Use of venous-thrombotic-embolic prophylaxis in patients undergoing surgery for renal tumors: a questionnaire survey in the Nordic countries (The NORENCA-2 study)

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Purpose: To examine the variation in venous thromboembolism prophylactic treatment (VTEP) among renal cancer patients undergoing surgery.

Materials and methods: An Internet-based questionnaire on renal tumor management before and after surgery was mailed to all Nordic departments of urology. The questions focused on the use of VTEP and were subdivided into different surgical modalities.

Results: Questionnaires were mailed to 91 institutions (response rate 53%). None of the centers used VTEP before surgery, unless the patient had a vena caval tumor thrombus. Overall, the VTEP utilized during hospitalization for patients undergoing renal surgery included early mobilization (45%), compression stockings (52%) and low-molecular-weight heparin (89%). In patients undergoing open radical Nx, 80% of institutions used VTEP during their hospitalization (23% compression stockings and 94% low-molecular-weight heparin). After leaving the hospital, the proportion and type of VTEP received varied considerably across institutions. The most common interval, used in 60% of the institutions, was for a period of 4 weeks. The restriction to the Nordic countries was a limitation and, therefore, may not reflect the practice patterns elsewhere. It is a survey study and, therefore, cannot measure the behaviors of those institutions that did not participate.

Conclusion: We found variation in the type and duration of VTEP use for each type of local intervention for renal cancer. These widely disparate variations in care strongly argue for the establishment of national and international guidelines regarding VTEP in renal surgery.

Keywords: venous-thrombotic-embolic prophylaxis kidney cancer, surgery, nephrectomy, mortality, complication, minimally invasive methods, thrombosis prophylaxis

Introduction

The development of venous thromboembolism (VTE) is due to an imbalance in hemostasis, which in Virchow’s triad is described as an interaction between the vessel wall, blood flow and blood composition.1 Prolonged immobilization or direct compression of the veins from a growing kidney tumor can give reduced flow in cancer patients and are well-known risk factors for VTE. Cancerous diseases can result in both direct and indirect effects on the vessel wall due to tumor invasion or increased inflammatory influence affecting both the vessel walls and the components of the coagulation cascade.

Cancer surgery is associated with a doubling of the risk of postoperative VTE compared with similar surgery for benign diseases.2 Patients with pancreatic or gastric cancer have the greatest risk, while pulmonary, bladder and gynecological cancers also...
have an increased risk.\(^3\) In a prospective cohort study, VTE was the most frequent cause of death within the first 30 days after cancer surgery for a variety of different malignancies and 40% of postoperative VTE occurred at least 21 days after surgical intervention.\(^4\) A high dose of low-molecular-weight heparin (LMWH; 5,000 vs 2,500 IU) results in significantly lower incidence of VTE in cancer surgery without an increased bleeding risk.\(^5\) A Cochrane review concluded that 4 weeks of thrombosis prophylaxis after major abdominal and pelvic surgery reduces the risk of postoperative VTE.\(^6\)

The five Nordic countries (Denmark, Finland, Iceland, Norway and Sweden) comprise 25 million people, and have quite similar treatment traditions and state-funded high-quality health care systems.

The Nordic Renal Cancer group is a newly formed working group with representatives from all the Nordic countries, with the aim of performing high-quality research on kidney cancer in the Nordic countries.\(^7\) The purpose of this study is to examine the use of VT prophylactic treatment (VTEP) among renal cancer patients undergoing curative therapy in the five Nordic countries prior to the publishing of the first European Association of Urology (EAU) Guidelines on thromboprophylaxis in urological surgery.\(^8\)

**Materials and methods**

**The study group**

The study was carried out by a collaborative group originating from the Scandinavian Association of Urology and is presented at the homepage of the Scandinavian Association of Urology (www.NUF.nu).

**The questionnaire**

A 21-question Internet-based questionnaire on renal tumor management before and after surgery was mailed to all Nordic departments of urology performing renal cancer surgery (Supplementary material). The questions addressed the use of VTEP and were subdivided into different surgical modalities: open radical and partial nephrectomy (Nx), laparoscopic/robotic Nx and laparoscopic and percutaneous thermal ablation of tumors.

In total, 91 institutions from all five countries were invited to participate. The questionnaires were mailed to the departments during late fall of 2016, and after two rounds of invitations, the database was closed on February 1, 2017. Six institutions did not perform renal surgery in 2016. We received responses from 45 of 85 hospitals performing any form of renal surgery (response rate 53%), as shown in Table 1. There were responses from 6 institutions from Denmark (75%), 12 from Finland (48%), 1 from Iceland (100%), 11 from Norway (73%) and 15 from Sweden (48%), as shown in Table 1.

The survey did not include patient-related data or affected patient treatment or care. Therefore, it did not require approval from the ethical committee. Completion of the survey was considered as informed consent from the participating departments.

**Results**

None of the 45 institutions used VTEP before surgery unless the patient had a vena caval tumor thrombus. Overall, the proportion of institutions that used VTEP during hospitalization for patients undergoing renal surgery included 45% using early mobilization, 52% using compression stockings and 89% using LMWH (Table 2). In patients undergoing open radical Nx, 80% of the institutions used VTEP during their hospitalization (23% compression stockings and 94% LMWH; Table 3). After leaving the hospital, the proportion and type of VTEP received varied considerably across institutions (5% for 1 week, 35% for 2 weeks and 60% for 4 weeks). For patients undergoing open partial Nx, 75% of institutions utilized VTEP (27% compression stockings and 91% LMWH; Table 3). After discharge, the VTEP regimen offered to patients was for 1 week in 12%, 2 weeks in 48% and 4 weeks in 40% of the institutions. For those who underwent laparoscopic/robotic partial Nx, 70% of the institutions utilized VTEP (29% compression stockings and 85% LMWH; Table 3). After hospital discharge, VTEP duration across institutions was 15% for 1 week, 39% for 2 weeks, 36% for 4 weeks and 10% for other. For those who underwent laparoscopic/robotic partial Nx, 70% of the institutions utilized VTEP (29% compression stockings and 85% LMWH; Table 3). VTEP was continued after discharge from the hospital for 1 week, 2 weeks and 4 weeks in 20%, 48% and 32% of the institutions.

<table>
<thead>
<tr>
<th>Total no. of departments</th>
<th>No. responding</th>
<th>Response rate (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Finland</td>
<td>25</td>
<td>12</td>
</tr>
<tr>
<td>Norway</td>
<td>15</td>
<td>11</td>
</tr>
<tr>
<td>Sweden</td>
<td>36</td>
<td>15</td>
</tr>
<tr>
<td>Denmark</td>
<td>8</td>
<td>6</td>
</tr>
<tr>
<td>Iceland</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>Total</td>
<td>85</td>
<td>45</td>
</tr>
</tbody>
</table>

Table 1 The response rate from the different Nordic countries among the institutions which perform therapeutic procedures for renal cancer
respectively. Five centers performed laparoscopic/robotic thermal ablation of tumors, and overall, 60% used compression stockings and 80% used LMWH. Two centers continued VTEP for 1 week (40%) and three centers continued it for 2 weeks (60%). Three centers performed percutaneous ablation and used VTEP using compression stockings (67%) and 33% used LMWM. After discharge, one facility continued VTEP for 1 week, while the others continued it for 2 weeks. The responses for individual questions among institutions from the different Nordic countries are shown in Table 2.

**Table 2** Tabulated data on what form(s) of VTEP is used by patients undergoing renal surgery while they are in the hospital and after they leave the hospital

<table>
<thead>
<tr>
<th>VTEP when hospitalized</th>
<th>Finland (n=12)</th>
<th>Norway (n=11)</th>
<th>Sweden (n=15)</th>
<th>Denmark (n=6)</th>
<th>Iceland (n=1)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Early ambulation</td>
<td>45%</td>
<td>40%</td>
<td>43%</td>
<td>60%</td>
<td>–</td>
</tr>
<tr>
<td>Compression stockings</td>
<td>–</td>
<td>40%</td>
<td>36%</td>
<td>60%</td>
<td>80%</td>
</tr>
<tr>
<td>Low-molecular-weight heparin</td>
<td>91%</td>
<td>100%</td>
<td>85%</td>
<td>60%</td>
<td>100%</td>
</tr>
</tbody>
</table>

**Table 3** What kind of VTEP is utilized in patients’ undergoing different modalities of surgery?

<table>
<thead>
<tr>
<th>Use of VTEP after open nephrectomy</th>
<th>Yes</th>
<th>No</th>
<th>Use of VTEP after open partial nephrectomy</th>
<th>Yes</th>
<th>No</th>
<th>Type of VTEP after open partial nephrectomy</th>
<th>Compression stockings</th>
<th>Subcutaneous heparin</th>
<th>Unfractionated heparin</th>
<th>Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)</th>
<th>LMWH</th>
</tr>
</thead>
<tbody>
<tr>
<td>Use of VTEP after laparoscopic/robot-assisted nephrectomy</td>
<td>Yes</td>
<td>No</td>
<td>Use of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Yes</td>
<td>No</td>
<td>Type of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Compression stockings</td>
<td>Subcutaneous heparin</td>
<td>Unfractionated heparin</td>
<td>Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)</td>
<td>LMWH</td>
</tr>
<tr>
<td>Use of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Yes</td>
<td>No</td>
<td>Type of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Compression stockings</td>
<td>Aspirin</td>
<td>Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td></td>
</tr>
<tr>
<td>Use of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Yes</td>
<td>No</td>
<td>Type of VTEP after laparoscopic/robot-assisted partial nephrectomy</td>
<td>Compression stockings</td>
<td>Aspirin</td>
<td>Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td>–</td>
<td></td>
</tr>
</tbody>
</table>

**Abbreviation:** VTEP, venous thromboembolism prophylaxis.

**Discussion**

Our survey results demonstrate wide variability in the type, duration and setting for VTEP among institutions performing therapeutic surgery/procedures for renal cancer in the Nordic countries. As just one example, the use of any VTEP after minimally invasive radical Nx ranged from 0% to 91% by the institutions within a given country (60%–91%, if the one institution that responded from Iceland was excluded). For minimally invasive partial Nx, the range was 40%–90%. Similar wide variability is seen in
the type of VTEP utilized and in the duration of VTEP use after discharge from the hospital. This is despite the fact that both the EAU and Danish Society for Hematostasis and Thrombosis recommend that all patients undergoing surgery for malignant solid tumors should have thrombosis prophylaxis, and that it ought to be given for up to 4 weeks after a major surgery.6,8

The variability in practice may reflect critical gaps in our knowledge on the use of VTEP in kidney cancer. Perhaps the most immediate and glaring gap is the prospective clinical trials on VTEP among patients undergoing surgery for renal cancer. Several studies have used the American College of Surgeons National Surgical Quality Improvement Program to estimate the rate of VTE within 30 days after renal cancer surgery and found rates between 0.7% and 1.7% (open radical Nx 1.7%, laparoscopic radical or partial Nx 0.7%).9–13 While several publications have explored the risk factors for the development of VTE,9,11,13–15 there are no prospective, randomized trials comparing different types, duration and therapeutic settings for VTEP among renal cancer surgery patients. A single-center, retrospective experience suggested that perioperative use of pharmacologic VTEP limited to postoperative hospitalization was not associated with lower VTE rates after robotic partial nephrectomy.14 However, in this study no investigations were done to find out whether the patients received any VTE prophylaxis after discharge from the hospital. Further, the study is based on a survey and has the potential to suffer from bias.

Another significant gap is the lack of specific guidelines for VTEP in urologic surgery. The American Urologic Association and the International Consultation on Urological Disease produced a collaborative review, Anticoagulation and Antiplatelet Therapy in Urologic Practice, which predominantly focused on the management of patients on chronic anticoagulation or antiplatelet therapy for differing medical comorbidities who required a urologic intervention. While there have been several review papers on the subject of VTEP in urology,16,17 urologic surgery-specific guidelines are largely lacking and where guidance exists, it is generally predicated on indirect evidence from other surgical procedures rather than procedure-specific studies within the urologic literature. For example, the EAU has created a guideline on VTEP (http://uroweb.org/guideline/thromboprophylaxis/#3) in which the quality and indirect nature of the evidence available to the panel meant that the overwhelming majority of recommendations were with low or very low quality of evidence and most of the recommendations had to be “weak”. This is not a reflection on the quality of the guideline per se, but on the quality of the evidence available in the literature that underlies the guideline process itself.

It is not surprising that given the gaps in evidence and specific guidance surrounding VTEP for urologic surgery, there have been large variations in the use of VTEP reported both for radical cystectomy16,18,19 and radical prostatectomy.16,18–20 Our study is the first report to survey the use of VTEP across institutions focused on surgery for renal tumors. As is the case with pelvic malignancies, there is widespread variation in the use of VTEP in terms of the type used, the surgical indications, whether it was used after hospital discharge and the total duration after surgery. The significant variability of VTEP across most urologic cancer surgery clearly highlights the need for urologic-specific VTEP guidelines such as the most recent one published by the EAU. Dissemination of these guidelines and more widespread adoption should help to reduce these variations in care and align practice patterns with the evidence available. There are several limitations of this study that should be acknowledged. This is a survey study and, therefore, cannot measure the behaviors of those institutions that did not fill out the instrument. However, the overall return rate, 53%, among those institutions performing kidney tumor surgery, was moderate for a survey-based study, which mitigates this to some degree. Additionally, the study was completed before the EAU guidelines on VTEP were published. However, the results of our study may still be used as baseline evaluation to which future practice patterns can be compared in Scandinavia. It would be useful to repeat this study at defined intervals to test the incorporation of the EAU guidelines into clinical practice. Finally, this survey was intentionally restricted to the Nordic countries and, therefore, may not reflect the practice patterns elsewhere.

Conclusion

The study shows large differences in peri- and postoperative prophylactic treatment for VTE after renal surgery. This variation in care often does not match current guidelines from either the EAU or the Danish Society for Hematostasis and Thrombosis and should be reevaluated as these guidelines become more widely disseminated. Further studies are needed to be able to give evidence-based recommendations.

Acknowledgment

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Disclosure
The authors report no conflicts of interest in this work.

References
**Supplementary material**

Venous-thrombotic-embolic (VTE) prophylaxis survey for renal surgery

1. Country in which you practice

2. What form(s) of VTE prophylaxis do you use in patients with renal tumors before surgery (check all that apply)?
   - None
   - Aspirin
   - Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
   - Subcutaneous heparin
   - Low molecular weight heparin
   - Unfractionated heparin
   - Other

3. What form(s) of VTE prophylaxis do you use in patients undergoing renal surgery while they are in the hospital (check all that apply)?
   - None
   - Early ambulation
   - Sequential compression device
   - Compression stockings
   - Aspirin
   - Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
   - Subcutaneous heparin
   - Low molecular weight heparin
   - Unfractionated heparin
   - Other

4. Do you use VTE prophylaxis in patients undergoing open nephrectomy after they leave the hospital?
   - Yes
   - No – if no go to question 7

5. If yes, what prophylaxis do you use (check all that apply)?
   - Compression stockings
   - Aspirin
   - Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
   - Subcutaneous heparin
   - Low molecular weight heparin
   - Unfractionated heparin
   - Other

6. How long do you use VTE prophylaxis after patients leave the hospital for nephrectomy?
   - One week
   - Two weeks
   - Four weeks
   - Six weeks
   - Other

7. Do you use VTE prophylaxis in patients undergoing open partial nephrectomy after they leave the hospital?
   - Yes
   - No – if no go to question 10

8. If yes, what prophylaxis do you use (check all that apply)?
   - Compression stockings
   - Aspirin
   - Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
   - Subcutaneous heparin
   - Low molecular weight heparin
   - Unfractionated heparin
   - Other

9. How long do you use VTE prophylaxis after patients leave the hospital for open partial nephrectomy?
   - One week
   - Two weeks
   - Four weeks
   - Six weeks
   - Other

10. Do you use VTE prophylaxis in patients undergoing laparoscopic/robot assisted nephrectomy after they leave the hospital?
    - Yes
    - No – if no go to question 13

11. If yes, what prophylaxis do you use (check all that apply)?
    - Compression stockings
    - Aspirin
    - Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
    - Subcutaneous heparin
    - Low molecular weight heparin
    - Unfractionated heparin
    - Other

12. How long do you use VTE prophylaxis after patients leave the hospital for laparoscopic/robot assisted nephrectomy?
    - One week
    - Two weeks
    - Four weeks
    - Six weeks
    - Other

13. Do you use VTE prophylaxis in patients undergoing laparoscopic/robot assisted partial nephrectomy after they leave the hospital?
Yes
No – if no go to question 16

14. If yes, what prophylaxis do you use (check all that apply)?
- Compression stockings
- Aspirin
- Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
- Subcutaneous heparin
- Low molecular weight heparin
- Unfractionated heparin
- Other

15. How long do you use VTE prophylaxis after patients leave the hospital for laparoscopic/robot assisted partial nephrectomy?
- One week
- Two weeks
- Four weeks
- Six weeks
- Other

16. Do you use VTE prophylaxis in patients undergoing laparoscopic/robotic thermal ablation of kidney tumors after they leave the hospital?
Yes
No – if no go to question 19

17. If yes, what prophylaxis do you use (check all that apply)?
- Compression stockings
- Aspirin
- Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
- Subcutaneous heparin
- Low molecular weight heparin
- Unfractionated heparin
- Other

18. How long do you use VTE prophylaxis after patients leave the hospital for laparoscopic/robotic thermal ablation of kidney tumors?
- One week
- Two weeks
- Four weeks
- Six weeks
- Other

19. Do you use VTE prophylaxis in patients undergoing percutaneous thermal ablation of kidney tumors after they leave the hospital?
Yes
No – if no finished

20. If yes, what prophylaxis do you use (check all that apply)?
- Compression stockings
- Aspirin
- Direct-acting oral anticoagulants (dabigatran, apixaban, rivaroxaban, edoxaban)
- Subcutaneous heparin
- Low molecular weight heparin
- Unfractionated heparin
- Other

21. How long do you use VTE prophylaxis after patients leave the hospital for percutaneous thermal ablation of kidney tumors?
- One week
- Two weeks
- Four weeks
- Six weeks
- Other