ANNEX I

SUMMARY OF PRODUCT CHARACTERISTICS
1. NAME OF THE MEDICINAL PRODUCT
Brilique 60 mg film-coated tablets

2. QUALITATIVE AND QUANTITATIVE COMPOSITION
Each film-coated tablet contains 60 mg ticagrelor.
For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM
Film-coated tablet (tablet).
Round, biconvex, pink tablets marked with ‘60’ above ‘T’ on one side and plain on the other.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications
Brilique, co-administered with acetylsalicylic acid (ASA), is indicated for the prevention of atherothrombotic events in adult patients with
- acute coronary syndromes (ACS) or
- a history of myocardial infarction (MI) and a high risk of developing an atherothrombotic event (see sections 4.2 and 5.1).

4.2 Posology and method of administration

Posology
Patients taking Brilique should also take a daily low maintenance dose of ASA 75-150 mg, unless specifically contraindicated.

Acute coronary syndromes
Brilique treatment should be initiated with a single 180 mg loading dose (two tablets of 90 mg) and then continued at 90 mg twice daily. Treatment with Brilique 90 mg twice daily is recommended for 12 months in ACS patients unless discontinuation is clinically indicated (see section 5.1).

History of myocardial infarction
Brilique 60 mg twice daily is the recommended dose when an extended treatment is required for patients with a history of MI of at least one year and a high risk of an atherothrombotic event (see section 5.1). Treatment may be started without interruption as continuation therapy after the initial one-year treatment with Brilique 90 mg or other adenosine diphosphate (ADP) receptor inhibitor therapy in ACS patients with a high risk of an atherothrombotic event. Treatment can also be initiated up to 2 years from the MI, or within one year after stopping previous ADP receptor inhibitor treatment. There are limited data on the efficacy and safety of Brilique beyond 3 years of extended treatment.
If a switch is needed, the first dose of Brilique should be administered 24 hours following the last
dose of the other antiplatelet medication.

Missed dose
Lapses in therapy should also be avoided. A patient who misses a dose of Brilique should take only
one tablet (their next dose) at its scheduled time.

Special populations
Elderly
No dose adjustment is required in elderly (see section 5.2).

Renal impairment
No dose adjustment is necessary for patients with renal impairment (see section 5.2). No
information is available concerning treatment of patients on renal dialysis and therefore ticagrelor
is not recommended in these patients.

Hepatic impairment
Ticagrelor has not been studied in patients with severe hepatic impairment and its use in these
patients is therefore contraindicated (see section 4.3). Only limited information is available in
patients with moderate hepatic impairment. Dose adjustment is not recommended, but ticagrelor
should be used with caution (see sections 4.4 and 5.2). No dose adjustment is necessary for patients
with mild hepatic impairment (see section 5.2).

Paediatric population
The safety and efficacy of ticagrelor in children below the age of 18 years have not been
established. No data are available.

Method of administration
For oral use.
Brilique can be administered with or without food.
For patients who are unable to swallow the tablet(s) whole, the tablets can be crushed to a fine
powder and mixed in half a glass of water and drunk immediately. The glass should be rinsed with
a further half glass of water and the contents drunk. The mixture can also be administered via a
nasogastric tube (CH8 or greater). It is important to flush the nasogastric tube through with water
after administration of the mixture.

4.3 Contraindications

- Hypersensitivity to the active substance or to any of the excipients listed in section 6.1 (see
section 4.8).
- Active pathological bleeding.
- History of intracranial haemorrhage (see section 4.8).
- Severe hepatic impairment (see sections 4.2, 4.4 and 5.2).
- Co-administration of ticagrelor with strong CYP3A4 inhibitors (e.g. ketoconazole,
clarithromycin, nefazodone, ritonavir, and atazanavir) is contraindicated, as
co-administration may lead to a substantial increase in exposure to ticagrelor (see
section 4.5).
4.4 Special warnings and precautions for use

Bleeding risk
The use of ticagrelor in patients at known increased risk for bleeding should be balanced against the benefit in terms of prevention of atherothrombotic events (see sections 4.8 and 5.1). If clinically indicated, ticagrelor should be used with caution in the following patient groups:

- Patients with a propensity to bleed (e.g. due to recent trauma, recent surgery, coagulation disorders, active or recent gastrointestinal bleeding). The use of ticagrelor is contraindicated in patients with active pathological bleeding, in those with a history of intracranial haemorrhage, and in patients with severe hepatic impairment (see section 4.3).
- Patients with concomitant administration of medicinal products that may increase the risk of bleeding (e.g. non-steroidal anti-inflammatory drugs (NSAIDs), oral anticoagulants and/or fibrinolytics) within 24 hours of ticagrelor dosing.

Platelet transfusion did not reverse the antiplatelet effect of ticagrelor in healthy volunteers and is unlikely to be of clinical benefit in patients with bleeding. Since co-administration of ticagrelor with desmopressin did not decrease template-bleeding time, desmopressin is unlikely to be effective in managing clinical bleeding events (see section 4.5).

Antifibrinolytic therapy (aminocaproic acid or tranexamic acid) and/or recombinant factor VIIa therapy may increase haemostasis. Ticagrelor may be resumed after the cause of bleeding has been identified and controlled.

Surgery
Patients should be advised to inform physicians and dentists that they are taking ticagrelor before any surgery is scheduled and before any new medicinal product is taken.

In PLATO patients undergoing coronary artery bypass grafting (CABG), ticagrelor had more bleeding than clopidogrel when stopped within 1 day prior to surgery but a similar rate of major bleeds compared to clopidogrel after stopping therapy 2 or more days before surgery (see section 4.8). If a patient is to undergo elective surgery and antiplatelet effect is not desired, ticagrelor should be discontinued 7 days prior to surgery (see section 5.1).

Patients with prior ischaemic stroke
ACS patients with prior ischaemic stroke can be treated with Brilique for up to 12 months (PLATO study).

In PEGASUS, patients with history of MI with prior ischaemic stroke were not included. Therefore, in the absence of data, treatment beyond one year is not recommended in these patients.

Hepatic impairment
Use of ticagrelor is contraindicated in patients with severe hepatic impairment (see sections 4.2 and 4.3). There is limited experience with ticagrelor in patients with moderate hepatic impairment, therefore, caution is advised in these patients (see sections 4.2 and 5.2).

Patients at risk for bradycardic events
Due to observations of mostly asymptomatic ventricular pauses in an earlier clinical study, patients with an increased risk of bradycardic events (e.g. patients without a pacemaker who have sick sinus syndrome, 2nd or 3rd degree AV block or bradycardic-related syncope) were excluded from the main studies evaluating the safety and efficacy of ticagrelor. Therefore, due to the limited clinical experience, ticagrelor should be used with caution in these patients (see section 5.1).
In addition, caution should be exercised when administering ticagrelor concomitantly with medicinal products known to induce bradycardia. However, no evidence of clinically significant adverse reactions was observed in the PLATO trial after concomitant administration with one or more medicinal products known to induce bradycardia (e.g. 96% beta blockers, 33% calcium channel blockers diltiazem and verapamil, and 4% digoxin) (see section 4.5).

During the Holter substudy in PLATO, more patients had ventricular pauses ≥3 seconds with ticagrelor than with clopidogrel during the acute phase of their ACS. The increase in Holter-detected ventricular pauses with ticagrelor was higher in patients with chronic heart failure (CHF) than in the overall study population during the acute phase of ACS, but not at one month with ticagrelor or compared to clopidogrel. There were no adverse clinical consequences associated with this imbalance (including syncope or pacemaker insertion) in this patient population (see section 5.1).

**Dyspnoea**

Dyspnoea was reported in patients treated with ticagrelor. Dyspnoea is usually mild to moderate in intensity and often resolves without need for treatment discontinuation. Patients with asthma/chronic obstructive pulmonary disease (COPD) may have an increased absolute risk of experiencing dyspnoea with ticagrelor. Ticagrelor should be used with caution in patients with history of asthma and/or COPD. The mechanism has not been elucidated. If a patient reports new, prolonged or worsened dyspnoea this should be investigated fully and if not tolerated, treatment with ticagrelor should be stopped. For further details see section 4.8.

**Creatinine elevations**

Creatinine levels may increase during treatment with ticagrelor. The mechanism has not been elucidated. Renal function should be checked according to routine medical practice. In patients with ACS, it is recommended that renal function is also checked one month after initiating the treatment with ticagrelor, paying special attention to patients ≥75 years, patients with moderate/severe renal impairment and those receiving concomitant treatment with an angiotensin receptor blocker (ARB).

**Uric acid increase**

Hyperuricaemia may occur during treatment with ticagrelor (see section 4.8). Caution is advised in patients with history of hyperuricaemia or gouty arthritis. As a precautionary measure, the use of ticagrelor in patients with uric acid nephropathy is discouraged.

**Other**

Based on a relationship observed in PLATO between maintenance ASA dose and relative efficacy of ticagrelor compared to clopidogrel, co-administration of ticagrelor and high maintenance dose ASA (>300 mg) is not recommended (see section 5.1).

**Premature discontinuation**

Premature discontinuation with any antiplatelet therapy, including Brilique, could result in an increased risk of cardiovascular (CV) death or MI due to the patient’s underlying disease. Therefore, premature discontinuation of treatment should be avoided.

### 4.5 Interaction with other medicinal products and other forms of interaction

Ticagrelor is primarily a CYP3A4 substrate and a mild inhibitor of CYP3A4. Ticagrelor is also a P-glycoprotein (P-gp) substrate and a weak P-gp inhibitor and may increase the exposure of P-gp substrates.
Effects of other medicinal products on ticagrelor

**Medicinal products metabolised by CYP3A4**

**CYP3A4 inhibitors**

- **Strong CYP3A4 inhibitors** – Co-administration of ketoconazole with ticagrelor increased the ticagrelor $C_{\text{max}}$ and AUC equal to 2.4-fold and 7.3-fold, respectively. The $C_{\text{max}}$ and AUC of the active metabolite were reduced by 89% and 56%, respectively. Other strong inhibitors of CYP3A4 (clarithromycin, nefazodone, ritonavir, and atazanavir) would be expected to have similar effects and therefore concomitant use of strong CYP3A4 inhibitors with ticagrelor is contraindicated (see section 4.3).

- **Moderate CYP3A4 inhibitors** – Co-administration of diltiazem with ticagrelor increased the ticagrelor $C_{\text{max}}$ by 69% and AUC to 2.7-fold and decreased the active metabolite $C_{\text{max}}$ by 38% and AUC was unchanged. There was no effect of ticagrelor on diltiazem plasma levels. Other moderate CYP3A4 inhibitors (e.g. amprenavir, aprepitant, erythromycin and fluconazole) would be expected to have a similar effect and can as well be co-administered with ticagrelor.

**CYP3A inducers**

Co-administration of rifampicin with ticagrelor decreased ticagrelor $C_{\text{max}}$ and AUC by 73% and 86%, respectively. The $C_{\text{max}}$ of the active metabolite was unchanged and the AUC was decreased by 46%, respectively. Other CYP3A inducers (e.g. phenytoin, carbamazepine and phenobarbital) would be expected to decrease the exposure to ticagrelor as well. Co-administration of ticagrelor with potent CYP3A inducers may decrease exposure and efficacy of ticagrelor, therefore, their concomitant use with ticagrelor is discouraged.

**Cyclosporine (P-gp and CYP3A inhibitor)**

Co-administration of cyclosporine (600 mg) with ticagrelor increased ticagrelor $C_{\text{max}}$ and AUC equal to 2.3-fold and 2.8-fold, respectively. The AUC of the active metabolite was increased by 32% and $C_{\text{max}}$ was decreased by 15% in the presence of cyclosporine.

No data are available on concomitant use of ticagrelor with other active substances that also are potent P-gp inhibitors and moderate CYP3A4 inhibitors (e.g. verapamil, quinidine) that also may increase ticagrelor exposure. If the association cannot be avoided, their concomitant use should be made with caution.

**Others**

Clinical pharmacology interaction studies showed that co-administration of ticagrelor with heparin, enoxaparin and ASA or desmopressin did not have any effect on the pharmacokinetics of ticagrelor or the active metabolite or on ADP-induced platelet aggregation compared with ticagrelor alone. If clinically indicated, medicinal products that alter haemostasis should be used with caution in combination with ticagrelor.

A 2-fold increase of ticagrelor exposure was observed after daily consumption of large quantities of grapefruit juice (3x200 ml). This magnitude of increased exposure is not expected to be clinically relevant to most patients.

Effects of ticagrelor on other medicinal products

**Medicinal products metabolised by CYP3A4**

- **Simvastatin** – Co-administration of ticagrelor with simvastatin increased simvastatin $C_{\text{max}}$ by 81% and AUC by 56% and increased simvastatin acid $C_{\text{max}}$ by 64% and AUC by 52% with
some individual increases equal to 2 to 3-fold. Co-administration of ticagrelor with doses of simvastatin exceeding 40 mg daily could cause adverse effects of simvastatin and should be weighed against potential benefits. There was no effect of simvastatin on ticagrelor plasma levels. Ticagrelor may have similar effect on lovastatin. The concomitant use of ticagrelor with doses of simvastatin or lovastatin greater than 40 mg is not recommended.

- **Atorvastatin** – Co-administration of atorvastatin and ticagrelor increased atorvastatin acid $C_{\text{max}}$ by 23% and AUC by 36%. Similar increases in AUC and $C_{\text{max}}$ were observed for all atorvastatin acid metabolites. These increases are not considered clinically significant.
- A similar effect on other statins metabolised by CYP3A4 cannot be excluded. Patients in PLATO receiving ticagrelor took a variety of statins, with no concern of an association with statin safety among the 93% of the PLATO cohort taking these medicinal products.

Ticagrelor is a mild CYP3A4 inhibitor. Co-administration of ticagrelor and CYP3A4 substrates with narrow therapeutic indices (i.e. cisapride or ergot alkaloids) is not recommended, as ticagrelor may increase the exposure to these medicinal products.

**P-gp substrates (including digoxin, cyclosporine)**
Concomitant administration of ticagrelor increased the digoxin $C_{\text{max}}$ by 75% and AUC by 28%. The mean trough digoxin levels were increased about 30% with ticagrelor co-administration with some individual maximum increases to 2-fold. In the presence of digoxin, the $C_{\text{max}}$ and AUC of ticagrelor and its active metabolite were not affected. Therefore, appropriate clinical and/or laboratory monitoring is recommended when giving narrow therapeutic index P-gp dependent medicinal products like digoxin concomitantly with ticagrelor.

There was no effect of ticagrelor on cyclosporine blood levels. Effect of ticagrelor on other P-gp substrates has not been studied.

**Medicinal products metabolised by CYP2C9**
Co-administration of ticagrelor with tolbutamide resulted in no change in the plasma levels of either medicinal product, which suggests that ticagrelor is not a CYP2C9 inhibitor and unlikely to alter the CYP2C9 mediated metabolism of medicinal products like warfarin and tolbutamide.

**Oral contraceptives**
Co-administration of ticagrelor and levonorgestrel and ethinyl estradiol increased ethinyl estradiol exposure approximately 20% but did not alter the pharmacokinetics of levonorgestrel. No clinically relevant effect on oral contraceptive efficacy is expected when levonorgestrel and ethinyl estradiol are co-administered with ticagrelor.

**Medicinal products known to induce bradycardia**
Due to observations of mostly asymptomatic ventricular pauses and bradycardia, caution should be exercised when administering ticagrelor concomitantly with medicinal products known to induce bradycardia (see section 4.4). However, no evidence of clinically significant adverse reactions was observed in the PLATO trial after concomitant administration with one or more medicinal products known to induce bradycardia (e.g. 96% beta blockers, 33% calcium channel blockers diltiazem and verapamil, and 4% digoxin).

**Other concomitant therapy**
In clinical studies, ticagrelor was commonly administered with ASA, proton pump inhibitors, statins, beta-blockers, angiotensin converting enzyme (ACE) inhibitors and angiotensin receptor blockers as needed for concomitant conditions for long-term and also heparin, low molecular weight heparin and intravenous GpIIb/IIIa inhibitors for short durations (see section 5.1). No evidence of clinically significant adverse interactions with these medicinal products was observed.
Co-administration of ticagrelor with heparin, enoxaparin or desmopressin had no effect on activated partial thromboplastin time (aPTT), activated coagulation time (ACT) or factor Xa assays. However, due to potential pharmacodynamic interactions, caution should be exercised with the concomitant administration of ticagrelor with medicinal products known to alter haemostasis.

Due to reports of cutaneous bleeding abnormalities with SSRIs (e.g. paroxetine, sertraline and citalopram), caution is advised when administering SSRIs with ticagrelor as this may increase the risk of bleeding.

4.6 Fertility, pregnancy and lactation

Women of childbearing potential
Women of childbearing potential should use appropriate contraceptive measures to avoid pregnancy during ticagrelor therapy.

Pregnancy
There are no or limited amount of data from the use of ticagrelor in pregnant women. Studies in animals have shown reproductive toxicity (see section 5.3). Ticagrelor is not recommended during pregnancy.

Breast-feeding
Available pharmacodynamic/toxicological data in animals have shown excretion of ticagrelor and its active metabolites in milk (see section 5.3). A risk to newborns/infants cannot be excluded. A decision must be made whether to discontinue breast-feeding or to discontinue/abstain from ticagrelor therapy taking into account the benefit of breast-feeding for the child and the benefit of therapy for the woman.

Fertility
Ticagrelor had no effect on male or female fertility in animals (see section 5.3).

4.7 Effects on ability to drive and use machines

Ticagrelor has no or negligible influence on the ability to drive and use machines. During treatment with ticagrelor, dizziness and confusion have been reported. Therefore, patients who experience these symptoms should be cautious while driving or using machines.

4.8 Undesirable effects

Summary of the safety profile
The safety profile of ticagrelor has been evaluated in two large phase 3 outcome trials (PLATO and PEGASUS) including more than 39,000 patients (see section 5.1).

In PLATO, patients on ticagrelor had a higher incidence of discontinuation due to adverse events than clopidogrel (7.4% vs. 5.4%). In PEGASUS, patients on ticagrelor had a higher incidence of discontinuation due to adverse events compared to ASA therapy alone (16.1% for ticagrelor 60 mg with ASA vs. 8.5% for ASA therapy alone). The most commonly reported adverse reactions in patients treated with ticagrelor were bleeding and dyspnoea (see section 4.4).

Tabulated list of adverse reactions
The following adverse reactions have been identified following studies or have been reported in post-marketing experience with ticagrelor (Table 1).
Adverse reactions are listed by MedDRA System Organ Class (SOC). Within each SOC the adverse reactions are ranked by frequency category. Frequency categories are defined according to the following conventions: Very common (≥1/10), common (≥1/100 to <1/10), uncommon (≥1/1,000 to <1/100), rare (≥1/10,000 to <1/1,000), very rare (<1/10,000), not known (cannot be estimated from the available data).

**Table 1 – Adverse reactions by frequency and system organ class (SOC)**

<table>
<thead>
<tr>
<th>System Organ Classification</th>
<th>Very Common</th>
<th>Common</th>
<th>Uncommon</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neoplasms benign, malignant and unspecified (including cysts and polyps)</td>
<td></td>
<td></td>
<td>Tumour bleedings&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td>Blood and lymphatic system disorders</td>
<td>Blood disorder bleedings&lt;sup&gt;b&lt;/sup&gt;</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Immune system disorders</td>
<td></td>
<td></td>
<td>Hypersensitivity including angioedema&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td>Metabolism and nutrition disorders</td>
<td>Hyperuricaemia&lt;sup&gt;d&lt;/sup&gt;</td>
<td>Gout/Gouty Arthritis</td>
<td></td>
</tr>
<tr>
<td>Psychiatric disorders</td>
<td></td>
<td></td>
<td>Confusion</td>
</tr>
<tr>
<td>Nervous system disorders</td>
<td>Dizziness, Syncope, Headache</td>
<td>Intracranial haemorrhage</td>
<td></td>
</tr>
<tr>
<td>Eye disorders</td>
<td></td>
<td></td>
<td>Eye haemorrhage&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td>Ear and labyrinth disorders</td>
<td>Vertigo</td>
<td>Ear haemorrhage</td>
<td></td>
</tr>
<tr>
<td>Vascular disorders</td>
<td></td>
<td></td>
<td>Hypotension</td>
</tr>
<tr>
<td>Respiratory, thoracic and mediastinal disorders</td>
<td>Dyspnoea</td>
<td>Respiratory system bleedings&lt;sup&gt;f&lt;/sup&gt;</td>
<td></td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td>Gastrointestinal haemorrhage&lt;sup&gt;g&lt;/sup&gt;, Diarrhoea, Nausea, Dyspepsia, Constipation</td>
<td></td>
<td>Retroperitoneal haemorrhage</td>
</tr>
<tr>
<td>Skin and subcutaneous tissue disorders</td>
<td>Subcutaneous or dermal bleeding&lt;sup&gt;h&lt;/sup&gt;, Rash, Pruritus</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Musculoskeletal connective tissue and bone</td>
<td></td>
<td></td>
<td>Muscular bleedings&lt;sup&gt;i&lt;/sup&gt;</td>
</tr>
<tr>
<td>Renal and urinary disorders</td>
<td>Urinary tract bleeding&lt;sup&gt;i&lt;/sup&gt;</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Reproductive system and breast disorders</td>
<td></td>
<td></td>
<td>Reproductive system bleedings&lt;sup&gt;k&lt;/sup&gt;</td>
</tr>
<tr>
<td>System Organ Classification</td>
<td>Very Common</td>
<td>Common</td>
<td>Uncommon</td>
</tr>
<tr>
<td>----------------------------</td>
<td>-------------</td>
<td>--------</td>
<td>----------</td>
</tr>
<tr>
<td>Investigations</td>
<td>Blood creatinine increased(^d)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Injury, poisoning and procedural complications</td>
<td>Post procedural haemorrhage, Traumatic bleedings(^1)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

\(^a\) e.g. bleeding from bladder cancer, gastric cancer, colon cancer  
\(^b\) e.g. increased tendency to bruise, spontaneous haematoma, haemorrhagic diathesis  
\(^c\) Identified in post-marketing experience  
\(^d\) Frequencies derived from lab observations (Uric acid increases to >upper limit of normal from baseline below or within reference range. Creatinine increases of >50% from baseline.) and not crude adverse event report frequency.  
\(^e\) e.g. conjunctival, retinal, intraocular bleeding  
\(^f\) e.g. epistaxis, haemoptysis  
\(^g\) e.g. gingival bleeding, rectal haemorrhage, gastric ulcer haemorrhage  
\(^h\) e.g. ecchymosis, skin haemorrhage, petechiae  
\(^i\) e.g. haemarthrosis, muscle haemorrhage  
\(^j\) e.g. haematuria, cystitis haemorrhagic  
\(^k\) e.g. vaginal haemorrhage, haematospermia, postmenopausal haemorrhage  
\(^l\) e.g. contusion, traumatic haematomata, traumatic haemorrhage

Description of selected adverse reactions

**Bleeding**

**Bleeding findings in PLATO**

Overall outcome of bleeding rates in the PLATO study are shown in Table 2.

**Table 2 – Analysis of overall bleeding events, Kaplan-Meier estimates at 12 months (PLATO)**

<table>
<thead>
<tr>
<th></th>
<th>Ticagrelor 90 mg twice daily</th>
<th>Clopidogrel</th>
<th>(p)-value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>PLATO Total Major</td>
<td>11.6</td>
<td>11.2</td>
<td>0.4336</td>
</tr>
<tr>
<td>PLATO Major Fatal/Life-Threatening</td>
<td>5.8</td>
<td>5.8</td>
<td>0.6988</td>
</tr>
<tr>
<td>Non-CABG PLATO Major</td>
<td>4.5</td>
<td>3.8</td>
<td>0.0264</td>
</tr>
<tr>
<td>Non-Procedural PLATO Major</td>
<td>3.1</td>
<td>2.3</td>
<td>0.0058</td>
</tr>
<tr>
<td>PLATO Total Major + Minor</td>
<td>16.1</td>
<td>14.6</td>
<td>0.0084</td>
</tr>
<tr>
<td>Non-Procedural PLATO Major + Minor</td>
<td>5.9</td>
<td>4.3</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>TIMI-defined Major</td>
<td>7.9</td>
<td>7.7</td>
<td>0.5669</td>
</tr>
<tr>
<td>TIMI-defined Major + Minor</td>
<td>11.4</td>
<td>10.9</td>
<td>0.3272</td>
</tr>
</tbody>
</table>

**Bleeding category definitions:**

**Major Fatal/Life-threatening Bleed:** Clinically apparent with >50 g/l decrease in haemoglobin or ≥4 red cell units transfused; or fatal; or intracranial; or intrapericardial with cardiac tamponade; or with hypovolaemic shock or severe hypotension requiring pressors or surgery.

**Major Other:** Clinically apparent with 30-50 g/L decrease in haemoglobin or 2-3 red cell units transfused; or significantly disabling.

**Minor Bleed:** Requires medical intervention to stop or treat bleeding.

**TIMI Major Bleed:** Clinically apparent with >50 g/l decrease in haemoglobin or intracranial haemorrhage.

**TIMI Minor Bleed:** Clinically apparent with 30-50 g/l decrease in haemoglobin.

\(*p\)-value calculated from Cox proportional hazards model with treatment group as the only explanatory variable.

Ticagrelor and clopidogrel did not differ in rates of PLATO Major Fatal/Life-threatening bleeding, PLATO total Major bleeding, TIMI Major bleeding, or TIMI Minor bleeding (Table 2). However, more PLATO combined Major + Minor bleeding occurred with ticagrelor compared with
Few patients in PLATO had fatal bleeds: 20 (0.2%) for ticagrelor and 23 (0.3%) for clopidogrel (see section 4.4).

Age, sex, weight, race, geographic region, concurrent conditions, concomitant therapy, and medical history, including a previous stroke or transient ischaemic attack, all did not predict either overall or non-procedural PLATO Major bleeding. Thus no particular group was identified at risk for any subset of bleeding.

*CABG-related bleeding:* In PLATO, 42% of the 1584 patients (12% of cohort) who underwent coronary artery bypass graft (CABG) surgery had a PLATO Major Fatal/Life-threatening bleeding with no difference between treatment groups. Fatal CABG bleeding occurred in 6 patients in each treatment group (see section 4.4).

*Non-CABG related bleeding and non-procedural related bleeding:* Ticagrelor and clopidogrel did not differ in non-CABG PLATO-defined Major Fatal/Life-threatening bleeding, but PLATO-defined Total Major, TIMI Major, and TIMI Major + Minor bleeding were more common with ticagrelor. Similarly, when removing all procedure related bleeds, more bleeding occurred with ticagrelor than with clopidogrel (Table 2). Discontinuation of treatment due to non-procedural bleeding was more common for ticagrelor (2.9%) than for clopidogrel (1.2%; p<0.001).

*Intracranial bleeding:* There were more intracranial non-procedural bleeds with ticagrelor (n=27 bleeds in 26 patients, 0.3%) than with clopidogrel (n=14 bleeds, 0.2%), of which 11 bleeds with ticagrelor and 1 with clopidogrel were fatal. There was no difference in overall fatal bleeds.

*Bleeding findings in PEGASUS*

Overall outcome of bleeding events in the PEGASUS study are shown in Table 3.

**Table 3 – Analysis of overall bleeding events, Kaplan-Meier estimates at 36 months (PEGASUS)**

<table>
<thead>
<tr>
<th>Safety Endpoints</th>
<th>Ticagrelor 60 mg twice daily + ASA N=6958</th>
<th>ASA alone N=6996</th>
</tr>
</thead>
<tbody>
<tr>
<td>KM%</td>
<td>Hazard Ratio (95% CI)</td>
<td>KM%</td>
</tr>
<tr>
<td><strong>TIMI-defined bleeding categories</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TIMI Major</td>
<td>2.3</td>
<td>2.32 (1.68, 3.21)</td>
</tr>
<tr>
<td>Fatal</td>
<td>0.3</td>
<td>1.00 (0.44, 2.27)</td>
</tr>
<tr>
<td>ICH</td>
<td>0.6</td>
<td>1.33 (0.77, 2.31)</td>
</tr>
<tr>
<td>Other TIMI Major</td>
<td>1.6</td>
<td>3.61 (2.31, 5.65)</td>
</tr>
<tr>
<td>TIMI Major or Minor</td>
<td>3.4</td>
<td>2.54 (1.93, 3.35)</td>
</tr>
<tr>
<td>TIMI Major or Minor or Requiring medical attention</td>
<td>16.6</td>
<td>2.64 (2.35, 2.97)</td>
</tr>
</tbody>
</table>

**PLATO-defined bleeding categories**
PLATO Major | 3.5 | 2.57 | 1.4 | <0.0001
| Fatal/Life-threatening | 2.4 | 2.38 | 1.1 | <0.0001
| Other PLATO Major | 1.1 | 3.37 | 0.3 | <0.0001
| PLATO Major or Minor | 15.2 | 2.71 | 6.2 | <0.0001

**Bleeding category definitions:**

**TIMI Major:** Fatal bleeding, OR any intracranial bleeding, OR clinically overt signs of haemorrhage associated with a drop in haemoglobin (Hgb) of ≥50 g/L, or when Hgb is not available, a fall in haematocrit (Hct) of 15%.

**Fatal:** A bleeding event that directly led to death within 7 days.

**ICH:** Intracranial haemorrhage.

**Other TIMI Major:** Non-fatal non-ICH TIMI Major bleeding.

**TIMI Minor:** Clinically apparent with 30-50 g/L decrease in haemoglobin.

**TIMI Requiring medical attention:** Requiring intervention, OR leading to hospitalization, OR prompting evaluation.

**PLATO Major Fatal/life-threatening:** Fatal bleeding, OR any intracranial bleeding, OR intrapericardial with cardiac tamponade, OR with hypovolaemic shock or severe hypotension requiring pressors/inotropes or surgery OR clinically apparent with >50 g/L decrease in haemoglobin or ≥4 red cell units transfused.

**PLATO Major Other:** Significantly disabling, OR clinically apparent with 30-50 g/L decrease in haemoglobin, OR 2-3 red cell units transfused.

**PLATO Minor:** Requires medical intervention to stop or treat bleeding.

In PEGASUS, TIMI Major bleeding for ticagrelor 60 mg twice daily was higher than for ASA alone. No increased bleeding risk was seen for fatal bleeding and only a minor increase was observed in intracranial haemorrhages, as compared to ASA therapy alone. There were few fatal bleeding events in the study, 11 (0.3%) for ticagrelor 60 mg and 12 (0.3%) for ASA therapy alone. The observed increased risk of TIMI Major bleeding with ticagrelor 60 mg was primarily due to a higher frequency of Other TIMI Major bleedings driven by events in the gastrointestinal SOC.

Increased bleeding patterns similar to TIMI Major were seen for TIMI Major or Minor and PLATO Major and PLATO Major or Minor bleeding categories (see Table 3). Discontinuation of treatment due to bleeding was more common with ticagrelor 60 mg compared to ASA therapy alone (6.2% and 1.5%, respectively). The majority of these bleedings were of less severity (classified as TIMI Requiring medical attention), e.g. epistaxis, bruising and haematomas.

The bleeding profile of ticagrelor 60 mg was consistent across multiple pre-defined subgroups (e.g. by age, gender, weight, race, geographic region, concurrent conditions, concomitant therapy, and medical history) for TIMI Major, TIMI Major or Minor and PLATO Major bleeding events.

**Intracranial bleeding:** Spontaneous ICHs were reported in similar rates for ticagrelor 60 mg and ASA therapy alone (n=13, 0.2% in both treatment groups). Traumatic and procedural ICHs showed a minor increase with ticagrelor 60 mg treatment, (n=15, 0.2%) compared with ASA therapy alone (n=10, 0.1%). There were 6 fatal ICHs with ticagrelor 60 mg and 5 fatal ICHs with ASA therapy alone. The incidence of intracranial bleeding was low in both treatment groups given the significant comorbidity and CV risk factors of the population under study.

**Dyspnoea**
Dyspnoea, a sensation of breathlessness, is reported by patients treated with Brilique. In PLATO, dyspnoea adverse events (AEs) (dyspnoea, dyspnoea at rest, dyspnoea exertional, dyspnoea paroxysmal nocturnal and nocturnal dyspnoea), when combined, was reported by 13.8% of patients.
treated with ticagrelor and by 7.8% of patients treated with clopidogrel. In 2.2% of patients taking ticagrelor and by 0.6% taking clopidogrel investigators considered the dyspnoea causally related to treatment in the PLATO study and few were serious (0.14% ticagrelor; 0.02% clopidogrel), (see section 4.4). Most reported symptoms of dyspnoea were mild to moderate in intensity, and most were reported as a single episode early after starting treatment.

Compared with clopidogrel, patients with asthma/COPD treated with ticagrelor may have an increased risk of experiencing non-serious dyspnoea (3.29% ticagrelor versus 0.53% clopidogrel) and serious dyspnoea (0.38% ticagrelor versus 0.00% clopidogrel). In absolute terms, this risk was higher than in the overall PLATO population. Ticagrelor should be used with caution in patients with history of asthma and/or COPD (see section 4.4).

About 30% of episodes resolved within 7 days. PLATO included patients with baseline congestive heart failure, COPD, or asthma; these patients, and the elderly, were more likely to report dyspnoea. For Brilique, 0.9% of patients discontinued study drug because of dyspnoea compared with 0.1% taking clopidogrel. The higher incidence of dyspnoea with Brilique is not associated with new or worsening heart or lung disease (see section 4.4). Brilique does not affect tests of pulmonary function.

In PEGASUS, dyspnoea was reported in 14.2% of patients taking ticagrelor 60 mg twice daily and in 5.5% of patients taking ASA alone. As in PLATO, most reported dyspnoea was mild to moderate in intensity (see section 4.4). Patients who reported dyspnoea tended to be older and more frequently had dyspnoea, COPD or asthma at baseline.

**Investigations**

Uric acid elevations: In PLATO, serum uric acid increased to more than upper limit of normal in 22% of patients receiving ticagrelor compared to 13% of patients receiving clopidogrel. The corresponding numbers in PEGASUS were 9.1%, 8.8% and 5.5% for ticagrelor 90 mg, 60 mg and placebo, respectively. Mean serum uric acid increased approximately 15% with ticagrelor compared to approximately 7.5% with clopidogrel and after treatment was stopped, decreased to approximately 7% on ticagrelor but with no decrease observed for clopidogrel. In PEGASUS, a reversible increase in mean serum uric acid levels of 6.3% and 5.6% was found for ticagrelor 90 mg and 60 mg, respectively, compared to a 1.5% decrease in the placebo group. In PLATO, the frequency of gouty arthritis was 0.2% for ticagrelor vs. 0.1% for clopidogrel. The corresponding numbers for gout/gouty arthritis in PEGASUS were 1.6%, 1.5% and 1.1% for ticagrelor 90 mg, 60 mg and placebo, respectively.

**Reporting of suspected adverse reactions**

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via the national reporting system listed in Appendix V.

**4.9 Overdose**

Ticagrelor is well tolerated in single doses up to 900 mg. Gastrointestinal toxicity was dose-limiting in a single ascending dose study. Other clinically meaningful adverse reactions which may occur with overdose include dyspnoea and ventricular pauses (see section 4.8).

In the event of an overdose, the above potential adverse reactions could occur and ECG monitoring should be considered.
There is currently no known antidote to reverse the effects of ticagrelor, and ticagrelor is not expected to be dialysable (see section 4.4). Treatment of overdose should follow local standard medical practice. The expected effect of excessive ticagrelor dosing is prolonged duration of bleeding risk associated with platelet inhibition. Platelet transfusion is unlikely to be of clinical benefit in patients with bleeding (see section 4.4). If bleeding occurs other appropriate supportive measures should be taken.

5. PHARMACOLOGICAL PROPERTIES

5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Platelet aggregation inhibitors excluding heparin, ATC code: B01AC24

Mechanism of action
Brilique contains ticagrelor, a member of the chemical class cyclopentyltriazolopyrimidines (CPTP), which is an oral, direct acting, selective and reversibly binding P2Y\textsubscript{12} receptor antagonist that prevents ADP-mediated P2Y\textsubscript{12} dependent platelet activation and aggregation. Ticagrelor does not prevent ADP binding but when bound to the P2Y\textsubscript{12} receptor prevents ADP-induced signal transduction. Since platelets participate in the initiation and/or evolution of thrombotic complications of atherosclerotic disease, inhibition of platelet function has been shown to reduce the risk of CV events such as death, MI or stroke.

Ticagrelor also increases local endogenous adenosine levels by inhibiting the equilibrative nucleoside transporter -1 (ENT-1).

Ticagrelor has been documented to augment the following adenosine-induced effects in healthy subjects and in patients with ACS: vasodilation (measured by coronary blood flow increases in healthy volunteers and ACS patients; headache), inhibition of platelet function (in human whole blood \textit{in vitro}) and dyspnoea. However, a link between the observed increases in adenosine and clinical outcomes (e.g. morbidity-mortality) has not been clearly elucidated.

Pharmacodynamic effects

\textbf{Onset of action}

In patients with stable coronary artery disease (CAD) on ASA, ticagrelor demonstrates a rapid onset of pharmacological effect as demonstrated by a mean inhibition of platelet aggregation (IPA) for ticagrelor at 0.5 hours after 180 mg loading dose of about 41%, with the maximum IPA effect of 89% by 2-4 hours post dose, and maintained between 2-8 hours. 90% of patients had final extent IPA>70% by 2 hours post dose.

\textbf{Offset of action}

If a CABG procedure is planned, ticagrelor bleeding risk is increased compared to clopidogrel when discontinued within less than 96 hours prior to procedure.

\textbf{Switching data}

Switching from clopidogrel 75 mg to ticagrelor 90 mg twice daily results in an absolute IPA increase of 26.4% and switching from ticagrelor to clopidogrel results in an absolute IPA decrease of 24.5%. Patients can be switched from clopidogrel to ticagrelor without any interruption of antiplatelet effect (see section 4.2).
Clinical efficacy and safety
The clinical evidence for the efficacy and safety of ticagrelor is derived from two phase 3 trials:

- The PLATO [PLATelet Inhibition and Patient Outcomes] study, a comparison of ticagrelor to clopidogrel, both given in combination with ASA and other standard therapy.
- The PEGASUS TIMI-54 [PrEvention with Ticagrelor of SecondAry Thrombotic Events in High-RiSk AcUte Coronary Syndrome Patients] study, a comparison of ticagrelor combined with ASA to ASA therapy alone.

PLATO study (Acute Coronary Syndromes)

The PLATO study included 18,624 patients who presented within 24 hours of onset of symptoms of unstable angina (UA), non ST elevation myocardial infarction (NSTEMI) or ST elevation myocardial infarction (STEMI), and were initially managed medically, or with percutaneous coronary intervention (PCI), or with CABG.

Clinical efficacy
On a background of daily ASA, ticagrelor 90 mg twice daily showed superiority to 75 mg daily clopidogrel in preventing the composite endpoint of CV death, MI, or stroke, with the difference driven by CV death and MI. Patients received a 300 mg loading dose of clopidogrel (600 mg possible if having PCI) or 180 mg of ticagrelor.

The result appeared early (absolute risk reduction [ARR] 0.6% and relative risk reduction [RRR] of 12% at 30 days), with a constant treatment effect over the entire 12-month period, yielding ARR 1.9% per year with RRR of 16%. This suggests it is appropriate to treat patients with ticagrelor 90 mg twice daily for 12 months (see section 4.2). Treating 54 ACS patients with ticagrelor instead of clopidogrel will prevent 1 atherothrombotic event; treating 91 will prevent 1 CV death (see Figure 1 and Table 4).

The treatment effect of ticagrelor over clopidogrel appears consistent across many subgroups, including weight; sex; medical history of diabetes mellitus, transient ischaemic attack or non-haemorrhagic stroke, or revascularisation; concomitant therapies including heparins, GpIIb/IIIa inhibitors and proton pump inhibitors (see section 4.5); final index event diagnosis (STEMI, NSTEMI, or UA); and treatment pathway intended at randomisation (invasive or medical).

A weakly significant treatment interaction was observed with region whereby the hazard ratio (HR) for the primary endpoint favours ticagrelor in the rest of world but favours clopidogrel in North America, which represented approximately 10% of the overall population studied (interaction p-value=0.045). Exploratory analyses suggest a possible association with ASA dose such that reduced efficacy was observed with ticagrelor with increasing ASA doses. Chronic daily ASA doses to accompany Brilique should be 75-150 mg (see sections 4.2 and 4.4).

Figure 1 shows the estimate of the risk to the first occurrence of any event in the composite efficacy endpoint.
Ticagrelor reduced the occurrence of the primary composite endpoint compared to clopidogrel in both the UA/NSTEMI and STEMI population (Table 4). Thus, Brilique 90 mg twice daily together with low-dose ASA can be used in patients with ACS (unstable angina, non ST elevation Myocardial Infarction [NSTEMI] or ST elevation Myocardial Infarction [STEMI]); including patients managed medically, and those who are managed with percutaneous coronary intervention (PCI) or coronary artery by-pass grafting (CABG).

**Table 4 - Analysis of primary and secondary efficacy endpoints (PLATO)**

<table>
<thead>
<tr>
<th>Endpoint</th>
<th>Ticagrelor 90 mg twice daily (% patients with event) N=9333</th>
<th>Clopidogrel 75 mg once daily (% patients with event) N=9291</th>
<th>ARR (%) (%/yr)</th>
<th>RRR (%) (95% CI)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CV death, MI (excl. silent MI) or stroke</td>
<td>9.3</td>
<td>10.9</td>
<td>1.9</td>
<td>16 (8, 23)</td>
<td>0.0003</td>
</tr>
<tr>
<td>Invasive intent</td>
<td>8.5</td>
<td>10.0</td>
<td>1.7</td>
<td>16 (6, 25)</td>
<td>0.0025</td>
</tr>
<tr>
<td>Medical intent</td>
<td>11.3</td>
<td>13.2</td>
<td>2.3</td>
<td>15 (0.3, 27)</td>
<td>0.0444d</td>
</tr>
<tr>
<td>CV death</td>
<td>3.8</td>
<td>4.8</td>
<td>1.1</td>
<td>21 (9, 31)</td>
<td>0.0013</td>
</tr>
<tr>
<td>MI (excl. silent MI)b</td>
<td>5.4</td>
<td>6.4</td>
<td>1.1</td>
<td>16 (5, 25)</td>
<td>0.0045</td>
</tr>
<tr>
<td>Stroke</td>
<td>1.3</td>
<td>1.1</td>
<td>-0.2</td>
<td>-17 (-52, 9)</td>
<td>0.2249</td>
</tr>
<tr>
<td>All-cause mortality, MI (excl. silent MI), or stroke</td>
<td>9.7</td>
<td>11.5</td>
<td>2.1</td>
<td>16 (8, 23)</td>
<td>0.0001</td>
</tr>
<tr>
<td>CV death, total MI, stroke, SRI, RI, TIA, or other ATE</td>
<td>13.8</td>
<td>15.7</td>
<td>2.1</td>
<td>12 (5, 19)</td>
<td>0.0006</td>
</tr>
<tr>
<td>---------------------------------------------------</td>
<td>------</td>
<td>------</td>
<td>-----</td>
<td>-----------</td>
<td>-------</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>4.3</td>
<td>5.4</td>
<td>1.4</td>
<td>22 (11, 31)</td>
<td>0.0003^d</td>
</tr>
<tr>
<td>Definite stent thrombosis</td>
<td>1.2</td>
<td>1.7</td>
<td>0.6</td>
<td>32 (8, 49)</td>
<td>0.0123d</td>
</tr>
</tbody>
</table>

^ARR = absolute risk reduction; RRR = relative risk reduction = (1-Hazard ratio) x 100%. A negative RRR indicates a relative risk increase.

^Excluding silent MI.

^SRI = serious recurrent ischaemia; RI = recurrent ischaemia; TIA = transient ischaemic attack; ATE = arterial thrombotic event. Total MI includes silent MI, with date of event set to date when discovered.

^Nominal significance value; all others are formally statistically significant by pre-defined hierarchical testing.

**PLATO genetic substudy**

CYP2C19 and ABCB1 genotyping of 10,285 patients in PLATO provided associations of genotype groups with PLATO outcomes. The superiority of ticagrelor over clopidogrel in reducing major CV events was not significantly affected by patient CYP2C19 or ABCB1 genotype. Similar to the overall PLATO study, total PLATO Major bleeding did not differ between ticagrelor and clopidogrel, regardless of CYP2C19 or ABCB1 genotype. Non-CABG PLATO Major bleeding was increased with ticagrelor compared clopidogrel in patients with one or more CYP2C19 loss of function alleles, but similar to clopidogrel in patients with no loss of function allele.

**Combined efficacy and safety composite**

A combined efficacy and safety composite (CV death, MI, stroke, or PLATO-defined ‘Total Major’ bleeding) indicates that the benefit in efficacy of ticagrelor compared to clopidogrel is not offset by the major bleeding events (ARR 1.4%, RRR 8%, HR 0.92; p=0.0257) over 12 months after ACS.

**Clinical safety**

**Holter substudy**

To study the occurrence of ventricular pauses and other arrhythmic episodes during PLATO, investigators performed Holter monitoring in a subset of nearly 3000 patients, of whom approximately 2000 had recordings both in the acute phase of their ACS and after one month. The primary variable of interest was the occurrence of ventricular pauses ≥3 seconds. More patients had ventricular pauses with ticagrelor (6.0%) than with clopidogrel (3.5%) in the acute phase; and 2.2% and 1.6% respectively after 1 month (see section 4.4). The increase in ventricular pauses in the acute phase of ACS was more pronounced in ticagrelor patients with history of CHF (9.2% versus 5.4% in patients without CHF history; for clopidogrel patients, 4.0% in those with versus 3.6% in those without CHF history). This imbalance did not occur at one month: 2.0% versus 2.1% for ticagrelor patients with and without CHF history respectively; and 3.8% versus 1.4% with clopidogrel. There were no adverse clinical consequences associated with this imbalance (including pacemaker insertions) in this population of patients.

**PEGASUS study (History of Myocardial Infarction)**

The PEGASUS TIMI-54 study was a 21,162 patient, event-driven, randomised, double-blind, placebo-controlled, parallel group, international multicentre study to assess the prevention of atherothrombotic events with ticagrelor given at 2 doses (either 90 mg twice daily or 60 mg twice daily) combined with low dose ASA (75-150 mg), compared to ASA therapy alone in patients with history of MI and additional risk factors for atherothrombosis.

Patients were eligible to participate if they were aged 50 years or over, with a history of MI (1 to 3 years prior to randomisation), and had at least one of the following risk factors for
atherothrombosis: age ≥65 years, diabetes mellitus requiring medication, a second prior MI, evidence of multivessel CAD, or chronic non-end-stage renal dysfunction.

Patients were ineligible if there was planned use of a P2Y12 receptor antagonist, dipyridamole, cilostazol, or anticoagulant therapy during the study period; if they had a bleeding disorder or a history of an ischaemic stroke or intracranial bleeding, a central nervous system tumour, or an intracranial vascular abnormality; if they had had gastrointestinal bleeding within the previous 6 months or major surgery within the previous 30 days.

Clinical efficacy

Figure 2 - Analysis of primary clinical composite endpoint of CV death, MI and stroke (PEGASUS)

![Cumulative % over Days from Randomisation graph]

Table 5 - Analysis of primary and secondary efficacy endpoints (PEGASUS)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Ticagrelor 60 mg twice daily +ASA</th>
<th>ASA alone</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Patients with events KM %</td>
<td>HR (95% CI)</td>
</tr>
<tr>
<td>Primary endpoint</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Characteristic</td>
<td>Patients with events</td>
<td>KM %</td>
</tr>
<tr>
<td>-------------------------------</td>
<td>----------------------</td>
<td>-------</td>
</tr>
<tr>
<td>Composite of CV Death/MI/Stroke</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ticagrelor 60 mg twice daily + ASA</td>
<td>487 (6.9%)</td>
<td>7.8%</td>
</tr>
<tr>
<td>ASA alone</td>
<td></td>
<td></td>
</tr>
<tr>
<td>CV death</td>
<td>174 (2.5%)</td>
<td>2.9%</td>
</tr>
<tr>
<td>MI</td>
<td>285 (4.0%)</td>
<td>4.5%</td>
</tr>
<tr>
<td>Stroke</td>
<td>91 (1.3%)</td>
<td>1.5%</td>
</tr>
</tbody>
</table>

Secondary endpoint

<table>
<thead>
<tr>
<th></th>
<th>Patients with events</th>
<th>KM %</th>
<th>HR (95% CI)</th>
<th>Patients with events</th>
<th>KM %</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CV death</td>
<td>174 (2.5%)</td>
<td>2.9%</td>
<td>0.83 (0.68, 1.01)</td>
<td>210 (3.0%)</td>
<td>3.4%</td>
<td>-</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>289 (4.1%)</td>
<td>4.7%</td>
<td>0.89 (0.76, 1.04)</td>
<td>326 (4.6%)</td>
<td>5.2%</td>
<td>-</td>
</tr>
</tbody>
</table>

Hazard ratio and p-values are calculated separately for ticagrelor vs. ASA therapy alone from Cox proportional hazards model with treatment group as the only explanatory variable. KM percentage calculated at 36 months. Note: the number of first events for the components CV death, MI and stroke are the actual number of first events for each component and do not add up to the number of events in the composite endpoint (s) Indicates statistical significance. CI = Confidence interval; CV = Cardiovascular; HR = Hazard ratio; KM = Kaplan-Meier; MI = Myocardial infarction; N = Number of patients.

Both 60 mg twice daily and 90 mg twice daily regimens of ticagrelor in combination with ASA were superior to ASA alone in the prevention of atherothrombotic events (composite endpoint: CV death, MI and stroke), with a consistent treatment effect over the entire study period, yielding a 16% RRR and 1.27% ARR for ticagrelor 60 mg and a 15% RRR and 1.19% ARR for ticagrelor 90 mg.

Although the efficacy profiles of 90 mg and 60 mg were similar, there is evidence that the lower dose has a better tolerability and safety profile in relation to risk of the bleeding and dyspnoea. Therefore, only Brilique 60 mg twice daily co-administered with ASA is recommended for the prevention atherothrombotic events (CV death, MI and stroke) in patients with a history of MI and a high risk of developing an atherothrombotic event.

Relative to ASA alone, ticagrelor 60 mg twice daily significantly reduced the primary composite endpoint of CV death, MI and stroke. Each of the components contributed to the reduction in the primary composite endpoint (CV death 17% RRR, MI 16% RRR, and stroke 25% RRR).

The RRR for the composite endpoint from 1 to 360 days (17% RRR) and from 361 days and onwards (16% RRR) was similar. There are limited data on the efficacy and safety of ticagrelor beyond 3 years of extended treatment.

There was no evidence of benefit (no reduction in the primary composite endpoint of CV death, MI and stroke, but an increase in major bleeding) when ticagrelor 60 mg twice daily was introduced in clinically stable patients >2 years from the MI, or more than one year after stopping previous ADP receptor inhibitor treatment (see also section 4.2).
Clinical safety
The rate of discontinuations with ticagrelor 60 mg due to bleeding and dyspnoea was higher in patients >75 years (42%) than in younger patients (range: 23-31%), with a difference versus placebo higher than 10% (42% vs. 29%) in patients >75 years.

Paediatric population
The European Medicines Agency has waived the obligation to submit the results of studies with Brilique in all subsets of the paediatric population in the granted indication (see section 4.2 for information on paediatric use).

5.2 Pharmacokinetic properties
Ticagrelor demonstrates linear pharmacokinetics and exposure to ticagrelor and the active metabolite (AR-C124910XX) are approximately dose proportional up to 1260 mg.

Absorption
Absorption of ticagrelor is rapid with a median \( t_{\text{max}} \) of approximately 1.5 hours. The formation of the major circulating metabolite AR-C124910XX (also active) from ticagrelor is rapid with a median \( t_{\text{max}} \) of approximately 2.5 hours. Following an oral ticagrelor 90 mg single dose under fasted conditions in healthy subjects, \( C_{\text{max}} \) is 529 ng/ml and AUC is 3451 ng*h/ml. The metabolite parent ratios are 0.28 for \( C_{\text{max}} \) and 0.42 for AUC. The pharmacokinetics of ticagrelor and AR-C124910XX in patients with a history of MI were generally similar to that in the ACS population. Based on a population pharmacokinetic analysis of the PEGASUS study the median ticagrelor \( C_{\text{max}} \) was 391 ng/ml and AUC was 3801 ng*h/ml at steady state for ticagrelor 60 mg. For ticagrelor 90 mg \( C_{\text{max}} \) was 627 ng/ml and AUC was 6255 ng*h/ml at steady state.

The mean absolute bioavailability of ticagrelor was estimated to be 36%. Ingestion of a high-fat meal resulted in a 21% increase in ticagrelor AUC and 22% decrease in the active metabolite \( C_{\text{max}} \) but had no effect on ticagrelor \( C_{\text{max}} \) or the AUC of the active metabolite. These small changes are considered of minimal clinical significance; therefore, ticagrelor can be given with or without food. Ticagrelor as well as the active metabolite are P-gp substrates.

Ticagrelor as crushed tablets mixed in water, given orally or administered through a nasogastric tube into the stomach, has a comparable bioavailability to whole tablets with regards to AUC and \( C_{\text{max}} \) for ticagrelor and the active metabolite. Initial exposure (0.5 and 1 hour post-dose) from crushed ticagrelor tablets mixed in water was higher compared to whole tablets, with a generally identical concentration profile thereafter (2 to 48 hours).

Distribution
The steady state volume of distribution of ticagrelor is 87.5 l. Ticagrelor and the active metabolite is extensively bound to human plasma protein (>99.0%).

Biotransformation
CYP3A4 is the major enzyme responsible for ticagrelor metabolism and the formation of the active metabolite and their interactions with other CYP3A substrates ranges from activation through to inhibition.

The major metabolite of ticagrelor is AR-C124910XX, which is also active as assessed by in vitro binding to the platelet P2Y_12 ADP-receptor. The systemic exposure to the active metabolite is approximately 30-40% of that obtained for ticagrelor.
Elimination
The primary route of ticagrelor elimination is via hepatic metabolism. When radiolabelled ticagrelor is administered, the mean recovery of radioactivity is approximately 84% (57.8% in faeces, 26.5% in urine). Recoveries of ticagrelor and the active metabolite in urine were both less than 1% of the dose. The primary route of elimination for the active metabolite is most likely via biliary secretion. The mean $t_{1/2}$ was approximately 7 hours for ticagrelor and 8.5 hours for the active metabolite.

Special populations

Elderly
Higher exposures to ticagrelor (approximately 25% for both $C_{\text{max}}$ and AUC) and the active metabolite were observed in elderly (≥75 years) ACS patients compared to younger patients by the population pharmacokinetic analysis. These differences are not considered clinically significant (see section 4.2).

Paediatric population
Ticagrelor has not been evaluated in a paediatric population (see sections 4.2 and 5.1).

Gender
Higher exposures to ticagrelor and the active metabolite were observed in women compared to men. These differences are not considered clinically significant.

Renal impairment
Exposure to ticagrelor was approximately 20% lower and exposure to the active metabolite was approximately 17% higher in patients with severe renal impairment (creatinine clearance <30 ml/min) compared to subjects with normal renal function (see section 4.2).

Hepatic impairment
$C_{\text{max}}$ and AUC for ticagrelor were 12% and 23% higher in patients with mild hepatic impairment compared to matched healthy subjects, respectively; however, the IPA effect of ticagrelor was similar between the two groups. No dose adjustment is needed for patients with mild hepatic impairment. Ticagrelor has not been studied in patients with severe hepatic impairment and there is no pharmacokinetic information in patients with moderate hepatic impairment. In patients that had moderate or severe elevation in one or more liver function tests at baseline, ticagrelor plasma concentrations were on average similar or slightly higher as compared to those without baseline elevations. No dose adjustment is recommended in patients with moderate hepatic impairment (see sections 4.2 and 4.4).

Ethnicity
Patients of Asian descent have a 39% higher mean bioavailability compared to Caucasian patients. Patients self-identified as black had an 18% lower bioavailability of ticagrelor compared to Caucasian patients, in clinical pharmacology studies, the exposure ($C_{\text{max}}$ and AUC) to ticagrelor in Japanese subjects was approximately 40% (20% after adjusting for body weight) higher compared to that in Caucasians. The exposure in patients self-identified as Hispanic or Latino was similar to that in Caucasians.

5.3 Preclinical safety data

Preclinical data for ticagrelor and its major metabolite have not demonstrated unacceptable risk for adverse effects for humans based on conventional studies of safety pharmacology, single and repeated dose toxicity and genotoxic potential.
Gastrointestinal irritation was observed in several animal species at clinical relevant exposure levels (see section 4.8).

In female rats, ticagrelor at high dose showed an increased incidence of uterine tumours (adenocarcinomas) and an increased incidence of hepatic adenomas. The mechanism for uterine tumours is likely hormonal imbalance which can lead to tumours in rats. The mechanism for the hepatic adenomas is likely due to a rodent-specific enzyme induction in the liver. Thus, the carcinogenicity findings are considered unlikely to be relevant for humans.

In rats minor developmental anomalies were seen at a maternal toxic dose (safety margin of 5.1). In rabbits a slight delay in hepatic maturity and skeletal development was seen in foetuses from dams at high dose without showing maternal toxicity (safety margin of 4.5).

Studies in rats and rabbits have shown reproductive toxicity, with slightly reduced maternal body weight gain and reduced neonatal viability and birth weight, with delayed growth. Ticagrelor produced irregular cycles (mostly extended cycles) in female rats, but did not affect overall fertility in male and female rats. Pharmacokinetic studies performed with radio-labelled ticagrelor have shown that the parent compound and its metabolites are excreted in the milk of rats (see section 4.6).

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Tablet core
Mannitol (E421)
Calcium hydrogen phosphate dihydrate
Magnesium stearate (E470b)
Sodium starch glycolate type A
Hydroxypropylcellulose (E463)

Tablet coating
Titanium dioxide (E171)
Iron oxide black (E172)
Iron oxide red (E172)
Macrogol 400
Hypromellose (E464)

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

3 years

6.4 Special precautions for storage

This medicinal product does not require any special storage conditions.

6.5 Nature and contents of container
- PVC-PVDC/Al transparent blister (with sun/moon symbols) of 10 tablets; cartons of 60 tablets (6 blisters) and 180 tablets (18 blisters).
- PVC-PVDC/Al transparent calendar blister (with sun/moon symbols) of 14 tablets; cartons of 14 tablets (1 blister), 56 tablets (4 blisters), and 168 tablets (12 blisters).

Not all pack sizes may be marketed.

6.6 Special precautions for disposal

No special requirements. Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

7. MARKETING AUTHORISATION HOLDER

AstraZeneca AB
SE-151 85 Södertälje
Sweden

8. MARKETING AUTHORISATION NUMBER(S)

EU/1/10/655/007-011

9. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 03 December 2010
Date of latest renewal: 17 July 2015

10. DATE OF REVISION OF THE TEXT

1. NAME OF THE MEDICINAL PRODUCT

Brilique 90 mg film-coated tablets

2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each film-coated tablet contains 90 mg ticagrelor.

For the full list of excipients, see section 6.1.

3. PHARMACEUTICAL FORM

Film-coated tablet (tablet).

Round, biconvex, yellow tablets marked with ‘90’ above ‘T’ on one side and plain on the other.

4. CLINICAL PARTICULARS

4.1 Therapeutic indications

Brilique, co-administered with acetylsalicylic acid (ASA), is indicated for the prevention of atherothrombotic events in adult patients with
- acute coronary syndromes (ACS) or
- a history of myocardial infarction (MI) and a high risk of developing an atherothrombotic event (see sections 4.2 and 5.1).

4.2 Posology and method of administration

Posology

Patients taking Brilique should also take a daily low maintenance dose of ASA 75-150 mg, unless specifically contraindicated.

Acute coronary syndromes
Brilique treatment should be initiated with a single 180 mg loading dose (two tablets of 90 mg) and then continued at 90 mg twice daily. Treatment with Brilique 90 mg twice daily is recommended for 12 months in ACS patients unless discontinuation is clinically indicated (see section 5.1).

History of myocardial infarction
Brilique 60 mg twice daily is the recommended dose when an extended treatment is required for patients with a history of MI of at least one year and a high risk of an atherothrombotic event (see section 5.1). Treatment may be started without interruption as continuation therapy after the initial one-year treatment with Brilique 90 mg or other adenosine diphosphate (ADP) receptor inhibitor therapy in ACS patients with a high risk of an atherothrombotic event. Treatment can also be initiated up to 2 years from the MI, or within one year after stopping previous ADP receptor inhibitor treatment. There are limited data on the efficacy and safety of Brilique beyond 3 years of extended treatment.
If a switch is needed, the first dose of Brilique should be administered 24 hours following the last dose of the other antiplatelet medication.

**Missed dose**

Lapses in therapy should also be avoided. A patient who misses a dose of Brilique should take only one tablet (their next dose) at its scheduled time.

**Special populations**

**Elderly**

No dose adjustment is required in elderly (see section 5.2).

**Renal impairment**

No dose adjustment is necessary for patients with renal impairment (see section 5.2). No information is available concerning treatment of patients on renal dialysis and therefore ticagrelor is not recommended in these patients.

**Hepatic impairment**

Ticagrelor has not been studied in patients with severe hepatic impairment and its use in these patients is therefore contraindicated (see section 4.3). Only limited information is available in patients with moderate hepatic impairment. Dose adjustment is not recommended, but ticagrelor should be used with caution (see sections 4.4 and 5.2). No dose adjustment is necessary for patients with mild hepatic impairment (see section 5.2).

**Paediatric population**

The safety and efficacy of ticagrelor in children below the age of 18 years have not been established. No data are available.

**Method of administration**

For oral use.

Brilique can be administered with or without food.

For patients who are unable to swallow the tablet(s) whole, the tablets can be crushed to a fine powder and mixed in half a glass of water and drunk immediately. The glass should be rinsed with a further half glass of water and the contents drunk. The mixture can also be administered via a nasogastric tube (CH8 or greater). It is important to flush the nasogastric tube through with water after administration of the mixture.

### 4.3 Contraindications

- Hypersensitivity to the active substance or to any of the excipients listed in section 6.1 (see section 4.8).
- Active pathological bleeding.
- History of intracranial haemorrhage (see section 4.8).
- Severe hepatic impairment (see sections 4.2, 4.4 and 5.2).
- Co-administration of ticagrelor with strong CYP3A4 inhibitors (e.g. ketoconazole, clarithromycin, nefazodone, ritonavir, and atazanavir) is contraindicated, as co-administration may lead to a substantial increase in exposure to ticagrelor (see section 4.5).
4.4 Special warnings and precautions for use

Bleeding risk
The use of ticagrelor in patients at known increased risk for bleeding should be balanced against
the benefit in terms of prevention of atherothrombotic events (see sections 4.8 and 5.1). If
clinically indicated, ticagrelor should be used with caution in the following patient groups:

- Patients with a propensity to bleed (e.g. due to recent trauma, recent surgery, coagulation
  disorders, active or recent gastrointestinal bleeding). The use of ticagrelor is contraindicated
  in patients with active pathological bleeding, in those with a history of intracranial
  haemorrhage, and in patients with severe hepatic impairment (see section 4.3).
- Patients with concomitant administration of medicinal products that may increase the risk of
  bleeding (e.g. non-steroidal anti-inflammatory drugs (NSAIDs), oral anticoagulants and/or
  fibrinolytics) within 24 hours of ticagrelor dosing.

Platelet transfusion did not reverse the antiplatelet effect of ticagrelor in healthy volunteers and is
unlikely to be of clinical benefit in patients with bleeding. Since co-administration of ticagrelor
with desmopressin did not decrease template-bleeding time, desmopressin is unlikely to be
effective in managing clinical bleeding events (see section 4.5).

Antifibrinolytic therapy (aminocaproic acid or tranexamic acid) and/or recombinant factor VIIa
therapy may increase haemostasis. Ticagrelor may be resumed after the cause of bleeding has been
identified and controlled.

Surgery
Patients should be advised to inform physicians and dentists that they are taking ticagrelor before
any surgery is scheduled and before any new medicinal product is taken.

In PLATO patients undergoing coronary artery bypass grafting (CABG), ticagrelor had more
bleeding than clopidogrel when stopped within 1 day prior to surgery but a similar rate of major
bleeds compared to clopidogrel after stopping therapy 2 or more days before surgery (see
section 4.8). If a patient is to undergo elective surgery and antiplatelet effect is not desired,
ticagrelor should be discontinued 7 days prior to surgery (see section 5.1).

Patients with prior ischaemic stroke
ACS patients with prior ischaemic stroke can be treated with Brilique for up to 12 months (PLATO
study).

In PEGASUS, patients with history of MI with prior ischaemic stroke were not included.
Therefore, in the absence of data, treatment beyond one year is not recommended in these patients.

Hepatic impairment
Use of ticagrelor is contraindicated in patients with severe hepatic impairment (see sections 4.2
and 4.3). There is limited experience with ticagrelor in patients with moderate hepatic impairment,
therefore, caution is advised in these patients (see sections 4.2 and 5.2).

Patients at risk for bradycardic events
Due to observations of mostly asymptomatic ventricular pauses in an earlier clinical study, patients
with an increased risk of bradycardic events (e.g. patients without a pacemaker who have sick
sinus syndrome, 2nd or 3rd degree AV block or bradycardic-related syncope) were excluded from
the main studies evaluating the safety and efficacy of ticagrelor. Therefore, due to the limited
clinical experience, ticagrelor should be used with caution in these patients (see section 5.1).
In addition, caution should be exercised when administering ticagrelor concomitantly with medicinal products known to induce bradycardia. However, no evidence of clinically significant adverse reactions was observed in the PLATO trial after concomitant administration with one or more medicinal products known to induce bradycardia (e.g. 96% beta blockers, 33% calcium channel blockers diltiazem and verapamil, and 4% digoxin) (see section 4.5).

During the Holter substudy in PLATO, more patients had ventricular pauses ≥3 seconds with ticagrelor than with clopidogrel during the acute phase of their ACS. The increase in Holter-detected ventricular pauses with ticagrelor was higher in patients with chronic heart failure (CHF) than in the overall study population during the acute phase of ACS, but not at one month with ticagrelor or compared to clopidogrel. There were no adverse clinical consequences associated with this imbalance (including syncope or pacemaker insertion) in this patient population (see section 5.1).

**Dyspnoea**

Dyspnoea was reported in patients treated with ticagrelor. Dyspnoea is usually mild to moderate in intensity and often resolves without need for treatment discontinuation. Patients with asthma/chronic obstructive pulmonary disease (COPD) may have an increased absolute risk of experiencing dyspnoea with ticagrelor. Ticagrelor should be used with caution in patients with history of asthma and/or COPD. The mechanism has not been elucidated. If a patient reports new, prolonged or worsened dyspnoea this should be investigated fully and if not tolerated, treatment with ticagrelor should be stopped. For further details see section 4.8.

**Creatinine elevations**

Creatinine levels may increase during treatment with ticagrelor. The mechanism has not been elucidated. Renal function should be checked according to routine medical practice. In patients with ACS, it is recommended that renal function is also checked one month after initiating the treatment with ticagrelor, paying special attention to patients ≥75 years, patients with moderate/severe renal impairment and those receiving concomitant treatment with an angiotensin receptor blocker (ARB).

**Uric acid increase**

Hyperuricaemia may occur during treatment with ticagrelor (see section 4.8). Caution is advised in patients with history of hyperuricaemia or gouty arthritis. As a precautionary measure, the use of ticagrelor in patients with uric acid nephropathy is discouraged.

**Other**

Based on a relationship observed in PLATO between maintenance ASA dose and relative efficacy of ticagrelor compared to clopidogrel, co-administration of ticagrelor and high maintenance dose ASA (>300 mg) is not recommended (see section 5.1).

**Premature discontinuation**

Premature discontinuation with any antiplatelet therapy, including Brilique, could result in an increased risk of cardiovascular (CV) death or MI due to the patient’s underlying disease. Therefore, premature discontinuation of treatment should be avoided.

**4.5 Interaction with other medicinal products and other forms of interaction**

Ticagrelor is primarily a CYP3A4 substrate and a mild inhibitor of CYP3A4. Ticagrelor is also a P-glycoprotein (P-gp) substrate and a weak P-gp inhibitor and may increase the exposure of P-gp substrates.
Effects of other medicinal products on ticagrelor

Medicinal products metabolised by CYP3A4

CYP3A4 inhibitors

- Strong CYP3A4 inhibitors – Co-administration of ketoconazole with ticagrelor increased the ticagrelor C\text{max} and AUC equal to 2.4-fold and 7.3-fold, respectively. The C\text{max} and AUC of the active metabolite were reduced by 89% and 56%, respectively. Other strong inhibitors of CYP3A4 (clarithromycin, nefazodone, ritonavir, and atazanavir) would be expected to have similar effects and therefore concomitant use of strong CYP3A4 inhibitors with ticagrelor is contraindicated (see section 4.3).

- Moderate CYP3A4 inhibitors – Co-administration of diltiazem with ticagrelor increased the ticagrelor C\text{max} by 69% and AUC to 2.7-fold and decreased the active metabolite C\text{max} by 38% and AUC was unchanged. There was no effect of ticagrelor on diltiazem plasma levels. Other moderate CYP3A4 inhibitors (e.g. amprenavir, aprepitant, erythromycin and fluconazole) would be expected to have a similar effect and can as well be co-administered with ticagrelor.

CYP3A inducers

Co-administration of rifampicin with ticagrelor decreased ticagrelor C\text{max} and AUC by 73% and 86%, respectively. The C\text{max} of the active metabolite was unchanged and the AUC was decreased by 46%, respectively. Other CYP3A inducers (e.g. phenytoin, carbamazepine and phenobarbital) would be expected to decrease the exposure to ticagrelor as well. Co-administration of ticagrelor with potent CYP3A inducers may decrease exposure and efficacy of ticagrelor, therefore, their concomitant use with ticagrelor is discouraged.

Cyclosporine (P-gp and CYP3A inhibitor)

Co-administration of cyclosporine (600 mg) with ticagrelor increased ticagrelor C\text{max} and AUC equal to 2.3-fold and 2.8-fold, respectively. The AUC of the active metabolite was increased by 32% and C\text{max} was decreased by 15% in the presence of cyclosporine.

No data are available on concomitant use of ticagrelor with other active substances that also are potent P-gp inhibitors and moderate CYP3A4 inhibitors (e.g. verapamil, quinidine) that also may increase ticagrelor exposure. If the association cannot be avoided, their concomitant use should be made with caution.

Others

Clinical pharmacology interaction studies showed that co-administration of ticagrelor with heparin, enoxaparin and ASA or desmopressin did not have any effect on the pharmacokinetics of ticagrelor or the active metabolite or on ADP-induced platelet aggregation compared with ticagrelor alone. If clinically indicated, medicinal products that alter haemostasis should be used with caution in combination with ticagrelor.

A 2-fold increase of ticagrelor exposure was observed after daily consumption of large quantities of grapefruit juice (3x200 ml). This magnitude of increased exposure is not expected to be clinically relevant to most patients.

Effects of ticagrelor on other medicinal products

Medicinal products metabolised by CYP3A4

- Simvastatin – Co-administration of ticagrelor with simvastatin increased simvastatin C\text{max} by 81% and AUC by 56% and increased simvastatin acid C\text{max} by 64% and AUC by 52% with
some individual increases equal to 2 to 3-fold. Co-administration of ticagrelor with doses of simvastatin exceeding 40 mg daily could cause adverse effects of simvastatin and should be weighed against potential benefits. There was no effect of simvastatin on ticagrelor plasma levels. Ticagrelor may have similar effect on lovastatin. The concomitant use of ticagrelor with doses of simvastatin or lovastatin greater than 40 mg is not recommended.

- **Atorvastatin** – Co-administration of atorvastatin and ticagrelor increased atorvastatin acid $C_{\text{max}}$ by 23% and AUC by 36%. Similar increases in AUC and $C_{\text{max}}$ were observed for all atorvastatin acid metabolites. These increases are not considered clinically significant.

- A similar effect on other statins metabolised by CYP3A4 cannot be excluded. Patients in PLATO receiving ticagrelor took a variety of statins, with no concern of an association with statin safety among the 93% of the PLATO cohort taking these medicinal products.

Ticagrelor is a mild CYP3A4 inhibitor. Co-administration of ticagrelor and CYP3A4 substrates with narrow therapeutic indices (i.e. cisapride or ergot alkaloids) is not recommended, as ticagrelor may increase the exposure to these medicinal products.

**P-gp substrates (including digoxin, cyclosporine)**

Concomitant administration of ticagrelor increased the digoxin $C_{\text{max}}$ by 75% and AUC by 28%. The mean trough digoxin levels were increased about 30% with ticagrelor co-administration with some individual maximum increases to 2-fold. In the presence of digoxin, the $C_{\text{max}}$ and AUC of ticagrelor and its active metabolite were not affected. Therefore, appropriate clinical and/or laboratory monitoring is recommended when giving narrow therapeutic index P-gp dependent medicinal products like digoxin concomitantly with ticagrelor.

There was no effect of ticagrelor on cyclosporine blood levels. Effect of ticagrelor on other P-gp substrates has not been studied.

**Medicinal products metabolised by CYP2C9**

Co-administration of ticagrelor with tolbutamide resulted in no change in the plasma levels of either medicinal product, which suggests that ticagrelor is not a CYP2C9 inhibitor and unlikely to alter the CYP2C9 mediated metabolism of medicinal products like warfarin and tolbutamide.

**Oral contraceptives**

Co-administration of ticagrelor and levonorgestrel and ethinyl estradiol increased ethinyl estradiol exposure approximately 20% but did not alter the pharmacokinetics of levonorgestrel. No clinically relevant effect on oral contraceptive efficacy is expected when levonorgestrel and ethinyl estradiol are co-administered with ticagrelor.

**Medicinal products known to induce bradycardia**

Due to observations of mostly asymptomatic ventricular pauses and bradycardia, caution should be exercised when administering ticagrelor concomitantly with medicinal products known to induce bradycardia (see section 4.4). However, no evidence of clinically significant adverse reactions was observed in the PLATO trial after concomitant administration with one or more medicinal products known to induce bradycardia (e.g. 96% beta blockers, 33% calcium channel blockers diltiazem and verapamil, and 4% digoxin).

**Other concomitant therapy**

In clinical studies, ticagrelor was commonly administered with ASA, proton pump inhibitors, statins, beta-blockers, angiotensin converting enzyme (ACE) inhibitors and angiotensin receptor blockers as needed for concomitant conditions for long-term and also heparin, low molecular weight heparin and intravenous GpIIb/IIIa inhibitors for short durations (see section 5.1). No evidence of clinically significant adverse interactions with these medicinal products was observed.
Co-administration of ticagrelor with heparin, enoxaparin or desmopressin had no effect on activated partial thromboplastin time (aPTT), activated coagulation time (ACT) or factor Xa assays. However, due to potential pharmacodynamic interactions, caution should be exercised with the concomitant administration of ticagrelor with medicinal products known to alter haemostasis.

Due to reports of cutaneous bleeding abnormalities with SSRIs (e.g. paroxetine, sertraline and citalopram), caution is advised when administering SSRIs with ticagrelor as this may increase the risk of bleeding.

4.6 Fertility, pregnancy and lactation

Women of childbearing potential
Women of childbearing potential should use appropriate contraceptive measures to avoid pregnancy during ticagrelor therapy.

Pregnancy
There are no or limited amount of data from the use of ticagrelor in pregnant women. Studies in animals have shown reproductive toxicity (see section 5.3). Ticagrelor is not recommended during pregnancy.

Breast-feeding
Available pharmacodynamic/toxicological data in animals have shown excretion of ticagrelor and its active metabolites in milk (see section 5.3). A risk to newborns/infants cannot be excluded. A decision must be made whether to discontinue breast-feeding or to discontinue/abstain from ticagrelor therapy taking into account the benefit of breast-feeding for the child and the benefit of therapy for the woman.

Fertility
Ticagrelor had no effect on male or female fertility in animals (see section 5.3).

4.7 Effects on ability to drive and use machines

Ticagrelor has no or negligible influence on the ability to drive and use machines. During treatment with ticagrelor, dizziness and confusion have been reported. Therefore, patients who experience these symptoms should be cautious while driving or using machines.

4.8 Undesirable effects

Summary of the safety profile
The safety profile of ticagrelor has been evaluated in two large phase 3 outcome trials (PLATO and PEGASUS) including more than 39,000 patients (see section 5.1).

In PLATO, patients on ticagrelor had a higher incidence of discontinuation due to adverse events than clopidogrel (7.4% vs. 5.4%). In PEGASUS, patients on ticagrelor had a higher incidence of discontinuation due to adverse events compared to ASA therapy alone (16.1% for ticagrelor 60 mg with ASA vs. 8.5% for ASA therapy alone). The most commonly reported adverse reactions in patients treated with ticagrelor were bleeding and dyspnoea (see section 4.4).

Tabulated list of adverse reactions
The following adverse reactions have been identified following studies or have been reported in post-marketing experience with ticagrelor (Table 1).
Adverse reactions are listed by MedDRA System Organ Class (SOC). Within each SOC the adverse reactions are ranked by frequency category. Frequency categories are defined according to the following conventions: Very common (≥1/10), common (≥1/100 to <1/10), uncommon (≥1/1,000 to <1/100), rare (≥1/10,000 to <1/1,000), very rare (<1/10,000), not known (cannot be estimated from the available data).

Table 1 – Adverse reactions by frequency and system organ class (SOC)

<table>
<thead>
<tr>
<th>System Organ Classification</th>
<th>Very Common</th>
<th>Common</th>
<th>Uncommon</th>
</tr>
</thead>
<tbody>
<tr>
<td>Neoplasms benign, malignant and unspecified (including cysts and polyps)</td>
<td></td>
<td>Tumour bleedings(a)</td>
<td></td>
</tr>
<tr>
<td>Blood and lymphatic system disorders</td>
<td>Blood disorder bleedings(^b)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Immune system disorders</td>
<td></td>
<td></td>
<td>Hypersensitivity including angioedema(^c)</td>
</tr>
<tr>
<td>Metabolism and nutrition disorders</td>
<td>Hyperuricaemia(^d)</td>
<td>Gout/Gouty Arthritis</td>
<td></td>
</tr>
<tr>
<td>Psychiatric disorders</td>
<td></td>
<td></td>
<td>Confusion</td>
</tr>
<tr>
<td>Nervous system disorders</td>
<td></td>
<td>Dizziness, Syncope, Headache</td>
<td>Intracranial haemorrhage</td>
</tr>
<tr>
<td>Eye disorders</td>
<td></td>
<td></td>
<td>Eye haemorrhage(^e)</td>
</tr>
<tr>
<td>Ear and labyrinth disorders</td>
<td></td>
<td>Vertigo</td>
<td>Ear haemorrhage</td>
</tr>
<tr>
<td>Vascular disorders</td>
<td></td>
<td>Hypotension</td>
<td></td>
</tr>
<tr>
<td>Respiratory, thoracic and mediastinal disorders</td>
<td>Dyspnoea</td>
<td>Respiratory system bleedings(^f)</td>
<td></td>
</tr>
<tr>
<td>Gastrointestinal disorders</td>
<td></td>
<td>Gastrointestinal haemorrhage(^g), Diarrhoea, Nausea, Dyspepsia, Constipation</td>
<td>Retroperitoneal haemorrhage</td>
</tr>
<tr>
<td>Skin and subcutaneous tissue disorders</td>
<td></td>
<td>Subcutaneous or dermal bleeding(^h), Rash, Pruritus</td>
<td></td>
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<tr>
<td>Musculoskeletal connective tissue and bone</td>
<td></td>
<td></td>
<td>Muscular bleedings(^i)</td>
</tr>
<tr>
<td>Renal and urinary disorders</td>
<td></td>
<td>Urinary tract bleeding(^j)</td>
<td></td>
</tr>
</tbody>
</table>
### System Organ Classification

<table>
<thead>
<tr>
<th>Reproductive system and breast disorders</th>
<th>Very Common</th>
<th>Common</th>
<th>Uncommon</th>
</tr>
</thead>
</table>

**Investigations**

<table>
<thead>
<tr>
<th>Blood creatinine increased&lt;sup&gt;d&lt;/sup&gt;</th>
</tr>
</thead>
</table>

**Injury, poisoning and procedural complications**

<table>
<thead>
<tr>
<th>Post procedural haemorrhage, Traumatic bleedings&lt;sup&gt;l&lt;/sup&gt;</th>
</tr>
</thead>
</table>

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**Description of selected adverse reactions**

**Bleeding**

**Bleeding findings in PLATO**

Overall outcome of bleeding rates in the PLATO study are shown in Table 2.

---

<sup>a</sup> e.g. bleeding from bladder cancer, gastric cancer, colon cancer

<sup>b</sup> e.g. increased tendency to bruise, spontaneous haematoma, haemorrhagic diathesis

<sup>c</sup> Identified in post-marketing experience

<sup>d</sup> Frequencies derived from lab observations (Uric acid increases to > upper limit of normal from baseline below or within reference range. Creatinine increases of > 50% from baseline.) and not crude adverse event report frequency.

<sup>e</sup> e.g. conjunctival, retinal, intraocular bleeding

<sup>f</sup> e.g. epistaxis, haemoptysis

<sup>g</sup> e.g. gingival bleeding, rectal haemorrhage, gastric ulcer haemorrhage

<sup>h</sup> e.g. ecchymosis, skin haemorrhage, petechiae

<sup>i</sup> e.g. haemarthrosis, muscle haemorrhage

<sup>j</sup> e.g. haematuria, cystitis haemorrhagic

<sup>k</sup> e.g. vaginal haemorrhage, haematospermia, postmenopausal haemorrhage

<sup>l</sup> e.g. contusion, traumatic haematoma, traumatic haemorrhage
Table 2 – Analysis of overall bleeding events, Kaplan-Meier estimates at 12 months (PLATO)

<table>
<thead>
<tr>
<th></th>
<th>Ticagrelor 90 mg twice daily N=9235</th>
<th>Clopidogrel N=9186</th>
<th>p-value*</th>
</tr>
</thead>
<tbody>
<tr>
<td>PLATO Total Major</td>
<td>11.6</td>
<td>11.2</td>
<td>0.4336</td>
</tr>
<tr>
<td>PLATO Major Fatal/Life-Threatening</td>
<td>5.8</td>
<td>5.8</td>
<td>0.6988</td>
</tr>
<tr>
<td>Non-CABG PLATO Major</td>
<td>4.5</td>
<td>3.8</td>
<td>0.0264</td>
</tr>
<tr>
<td>Non-Procedural PLATO Major</td>
<td>3.1</td>
<td>2.3</td>
<td>0.0058</td>
</tr>
<tr>
<td>PLATO Total Major + Minor</td>
<td>16.1</td>
<td>14.6</td>
<td>0.0084</td>
</tr>
<tr>
<td>Non-Procedural PLATO Major + Minor</td>
<td>5.9</td>
<td>4.3</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>TIMI-defined Major</td>
<td>7.9</td>
<td>7.7</td>
<td>0.5669</td>
</tr>
<tr>
<td>TIMI-defined Major + Minor</td>
<td>11.4</td>
<td>10.9</td>
<td>0.3272</td>
</tr>
</tbody>
</table>

Bleeding category definitions:

Major Fatal/Life-threatening Bleed: Clinically apparent with >50 g/l decrease in haemoglobin or ≥4 red cell units transfused; or fatal; or intracranial; or intrapericardial with cardiac tamponade; or with hypovolaemic shock or severe hypotension requiring pressors or surgery.

Major Other: Clinically apparent with 30 50 g/L decrease in haemoglobin or 2-3 red cell units transfused; or significantly disabling.

Minor Bleed: Requires medical intervention to stop or treat bleeding.

TIMI Major Bleed: Clinically apparent with >50 g/L decrease in haemoglobin or intracranial haemorrhage.

TIMI Minor Bleed: Clinically apparent with 30 50 g/l decrease in haemoglobin.

*p-value calculated from Cox proportional hazards model with treatment group as the only explanatory variable.

Ticagrelor and clopidogrel did not differ in rates of PLATO Major Fatal/Life-threatening bleeding, PLATO total Major bleeding, TIMI Major bleeding, or TIMI Minor bleeding (Table 2). However, more PLATO combined Major + Minor bleeding occurred with ticagrelor compared with clopidogrel. Few patients in PLATO had fatal bleeds: 20 (0.2%) for ticagrelor and 23 (0.3%) for clopidogrel (see section 4.4).

Age, sex, weight, race, geographic region, concurrent conditions, concomitant therapy, and medical history, including a previous stroke or transient ischaemic attack, all did not predict either overall or non-procedural PLATO Major bleeding. Thus no particular group was identified at risk for any subset of bleeding.

CABG-related bleeding: In PLATO, 42% of the 1584 patients (12% of cohort) who underwent coronary artery bypass graft (CABG) surgery had a PLATO Major Fatal/Life-threatening bleeding with no difference between treatment groups. Fatal CABG bleeding occurred in 6 patients in each treatment group (see section 4.4).

Non-CABG related bleeding and non-procedural related bleeding: Ticagrelor and clopidogrel did not differ in non-CABG PLATO-defined Major Fatal/Life-threatening bleeding, but PLATO-defined Total Major, TIMI Major, and TIMI Major + Minor bleeding were more common with ticagrelor. Similarly, when removing all procedure related bleeds, more bleeding occurred with ticagrelor than with clopidogrel (Table 2). Discontinuation of treatment due to non-procedural bleeding was more common for ticagrelor (2.9%) than for clopidogrel (1.2%; p<0.001).

Intracranial bleeding: There were more intracranial non-procedural bleeds with ticagrelor (n=27 bleeds in 26 patients, 0.3%) than with clopidogrel (n=14 bleeds, 0.2%), of which 11 bleeds with ticagrelor and 1 with clopidogrel were fatal. There was no difference in overall fatal bleeds.

Bleeding findings in PEGASUS

Overall outcome of bleeding events in the PEGASUS study are shown in Table 3.
Table 3 – Analysis of overall bleeding events, Kaplan-Meier estimates at 36 months (PEGASUS)

<table>
<thead>
<tr>
<th>Safety Endpoints</th>
<th>KM%</th>
<th>Hazard Ratio (95% CI)</th>
<th>KM%</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>TIMI-defined bleeding categories</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>TIMI Major</td>
<td>2.3</td>
<td>2.32 (1.68, 3.21)</td>
<td>1.1</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Fatal</td>
<td>0.3</td>
<td>1.00 (0.44, 2.27)</td>
<td>0.3</td>
<td>1.0000</td>
</tr>
<tr>
<td>ICH</td>
<td>0.6</td>
<td>1.33 (0.77, 2.31)</td>
<td>0.5</td>
<td>0.3130</td>
</tr>
<tr>
<td>Other TIMI Major</td>
<td>1.6</td>
<td>3.61 (2.31, 5.65)</td>
<td>0.5</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>TIMI Major or Minor</td>
<td>3.4</td>
<td>2.54 (1.93, 3.35)</td>
<td>1.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>TIMI Major or Minor or Requiring medical attention</td>
<td>16.6</td>
<td>2.64 (2.35, 2.97)</td>
<td>7.0</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td><strong>PLATO-defined bleeding categories</strong></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>PLATO Major</td>
<td>3.5</td>
<td>2.57 (1.95, 3.37)</td>
<td>1.4</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Fatal/Life-threatening</td>
<td>2.4</td>
<td>2.38 (1.73, 3.26)</td>
<td>1.1</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Other PLATO Major</td>
<td>1.1</td>
<td>3.37 (1.95, 5.83)</td>
<td>0.3</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>PLATO Major or Minor</td>
<td>15.2</td>
<td>2.71 (2.40, 3.08)</td>
<td>6.2</td>
<td>&lt;0.0001</td>
</tr>
</tbody>
</table>

Bleeding category definitions:

TIMI Major: Fatal bleeding, OR any intracranial bleeding, OR clinically overt signs of haemorrhage associated with a drop in haemoglobin (Hgb) of ≥50 g/L, or when Hgb is not available, a fall in haematocrit (Hct) of 15%.

Fatal: A bleeding event that directly led to death within 7 days.

ICH: Intracranial haemorrhage.

Other TIMI Major: Non-fatal non-ICH TIMI Major bleeding.

TIMI Minor: Clinically apparent with 30-50 g/L decrease in haemoglobin.

TIMI Requiring medical attention: Requiring intervention, OR leading to hospitalization, OR prompting evaluation.

PLATO Major Fatal/life-threatening: Fatal bleeding, OR any intracranial bleeding, OR intrapericardial with cardiac tamponade, OR with hypovolaemic shock or severe hypotension requiring pressors/inotropes or surgery OR clinically apparent with >50 g/L decrease in haemoglobin or ≥4 red cell units transfused.

PLATO Major Other: Significantly disabling, OR clinically apparent with 30-50 g/L decrease in haemoglobin, OR 2-3 red cell units transfused.

PLATO Minor: Requires medical intervention to stop or treat bleeding.

In PEGASUS, TIMI Major bleeding for ticagrelor 60 mg twice daily was higher than for ASA alone. No increased bleeding risk was seen for fatal bleeding and only a minor increase was
observed in intracranial haemorrhages, as compared to ASA therapy alone. There were few fatal bleeding events in the study, 11 (0.3%) for ticagrelor 60 mg and 12 (0.3%) for ASA therapy alone. The observed increased risk of TIMI Major bleeding with ticagrelor 60 mg was primarily due to a higher frequency of Other TIMI Major bleedings driven by events in the gastrointestinal SOC.

Increased bleeding patterns similar to TIMI Major were seen for TIMI Major or Minor and PLATO Major and PLATO Major or Minor bleeding categories (see Table 3). Discontinuation of treatment due to bleeding was more common with ticagrelor 60 mg compared to ASA therapy alone (6.2% and 1.5%, respectively). The majority of these bleedings were of less severity (classified as TIMI Requiring medical attention), e.g. epistaxis, bruising and haematomas.

The bleeding profile of ticagrelor 60 mg was consistent across multiple pre-defined subgroups (e.g. by age, gender, weight, race, geographic region, concurrent conditions, concomitant therapy, and medical history) for TIMI Major, TIMI Major or Minor and PLATO Major bleeding events.

**Intracranial bleeding:** Spontaneous ICHs were reported in similar rates for ticagrelor 60 mg and ASA therapy alone (n=13, 0.2% in both treatment groups). Traumatic and procedural ICHs showed a minor increase with ticagrelor 60 mg treatment, (n=15, 0.2%) compared with ASA therapy alone (n=10, 0.1%). There were 6 fatal ICHs with ticagrelor 60 mg and 5 fatal ICHs with ASA therapy alone. The incidence of intracranial bleeding was low in both treatment groups given the significant comorbidity and CV risk factors of the population under study.

**Dyspnoea**

Dyspnoea, a sensation of breathlessness, is reported by patients treated with Brilique. In PLATO, dyspnoea adverse events (AEs) (dyspnoea, dyspnoea at rest, dyspnoea exertional, dyspnoea paroxysmal nocturnal and nocturnal dyspnoea), when combined, was reported by 13.8% of patients treated with ticagrelor and by 7.8% of patients treated with clopidogrel. In 2.2% of patients taking ticagrelor and by 0.6% taking clopidogrel investigators considered the dyspnoea causally related to treatment in the PLATO study and few were serious (0.14% ticagrelor; 0.02% clopidogrel), (see section 4.4). Most reported symptoms of dyspnoea were mild to moderate in intensity, and most were reported as a single episode early after starting treatment.

Compared with clopidogrel, patients with asthma/COPD treated with ticagrelor may have an increased risk of experiencing non-serious dyspnoea (3.29% ticagrelor versus 0.53% clopidogrel) and serious dyspnoea (0.38% ticagrelor versus 0.00% clopidogrel). In absolute terms, this risk was higher than in the overall PLATO population. Ticagrelor should be used with caution in patients with history of asthma and/or COPD (see section 4.4).

About 30% of episodes resolved within 7 days. PLATO included patients with baseline congestive heart failure, COPD, or asthma; these patients, and the elderly, were more likely to report dyspnoea. For Brilique, 0.9% of patients discontinued study drug because of dyspnoea compared with 0.1% taking clopidogrel. The higher incidence of dyspnoea with Brilique is not associated with new or worsening heart or lung disease (see section 4.4). Brilique does not affect tests of pulmonary function.

In PEGASUS, dyspnoea was reported in 14.2% of patients taking ticagrelor 60 mg twice daily and in 5.5% of patients taking ASA alone. As in PLATO, most reported dyspnoea was mild to moderate in intensity (see section 4.4). Patients who reported dyspnoea tended to be older and more frequently had dyspnoea, COPD or asthma at baseline.
**Investigations**

Uric acid elevations: In PLATO, serum uric acid increased to more than upper limit of normal in 22% of patients receiving ticagrelor compared to 13% of patients receiving clopidogrel. The corresponding numbers in PEGASUS were 9.1%, 8.8% and 5.5% for ticagrelor 90 mg, 60 mg and placebo, respectively. Mean serum uric acid increased approximately 15% with ticagrelor compared to approximately 7.5% with clopidogrel and after treatment was stopped, decreased to approximately 7% on ticagrelor but with no decrease observed for clopidogrel. In PEGASUS, a reversible increase in mean serum uric acid levels of 6.3% and 5.6% was found for ticagrelor 90 mg and 60 mg, respectively, compared to a 1.5% decrease in the placebo group. In PLATO, the frequency of gouty arthritis was 0.2% for ticagrelor vs. 0.1% for clopidogrel. The corresponding numbers for gout/gouty arthritis in PEGASUS were 1.6%, 1.5% and 1.1% for ticagrelor 90 mg, 60 mg and placebo, respectively.

**Reporting of suspected adverse reactions**

Reporting suspected adverse reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions via the national reporting system listed in Appendix V.

**4.9 Overdose**

Ticagrelor is well tolerated in single doses up to 900 mg. Gastrointestinal toxicity was dose-limiting in a single ascending dose study. Other clinically meaningful adverse reactions which may occur with overdose include dyspnoea and ventricular pauses (see section 4.8).

In the event of an overdose, the above potential adverse reactions could occur and ECG monitoring should be considered.

There is currently no known antidote to reverse the effects of ticagrelor, and ticagrelor is not expected to be dialysable (see section 4.4). Treatment of overdose should follow local standard medical practice. The expected effect of excessive ticagrelor dosing is prolonged duration of bleeding risk associated with platelet inhibition. Platelet transfusion is unlikely to be of clinical benefit in patients with bleeding (see section 4.4). If bleeding occurs other appropriate supportive measures should be taken.

**5. PHARMACOLOGICAL PROPERTIES**

**5.1 Pharmacodynamic properties**

Pharmacotherapeutic group: Platelet aggregation inhibitors excluding heparin, ATC code: B01AC24

**Mechanism of action**

Brilique contains ticagrelor, a member of the chemical class cyclopentyltriazolopyrimidines (CPTP), which is an oral, direct acting, selective and reversibly binding P2Y$_12$ receptor antagonist that prevents ADP-mediated P2Y$_12$ dependent platelet activation and aggregation. Ticagrelor does not prevent ADP binding but when bound to the P2Y$_12$ receptor prevents ADP-induced signal transduction. Since platelets participate in the initiation and/or evolution of thrombotic complications of atherosclerotic disease, inhibition of platelet function has been shown to reduce the risk of CV events such as death, MI or stroke.
Ticagrelor also increases local endogenous adenosine levels by inhibiting the equilibrative nucleoside transporter -1 (ENT-1).

Ticagrelor has been documented to augment the following adenosine-induced effects in healthy subjects and in patients with ACS: vasodilation (measured by coronary blood flow increases in healthy volunteers and ACS patients; headache), inhibition of platelet function (in human whole blood \textit{in vitro}) and dyspnoea. However, a link between the observed increases in adenosine and clinical outcomes (e.g. morbidity-mortality) has not been clearly elucidated.

\textbf{Pharmacodynamic effects}

\textit{Onset of action}
In patients with stable coronary artery disease (CAD) on ASA, ticagrelor demonstrates a rapid onset of pharmacological effect as demonstrated by a mean inhibition of platelet aggregation (IPA) for ticagrelor at 0.5 hours after 180 mg loading dose of about 41%, with the maximum IPA effect of 89% by 2-4 hours post dose, and maintained between 2-8 hours. 90% of patients had final extent IPA>70% by 2 hours post dose.

\textit{Offset of action}
If a CABG procedure is planned, ticagrelor bleeding risk is increased compared to clopidogrel when discontinued within less than 96 hours prior to procedure.

\textit{Switching data}
Switching from clopidogrel 75 mg to ticagrelor 90 mg twice daily results in an absolute IPA increase of 26.4% and switching from ticagrelor to clopidogrel results in an absolute IPA decrease of 24.5%. Patients can be switched from clopidogrel to ticagrelor without any interruption of antiplatelet effect (see section 4.2).

\textbf{Clinical efficacy and safety}
The clinical evidence for the efficacy and safety of ticagrelor is derived from two phase 3 trials:

- The PLATO [PLATelet Inhibition and Patient Outcomes] study, a comparison of ticagrelor to clopidogrel, both given in combination with ASA and other standard therapy.
- The PEGASUS TIMI-54 [PrEvention with TicaGrelor of SecondAry Thrombotic Events in High-RiSk AcUte Coronary Syndrome Patients] study, a comparison of ticagrelor combined with ASA to ASA therapy alone.

\textbf{PLATO study (Acute Coronary Syndromes)}
The PLATO study included 18,624 patients who presented within 24 hours of onset of symptoms of unstable angina (UA), non ST elevation myocardial infarction (NSTEMI) or ST elevation myocardial infarction (STEMI), and were initially managed medically, or with percutaneous coronary intervention (PCI), or with CABG.

\textit{Clinical efficacy}
On a background of daily ASA, ticagrelor 90 mg twice daily showed superiority to 75 mg daily clopidogrel in preventing the composite endpoint of CV death, MI, or stroke, with the difference driven by CV death and MI. Patients received a 300 mg loading dose of clopidogrel (600 mg possible if having PCI) or 180 mg of ticagrelor.

The result appeared early (absolute risk reduction [ARR] 0.6% and relative risk reduction [RRR] of 12% at 30 days), with a constant treatment effect over the entire 12 month period, yielding ARR 1.9% per year with RRR of 16%. This suggests it is appropriate to treat patients with ticagrelor 90
mg twice daily for 12 months (see section 4.2). Treating 54 ACS patients with ticagrelor instead of clopidogrel will prevent 1 atherothrombotic event; treating 91 will prevent 1 CV death (see Figure 1 and Table 4).

The treatment effect of ticagrelor over clopidogrel appears consistent across many subgroups, including weight; sex; medical history of diabetes mellitus, transient ischaemic attack or non-haemorrhagic stroke, or revascularisation; concomitant therapies including heparins, GpIIb/IIIa inhibitors and proton pump inhibitors (see section 4.5); final index event diagnosis (STEMI, NSTEMI, or UA); and treatment pathway intended at randomisation (invasive or medical).

A weakly significant treatment interaction was observed with region whereby the hazard ratio (HR) for the primary endpoint favours ticagrelor in the rest of world but favours clopidogrel in North America, which represented approximately 10% of the overall population studied (interaction $p$-value=0.045). Exploratory analyses suggest a possible association with ASA dose such that reduced efficacy was observed with ticagrelor with increasing ASA doses. Chronic daily ASA doses to accompany Brilique should be 75-150 mg (see sections 4.2 and 4.4).

Figure 1 shows the estimate of the risk to the first occurrence of any event in the composite efficacy endpoint.

**Figure 1 – Analysis of primary clinical composite endpoint of CV death, MI and stroke (PLATO)**

![Graph showing Kaplan-Meier percentage](image)

<table>
<thead>
<tr>
<th>N at risk</th>
<th>Days from Randomisation</th>
</tr>
</thead>
<tbody>
<tr>
<td>T</td>
<td>9333 8628 8460 8219 6743 5161 4147</td>
</tr>
<tr>
<td>C</td>
<td>9291 8521 8362 8124 6650 5096 4074</td>
</tr>
</tbody>
</table>
Ticagrelor reduced the occurrence of the primary composite endpoint compared to clopidogrel in both the UA/NSTEMI and STEMI population (Table 4). Thus, Brilique 90 mg twice daily together with low-dose ASA can be used in patients with ACS (unstable angina, non ST elevation Myocardial Infarction [NSTEMI] or ST elevation Myocardial Infarction [STEMI]); including patients managed medically, and those who are managed with percutaneous coronary intervention (PCI) or coronary artery by-pass grafting (CABG).

Table 4 - Analysis of primary and secondary efficacy endpoints (PLATO)

<table>
<thead>
<tr>
<th>Event</th>
<th>Ticagrelor 90 mg twice daily (% patients with event)</th>
<th>Clopidogrel 75 mg once daily (% patients with event)</th>
<th>ARR² (%/yr)</th>
<th>RRR² (%) (95% CI)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>CV death, MI (excl. silent MI) or stroke</td>
<td>9.3</td>
<td>10.9</td>
<td>1.9</td>
<td>16 (8, 23)</td>
<td>0.0003</td>
</tr>
<tr>
<td>Invasive intent</td>
<td>8.5</td>
<td>10.0</td>
<td>1.7</td>
<td>16 (6, 25)</td>
<td>0.0025</td>
</tr>
<tr>
<td>Medical intent</td>
<td>11.3</td>
<td>13.2</td>
<td>2.3</td>
<td>15 (0.3, 27)</td>
<td>0.0444d</td>
</tr>
<tr>
<td>CV death</td>
<td>3.8</td>
<td>4.8</td>
<td>1.1</td>
<td>21 (9, 31)</td>
<td>0.0013</td>
</tr>
<tr>
<td>MI (excl. silent MI)º</td>
<td>5.4</td>
<td>6.4</td>
<td>1.1</td>
<td>16 (5, 25)</td>
<td>0.0045</td>
</tr>
<tr>
<td>Stroke</td>
<td>1.3</td>
<td>1.1</td>
<td>-0.2</td>
<td>-17 (-52, 9)</td>
<td>0.2249</td>
</tr>
<tr>
<td>All-cause mortality, MI (excl. silent MI), or stroke</td>
<td>9.7</td>
<td>11.5</td>
<td>2.1</td>
<td>16 (8, 23)</td>
<td>0.0001</td>
</tr>
<tr>
<td>CV death, total MI, stroke, SRI, RI, TIA, or other ATEº</td>
<td>13.8</td>
<td>15.7</td>
<td>2.1</td>
<td>12 (5, 19)</td>
<td>0.0006</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>4.3</td>
<td>5.4</td>
<td>1.4</td>
<td>22 (11, 31)</td>
<td>0.0003d</td>
</tr>
<tr>
<td>Definite stent thrombosis</td>
<td>1.2</td>
<td>1.7</td>
<td>0.6</td>
<td>32 (8, 49)</td>
<td>0.0123d</td>
</tr>
</tbody>
</table>

ARR = absolute risk reduction; RRR = relative risk reduction = (1-Hazard ratio) x 100%. A negative RRR indicates a relative risk increase.
Excluding silent MI.
SRI = serious recurrent ischaemia; RI = recurrent ischaemia; TIA = transient ischaemic attack; ATE = arterial thrombotic event. Total MI includes silent MI, with date of event set to date when discovered.
Nominal significance value; all others are formally statistically significant by pre-defined hierarchical testing.

PLATO genetic substudy
CYP2C19 and ABCB1 genotyping of 10,285 patients in PLATO provided associations of genotype groups with PLATO outcomes. The superiority of ticagrelor over clopidogrel in reducing major CV events was not significantly affected by patient CYP2C19 or ABCB1 genotype. Similar to the overall PLATO study, total PLATO Major bleeding did not differ between ticagrelor and clopidogrel, regardless of CYP2C19 or ABCB1 genotype. Non-CABG PLATO Major bleeding was increased with ticagrelor compared clopidogrel in patients with one or more CYP2C19 loss of function alleles, but similar to clopidogrel in patients with no loss of function allele.

Combined efficacy and safety composite
A combined efficacy and safety composite (CV death, MI, stroke, or PLATO-defined ‘Total Major’ bleeding) indicates that the benefit in efficacy of ticagrelor compared to clopidogrel is not offset by the major bleeding events (ARR 1.4%, RRR 8%, HR 0.92; p=0.0257) over 12 months after ACS.
Clinical safety

Holter substudy

To study the occurrence of ventricular pauses and other arrhythmic episodes during PLATO, investigators performed Holter monitoring in a subset of nearly 3000 patients, of whom approximately 2000 had recordings both in the acute phase of their ACS and after one month. The primary variable of interest was the occurrence of ventricular pauses ≥3 seconds. More patients had ventricular pauses with ticagrelor (6.0%) than with clopidogrel (3.5%) in the acute phase; and 2.2% and 1.6% respectively after 1 month (see section 4.4). The increase in ventricular pauses in the acute phase of ACS was more pronounced in ticagrelor patients with history of CHF (9.2% versus 5.4% in patients without CHF history; for clopidogrel patients, 4.0% in those with versus 3.6% in those without CHF history). This imbalance did not occur at one month: 2.0% versus 2.1% for ticagrelor patients with and without CHF history respectively; and 3.8% versus 1.4% with clopidogrel. There were no adverse clinical consequences associated with this imbalance (including pacemaker insertions) in this population of patients.

PEGASUS study (History of Myocardial Infarction)

The PEGASUS TIMI-54 study was a 21,162 patient, event-driven, randomised, double-blind, placebo-controlled, parallel group, international multicentre study to assess the prevention of atherothrombotic events with ticagrelor given at 2 doses (either 90 mg twice daily or 60 mg twice daily) combined with low dose ASA (75-150 mg), compared to ASA therapy alone in patients with history of MI and additional risk factors for atherothrombosis.

Patients were eligible to participate if they were aged 50 years or over, with a history of MI (1 to 3 years prior to randomisation), and had at least one of the following risk factors for atherothrombosis: age ≥65 years, diabetes mellitus requiring medication, a second prior MI, evidence of multivessel CAD, or chronic non-end-stage renal dysfunction.

Patients were ineligible if there was planned use of a P2Y12 receptor antagonist, dipyridamole, cilostazol, or anticoagulant therapy during the study period; if they had a bleeding disorder or a history of an ischaemic stroke or intracranial bleeding, a central nervous system tumour, or an intracranial vascular abnormality; if they had had gastrointestinal bleeding within the previous 6 months or major surgery within the previous 30 days.

Clinical efficacy
Figure 2 - Analysis of primary clinical composite endpoint of CV death, MI and stroke (PEGASUS)

Table 5 - Analysis of primary and secondary efficacy endpoints (PEGASUS)

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Ticagrelor 60 mg twice daily +ASA</th>
<th>ASA alone</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>N = 7045</td>
<td>N = 7067</td>
</tr>
<tr>
<td></td>
<td>Patients with events KM % HR (95% CI)</td>
<td>Patients with events KM %</td>
</tr>
<tr>
<td>Primary endpoint</td>
<td></td>
<td>p-value</td>
</tr>
<tr>
<td>Composite of CV Death/MI/Stroke</td>
<td>487 (6.9%) 7.8% 0.84 (0.74, 0.95)</td>
<td>578 (8.2%) 9.0%</td>
</tr>
<tr>
<td>CV death</td>
<td>174 (2.5%) 2.9% 0.83 (0.68, 1.01)</td>
<td>210 (3.0%) 3.4%</td>
</tr>
<tr>
<td>MI</td>
<td>285 (4.0%) 4.5% 0.84 (0.72, 0.98)</td>
<td>338 (4.8%) 5.2%</td>
</tr>
<tr>
<td>Stroke</td>
<td>91 (1.3%) 1.5% 0.75 (0.57, 0.98)</td>
<td>122 (1.7%) 1.9%</td>
</tr>
</tbody>
</table>
### Ticagrelor 60 mg twice daily + ASA

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Patients with events</th>
<th>KM %</th>
<th>HR (95% CI)</th>
<th>Patients with events</th>
<th>KM %</th>
<th>p-value</th>
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<tr>
<td>Secondary endpoint</td>
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<tr>
<td>CV death</td>
<td>174 (2.5%)</td>
<td>2.9%</td>
<td>0.83 (0.68, 1.01)</td>
<td>210 (3.0%)</td>
<td>3.4%</td>
<td>-</td>
</tr>
<tr>
<td>All-cause mortality</td>
<td>289 (4.1%)</td>
<td>4.7%</td>
<td>0.89 (0.76, 1.04)</td>
<td>326 (4.6%)</td>
<td>5.2%</td>
<td>-</td>
</tr>
</tbody>
</table>

Hazard ratio and p-values are calculated separately for ticagrelor vs. ASA therapy alone from Cox proportional hazards model with treatment group as the only explanatory variable.

KM percentage calculated at 36 months.

Note: the number of first events for the components CV death, MI and stroke are the actual number of first events for each component and do not add up to the number of events in the composite endpoint

(s) Indicates statistical significance.

CI = Confidence interval; CV = Cardiovascular; HR = Hazard ratio; KM = Kaplan-Meier; MI = Myocardial infarction; N = Number of patients.

Both 60 mg twice daily and 90 mg twice daily regimens of ticagrelor in combination with ASA were superior to ASA alone in the prevention of atherothrombotic events (composite endpoint: CV death, MI and stroke), with a consistent treatment effect over the entire study period, yielding a 16% RRR and 1.27% ARR for ticagrelor 60 mg and a 15% RRR and 1.19% ARR for ticagrelor 90 mg.

Although the efficacy profile of 90 mg and 60 mg were similar, there is evidence that the lower dose has a better tolerability and safety profile in relation to risk of the bleeding and dyspnoea. Therefore only Brilique 60 mg twice daily co-administered with ASA is recommended for the prevention atherothrombotic events (CV death, MI and stroke) in patients with a history of MI and a high risk of developing an atherothrombotic event.

Relative to ASA alone, ticagrelor 60 mg twice daily significantly reduced the primary composite endpoint of CV death, MI and stroke. Each of the components contributed to the reduction in the primary composite endpoint (CV death 17% RRR, MI 16% RRR, and stroke 25% RRR).

The RRR for the composite endpoint from 1 to 360 days (17% RRR) and from 361 days and onwards (16% RRR) was similar. There are limited data on the efficacy and safety of ticagrelor beyond 3 years of extended treatment.

There was no evidence of benefit (no reduction in the primary composite endpoint of CV death, MI and stroke, but an increase in major bleeding) when ticagrelor 60 mg twice daily was introduced in clinically stable patients >2 years from the MI, or more than one year after stopping previous ADP receptor inhibitor treatment (see also section 4.2).

**Clinical safety**

The rate of discontinuations with ticagrelor 60 mg due to bleeding and dyspnoea was higher in patients >75 years (42%) than in younger patients (range: 23-31%), with a difference versus placebo higher than 10% (42% vs. 29%) in patients >75 years.

**Paediatric population**

The European Medicines Agency has waived the obligation to submit the results of studies with Brilique in all subsets of the paediatric population in the granted indication (see section 4.2 for information on paediatric use).
5.2 Pharmacokinetic properties

Ticagrelor demonstrates linear pharmacokinetics and exposure to ticagrelor and the active metabolite (AR-C124910XX) are approximately dose proportional up to 1260 mg.

Absorption
Absorption of ticagrelor is rapid with a median $t_{\text{max}}$ of approximately 1.5 hours. The formation of the major circulating metabolite AR-C124910XX (also active) from ticagrelor is rapid with a median $t_{\text{max}}$ of approximately 2.5 hours. Following an oral ticagrelor 90 mg single dose under fasted conditions in healthy subjects, $C_{\text{max}}$ is 529 ng/ml and AUC is 3451 ng*h/ml. The metabolite parent ratios are 0.28 for $C_{\text{max}}$ and 0.42 for AUC. The pharmacokinetics of ticagrelor and AR-C124910XX in patients with a history of MI were generally similar to that in the ACS population. Based on a population pharmacokinetic analysis of the PEGASUS study the median ticagrelor $C_{\text{max}}$ was 391 ng/ml and AUC was 3801 ng*h/ml at steady state for ticagrelor 60 mg. For ticagrelor 90 mg $C_{\text{max}}$ was 627 ng/ml and AUC was 6255 ng*h/ml at steady state.

The mean absolute bioavailability of ticagrelor was estimated to be 36%. Ingestion of a high-fat meal resulted in a 21% increase in ticagrelor AUC and 22% decrease in the active metabolite $C_{\text{max}}$ but had no effect on ticagrelor $C_{\text{max}}$ or the AUC of the active metabolite. These small changes are considered of minimal clinical significance; therefore, ticagrelor can be given with or without food. Ticagrelor as well as the active metabolite are P-gp substrates.

Ticagrelor as crushed tablets mixed in water, given orally or administered through a nasogastric tube into the stomach, has a comparable bioavailability to whole tablets with regards to AUC and $C_{\text{max}}$ for ticagrelor and the active metabolite. Initial exposure (0.5 and 1 hour post-dose) from crushed ticagrelor tablets mixed in water was higher compared to whole tablets, with a generally identical concentration profile thereafter (2 to 48 hours).

Distribution
The steady state volume of distribution of ticagrelor is 87.5 l. Ticagrelor and the active metabolite is extensively bound to human plasma protein (>99.0%).

Biotransformation
CYP3A4 is the major enzyme responsible for ticagrelor metabolism and the formation of the active metabolite and their interactions with other CYP3A substrates ranges from activation through to inhibition.

The major metabolite of ticagrelor is AR-C124910XX, which is also active as assessed by in vitro binding to the platelet P2Y$_{12}$ ADP-receptor. The systemic exposure to the active metabolite is approximately 30-40% of that obtained for ticagrelor.

Elimination
The primary route of ticagrelor elimination is via hepatic metabolism. When radiolabelled ticagrelor is administered, the mean recovery of radioactivity is approximately 84% (57.8% in faeces, 26.5% in urine). Recoveries of ticagrelor and the active metabolite in urine were both less than 1% of the dose. The primary route of elimination for the active metabolite is most likely via biliary secretion. The mean $t_{1/2}$ was approximately 7 hours for ticagrelor and 8.5 hours for the active metabolite.

Special populations

Elderly

43
Higher exposures to ticagrelor (approximately 25% for both \(C_{\text{max}}\) and AUC) and the active metabolite were observed in elderly (≥75 years) ACS patients compared to younger patients by the population pharmacokinetic analysis. These differences are not considered clinically significant (see section 4.2).

**Paediatric population**
Ticagrelor has not been evaluated in a paediatric population (see sections 4.2 and 5.1).

**Gender**
Higher exposures to ticagrelor and the active metabolite were observed in women compared to men. These differences are not considered clinically significant.

**Renal impairment**
Exposure to ticagrelor was approximately 20% lower and exposure to the active metabolite was approximately 17% higher in patients with severe renal impairment (creatinine clearance <30 ml/min) compared to subjects with normal renal function (see section 4.2).

**Hepatic impairment**
\(C_{\text{max}}\) and AUC for ticagrelor were 12% and 23% higher in patients with mild hepatic impairment compared to matched healthy subjects, respectively, however, the IPA effect of ticagrelor was similar between the two groups. No dose adjustment is needed for patients with mild hepatic impairment. Ticagrelor has not been studied in patients with severe hepatic impairment and there is no pharmacokinetic information in patients with moderate hepatic impairment. In patients that had moderate or severe elevation in one or more liver function tests at baseline, ticagrelor plasma concentrations were on average similar or slightly higher as compared to those without baseline elevations. No dose adjustment is recommended in patients with moderate hepatic impairment (see sections 4.2 and 4.4).

**Ethnicity**
Patients of Asian descent have a 39% higher mean bioavailability compared to Caucasian patients. Patients self-identified as black had an 18% lower bioavailability of ticagrelor compared to Caucasian patients, in clinical pharmacology studies, the exposure (\(C_{\text{max}}\) and AUC) to ticagrelor in Japanese subjects was approximately 40% (20% after adjusting for body weight) higher compared to that in Caucasians. The exposure in patients self-identified as Hispanic or Latino was similar to that in Caucasians.

5.3 **Preclinical safety data**

Preclinical data for ticagrelor and its major metabolite have not demonstrated unacceptable risk for adverse effects for humans based on conventional studies of safety pharmacology, single and repeated dose toxicity and genotoxic potential.

Gastrointestinal irritation was observed in several animal species at clinical relevant exposure levels (see section 4.8).

In female rats, ticagrelor at high dose showed an increased incidence of uterine tumours (adenocarcinomas) and an increased incidence of hepatic adenomas. The mechanism for uterine tumours is likely hormonal imbalance which can lead to tumours in rats. The mechanism for the hepatic adenomas is likely due to a rodent-specific enzyme induction in the liver. Thus, the carcinogenicity findings are considered unlikely to be relevant for humans.
In rats minor developmental anomalies were seen at a maternal toxic dose (safety margin of 5.1). In rabbits a slight delay in hepatic maturity and skeletal development was seen in foetuses from dams at high dose without showing maternal toxicity (safety margin of 4.5).

Studies in rats and rabbits have shown reproductive toxicity, with slightly reduced maternal body weight gain and reduced neonatal viability and birth weight, with delayed growth. Ticagrelor produced irregular cycles (mostly extended cycles) in female rats, but did not affect overall fertility in male and female rats. Pharmacokinetic studies performed with radio-labelled ticagrelor have shown that the parent compound and its metabolites are excreted in the milk of rats (see section 4.6).

6. PHARMACEUTICAL PARTICULARS

6.1 List of excipients

Tablet core
- Mannitol (E421)
- Calcium hydrogen phosphate dihydrate
- Magnesium stearate (E470b)
- Sodium starch glycolate type A
- Hydroxypropylcellulose (E463)

Tablet coating
- Talc
- Titanium dioxide (E171)
- Iron oxide yellow (E172)
- Macrogol 400
- Hypromellose (E464)

6.2 Incompatibilities

Not applicable.

6.3 Shelf life

3 years

6.4 Special precautions for storage

This medicinal product does not require any special storage conditions.

6.5 Nature and contents of container

- PVC-PVDC/Al transparent blister (with sun/moon symbols) of 10 tablets; cartons of 60 tablets (6 blisters) and 180 tablets (18 blisters).
- PVC-PVDC/Al transparent calendar blister (with sun/moon symbols) of 14 tablets; cartons of 14 tablets (1 blister), 56 tablets (4 blisters), and 168 tablets (12 blisters).
- PVC-PVDC/Al perforated unit dose transparent blister of 10 tablets; cartons of 100x1 tablets (10 blisters).

Not all pack sizes may be marketed.
6.6 Special precautions for disposal

No special requirements. Any unused medicinal product or waste material should be disposed of in accordance with local requirements.

7. MARKETING AUTHORISATION HOLDER

AstraZeneca AB
SE-151 85 Södertälje
Sweden

8. MARKETING AUTHORISATION NUMBER(S)

EU/1/10/655/001-006

9. DATE OF FIRST AUTHORISATION/RENEWAL OF THE AUTHORISATION

Date of first authorisation: 03 December 2010
Date of latest renewal: 17 July 2015

10. DATE OF REVISION OF THE TEXT

Detailed information on this medicinal product is available on the website of the European Medicines Agency http://www.ema.europa.eu.
ANNEX II

A. MANUFACTURER(S) RESPONSIBLE FOR BATCH RELEASE

B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE

C. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING AUTHORISATION

D. CONDITIONS OR RESTRICTIONS WITH REGARD TO THE SAFE AND EFFECTIVE USE OF THE MEDICINAL PRODUCT
A. MANUFACTURER(S) RESPONSIBLE FOR BATCH RELEASE

Name and address of the manufacturer(s) responsible for batch release

AstraZeneca AB
Gärtunavägen
SE-151 85 Södertälje
Sweden

AstraZeneca UK Limited
Silk Road Business Park
Macclesfield, Cheshire, SK10 2NA
United Kingdom

The printed package leaflet of the medicinal product must state the name and address of the manufacturer responsible for the release of the concerned batch.

B. CONDITIONS OR RESTRICTIONS REGARDING SUPPLY AND USE

Medicinal product subject to medical prescription.

C. OTHER CONDITIONS AND REQUIREMENTS OF THE MARKETING AUTHORISATION

- Periodic Safety Update Reports

The requirements for submission of periodic safety update reports for this medicinal product are set out in the list of Union reference dates (EURD list) provided for under Article 107c(7) of Directive 2001/83/EC and any subsequent updates published on the European medicines web-portal.

D. CONDITIONS OR RESTRICTIONS WITH REGARD TO THE SAFE AND EFFECTIVE USE OF THE MEDICINAL PRODUCT

- Risk Management Plan (RMP)

The MAH shall perform the required pharmacovigilance activities and interventions detailed in the agreed RMP presented in Module 1.8.2 of the Marketing Authorisation and any agreed subsequent updates of the RMP.

An updated RMP should be submitted:

- At the request of the European Medicines Agency;
- Whenever the risk management system is modified, especially as the result of new information being received that may lead to a significant change to the benefit/risk profile or as the result of an important (pharmacovigilance or risk minimisation) milestone being reached.
ANNEX III

LABELLING AND PACKAGE LEAFLET
A. LABELLING
PARTICULARS TO APPEAR ON THE OUTER PACKAGING CARTON

1. NAME OF THE MEDICINAL PRODUCT

Brilique 60 mg film-coated tablets
ticagrelor

2. STATEMENT OF ACTIVE SUBSTANCE(S)

Each film-coated tablet contains 60 mg ticagrelor

3. LIST OF EXCIPIENTS

4. PHARMACEUTICAL FORM AND CONTENTS

14 film-coated tablets
56 film-coated tablets
60 film-coated tablets
168 film-coated tablets
180 film-coated tablets

5. METHOD AND ROUTE(S) OF ADMINISTRATION

Read the package leaflet before use.
Oral use

6. SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN

Keep out of the sight and reach of children.

7. OTHER SPECIAL WARNING(S), IF NECESSARY

8. EXPIRY DATE

EXP
9. SPECIAL STORAGE CONDITIONS

10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE

11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

AstraZeneca AB
SE-151 85
Södertälje
Sweden

12. MARKETING AUTHOURISATION NUMBER(S)

EU/1/10/655/007 14 film-coated tablets
EU/1/10/655/008 56 film-coated tablets
EU/1/10/655/009 60 film-coated tablets
EU/1/10/655/010 168 film-coated tablets
EU/1/10/655/011 180 film-coated tablets

13. BATCH NUMBER

Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

brilique 60 mg

17. UNIQUE IDENTIFIER – 2D BARCODE

2D barcode carrying the unique identifier included.
18. UNIQUE IDENTIFIER - HUMAN READABLE DATA

PC:
SN:
NN:
## MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS

### BLISTER

<table>
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MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS

CALENDAR BLISTER

1. NAME OF THE MEDICINAL PRODUCT

Brilique 60 mg tablets
ticagrelor

2. NAME OF THE MARKETING AUTHORISATION HOLDER

AstraZeneca AB

3. EXPIRY DATE

EXP

4. BATCH NUMBER

Lot

5. OTHER

Mon Tue Wed Thu Fri Sat Sun
Sun/Moon symbol
### PARTICULARS TO APPEAR ON THE OUTER PACKAGING CARTON

1. **NAME OF THE MEDICINAL PRODUCT**
   - Brilique 90 mg film-coated tablets
   - ticagrelor

2. **STATEMENT OF ACTIVE SUBSTANCE(S)**
   - Each film-coated tablet contains 90 mg ticagrelor

3. **LIST OF EXCipients**

4. **PHARMACEUTICAL FORM AND CONTENTS**
   - 14 film-coated tablets
   - 56 film-coated tablets
   - 60 film-coated tablets
   - 100x1 film-coated tablets
   - 168 film-coated tablets
   - 180 film-coated tablets

5. **METHOD AND ROUTE(S) OF ADMINISTRATION**
   - Read the package leaflet before use.
   - Oral use

6. **SPECIAL WARNING THAT THE MEDICINAL PRODUCT MUST BE STORED OUT OF THE SIGHT AND REACH OF CHILDREN**
   - Keep out of the sight and reach of children.

7. **OTHER SPECIAL WARNING(S), IF NECESSARY**

8. **EXPIRY DATE**
   - EXP
9. SPECIAL STORAGE CONDITIONS

10. SPECIAL PRECAUTIONS FOR DISPOSAL OF UNUSED MEDICINAL PRODUCTS OR WASTE MATERIALS DERIVED FROM SUCH MEDICINAL PRODUCTS, IF APPROPRIATE

11. NAME AND ADDRESS OF THE MARKETING AUTHORISATION HOLDER

AstraZeneca AB
SE-151 85
Södertälje
Sweden

12. MARKETING AUTHORISATION NUMBER(S)

EU/1/10/655/001 60 film-coated tablets
EU/1/10/655/002 180 film-coated tablets
EU/1/10/655/003 14 film-coated tablets
EU/1/10/655/004 56 film-coated tablets
EU/1/10/655/005 168 film-coated tablets
EU/1/10/655/006 100x1 film-coated tablets

13. BATCH NUMBER

Lot

14. GENERAL CLASSIFICATION FOR SUPPLY

Medicinal product subject to medical prescription.

15. INSTRUCTIONS ON USE

16. INFORMATION IN BRAILLE

brilique 90 mg

17. UNIQUE IDENTIFIER – 2D BARCODE

2D barcode carrying the unique identifier included.
18. UNIQUE IDENTIFIER - HUMAN READABLE DATA

PC:
SN:
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<th><strong>MINIMUM PARTICULARS TO APPEAR ON BLISTERS OR STRIPS</strong></th>
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1. **NAME OF THE MEDICINAL PRODUCT**

   Brilique 90 mg tablets
ticagrelor

2. **NAME OF THE MARKETING AUTHORIZING HOLDER**

   AstraZeneca AB

3. **EXPIRY DATE**

   EXP

4. **BATCH NUMBER**

   Lot

5. **OTHER**
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B. PACKAGE LEAFLET
Read all of this leaflet carefully before you start taking this medicine because it contains important information for you.
- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This medicine has been prescribed for you only. Do not pass it on to others. It may harm them, even if their signs of illness are the same as yours.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet
1. What Brilique is and what it is used for
2. What you need to know before you take Brilique
3. How to take Brilique
4. Possible side effects
5. How to store Brilique
6. Contents of the pack and other information

1. What Brilique is and what it is used for

What Brilique is
Brilique contains an active substance called ticagrelor. This belongs to a group of medicines called antiplatelet medicines.

What Brilique is used for
Brilique in combination with acetylsalicylic acid (another antiplatelet agent) is to be used in adults only. You have been given Brilique because you have had:
- a heart attack, over a year ago.
It reduces the chances of you having another heart attack, stroke or dying from a disease related to your heart or blood vessels.

How Brilique works
Brilique affects cells called ‘platelets’ (also called thrombocytes). These very small blood cells help stop bleeding by clumping together to plug tiny holes in blood vessels that are cut or damaged.

However, platelets can also form clots inside diseased blood vessels in the heart and brain. This can be very dangerous because:
- the clot can cut off the blood supply completely; this can cause a heart attack (myocardial infarction) or stroke, or
- the clot can partly block the blood vessels to the heart; this reduces the blood flow to the heart and can cause chest pain which comes and goes (called ‘unstable angina’).

Brilique helps stop the clumping of platelets. This reduces the chance of a blood clot forming that can reduce blood flow.
2. **What you need to know before you take Brilique**

**Do not take Brilique if:**
- You are allergic to ticagrelor or any of the other ingredients of this medicine (listed in section 6).
- You are bleeding now.
- You have had a stroke caused by bleeding in the brain.
- You have severe liver disease.
- You are taking any of the following medicines:
  - ketoconazole (used to treat fungal infections)
  - clarithromycin (used to treat bacterial infections)
  - nefazodone (an antidepressant)
  - ritonavir and atazanavir (used to treat HIV infection and AIDS).

Do not take Brilique if any of the above applies to you. If you are not sure, talk to your doctor or pharmacist before taking this medicine.

**Warnings and precautions**

Talk to your doctor or pharmacist before taking Brilique if:
- You have an increased risk of bleeding because of:
  - a recent serious injury
  - recent surgery (including dental work, ask your dentist about this)
  - you have a condition that affects blood clotting
  - recent bleeding from your stomach or gut (such as a stomach ulcer or colon ‘polyps’)
- You are due to have surgery (including dental work) at any time while taking Brilique. This is because of the increased risk of bleeding. Your doctor may want you to stop taking this medicine 7 days prior to surgery.
- Your heart rate is abnormally low (usually lower than 60 beats per minute) and you do not already have in place a device that paces your heart (pacemaker).
- You have asthma or other lung problems or breathing difficulties.
- You have had any problems with your liver or have previously had any disease which may have affected your liver.
- You have had a blood test that showed more than the usual amount of uric acid.

If any of the above apply to you (or you are not sure), talk to your doctor or pharmacist before taking this medicine.

**Children and adolescents**

Brilique is not recommended for children and adolescents under 18 years.

**Other medicines and Brilique**

Please tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines. This is because Brilique can affect the way some medicines work and some medicines can have an effect on Brilique.

Tell your doctor or pharmacist if you are taking any of the following medicines:
- more than 40 mg daily of either simvastatin or lovastatin (medicines used to treat high cholesterol)
- rifampicin (an antibiotic)
- phenytoin, carbamazepine and phenobarbital (used to control seizures)
- digoxin (used to treat heart failure)
- cyclosporine (used to lessen your body’s defenses)
- quinidine and diltiazem (used to treat abnormal heart rhythms)
- beta blockers and verapamil (used to treat high blood pressure).
In particular, tell your doctor or pharmacist if you are taking any of the following medicines that increase your risk of bleeding:

- ‘oral anticoagulants’ often referred to as ‘blood thinners’ which include warfarin.
- Non-Steroidal Anti-Inflammatory Drugs (abbreviated as NSAIDs) often taken as painkillers such as ibuprofen and naproxen.
- Selective Serotonin Reuptake Inhibitors (abbreviated as SSRIs) taken as antidepressants such as paroxetine, sertraline and citalopram.
- other medicines such as ketoconazole (used to treat fungal infections), clarithromycin (used to treat bacterial infections), nefazodone (an antidepressant), ritonavir and atazanavir (used to treat HIV infection and AIDS), cisapride (used to treat heartburn), ergot alkaloids (used to treat migraines and headaches).

Also tell your doctor that because you are taking Brilique, you may have an increased risk of bleeding if your doctor gives you fibrinolytics, often called ‘clot dissolvers’, such as streptokinase or alteplase.

**Pregnancy and breast-feeding**

It is not recommended to use Brilique if you are pregnant or may become pregnant. Women should use appropriate contraceptive measures to avoid pregnancy while taking this medicine.

Talk to your doctor before taking Brilique if you are breast-feeding. Your doctor will discuss with you the benefits and risks of taking Brilique during this time.

If you are pregnant or breast-feeding, think you may be pregnant or are planning to have a baby, ask your doctor or pharmacist for advice before taking this medicine.

**Driving and using machines**

Brilique is not likely to affect your ability to drive or use machines. If you feel dizzy or confusion while taking this medicine, be careful while driving or using machines.

3. **How to take Brilique**

Always take this medicine exactly as your doctor has told you. Check with your doctor or pharmacist if you are not sure.

**How much to take**

- The usual dose is one tablet of 60 mg twice a day. Continue taking Brilique as long as your doctor tells you.
- Take this medicine around the same time every day (for example, one tablet in the morning and one in the evening).

**Taking Brilique with other medicines for blood clotting**

Your doctor will usually also tell you to take acetylsalicylic acid. This is a substance present in many medicines used to prevent blood clotting. Your doctor will tell you how much to take (usually between 75-150 mg daily).

**How to take Brilique**

- You can take the tablet with or without food.
- You can check when you last took a tablet of Brilique by looking on the blister. There is a sun (for the morning) and a moon (for the evening). This will tell you whether you have taken the dose.
If you have trouble swallowing the tablet
If you have trouble swallowing the tablet you can crush it and mix with water as follows:
- Crush the tablet to a fine powder.
- Pour the powder into half a glass of water.
- Stir and drink immediately.
- To make sure there is no medicine left, rinse the empty glass with another half a glass of water and drink it.

If you take more Brilique than you should
If you take more Brilique than you should, talk to a doctor or go to hospital straight away. Take the medicine pack with you. You may be at increased risk of bleeding.

If you forget to take Brilique
- If you forget to take a dose, just take your next dose as normal.
- Do not take a double dose (two doses at the same time) to make up for the forgotten dose.

If you stop taking Brilique
Do not stop taking Brilique without talking to your doctor. Take this medicine on a regular basis and for as long as your doctor keeps prescribing it. If you stop taking Brilique, it may increase your chances of having another heart attack or stroke or dying from a disease related to your heart or blood vessels.

If you have any further questions on the use of this medicine, ask your doctor or pharmacist.

4. Possible side effects
Like all medicines, this medicine can cause side effects, although not everybody gets them. The following side effects may happen with this medicine:

Brilique affects blood clotting, so most side effects are related to bleeding. Bleeding may occur in any part of the body. Some bleeding is common (like bruising and nosebleeds). Severe bleeding is uncommon, but can be life threatening.

See a doctor straight away if you notice any of the following – you may need urgent medical treatment:
- **Bleeding into the brain or inside the skull is an uncommon side effect, and may cause signs of a stroke such as:**
  - sudden numbness or weakness of your arm, leg or face, especially if only on one side of the body
  - sudden confusion, difficulty speaking or understanding others
  - sudden difficulty in walking or loss of balance or co-ordination
  - suddenly feeling dizzy or sudden severe headache with no known cause

- **Signs of bleeding such as:**
  - bleeding that is severe or that you cannot control
  - unexpected bleeding or bleeding that lasts a long time
  - pink, red, or brown urine
  - vomiting red blood or your vomit looks like ‘coffee grounds’
  - red or black stools (look like tar)
  - coughing up or vomiting blood clots

- **Fainting (syncope)**
- a temporary loss of consciousness due to sudden drop in blood flow to the brain (common)

Discuss with your doctor if you notice any of the following:
- **Feeling short of breath - this is very common.** It might be due to your heart disease or another cause, or it might be a side effect of Brilique. Brilique-related breathlessness is generally mild and characterised as a sudden, unexpected hunger for air usually occurring at rest and may appear in the first weeks of therapy and for many may disappear. If your feeling of shortness of breath gets worse or lasts a long time, tell your doctor. Your doctor will decide if it needs treatment or further investigations.

Other possible side effects

**Very common (may affect more than 1 in 10 people)**
- High level of uric acid in your blood (as seen in tests)
- Bleeding caused by blood disorders

**Common (may affect up to 1 in 10 people)**
- Bruising
- Headache
- Feeling dizzy or like the room is spinning
- Diarrhoea or indigestion
- Feeling sick (nausea)
- Constipation
- Rash
- Itching
- Severe pain and swelling in your joints – these are signs of gout
- Feeling dizzy or light-headed, or having blurred vision – these are signs of low blood pressure
- Nosebleed
- Bleeding after surgery or from cuts (for example while shaving) and wounds more than is normal
- Bleeding from your stomach lining (ulcer)
- Bleeding gums

**Uncommon (may affect up to 1 in 100 people)**
- Allergic reaction – a rash, itching, or a swollen face or swollen lips/tongue may be signs of an allergic reaction.
- Confusion
- Visual problems caused by blood in your eye
- Vaginal bleeding that is heavier, or happens at different times, than your normal period (menstrual) bleeding
- Bleeding into your joints and muscles causing painful swelling
- Blood in your ear
- Internal bleeding, this may cause dizziness or light-headedness

**Reporting of side effects**
If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via the national reporting system listed in Appendix V. By reporting side effects you can help provide more information on the safety of this medicine.
5. **How to store Brilique**

Keep this medicine out of the sight and reach of children.
Do not use this medicine after the expiry date which is stated on the blister and carton after EXP.
The expiry date refers to the last day of that month.
This medicine does not require any special storage conditions.
Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help to protect the environment.

6. **Contents of the pack and other information**

**What Brilique contains**
- The active substance is ticagrelor. Each film-coated tablet contains 60 mg of ticagrelor.
- The other ingredients are:
  - **Tablet core**: mannitol (E421), calcium hydrogen phosphate dihydrate, sodium starch glycolate type A, hydroxypropylcellulose (E463), magnesium stearate (E470b).
  - **Tablet film-coating**: hypromellose (E464), titanium dioxide (E171), macrogol 400, iron oxide black (E172) and iron oxide red (E172).

**What Brilique looks like and contents of the pack**
Film-coated tablet (tablet): The tablets are round, biconvex, pink, film-coated marked with a “60” above “T” on one side.

Brilique is available in:
- standard blisters (with sun/moon symbols) in cartons of 60 and 180 tablets
- calendar blisters (with sun/moon symbols) in cartons of 14, 56 and 168 tablets
Not all pack sizes may be marketed.

**Marketing Authorisation Holder and Manufacturer**

Marketing Authorisation Holder:
AstraZeneca AB
SE-151 85 Södertälje
Sweden

Manufacturer:
AstraZeneca AB
Gärtunavägen
SE-151 85 Södertälje
Sweden

Manufacturer:
AstraZeneca UK Limited
Silk Road Business Park
Macclesfield, Cheshire, SK10 2NA
United Kingdom
For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

**Belgïë/Belgique/Belgien**  
AstraZeneca S.A./N.V.  
Tel: +32 2 370 48 11

**България**  
АстраZeneca България ЕООД  
Tel.: +359 2 44 55 000

**Česká republika**  
AstraZeneca Czech Republic s.r.o  
Tel: +420 222 807 111

**Danmark**  
AstraZeneca A/S  
Tel: +45 43 66 64 62

**Deutschland**  
AstraZeneca GmbH  
Tel: +49 41 03 7080

**Ελλάδα**  
AstraZeneca A.E.  
Τηλ.: +30 2 106871500

**España**  
AstraZeneca Farmacéutica Spain, S.A.  
Tel: +34 91 301 91 00

**France**  
AstraZeneca  
Tél: +33 1 41 29 40 00

**Hrvatska**  
AstraZeneca d.o.o.  
Tel: +385 1 4628 000

**Ireland**  
AstraZeneca Pharmaceuticals (Ireland) Ltd  
Tel: +353 1609 7100

**Ísland**  
Vistor hf.  
Sími: +354 535 7000

**Italia**  
AstraZeneca S.p.A.

**Lietuva**  
UAB AstraZeneca Lietuva  
Tel: +370 5 2660550

**Luxembourg/Luxemburg**  
AstraZeneca S.A./N.V.  
Tel/Tel: +32 2 370 48 11

**Magyarország**  
AstraZeneca Kft.  
Tel.: +36 1 883 6500

**Malta**  
Associated Drug Co. Ltd  
Tel: +356 2277 8000

**Nederland**  
AstraZeneca BV  
Tel: +31 79 363 2222

**Norge**  
AstraZeneca AS  
Tel: +47 21 00 64 00

**Österreich**  
AstraZeneca Österreich GmbH  
Tel: +43 1 711 31 0

**Polska**  
AstraZeneca Pharma Poland Sp. z o.o.  
Tel.: +48 22 245 73 00

**Portugal**  
AstraZeneca Produtos Farmacêuticos, Lda.  
Tel: +351 21 434 61 00

**România**  
AstraZeneca Pharma SRL  
Tel: +40 21 317 60 41

**Slovenija**  
AstraZeneca UK Limited  
Tel: +386 1 51 35 600

**Slovenská republika**  
AstraZeneca AB, o.z.  
Tel: +421 2 5737 7777

**Suomi/Finland**  
AstraZeneca Oy
This leaflet was last revised in

**Other sources of information**
Read all of this leaflet carefully before you start taking this medicine because it contains important information for you.
- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor or pharmacist.
- This medicine has been prescribed for you only. Do not pass it on to others. It may harm them, even if their signs of illness are the same as yours.
- If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet
1. What Brilique is and what it is used for
2. What you need to know before you take Brilique
3. How to take Brilique
4. Possible side effects
5. How to store Brilique
6. Contents of the pack and other information

1. What Brilique is and what it is used for

What Brilique is
Brilique contains an active substance called ticagrelor. This belongs to a group of medicines called antiplatelet medicines.

What Brilique is used for
Brilique in combination with acetylsalicylic acid (another antiplatelet agent) is to be used in adults only. You have been given Brilique because you have had:
- a heart attack, or
- unstable angina (angina or chest pain that is not well controlled).
It reduces the chances of you having another heart attack, stroke or dying from a disease related to your heart or blood vessels.

How Brilique works
Brilique affects cells called ‘platelets’ (also called thrombocytes). These very small blood cells help stop bleeding by clumping together to plug tiny holes in blood vessels that are cut or damaged.

However, platelets can also form clots inside diseased blood vessels in the heart and brain. This can be very dangerous because:
- the clot can cut off the blood supply completely; this can cause a heart attack (myocardial infarction) or stroke, or
- the clot can partly block the blood vessels to the heart; this reduces the blood flow to the heart and can cause chest pain which comes and goes (called ‘unstable angina’).

Brilique helps stop the clumping of platelets. This reduces the chance of a blood clot forming that can reduce blood flow.
2. What you need to know before you take Brilique

Do not take Brilique if:

- You are allergic to ticagrelor or any of the other ingredients of this medicine (listed in section 6).
- You are bleeding now.
- You have had a stroke caused by bleeding in the brain.
- You have severe liver disease.
- You are taking any of the following medicines:
  - ketoconazole (used to treat fungal infections)
  - clarithromycin (used to treat bacterial infections)
  - nefazodone (an antidepressant)
  - ritonavir and atazanavir (used to treat HIV infection and AIDS).

Do not take Brilique if any of the above applies to you. If you are not sure, talk to your doctor or pharmacist before taking this medicine.

Warnings and precautions

Talk to your doctor or pharmacist before taking Brilique if:

- You have an increased risk of bleeding because of:
  - a recent serious injury
  - recent surgery (including dental work, ask your dentist about this)
  - you have a condition that affects blood clotting
  - recent bleeding from your stomach or gut (such as a stomach ulcer or colon ‘polyps’)
- You are due to have surgery (including dental work) at any time while taking Brilique. This is because of the increased risk of bleeding. Your doctor may want you to stop taking this medicine 7 days prior to surgery.
- Your heart rate is abnormally low (usually lower than 60 beats per minute) and you do not already have in place a device that paces your heart (pacemaker).
- You have asthma or other lung problems or breathing difficulties.
- You have had any problems with your liver or have previously had any disease which may have affected your liver.
- You have had a blood test that showed more than the usual amount of uric acid.

If any of the above apply to you (or you are not sure), talk to your doctor or pharmacist before taking this medicine.

Children and adolescents

Brilique is not recommended for children and adolescents under 18 years.

Other medicines and Brilique

Please tell your doctor or pharmacist if you are taking, have recently taken or might take any other medicines. This is because Brilique can affect the way some medicines work and some medicines can have an effect on Brilique.

Tell your doctor or pharmacist if you are taking any of the following medicines:

- more than 40 mg daily of either simvastatin or lovastatin (medicines used to treat high cholesterol)
- rifampicin (an antibiotic)
- phenytoin, carbamazepine and phenobarbital (used to control seizures)
- digoxin (used to treat heart failure)
- cyclosporine (used to lessen your body’s defenses)
- quinidine and diltiazem (used to treat abnormal heart rhythms)
- beta blockers and verapamil (used to treat high blood pressure).
In particular, tell your doctor or pharmacist if you are taking any of the following medicines that increase your risk of bleeding:

- ‘oral anticoagulants’ often referred to as ‘blood thinners’ which include warfarin.
- Non-Steroidal Anti-Inflammatory Drugs (abbreviated as NSAIDs) often taken as painkillers such as ibuprofen and naproxen.
- Selective Serotonin Reuptake Inhibitors (abbreviated as SSRIs) taken as antidepressants such as paroxetine, sertraline and citalopram.
- other medicines such as ketoconazole (used to treat fungal infections), clarithromycin (used to treat bacterial infections), nefazodone (an antidepressant), ritonavir and atazanavir (used to treat HIV infection and AIDS), cisapride (used to treat heartburn), ergot alkaloids (used to treat migraines and headaches).

Also tell your doctor that because you are taking Brilique, you may have an increased risk of bleeding if your doctor gives you fibrinolytics, often called ‘clot dissolvers’, such as streptokinase or alteplase.

**Pregnancy and breast-feeding**

It is not recommended to use Brilique if you are pregnant or may become pregnant. Women should use appropriate contraceptive measures to avoid pregnancy while taking this medicine.

Talk to your doctor before taking Brilique if you are breast-feeding. Your doctor will discuss with you the benefits and risks of taking Brilique during this time.

If you are pregnant or breast-feeding, think you may be pregnant or are planning to have a baby, ask your doctor or pharmacist for advice before taking this medicine.

**Driving and using machines**

Brilique is not likely to affect your ability to drive or use machines. If you feel dizzy or confusion while taking this medicine, be careful while driving or using machines.

3. **How to take Brilique**

Always take this medicine exactly as your doctor has told you. Check with your doctor or pharmacist if you are not sure.

**How much to take**

- The starting dose is two tablets at the same time (loading dose of 180 mg). This dose will usually be given to you in the hospital.
- After this starting dose, the usual dose is one tablet of 90 mg twice a day for up to 12 months unless your doctor tells you differently.
- Take this medicine around the same time every day (for example, one tablet in the morning and one in the evening).

**Taking Brilique with other medicines for blood clotting**

Your doctor will usually also tell you to take acetylsalicylic acid. This is a substance present in many medicines used to prevent blood clotting. Your doctor will tell you how much to take (usually between 75-150 mg daily).

**How to take Brilique**

- You can take the tablet with or without food.
• You can check when you last took a tablet of Brilique by looking on the blister. There is a sun (for the morning) and a moon (for the evening). This will tell you whether you have taken the dose.

If you have trouble swallowing the tablet
If you have trouble swallowing the tablet you can crush it and mix with water as follows:
• Crush the tablet to a fine powder.
• Pour the powder into half a glass of water.
• Stir and drink immediately.
• To make sure there is no medicine left, rinse the empty glass with another half a glass of water and drink it.

If you take more Brilique than you should
If you take more Brilique than you should, talk to a doctor or go to hospital straight away. Take the medicine pack with you. You may be at increased risk of bleeding.

If you forget to take Brilique
• If you forget to take a dose, just take your next dose as normal.
• Do not take a double dose (two doses at the same time) to make up for the forgotten dose.

If you stop taking Brilique
Do not stop taking Brilique without talking to your doctor. Take this medicine on a regular basis and for as long as your doctor keeps prescribing it. If you stop taking Brilique, it may increase your chances of having another heart attack or stroke or dying from a disease related to your heart or blood vessels.

If you have any further questions on the use of this medicine, ask your doctor or pharmacist.

4. Possible side effects

Like all medicines, this medicine can cause side effects, although not everybody gets them. The following side effects may happen with this medicine:

Brilique affects blood clotting, so most side effects are related to bleeding. Bleeding may occur in any part of the body. Some bleeding is common (like bruising and nosebleeds). Severe bleeding is uncommon, but can be life threatening.

See a doctor straight away if you notice any of the following – you may need urgent medical treatment:
• Bleeding into the brain or inside the skull is an uncommon side effect, and may cause signs of a stroke such as:
  - sudden numbness or weakness of your arm, leg or face, especially if only on one side of the body
  - sudden confusion, difficulty speaking or understanding others
  - sudden difficulty in walking or loss of balance or co-ordination
  - suddenly feeling dizzy or sudden severe headache with no known cause

• Signs of bleeding such as:
  - bleeding that is severe or that you cannot control
  - unexpected bleeding or bleeding that lasts a long time
  - pink, red, or brown urine
  - vomiting red blood or your vomit looks like ‘coffee grounds’
- red or black stools (look like tar)
- coughing up or vomiting blood clots

- **Fainting (syncope)**
  - a temporary loss of consciousness due to sudden drop in blood flow to the brain (common)

**Discuss with your doctor if you notice any of the following:**
- **Feeling short of breath - this is very common.** It might be due to your heart disease or another cause, or it might be a side effect of Brilique. Brilique-related breathlessness is generally mild and characterised as a sudden, unexpected hunger for air usually occurring at rest and may appear in the first weeks of therapy and for many may disappear. If your feeling of shortness of breath gets worse or lasts a long time, tell your doctor. Your doctor will decide if it needs treatment or further investigations.

**Other possible side effects**

**Very common (may affect more than 1 in 10 people)**
- High level of uric acid in your blood (as seen in tests)
- Bleeding caused by blood disorders

**Common (may affect up to 1 in 10 people)**
- Bruising
- Headache
- Feeling dizzy or like the room is spinning
- Diarrhoea or indigestion
- Feeling sick (nausea)
- Constipation
- Rash
- Itching
- Severe pain and swelling in your joints – these are signs of gout
- Feeling dizzy or light-headed, or having blurred vision – these are signs of low blood pressure
- Nosebleed
- Bleeding after surgery or from cuts (for example while shaving) and wounds more than is normal
- Bleeding from your stomach lining (ulcer)
- Bleeding gums

**Uncommon (may affect up to 1 in 100 people)**
- Allergic reaction – a rash, itching, or a swollen face or swollen lips/tongue may be signs of an allergic reaction.
- Confusion
- Visual problems caused by blood in your eye
- Vaginal bleeding that is heavier, or happens at different times, than your normal period (menstrual) bleeding
- Bleeding into your joints and muscles causing painful swelling
- Blood in your ear
- Internal bleeding, this may cause dizziness or light-headedness

**Reporting of side effects**
If you get any side effects, talk to your doctor or pharmacist. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via the national reporting system listed in Appendix V. By reporting side effects you can help provide more information on the safety of this medicine.

5. **How to store Brilique**

Keep this medicine out of the sight and reach of children.
Do not use this medicine after the expiry date which is stated on the blister and carton after EXP. The expiry date refers to the last day of that month.
This medicine does not require any special storage conditions.
Do not throw away any medicines via wastewater or household waste. Ask your pharmacist how to throw away medicines you no longer use. These measures will help to protect the environment.

6. **Contents of the pack and other information**

**What Brilique contains**
- The active substance is ticagrelor. Each film-coated tablet contains 90 mg of ticagrelor.
- The other ingredients are:
  
  **Tablet core:** mannitol (E421), calcium hydrogen phosphate dihydrate, sodium starch glycolate type A, hydroxypropylcellulose (E463), magnesium stearate (E470b).

  
  **Tablet film-coating:** hypromellose (E464), titanium dioxide (E171), talc, macrogol 400, and iron oxide yellow (E172).

**What Brilique looks like and contents of the pack**
Film-coated tablet (tablet): The tablets are round, biconvex, yellow, film-coated marked with a “90” above “T” on one side.

Brilique is available in:
- standard blisters (with sun/moon symbols) in cartons of 60 and 180 tablets
- calendar blisters (with sun/moon symbols) in cartons of 14, 56 and 168 tablets
- perforated unit-dosed blisters in a carton of 100x1 tablets

Not all pack sizes may be marketed.

**Marketing Authorisation Holder and Manufacturer**

Marketing Authorisation Holder: AstraZeneca AB
SE-151 85 Södertälje
Sweden

Manufacturer: AstraZeneca AB
Gärtsnavägen
SE-151 85 Södertälje
Sweden

Manufacturer: AstraZeneca UK Limited
For any information about this medicine, please contact the local representative of the Marketing Authorisation Holder:

**België/Belgique/Belgien**
AstraZeneca S.A./N.V.
Tel: +32 2 370 48 11

**България**
АстраЗенека България ЕООД
Tel.: +359 2 44 55 000

**Česká republika**
AstraZeneca Czech Republic s.r.o
Tel: +420 222 807 111

**Danmark**
AstraZeneca A/S
Tel: +45 43 66 64 62

**Deutschland**
AstraZeneca GmbH
Tel: +49 41 03 7080

**Eesti**
AstraZeneca
Tel: +372 6549 600

**Ελλάδα**
AstraZeneca A.E.
Tel: +30 2 106871500

**España**
AstraZeneca Farmacéutica Spain, S.A.
Tel: +34 91 301 91 00

**France**
AstraZeneca
Tél: +33 1 41 29 40 00

**Hrvatska**
AstraZeneca d.o.o.
Tel: +385 1 4628 000

**Ireland**
AstraZeneca Pharmaceuticals (Ireland) Ltd
Tel: +353 1609 7100

**Ísland**

**Ísland**

**Luxembourg/Luxemburg**
AstraZeneca S.A./N.V.
Tél/Tel: +32 2 370 48 11

**Magyarország**
AstraZeneca Kft.
Tel.: +36 1 883 6500

**Malta**
Associated Drug Co. Ltd
Tel: +356 2277 8000

**Nederland**
AstraZeneca BV
Tel: +31 79 363 2222

**Norge**
AstraZeneca AS
Tel: +47 21 00 64 00

**Österreich**
AstraZeneca Österreich GmbH
Tel: +43 1 711 31 0

**Polska**
AstraZeneca Pharma Poland Sp. z o.o.
Tel.: +48 22 245 73 00

**Portugal**
AstraZeneca Produtos Farmacêuticos, Lda.
Tel: +351 21 434 61 00

**România**
AstraZeneca Pharma SRL
Tel: +40 21 317 60 41

**Slovenija**
AstraZeneca UK Limited
Tel: +386 1 51 35 600

**Slovenská republika**
This leaflet was last revised in

Other sources of information

Detailed information on this medicine is available on the European Medicines Agency web site: http://www.ema.europa.eu.