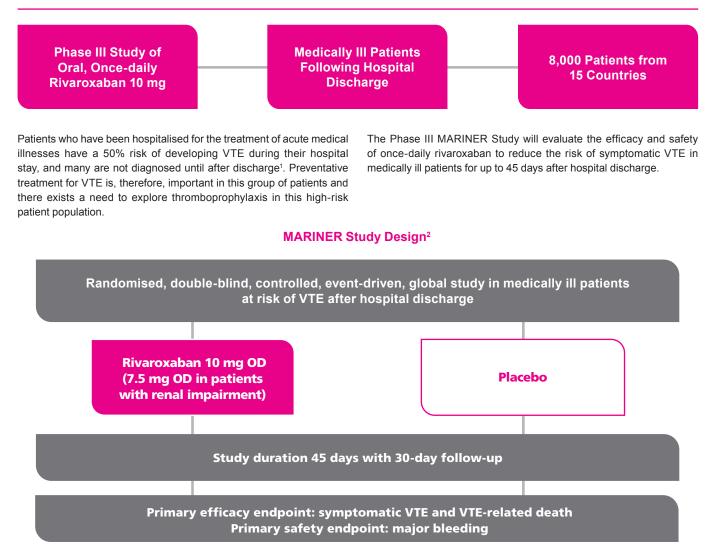
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PHASE III

MARINER Will Provide Insights into the Potential Benefit of Once-daily Rivaroxaban in Protecting Medically III Patients from Venous Thromboembolism (VTE) Following Hospital Discharge



The extensive evaluation of rivaroxaban to protect different patient populations at risk of venous and arterial thromboembolism (VAT), makes it the most studied novel OAC in the world. Rivaroxaban (Xarelto[®]) is already approved for five indications in seven areas of use and its investigation - both completed and ongoing - will include more than 275,000 patients in clinical trials and real world settings.



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About the Study Patient Population

Key inclusion criteria:

Patients must be 40 years and older, with an increased risk of VTE and must have been hospitalised for at least three consecutive days and up to 14 days for an acute medical condition such as, but not limited to:

- Congestive heart failure
- Acute respiratory insufficiency
- Acute exacerbation of COPD
- Acute ischaemic stroke
- Acute infectious or inflammatory diseases

Key exclusion criteria:

- Serious bleeding or severe head trauma within the last three months prior to entering the study
- Serious trauma within four weeks prior to entering the study
- History of haemorrhagic stroke
- Any medical condition that requires chronic use of anticoagulation

For more details, please visit www.clinicaltrials.gov

References

1) Streiff MB, Brady JP, Grant AM, et al. CDC Grand Rounds: Preventing Hospital-Associated Venous Thromboembolism. Centers for Disease Control and Prevention. March 7, 2014; 63(09);190-193. 2) Clinical Trials.gov. A Study of Rivaroxaban (JNJ-39039039) on the Venous Thromboembolic Risk in Post-Hospital Discharge Patients (MARINER). Available at: http://clinicaltrials.gov/ct2/show/NCT02111564?term=mariner&rank=1. Accessed July 2014.





PHASE III