

Are graduated compression stockings still essential for VTE prophylaxis in general surgical patients?*

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References

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weight tapes, printed cassette decks,
silence refills.

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Background: Current guidelines for the prevention of severe rheumatoid arthritis (RA) suggest primary corticoids to reduce the effects of sustained symptomatic nonsteroidal anti-inflammatory drugs (NSAIDs) on the gastrointestinal tract. However, NSAIDs are also known to be associated with an increased risk of cardiovascular disease. The aim of this study was to evaluate the effect of NSAIDs on the cardiovascular system in patients with RA.

Patient methods: We studied the rate of asymptomatic venous thromboembolism (VTE) according to patterns of a general surgery department during two consecutive five periods of 24 months with different prophylactic regimes. During the first 24 months (Group A), parenteral heparins were used (1000 IU) but postulated inappropriate (Group B). During the second 24 months (Group C) only LMWH was used. **Results:** In Group A there were 21 90 patients, with a total of 3028 operations; in Group B there were 2796 patients with a total of 3977 operations. 82.5% of the patients in Group A and 84.3% in Group B received VTE prophylaxis (heparins) and 1000 IU in A, 1000 IU only in B). Symptoms suggesting venous thromboembolism occurred in 45 patients in Group A and in 17 in Group B. Examination of these patients revealed a VTE in 14 and 5 patients.

Conclusion: The widespread asymptomatic virus seroconversion is a feature of a general surgery department which has both a high, routinely applied, patient to patient rate of MSM. Co-dated asymptomatic serology did not yield additional benefits to our patients.

References

**Spring Therapeutics, UT Physicians, nationwide
and Hospital Organizations, Hospital Administrators**

Transcription

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Welche Konsequenzen ergeben sich aus Ihrer Analyse für die Altersvorsicherung im öffentlichen Dienst?

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Résumé: Les diverticels colocoliques sont la prévalence de la Maladie Chronique Intestinale Sévère (MCIS) ou d'une charge continue de surcroissance l'usage de l'acide aminé d'origine digestive-muqueuse. Les patients souffrant de MCIS ont des symptômes gastro-intestinaux (GI), la plus fréquente de ces études, qui ont montré un lien entre la consommation d'aliments, et les symptômes gastro-intestinaux (GI). **But de l'étude:** L'objectif de l'étude était d'évaluer la relation entre la consommation d'aliments et les symptômes gastro-intestinaux (GI) chez les patients souffrant de MCIS. **Méthode:** Les patients souffrant de MCIS ont été recrutés dans un hôpital de soins de 2e année, spécialisée de 24 ans avec différentes maladies gastro-intestinales. Au cours des 24 semaines, les patients ont été divisés en deux groupes: le groupe A (groupe A) et le groupe B (groupe B). Les patients du groupe A ont été divisés en deux sous-groupes: le groupe A1 (groupe A1) et le groupe A2 (groupe A2). Les patients du groupe B ont été divisés en deux sous-groupes: le groupe B1 (groupe B1) et le groupe B2 (groupe B2). **Résultats:** Chez le groupe A, les patients ont été divisés en deux sous-groupes: le groupe A1 (groupe A1) et le groupe A2 (groupe A2). Les patients du groupe A1 ont été divisés en deux sous-groupes: le groupe A1.1 (groupe A1.1) et le groupe A1.2 (groupe A1.2). Les patients du groupe A2 ont été divisés en deux sous-groupes: le groupe A2.1 (groupe A2.1) et le groupe A2.2 (groupe A2.2). Les patients du groupe B ont été divisés en deux sous-groupes: le groupe B1 (groupe B1) et le groupe B2 (groupe B2). Les patients du groupe B1 ont été divisés en deux sous-groupes: le groupe B1.1 (groupe B1.1) et le groupe B1.2 (groupe B1.2). Les patients du groupe B2 ont été divisés en deux sous-groupes: le groupe B2.1 (groupe B2.1) et le groupe B2.2 (groupe B2.2). **Conclusion:** Les patients souffrant de MCIS ont des symptômes gastro-intestinaux (GI) et les symptômes gastro-intestinaux (GI) sont liés à la consommation d'aliments. Les patients souffrant de MCIS ont des symptômes gastro-intestinaux (GI) et les symptômes gastro-intestinaux (GI) sont liés à la consommation d'aliments.

Les bas à compression élastique dirigent-ils le sang vers le cœur pour la prophylaxie de la maladie thrombo-embolique ou du varicose veineux ?

Venous thromboembolism (VTE) continues to account for considerable morbidity, mortality and medical care expense. In the USA the incidence of deep venous thrombosis (DVT) is reported to be approximately 450 000 cases per year and the overall incidence of pulmonary embolism (PE) to be about 355 000 cases per year (8). Numerous studies have indicated that especially surgical and trauma patients are at significant risk for developing venous thromboembolic complications, including pulmonary embolism. Without prophylactic treatment, the risk of VTE (2, 19, 50) is considered to be:

- 25–30% in general surgery (8, 13, 46);
- 60% after major trauma (22, 23) and
- up to 45–70% with hip or knee replacements.

Both physical and pharmacological measures aimed at preventing thromboembolic events have been previously evaluated. In earlier studies graduated compression stockings were found to substantially reduce the incidence of VTE in surgical patients (2, 13, 30, 46, 60). During recent years, however, notable developments in pharmacological thromboprophylaxis have been achieved. The introduction of low molecular weight heparins (LMWH), in particular, has resulted in significant advances in treatment modalities so that today they have become the standard for VTE prophylaxis in most countries. The prophylactic properties of LMWHs have yielded equal or even slightly better outcomes than unfractionated heparins. Moreover, they have fewer side effects, do not require laboratory monitoring, and possess pharmacokinetic characteristics which permit their use as a subcutaneously administered, fixed daily dose. LMWHs are reported to reduce the overall incidence of deep vein thrombosis after general surgery by up to 70% (2, 33).

In light of this progress, reevaluation of the common use of graduated compression stockings, as it is still recommended in this setting, is warranted.

Patients, methods

Study population and design

The study population consisted of all patients treated in a general surgery department over a period of four years.

With a prospective observational study design, the incidence of symptomatic deep venous thrombosis and pulmonary embolism was investigated with two different VTE prophylaxis regimens. During the

- first 24 months (group A) patients at risk received combined treatment for the prevention of thromboembolism consisting of LMWH and graduated compression stockings;
- second 24 month period (group B) only LMWH was used for prophylaxis.

Treatment

Prophylactic treatment was given to all patients undergoing major surgery (>30 min). The treatment was initiated the day before surgery and was continued for five days thereafter, or until complete remobilization was achieved. The prophylactic measures were also taken in patients with minor surgery and patients who were not actually operated on if they were expected to be immobile for more than 6 hours during the day for more than two days. Patients under the age of 14 did not receive VTE prophylaxis.

The thigh-length compression stockings (TED[®] Kendall) in group A were fitted to each patient's legs according to the manufacturer's instructions. The patients were advised to wear them 24 hours a day. Compliance with the wearing of stockings was checked at least twice daily. Patients found without stockings or not wearing them properly on more than two occasions were categorized as noncompliant. Patients with known significant peripheral artery disease (intermittent claudication and/or ABI <0.7) did not receive stockings.

The LMWH employed in both groups was enoxaparin. It was administered according to a simple weight-adjusted protocol. Patients with a body mass index (BMI)

of up to 30 kg/m² received 20 mg of enoxaparin and patients with BMI >30 kg/m² received 40 mg – subcutaneously, once daily. Patients in whom the use of LMWH was contraindicated (those with a history of or suspected active HIT II or allergic skin reactions) were treated with 15 mg of fondaparinux subcutaneously, twice daily.

In group B, no stockings were used. The prophylactic treatment consisted of LMWH (or fondaparinux alone, administered following the same protocol as in group A).

All patients in groups A and B were strongly encouraged to get out of bed and walk as early as possible.

Outcomes

In all patients with clinical symptoms suggesting deep vein thrombosis colour-Doppler ultrasonography of the pelvic and leg veins was done, including compression testing of the leg veins. The leg examination included all veins from the thigh down to the ankle. All sonographic examinations were carried out by two highly-skilled examiners with an ATL / Philips HDI 3000 machine, using a 3–5 mHz curved array scanner for the pelvic vein and a 5–7 mHz phased array scanner for the leg veins. If this examination was temporarily not possible, or if the results were inconclusive, ascending phlebography followed.

Patients with symptoms of pulmonary embolism underwent spiral computed tomography of the lungs. This was judged positive if it revealed distinct filling defects in large vessels. If the diagnosis remained inconclusive after this examination, a ventilation-perfusion lung scan (judged positive only in cases of „high probability“ results) and/or a pulmonary angiography was performed.

Results

A total of 6187 patients (Tab. I) were treated in the general surgery department during the study period, with a total of 5861 operations.

Treatment group characteristics

In group A (treated with stockings and LMWH), there were 3181 patients: 1742 men and 1439 women, with a mean age of 48 years. The mean duration of hospital stay was 7.3 days; 2720 of the A patients had surgery with a total of 3890 operations (Tab. 1, Tab. 2). Four hundred sixty-one patients had no surgery. VTE prophylaxis (Tab. 3) was applied to a total of 2623 (82.5%) of the 3181 patients of period A: to 2422 (89.1%) patients with surgery and to 202 (43.8%) patients without surgery. Fifteen patients in group A could not wear compression stockings due to advanced occlusive artery disease and in others because of extreme obesity. Twenty-eight patients were non-compliant with wearing stockings. Three patients in group A needed to be treated with hirudin, instead of LMWH, due to contraindications.

In group B (only LMWH), 2986 patients were treated: 1691 men and 1295 women, with a mean age 49 years. The mean duration of hospital stay was 6.2 days and 2551 patients had a total of 2911 operations. Four hundred thirty-three patients had no surgery. VTE prophylaxis was provided to 2512 (84.0%) of the 2986 patients in period B: to 2305 (90.3%) patients with surgery and to 207 (46.6%) patients without surgery. Two patients had to be treated with hirudin, instead of LMWH.

Tab. 1
 Patient characteristics,
 type of surgery

		group A	group B
		LMWH + stockings	LMWH only
patients	patient number, sex	3181	2986
	men	1742	1691
	women	1439	1295
	age (years)	48	49
	days of stay	7.3	6.2
type of surgery	vascular	261	157
	thyroid	149	104
	choleci	119	62
	anal related	49	65
	upper GI tract	227	153
	urologic	99	164
	muscul	176	108
	skin	215	206
	low	5	4
	gastrointestinal	44	29
	gastrointestinal	290	284
	gastrointestinal	5	3
	urologic	257	181
	urologic	58	44
	urologic	42	33
	urologic	190	93
	choleci	226	102
	choleci (gastrointestinal)	45	29
	choleci (gastrointestinal)	83	81
	choleci (gastrointestinal)	167	173
	choleci	25	29
	choleci	257	181
	choleci	58	44
	choleci	42	33
	choleci	190	93
	choleci	226	102
	choleci (gastrointestinal)	45	29
	choleci (gastrointestinal)	83	81
	choleci (gastrointestinal)	167	173
	choleci	25	29
total		3890	2911

Outcomes (Tab. 4)

Deep vein thrombosis

Thirty-eight patients in group A and 40 in group B developed clinical symptoms suggestive of a deep vein thrombosis which required further investigation. Thirty patients in A and 31 patients in B received color-duplex assisted compression sonography and eight patients in A and nine in B received phlebography. A DVT was confirmed in six patients in group A and in three patients in group B.

Tab. 2
 Number of operations
 per patient

number of operations	group A	group B	p value
	LMWH + stockings	LMWH only	
1	2517	1349	0.658
2	127	124	0.718
3	44	45	0.688
4	16	19	0.596
5	9	9	0.893
6	5	10	0.157
8	0	1	0.262
10	0	1	0.262

	VTE prophylaxis	group A LMWH + stockings	group B LMWH only	P value
patients		2181	2166	
with surgery	—	1726	2053	0.002
	+	1429 (83%)	2014 (98%)	0.001
without surgery	—	455	113	0.002
	+	202 (45%)	202 (94%)	0.001
total	+	2629 (82.3%)	2217 (94%)	0.001

—, not treated; +, treated.

Table 3
VTE prophylaxis

	group A LMWH + stockings	group B LMWH only	P value
operative patients	1612/1612	1612/1612	
hip	70/5	7/7	
phlebectomy	6/3	0/0	
joint CT	5/1	4/2	
joint arthroscopy	1/0	0/0	
joint arthroscopy	3/0	3/3	
total knee arthroscopy	44/7	4/0	0.001
patients with VTE	7	5	0.445
total rate	0.22	0.20	0.87

Table 4
Ischemic events and VTE rates

Pulmonary embolism

A pulmonary embolism was suspected in six patients in group A and seven in group B during their hospital stay. 12 of these patients underwent spiral computed tomography of the lungs, one patient had a ventilation-perfusion scan and one needed an additional pulmonary angiogram. The pulmonary embolism was confirmed in one patient in group A and in three in group B.

Characteristics of patients with VTE

All the 12 patients (7 in group A, 5 in group B) with confirmed VTE were receiving prophylaxis at the time the VTE event occurred. The mean duration of hospital stay in these patients was 38 days in group A and 39 days in group B. All patients with PE in both groups had malignancy disease.

Discussion

This study, which took place over a period of four years and involved 6167 patients on a general surgery ward, revealed no significant differences in the incidence of symptomatic venous thromboembolism whether graduated compression stockings were used in addition to LMWH prophylaxis, or not.

A review of the literature over the last three decades shows that, despite significant advancements in the prevention of venous thromboembolism, there is still a considerable degree of variation in regard to risk group definition, the number of patients receiving prophylaxis, and prophylactic modalities.

To achieve better standardization in VTE prophylaxis, several consensus conferences have generated guidelines which are aimed at promoting and attaining adequate VTE prophylaxis for the individual patient. Yet, even the latest guidelines concerning the

prevention of thromboembolism in surgical patients, namely the updated ACCP guidelines from 2004 (24) and those of the International Consensus Statement dating from 2006 (46), are not completely concordant. In regard to general surgery patients, the former do not recommend a specific prophylaxis in low-risk patients, propose the alternative use of stockings or heparin in moderate- and high-risk patients, and recommend combination therapy only in very high-risk patients. The latter consider the use of compression stockings in low risk patients, recommend stockings with intermittent pneumatic compression as an alternative to LMWH in moderate risk patients with bleeding tendency and only consider compression stockings as an adjunct to LMWH in high-risk patients. In other fields of surgery (for example urologic surgery or gynecologic surgery) there is even less evidence for the necessity of compression stockings. In these fields the recommendations are either extrapolated from general surgery or consensus concerning the use of compression stockings are lacking – as is also the case in all recently published studies on VTE prophylaxis in medical patients (36, 41, 52).

There are generally nine studies cited which substantiate significant VTE risk reduction following the use of graduated compression stockings in general surgery (65). All of them were carried out 10 or more years ago (3, 5, 28, 43, 55, 56, 58, 63, 64). The study designs are rather heterogeneous and most of them included a total of less than 100 patients (5, 28, 55, 56, 58). Only two out of the nine studies (3, 23) compared the use of stockings with the absence of prophylactic measures. These found a 50% reduction in the rate of venous thromboembolism in patients who wore stockings. In the seven other studies, a combination therapy was used consisting of either intermittent pneumatic compression (43, 56) or pharmacological methods (dextran, low dose heparin, dithyroglycerin-heparin) together with graduated compression stockings (5, 43, 58, 63, 64). In five of the studies the effectiveness of compression stockings was investigated on only one of the patient's legs; the other leg served as a control (5, 43, 55, 56, 58).

Several studies in the following years have provided evidence that low dose heparin significantly reduced the incidence of postoperative VTE, and the benefit appeared to increase with the additional use of compression stockings (11, 15, 31, 32, 45). Subsequently, low dose heparin in combination with compressive stockings became the most widely adopted measures for VTE prophylaxis in general surgery – a regimen which has been used in many facilities until today (4, 10, 12, 65).

When LMWHs were introduced into VTE prophylaxis in the mid-1990s, they first proved their effectiveness in orthopedic surgery, namely with hip and knee replacements. Later LMWHs were adopted in general surgery and were combined with the compression stocking prophylaxis already in use. Only four studies can be found which show an advantage to the combination of LMWH with compression stockings over stockings alone – none of them in general surgery (1, 38, 42, 47). In all four studies the group employing compression stockings alone is categorized as a placebo group. However, the incidence of VTE in these placebo groups is actually about the same as in previous studies where no prophylactic measures were used. Only one single, rather small study has compared, as we have, the use of LMWH in combination with compression stockings to LMWH alone for VTE prophylaxis. Kaloiki et al. (34), randomized 78 patients undergoing a total hip replacement into three groups:

- a control group without prophylaxis,
- a group which received enoxaparin once daily,
- a group which received enoxaparin and wore graduated compression stockings (TED) for a period of 8–10 days.

All patients had a preoperative perfusion lung scan and a postoperative perfusion-ventilation scan, as well as bilateral ascending phlebography between days 8 and 12. The DVT rates were 93%, 38% and 25%, respectively. No other study comparing LMWH with and without stockings has been published as of today – neither in general surgery nor in any other medical specialty.

The mechanism of action through which compression stockings prevent thrombo-

embolism remains unclear, though it is thought to be multi-functional with a reduction in venous diameter playing the key role. External compression was shown to reduce the overall cross-sectional area of the lower limb and increase the flow velocity within the veins. Compression also appears to improve the evacuation and reabsorption of incompetent and incompletely emptied valvular cusps. Reduction in reflux-related stasis and augmented flow velocity could decrease the risk of thrombus formation by reducing venous wall distension, local contact time, and the concentration of coagulation reactants (2, 14, 57, 61, 66).

Despite their widespread use, there is also no agreement as to the most effective stocking design to achieve the above mentioned effects. High-length TED stockings, which are probably employed in most studies, have intended pressures of 18 mmHg at the ankle, 14 mmHg at the calf, 8 mmHg at the knee, 10 mmHg above the knee, and 8 mmHg at the thigh. It is not known whether these pressures are optimal. Moreover, some authors estimate calf-length stockings to be more effective than thigh-length and some prefer non-elastic ones (2, 9, 26). Stocking design is not always specified in the study protocols and, particularly in some of the newer studies, it is not clear whether or not stockings were used at all (18, 27, 30, 44, 59, 41, 35, 52).

Even those stockings believed to be ideally designed may not achieve the warranted effects. A study measuring pressure gradients in patients wearing prophylaxis stockings showed that 90% of the stockings failed to produce the ideal pressure gradient of 18–14–8 mmHg from the ankle to the knee; 54% even produced an inverse gradient (7, 67).

Given the recommended pressure gradients intended to be realized by the design of prophylaxis stockings, it remains obscure why other types of compression stockings, e.g., those recommended for the prevention of post-thrombotic syndrome following a deep vein thrombosis, achieve significantly higher pressure ranges (mostly 40–30–21 mmHg) although they must be worn for much longer time periods. These higher pressure compression stockings fail, however, to prevent DVT recurrence, whereas

prophylaxis stockings are claimed to reduce DVT incidence by 50% or more (9, 37).

Finally, prophylaxis stockings were conceived for supine patients (61). However, today most patients are mobilized within only a few hours after surgery. Nevertheless guidelines recommend stockings for 3–7 days (61).

Unlike most other studies, ours did not serially screen all our study patients for deep venous thrombosis or pulmonary embolism. The significance of an asymptomatic VTE detected by screening methods such as ¹²⁵I fibrinogen marking, plethysmography, contrast phlebography, compression sonography, D-dimer testing, or ventilation-perfusion scans is a matter of debate (6, 35, 40, 54). Conclusive evidence that treatment of asymptomatic patients with VTE detected by these methods results in better outcomes is lacking, not only in regard to nonfatal or fatal pulmonary embolism (35), but also in regard to postthrombotic syndrome (25) and pulmonary hypertension (48). Recent study designs have begun to focus more, as we have, on symptomatic thromboembolic events as clinically relevant endpoints (40).

Limitations

We cannot rule out the possibility of diagnostic bias in our study, since the use of stockings cannot be blinded and for organizational reasons we were not able to perform randomization. It is possible that physicians were more likely to order an imaging test for patients who had not received compression stockings than for those who had. Nevertheless, the incidence of symptomatic VTE in our surgical patients was very low with both prophylaxis regimens.

A decline in VTE rates in clinical medicine has been anticipated in recent years (17, 29). Research in this area is lacking, but considering the still doubtless significance of the Virchow triad in regard to VTE pathogenesis, the current prevailing concept of early mobilization is likely to be the most effective measure contributing towards the prevention of thromboembolism in all areas of medicine. New surgical techniques, especially endoscopic and fast track surgery and modern anesthesiologic methods, are

also reported to reduce VTE rates (49, 51, 53). Furthermore, complicated risk stratification concepts are increasingly being disregarded in favor of simple and effective protocols like the one we used with our surgical patients (24) and finally, new, even more potent, anti-thrombotic agents are being developed (16, 20, 21, 49).

Conclusion

Our data showed that the rate of symptomatic venous thromboembolism in patients on a general surgery ward is kept low when a concept of early mobilization and a simple, but strictly applied protocol of preventative low molecular weight heparin is implemented. Graduated compression stockings did not yield an additional preventive effect with this regimen.

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