

When to start anticoagulation after acute ischemic stroke

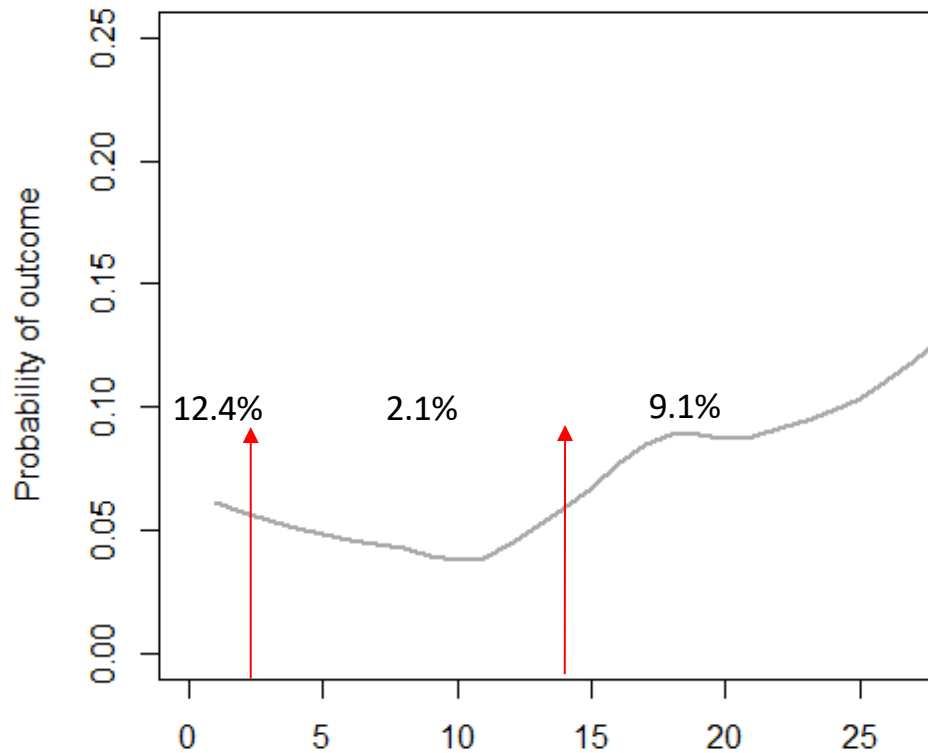
Valeria Caso

Stroke Unit

Ospedale Santa Maria della Misericordia, Perugia

RAF NOACs study: Outcome events (ischemic and hemorrhagic) depending on the time between onset and initiation of therapy with NOACs.

- 1127 patients with acute stroke and AF
- 32 (2.8%) ischemic recurrences (IS –TIA – SE) at 90 days
- 18 (1.6%) symptomatic ICH



Risk of combined outcome events based upon the day of initiating NOAC



ORIGINAL RESEARCH ARTICLE



Early Versus Delayed Non–Vitamin K Antagonist
Oral Anticoagulant Therapy After Acute Ischemic
Stroke in Atrial Fibrillation (TIMING):
A Registry-Based Randomized Controlled
Noninferiority Study

Jonas Oldgren¹, MD, PhD*; Signild Åsberg², MD, PhD*; Ziad Hijazi³, MD, PhD; Per Wester⁴, MD, PhD; Maria Bertilsson, MSc;
Bo Norving⁵, MD, PhD; for the National TIMING Collaborators

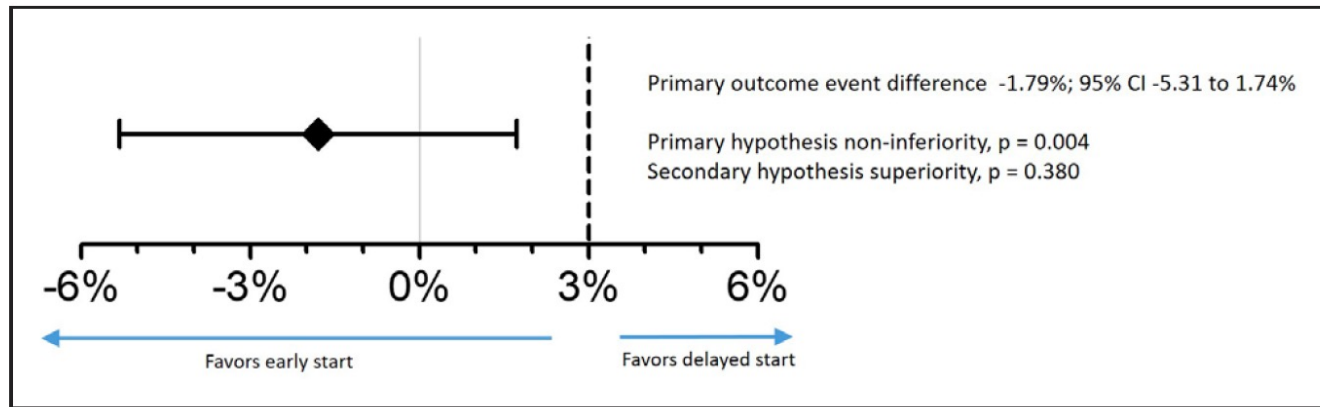


Figure 2. Risk difference in the primary composite outcome for early vs delayed initiation of NOAC at 90 days.

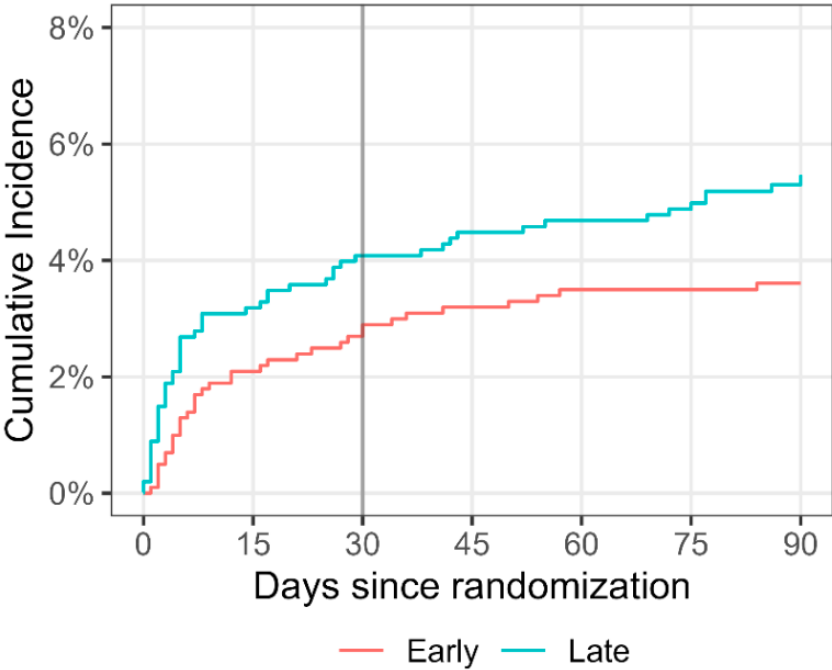
Primary outcome was a composite of ischemic stroke, symptomatic intracerebral hemorrhage, or all-cause mortality. Primary hypothesis testing for noninferiority at an absolute 3% margin, and secondary hypothesis testing for superiority. NOAC indicates non-vitamin K antagonist oral anticoagulant.

ORIGINAL ARTICLE

Early versus Later Anticoagulation for Stroke with Atrial Fibrillation

U. Fischer, M. Koga, D. Strbian, M. Branca, S. Abend, S. Trelle, M. Paciaroni, G. Thomalla, P. Michel, K. Nedeltchev, L.H. Bonati, G. Ntaios, T. Gattlinger, E.-C. Sandset, P. Kelly, R. Lemmens, P.N. Sylaja, D. Aguiar de Sousa, N.M. Bornstein, Z. Gdovinova, T. Yoshimoto, M. Tiainen, H. Thomas, M. Krishnan, G.C. Shim, C. Gumbinger, J. Vehoff, L. Zhang, K. Matsuzono, E. Kristoffersen, P. Desfontaines, P. Vanacker, A. Alonso, Y. Yakushiji, C. Kulyk, D. Hemelsoet, S. Poli, A. Paiva Nunes, N. Caracciolo, P. Slade, J. Demeestere, A. Salerno, M. Kneihsl, T. Kahles, D. Giudici, K. Tanaka, S. Rätty, R. Hidalgo, D.J. Werring, M. Gödlin, M. Arnold, C. Ferrari, S. Beyeler, C. Fung, B.J. Weder, T. Tatlisumak, S. Fenzl, B. Rezny-Kasprzak, A. Hakim, G. Salanti, C. Bassetti, J. Gralla, D.J. Seiffge, T. Horvath, and J. Dawson, for the ELAN Investigators*

Primary outcome at 30 and 90 days



At Risk

Early	1006	975	957	946	940	933	583
Late	1007	969	949	938	929	919	558