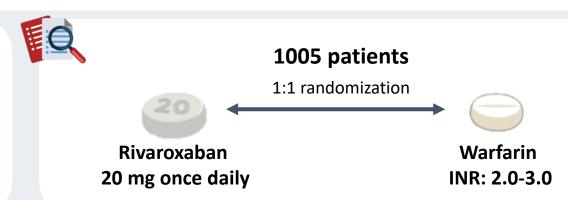
Rivaroxaban in Patients with Atrial Fibrillation and a Bioprosthetic Mitral Valve: The RIVER Trial

Background: The effects of rivaroxaban in patients with atrial fibrillation and a bioprosthetic mitral valve remain uncertain. RIVER trial was conducted to assess the efficacy and safety of rivaroxaban as compared with warfarin in patients with AF and a bioprosthetic mitral valve.



- Multicenter, open-label, randomized, noninferiority trial in Brazil
- Patients with AF and a bioprosthetic mitral valve were included



Mean time until primary-outcome:

Rivaroxaban group 347.5 days

P<0.001 for noninferiority

Warfarin group 340.1 days

The primary outcome: Composite of death, major cardiovascular events or major bleeding at 12 months

Major bleeding events:

Rivaroxaban group 7 patients (1.4%)

Warfarin group 13 patients (2.6%)

HR, 0.54; 95% CI, 0.21 to 1.35

In patients with atrial fibrillation and a bioprosthetic mitral valve, rivaroxaban was noninferior to warfarin with respect to the mean time until the primary outcome of death, major cardiovascular events, or major bleeding at 12 months.

