**Men had greater risk of recurrent venous thromboembolism than women**


Clinical impact rating GP/FP/Primary care ★★★★★☆ IM/Ambulatory care ★★★★★☆ Haematology ★★★★★☆

![Question](https://example.com/question.png)

**Does the risk of recurrent venous thromboembolism (VTE) differ between men and women?**

**METHODS**

**Design:** inception cohort followed for a median of 26 months (Austrian Study on Recurrent Venous Thromboembolism).

**Setting:** 4 thrombosis centres in Vienna.

**Patients:** 826 patients ≥18 years of age (mean age 48 y, 55% women) who had a first episode of VTE and who had been treated with oral anticoagulants for ≥3 months. Patients who had received standard heparin or doses designed to keep the activated partial thromboplastin time 1.5–2.0 times that of the control value or had received subcutaneous low molecular weight heparin at therapeutic doses. Exclusion criteria: surgery, trauma, or pregnancy in the previous 3 months; known deficiency of antithrombin, protein C, or protein S; lupus anticoagulant; cancer; or need for long term treatment with antithrombotic drugs. Women were discouraged from using oral contraceptives or hormone therapy.

**Prognostic factors:** sex; age; presence or absence of symptomatic pulmonary embolism at the time of first VTE; duration of anticoagulation; and presence or absence of factor V Leiden, factor II G20210A, and elevated concentrations of factors VIII and IX.

**Outcomes:** recurrence of symptomatic VTE. The diagnosis of deep venous thrombosis was confirmed by venography or colour coded duplex sonography. The diagnosis of pulmonary embolism was confirmed by ventilation perfusion lung scanning.

For correspondence: Dr P A Kyrle, Ludwig Boltzmann-Institut für Thromboseforschung, Vienna, Austria. paul.kyrle@meduniwien.ac.at
Sources of funding: Jubilaeumsfonds of the österreichische Nationalbank.

**MAIN RESULTS**

102 patients (12%) had recurrent VTE: 74 men (73%) and 28 women (27%). Men had a greater risk of recurrence than did women (5 y cumulative recurrence 30.7% v 8.5%) (table). For men and women, risk of recurrent VTE was associated with an elevated concentration of factor VIII (≥234 IU/dl [23.4 U/l]), presence of factor II G20210A, and having symptomatic pulmonary embolism or deep venous thrombosis as the first VTE event (table).

**CONCLUSION**

After a first episode of venous thromboembolism, men were at greater risk than women of a recurrent episode.

Abstract and commentary also appear in ACP Journal Club.

**Commentary**

Kyrle et al compared the incidence of recurrent VTE among men and women with incident unprovoked VTE (eg, no previous surgery or trauma) after a mean of 8–9 months of oral anticoagulant therapy. After excluding patients with high risk thrombophilic disorders (deficiencies of protein C, protein S, and antithrombin, or presence of lupus anticoagulant), they found that men with unprovoked VTE had a ≥3 fold higher incidence of recurrence than women. This result was surprising and unforeseen and did not appear to occur because women with unprovoked VTE taking oral contraceptives or hormone therapy simply stopped taking these hormones. In a stratified multivariate analysis, women with incident VTE who were not taking any hormones still had a 3 fold lower incidence of recurrent VTE than men.

Why do women have such a strikingly lower incidence of VTE when the incidence of first time VTE is the same between men and women? Possibilities include the following: (1) Men may not lyse a thrombus as readily as women, leading to more residual changes in the veins and a higher recurrence rate; (2) men might develop more comorbid conditions after the initial VTE event (eg, more hospital admissions or worse heart failure); and (3) men might have more underlying occult malignancies. The finding may also be due to chance or bias. This last possibility seems unlikely because another large study also found that women with unprovoked VTE had a significantly lower incidence of recurrent VTE than men [Baglin T. Personal communication], although this finding was not stated in the report of the study.

A pressing need exists for prospective studies that more precisely define the risk factors associated with recurrent VTE, particularly among men and women with unprovoked VTE. The ultimate goal is to determine which patients receive more benefit than harm when prescribed long term anticoagulant therapy. Based on this study, it is still not clear exactly which women and which men are at such high risk of recurrent thromboembolism that they are appropriate candidates for extended anticoagulant treatment.

Richard H White, MD
University of California, Davis
Sacramento, California, USA


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**Prognostic factors for recurrent venous thromboembolism**

<table>
<thead>
<tr>
<th>Prognostic factors</th>
<th>Relative risk (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Male sex (v female sex)</td>
<td>3.6 (2.3 to 5.5)</td>
</tr>
<tr>
<td>Age (per 10 y increase)</td>
<td>1.1 (0.9 to 1.3)†</td>
</tr>
<tr>
<td>Symptomatic pulmonary embolism (v deep venous thrombosis)</td>
<td>1.7 (1.1 to 2.5)</td>
</tr>
<tr>
<td>Factor V Leiden (v absence)</td>
<td>1.2 (0.8 to 1.8)†</td>
</tr>
<tr>
<td>Factor II G20210A (v absence)</td>
<td>2.1 (1.1 to 3.8)</td>
</tr>
<tr>
<td>Factor VIII (v ≥234 IU/dl [23.4 U/l])</td>
<td>2.9 (1.6 to 5.1)</td>
</tr>
<tr>
<td>Factor IX (v ≥138 IU/dl [13.8 U/l])</td>
<td>1.0 (0.8 to 1.2)†</td>
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</tbody>
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<tr>
<th>Prognostic factors</th>
<th>Relative risk (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Duration of anticoagulation (per 3 mo increase)</td>
<td>1.0 (0.98 to 1.05)†</td>
</tr>
</tbody>
</table>

*Relative risks were adjusted for age, presence or absence of a first symptomatic pulmonary embolism, factor V Leiden, factor II G20210A, elevated factor VIII or factor IX concentrations, and duration of anticoagulation.
†Not significant.
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*Evid Based Med* 2004 9: 187
doi: 10.1136/ebm.9.6.187

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