



Supplementary figure 1 Hospital sample and patient record flowchart

**Supplementary table 1** Thromboembolic risk stratification used. Based on ACCP 2012 guideline <sup>13</sup>

Risk stratum	VKA indication		
	Mechanical heart valve	Atrial fibrillation	Venous thromboembolism (VTE)
<b>High</b>	Any mitral valve prosthesis Any caged-ball or tilting disk aortic valve prosthesis iCVA or TIA within 6 months Previous thromboembolic event during VKA interruption	CHA <sub>2</sub> DS <sub>2</sub> -VASC score 8 or 9 iCVA or TIA within 3 months CHA <sub>2</sub> DS <sub>2</sub> -VASC score <8 with iCVA or TIA >3 months ago Rheumatic valvular heart disease Previous thromboembolic event during VKA interruption	VTE within 3 months Severe thrombophilia (e.g. deficiency of protein C or S or anti-thrombin, anti-phospholipid antibodies; multiple abnormalities) Previous thromboembolic event during VKA interruption
<b>Moderate</b>	Bileaflet aortic valve prosthesis and one or more riskfactors: <ul style="list-style-type: none"> <li>• Atrial fibrillation</li> <li>• iCVA or TIA</li> <li>• Hypertension</li> <li>• Diabetes</li> <li>• Heart failure</li> <li>• Age &gt;75 years</li> </ul>	CHA <sub>2</sub> DS <sub>2</sub> -VASC score 5 to 7	VTE within the past 3-12 months Recurrent VTE Non-severe thrombophilia (e.g. heterozygous factor V or prothrombin gene mutation) Active cancer
<b>Low</b>	Bileaflet aortic valve without riskfactors	CHA <sub>2</sub> DS <sub>2</sub> -VASC score <5	VTE > 12 months and no other risk factors

CHA<sub>2</sub>DS<sub>2</sub>-VASC= Congestive heart failure, hypertension, age ≥75, diabetes, iCVA/TIA, vascular disease; iCVA: ischaemic cerebrovascular accident; TIA: transient ischaemic attack

**Supplementary table 2** LMWH dose thresholds used for sub therapeutic and therapeutic dose classification. Based on the Dutch antithrombotic policy guideline <sup>31</sup>

<b>LMWH</b>	<b>Sub therapeutic dose threshold</b>	<b>Weight adjusted therapeutic dose threshold</b>
<b>Dalteparin</b>	<b>&gt; 5.000 IE/day</b>	< 55 kg: ≥ 10.000 IE/day ≥ 55 < 65 kg: ≥ 12.500 IE/day ≥ 65 < 85 kg: ≥ 15.000 IE/day ≥ 85 kg: ≥ 18.000 IE/day
<b>Nadroparin</b>	<b>&gt; 5.700 IE/day</b>	< 50 kg: ≥ 7.600 IE/day ≥ 50 < 70 kg: ≥ 11.400 IE/day ≥ 70 < 90 kg: ≥ 15.200 IE/day ≥ 90 kg: ≥ 19.000 IE/day
<b>Tinzaparin</b>	<b>&gt; 4.500 IE/day</b>	< 60 kg: ≥ 10.000 IE/day ≥ 60 < 80 kg: ≥ 14.000 IE/day ≥ 80 kg: ≥ 18.000 IE/day

**Supplementary table 3** Use of postoperative bridging anticoagulation regimen for patients' bodyweight groups

	Postoperative bridging anticoagulation regimen <sup>a</sup>		<i>P-value</i> <sup>b</sup>
	Sub therapeutic dose N=54	Therapeutic dose N=45	
<b>Patient bodyweight (kg), n (%)</b>			<.001
< 50	0 (0)	3 (100)	
50-70	5 (23.8)	16 (76.2)	
70-90	27 (58.7)	19 (41.3)	
>90	22 (75.9)	7 (24.1)	

<sup>a</sup> Excluding patients who received both therapeutic and intermediate doses (n=4) and patients with unknown doses (n=4)

<sup>b</sup> Fisher's exact test