

# dal-Outcomes

**Principal Investigator:**

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**Sponsor:**

F. Hoffmann-La Roche Ltd.

**Study Purpose:**

NC20971RG: Roche Clinical Repository in association with protocol NC20971

**Objective:**

To obtain a single blood sample from consenting subjects enrolled in associated study NC20971 for pharmacogenetic and genetic research analysis.

**Inclusion Criteria:**

- adult patients,  $\geq 45$  years of age;
- recently hospitalized for ACS;
- clinically stable;
- receiving evidence-based medical and dietary management of dyslipidemia.

**Exclusion Criteria:**

- uncontrolled diabetes;
- clinically unstable;
- severe anemia;
- uncontrolled hypertension;
- concomitant treatment with any other drug raising HDL-C (eg niacin, fibrates).

**Status:**

Active enrollment

If you have any questions, please feel free to contact the coordinators, and they will be happy to answer any questions you have regarding this study.