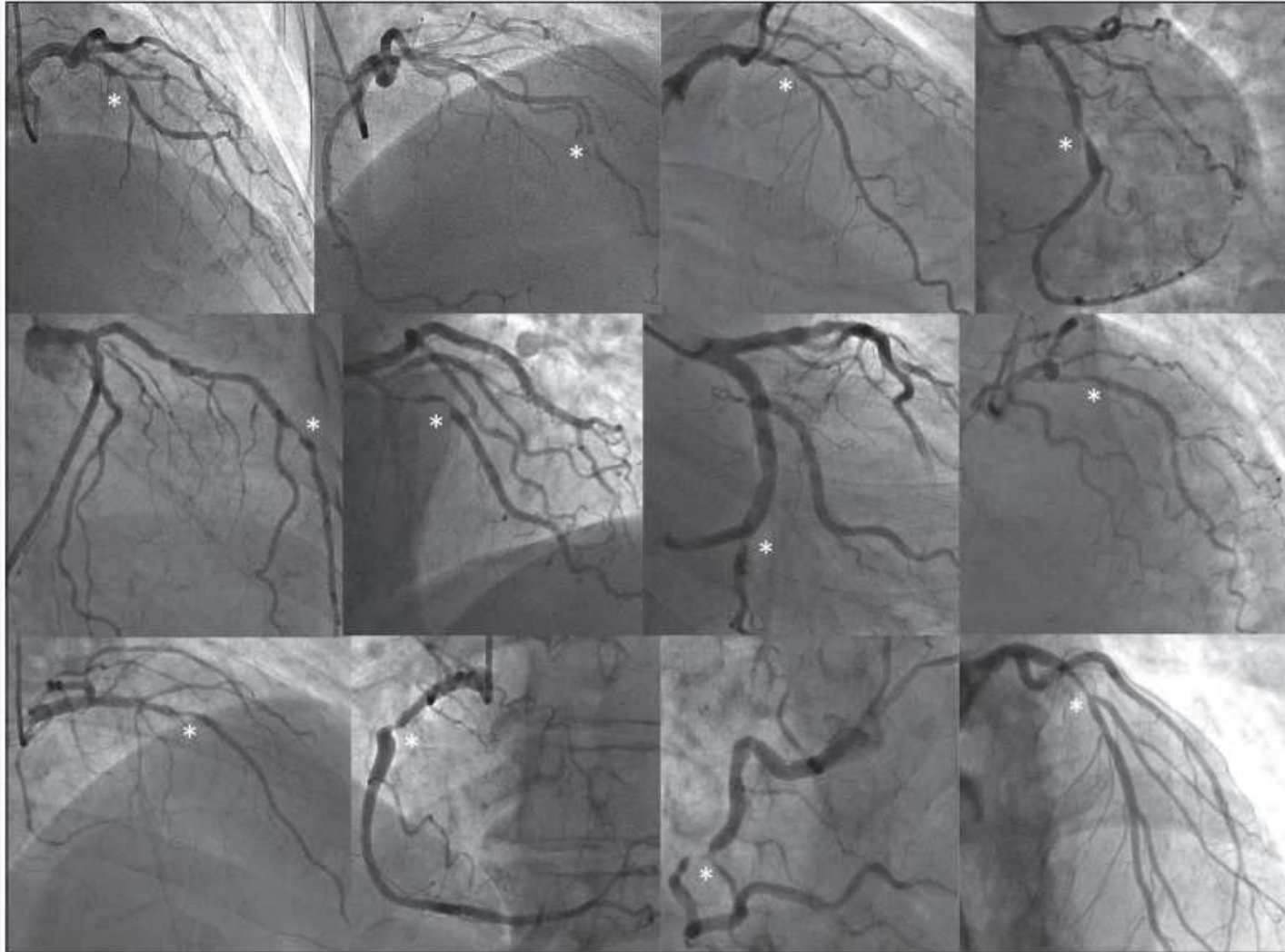


Patients with more severe angina were more likely to achieve (odds ratio [OR] = 4.3) a better placebo-controlled “health state.”

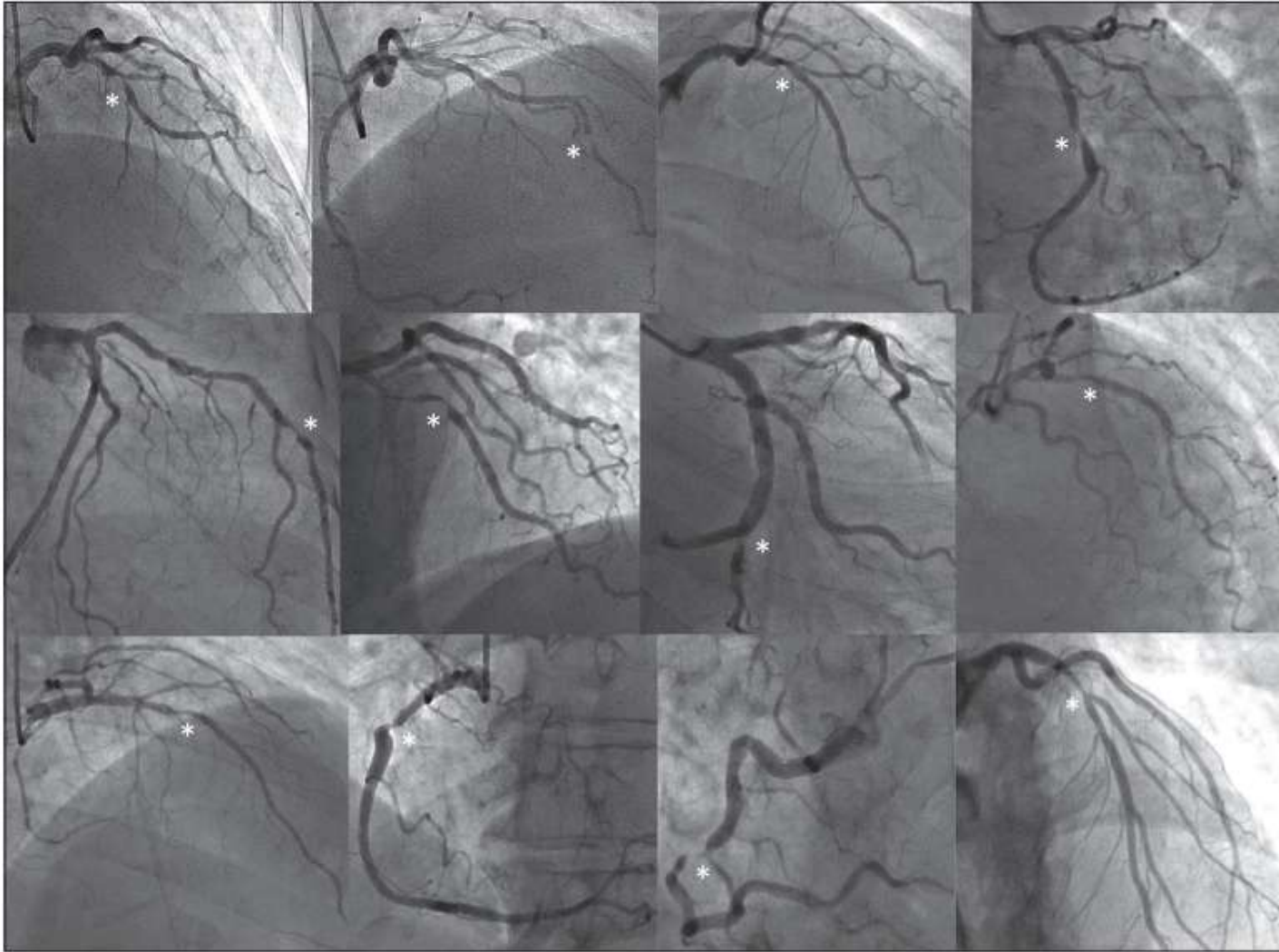


Patients who met criteria for Rose angina (vs those who did not) were more likely (OR 1.9) to have a placebo-controlled benefit from PCI on the angina symptom score and the number of angina episodes.

???



??? - ORBITA tədqiqatı – ilk 12 pasientin angiogramı



✓ ORBITA-2 : PCI qrupunda işemiya ortadan qalxsa da 60% pasientin simptomları davam etmişdir.

✓ Angioqrafiya göstərişləri? KAX skrininqi?

✓ Gündəlik həyatda bu ehtimal necədir?

✓ Tənliyi həll etməyə biz həkimlər istəmədən mane oluruq?

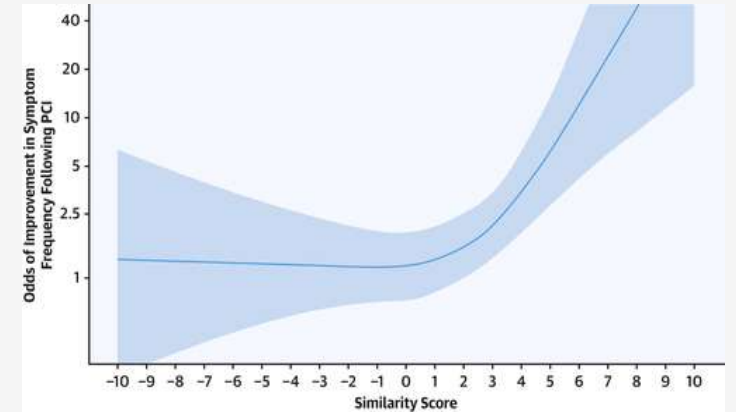
ORBITA -STAR

- Tək damar KAX – 51 xəstə.
- Kat lab – stenoz bölgəsində 4 dəfə 60sn-lik balon
- 4 plasebo və ya sham balon şişmə
- Balon okkluziyadan sonra simptomlar bənzərliyi ?
- Gündəlik həyatdakı simptom – balon okkluziyada simptom bənzərliyi
- Bu bənzərliyin PCI sonrası simptomlarda azalmanı prediktə edə bilməsi

Simptom skoru -10 ilə +10 arasında dəyişib.
10 - - - plasebo balon istifadəsində balon okkluziya ilə eyni simptomlar
+ 10 - - - balon okkluziyası ilə gündəlik simptomlar eynidir.

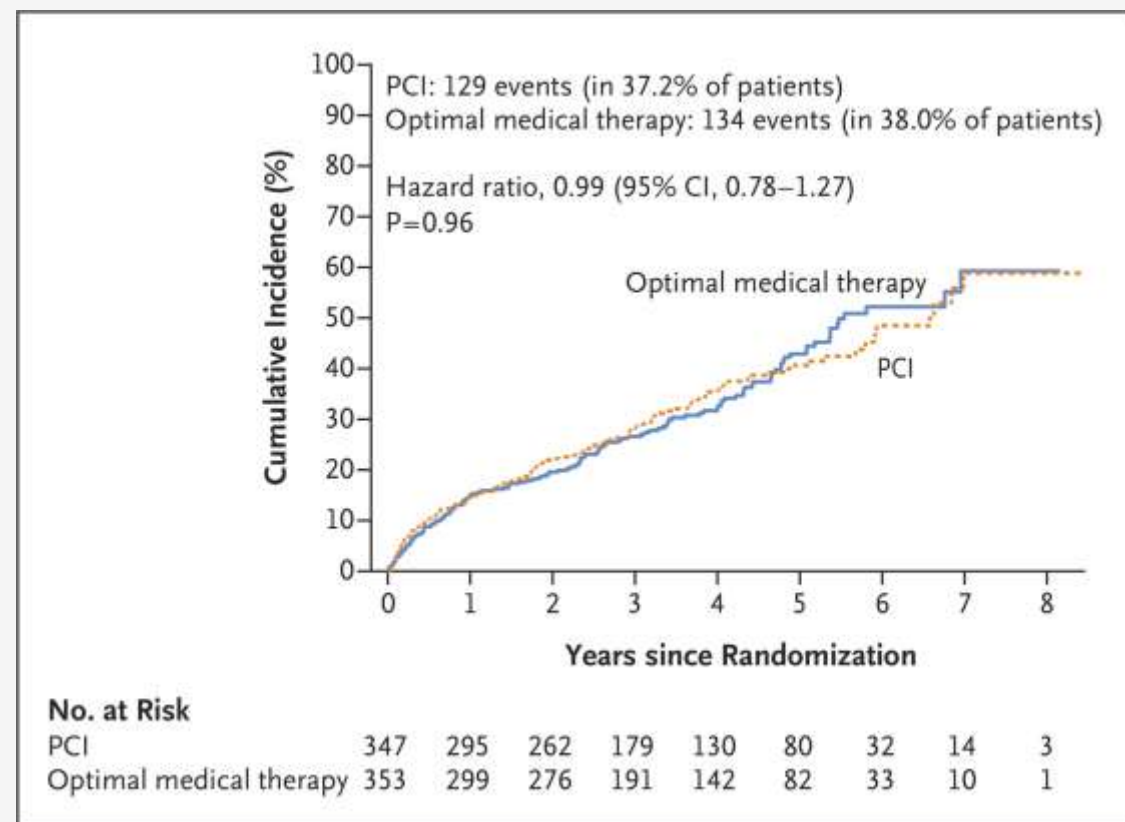
**Median
bənzərlik
skoru**

3



REVIVED-BCIS

- ✓ LV disfunksiya + PCI uyğun KAX + canlı miokard
- ✓ PCI vs medikal müalicə
- ✓ 3.4 il təqib
- ✓ Ölüm və ya ÜÇ hospitalizasiyasında fərq olmamışdır.
- ✓ LVEF-də yaxşılaşma olmamışdır.
- ✓ İşemik LV disfunksiyasında PCI müalicəsinin medikamentoz müalicədən üstün olduğu altqrup olmamışdır.
- ✓ Tam revaskularizasiya qrupunda belə dərmanla fərq olmamışdır. (HR 0.90, CI 0.62-1.32).



REVIVED-BCIS

✓ PCI – stent və ya prosedurun texnikası illər ərzində çox təkmilləşib. Problem hardadır?
 Problem – sistemik xəstəliyi fokal metodlarla həllindən çox şeylər gözləməyimizdir.

✓ İшемik KMP qrupu pasientlər 3 qrup müalicədən faydalanır:

✓ Yüksək effektivlikli statinlər + antiplatelet müalicə

✓ Ürək çatışmazlığının 4-lü müalicəsi

✓ ICD (CRT)

NYHA functional class — no./total no. (%)§

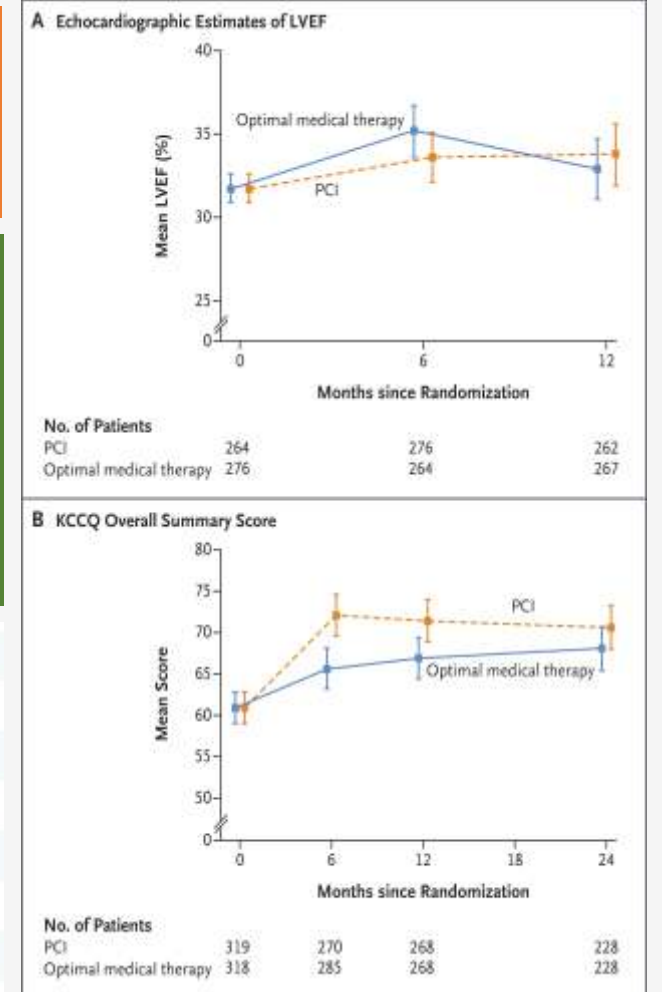
| | | |
|-----------|--------------|--------------|
| I or II | 265/345 (77) | 248/350 (71) |
| III or IV | 80/345 (23) | 102/350 (29) |

CCS angina class — no./total no. (%)¶

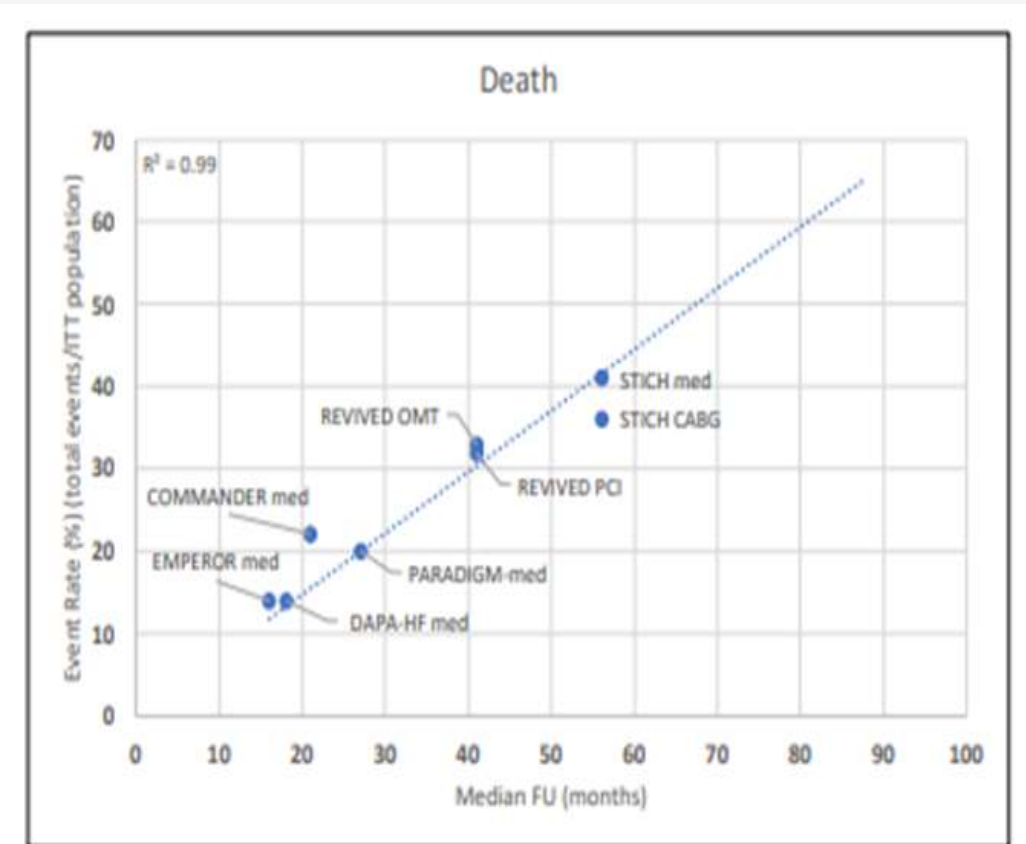
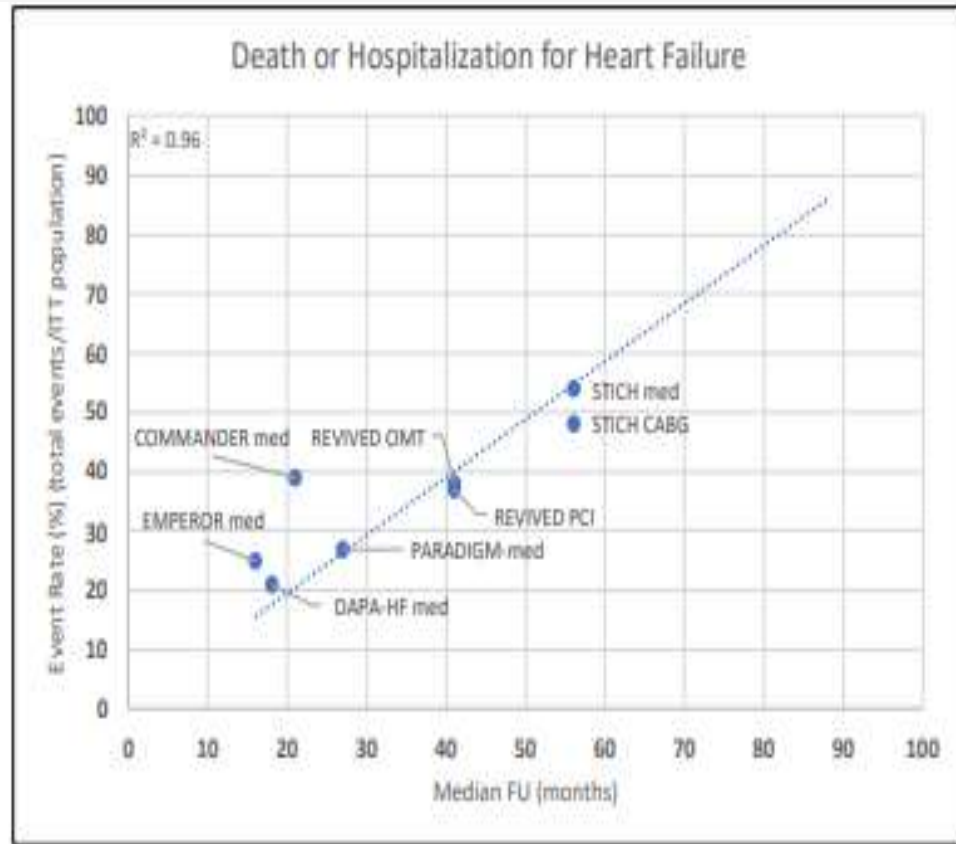
| | | |
|-----------|--------------|--------------|
| No angina | 228/346 (66) | 236/351 (67) |
| I or II | 111/346 (32) | 107/351 (30) |
| III | 7/346 (2) | 8/351 (2) |

Left ventricular ejection fraction — %||

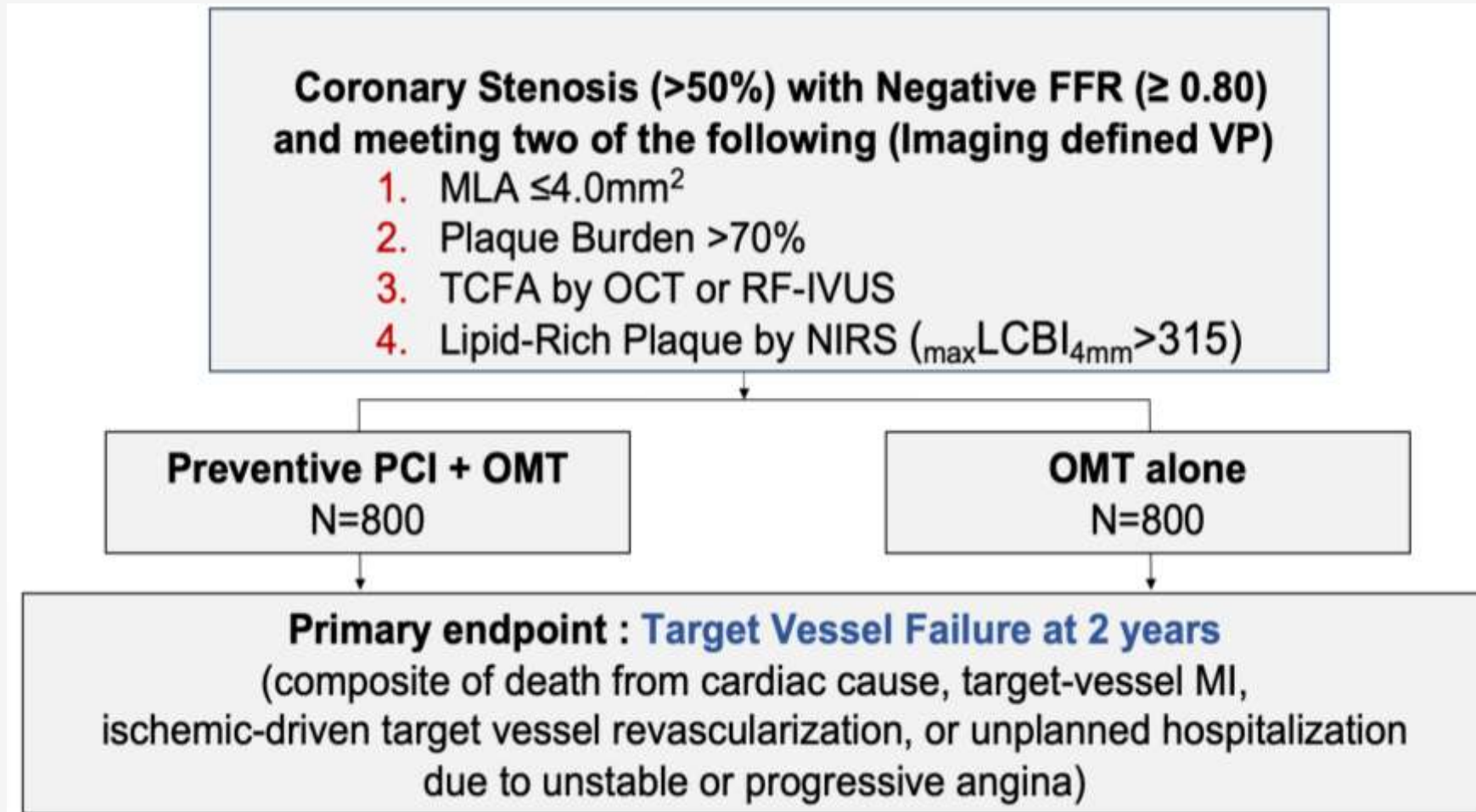
| | | |
|--|----------|----------|
| | 27.0±6.6 | 27.0±6.9 |
|--|----------|----------|



REVIVED-BCIS vs digər tədqiqatlar



PREVENT tədqiqatı

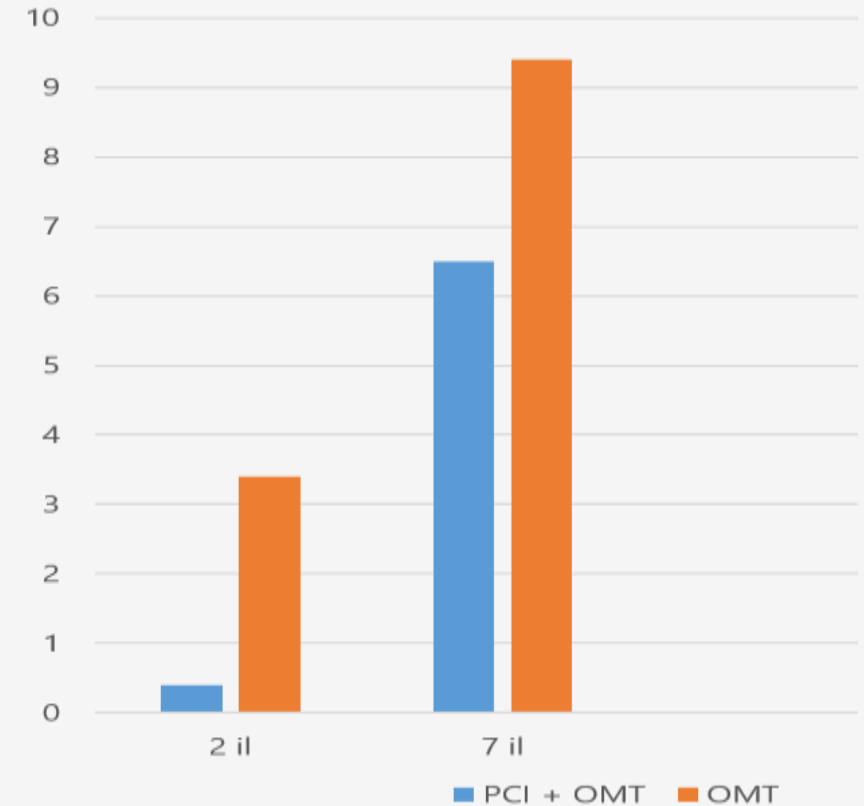


PREVENT tədqiqatı

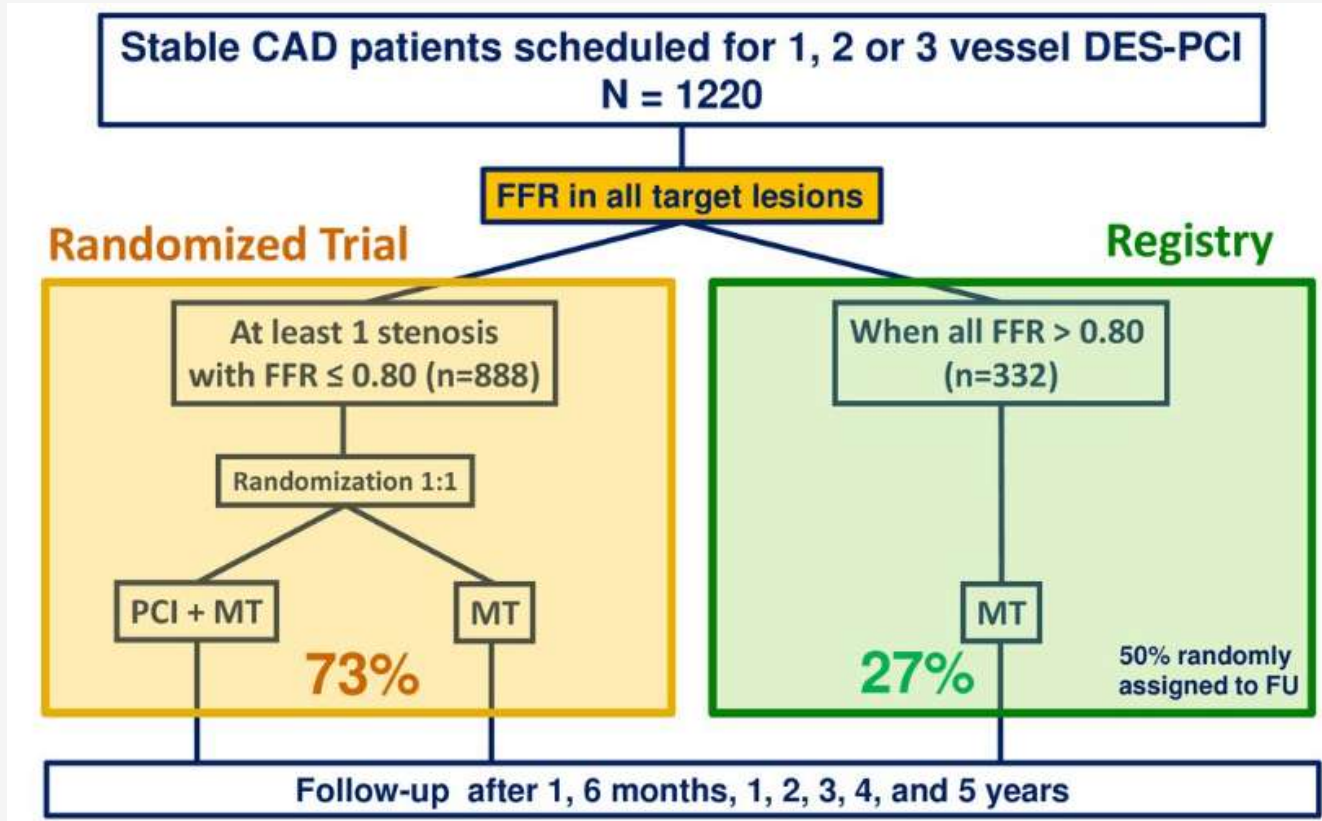
Birincili sonlanma nöqtəsi

2 ildə hədəf damarla əlaqədar ağırlaşmaların cəmi

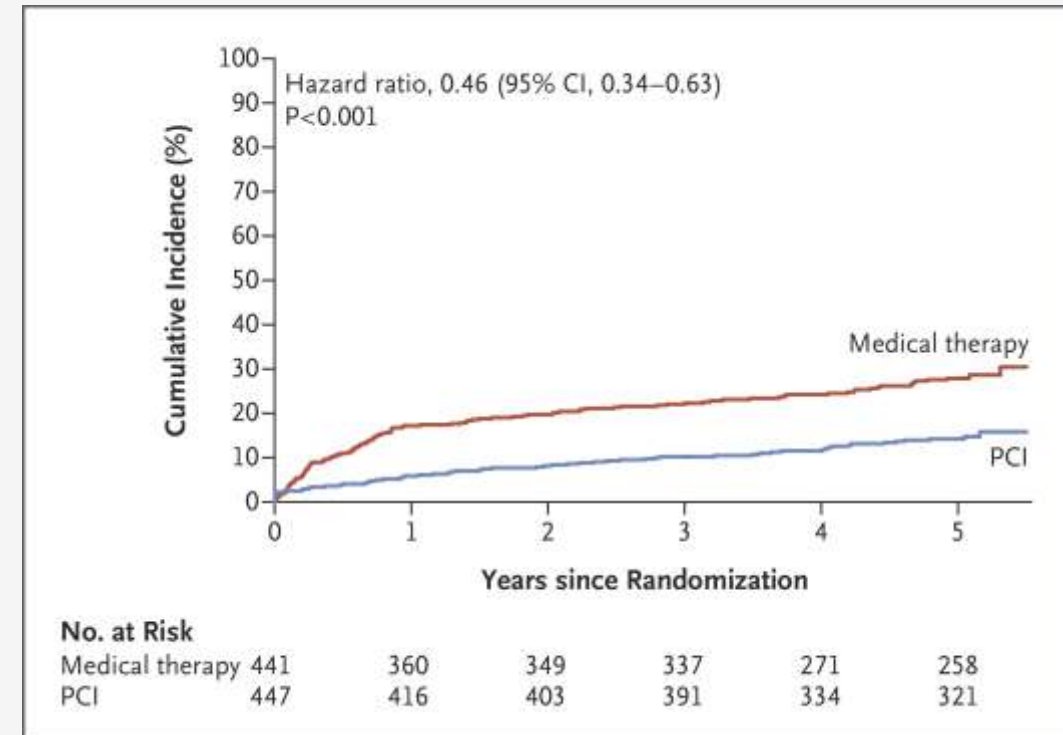
- ✓ kardiak səbəbli ölüm
- ✓ hədəf damarla əlaqədar MI (TV-MI)
- ✓ İşemiya səbəbli hədəf damar müdaxiləsi (ID-TLR)
- ✓ USAP/proqressiv angina səbəbli hospitalizasiya



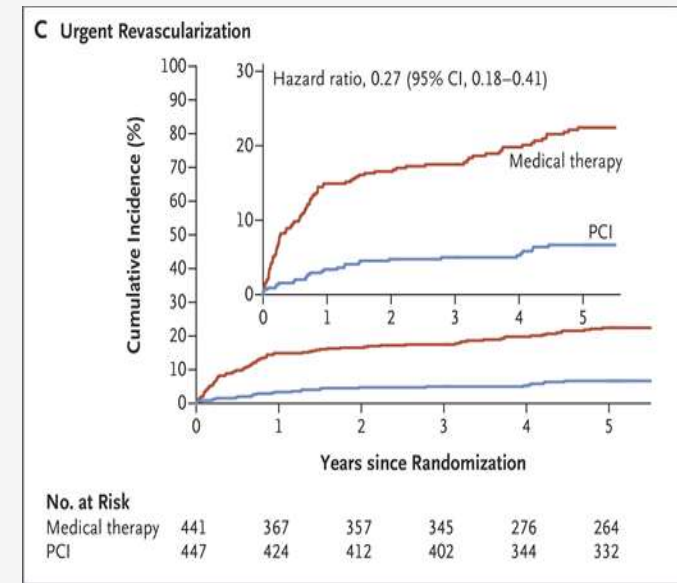
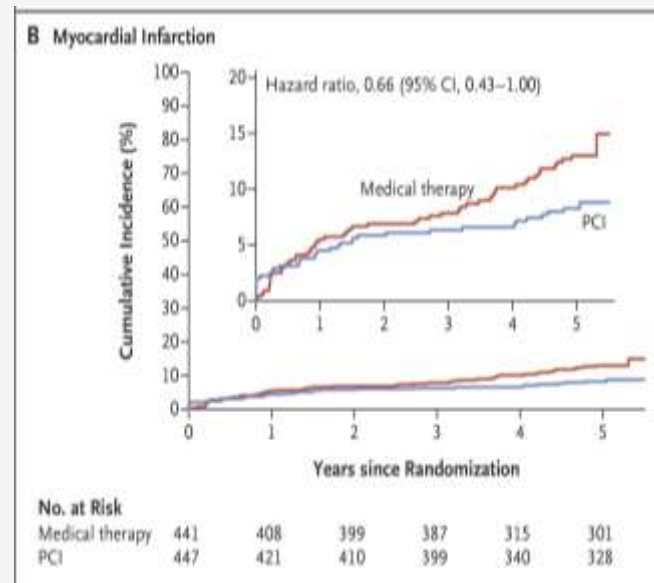
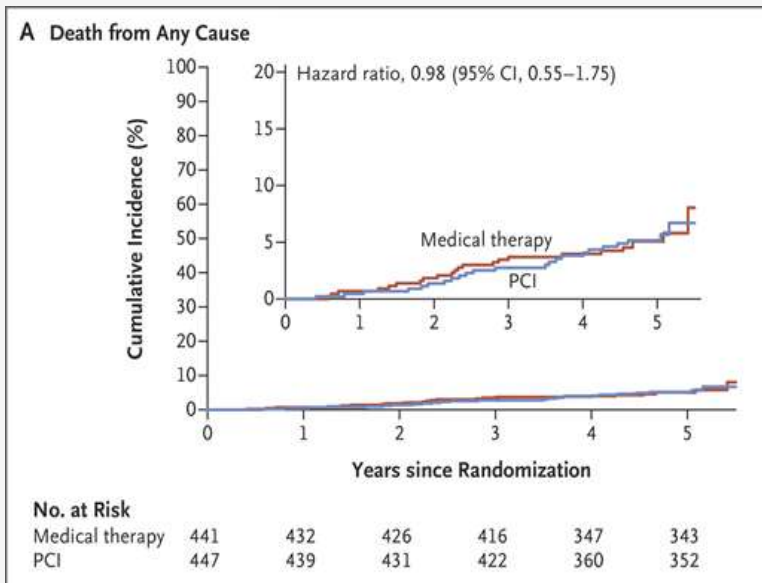
FAME 2



- MACE (death, MI, unplanned hospitalization leading to revascularization) at 24 months)

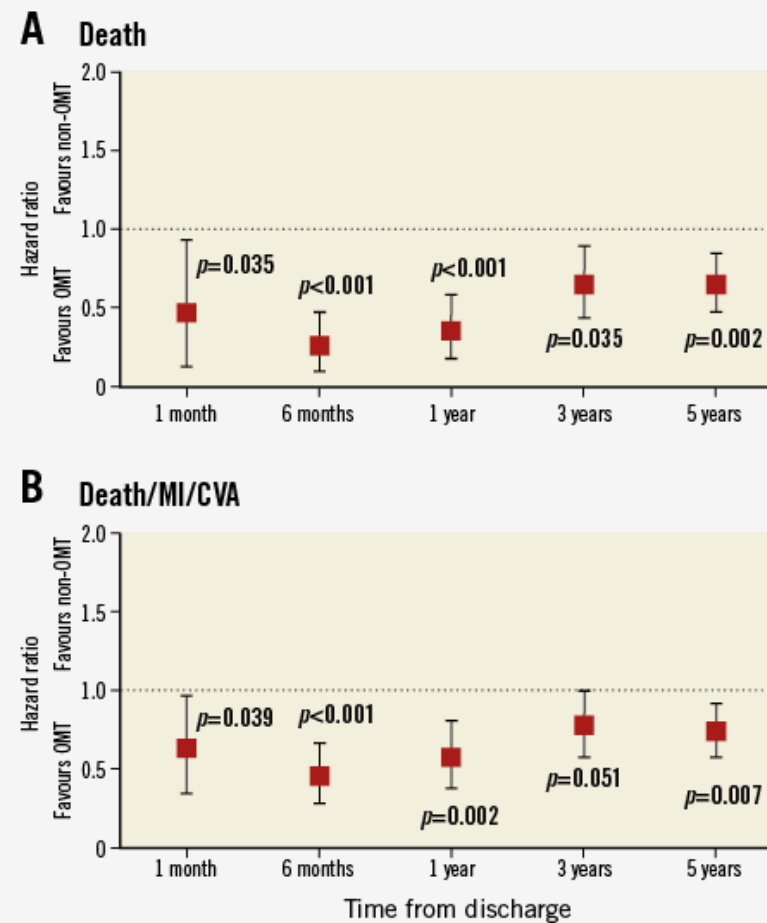


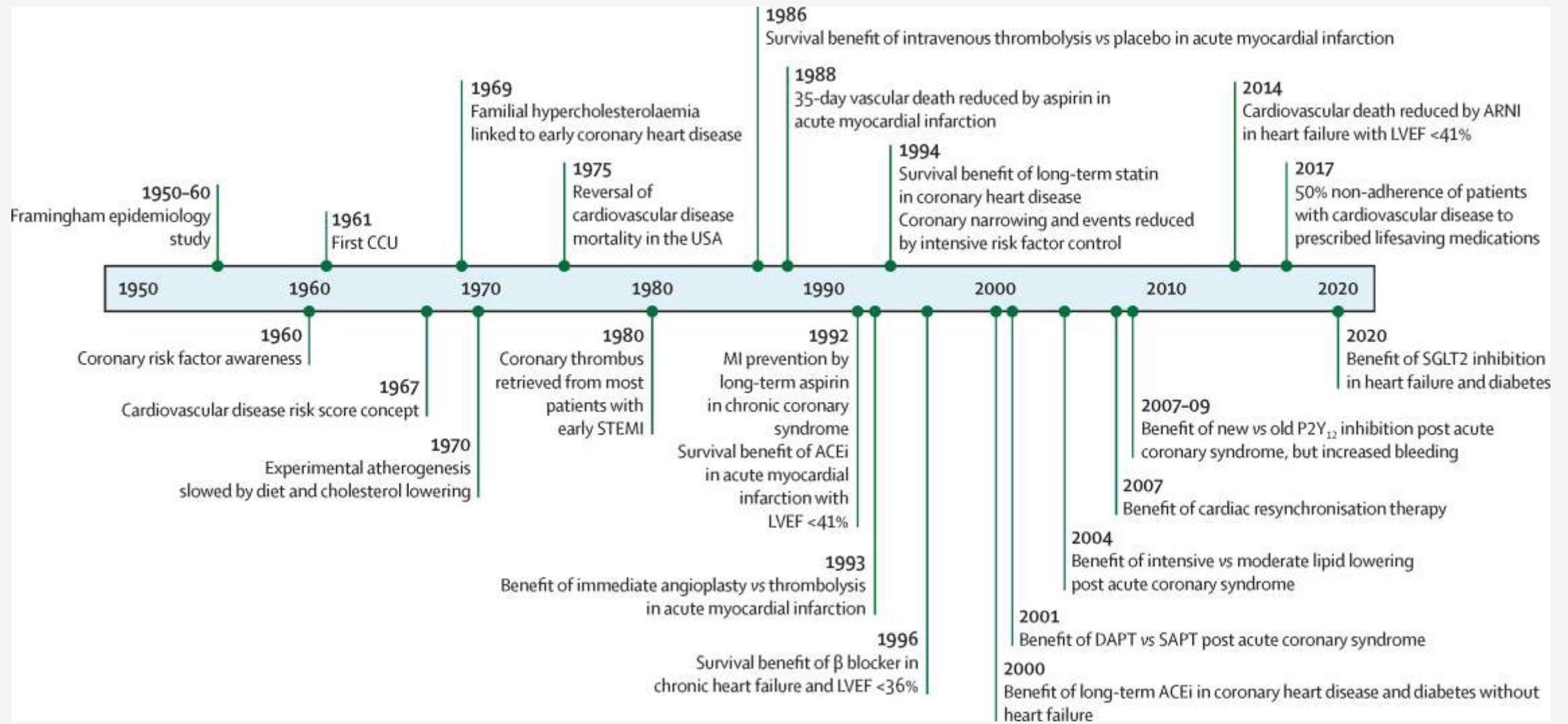
FAME 2



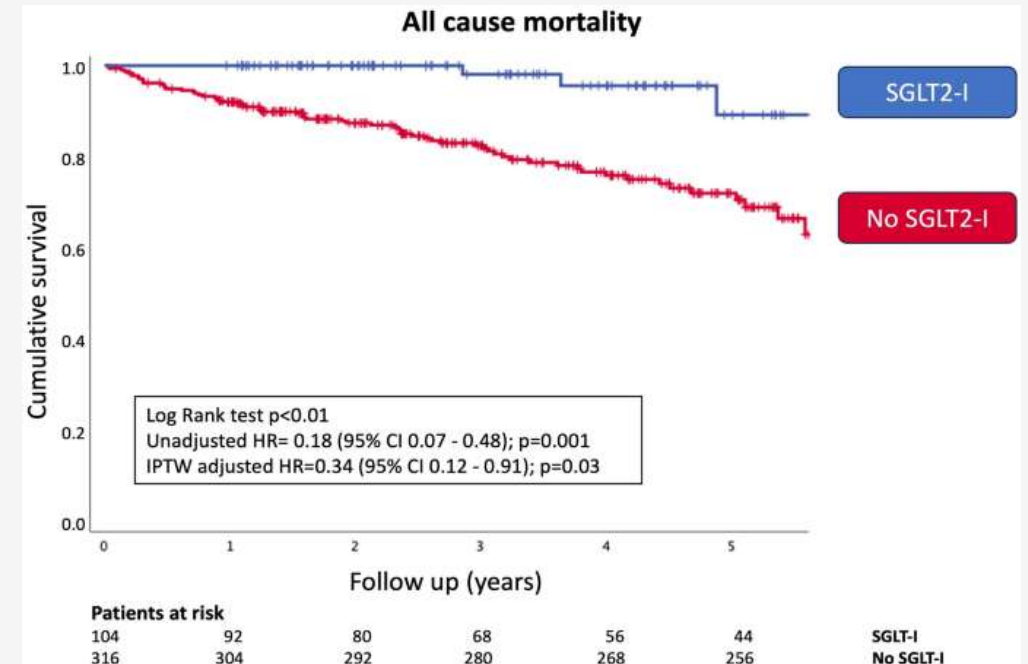
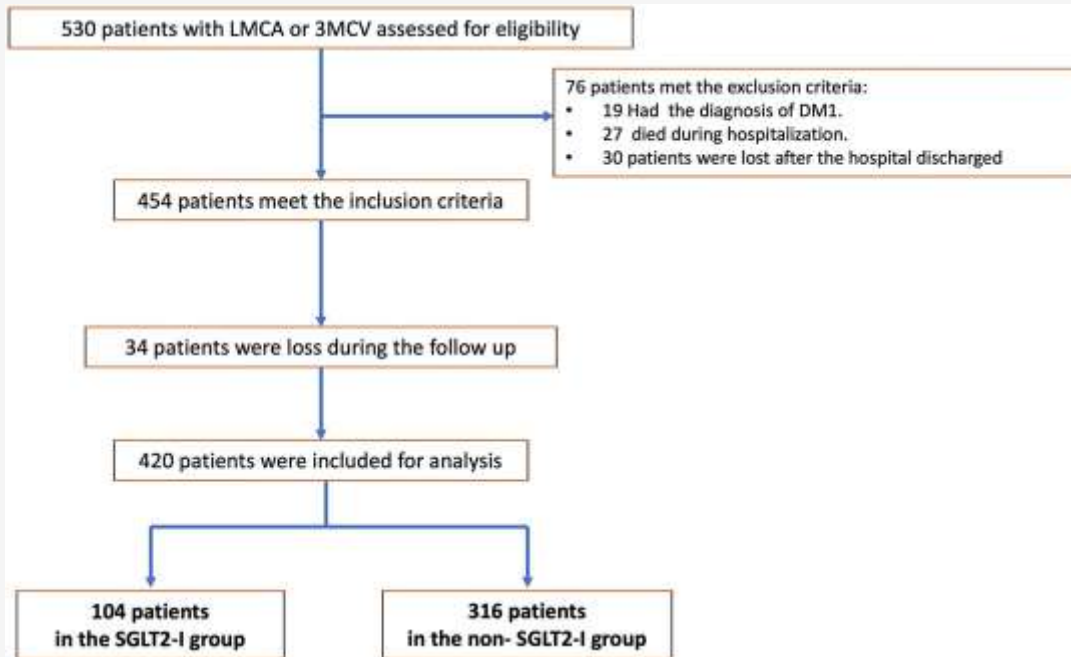
Revaskularizasiya sonrası OMT ?

- ✓ SYNTAX tədqiqatının *post hoc* analizi
- ✓ Evə yazılarkən 41% xəstə OMT alıb
- ✓ OMT – antiplatelet + bb + statin + ace inh
- ✓ 5 il sonra PCI qolunda bu göstərici 40%, CABG 36%

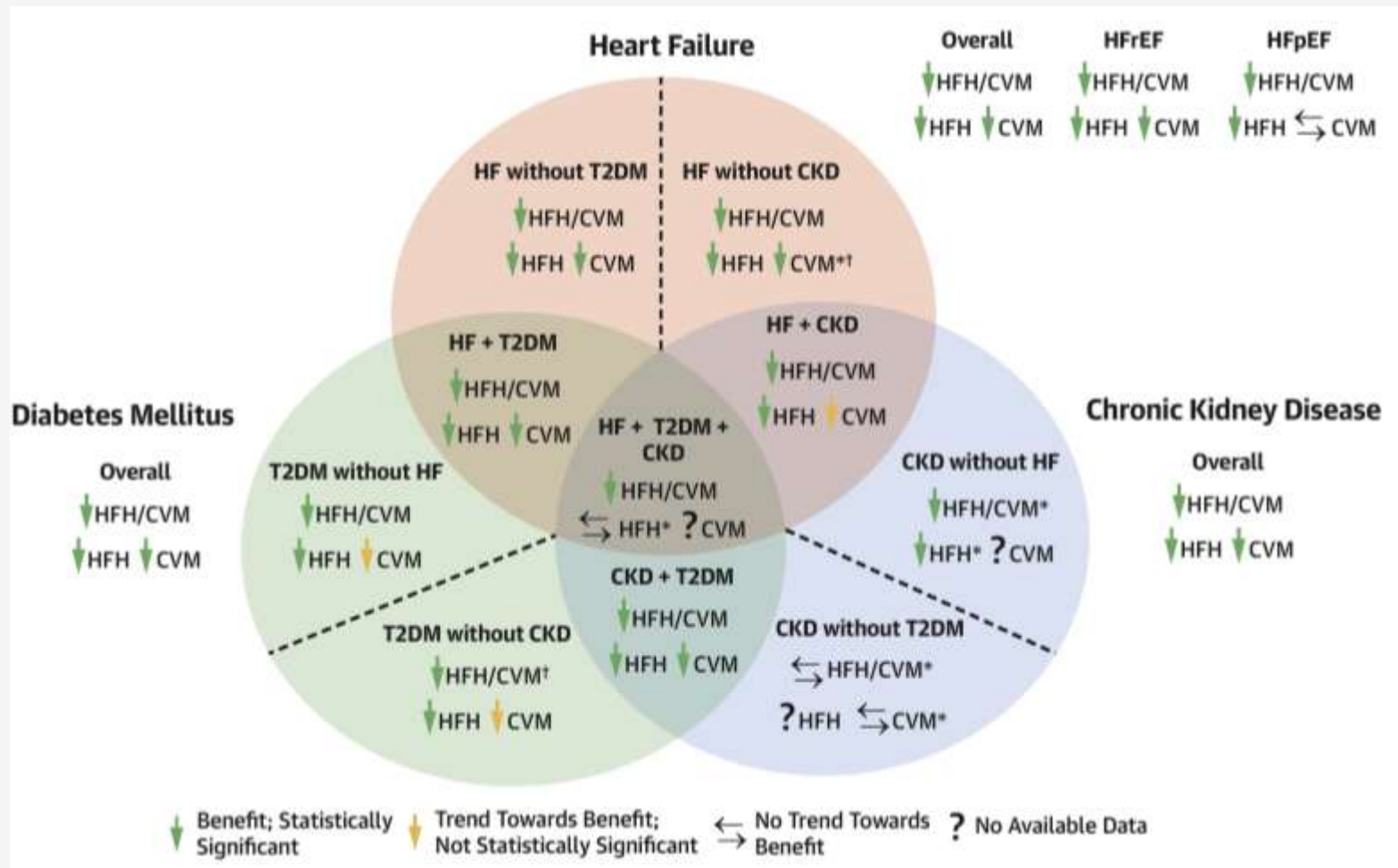




Treatment with SGLT2 Inhibitors in Patients with Diabetes Mellitus and Extensive Coronary Artery Disease: Mortality and Cardiovascular Outcomes



| | All patients n = 420 | SGLT2-I n = 104 | Non-SGLT2-I n = 316 | p |
|-----------------------------------|-------------------------|--------------------|------------------------|-------|
| Clinical presentation | | | | |
| ACS | 44.3% | 47.1% | 43.4% | 0.503 |
| CCS | 55.7% | 52.9% | 56.6% | |
| Revascularization strategy | | | | |
| PCI + OMT | 48.3% | 37.3% | 50.6% | 0.193 |
| CABG + OMT | 39.8% | 47.1% | 37.3% | |
| OMT only | 11.9% | 11.5% | 12% | |
| Complete revascularization (yes) | 28.8% | 31.7% | 27.8% | 0.448 |



SELECT tədqiqatı

Semaglutide effects on cardiovascular outcomes in people with overweight or obesity

SELECT Cardiovascular Outcomes Trial

Semaglutide 2.4 mg



SELECT is a Phase 3b, multinational, randomized, double-blind, placebo controlled, event-driven superiority trial that evaluated the effect of semaglutide 2.4 mg vs placebo, both in addition to SoC for CVD, in patients with established CVD and overweight or obesity on CV outcomes.¹

Key Eligibility Criteria²



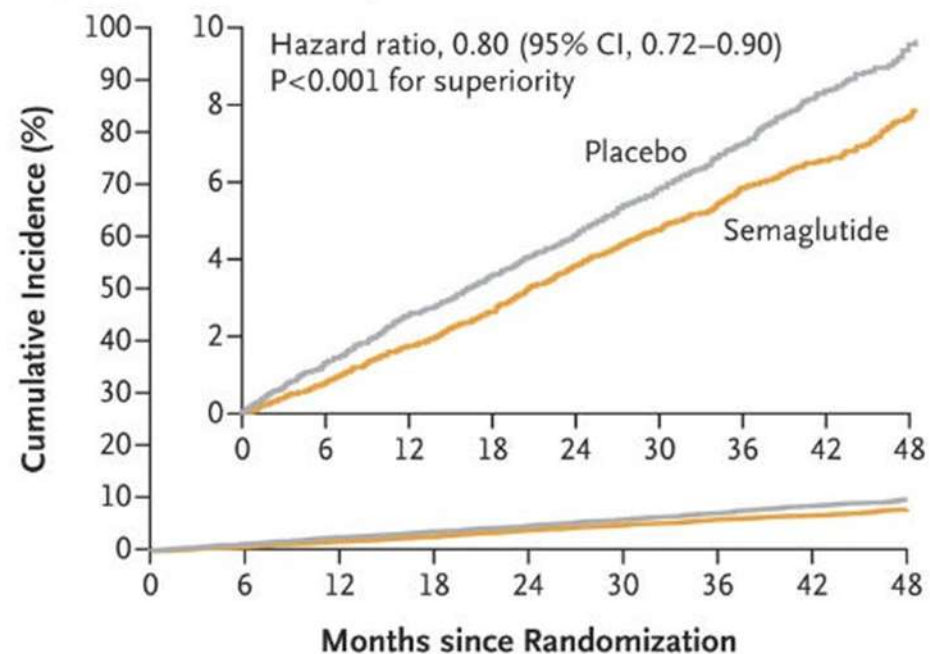
- Adults (age ≥ 45 years) with a BMI ≥ 27 kg/m²
- A1C $< 6.5\%$, and no history of type 1 or 2 diabetes
- Established CVD, defined as prior MI, prior stroke (ischemic or hemorrhagic) or symptomatic PAD (as evidenced by ≥ 1 of the following; intermittent claudication with ABI < 0.85 at rest, history of peripheral arterial revascularization or amputation due to atherosclerotic disease)
- Key exclusion criteria: history of either type 1 or type 2 diabetes, A1C $\geq 6.5\%$ at screening or treatment with glucose-lowering agents or GLP-1 RAs within previous 90 days, NYHA Class IV HF, a CV or neurological event within previous 60 days, or ESRD or hemodialysis (chronic or intermittent) or peritoneal dialysis.

Study Design²



Patients were randomized (1:1) to semaglutide 2.4 mg or placebo, both in addition to SoC for CVD.

A Primary Cardiovascular Composite End Point



No. at Risk

| | | | | | | | | | |
|-------------|------|------|------|------|------|------|------|------|------|
| Placebo | 8801 | 8652 | 8487 | 8326 | 8164 | 7101 | 5660 | 4015 | 1672 |
| Semaglutide | 8803 | 8695 | 8561 | 8427 | 8254 | 7229 | 5777 | 4126 | 1734 |