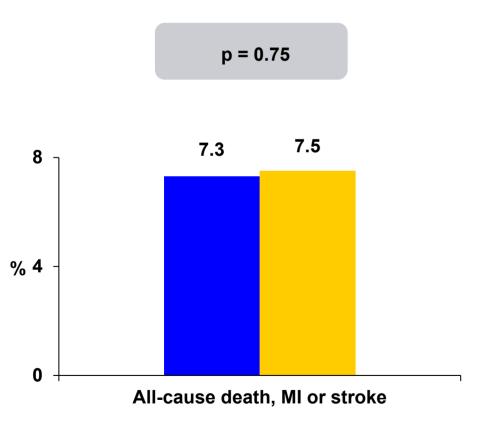
ADAPTABLE #ACC21



Trial Description: Patients with ASCVD were randomized to aspirin 81 mg vs. aspirin 325 mg.



RESULTS

- Primary effectiveness outcome of all-cause death, MI, or stroke at 12 months: 7.3% of the aspirin 81 mg group vs. 7.5% of the aspirin 325 mg group (p = 0.75)
- Primary safety outcome of major bleeding requiring blood transfusion at 12 months:
 0.6% of the aspirin 81 mg group vs. 0.6% of the aspirin 325 mg group (p = 0.41)

CONCLUSIONS

- Among patients with ASCVD, aspirin 325 mg was not superior compared with 81 mg
- Aspirin 325 mg was not associated with a reduction in all-cause death, MI, or stroke compared with 81 mg daily
- Major bleeding was similar between the treatment groups

Aspirin 81 mg (n = 7,540) Aspirin 325 mg (n = 7,536)

Jones WS, et al. *N Engl J Med* 2021;384(21):1981-1990