Intermittent pneumatic compression on top of pharmacological thromboprophylaxis in intensive care: added value or added cost?

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Venous thromboembolism (VTE) is a leading cause of in-hospital morbidity and mortality, particularly in patients at intensive care units (ICU) (1). Hypercoagulability, venous stasis and vascular injury, but also the frequent use of highly coagulable catheters and extracorporeal support devices predispose this patient group to thromboembolic complications. Therefore, efficient strategies for the prevention of thrombosis that do not cause excessive bleeding are paramount in critically ill patients.

In view of the multi-causal pathogenesis of venous thrombosis, both pharmacologic strategies with anticoagulant drugs and mechanical strategies with devices that decrease lower-limb venous stasis by blood displacing (2) reduce the rate of VTE and have been extensively evaluated in both surgical and non-surgical patients. An overview of current recommended strategies for in hospital thromboprophylaxis is provided in Table 1 and Figure 1.

According to the guidelines of the American College of Chest Physicians, pharmacologic thromboprophylaxis with low-molecular-weight-heparin (LMWH) or low-dose unfractionated heparin (UFH) is the recommended strategy in ICU-patients, even though the number of well-performed randomized clinical trials are limited (4). Mechanical alternatives, with IPC preferred over graduated compression stockings, are first choice in patients at very high risk for VTE who have contra-indications for prophylactic anticoagulant therapy (e.g., active bleeding or high risk for major bleeding) (5,6). Indeed, the difficult balance between bleeding and thrombosis is a daily struggle for physicians working at ICU-departments. Hence, institutional protocols regarding thromboprophylaxis should include both pharmacological and mechanical strategies, each with their own merits and risks (1,4,7,8).

Although pharmacologic thromboprophylaxis reduced the incidence of deep-vein thrombosis by 50% compared with no prophylaxis in critically ill patients, still 5% to 20% of this pharmacologically treated patient group develops deep-vein thrombosis (6,9,10). However, no previous clinical trials addressed whether IPC provides a meaningful benefit when added on top of anticoagulant prophylaxis therapy in critically ill patients. Vice versa, no clinical trials in this patient group evaluated whether pharmacological prophylaxis on top of IPC reduces clinically meaningful thrombotic events.

Therefore, Arabi et al. investigated if adjunctive IPC reduced the risk of VTE in critically ill patients receiving pharmacologic thromboprophylaxis (3). This is an important question because such combined strategy has been implemented in many critical care units in spite of absence of clinical data. Moreover, IPC immobilizes patients yet mobilizes precious resources at ICU (nursing time and cost of goods), and may cause some skin-injuries.

The PREVENT (The Pneumatic Compression for Preventing Venous Thromboembolism) trial was...
Table 1 Thromboprophylaxis according to VTE risk

<table>
<thead>
<tr>
<th>Inpatient baseline VTE risk</th>
<th>Pharmacological thromboprophylaxis</th>
<th>Mechanical thromboprophylaxis</th>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>Applicable to both surgical as well as nonsurgical patients</td>
<td>low-dose UFH or LMWH</td>
<td>IPC and/or GCS</td>
<td>European guidelines on perioperative venous thromboembolism prophylaxis; mechanical prophylaxis (1)</td>
</tr>
<tr>
<td>Low risk patients</td>
<td>No</td>
<td>No</td>
<td>In all, an individualized assessment should be performed regarding the thrombotic and (potential) bleeding risk. Early ambulation should be promoted actively</td>
</tr>
<tr>
<td>Non-high risk patients with a contra-indication for pharmacological thromboprophylaxis</td>
<td>No</td>
<td>No</td>
<td>No GCS in non-high risk (grade 2C)</td>
</tr>
<tr>
<td>Intermediate to high risk patients</td>
<td>Yes</td>
<td>No</td>
<td>Pharmacological prophylaxis should be considered as mainstay in a majority of inpatients and should be preferred over mechanical prophylaxis</td>
</tr>
<tr>
<td>Selected (very) high risk patients</td>
<td>Yes</td>
<td>Yes (add-on)</td>
<td>Given the results of PREVENT, this does not include routine ICU-patients (3). IPC might be preferred over GCS</td>
</tr>
<tr>
<td>High risk with contra-indication for pharmacological thromboprophylaxis</td>
<td>No</td>
<td>Yes, IPC preferred over GCS</td>
<td>Combination therapy (grade 2B); preference for IPC (grade 2B)</td>
</tr>
</tbody>
</table>

VTE, venous thromboembolism; UFH, unfractionated heparin; LMWH, low molecular weight heparin; IPC, intermittent pneumatic compression; GCS, graduated compression stockings.

In conclusion, the PREVENT-trial showed that the routine use of adjunctive IPC did not provide a clinically meaningful effect on the incidence of proximal DVT in critically ill patients at ICU (mean APACHE-II 20) who were concomitantly receiving pharmacologic thromboprophylaxis. The PREVENT-trial will help to refine international guidelines and institutional protocols on VTE thromboprophylaxis, especially in the money saving, time consuming and protocol driven environment of critical care (11-14).

an international, randomized controlled trial in a heterogeneous group of 2003 medical, surgical and trauma intensive care patients, although the latter only represented 8% of the study population. The trial compared IPC for at least 18 hours per day in addition to pharmacologic thromboprophylaxis (LMWH or UFH) with pharmacologic thromboprophylaxis alone. The trial was not double blind (for obvious reasons), but the risk of bias was mitigated because of its randomized design, high adherence to the assigned treatment (22 hours of IPC per day, median duration of 7 days), low incidence of concomitant use of compression stockings (0.9%) and minimal loss to follow-up (3%). Although the trial is underpowered as a result of the lower incidence of DVT in both groups (approximately one third of the predicted primary outcome events were achieved), the event rates in both groups were nearly identical (3.9% in the IPC-arm vs. 4.2% in the control; \( P=ns \)).
Figure 1 An overview of current recommended strategies for in hospital thromboprophylaxis each targeting a specific component of Virchow’s triad.

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None.

Footnote
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