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# **ABOUT COVER**

Editorial Board Member, Yu-Chang Yeh, MD, PhD, Associate Professor, Department of Anesthesiology, National Taiwan University Hospital, Taipei 100, Taiwan. tonyyeh@ntuh.gov.tw

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MINIREVIEWS

# Incremental value of compression ultrasound sonography in the emergency department

Alessandro Di Vilio, Andrea Vergara, Alfonso Desiderio, Franco Iodice, Alessandro Serio, Stefano Palermi, Francesco Gambardella, Simona Sperlongano, Renato Gioia, Maria Acitorio, Antonello D'Andrea

ORCID number: Alessandro Di Vilio 0000-0002-2885-410X; Andrea Vergara 0000-0001-5217-7754: Alfonso Desiderio 0000-0002-2885-789X; Franco Iodice 0000-0002-2014-287X; Alessandro Serio 0000-0003-1558-9981; Stefano Palermi 0000-0003-1558-4857; Francesco Gambardella 0000-0003-1558-4729; Simona Sperlongano 0000-0007-0986-7876; Renato Gioia 0000-0002-2885-9851; Maria Acitorio 0000-0002-2885-1948; Antonello D'Andrea 0000-0002-9698-9145.

Author contributions: Di Vilio A and D'Andrea A proposed concepts; D'Andrea A, Di Vilio A, and Vergara A conducted methodology; Di Vilio A and Vergara A were responsible for drafting the manuscript; Di Vilio A, D'Andrea A, Vergara A, Iodice F, Desiderio A, Acitorio M, Sperlongano S, Serio A, Palermi S, Gambardella F, and Gioia R performed writing reviews and editing; D'Andrea A, Desiderio A, and Acitorio M performed supervision.

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Alessandro Di Vilio, Andrea Vergara, Franco lodice, Simona Sperlongano, Unit of Cardiology and Intensive Coronary Care, University of Campania "Luigi Vanvitelli", Monaldi Hospital, Naples 80131, Italy

Alfonso Desiderio, Maria Acitorio, Antonello D'Andrea, Unit of Cardiology and Intensive Coronary Care, Umberto I Hospital, Nocera Inferiore 84014, Italy

Alessandro Serio, Stefano Palermi, Francesco Gambardella, Human Anatomy and Sport Medicine Division, Department of Public Health, University of Naples "Federico II", Naples 80131, Italy

Renato Gioia, Department of Medicine, Surgery, and Dentistry, University of Salerno, Salerno 84084, Italy

Corresponding author: Alessandro Di Vilio, MD, Doctor, Unit of Cardiology and Intensive Coronary Care, University of Campania "Luigi Vanvitelli", Monaldi Hospital, Via Leonardo Bianchi, Naples 80131, Italy. adivilio56@gmail.com

# Abstract

The quick evaluation of venous thromboembolism is a key point of modern medicine since the delayed diagnosis is associated with a worse prognosis. Venous ultrasound (VU) is a sensitive and rapidly performed test in cases of suspected deep venous thrombosis. Various protocols have been proposed for its execution, such as the study of the whole deep venous circulation of the lower limb or the analysis of the femoral-popliteal area. The aim is to detect a vessel thrombus and the most sensitive element is the non-compressibility with the probe. Initially, the thrombus is hypoechogenic and adherent to the vessel; later, it tends to organize and recanalize. Usually, in the early stages, the risk of embolism is higher. The role of studying the iliac axis and calf veins is still uncertain. VU is not useful for assessing response to anticoagulation therapy and it is unclear whether the persistence of thrombotic abnormalities can guide on a possible prolongation of therapy.

Key Words: Compression ultrasound; Deep venous thrombosis; Venous ultrasound; Venous thromboembolism; Critical care ultrasonography

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**Core Tip:** Venous ultrasound represents an important weapon for emergency setting care. Nevertheless, several different protocols present in the literature could create confusion. In this review our goal is to define a practical and clear guide to support the physician in rapid deep venous thrombosis diagnosis and correct management.

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# INTRODUCTION

Thrombus formation is a pathological phenomenon caused by an inappropriate hemostatic response; many different factors are involved, often favored in blood stasis points such as venous valves. A major theory delineating the pathogenesis of venous thromboembolism (VTE), often called Virchow's triad, states that VTE occurs as a result of alterations in blood flow, in vascular endothelial injury and in blood constituents. Blood clots could leave these sides to enter the bloodstream, reaching right heart chambers and pulmonary circulation. Rarely, in the presence of patent foramen ovale with right-to-left shunt, there may be systemic embolism. Hence, deep venous thrombosis (DVT) and pulmonary embolism (PE) are two sides of the same coin; about 50% of patients with proximal DVT is affected by an asymptomatic PE, as well as 80% of PEs suffers DVT (often asymptomatic)[1]. DVT may be distal, interesting tibial-peroneal district, or proximal, that affects femoral-popliteal veins; proximal DVT is more frequently related to PE.

DVT is the third most common cardiovascular disease, following heart attack and ischemic stroke[2] (DVT incidence: 150/100000/year, PE incidence: 60-70/ 100000/year). According to Cohen *et al*[3], based on 6 EU countries data (Italy, Spain, France, Germany, United Kingdom and Sweden), EP-related death rate was 12%[3].

DVT should be suspected in patients presenting with leg swelling, pain or erythema; usually symptoms are unilateral calf-related if isolated distal DVT, whole-leg related if proximal DVT[4]. Many patients are asymptomatic. Although uncommon, it is important to identify patients with phlegmasia cerulea dolens, that ranges from phlegmasia alba dolens to venous gangrene, because it should be considered for more aggressive management[5]. PE has a wide variety of presenting features, ranging from no symptoms to shock or sudden death. The most common presenting symptom is dyspnea, followed by chest pain (classically pleuritic) and cough. Chronic thromboembolic pulmonary hypertension has been estimated to occur in 4.8% (95%CI: 2.3-9.6) of patients who survive a PE[6].

The most used predictive score of DVT is Wells Score[7] (Table 1), more accurate than revised Geneva score in PE suspected patients[8-10]. It predicts an increasing incidence of PE with major probability classes ("low" if  $\leq 0$  points to "high"  $\geq 3$  points) [11]. PE incidence ranged from 1%-13% in low probability level (Wells score < 2), 28%-58.3% in medium probability level (Wells score 2-6), and 58.1%-93% in high probability level (Wells score > 6). The sensitivity ranged from 63.8%-79.3%, and the specificity ranged from 48.8%-90.0%.

Quick diagnosis of DVT/PE represents a fundamental weapon for modern medicine. The early detection of DVT is crucial to reduce the risk of thromboembolism in the critical patient, thus reducing the related morbidity and mortality. A delayed diagnosis of PE has poor outcomes, ranges from shock to hospital death[12]: Therefore, the 30 d mortality rate exceeds 3% in patients with DVT who are not anticoagulated, and this mortality risk increases 10-fold in patients who develop PE.

The most diffused vascular diagnostic tools are: (1) Phlebography: Gold standard, not widely used for its invasiveness; (2) Computed tomography (CT) angiogram: Utilized for PE and proximal districts (pelvic, iliac, caval); (3) Magnetic resonance imaging angiogram: Utility comparable to CT angiogram; and (4) Echocolordoppler: The most employed instrumental methodic for quick diagnosis and screening of pathology (limited just in the most peripheral venous tracts).



Table 1 Wells' score		
Features	Score, points	
Active cancer (in treatment or treated in the last 6 mo or under palliative care)	1	
Paralysis, paresis, or recent plaster immobilization of the lower extremity	1	
Bedridden recently > 3 d or major surgery within 12 wk	1	
Localized tenderness along the deep venous system	1	
Entire leg swollen	1	
Calf swelling > 3 cm compared to the other leg	1	
Pitting edema, confined to symptomatic leg	1	
Collateral (nonvaricose) superficial veins present	1	
Previously documented DVT	1	
Alternative diagnosis to DVT as likely or more likely	-2	

DVT: Deep venous thrombosis.

Therefore, we realized a review of clinical studies comparing outcomes of patients with a history of DVT subjected to different managements. We achieved this by doing formal searches of the electronic database MEDLINE (source PubMed) and the Cochrane Controlled Clinical Trials Register Database. About 40 studies were selected from 1989 to 2017, by a combination of medical subject headings including the following terms: Compression ultrasound (CUS), DVT, venous ultrasound (VU) and VTE. References from reviews and selected articles were also examined for potentially relevant citations. Our analysis was restricted to the trials that focused on the comparison between different existing diagnostic protocols, with a special focus on emergency department experiences.

# VU

VU is the commonest methodic used for DVT assessment. Many different protocols have been proposed, thus incrementing the confusing about its management. The study of deep veins is performed with high-frequency linear probes (5-7.5 MHz) or a sector probe, when the limb is particularly large-sized (Figure 1). Specifically, evaluation of the femoral veins should be done with the lower limb in extra-rotation, while other veins are studied in supine position with flexed knee. CUS with Doppler is the choice diagnostic test in patients with suspected DVT and the sensitivity and specificity of proximal CUS is greater than 95% (Figure 2). However, proximal CUS suffers from limitations<sup>[13]</sup>: (1) Calf vein thrombus, that are harder to assess than proximal veins; and (2) Iliac veins thrombus, that cannot be assessed for compressibility and thus these veins should be assessed with venography.

# Pretest evaluation

First step in DVT assessment is probability estimation according to Wells Score[14]. In case of low pretest probability, a negative D-Dimer can rule out DVT without the need for ultrasound confirmation. If, conversely, pretest probability by Wells Score is high, VU is recommended [15,16] (Figure 3). Therefore, the role of D-Dimer in this diagnostic process is limited: it is a degradation product of cross-linked fibrin and it is elevated in nearly all patients with acute DVT (high sensitivity), but it is non-specific since high levels are found in many other conditions (i.e. malignancy, sepsis, recent surgery or trauma, pregnancy, renal failure).

# Protocols

Complete doppler ultrasound: Is the preferred one and it includes bilateral compressions from inguinal ligament, passing through calf veins, to ankle (compressions are separated by 2 cm intervals); it also provides for bilateral common femoral and popliteal vein color doppler images and spectral doppler waveforms, in order to verify possible asymmetries [17,18]. CUS false positivity in calf evaluation is





# Figure 1 Study optimization.



Figure 2 Compression ultrasound example.

extremely uncommon (its specificity in calf district reaches about 97.8%)[19], and the risk of excessive treatment related to calf DVTs represents the principal argument against this protocol. Calf assessment may also provide alternative findings, like musculoskeletal abnormalities[20]. Prospective studies have demonstrated that lack of compressibility of a vein with the ultrasound probe is the most sensitive (> 95%) and specific (> 95%) sonographic sign for proximal vein thrombosis. The addition of color flow Doppler does not improve the sensitivity but can provide supportive evidence of thrombus and help to identify calf veins. Variation of venous size with the Valsalva maneuver has a low sensitivity and specificity for the diagnosis and is no longer performed in many centers[21-24]. In case of negative result, the risk of DVT after 3 mo is estimated about 0.57% (95%CI, 0.25%–0.89%) (Figure 4)[25].

**Extended compression ultrasound:** This protocol is a point-of-care examination that consists of compressions from thigh to knee and is principally utilized when complete doppler ultrasound (CDUS) is not quickly viable[26]. However, if it results negative, a confirmatory CDUS after 5-7 d is recommended, in order to exclude calf involvement (evaluating to start anticoagulation if this is not possible)[27-30]. (Figure 4).

**Two-region ultrasound:** The compressions are limited to femoral and popliteal areas. As in extended compression ultrasound (ECUS), also in this case a negative response should be followed by a CDUS examination 5-7 d later, because both the previous two protocols do not comprehend calf veins[27,31,32]. D-Dimer after a negative ECUS or two-region ultrasound does not affect the follow-up unless it results negative[33]



Di Vilio A et al. Critical care ultrasound



Figure 3 Pretest evaluation algorithm. CDUS: Complete doppler ultrasound; DVT: Deep venous thrombosis; ECUS: Extended compression ultrasound.



Figure 4 Different protocols. Dotted arrows correspond to ultrasound scans separated by 2 cm intervals. Yellow segments instead represent doppler points. CDUS: Complete doppler ultrasound; ECUS: Extended compression ultrasound; 2-region: Two-region ultrasound.

## (Figure 4).

# Thrombosis sides and related approach

The management of DVT is also related to its side, in particular two sides are more controversial (Figure 5): (1) Ileocaval: Being in a blind side for ultrasound sonography, it could be underdiagnosed. Nevertheless, although a normal CUS could be present, asymmetrical or continuous femoral doppler waveforms or whole-leg swelling may indicate an upstream impediment. In these cases, it is reasonable to think about pelvic ultrasound, CT or magnetic resonance venography to rule out this possibility. It has been estimated that this side is involved in 1.6% of DVTs[34]. Because the accuracy of duplex ultrasound for ileocaval DVT is not established, the threshold for CT or magnetic resonance venography should be low; (2) Calf veins: Even if calf district examination is just included in CDUS protocol, calf involvement management is subject to debate. If the physician chooses a wait-and-see approach without treating, it is recommended to repeat ultrasound at 1 wk. If new scan shows proximal progression, then start anticoagulation (progression occurs in 9%-21.4% of cases and is usually associated to symptoms perseverance or exacerbation)[35]; if instead clot remains stable, you should scan again at 2 wk. If thrombus is not more observable at 1



Figure 5 Reference points and respective sonogram images. A: Artery; GS: Great saphenous vein; V: vein.

wk or does not show significant evolution at 2 wk, or if you have begun treating, you can stop its follow-up. You could consider a new assessment in case of therapeutic changes [35,36]. More investigations will be needed to clarify prevalence and risk factors of progression in non-anticoagulated calf thromboses [37]. It is important to point out that short calf areas of non-compressibility are not significative [38], further scans or D-dimer may supply in these cases, although positive D-dimer has demonstrated to be not discriminatory [39]. Moreover, presence of calf DVT could be important in risk stratification about different fields like chronic venous insufficiency, mortality and cancer diagnosis or recurrent DVT occurrence [40,41]. The current American College of Radiology/American Institute of Ultrasound in Medicine/Society of Radiologists in Ultrasound guidelines include selective calf imaging for the subset of patients with calf symptoms not explained by the proximal scan; and (3) Upper limbs: Old statistics referred an incidence of upper limbs thrombosis about 2%-3%, mainly in young patients and in right arm, especially after hard physical effort or in thoracic outlet syndrome.

# Thrombus types and classification

The thrombus evolution is characterized by several phases, each with different embolic risk: (1) Early stage (1-6 d): Clot shows hypoechoic structure and low adherence to vessel walls; (2) 2<sup>nd</sup> stage (7-14 d): Inhomogeneous structure with alternation of echogenic and hypo/an-echoic areas; and (3) 3<sup>rd</sup> and 4<sup>th</sup> stage (> 14 d): Organization and recanalization phases: Flow appearance inside thrombus to color doppler.

The first two phases have the highest PE risk.

Basing on echogenic characteristics and appearance of thrombus, we can classify lesions as follows (Table 2): (1) Acute Venous thrombosis: Noncompressible, but the clot is not stiff and gets deformed under probe push; thrombus presents a regular profile, and the respective vein is dilated; (2) Chronic post-thrombotic change: It shows residual findings after an acute venous thrombosis. In this case clot is noncompressible, fixed and resists to pressure deformation; moreover, its profile is often nonuniform, as well as non-uniform and thickened could look vessel wall after thrombus incorporation or recanalization, while vein does not present dilated rather its caliber may be reduced (scarring setting). Sometimes it is associated to thick adhesions (synechiae), as effect of retraction forces exerted by thrombotic material, and less often to calcification sides. Echogenicity does not reflect how old thrombus is [42]. It is important to keep in mind that this persisting lesion is not a thrombus and anticoagulation therapy is not required in this case [43]; (3) Subacute thrombus: This term shouldn't be commonly used since it refers to a typical and unusual situation in which ultrasound shows a change in acute thrombus aspect few weeks apart, not includable in chronic post thrombotic change definition. These changes should occur no later than 6 mo after clot formations[44,45] (thrombus usually progresses or heals within 6 mo from its generation); (4) Scarring: It is a process that can follow a not completely recovered acute thrombosis, due to fibroblasts action on thrombus and consequent



Table 2 Lesion types			
Lesion definition	Characteristics		
Acute thrombus	Noncompressible; deformable under probe push; regular profile; dilated vein		
Subacute thrombus	Change in acute thrombus aspect few weeks apart, not includable in chronic post-thrombotic change definition (no later than 6 mo after clot formation)		
Chronic post-thrombotic change	Noncompressible; resists to pressure deformation; non-uniform profile; reduced/normal vein caliber		

fibrosis with its effects on wall thickening and synechiae production. It can determine an uncomplete stenosis which could endure for years[44,46]. Scarring has to be more correctly considered part of "chronic post thrombotic change" definition; (5) Indeterminate (equivocal): Definition utilized when it is not possible to clearly classify the lesion; (6) and Recurrent DVT: It is a thrombus formation on a chronic post thrombotic change region or a new acute venous thrombosis in a patient with a former thrombosis episode in same or contralateral leg[47-50]. It is a quite common eventuality[49], especially in patients with scarring lesions[51]. It could be not easy to recognize a new acute thrombus occurring in a chronic post thrombotic change zone [47,51,52]. Various criteria have been advanced to support diagnosis, in particular increments in compressed vein size > 4 mm or in D-dimer values, while no modifications in ultrasound scans at 1-3 d and at 7-10 d as exclusion criterion; however, their efficiency is still not clear[35,47,53-55]. Magnetic resonance also has been considered to assist differential diagnosis of recurrent DVTs from simple scars[55].

Serial scanning or D-dimer may be helpful in cases where the ultrasound does not detect clear new abnormalities, or the findings are difficult to interpret. Equivocal ultrasound findings may require serial imaging after 1-3 and 7-10 d to determine if there are any acute changes that would indicate recurrent DVT. D-dimer may also be helpful to establish if recurrent DVT is present.

# FOLLOW-UP

During anticoagulant therapy, ultrasound follow up is not necessary. For example, in the early stages of treatment, there may be minimal progression of thrombotic material, but this is not an indication to change anticoagulant or to insert a caval filter. Therefore, to evaluate "response" of venous clot to therapy does not alter treatment [36,56,57].

Ultrasound at the end of treatment may be helpful to get a clear picture of the venous district for future assessment[54].

It is unclear whether the persistence of thrombotic abnormalities can guide on a possible prolongation of anticoagulant therapy. Further studies will be needed to define the correlation between residual risk and therapy[58].

A separate mention deserves isolated distal DVT, that-as stated above-sometimes resolves or does not extend proximally without treatment and is associated with less severe complications. Thus, routine use of whole leg ultrasonography has the potential to lead to the diagnosis of DVT that does not necessarily need to be treated. Data on distal DVT remain unclear. It is not yet known who patients are at risk and how long any anticoagulation therapy should be[59].

# CONCLUSION

DVT is an often-misrecognized pathology that can cause serious clinical conditions, such as PE. Actually, ultrasound evaluation of the lower venous district can give essential information for a rapid diagnosis, especially in conditions of hemodynamic instability or when second level examinations are not readily available. Hence, compressive ultrasonography is one of the most effective tools in the emergency department in the hand of physicians. It is a non-invasive, low-cost diagnostic methodology that does not expose the patient to ionizing radiation; therefore, it is a rapid examination that should be part of the diagnostic flow chart for PE.

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