PRODUCT INFORMATION
CLOPIDOGREL WINTHROP PLUS ASPIRIN® 75MG/75MG & CLOPIDOGREL WINTHROP PLUS ASPIRIN® 75MG/100MG

NAME OF THE MEDICINE
Non-proprietary Name
Clopidogrel Winthrop Plus Aspirin® 75mg/75mg containing clopidogrel 75 mg (as clopidogrel hydrogen sulfate) and aspirin 75 mg
Clopidogrel Winthrop Plus Aspirin® 75mg/100mg containing clopidogrel 75 mg (as clopidogrel hydrogen sulfate) and aspirin 100 mg

Chemical Structure
Clopidogrel
Clopidogrel hydrogen sulfate is designated chemically as methyl (+)-(S)-α-(2-chlorophenyl)-6,7-dihydrothieno[3,2-c] pyridine-5(4H)-acetate sulfate (1:1).
The empirical formula of clopidogrel hydrogen sulfate is C_{16}H_{16}ClNO_{2}S.H_{2}SO_{4} and its molecular weight is 419.9.
Clopidogrel hydrogen sulfate has the following chemical structure:

![Clopidogrel Chemical Structure](image)

CAS Number
120202-66-6 (Clopidogrel hydrogen sulfate),
113 665-84-2 (Clopidogrel base).

Aspirin
Aspirin (or acetylsalicylic acid) is designated chemically as 2-acetoxybenzoic acid and has the following chemical structure.

![Aspirin Chemical Structure](image)

The empirical formula is C_{9}H_{8}O_{4} and its molecular weight is 180.2.

CAS Number
50-78-2.
DESCRIPTION

Clopidogrel

Clopidogrel hydrogen sulfate is a white to off-white powder. It is practically insoluble in water at neutral pH but freely soluble at pH 1. It is freely soluble in methanol, sparingly soluble in methylene chloride and is practically insoluble in ethyl ether. It has a specific optical rotation of about +56°.

Aspirin

Aspirin is a white crystalline powder or colourless crystals, odourless or almost odourless, slightly soluble in water, freely soluble in alcohol, soluble in chloroform and in ether. It melts at about 135°C.

Clopidogrel Winthrop Plus Aspirin

Clopidogrel Winthrop Plus Aspirin tablets are film coated and for both strengths each tablet contains mannitol, macrogol 6000, cellulose – microcrystalline, castor oil – hydrogenated, hydroxypropylcellulose, maize starch, stearic acid, silica – colloidal anhydrous. The coating contains lactose, hypromellose, titanium dioxide, glycerol triacetate, a colourant and carnauba wax. The colourant is yellow iron oxide in Clopidogrel Winthrop Plus Aspirin 75mg/75mg and red iron oxide in Clopidogrel Winthrop Plus Aspirin 75mg/100mg.

PHARMACOLOGY

Pharmacodynamics

Clopidogrel

Clopidogrel is a specific and potent inhibitor of platelet aggregation. Platelets have an established role in the pathophysiology of atherosclerotic disease and thrombotic events. Long term use of anti-platelet drugs has shown consistent benefit in the prevention of ischaemic stroke, myocardial infarction and vascular death in patients at increased risk of such outcomes, including those with established atherosclerosis or a history of atherothrombosis.

Clopidogrel selectively inhibits the binding of adenosine diphosphate (ADP) to its platelet receptor, and the subsequent ADP-mediated activation of the GPIIb/IIIa complex, thereby inhibiting platelet aggregation. Biotransformation of clopidogrel is necessary to produce inhibition of platelet aggregation. The active metabolite, a thiol derivative, is formed by oxidation of clopidogrel to 2-oxoclopidogrel and subsequent hydrolysis. The active thiol metabolite, which has been isolated in vitro, binds rapidly and irreversibly to platelet ADP receptors, P2Y12, thus inhibiting platelet aggregation. Clopidogrel also inhibits platelet aggregation induced by other agonists by blocking the amplification of platelet activation by released ADP. Due to the irreversible binding, platelets exposed are affected for the remainder of their lifespan and recovery of normal platelet function occurs at a rate consistent with platelet turnover (approximately 7 days).

Statistically significant and dose-dependent inhibition of platelet aggregation was noted 2 hours after single oral doses of clopidogrel. Repeated doses of 75 mg per day produced substantial inhibition of ADP-induced platelet aggregation from the first day; this increased progressively and reached steady state between Day 3 and Day 7. At steady state, the average inhibition level observed with a dose of 75 mg per day was between 40% and 60%. Platelet aggregation and bleeding time gradually returned to baseline values, generally within 7 days after treatment was discontinued.

Aspirin

Aspirin inhibits platelet aggregation by irreversible inhibition of platelet cyclo-oxygenase and the production of thromboxane A2, an inducer of platelet aggregation and vasoconstriction. This effect lasts for the life of the platelet.
**Pharmacokinetics**

**Clopidogrel**

Clopidogrel After single and repeated oral doses of 75 mg per day, clopidogrel is rapidly absorbed. Clopidogrel mean peak plasma levels (approximately 2.2 - 2.5 ng/ml after a single 75 mg oral dose) occurred approximately 45 minutes after dosing. Absorption is at least 50%, based on urinary excretion of clopidogrel metabolites.

Clopidogrel is a prodrug which is extensively hydrolysed in the liver by HCE1 (human carboxylesterase1). In vitro and in vivo, clopidogrel is metabolised according to two main metabolic pathways:

- One mediated by esterases and leading to hydrolysis into its carboxylic acid derivative, which is inactive and is the main circulating metabolite (about 85% of the circulating compound in plasma). Mean peak plasma levels of this metabolite (approx. 3600 ng/ml after single 75 mg oral dose) occurred approximately 45 minutes after dosing. In vitro in the presence of ethyl alcohol, the rate of clopidogrel hydrolysis was decreased, and some of the clopidogrel was converted to ethyl clopidogrel.

- One mediated by multiple cytochromes P450 in the gastrointestinal tract and liver leading to the active metabolite(s) of clopidogrel, a thiol derivative, which is generated through formation of 2 oxo clopidogrel. This metabolic pathway is mediated by multiple Cytochrome P450 isoenzymes, i.e., CYP3A4, CYP3A5, CYP2C19, CYP1A2, CYP2C9, CYP2E1 and CYP2B6. The active thiol metabolite, which has been isolated in vitro, binds rapidly and irreversibly to platelet receptors, thus inhibiting platelet aggregation. Clinical studies have indicated that individuals with loss of function variants of CYP2C9 and CYP2C19 are more likely to have lower concentrations of the active metabolite and higher residual platelet activity; clopidogrel is therefore less likely to be efficacious in these poor metabolisers.

The kinetics of the main circulating metabolite were linear (plasma concentrations increased in proportion to dose) in the dose range of 50 to 150 mg of clopidogrel.

Clopidogrel and the main circulating metabolite bind reversibly in vitro to human plasma proteins (98% and 94% respectively). The binding is non-saturable in vitro over a wide concentration range.

Following an oral dose of $^{14}$C-labelled clopidogrel in man, approximately 50% was excreted in the urine and approximately 46% in the faeces in the 120 hour interval after dosing. After a single oral dose of 75 mg, clopidogrel has a half-life of approximately 6 hours. The elimination half-life of the main circulating metabolite was 8 hours after single and repeated administration. Plasma concentrations of the main circulating metabolite were significantly higher in elderly subjects (≥75 years) as compared to young healthy volunteers. However, these higher plasma levels were not associated with differences in platelet aggregation and bleeding time.

Plasma levels of the main circulating metabolite were lower in subjects with severe renal disease (creatinine clearance from 5 to 15 mL/min) compared to subjects with moderate renal disease (creatinine clearance from 30 to 60 mL/min) and healthy subjects, after repeated doses of 75 mg/day. Although inhibition of ADP-induced platelet aggregation was lower (25%) than that observed in healthy subjects, the prolongation of bleeding was similar to that seen in healthy subjects receiving 75 mg of clopidogrel per day.

**Aspirin**

**Absorption**

Following absorption, the aspirin in Clopidogrel Winthrop Plus Aspirin is hydrolysed to salicylic acid, with peak plasma levels of salicylic acid occurring within 1 hour of dosing, such that plasma levels of aspirin are essentially undetectable 1.5 to 4 hours after dosing. Administration of aspirin with meals did not significantly modify its bioavailability.
Distribution

Based on available data, aspirin is poorly bound to plasma proteins and its apparent volume of distribution is low (10 L). Its metabolite, salicylic acid, is highly bound to plasma proteins, but its binding is concentration dependent (nonlinear). At low concentrations (<100 μg/mL), approximately 90% of salicylic acid is bound to albumin. Salicylic acid is widely distributed to all tissues and fluids in the body, including the central nervous system, breast milk and foetal tissues.

Metabolism and Elimination

The aspirin in Clopidogrel Winthrop Plus Aspirin is rapidly hydrolysed by HCE2 (human carboxylesterase 2) in the intestine and the liver to salicylic acid, with a half-life of 0.3 to 0.4 hours for aspirin doses from 75 to 100 mg. This salicylic acid has a plasma half-life of approximately 2 hours. Salicylic acid is primarily conjugated in the liver to form salicylic acid, a phenolic glucuronide, an acyl glucuronide and a number of minor metabolites. Salicylate metabolism is saturable and total body clearance decreases at higher serum concentrations, due to the limited ability of the liver to form both salicylic acid and phenolic glucuronide. Following toxic aspirin doses (10 to 20 g), the plasma half-life may be increased to over 20 hours. At high aspirin doses, the elimination of salicylic acid follows zero-order kinetics (i.e. the rate of elimination is constant in relation to plasma concentration), with an apparent half-life of 6 hours or higher. Renal excretion of unchanged drug depends upon urinary pH. As urinary pH rises above 6.5, the renal clearance of free salicylate increases from <5% to >80%. Following therapeutic doses, approximately 10% is found excreted in the urine as salicylic acid, 75% as salicylic acid, 10% phenolic- and 5% acyl-glucuronides of salicylic acid.

Clopidogrel/Aspirin Bioequivalence

Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg and 75/100 mg were demonstrated to be bioequivalent to the clopidogrel 75 mg tablets in terms of clopidogrel C_max and AUC, and/or carboxylic acid metabolite. For aspirin, Clopidogrel Winthrop Plus Aspirin® 75 mg/75 mg and 75/100 mg were demonstrated to be bioequivalent to aspirin 75 mg and 100 mg, respectively, in terms of aspirin AUC, and salicylic acid C_max and AUC. In terms of C_max, aspirin was not bioequivalent with the C_max being 1.3- to 1.6-fold higher for Clopidogrel Winthrop Plus Aspirin than for the aspirin tablets. However, a slight difference in aspirin C_max is not considered to be clinically significant (See CLINICAL TRIALS).

Pharmacogenetics

CYP2C19 is involved in the formation of both the active metabolite and the 2-oxo-clopidogrel intermediate metabolite. Clopidogrel active metabolite pharmacokinetics and antiplatelet effects, as measured by ex vivo platelet aggregation assays, differ according to CYP2C19 genotype. The CYP2C19*1 allele corresponds to fully functional metabolism while the CYP2C19*2 and CYP2C19*3 alleles are nonfunctional. The CYP2C19*2 and CYP2C19*3 alleles account for the majority of reduced function alleles in Caucasian (85%) and Asian (99%) poor metabolisers. Other alleles associated with absent or reduced metabolism are less frequent and include CYP2C19*4, *5, *6, *7, and *8. A patient with poor metaboliser status will possess two loss-of-function alleles as defined above. Published frequencies for the poor CYP2C19 metaboliser genotypes are approximately 2% for Caucasians, 4% for Blacks and 14% for Chinese. Tests are available to determine a patient's CYP2C19 genotype.

A crossover study in 40 healthy subjects, 10 each in the four CYP2C19 metaboliser groups (ultrarapid, extensive, intermediate and poor), evaluated pharmacokinetic and antiplatelet responses using 300 mg followed by 75 mg/day and 600 mg followed by 150 mg/day, each for a total of 5 days (steady state). No substantial differences in active metabolite exposure and mean inhibition of platelet aggregation (IPA) were observed between ultrarapid, extensive and intermediate metabolisers. In poor metabolisers, active metabolite exposure was decreased by 63-71% compared to extensive metabolisers. After the 300 mg/75 mg dose regimen, antiplatelet responses were decreased in the poor metabolisers with mean IPA (5 μM ADP) of 24% (24 hours) and 37% (Day 5) as compared to IPA of 39% (24 hours) and 58% (Day 5) in the extensive metabolisers and 37% (24 hours) and 60% (Day 5) in the intermediate metabolisers.
When poor metabolisers received the 600 mg/150 mg regimen, active metabolite exposure was greater than with the 300 mg/75 mg regimen. In addition, IPA was 32% (24 hours) and 61% (Day 5), which were greater than in the poor metabolisers receiving the 300 mg/75 mg regimen, and were similar to the other CYP2C19 metaboliser groups receiving the 300 mg/75 mg regimen. An appropriate dose regimen for this patient population has not been established in clinical outcome trials.

Table 1: Active Metabolite Pharmacokinetics and Antiplatelet Responses by CYP2C19 Metaboliser Status

<table>
<thead>
<tr>
<th>DOSE</th>
<th>ULTRARAPID (N=10)</th>
<th>EXTENSIVE (N=10)</th>
<th>INTERMEDIATE (N=10)</th>
<th>POOR (N=10)</th>
</tr>
</thead>
<tbody>
<tr>
<td>AUC_{last} (ng.h/mL)</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>300 mg (Day 1)</td>
<td>33 (11)</td>
<td>39 (24)</td>
<td>31 (14)</td>
<td>14 (6)</td>
</tr>
<tr>
<td>600 mg (Day 1)</td>
<td>56 (22)</td>
<td>70 (46)</td>
<td>56 (27)</td>
<td>23 (7)</td>
</tr>
<tr>
<td>75 mg (Day 5)</td>
<td>11 (5)</td>
<td>12 (6)</td>
<td>9.9 (4)</td>
<td>3.2 (1)</td>
</tr>
<tr>
<td>150 mg (Day 5)</td>
<td>18 (8)</td>
<td>19 (8)</td>
<td>16 (7)</td>
<td>7 (2)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>IPA (%)\textsuperscript{a}</th>
<th>300 mg (24 h)</th>
<th>40 (21)</th>
<th>39 (28)</th>
<th>37 (21)</th>
<th>24 (26)</th>
</tr>
</thead>
<tbody>
<tr>
<td>600 mg (24 h)</td>
<td>51 (28)</td>
<td>49 (23)</td>
<td>56 (22)</td>
<td>32 (25)</td>
<td></td>
</tr>
<tr>
<td>75 mg (Day 5)</td>
<td>56 (13)</td>
<td>58 (19)</td>
<td>60 (18)</td>
<td>37 (23)</td>
<td></td>
</tr>
<tr>
<td>150 mg (Day 5)</td>
<td>68 (18)</td>
<td>73 (9)</td>
<td>74 (14)</td>
<td>61 (14)</td>
<td></td>
</tr>
</tbody>
</table>

Values are mean (SD)
\textsuperscript{a} Inhibition of platelet aggregation with 5μM ADP; larger value indicates greater platelet inhibition

Consistent with the above results, in a meta-analysis including 6 studies of 335 clopidogrel-treated subjects at steady state, it was shown that active metabolite exposure was decreased by 28% for intermediate metabolisers, and 72% for poor metabolisers while platelet aggregation inhibition (5 μM ADP) was decreased with differences in IPA of 5.9% and 21.4%, respectively, when compared to extensive metabolisers.

The influence of CYP2C19 genotype on clinical outcomes in patients treated with clopidogrel has not been evaluated in prospective, randomised, controlled trials. There have been a number of retrospective analyses, however, to evaluate this effect in patients treated with clopidogrel for whom there are genotyping results: CURE (n=2721), CHARISMA (n=2428), CLARITY-TIMI 28 (n=227), and TRITON-TIMI 38 (n=1477) and ACTIVE-A (n=601), as well as a number of published cohort studies.

In TRITON-TIMI 38 and 3 of the cohort studies (Collet, Sibbing, Giusti) the combined group of patients with either intermediate or poor metaboliser status had a higher rate of cardiovascular events (death, myocardial infarction, and stroke) or stent thrombosis compared to extensive metabolisers.

In CHARISMA and one cohort study (Simon), an increased event rate was observed only in poor metabolisers when compared to extensive metabolisers.

In CURE, CLARITY, ACTIVE-A and one of the cohort studies (Trenk), no increased event rate was observed based on metaboliser status.

None of these analyses were adequately sized to detect differences in outcome in poor metabolisers.

Special Populations

Geriatric Patients

Plasma concentrations of the main circulating metabolite of clopidogrel are significantly higher in the elderly (≥ 75 years) compared to young healthy volunteers but these higher plasma levels were not associated with differences in platelet aggregation and bleeding time. No dosage adjustment is needed for the elderly.
**Renally Impaired Patients**

Clopidogrel Winthrop Plus Aspirin is contraindicated in severe renal impairment. After repeated doses of 75 mg clopidogrel per day, plasma levels of the main circulating metabolite were lower in patients with severe renal impairment (creatinine clearance from 5 to 15 mL/min) compared to subjects with moderate renal impairment (creatinine clearance 30 to 60 mL/min) or healthy subjects. Although inhibition of ADP-induced platelet aggregation was lower (25%) than that observed in healthy volunteers, the prolongation of bleeding time was similar in healthy volunteers receiving 75 mg of clopidogrel per day. Experience with clopidogrel plus aspirin is limited in patients with mild to moderate renal impairment. Therefore Clopidogrel Winthrop Plus Aspirin should be used with caution in this population. (see ‘PRECAUTIONS’).

**Patients with Hepatic Impairment**

Clopidogrel Winthrop Plus Aspirin is contraindicated in severe hepatic impairment. Experience is limited in patients with moderate hepatic disease who may have bleeding diatheses. Clopidogrel Winthrop Plus Aspirin should therefore be used with caution in this population (see ‘PRECAUTIONS’).

**CYP2C9 AND CYP2C19 poor metabolisers**

Clinical studies have indicated that individuals with loss of function variants of CYP2C9 and CYP2C19 are more likely to have lower concentrations of the active metabolite and higher residual platelet activity; clopidogrel is therefore less likely to be efficacious in these poor metabolisers.

**Gender**

No significant difference was observed in the plasma levels of the main circulating metabolite of clopidogrel between males and females. In a small study comparing men and women, less inhibition of ADP-induced platelet aggregation was observed in women, but there was no difference in prolongation of bleeding time. In the large, controlled clinical study (Clopidogrel vs. Aspirin in Patients at Risk of Ischemic Events; CAPRIE), the incidence of clinical outcome events, other adverse clinical events, and abnormal clinical laboratory parameters was similar in men and women.

**Ethnicity**

The prevalence of CYP2C19 alleles that result in intermediate and poor CYP2C19 metabolism differs according to race/ethnicity (see PHARMACOLOGY, Pharmacogenetics). From literature, limited data in Asian populations are available to assess the clinical implication of genotyping of this CYP on clinical outcome events.

**CLINICAL TRIALS**

The safety and efficacy of clopidogrel and aspirin has been evaluated in patients in three double-blind studies: the CURE, CLARITY, and COMMIT studies, which compared clopidogrel to placebo, both given in combination with aspirin and other standard therapy.

The CURE study included 12,562 patients with acute coronary syndrome (unstable angina or non-ST-elevation myocardial infarction), and presenting within 24 hours of onset of the most recent episode of chest pain or symptoms consistent with ischaemia. Patients were required to have either ECG changes compatible with new ischaemia or elevated cardiac enzymes or troponin I or T to at least twice the upper limit of normal. Patients were randomised to clopidogrel (300 mg loading dose followed by 75 mg/day, n=6244) or placebo (n=6287), both given in combination with aspirin (75-325 mg once daily) and other standard therapies (oral anti-coagulants and long term NSAIDs were not permitted). Patients were treated for up to one year. The number of patients experiencing the primary endpoint [cardiovascular (CV) death, myocardial infarction (MI), or stroke] was 582 (9.3%) in the clopidogrel-treated group and 719 (11.4%) in the placebo-treated group, a 20% relative risk reduction (95% CI of 10%-28%; p = 0.00009) for the clopidogrel-treated group. The benefits of clopidogrel were seen within a few hours and maintained throughout the course of the study (up to 12 months). The primary outcome was reduced to a similar extent within the first 30 days (relative risk reduction of 22%).
from 30 days to one year (relative risk reduction of 19%), and for the entire one year study
(relative risk reduction of 20%).

The number of patients experiencing the co-primary endpoint (CV death, MI, stroke or refractory
ischaemia) was 1035 (16.5%) in the clopidogrel-treated group and 1187 (18.8%) in the placebo-
treated group, a 14% relative risk reduction (95% CI of 6%-21%, p = 0.0005) for the clopidogrel-
treated group, a benefit which was consistent for each component, indicating that clopidogrel
reduced a range of atherothrombotic events.

In the course of the study, patients who underwent cardiac revascularisation (surgical or
percutaneous coronary intervention with or without coronary stent implantation), received similar
benefit from clopidogrel + aspirin (including standard therapies) as those who did not have a
cardiac revascularisation.

The results obtained in populations with different characteristics (e.g. unstable angina or non-
ST-elevation MI, low to high risk levels, diabetes, need for revascularisation, age, gender, etc.)
were consistent with the results of the primary analysis.

The benefits observed with clopidogrel were independent of other acute and long-term
cardiovascular therapies (such as heparin/LMWH, GPIIb/IIIa antagonists, lipid lowering drugs,
beta blockers, and ACE-inhibitors). The efficacy of clopidogrel was observed independently of
the dose of aspirin (75-325 mg once daily).

In patients with ST-segment elevation acute myocardial infarction, safety and efficacy of
clopidogrel have been evaluated in two randomised, placebo-controlled, double-blind studies,
CLARITY and COMMIT.

The randomised, double-blind, placebo-controlled CLARITY trial included 3,491 patients
presenting within 12 hours of the onset of a ST elevation myocardial infarction and planned for
thrombolytic therapy. Patients were randomised to receive either clopidogrel (300 mg loading
dose, followed by 75 mg/day; n = 1752) or placebo (n = 1739), together with aspirin (150 to 325
mg loading dose followed by 75 to 162 mg/day), a fibrinolytic agent and, when appropriate,
heparin for 48 hours. The patients were followed for 30 days.

The primary endpoint was the occurrence of the composite of an occluded infarct-related artery
(defined as TIMI Flow Grade 0 or 1) on the predischarge angiogram, or death or recurrent
myocardial infarction by the time of the start of coronary angiography. For patients who did not
undergo angiography, the primary endpoint was death or recurrent myocardial infarction by Day
8 or by hospital discharge, if prior to Day 8.

The patient population was mostly Caucasian (89.5%) and included 19.7% women and 29.2%
were 65 years or over. A total of 99.7% of patients received fibrinolytics (fibrin specific: 68.7%,
non-fibrin specific: 31.1%), 89.5% heparin, 78.7% beta-blockers, 54.7% ACE inhibitors and 63%
statins.

The number of patients who reached the primary endpoint was 262 (15.0%) in the clopidogrel-
treated group and 377 (21.7%) in the placebo group, representing an absolute reduction of
6.7% and a 36% reduction in the odds of the endpoint in favour of treatment with clopidogrel
(95% CI: 0.53, 0.76; p<0.001), as shown in Table 2, mainly related to a reduction in occluded
infarct-related arteries.

The benefit of clopidogrel on the primary endpoint was consistent across all prespecified
subgroups, including patients’ age, gender, infarct location and type of fibrinolytic or heparin
used.
Table 2: Event Rates for the Primary Composite Endpoint in the CLARITY Study

<table>
<thead>
<tr>
<th>Event</th>
<th>Clopidogrel + aspirin N = 1753</th>
<th>Placebo + aspirin N = 1739</th>
<th>OR</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Number (%) of patients reporting the composite endpoint</td>
<td>262 (15.0%)</td>
<td>377 (21.7%)</td>
<td>0.64</td>
<td>0.53, 0.76</td>
</tr>
<tr>
<td>Occluded IRA</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N (subjects undergoing angiography)</td>
<td>1640</td>
<td>1634</td>
<td>0.59</td>
<td>0.48, 0.72</td>
</tr>
<tr>
<td>n (%) patients reporting endpoint</td>
<td>192 (11.7%)</td>
<td>301 (18.4%)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Death</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n (%) patients reporting endpoint</td>
<td>45 (2.6%)</td>
<td>38 (2.2%)</td>
<td>1.18</td>
<td>0.76, 1.83</td>
</tr>
<tr>
<td>Recurrent MI</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>n (%) patients reporting endpoint</td>
<td>44 (2.5%)</td>
<td>62 (3.6%)</td>
<td>0.69</td>
<td>0.47, 1.02</td>
</tr>
</tbody>
</table>

The total number of patients with a component event (occluded IRA, death or recurrent MI) is greater than the number of patients with a composite event because some patients had more than a single type of component event.

The randomised, double-blind, placebo-controlled, 2x2 factorial design COMMIT trial included 45,852 patients presenting within 24 hours of the onset of the symptoms of suspected myocardial infarction with supporting ECG abnormalities (i.e. ST elevation, ST depression or left bundle-branch block). Patients were randomised to receive clopidogrel (75 mg/day) or placebo, in combination with aspirin (162 mg/day), for 28 days or until hospital discharge, whichever came first.

The co-primary endpoints were death from any cause and the first occurrence of re-infarction, stroke or death. The patient population included 27.8% women, 58.4% 60 years or over (26% 70 years or over) and 54.5% patients who received fibrinolytics, 68% who received ACE-inhibitors and 10.9% who received non-trial beta-blockers (as well as half of the patients who received metoprolol as study medication).

As shown in the Table 3 and Figures 1 and 2 below, clopidogrel significantly reduced the relative risk of death from any cause by 7% (p = 0.029) and the relative risk of the combination of re-infarction, stroke or death by 9% (p = 0.002), representing an absolute risk reduction of 5 and 9 patients per 1000 treated (0.5 and 0.9%), respectively.

Table 3: Outcome Events in the COMMIT Analysis

<table>
<thead>
<tr>
<th>Event</th>
<th>Clopidogrel + aspirin n = 22961</th>
<th>Placebo + aspirin n = 22891</th>
<th>Odds ratio (95% CI)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Composite endpoint:</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Death, MI or Stroke</td>
<td>2121 (9.2%)</td>
<td>2310 (10.1%)</td>
<td>0.91 (0.86, 0.97)</td>
<td>0.002</td>
</tr>
<tr>
<td>Death</td>
<td>1726 (7.5%)</td>
<td>1845 (8.1%)</td>
<td>0.93 (0.87, 0.99)</td>
<td>0.029</td>
</tr>
<tr>
<td>Non-fatal MI</td>
<td>270 (1.2%)</td>
<td>330 (1.4%)</td>
<td>0.81 (0.69, 0.95)</td>
<td>0.011</td>
</tr>
<tr>
<td>Non-fatal Stroke</td>
<td>127 (0.6%)</td>
<td>142 (0.6%)</td>
<td>0.89 (0.70, 1.13)</td>
<td>0.33</td>
</tr>
</tbody>
</table>

Note: 9 patients (2 clopidogrel and 7 placebo) suffered from both a non-fatal stroke and a non-fatal MI, hence the apparent disparity between composite endpoint and the sum of death, non-fatal MI and non-fatal stroke. Values for non-fatal MI and non-fatal stroke exclude patients who died of any cause.
The benefit associated with clopidogrel on the combined endpoint was consistent across age, gender and with or without fibrinolytics and was observed as early as 24 hours.

The bioequivalence of Clopidogrel Winthrop Plus Aspirin to reference clopidogrel and aspirin tablets has been demonstrated in three open-label, randomized, single-dose, 2-sequence, 2-period, 2-treatment crossover studies. One study was performed with Clopidogrel Winthrop Plus...
Aspirin 75 mg/75 mg (BDR4659) and two with Clopidogrel Winthrop Plus Aspirin 75 mg/100 mg (BDR5000 and BEQ10600). Study BEQ10600 (Clopidogrel Winthrop Plus Aspirin 75 mg/100 mg) evaluated bioequivalence in 40 young healthy subjects based on clopidogrel and its inactive carboxylic acid metabolite, and aspirin and salicylic acid. Studies BDR4659 (Clopidogrel Winthrop Plus Aspirin \textsuperscript{®} 75 mg/75 mg) and BDR5000 (Clopidogrel Winthrop Plus Aspirin 75 mg/100 mg) evaluated bioequivalence in 121 young healthy subjects based on clopidogrel inactive carboxylic acid metabolite, and aspirin and salicylic acid. 

Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg and 75/100 mg were demonstrated to be bioequivalent to the clopidogrel 75 mg tablets in terms of clopidogrel \( C_{\text{max}} \) and AUC, and/or carboxylic acid metabolite. For aspirin, Clopidogrel Winthrop Plus Aspirin \textsuperscript{®} 75 mg/75 mg and 75/100 mg were demonstrated to be bioequivalent to aspirin 75 mg and 100 mg, respectively, in terms of aspirin AUC, and salicylic acid \( C_{\text{max}} \) and AUC. The 90% CIs for these parameters were entirely within the bioequivalence interval [0.80-1.25].

In terms of \( C_{\text{max}} \), aspirin was not bioequivalent in the 3 studies, with the \( C_{\text{max}} \) being 1.3- to 1.6-fold higher for Clopidogrel Winthrop Plus Aspirin than for the aspirin tablets. However, considering the large number of aspirin formulations on the market and the clinical studies evaluating the benefit/risk of clopidogrel in combination with ASA (see above), a slight difference in ASA \( C_{\text{max}} \) is not considered to be clinically significant.

Table 4: Mean (coefficient of variation %) exposure of clopidogrel and its inactive carboxylic acid metabolite after a single oral dose of Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg or 75 mg/100 mg and Plavix 75 mg

<table>
<thead>
<tr>
<th>Compound</th>
<th>PK parameter</th>
<th>Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg</th>
<th>Clopidogrel Winthrop Plus Aspirin 75 mg/100 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>BDR4659</td>
<td>BDR5000</td>
</tr>
<tr>
<td>Clopidogrel</td>
<td>( C_{\text{max}} ) (ng/mL)</td>
<td>Not assessed</td>
<td>Not assessed</td>
</tr>
<tr>
<td></td>
<td>AUC (ng*h/mL)</td>
<td>Not assessed</td>
<td>Not assessed</td>
</tr>
<tr>
<td>Carboxylic acid metabolite</td>
<td>( C_{\text{max}} ) (ng/mL)</td>
<td>3319 (26)</td>
<td>3042 (26)</td>
</tr>
<tr>
<td></td>
<td>AUC (ng*h/mL)</td>
<td>9215 (23)</td>
<td>8069 (27)</td>
</tr>
</tbody>
</table>

Table 5: Mean (coefficient of variation %) exposure of aspirin and salicylic acid after a single oral dose of Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg or 75 mg/100 mg and aspirin 75 mg or 100 mg

<table>
<thead>
<tr>
<th>Compound</th>
<th>PK parameter</th>
<th>Clopidogrel Winthrop Plus Aspirin 75 mg/75 mg</th>
<th>Clopidogrel Winthrop Plus Aspirin 75 mg/100 mg</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>BDR4659</td>
<td>BDR5000</td>
</tr>
<tr>
<td>Aspirin</td>
<td>( C_{\text{max}} ) (ng/mL)</td>
<td>1207 (23)</td>
<td>1482 (25)</td>
</tr>
<tr>
<td></td>
<td>AUC (ng*h/mL)</td>
<td>936 (17)</td>
<td>1131 (22)</td>
</tr>
<tr>
<td>Salicylic acid</td>
<td>( C_{\text{max}} ) (ng/mL)</td>
<td>3533 (16)</td>
<td>4878 (16)</td>
</tr>
<tr>
<td></td>
<td>AUC (ng*h/mL)</td>
<td>12217 (21)</td>
<td>17791 (19)</td>
</tr>
</tbody>
</table>

\( n=39; \) \( n=110; \) \( n=111 \)

\( n=39; \) \( n=37; \) \( n=116; \) \( n=111 \)
INDICATIONS
Clopidogrel Winthrop Plus Aspirin is a fixed-dose combination product. Clopidogrel Winthrop Plus Aspirin is intended as continuation of therapy in patients with acute coronary syndrome already initiated with separate clopidogrel and aspirin products:

- Unstable angina or non-ST-elevation myocardial infarction in order to prevent early and long-term atherothrombotic events (myocardial infarction, stroke, vascular death or refractory ischaemia). Clopidogrel Winthrop Plus Aspirin is indicated for the treatment of acute coronary syndrome whether or not patients undergo cardiac revascularisation (surgical or PCI, with or without stent).
- ST-segment elevation acute myocardial infarction in order to prevent atherothrombotic events. In this population, Clopidogrel Winthrop Plus Aspirin has been shown to reduce the rate of death from any cause and the rate of a combined endpoint of death, re-infarction or stroke in medically treated patients eligible for thrombolytic therapy.

CONTRAINDICATIONS
Due to the presence of both components of the product, Clopidogrel Winthrop Plus Aspirin is contraindicated in case of:

- Hypersensitivity to clopidogrel, salicylates or any of the excipients.
- Severe liver impairment.
- Active pathological bleeding such as haemophilia, intracranial haemorrhage or gastrointestinal bleeding.
- Peptic ulcer or erosive gastritis
- Breast-feeding (see ‘PRECAUTIONS’-Use in Lactation).

In addition, due to the presence of aspirin, its use is also contraindicated in case of:

- Known allergy to Non-Steroidal Anti-Inflammatory Drugs (NSAIDs) and in patients with the syndrome of asthma with rhinitis and/or nasal polyps.
- Severe renal impairment.
- Third trimester of pregnancy (see Pregnancy).

PRECAUTIONS
General
Clopidogrel and aspirin prolong bleeding time, and should be used with caution in patients who may be at risk of increased bleeding from trauma, surgery or other pathological conditions, as follows:

- If a patient is to undergo elective surgery and an anti-platelet effect is not desired, Clopidogrel Winthrop Plus Aspirin should be discontinued 7 days prior to surgery.
- If the patient is at high risk of ophthalmic bleeding due to intraocular lesions clopidogrel should be used with extra caution.

Clopidogrel Winthrop Plus Aspirin should be used with caution in patients who have lesions with a propensity to bleed. Drugs that might induce such lesions (such as NSAIDs) are not recommended in patients taking Clopidogrel Winthrop Plus Aspirin (see ‘PRECAUTIONS’-Interactions with other medicines).

Clopidogrel Winthrop Plus Aspirin should be used with caution in patients with a history of peptic ulcer or gastroduodenal haemorrhage or minor upper gastrointestinal symptoms, as this may be due to gastric ulceration which may lead to gastric bleeding.

Gastrointestinal side effects, including stomach pain, heartburn, nausea, vomiting and gastrointestinal bleeding, may occur. Although minor upper gastrointestinal symptoms, such as dyspepsia, are common and can occur anytime during therapy, physicians should remain alert for signs of ulceration and bleeding, even in the absence of previous gastrointestinal symptoms. Patients should be told about the signs and symptoms of gastrointestinal side effects and what steps to take if they occur. Patients should be told that it may take longer than usual for
bleeding to stop when they take Clopidogrel Winthrop Plus Aspirin, and that they should report any unusual bleeding (site or duration) to their physician. Patients should inform physicians and dentists that they are taking Clopidogrel Winthrop Plus Aspirin before any surgery is scheduled and before any new drug is taken.

In patients with recent transient ischaemic attack or stroke who are at high risk of recurrent ischaemic events, the combination of aspirin and clopidogrel has been shown to increase major bleeding. Therefore, use of the combination of clopidogrel and aspirin should be undertaken with caution outside of clinical situations where the combination has proven to be beneficial.

To prevent gastric irritation due to aspirin, Clopidogrel Winthrop Plus Aspirin should be taken with or after food.

Due to the presence of aspirin, caution is required in patients with a history of asthma or allergic disorders (as they are at increased risk of hypersensitivity reactions) or with gout (as low doses of aspirin increase urate concentrations).

Patients with rare hereditary problems of galactose intolerance, the Lapp lactase deficiency or glucose-galactose malabsorption should not take this medicine.

The hypoglycaemic effect of chlorpropamide may be enhanced by the concurrent administration of aspirin. Large doses of aspirin may have intrinsic hypoglycaemic activity when given to diabetic patients, but the effects on carbohydrate metabolism are complex and it may cause hyperglycaemia.

Tinnitus is a premonitory sign of salicylism but may not be detected in patients with hearing loss.

This medicinal product also contains hydrogenated castor oil which may cause stomach upset and diarrhoea.

Clopidogrel Winthrop Plus Aspirin is to be used under medical supervision only.

**Coronary Artery Bypass Surgery**

When coronary artery bypass surgery is to be performed, clopidogrel and aspirin should be suspended at least 7 days before surgery to reduce the risk of bleeding (see 'ADVERSE EFFECTS').

**Cytochrome P450 2C19 (CYP2C19)**

Pharmacogenetics: In patients who are CYP2C19 poor metabolizers clopidogrel at recommended doses forms less of the active metabolite of clopidogrel and has a smaller effect on platelet function. Poor metabolisers with acute coronary syndrome or undergoing percutaneous coronary intervention treated with clopidogrel at recommended doses may exhibit higher cardiovascular event rates than do patients with normal CYP2C19 function (see Pharmacokinetics, Pharmacogenetics). Tests are available to identify a patient's CYP2C19 genotype; these tests can be used as an aid in determining therapeutic strategy. Consider the alternative treatment strategies in patients identified as CYP2C19 poor metabolisers (see Dosage and Administration, Pharmacogenetics).

**Ischaemic Stroke**

In view of the lack of data, clopidogrel cannot be recommended in acute ischaemic stroke (less than 7 days).

**Haematological**

Thrombotic Thrombocytopenic Purpura (TTP) has been reported very rarely following the use of clopidogrel, sometimes after a short exposure. It is characterised by thrombocytopenia and microangiopathic haemolytic anaemia associated with either neurological findings, renal dysfunction or fever. TTP is a potentially fatal condition requiring prompt treatment, including plasmapheresis (plasma exchange).

Thrombocytopenia, neutropenia, aplastic anaemia and pancytopenia have also been reported very rarely in patients taking clopidogrel (see ‘ADVERSE EFFECTS’).

Due to the risk of bleeding and haematological undesirable effects, blood cell count determination and/or other appropriate testing should be promptly considered whenever clinical symptoms suggestive of bleeding arise during the course of treatment. With chronic
administration, occult blood loss may lead to iron deficiency anaemia. As a dual anti-platelet agent, Clopidogrel Winthrop Plus Aspirin should be used with caution in patients who may be at risk of increased bleeding from trauma, surgery or other pathological conditions and in patients receiving treatment with other NSAIDs including Cox-2 inhibitors, heparin, glycoprotein IIb/IIIa inhibitors or thrombolytics. Patients should be followed carefully for any signs of bleeding including occult bleeding, especially during the first weeks of treatment and/or after invasive cardiac procedures or surgery.

**Renal Impairment**

Experience with clopidogrel plus aspirin is limited in patients with mild to moderate renal impairment. Therefore Clopidogrel Winthrop Plus Aspirin should be used with caution in this population. Patients should be observed closely for signs of salicylism. See also ‘CONTRAINDICATIONS’ for severe renal impairment.

**Hepatic Impairment**

Experience is limited in patients with moderate hepatic disease who may have bleeding diatheses. Clopidogrel Winthrop Plus Aspirin should therefore be used with caution in this population. See also ‘CONTRAINDICATIONS’ for severe hepatic impairment.

**Carcinogenicity**

There was no evidence of carcinogenic effects when clopidogrel was given in the diet for 78 weeks to mice and 104 weeks to rats at doses up to 77 mg/kg per day (representing an exposure \( \approx 18 \) times the anticipated patient exposure, based on plasma AUC for the main circulating metabolite in elderly subjects).

Carcinogenicity studies have not been conducted with aspirin.

**Genotoxicity**

Clopidogrel was not genotoxic in four *in vitro* tests (Ames test, DNA-repair test in rat hepatocytes, gene mutation assay in Chinese hamster fibroblasts and metaphase chromosome analysis of human lymphocytes) and in one *in vivo* test (micronucleus test by the oral route in mice).

Aspirin was not genotoxic in bacterial reverse mutation assays or in a recessive lethal mutation assay in Drosophila. However, there are conflicting results on the clastogenicity of aspirin in mammalian cells.

**Effects on Fertility**

Clopidogrel was found to have no effect on fertility of male and female rats at oral doses up to 400 mg/kg per day.

Aspirin had antispermatogenic effects by inhibiting prostaglandin formation in Long–Evans rats at 250 mg/kg/day PO, but did not affect the fertility of male Wistar rats at 300 mg/kg/day IP. The clinical relevance of these observations is unknown.

**Use in Pregnancy (Category C)**

No clinical data on exposure to clopidogrel plus aspirin during pregnancy are available. Clopidogrel plus aspirin should not be used during the first two trimesters of pregnancy unless the clinical condition of the woman requires treatment with clopidogrel in combination with aspirin. Due to the present of aspirin clopidogrel plus aspirin is contraindicated during the third trimester of pregnancy.

Clopidogrel and/or its metabolites are known to cross the placenta in pregnant rats and rabbits. However, teratology studies in rats and rabbits at doses up to 500 mg and 300 mg/kg/day PO, respectively, revealed no evidence of embryotoxicity or teratogenicity.

Reproduction toxicity data show that aspirin is teratogenic in several laboratory animals.
Aspirin inhibits prostaglandin synthesis. When given late in pregnancy, it may cause premature closure of the foetal ductus arteriosus, prolong labour and delay birth. Aspirin increases bleeding time both in the newborn infant and in the mother because of its anti-platelet effects. Clopidogrel Winthrop Plus Aspirin should not be used in women during pregnancy unless the potential benefits outweigh the risks.

Use in Lactation
Breast-feeding is contraindicated during treatment with Clopidogrel Winthrop Plus Aspirin (see ‘CONTRAINDICATIONS’). Studies in rats have shown that clopidogrel and/or its metabolites are excreted in breast milk. Salicylates are excreted in breast milk. Chronic high doses of aspirin can cause adverse effects in the infant.

Interactions with alcohol
The effect of alcohol on the safety and efficacy of the combination of clopidogrel and aspirin has not been investigated in clinical trials. Concurrent ingestion of alcohol and aspirin may enhance occult blood loss and gastric irritation. In prolonged aspirin administration, occult blood loss may lead to iron deficiency anaemia. Aspirin inhibits ethanol dehydrogenase, a major enzyme in the first pass elimination of alcohol. 
In vitro, the metabolism of clopidogrel has been shown to be altered in the presence of ethanol, such that clopidogrel is hydrolysed (inactivated) more slowly, and ethyl clopidogrel formed; the toxicity of ethyl clopidogrel has not been fully investigated.

INTERACTIONS WITH OTHER MEDICINES
Aspirin
A pharmacodynamic interaction between clopidogrel and aspirin is possible, leading to increased risk of bleeding. Therefore, concomitant use should be undertaken with caution. However, clopidogrel and aspirin have been administered together for up to one year. See also ‘PRECAUTIONS’ - General
Clopidogrel Winthrop Plus Aspirin should not be administered simultaneously with other salicylate containing preparations, uricosuric agents or NSAIDs.
Oral Anticoagulants (including warfarin)
The concomitant administration of Clopidogrel Winthrop Plus Aspirin with oral anticoagulants, including warfarin, is not recommended since it may increase the intensity of bleeding (see PRECAUTIONS).
Glycoprotein IIb/IIIa inhibitors
Clopidogrel Winthrop Plus Aspirin should be used with caution in patients who may be at risk of increased bleeding from trauma, surgery or other pathological conditions that receive concomitant glycoprotein IIb/IIIa inhibitors (see PRECAUTIONS). As a pharmacodynamic interaction between Clopidogrel Winthrop Plus Aspirin and glycoprotein IIb/IIIa inhibitors is possible, concomitant use should be undertaken with caution.
Injectable Anticoagulants
A pharmacodynamic interaction between Clopidogrel Winthrop Plus Aspirin and heparin is possible, leading to increased risk of bleeding. Therefore, concomitant use should be undertaken with caution.
Anti-platelet agents (such as eptifibatide, ticlopidine, tirofiban)
The effects of Clopidogrel Winthrop Plus Aspirin and other drugs which inhibit platelet aggregation may be additive, leading to an increased risk of bleeding.
Thrombolytics
The safety of the concomitant administration of clopidogrel, fibrin or non-fibrin specific thrombolytic agents and heparins was assessed in patients with acute myocardial infarction. The incidence of clinically significant bleeding was similar to that observed when thrombolytic
agents and heparins are co-administered with aspirin. The safety of concomitant administration of Clopidogrel Winthrop Plus Aspirin with thrombolytic agents has not been formally established and should be undertaken with caution.

**Methotrexate**

Due to the presence of aspirin, methotrexate and Clopidogrel Winthrop Plus Aspirin should be used together with caution, as aspirin can inhibit renal clearance of methotrexate, which may lead to bone marrow toxicity. Salicylates can also displace methotrexate from albumin.

**Non Steroidal Anti-inflammatory Drugs (NSAIDs)**

Aspirin may increase the risk of gastrointestinal side effects, including bleeding, when administered with NSAIDs. Aspirin displaces diclofenac from its binding sites, reducing diclofenac effectiveness.

The concomitant administration of ibuprofen with aspirin may limit the beneficial cardiovascular effects of aspirin in patients with increased cardiovascular risk.

In a clinical study conducted in healthy volunteers, the concomitant administration of clopidogrel and naproxen increased occult gastrointestinal blood loss. Consequently, the concomitant use of NSAIDs including Cox-2 inhibitors is not recommended with Clopidogrel Winthrop Plus Aspirin (see 'PRECAUTIONS').

**Uricosuric agents (e.g. probenecid)**

Caution is required because aspirin may inhibit the effect of uricosuric agents through competitive elimination of uric acid.

**Drugs metabolised by Cytochrome P450 2C9**

At high concentrations *in vitro*, clopidogrel inhibits cytochrome P450 2C9 and at lower concentrations inhibits CYP2B6 and CYP2C19. Accordingly, Clopidogrel Winthrop Plus Aspirin may interfere with the metabolism of bupropion, lansoprazole, omeprazole, pantoprazole, diazepam, phenytoin, tamoxifen, tolbutamide, warfarin, fluvastatin, and many NSAIDs, but there are no data with which to predict the magnitude of these interactions. Caution should be used when any of these drugs is co-administered with Clopidogrel Winthrop Plus Aspirin.

**Other concomitant therapy**

Since clopidogrel is metabolised to its active metabolite partly by CYP2C19, use of drugs that inhibit the activity of this enzyme would be expected to result in reduced drug levels of the active metabolite of clopidogrel. The clinical relevance of this is uncertain. Concomitant use of drugs that inhibit CYP2C19 (e.g., omeprazole) should be discouraged.

Medicinal products that inhibit CYP2C19 include omeprazole and esomeprazole, fluvoxamine, fluoxetine, moclobemide, voriconazole, flucanazole, ticlopidine, ciprofloxacin, cimetidine, carbamazepine, oxcarbazepine and chloramphenicol.

Care should be observed when coadministering aspirin and methotrexate, chlorpropamide, corticosteroids, sulfinpyrazone, probenecid and spironolactone. The hypoglycaemic effect of chlorpropamide may be enhanced by the concurrent administration of aspirin.

Hydrocortisone may increase the renal clearance of salicylate and when hydrocortisone is discontinued, serum salicylate levels may rise significantly. Aspirin may antagonise the diuretic effect of spironolactone. The rate and extent of aspirin absorption is increased by caffeine. The rate of excretion is increased by urinary alkalinisers. Aspirin at high doses reduces the uricosuric effects of probenecid and sulfinpyrazone.

A number of other clinical studies have been conducted with clopidogrel and other concomitant medications to investigate the potential for pharmacodynamic and pharmacokinetic interactions. No clinically significant pharmacodynamic interactions were observed when clopidogrel was co-administered with atenolol, nifedipine, or both atenolol and nifedipine. Furthermore, the pharmacodynamic activity of clopidogrel was not significantly influenced by the co-administration of phenobarbital, cimetidine, or oestrogen.

The pharmacokinetics of digoxin or theophylline were not modified by the co-administration of clopidogrel. Antacids did not modify the extent of clopidogrel absorption.
Combination use of ACE inhibitors or angiotensin receptor antagonists, anti-inflammatory drugs and thiazide diuretics

Concomitant use of a renin-angiotensin system inhibiting drug (angiotensin converting enzyme (ACE) inhibitor or angiotensin receptor antagonist), an anti-inflammatory drug (NSAID, including aspirin or COX-2 inhibitors) and a thiazide diuretic may increase the risk of renal impairment. This includes use in fixed-combination products containing more than one class of drug. The combination of these agents should be administered with caution, especially in the elderly and in patients with pre-existing renal impairment. Renal function (serum creatinine) should be monitored after initiation of concomitant therapy and periodically thereafter.

Interactions with higher dose aspirin

Interactions with the following medicines with higher (anti-inflammatory) doses of aspirin have been reported: alendronate, ACE inhibitors, anticonvulsants (phenytoin and valproic acid), beta blockers, systemic corticosteroids, diuretics, selective serotonin reuptake inhibitors (SSRIs), spironolactone, verapamil, hypoglycaemic agents and zafirlukast.

More than 30,000 patients entered into clinical trials with clopidogrel plus aspirin at maintenance doses lower than or equal to 325 mg, received a variety of concomitant medications, including diuretics, beta blockers, ACE inhibitors, calcium channel antagonists, cholesterol lowering agents, coronary vasodilators, anti-diabetic agents (including insulin), anti-epileptic agents, hormone replacement therapy and GPIIb/IIIa antagonists, without evidence of clinically significant adverse interactions.

Effects on ability to drive and use machines

Clopidogrel Winthrop Plus Aspirin has no or negligible influence on the ability to drive and use machines.

ADVERSE EFFECTS

Clopidogrel

Clinical Studies Experience

Clopidogrel has been evaluated for safety in more than 44,000 patients, including over 30,000 patients treated with clopidogrel plus aspirin, and over 12,000 patients treated for 1 year or more. The clinically relevant adverse events observed in CURE, CLARITY, COMMIT, CHARISMA, ACTIVE-A and ACTIVE-W are discussed below.

CURE, CLARITY AND COMMIT

Haemorrhagic disorders

In CURE, there was a significant difference between the two treatment groups for non life-threatening major bleeds (1.6% clopidogrel + aspirin vs. 1.0% placebo + aspirin), primarily gastrointestinal and at arterial puncture sites, and minor bleeds (5.1% clopidogrel + aspirin vs. 2.4% placebo + aspirin). The major bleeding event rate for clopidogrel + aspirin was dose-dependent on aspirin (<100 mg: 2.6%; 100-200 mg: 3.5%; >200 mg: 4.9%) as was the major bleeding event rate for placebo + aspirin (<100 mg: 2.0%; 100-200 mg: 2.3%; >200 mg: 4.0%).

The administration of clopidogrel + aspirin as compared to placebo + aspirin, was not associated with an increase in life-threatening or fatal bleeds (event rates 2.2% vs. 1.8% and 0.2% vs. 0.2%, respectively). The incidence of intra-cranial bleeding was 0.1% in both groups.

There was no excess in major bleeds within 7 days after coronary bypass graft surgery in patients who stopped therapy more than five days prior to surgery (4.4% clopidogrel + aspirin vs. 5.3% placebo + aspirin). In patients who remained on therapy within five days of bypass graft surgery, the event rate was 9.6% for clopidogrel + aspirin, and 6.3% for placebo + aspirin.

In CLARITY, there was an overall increase in bleeding in the clopidogrel + aspirin group (17.4%) versus the placebo + aspirin group (12.9%), with the incidence of major bleeding (defined as intracranial bleeding or bleeding associated with a fall in haemoglobin > 5 g/dL) being similar between groups (1.3% versus 1.1% in the clopidogrel + aspirin and the placebo + aspirin groups, respectively). This was consistent across subgroups of patients defined by
baseline characteristics, and type of fibrinolytics or heparin therapy. The incidence of fatal bleeding (0.8% versus 0.6% in the clopidogrel + aspirin and in the placebo + aspirin groups, respectively) and intracranial haemorrhage (0.5% versus 0.7%, respectively) was low and similar in both groups.

The overall rate of non-cerebral major bleeding or cerebral bleeding in COMMIT was low and similar in both groups, as shown in Table 6.

### Table 6: Number of Patients with Bleeding Events in COMMIT

<table>
<thead>
<tr>
<th>Type of bleeding</th>
<th>Clopidogrel + aspirin (n = 22,961)</th>
<th>Placebo + aspirin (n = 22,891)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Major * non-cerebral or cerebral bleeding</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Major non-cerebral</td>
<td>134 (0.6%)</td>
<td>125 (0.5%)</td>
<td>0.59</td>
</tr>
<tr>
<td>Fatal</td>
<td>82 (0.4%)</td>
<td>73 (0.3%)</td>
<td>0.48</td>
</tr>
<tr>
<td>Haemorrhagic stroke</td>
<td>55 (0.2%)</td>
<td>56 (0.2%)</td>
<td>0.91</td>
</tr>
<tr>
<td>Fatal</td>
<td>39 (0.2%)</td>
<td>41 (0.2%)</td>
<td>0.81</td>
</tr>
<tr>
<td>Other non-cerebral bleeding (non major)</td>
<td>831 (3.6%)</td>
<td>721 (3.1%)</td>
<td>0.005</td>
</tr>
<tr>
<td>Any non-cerebral bleeding</td>
<td>896 (3.9%)</td>
<td>777 (3.4%)</td>
<td>0.004</td>
</tr>
</tbody>
</table>

*Major bleeds are cerebral bleeds or non-cerebral bleeds thought to have caused death or that required transfusion

**Haematological disorders**

In CURE and CLARITY, the numbers of patients with thrombocytopenia or neutropenia were similar in both groups.

Although the risk of myelotoxicity with clopidogrel appears to be quite low, this possibility should be considered when a patient receiving clopidogrel demonstrates fever or other signs of infection.

**Gastrointestinal**

In CURE, there was no significant difference in the incidence of non-haemorrhagic gastrointestinal effects in the clopidogrel or placebo groups.

In CLARITY, the incidence of gastrointestinal adverse events was 6.9% for clopidogrel treated patients, compared to 7.2% in placebo treated patients.

In COMMIT, 2 patients reported gastrointestinal adverse events in the clopidogrel treated group, compared to one in the placebo treated group.

**Rash**

In CURE, rash occurred in more patients in the clopidogrel group. In CLARITY, 0.7% of patients in the clopidogrel group reported a rash, compared to 0.5% in the placebo group.

**Treatment Discontinuation**

In CURE, the overall incidence of discontinuation due to adverse events was greater in the clopidogrel group than in the placebo group (366 [5.8%] and 247 [3.9%] patients, respectively), with the main differences being in events in the platelet, bleeding and clotting disorders (1.1% versus 0.7%) and skin disorders (0.7% versus 0.3%). The increase in the rate of study drug discontinuation due to non-haemorrhagic adverse events was primarily due to the increase in rash seen in the clopidogrel group. There was no apparent difference between the 2 treatment groups in the rates of discontinuations due to other adverse events.

In CLARITY, the overall incidence of discontinuation due to adverse events was greater in the placebo group compared with the clopidogrel group (6.9% for clopidogrel treated patients compared to 8.6% for placebo treated patients).
In COMMIT, the overall incidence of discontinuation due to adverse events was similar in each treatment group (2.4% for clopidogrel treated patients compared to 2.2% for placebo treated patients).

**ACTIVE Studies**

The ACTIVE-W and ACTIVE-A studies, separate trials in the ACTIVE program, included patients with atrial fibrillation (AF) who had at least one risk factor for vascular events. Based on enrollment criteria, physicians enrolled patients in ACTIVE-W if they were candidates for vitamin K antagonist (VKA) therapy (such as warfarin). The ACTIVE-A study included patients who could not receive VKA therapy because they were unable or unwilling to receive the treatment.

The ACTIVE-W study demonstrated that treatment with VKA was more effective than the combination of clopidogrel and aspirin. The rate of major bleeding episodes was higher in the clopidogrel + aspirin group than in the VKA group: 101 (3.03%) subjects compared with 93 (2.76%).

The ACTIVE-A study demonstrated when preventing atherothrombotic and thromboembolic events including stroke, the rate of major bleeding was greater in the clopidogrel + aspirin group 251 (6.7%) than in the placebo + aspirin group 162 (4.3%).

**CAPRIE & CHARISMA**

The following safety data is extracted from clinical studies for different indications of clopidogrel.

**Haemorrhagic disorders**

In CAPRIE a study conducted in 19,185 patients with established atherosclerosis or history of atherothrombosis as manifested by myocardial infarction, ischaemic stroke or peripheral arterial disease, who randomised to clopidogrel 75 mg/day or aspirin 325 mg/day, and followed for 1 to 3 years, the overall incidence of any bleeding in patients treated with either clopidogrel or aspirin was similar (9.3%). The incidence of severe bleeds was 1.4% in the clopidogrel group and 1.6% in the aspirin group.

The overall incidence of other bleeding disorders was higher in the clopidogrel group (7.3%) compared to aspirin (6.5%). However, the incidence of severe events was similar in both treatment groups (0.6% vs. 0.4%). The most frequent events reported were purpura/bruising and epistaxis. Other less frequently reported events were haematoma, haematuria and eye bleeding (mainly conjunctival).

Gastrointestinal haemorrhage was significantly less frequent with clopidogrel (1.99%) compared to aspirin (2.66%). The incidence of intracranial haemorrhage was 0.35% for clopidogrel compared to 0.49% for aspirin.

In CHARISMA, a study conducted in patients with coronary artery disease, cerebrovascular disease or peripheral arterial disease as well as patients with a combination of atherothrombotic risk factors only, all receiving a background therapy with low dose aspirin (75-162 mg), there was an excess in moderate and severe bleeding, as adjudicated to the GUSTO definitions, in the clopidogrel group (see Table 7). This represented a number needed to treat, to harm, of 84 in 23 months of follow-up.

<table>
<thead>
<tr>
<th>Type of bleeding (GUSTO)</th>
<th>Number (%) with event</th>
<th>Difference Clopidogrel – Placebo (%) (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Any</td>
<td>Any</td>
<td>Any</td>
</tr>
<tr>
<td>Clopidogrel + aspirin</td>
<td>2827 (36.2)</td>
<td>1616 (20.7)</td>
</tr>
<tr>
<td>Placebo + aspirin</td>
<td>1616 (20.7)</td>
<td></td>
</tr>
<tr>
<td>Severe/moderate</td>
<td>290 (3.7)</td>
<td>197 (2.5)</td>
</tr>
</tbody>
</table>

**Table 7: Number of Patients with Bleeding Events in CHARISMA**
Haematological disorders

In CAPRIE, Clopidogrel was not associated with an increase in the incidence of thrombocytopenia compared to aspirin. Very rare cases of platelet count <30 × 10^9/L have been reported.

Severe neutropenia (<0.45 × 10^9/L) was observed in four patients (0.04%) that received clopidogrel and in two patients that received aspirin. Two of the 9599 patients who received clopidogrel and none of the patients who received aspirin had a neutrophil count of zero. One of the clopidogrel treated patients was receiving cytostatic chemotherapy, and another recovered and returned to the trial after only temporarily interrupting treatment with clopidogrel. One case of aplastic anaemia occurred on clopidogrel treatment. The incidence of severe thrombocytopenia (<80 G/L) was 0.2% on clopidogrel and 0.1% on aspirin; very rare cases of platelet count <=30 G/L have been reported.

Gastrointestinal

In CAPRIE, overall the incidence of gastrointestinal events (e.g. abdominal pain, dyspepsia, gastritis and constipation) in patients receiving clopidogrel was significantly lower than in those receiving aspirin. The incidence of peptic, gastric, or duodenal ulcers was 0.68% for clopidogrel and 1.15% for aspirin. Cases of diarrhoea were reported at a higher frequency in the clopidogrel group (4.46%) compared to the aspirin group (3.36%).

Rash

In CAPRIE, there were significantly more patients with rash in the clopidogrel group (4.2%) compared to the aspirin group (3.5%).

Treatment Discontinuation

In the clopidogrel and aspirin treatment groups of the CAPRIE study, discontinuation due to adverse events occurred in approximately 13% of patients after 2 years of treatment.

Adverse events occurring in ≥ 2.5% of patients on clopidogrel in the CAPRIE and CURE controlled clinical trials are shown in Table 8 regardless of relationship to clopidogrel.

<table>
<thead>
<tr>
<th>BODY SYSTEM/EVENT</th>
<th>CAPRIE</th>
<th>CURE</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>% Incidence (% discontinuation)</td>
<td>% Incidence (% discontinuation)</td>
</tr>
<tr>
<td><strong>Clopidogrel</strong> n = 9599</td>
<td>Aspirin n = 9586</td>
<td>Clopidogrel + aspirin n = 6259</td>
</tr>
<tr>
<td>Body as a Whole - general disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Chest pain</td>
<td>8.3 (0.2)</td>
<td>8.3 (0.3)</td>
</tr>
<tr>
<td>Accidental/inflicted injury</td>
<td>7.9 (0.1)</td>
<td>7.3 (0.1)</td>
</tr>
<tr>
<td>Influenza like symptoms</td>
<td>7.5 (&lt;0.1)</td>
<td>7.0 (&lt;0.1)</td>
</tr>
<tr>
<td>Pain</td>
<td>6.4 (0.1)</td>
<td>6.3 (0.1)</td>
</tr>
<tr>
<td>Fatigue</td>
<td>3.3 (0.1)</td>
<td>3.4 (0.1)</td>
</tr>
<tr>
<td>Cardiovascular disorders - general</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypertension</td>
<td>4.3 (&lt;0.1)</td>
<td>5.1* (&lt;0.1)</td>
</tr>
<tr>
<td>Central and peripheral nervous system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Headache</td>
<td>7.6 (0.3)</td>
<td>7.2 (0.2)</td>
</tr>
<tr>
<td>Dizziness</td>
<td>6.2 (0.2)</td>
<td>6.7 (0.3)</td>
</tr>
<tr>
<td>BODY SYSTEM/EVENT</td>
<td>CAPRIE</td>
<td>CURE</td>
</tr>
<tr>
<td>-----------------------------------------------</td>
<td>------------------------------------------------------------------------</td>
<td>----------------------------------------------------------------------</td>
</tr>
<tr>
<td></td>
<td>% Incidence (% discontinuation)</td>
<td>% Incidence (% discontinuation)</td>
</tr>
<tr>
<td></td>
<td>Clopidogrel n = 9599 Aspirin n = 9586</td>
<td>Clopidogrel + aspirin n = 6259 Placebo + aspirin n = 6303</td>
</tr>
<tr>
<td>Gastrointestinal</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Abdominal pain</td>
<td>5.6 (0.7)</td>
<td>7.1* (1.0)</td>
</tr>
<tr>
<td>Dyspepsia</td>
<td>5.2 (0.6)</td>
<td>6.1* (0.7)</td>
</tr>
<tr>
<td>Diarrhoea</td>
<td>4.5* (0.4)</td>
<td>3.4 (0.3)</td>
</tr>
<tr>
<td>Nausea</td>
<td>3.4 (0.5)</td>
<td>3.8 (0.4)</td>
</tr>
<tr>
<td>Metabolic and nutritional disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypercholesterolemia</td>
<td>4.0 (0)</td>
<td>4.4 (&lt;0.1)</td>
</tr>
<tr>
<td>Musculoskeletal system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Arthralgia</td>
<td>6.3 (0.1)</td>
<td>6.2 (0.1)</td>
</tr>
<tr>
<td>Back pain</td>
<td>5.8 (0.1)</td>
<td>5.3 (&lt;0.1)</td>
</tr>
<tr>
<td>Myo-, endo-, pericardial and valve disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Angina pectoris</td>
<td>10.1 (0.6)</td>
<td>10.7 (0.4)</td>
</tr>
<tr>
<td>Coronary artery disorder</td>
<td>6.2 (0.3)</td>
<td>5.6 (0.3)</td>
</tr>
<tr>
<td>Platelet, bleeding and clotting disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Purpura</td>
<td>5.3* (0.3)</td>
<td>3.7 (0.1)</td>
</tr>
<tr>
<td>Epistaxis</td>
<td>2.9 (0.2)</td>
<td>2.5 (0.1)</td>
</tr>
<tr>
<td>Psychiatric disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Depression</td>
<td>3.6 (0.1)</td>
<td>3.9 (0.2)</td>
</tr>
<tr>
<td>Resistance mechanism disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Infection</td>
<td>4.7 (&lt;0.1)</td>
<td>4.2 (0.1)</td>
</tr>
<tr>
<td>Respiratory system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>8.7 (&lt;0.1)</td>
<td>8.3 (&lt;0.1)</td>
</tr>
<tr>
<td>Dyspnoea</td>
<td>4.5 (0.1)</td>
<td>4.2 (0.1)</td>
</tr>
<tr>
<td>Rhinitis</td>
<td>4.2 (0.1)</td>
<td>4.2 (&lt;0.1)</td>
</tr>
<tr>
<td>Bronchitis</td>
<td>3.7 (0.1)</td>
<td>3.7 (0)</td>
</tr>
<tr>
<td>Coughing</td>
<td>3.1 (&lt;0.1)</td>
<td>2.7 (&lt;0.1)</td>
</tr>
<tr>
<td>Skin and appendage disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rash</td>
<td>4.2* (0.5)</td>
<td>3.5 (0.2)</td>
</tr>
<tr>
<td>Pruritis</td>
<td>3.3* (0.3)</td>
<td>1.6 (0.1)</td>
</tr>
<tr>
<td>Urinary system disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Urinary tract infection</td>
<td>3.1 (0)</td>
<td>3.5 (0.1)</td>
</tr>
<tr>
<td>Vascular (extracardiac) disorders</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Claudication intermittent</td>
<td>3.8 (0.2)</td>
<td>3.8 (0.2)</td>
</tr>
<tr>
<td>Peripheral ischaemia</td>
<td>3.2 (0.2)</td>
<td>3.4 (0.2)</td>
</tr>
<tr>
<td>Cerebrovascular disorder</td>
<td>2.6 (0.3)</td>
<td>2.9 (0.3)</td>
</tr>
</tbody>
</table>

* indicates statistical significance (p ≤ 0.05)
Incidence of discontinuation, regardless of relationship to therapy is shown in parentheses.

Clinically relevant adverse reactions not listed above pooled from CAPRIE, CURE, CLARITY and COMMIT studies with an incidence of ≥ 0.1% as well as all serious and clinically relevant adverse reactions are listed below according to the World Health Organisation classification. Their frequency is defined using the following conventions: common: > 1/100 (1%) and < 1/10
(10%); uncommon: ≥ 1/1000 (0.1%) and < 1/100 (1%) and rare: ≥ 1/10000 (0.01%) and < 1/1000 (0.1%).

**Central and peripheral nervous system disorders**

*Uncommon:* headache, dizziness, paraesthesia,
*Rare:* Vertigo

**Gastrointestinal system disorders**

*Common:* dyspepsia, abdominal pain, diarrhoea
*Uncommon:* nausea, gastritis, flatulence, constipation, vomiting, gastric, peptic or duodenal ulcer

**Platelet, bleeding and clotting disorders**

*Uncommon:* Bleeding time increased, decreased platelets

**Skin and appendages disorders**

*Uncommon:* rash, pruritus

**White cell and RES disorders**

*Uncommon:* Leucopenia, decreased neutrophils, eosinophilia

**Post-Marketing Experience**

In addition to clinical study experience with clopidogrel either alone or in combination with aspirin, the following is a list of adverse reactions reported with clopidogrel or aspirin.

Bleeding is the most common reaction reported in the post-marketing experience with clopidogrel or aspirin.

The following have been reported spontaneously from worldwide post-marketing experience with clopidogrel:

<table>
<thead>
<tr>
<th>Note</th>
<th>Frequency</th>
</tr>
</thead>
<tbody>
<tr>
<td>very common</td>
<td>≥ 1/10 (≥ 10%)</td>
</tr>
<tr>
<td>common</td>
<td>≥ 1/100 and &lt; 1/10 (≥ 1% and &lt; 10%)</td>
</tr>
<tr>
<td>uncommon</td>
<td>≥ 1/1000 and &lt; 1/100 (≥ 0.1% and &lt; 1.0%)</td>
</tr>
<tr>
<td>rare</td>
<td>≥ 1/10,000 and &lt; 1/1000 (≥ 0.01% and &lt; 0.1%)</td>
</tr>
<tr>
<td>very rare</td>
<td>&lt; 1/10,000 (&lt; 0.01%)</td>
</tr>
</tbody>
</table>

**Musculoskeletal, connective and bone**

*Very rare:* Arthralgia, arthritis, myalgia

**Immune system disorders**

*Very rare:* anaphylactoid reactions, serum sickness

**Vascular disorders**

*Very rare:* vasculitis, hypotension

**Blood and lymphatic system disorders**

*Very rare:* serious cases of bleeding, mainly skin, musculo-skeletal (haemarthrosis, haematoma), eye (conjunctival, ocular, retinal) and respiratory tract bleeding (haemoptysis, pulmonary haemorrhage), epistaxis, haematuria and haemorrhage of operative wound. Fatal haemorrhage, especially intracranial, gastrointestinal and retroperitoneal haemorrhage. Cases of serious haemorrhage have been reported in patients taking clopidogrel concomitantly with aspirin or clopidogrel with aspirin and heparin (see 'Interactions with Other Medicines').

*Very rare cases of thrombotic thrombocytopenic purpura (TTP) have been reported*
Aplastic anaemia, neutropenia, pancytopenia, agranulocytosis, granulocytopenia, anaemia

*Uncommon*: eosinophilia, leucopenia, decreased neutrophils, decreased platelets, increased bleeding time

**Skin and Subcutaneous tissue disorders**

*Very rare*: maculopapular or erythematous rash, urticaria, pruritus, angioedema, bullous dermatitis (erythema multiforme, Stevens-Johnson syndrome, toxic epidermal necrolysis), eczema, lichen planus

**Psychiatric**

*Very rare*: confusion, hallucinations

**Nervous System disorders**

*Very rare*: taste disturbances

**Hepatobiliary disorders**

*Very rare*: hepatitis, acute liver failure

**Gastrointestinal disorders**

*Very rare*: colitis (including ulcerative or lymphocytic colitis), pancreatitis, stomatitis

**Respiratory, thoracic and mediastinal disorders**

*Very rare*: bronchospasm, interstitial pneumonitis

**Renal and Urinary disorders**

*Very rare*: glomerulopathy

**Investigations**

*Very rare*: blood creatinine increase, abnormal liver function tests

**General disorders and administration site conditions**

*Very rare*: fever, syncope

**Aspirin**

In addition to some of the adverse reactions listed above, aspirin is associated with the following adverse effects.

Aspirin produces a prolongation of the bleeding time and may produce epigastric distress, nausea and vomiting, gastric or duodenal ulcers and erosive gastritis which may lead to serious gastrointestinal bleeding. These side effects are more likely to occur when higher doses are administered, although they may also occur when low doses are used.

Gastro-duodenal ulcer/perforations, upper gastro-intestinal symptoms such as gastralgia (see Precautions).

Iron deficiency anaemia may develop as a result of occult gastrointestinal bleeding when aspirin is used for long periods of time.

Aspirin may cause haemolysis in patients with glucose-6-phosphate dehydrogenase deficiency. Aspirin may cause tinnitus, dizziness, vertigo or hearing loss.

Aspirin sensitivity is most commonly manifested by asthma, vasomotor rhinitis, urticaria, angioneurotic oedema and allergic dermatological reactions, hypoglycaemia, gout. As well as anaphylactic shock, aggravation of allergic symptoms of food allergy.

Low doses of aspirin have been reported to cause retention of uric acid, whereas high dosage may increase excretion.

Aspirin may cause acute renal impairment (especially in patients with existing renal impairment, heart decompensation, nephritic syndrome, or concomitant treatment with diuretics).
DOSAGE AND ADMINISTRATION

Adults

Clopidogrel Winthrop Plus Aspirin is given as a single tablet (75mg/75mg or 75mg/100mg) once a day taken with adequate water.

Acute Coronary Syndrome

Clopidogrel Winthrop Plus Aspirin is used following an initial loading dose of 300 mg clopidogrel in combination with aspirin in patients with acute coronary syndrome:

- Unstable angina or non-ST-elevation myocardial infarction:
  - Treatment should be initiated with a single 300 mg loading dose of clopidogrel plus aspirin (75 mg to 325 mg).
  - Long-term daily treatment should be continued with one Clopidogrel Winthrop Plus Aspirin tablet (75mg/75mg or 75mg/100mg) once a day taken with adequate water.

- ST-segment elevation acute myocardial infarction:
  - Treatment should be initiated with or without a 300-mg loading dose of clopidogrel in combination with aspirin and with or without thrombolytics as soon as possible after symptoms start. There are no data on the use of a 300-mg loading dose in elderly patients (aged 75 years or more) with ST-segment acute myocardial infarction, as no patients over 75 years old were included in the CLARITY study and no loading dose was used in the COMMIT study.
  - Daily treatment should continue with one Clopidogrel Winthrop Plus Aspirin tablet (75mg/75mg or 75mg/100mg) once a day with adequate water. The benefit of the combination of clopidogrel with aspirin beyond four weeks has not been studied in this setting.

In patients who have had percutaneous coronary intervention with stent insertion, clopidogrel and aspirin should be continued for as long as is currently recommended in evidence-based guidelines for the type of stent and circumstances of implantation or for as long as otherwise indicated, taking into account the overall atherothrombotic risk profile of the patient.

Should doses of aspirin greater than 100 mg be required for daily maintenance therapy, clopidogrel and aspirin products should be administered separately.

Pharmacogenetics

CYP2C19 poor metaboliser status is associated with diminished antiplatelet response to clopidogrel. A higher dose of clopidogrel (600 mg loading dose followed by 150 mg once daily) in poor metabolisers increases antiplatelet response (see Pharmacokinetics, Pharmacogenetics). Consider the use of higher clopidogrel doses in patients who are poor CYP2C19 metabolisers. An appropriate dose regimen for this patient population has not been established in clinical outcome trials.

Renal Impairment

Experience is limited in patients with mild to moderate renal impairment (see ‘PRECAUTIONS’). Clopidogrel Winthrop Plus Aspirin should not be used in patients with severe renal impairment (see ‘CONTRAINDICATIONS’).

Hepatic Impairment

Experience is limited in patients with moderate hepatic disease who may have bleeding diatheses (see ‘PRECAUTIONS’). Clopidogrel Winthrop Plus Aspirin should not be used in patients with severe hepatic impairment (see ‘CONTRAINDICATIONS’).

No dosage adjustment is necessary for elderly patients (see ‘PHARMACOLOGY’-Special Populations).

Children and Adolescents

Safety and efficacy in subjects below the age of 18 have not been established. There is a possible association between aspirin and Reye’s syndrome when aspirin is given to children. Reye’s syndrome is a very rare disease which can be fatal.
OVERDOSAGE
There is no information concerning overdose with Clopidogrel Winthrop Plus Aspirin.
In animals, clopidogrel at single oral doses ≥ 1500 mg/kg caused necrotic-haemorrhagic gastritis, oesophagitis and enteritis in mice, rats and baboons. Necrotic tubulopathy and tubulo-interstitial nephritis were also noted in mice.

Overdose following clopidogrel administration may lead to prolonged bleeding time and subsequent bleeding complications. Appropriate therapy should be considered if bleeding is observed. No antidote to the pharmacological activity of clopidogrel has been found. If prompt correction of prolonged bleeding time is required, platelet transfusion may reverse the effects of clopidogrel.

Aspirin overdose is manifested by the following symptoms:
Moderate overdose: tinnitus, hearing loss, dizziness, headaches, vertigo, confusion and gastrointestinal symptoms (nausea, vomiting and gastric pain).
Severe overdose: fever, hyperventilation, ketosis, respiratory alkalosis, metabolic acidosis, coma, cardiovascular collapse, respiratory failure, severe hypoglycaemia, haemorrhage

In case of severe aspirin overdose, the following actions should be undertaken: admission to hospital is necessary, control of acid-base balance, possibility of haemodialysis or peritoneal dialysis if necessary.

Apart from general measures, treatment of aspirin overdose consists chiefly of measures to accelerate the excretion (forced alkaline diuresis) and to restore the acid-base and electrolyte balance. Infusions of sodium bicarbonate and potassium chloride solutions may be given.

For advice on the management of overdose, please contact the Poisons Information Centre on 13 11 26.

PRESENTATION AND STORAGE CONDITIONS
Clopidogrel Winthrop Plus Aspirin 75mg/75mg tablets are yellow, oval, slightly biconvex, film-coated, engraved with “C75” on one side and “A75” on the other side.
Clopidogrel Winthrop Plus Aspirin 75mg/100mg tablets are light pink, oval, slightly biconvex, film-coated, engraved with “C75” on one side and “A100” on the other side.
Store below 25°C.

Clopidogrel Winthrop Plus Aspirin 75mg/75mg tablets are registered in blister packs containing 2*, 4*, 7*, 14*, 28*, 30*, 50*, 56*, 84*, 98*, 100*, 112* and 280* tablets.

Clopidogrel Winthrop Plus Aspirin 75mg/100mg tablets are registered in blister packs containing 2*, 4*, 7*, 14*, 28*, 30, 50*, 56*, 84*, 98*, 100*, 112* and 280* tablets.
Store below 25°C
* Presentations currently not marketed

NAME AND ADDRESS OF SPONSOR
sanofi-aventis australia pty ltd
12-24 Talavera Road
Macquarie Park, NSW 2113

POISON SCHEDULE OF THE MEDICINE
Schedule 4.

DATE OF FIRST INCLUSION IN THE ARTG: 24 September 2009

DATE OF MOST RECENT AMENDMENT: 19 August 2011

Clopidogrel Winthrop Plus Aspirin is a registered trademark of sanofi-aventis.