Aspirin for venous thromboembolism prevention and treatment: a renewal?

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PMID : 23123525

Available at: http://archive-ouverte.unige.ch/unige:26367

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Aspirin for venous thromboembolism prevention and treatment: a renewal?

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Over the past months, aspirin has suddenly moved from the backstage to the front of the scene of venous thromboembolism. Indeed, two features raised interest of the medical community. First, in 2012, the American College of Chest Physicians (ACCP) evidence-based clinical practice guidelines changed its former strong recommendation (Grade 1A) against use of aspirin for preventing venous thromboembolism after major orthopedic surgery for a conditional Grade 1B recommendation in favor of its use in this particular indication. Second, The New England Journal of Medicine has recently published the results of the Warfarin and Aspirin (WARFASA) study that convincingly demonstrated the efficacy of low-dose aspirin for preventing the long-term recurrence of venous thromboembolism in patients who received a course of 6 to 18 months of oral anticoagulant treatment for the initial event. These features are in line with recent basic science data that emphasize the role of platelets in the pathogenesis of venous thrombosis. Indeed, the histological inspection of venous thrombi shows that cells such as red blood cells and leukocytes are in close vicinity of platelets. Of note, neutrophils stimulated with proinflammatory agents or activated platelets release their nuclear content (including histones) that form web-like structures designated as neutrophil extracellular traps, which in turn promote the generation of thrombin through platelet-dependent mechanisms as well as vein thrombi, at least in a murine model. Taken together, these findings provide a rationale for clinical studies addressing the issue of the prevention of venous thrombosis by platelet function inhibitors, particularly aspirin.

However, both the new ACCP recommendation about thromboprophylaxis in major orthopedic surgery with the suggestion of aspirin as an alternative to anticoagulants, and the superiority of aspirin over placebo for secondary prophylaxis after a first venous thromboembolic event are essentially each based on a single study. In 2000, the Pulmonary Embolism Prevention (PEP) investigators reported a significant risk reduction with aspirin 160 mg daily compared with placebo for symptomatic venous thromboembolism in a very large population undergoing surgery for hip fractures or elective hip arthroplasty. In the WARFASA multicenter, investigator-initiated, double-blind study, Becattini et al. concluded that aspirin is efficacious and safe for secondary long-term prophylaxis of venous thromboembolism in patients who had experienced a first unprovoked deep vein thrombosis (DVT) or pulmonary embolism.

Let us first consider the issue of aspirin for thromboprophylaxis in major orthopedic surgery. As early as 1994, the Antiplatelet Trialists’ Collaboration meta-analysis pooled data from 62 randomized studies totaling over 8000 patients and concluded that aspirin (various dosages) reduced the frequency of both pulmonary embolism (by two-thirds) and DVT (by 40%). However, the methodological quality of many studies included in the meta-analysis was poor, and diagnostic methods were far from optimal, which resulted in mixed opinions on its results. Therefore, the PEP trial was launched in the mid-1990s to test aspirin (160 mg daily for 35 days) vs. placebo in 17,444 patients with hip fracture or elective hip arthroplasty. The study had several methodological strengths, including a central randomization, a double-blind design, an independent blinded committee for adjudicating objectively confirmed endpoints, and a follow-up that was close to 100%. There was a statistically significant 28% but limited relative risk reduction in symptomatic DVT (relative risk [RR], 0.72; 95% confidence interval [CI], 0.53–0.96) with no beneficial effect on nonfatal pulmonary embolism, and no significant effect on overall mortality. The number needed to treat (NNT) to prevent one symptomatic DVT was 238 (P = 0.03). Despite the large sample size, major nonfatal bleeding was not significantly increased in patients given aspirin even...
ACCP consensus recommendations thus seems 1.12; 95% CI, 0.94–1.34). Of note, a substan‑

6‑ to 18‑month period of anticoagulant treatment after an initial course of anticoagulant treatment

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Abbreviations: CI – confidence interval, INR – international normalized ratio, NA – not applicable, NNH – number needed to harm, NNT – number needed to treat, RR – relative risk, RRR – relative risk reduction (compared with placebo), VTE – venous thromboembolism

though there was a trend in this direction (RR, 1.12; 95% CI, 0.94–1.34). Of note, a substantial proportion of the PEP population also received some sort of heparin, which may have offset the effect of aspirin. Altogether, the PEP trial confirmed a statistically significant but small thromboprophylactic effect of aspirin in major orthopedic surgery (mainly hip fracture) compared with placebo. Based on this evidence, the 8th ACCP consensus1 admitted that aspirin provides some protection against venous thromboembolism, but did not recommend its use alone as prophylaxis primarily because more effective methods are readily available, e.g., low‑dose unfractionated heparin and low‑molecular‑weight heparins (LMWH). This nuanced appreciation was, however, translated into a strong recommendation against aspirin. In turn, the 9th edition of the ACCP consensus concentrated their analysis on the so called patient‑important (say symptomatic) outcomes and, therefore, rehabilitated aspirin as one prophylactic agent among others for patients undergoing major orthopedic surgery. However, the panelists state in the text that indirect evidence from trials of LMWH and aspirin against placebo also shows greater relative efficacy of LMWH and, in the end, suggest the use of LMWH in preference to the other options including aspirin. The controversy raised around the contradiction between the 8th and the 9th ACCP consensus recommendations thus seems to be largely artificial.

Let us now consider the issue of aspirin for long‑term secondary prophylaxis after an initial 6‑ to 18‑month period of anticoagulant treatment in patients with a first idiopathic DVT and/or pulmonary embolism, as tested in the WARFASA study. In this trial, 205 patients were randomized to receive aspirin 100 mg daily and 197 patients were given placebo. Twenty‑eight (6.6% per year) and 43 (11.2% per year) symptomatic venous thromboembolic events were adjudicated in the two groups, respectively, during a median follow‑up of 2 years, resulting in a hazard ratio of 0.58 (95% CI, 0.36–0.93) (P = 0.02). Only 1 patient in each group experienced a major bleeding episode.

These results need to be viewed in the perspective of other large‑scale, randomized controlled studies of long‑term secondary prevention of venous thromboembolism recurrence – the Prevention of Recurrent Venous Thromboembolism (PREVENT) study, the EINSTEIN Extension study, and the Extended Low‑Intensity Anticoagulation for Thromboembolism (ELATE) study. Both PREVENT and EINSTEIN Extension were placebo‑controlled trials while ELATE compared 2 intensities of warfarin therapy (TABLE). In all these studies, the main efficacy outcome was recurrent symptomatic, objectively confirmed DVT or pulmonary embolism. The active regimen in PREVENT and EINSTEIN Extension was warfarin with a target international normalized ratio (INR) of 1.5–2.0 and rivaroxaban (20 mg once daily, p.o), respectively. While the relative risk reduction of venous thromboembolism recurrence was 64% (95% CI, 19%–67%) and 82% (95% CI, 61%–91%), respectively, in these 2 trials, it was only 42% in WARFASA with a large CI (95% CI, 7%–64%). Even though the CIs in the 3 studies largely
Aspirin is a commonly used medication worldwide, especially in primary and secondary prevention of cardiovascular disease. However, its role in the prevention of venous thromboembolism (VTE) has been a subject of debate. A recent editorial in the *New England Journal of Medicine* discusses the ongoing placebo-controlled Aspirin to Prevent Recurrent Venous Thromboembolism (ASPIRE) study versus the Preventing Events That preceded Recurrence with Extended Use of Low-Intensity Warfarin (PREVENT) and EINSTEIN Extension studies, which included patients with first idiopathic venous thromboembolism.

The editorial highlights the challenges in comparing results from different studies due to variations in study populations, study design, and endpoints. Despite these differences, the authors argue that aspirin remains a viable option for the prevention of VTE, especially in high-risk patients. They recommend that practitioners consider aspirin as a viable prophylactic option based on the available evidence and the physician's judgment of the patient's risk.

**References**