Review: thrombolytic treatment does not reduce the risk of recurrent pulmonary embolism and death more than heparin


Clinical impact ratings IM/Ambulatory care ★★★★★★ Internal medicine ★★★★★★ Emergency Medicine★★★★★★ Respirology ★★★★★ Critical Care ★★★★★ Haematology ★★★★★ Cardiology ★★★★★

In patients with acute pulmonary embolism (PE), is initial treatment with thrombolysis more effective than heparin for reducing risk of recurrent PE and death?

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METHODS

Data sources: Medline and EMBASE/Excerpta Medica (January 1980 to January 2003), Cochrane Library (2003, Issue 1), and hand searches of reference lists of retrieved articles and abstracts of conference proceedings.

Study selection and assessment: published and unpublished randomised controlled trials (RCTs) that were properly randomised, compared thrombolysis with heparin for initial treatment of patients with objectively diagnosed symptomatic PE, and used objective methods to assess clinical outcomes. 2 independent reviewers assessed methodological quality (allocation sequence and concealment, blinding, and follow up).

Outcomes: a composite of recurrent PE or death. Secondary outcomes were PE, death, major bleeding, non-major bleeding, and intracranial haemorrhage.

MAIN RESULTS

11 trials (n = 748) met the selection criteria. 3 RCTs used random number tables or programs for generating randomised allocation sequences, 5 RCTs provided information on concealment, 3 RCTs had blinding of patients and investigators, and no trial reported loss to follow up. All studies included patients with symptomatic PE, and 5 trials included patients with major PE (haemodynamic instability). Thrombolysis included urokinase, streptokinase, and tissue plasminogen activator. Meta-analysis (fixed effects model) of 11 trials included patients with major PE (haemodynamic instability). The meta-analysis also underscored the importance of risk stratification for patients with acute PE. However, important questions remain. For example, how should we identify patients at high risk? We know that patients with right ventricular hypokinesis on echocardiography have increased mortality, but we do not know whether thrombolysis reduces their mortality. Further meta-analyses of existing data are unlikely to be helpful. A large clinical trial that enrols high risk patients with acute PE is needed to address the persistent controversy over the exact role of thrombolysis for these patients.

CONCLUSIONS

In unselected patients with acute pulmonary embolism (PE), thrombolysis does not reduce the risk of recurrent PE and death more than heparin. Thrombolysis increases the risk of non-major bleeding.

Commentary

The exact role of thrombolytic treatment in the management of acute PE remains controversial. Uncertainty over the balance between risk and benefit fuels this controversy. Thrombolytic agents accelerate the dissolution of venous thrombi, but they also increase the risk of serious bleeding. The authors of the 7th consensus statement of the American College of Chest Physicians recommend that clinicians not use systemic thrombolytic therapy for most patients with acute PE.1 The meta-analysis by Wan et al supports this position.

The principal result of the review is no surprise since 6 of 11 trials in the meta-analysis excluded patients who were haemodynamically unstable. A more telling observation lies in the subgroup analysis of 5 trials that did include haemodynamically unstable patients. This analysis suggests that thrombolysis reduces the likelihood of the composite outcome of recurrent PE or death. It does not show a survival benefit (odds ratio 0.47, 95% CI 0.20 to 1.10). However, the number of patients who were randomised was small (n = 254), and even fewer patients were haemodynamically unstable. Thus, the review by Wan et al has limited power to detect a survival benefit.

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Outcomes during hospital stay or at 30 days

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>Thrombolysis</th>
<th>Heparin</th>
<th>RRR (95% CI)</th>
<th>NNT (CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Recurrent PE or death</td>
<td>6.7%</td>
<td>9.6%</td>
<td>31% (–11 to 56)</td>
<td>Not significant</td>
</tr>
<tr>
<td>Recurrent PE</td>
<td>2.7%</td>
<td>4.3%</td>
<td>32% (–35 to 64)</td>
<td>Not significant</td>
</tr>
<tr>
<td>Death</td>
<td>4.3%</td>
<td>5.9%</td>
<td>29% (–28 to 62)</td>
<td>Not significant</td>
</tr>
<tr>
<td>Major bleeding</td>
<td>9.1%</td>
<td>6.1%</td>
<td>38% (–18 to 126)</td>
<td>Not significant</td>
</tr>
<tr>
<td>Non-major bleeding</td>
<td>23%</td>
<td>10%</td>
<td>140% (48 to 276)</td>
<td>13 (7 to 36)</td>
</tr>
<tr>
<td>Intracranial haemorrhage</td>
<td>0.5%</td>
<td>0.3%</td>
<td>4% (–64 to 202)</td>
<td>Not significant</td>
</tr>
</tbody>
</table>

*Abbreviations defined in glossary; RRR, RRI, NNT, NNH, and CI calculated from control event rate and odds ratio in article.