



A World of Experience and Confidence in One Anticoagulant

SANOFI 🗳



רח' בני גאון 10, פארק פולג נתניה דרום, טל': 09-8633700

ASSESSING THE BENEFITS AND RISKS

A risk assessment model for the identification of hospitalized medical patients at risk for venous thromboembolism:

The Padua Prediction Score¹

Risk assessment model (high risk of VTE: ≥4)

Baseline features	Score
Active cancer ^A	3
Previous VTE (with the exclusion of	3
superficial vein thrombosis)	
Reduced mobility ⁸	3
Already known thrombophilic condition ^c	3
Recent (≤1 month) trauma and/or surgery	2
Elderly age (≥ 70 years)	1
Heart and/or respiratory failure	1
Acute myocardial infarction or ischemic stroke	1
Acute infection and/or rheumatologic disorder	1
Obesity (BMI ≥ 30)	1
Ongoing hormonal treatment	1

A Patients with local or distant metastases and/or in whom chemotherapy or radiotherapy had been performed in the previous 6 months.

Possible exclusion criteria for pharmacological VTE prophylaxis²:

Bleeding (active)
Hypersensitivity to UFH or LMWH
Uncontrolled hypertension

Coagulopathy

Heparin-induced thrombocytopenia

Other relative or absolute exclusion criteria for phamacological prophylaxis, including surgical procedures placing patients at high risk for bleeding

Severe renal impairment³:

A dosage adjustment is required for patients with severe renal impairment (creatinine clearance < 30 ml/min),

The following dosage adjustments are recommended for the **prophylactic dosage ranges:**

Normal Dosing	Severe renal impairment
40 mg SC once daily	20 mg SC once daily
20 mg SC once daily	20 mg SC once daily

^{2.} Cohen et al. Thromb Haemost 2005; 94: 750-9

⁸ Bedrest with bathroom privileges (either due to patient's limitations or on physicians order) for at least 3 days.

^c Carriage of defects of antithrombin, protein C or S, factor V Leiden, G20210A prothrombin mutation, antiphospholipid syndrome.

Barbar S. et al. Journal of Thrombosis and Haemostasis, 8: 2450-2457

^{*} Applied in a prospective cohort study of 1180 consecutive patients admitted to a department of internal medicine in a 2-year period and followed up for up to 90 days

^{3.} Clexane prescribing Information, Israel ,October 2013