

28 given the opportunity to express their concerns and to receive information
29 about alternative options.

30 **1 What the guideline is about**

31 **1.1 Who is the focus?**

32 **Groups that will be covered**

- 33 • Adults and young people (16 years and older) admitted to hospital as
34 inpatients, including:
 - 35 – groups considered in last version of this guideline, CG92
 - 36 – people having long-term care in hospital such as those who have
37 experienced a major traumatic event. **(new area)**
- 38
- 39 • Adults and young people (16 years and older) discharged from hospital
40 with lower-limb devices such as plaster casts and braces.
- 41 • Adults and young people (16 years and older) attending hospital for day
42 procedures including cancer treatment **(new area)** and surgery.
- 43 • Adults and young people (16 years and older) with psychiatric illness
44 admitted to community mental health hospitals or units. **(new area)**
- 45 • Special consideration will be given to:
 - 46 – pregnant women admitted to hospital and midwife units (and up to 6
47 weeks after giving birth)
 - 48 – people in whom pharmacological prophylaxis is contraindicated **(new**
49 **area)**
 - 50 – people in whom mechanical prophylaxis is contraindicated **(new area)**
 - 51 – people needing anticoagulants or antiplatelets for bridging prophylaxis or
52 other reasons. **(new area)**

53 **Groups that will not be covered**

- 54 • People with suspected or confirmed venous thromboembolism (VTE).

55 **1.2 Settings**

56 **Settings that will be covered**

- 57 • Primary and community care after hospital discharge.
- 58 • Secondary care.

59 **Settings that will not be covered**

- 60 • Community settings and hospices, except when continuing prophylaxis that
61 has been started in hospital.

62 **1.3 Activities, services or aspects of care**

63 **Key areas that will be covered**

64 Note that guideline recommendations will normally fall within licensed
65 indications; exceptionally, and only if clearly supported by evidence, use
66 outside a licensed indication may be recommended. The guideline will
67 assume that prescribers will use a medicine's summary of product
68 characteristics to inform decisions made with individual patients.

69 **Areas from the published guideline that will be updated**

- 70 1. Risk assessment
 - 71 – Patient risk factors for venous thromboembolism (VTE)
- 72 2. Methods of prophylaxis for reducing the incidence of VTE:
 - 73 – Mechanical prophylaxis including graduated compression stockings
74 (above or below the knee), intermittent pneumatic compression
75 devices (above or below the knee), electrical stimulation, continuous
76 passive motion and vena caval filters
 - 77 – Pharmacological prophylaxis including aspirin, dabigatran,
78 fondaparinux, unfractionated heparin, low molecular weight heparin
79 (LMWH), rivoroxaban and vitamin k antagonists [for example
80 warfarin]
 - 81 – Timing of prophylaxis
 - 82 – Duration of prophylaxis
- 83 3. Information and support

- 84 – Content of information on prophylaxis methods and VTE provided to
85 patients and their family members or carers.

86 ***Areas not in the published guideline that will be included in the update***

- 87 1. Risk assessment
88 – Risk prediction tools (for bleeding or VTE)
89 – Reassessment of risk
90 2. Methods of prophylaxis
91 – New interventions (apixaban and geko devices)
92 – Bridging prophylaxis

93 **Areas that will not be covered**

94 ***Areas from the published guideline that will not be updated***

- 95 1 Methods of prophylaxis
96 – Early mobilisation and leg exercises
97 – Physiotherapy
98 – Hydration
99 – Regional compared with general anaesthetic.

100 ***Areas from the published guideline that will be removed***

- 101 1. Methods of prophylaxis
102 – Leg elevation

103 ***Areas not covered by the published guideline or the update***

- 104 1. Secondary prevention of VTE
105

106 Recommendations in areas that are not being updated may be edited to
107 ensure that they meet current editorial standards, and reflect the current policy
108 and practice context.

109 **1.4 *Economic aspects***

- 110 We will take economic aspects into account when making recommendations.
111 We will develop an economic plan that states for each review question (or key

112 area in the scope) whether economic considerations are relevant, and if so
113 whether this is an area that should be prioritised for economic modelling and
114 analysis. We will review the economic evidence and carry out economic
115 analyses, using an NHS and personal social services (PSS) perspective, as
116 appropriate.

117 **1.5 Key issues and questions**

118 While writing this scope we have identified the following key issues, and key
119 questions related to them:

120 1. Risk assessment:

121 1.1 What is the accuracy of individual risk assessment or prediction tools
122 in predicting the likelihood of VTE (deep vein thrombosis [DVT] or
123 pulmonary embolism [PE]) in a patient who is admitted to hospital?

124 1.2 What is the accuracy of individual risk assessment or prediction tools
125 in predicting the likelihood of VTE (DVT or PE) in patients who are
126 having day procedures (including surgery and chemotherapy) at
127 hospital?

128 1.3 What is the accuracy of individual risk assessment or prediction tools
129 in predicting the likelihood of major bleeding or the risk of bleeding in a
130 patient who is admitted to hospital?

131 1.4 What is the accuracy of individual risk assessment or prediction tools
132 in predicting the likelihood of major bleeding or the risk of in patients who
133 are having day procedures (including surgery and chemotherapy) at
134 hospital?

135 1.5 How clinically and cost effective are risk tools at reducing the rates of
136 VTE (DVT or PE) in patients who are admitted to hospital?

137 1.6 How clinically and cost effective are risk tools at reducing the rates of
138 VTE (DVT or PE) in patients who are having day procedures (including
139 surgery and chemotherapy) at hospital?

140 1.7 How effective is reassessment of patients who are admitted to or
141 having day procedures at hospital?

142

143 If appropriate evidence is not identified from the questions above (1.1 to 1.7)
 144 the following 2 questions may also be considered:

145 1.8 What are the individual risk factors for VTE (DVT or PE) in patients
 146 who are admitted to hospital?

147 1.9 What are the individual risk factors for VTE (DVT or PE) in patients
 148 who are having day procedures (including surgery and chemotherapy) at
 149 hospital?

150

151 2. Prophylaxis:

152 Each of the following questions will investigate individual populations
 153 separately. Populations include:

154 • people having the following types of surgery:

- 155 – elective hip surgery
- 156 – elective knee surgery
- 157 – hip fracture
- 158 – knee arthroscopy
- 159 – other orthopaedic surgery
- 160 – abdominal surgery (bariatric, liver, gastrointestinal, gynaecological,
 161 laparoscopic, thoracic and urological)
- 162 – cranial or spinal surgery
- 163 – cardiac surgery
- 164 – vascular surgery

165 • people discharged wearing lower-limb devices

166 • people being treated for:

- 167 – major trauma
- 168 – spinal injury
- 169 – stroke
- 170 – acute coronary syndromes
- 171 – cancer

172 • people attending hospital as acute or chronic medical admissions

173 • people with central venous catheters

174 • people having palliative care

- 175 • pregnant women
176 • psychiatric patients
177 • people in hospital for long-term care.

178 Each of the questions will consider the following settings, if appropriate:
179 people in hospital and those having day procedures (including surgery,
180 chemotherapy)

181 Each of the questions will include the following prophylaxis methods, if
182 applicable:

- 183 • mechanical prophylaxis, including:
184 – graduated compression stockings (above or below knee)
185 – intermittent pneumatic compression devices (above or below knee)
186 – electrical stimulation (including geko devices)
187 – continuous passive motion
188 – vena caval filters.

189

- 190 • pharmacological prophylaxis, including:
191 – apixaban
192 – aspirin
193 – dabigatran
194 – fondaparinux
195 – unfractionated heparin
196 – low molecular weight heparin (LMWH)
197 – rivaroxaban
198 – vitamin k antagonists (for example warfarin).

199 2.1 What is the effectiveness of different pharmacological and
200 mechanical prophylaxis strategies (alone or in combination)?

201 2.2 What is the effectiveness of vena caval filters in people admitted to
202 hospital who are at high risk of DVT or PE admitted to hospital?

203 2.3 What is the most effective timing for starting prophylaxis with LMWH
204 for people having surgery?

- 205 2.4 What is the most effective prophylaxis duration (covering the time in
206 hospital only or continuing after discharge)?
- 207 2.5 What is the most effective prophylaxis strategy for inpatients in
208 whom pharmacological prophylaxis is contraindicated?
- 209 2.6 What is the most effective prophylaxis strategy for inpatients in
210 whom mechanical prophylaxis is contraindicated?
- 211 2.7 How should VTE be prevented for patients in whom both mechanical
212 and pharmacological prophylaxis are contraindicated?
- 213 2.8 What is the most effective prophylaxis strategy in bridging patients
214 who are already using anticoagulants or antiplatelets for other reasons in
215 reducing the incidence of VTE?
216
- 217 3. Information for patients, family members and carers:
- 218 3.1 What specific information should be provided to people who need
219 VTE prophylaxis?
- 220 3.2 What information do patients, their family members and carers say
221 they want about VTE prophylaxis?

222 **1.6 Main outcomes**

223 The main outcomes that will be considered when searching for and assessing
224 the evidence are:

- 225 1. All-cause mortality
- 226 2. Pulmonary embolism
- 227 3. Fatal pulmonary embolism
- 228 4. Deep vein thrombosis (symptomatic or asymptomatic)
- 229 5. Major bleeding
- 230 6. Fatal bleeding
- 231 7. Heparin-induced thrombocytopenia
- 232 8. Post-thrombotic syndrome
- 233 9. Pulmonary hypertension
- 234 10. Quality of life (validated scores)
- 235 11. Hospital length of stay
- 236 12. Readmission

237 13. Neurological events (for example haemorrhagic stroke)

238 **2 Links with other NICE guidance, NICE quality** 239 **standards and NICE Pathways**

240 **2.1 NICE guidance**

- 241 • [Venous thromboembolism in adults admitted to hospital: reducing the risk](#)
242 (2010) NICE guideline CG92
- 243 • [Venous thromboembolic diseases: the management of venous](#)
244 [thromboembolic diseases and the role of thrombophilia testing](#) (2012)
245 NICE clinical guideline 144
- 246 • [Caesarean section](#) (2011) NICE clinical guideline 132
- 247 • [Stroke: Diagnosis and initial management of acute stroke and transient](#)
248 [ischaemic attack \(TIA\)](#) (2008) NICE clinical guideline 68.
- 249 • [Apixaban for the prevention of venous thromboembolism after total hip or](#)
250 [knee replacement in adults \(2012\)](#) NICE technology appraisal 245
- 251 • [Dabigatran etexilate for the prevention of venous thromboembolism after](#)
252 [hip or knee replacement surgery in adults](#) (2008) NICE technology
253 appraisal 157.
- 254 • [Rivaroxaban for the prevention of venous thromboembolism after total hip](#)
255 [or total knee replacement in adults](#) (2009) NICE technology appraisal 170
- 256 • [The geko device for reducing the risk of venous thromboembolism](#) (2014)
257 NICE medical technology guidance 19.

258 **NICE guidance that will be updated by this guideline**

- 259 • [Venous thromboembolism in adults admitted to hospital: reducing the risk](#)
260 (2010) NICE guideline CG92

261 **NICE guidance about the experience of people using NHS services**

262 NICE has produced the following guidance on the experience of people using
263 the NHS. This guideline will not include additional recommendations on these
264 topics unless there are specific issues related to VTE:

- 265 • [Patient experience in adult NHS services](#) (2012) NICE guideline CG138

- 266 • [Service user experience in adult mental health](#) (2011) NICE guideline
 267 CG136
- 268 • [Medicines adherence](#) (2009) NICE guideline CG76

269 **2.2 NICE quality standards**

270 **NICE quality standards that may need to be revised or updated when** 271 **this guideline is published**

- 272 • [Venous thromboembolism prevention](#) (2010) NICE quality standard 3.

273 **2.3 NICE Pathways**

274 When this guideline is published it will update the existing NICE pathway on
 275 [venous thromboembolism](#). NICE Pathways bring together all related NICE
 276 guidance and associated products on a topic in an interactive topic-based flow
 277 chart.

278 **3 Context**

279 **3.1 Key facts and figures**

280 [Hospital Episode Statistics](#) showed that in 2013–14 there were 24,725
 281 admissions for pulmonary embolism and 19,463 for DVT in England, resulting
 282 in 205,448 and 67,028 bed-days and 47,594 and 25,958 finished consultant
 283 episodes respectively. In 2013, in England and Wales there were 2,191
 284 deaths recorded as due to pulmonary embolism (PE) and 2,816 due to deep
 285 vein thrombosis (DVT), but the actual number of people dying from these
 286 conditions is likely to be higher because of misdiagnosis and the failure to
 287 recognise VTE as the underlying cause. These figures relate to all VTE;
 288 hospital-acquired VTE will account for a proportion of them.

289 **3.2 Current practice**

290 In 2010, the CQUIN target introduced a payment linked to at least 90% of
 291 adults being risk assessed on admission to hospital. Figures reporting the
 292 uptake of some of the recommendations in CG92 are reported on [NICE's](#)
 293 [website](#). Recent evidence also estimates that the national mortality rate from

294 VTE has fallen by 8–9% since the recommendations in CG92 were
295 introduced.

296 In addition, since the publication of the last version of the guideline, [CG92](#),
297 two new interventions for preventing venous thromboembolism (VTE) have
298 become available: apixaban and geko devices.

299 **3.3 Policy, legislation, regulation and commissioning**

300 **Policy**

301 The [National VTE prevention programme](#) was launched in England in 2010 by
302 the Department of Health. This included the mandatory VTE risk assessment
303 of 95% of all people admitted to hospital. A risk assessment tool was created
304 by the Department of Health and this was incorporated into the last version of
305 this guideline. Risk assessment will be a key part of this update.

306 **4 Further information**

This is the draft scope for consultation with registered stakeholders. The
consultation dates are 11 December 2015 to 20 January 2016.

The guideline is expected to be published in February 2018.

You can follow progress of the [guideline](#).

Our website has information about how [NICE guidelines](#) are developed.

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