a

Acute MI ≤ 8 weeks (n=1400)
Inclusion/exclusion criteria
Informed consent
Randomization (1:1)
Baseline registration
Laboratory assessment, ECG, fat biopsy

n-3 PUFA
n = 700

Placebo
n = 700

3 month follow-up
Registration of clinical endpoints, laboratory assessment, food questionnaire.

12 month follow-up
Registration of clinical endpoints, atrial fibrillation registration device, laboratory assessment, ECG.

24 month follow-up
Registration of clinical endpoints, laboratory assessment, ECG, fat biopsy.

b

Enrollment

Assessed for eligibility (n=397)

Excluded (n=282)
- Not meeting inclusion criteria (n=52)
- Declined to participate (n=218)
- Other reasons (n=12)

Randomized (n=115)

Allocation

Allocated to intervention (n=58)
- Received allocated intervention (n=58)

Allocated to intervention (n=57)
- Received allocated intervention (n=57)

Follow-Up

Lost to follow-up (give reasons) (n=0)

Lost to follow-up (give reasons) (n=0)

Analysis

Trial not completed.

Trial not completed.