Dissolution test of various low-dose acetylsalicylic acid preparations marketed in Indonesia

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Abstrak

Tujuan Membandingkan profil disolusi berbagai tablet asam asetilsalizilat dosis rendah salut enterik yang dipasarkan di Indonesia.

Metode Studi disolusi dilakukan mengikuti Farmakope Amerika (USP)/Eropa, metode A, menggunakan alat 1 USP (keranjang) 100 rpm, dengan 2 media: 0,1 N HCl, 120 menit untuk stadium asam, dan buffer fosfat pH 6,8, 90 menit untuk stadium buffer. Sampel diambil pada 120 menit untuk stadium asam, dan setiap 10 menit sampai dengan 90 menit untuk stadium buffer. Asam asetilsalizilat diukur dengan spektrofotometer pada 280 nm untuk stadium asam, dan pada 265 nm untuk stadium buffer. Asam asetilsalizilat bebas diukur pada akhir stadium dengan metode HPLC. Ada 6 produk uji (Cardio Aspirin® 100 mg, Aprot® 100 mg, Ascardia® 80 mg, Thrombo Aspilet® 80 mg, Astika® 100 mg dan Farmasal® 100 mg), 3 batch untuk setiap produk, dan 6 unit untuk setiap batch.

Hasil Jumlah asam asetilsalizilat yang dilepaskan dari setiap produk asam asetilsalizilat yang diuji pada akhir stadium asam (120 menit) berkisar dari 1,79% untuk Cardio Aspirin® sampai 6,92% untuk Thrombo Aspilet®, semua produk memenuhi persyaratan farmakope sebagai produk salut enterik (< 10%). Jumlah asam asetilsalizilat yang dilepaskan dari setiap produk pada akhir stadium buffer berkisar dari 3,47% untuk Cardio Aspirin® sampai 10,90% untuk Astika® dan 11,90% untuk Thrombo Aspilet®. Thrombo Aspilet® menunjukkan sifat lepas lambat, yang menyebabkan variabilitas yang sedemikian tinggi dalam melepaskan asam asetilsalizilat, sehingga salah satu dari 3 batch yang diuji tidak memenuhi persyaratan kompendial yaitu lebih dari 75% (pelepasan hanya 55,11%). Variabilitas yang tinggi dalam melepaskan asam asetilsalizilat antar batch juga ditemukan pada Farmasal® pada 10, 20, dan 30 menit dalam medium buffer. Dosis efektif asam asetilsalizilat terendah sebagai obat antiplatelet untuk penggunaan jangka panjang adalah 75 mg asam asetilsalizilat sebagai tablet biasa, yang ekivalen dengan 100 mg asam asetilsalizilat sebagai tablet salut enterik.


Abstrak

Aim To compare the dissolution profiles of various enteric-coated low-dose acetylsalicylic acid (ASA) tablets marketed in Indonesia.

Methods The dissolution study was carried out according to US Pharmacopoeia (USP)/European Pharmacopoeia, method A, using USP apparatus 1 (basket) 100 rpm, with 2 media: 0.1 N HCl, 120 minutes for acid stage, and phosphate buffer pH 6.8, 90 minutes for buffer stage. The sampling points were 120 minutes for the acid stage, and every 10 minutes until 90 minutes for the buffer stage. The acetylsalicylic acid was assayed using spectrophotometry at 280 nm for the acid stage, and at 265 nm for the buffer stage. The free salicylic acid was determined only at the end of the buffer stage with HPLC. There were 6 test products (Cardio Aspirin® 100 mg, Aprot® 100 mg, Ascardia® 80 mg, Thrombo Aspilet® 80 mg, Astika® 100 mg and Farmasal® 100 mg), 3 batches for each product, and 6 units for each batch.

Results The amount of ASA released from each ASA product tested at the end of acid stage (120 minutes) ranged from 1.79% for Cardio Aspirin® to 6.92% for Thrombo Aspilet®, all conformed to the compendial requirement for enteric-coated product (< 10%). The amount of salicylic acid observed at the end of the dissolution test ranged from 3.47% for Cardio Aspirin® to 10.90% for Astika® and 11.90% for Thrombo Aspilet®. Thrombo Aspilet® showed sustained-release properties, causing high variability in ASA release, such that one of the 3 batches tested did not fulfill the compendial requirement of more than 75% (the release was only 55.11%). High variability in ASA release between batches was also found with Farmasal® at 10, 20, and 30 minutes in buffer medium. The lowest effective dose of ASA as an antiplatelet drug for long-term use is 75 mg of plain ASA, and this is equivalent to 100 mg of enteric-coated ASA.

Conclusions All of the low-dose ASA preparations marketed in Indonesia are enteric-coated products, while Thrombo Aspilet® is not only an enteric-coated but also a sustained-release product. Cardio Aspirin®, followed by Aprot®, has the right dose for low-dose enteric-coated preparation (100 mg), produces consistent ASA release between batches, and the most stable towards deacetylation (antiplatelet inactivation). (Med J Indones 2009; 18: 161-6)

Key words: Dissolution profile, enteric coated, deacetylation
Low-dose acetylsalicylic acid (ASA) has been used worldwide as an antiplatelet agent for prevention of cardiovascular death, myocardial infarction, and stroke in patients at high risk of occlusive vascular events. Among various antiplatelet agents, ASA is the oldest, the cheapest, and the most established, and therefore it becomes the standard drug to which other antiplatelets are compared.

Among the above-mentioned high risk patients, antiplatelet therapy, especially acetylsalicylic acid (ASA), has been found to reduce the combined outcome of any serious vascular event by about one quarter, nonfatal myocardial infarction by one third, nonfatal stroke by one quarter, and vascular mortality by one sixth in a meta-analysis of >100 randomized trials. For this purpose, ASA has been used as life-long secondary and primary prevention medication; the most effective dose according to the meta-analysis is 75-150 mg daily. The higher doses are less effective because higher doses also block the synthesis of prostacyclin which has platelet antiaggregation and vasodilator effects, while the lower doses are reported to “spare” prostacyclin. Higher doses also produce more gastrointestinal side effects and have higher risks of major bleeding. To reduce the gastrointestinal side effects, ASA is formulated as enteric-coated tablets, which prevent the release of ASA in the stomach, and thereby preventing its perfusion into gastric mucosal cells which causes gastric injury.

The present study was conducted to compare the dissolution profiles of various enteric-coated low-dose ASA tablets marketed in Indonesia.

METHODS

Products tested

The products tested (3 batches for each product) were: Cardio Aspirin®, 100 mg tablet (Batch number/BN: BXC7A91, BXBVUP2, BXC8EZ1), Aptom®, 100 mg tablet (BN: 2D1137B, 7B1030A, 611584), Ascardia®, 80 mg tablet (BN: B7G676G, B7C973F, B7G674G), Astika®, 100 mg tablet (BN: 760306, 761203, 761216), Farmasa®, 100 mg tablet (BN: TH 0921, PJ 0701, TC 0021), and Thrombo Aspilets® 80 mg tablet (BN: 7032501, 6063102, 7032401). All of the products were low-dose ASA that were produced in Indonesia. These products were supplied by PT Bayer Indonesia.

Instruments

The main instruments used were dissolution tester (United States Pharmacopoeia/USP type) Erweka DT6, high performance liquid chromatograph (HPLC) with variable UV-Vis detector (HP 1100 series), spectrophotometer Beckman DU 7500i, and pH-meter Beckman™ 50.

Dissolution study

This study was conducted at School of Pharmacy, Bandung Institute of Technology, Bandung, in the first quarter of 2008. There are two dissolution methods for ASA that are used and accepted in Indonesia; one of these methods is the method of USP/European Pharmacopoeia.

The apparatus used was USP apparatus 1 (basket) 100 rpm. The method was according to USP/European Pharmacopoeia, method A, using 2 media: medium 1 was 0.1N HCl (pH=1), for acid stage, and medium 2 was phosphate buffer pH 6.8, for buffer stage. The sampling time for acid stage was 120 minutes, while for buffer stage were every 10 minutes until 90 minutes. Spectrophotometry was used for the assay of ASA, at 280 nm for acid stage, and at 265 nm for buffer stage. The free salicylic acid was determined only at the end of buffer stage with HPLC method. There were 3 batches for each product tested, and 6 units for each batch. These batches and units were randomly taken from the market (a pharmacy in Jakarta) by PT Bayer Indonesia.

Procedure

Seven hundred fifty mL of medium 1 (0.1 N HCl) was transferred to the dissolution vessel and warmed to 37°C (+ 0.5°C). One tablet was placed in the basket, and the basket was put in the dissolution vessel, the stirrer was switched to 100 rpm and run for 120 minutes. Five mL of the dissolution medium was taken as the first sample and the withdrawn dissolution medium was replaced with the same quantity of medium 1. The aspirin concentration in the sample was determined spectrophotometrically at 280 nm.

Immediately after sampling and medium replacement at the end of acid stage, 250 mL of medium 2 (0.20 M tribasic sodium phosphate, previously heated to 37°C) was added. The pH was checked and was adjusted if necessary to 6.8 ± 0.05 with 2N HCl or 2N NaOH. Five mL of the dissolution medium was withdrawn regularly every 10 minutes until 90 minutes. Each time of sampling, withdrawn medium was replaced with the same quantity of the same medium. The ASA concentrations in the samples were determined spectrophotometrically at 265 nm (at which the absorption is the same for both ASA and salicylic acid).
Free salicylic acid concentration was determined only at
the end of buffer stage (90 minutes) with validated HPLC
method (with this method, the concentrations of ASA
and salicylic acid can be determined simultaneously).
The HPLC condition was as follows: the column was
SGE, Wakosil C-18, 250 x 4.6 mm, 5 µm; the mobile
phase was phosphate buffer-acetonitrile (60 : 40) pH 2.5,
1 mL/minute, with UV detector at 237 nm.

**Estimation of drug dissolution rates**

Drug dissolution rate was estimated during the main
dissolution phase (between 10% and 90% of drug
release)

**Estimation of similarity factors**

Similarity factor (f2) was estimated using time points
from 10 to 40 minutes. Those points were taken in
accordance to the general guideline in calculating
similarity factor (not more than one point that was
greater than 85%). The similarity factors would show
the similarity or the difference between the dissolution
profiles of the products tested.

**Analytical methods**

The analytical methods (spectrophotometric and HPLC
methods) were verified for selectivity, linearity, and
precision. The selectivities of both methods were tested
for possible interference from dissolution medium, not
from tablet excipients since the excipients of each tablet
tested were not available.

**RESULTS**

No interference was found from the dissolution
medium. The standard curve of ASA concentration
vs absorbance by spectrophotometric method in acid
medium was linear with $R^2 = 0.9995$, and that in buffer
medium was also linear with $R^2 = 1$. The standard curve
of salicylic acid concentration vs peak area by HPLC
method was linear with $R^2 = 0.9996$, and that of ASA
concentration was also linear with $R^2 = 0.9999$. The
precision of ASA assay by spectrophotometric method
in acid medium was high (coefficient of variation/CV=
0.70% and 1.25% for absorbance of 1 ppm and 4 ppm,
respectively); and that in buffer medium was also high
(CV= 1.09, 2.37, and 1.17% for absorbance of 0.5 ppm,
45 ppm, and 90 ppm, respectively). The precision of
salicylic acid assay by HPLC method was also high
(CV= 5.58, 3.54, and 2.88% for peak area of 1 ppm, 50
ppm, and 150 ppm, respectively).

**Dissolution results**

The results of the dissolution study of all ASA products
tested are presented in Table 1 and Figure 1.

<table>
<thead>
<tr>
<th>Time (minute)</th>
<th>Cardio Aspirin® (100 mg)</th>
<th>Aptor® (100 mg)</th>
<th>Ascardia® (80 mg)</th>
<th>Thro.Aspilet® (80 mg)</th>
<th>Astika® (100 mg)</th>
<th>Farmasal® (100 mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acid medium (0.1 N HCl) 120</td>
<td>1.79 ± 1.47 (1.79%)</td>
<td>3.70 ± 1.11 (3.70%)</td>
<td>2.61 ± 1.41 (3.26%)</td>
<td>5.53 ± 3.00 (6.92%)</td>
<td>2.74 ± 0.80 (2.74%)</td>
<td>3.25 ± 1.71 (3.25%)</td>
</tr>
<tr>
<td>Buffer medium (pH 6.8)</td>
<td>0</td>
<td>1.79 ± 1.47</td>
<td>3.70 ± 1.11</td>
<td>2.61 ± 1.41</td>
<td>5.53 ± 3.00</td>
<td>2.74 ± 0.80</td>
</tr>
<tr>
<td>10</td>
<td>1.53 ± 0.61</td>
<td>11.43 ± 6.62</td>
<td>48.38 ± 13.57</td>
<td>14.08 ± 2.35</td>
<td>12.13 ± 2.90</td>
<td>38.84 ± 34.66</td>
</tr>
<tr>
<td>20</td>
<td>14.86 ± 1.96</td>
<td>83.22 ± 3.14</td>
<td>86.46 ± 4.70</td>
<td>22.29 ± 6.09</td>
<td>65.93 ± 5.00</td>
<td>75.52 ± 37.60</td>
</tr>
<tr>
<td>30</td>
<td>54.44 ± 8.89</td>
<td>103.87 ± 1.53</td>
<td>92.62 ± 1.31</td>
<td>30.97 ± 7.47</td>
<td>93.87 ± 3.43</td>
<td>94.84 ± 20.08</td>
</tr>
<tr>
<td>40</td>
<td>90.25 ± 0.56</td>
<td>107.88 ± 1.95</td>
<td>91.94 ± 1.38</td>
<td>38.15 ± 9.30</td>
<td>100.05 ± 1.83</td>
<td>100.56 ± 13.14</td>
</tr>
<tr>
<td>50</td>
<td>101.22 ± 3.60</td>
<td>108.34 ± 2.43</td>
<td>90.87 ± 0.58</td>
<td>44.59 ± 11.35</td>
<td>100.24 ± 2.46</td>
<td>102.59 ± 9.07</td>
</tr>
<tr>
<td>60</td>
<td>102.06 ± 4.28</td>
<td>107.78 ± 2.22</td>
<td>89.81 ± 0.44</td>
<td>49.99 ± 13.26</td>
<td>99.70 ± 1.45</td>
<td>103.07 ± 8.51</td>
</tr>
<tr>
<td>70</td>
<td>102.15 ± 5.37</td>
<td>107.12 ± 2.51</td>
<td>89.08 ± 0.50</td>
<td>54.75 ± 15.26</td>
<td>99.64 ± 1.73</td>
<td>103.51 ± 7.89</td>
</tr>
<tr>
<td>80</td>
<td>102.84 ± 4.67</td>
<td>106.95 ± 2.34</td>
<td>88.92 ± 0.97</td>
<td>59.27 ± 16.31</td>
<td>99.13 ± 1.71</td>
<td>103.35 ± 7.93</td>
</tr>
<tr>
<td>90</td>
<td>100.90 ± 4.70</td>
<td>105.26 ± 2.60</td>
<td>88.02 ± 1.26</td>
<td>62.76 ± 17.54</td>
<td>98.81 ± 2.80</td>
<td>102.99 ± 7.52</td>
</tr>
</tbody>
</table>

(100.90%) (105.26%) (109.80%) (78.46%) (98.81%) (102.99%)
Acid resistance

According to the compendial requirement, less than 10% of ASA should be released from enteric-coated formulations after 120 minutes in acid medium.

The amount of ASA release from each ASA product that was tested at the end of acid stage (120 minutes) can be seen in Table 1, and these were less than 10% of the stated potencies. Therefore, all of the tested drugs conformed to the compendial requirement for enteric-coated product. Cardio Aspirin® tablet released the smallest amount of ASA during the acid stage, less than 2% of ASA (1.79%) were released after 120 minutes of dissolution in acid medium. Meanwhile, the other drug products released more ASA which varied from 2.74% to 6.92%, but the differences were not significant.

Dissolution profiles in buffer stage

The results showed that the 6 ASA products had different drug release profiles. In buffer medium, ASA was released immediately from Ascardia® and Farmasal® tablets, but there was a lag time of about 10 minutes for Aptor® and Astika® tablets, and a longer lag time of about 20 minutes for Cardio Aspirin® tablet, which then fastly released ASA after 20 minutes of dissolution (see Figure 1).

A very different release profile was observed for Thrombo Aspilet® tablet, which showed sustained or slow release properties (see Figure 1).

Table 2. Amount (in %) of ASA released at 90 minutes in buffer stage

<table>
<thead>
<tr>
<th>Batch</th>
<th>Cardio Aspirin® (100 mg)</th>
<th>Aptor® (100 mg)</th>
<th>Ascardia® (100 mg)</th>
<th>Th.Aspilet® (100 mg)</th>
<th>Astika® (100 mg)</th>
<th>Farmasal® (100 mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>105.95</td>
<td>106.92</td>
<td>108.88</td>
<td>98.62</td>
<td>100.32</td>
<td>106.20</td>
</tr>
<tr>
<td>2</td>
<td>96.67</td>
<td>106.61</td>
<td>111.82</td>
<td>55.11</td>
<td>100.53</td>
<td>108.38</td>
</tr>
<tr>
<td>3</td>
<td>100.09</td>
<td>102.27</td>
<td>108.69</td>
<td>81.63</td>
<td>95.57</td>
<td>94.41</td>
</tr>
<tr>
<td>Mean</td>
<td>100.90</td>
<td>105.26</td>
<td>109.80</td>
<td>78.46</td>
<td>98.81</td>
<td>102.99</td>
</tr>
</tbody>
</table>
All of the products tested released more than 75% of ASA in 90 minutes of dissolution (compendial requirement), except for one of the 3 batches of Thrombo Aspilet® which showed less than 75% of drug released in 90 minutes (see Table 2).

**Free salicylic acid**

At the end of dissolution test (90 minutes of ASA in buffer phase), the amount of salicylic acid release from ASA was measured. Deacetylation of ASA to salicylic acid causes ASA to lose its activity to acetylate COX-1 enzyme and to produce persistent antiplatelet effect.

The results of salicylic acid assay at the end of dissolution test are presented in Table 3. The amount of salicylic acid found during the dissolution test was smallest with Cardio Aspirin® tablet, followed by Aptor®, Tablet, Farmasal®, Ascardia®, Astika® and Thrombo Aspilet®.

Table 3. The amount of salicylic acid (in %*) observed at the end of the dissolution test

<table>
<thead>
<tr>
<th>Tested product</th>
<th>Cardio Aspirin® (100 mg)</th>
<th>Aptor® (100 mg)</th>
<th>Ascardia® (80 mg)</th>
<th>Th. Aspilet® (80 mg)</th>
<th>Astika® (100 mg)</th>
<th>Farmasal® (100 mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean</td>
<td>3.47</td>
<td>4.72</td>
<td>6.25</td>
<td>11.90</td>
<td>10.90</td>
<td>5.73</td>
</tr>
<tr>
<td>SD</td>
<td>0.85</td>
<td>0.51</td>
<td>0.93</td>
<td>5.36</td>
<td>3.94</td>
<td>1.49</td>
</tr>
</tbody>
</table>

*of total ASA content

**DISCUSSION**

All the investigated low-dose ASA products in this study were enteric-coated formulations, and all of the products conformed to the compendial requirement of releasing ASA less than 10% after 120 minutes in acid medium. Cardio Aspirin® tablet released the lowest amount of ASA (less than 2%), which suggested that it was the most acid resistant compared to the other products (see Table 1), but the differences were not significant.

As enteric-coated products, the ASA should be released in buffer medium, and the compendial requirement requires that more than 75% of ASA should be released in 90 minutes. Again, our data showed that all of the products fulfilled this requirement, except Thrombo Aspilet®, as one of the 3 tested batches released ASA only 55% at the end of the dissolution test (see Table 2).

Enteric-coated low-dose ASA has been shown to have less antiplatelet activity compared to equivalent doses of plain ASA. Lower bioavailability of these preparations and poor absorption from the higher pH environment of the small intestine may result in inadequate platelet inhibition. Enteric-coated ASA preparations release ASA into the upper small intestine, which has a near neutral pH, causing ASA to become unstable and some are deacetylated to salicylic acid. Deacetylation of ASA to salicylic acid causes ASA to lose its activity to acetylate serine 529 at Cox-1 enzyme, which produces irreversible inhibition of the enzyme in platelet, and thereby causes persistent antiplatelet effect. Therefore, deacetylation causes inactivation of ASA in producing persistent antiplatelet effect

Enteric-coated preparations may also differ in their rate of dissolution at the intestinal pH (pH = 6). Furthermore, enteric-coated 75 mg ASA preparation was estimated to deliver a dose equivalent to 50 mg plain ASA. This may be true for enteric-coated 80 mg ASA preparations as was shown in the present study for Ascardia® and Thrombo Aspilet®. To ensure adequate inhibition of thromboxane production, plain 75 mg ASA or an equivalent dose of enteric-coated ASA (estimated to be around 100 mg) should be used. The most effective dose of ASA as an antiplatelet drug for long-term use is 75 to 150 mg. Since the side effects of ASA is dose-dependent, then 75 mg of plain ASA would be the right choice, and this is equivalent to 100 mg of enteric-coated ASA.

Farmasal® is an enteric-coated 100 mg ASA preparation, but it had a very high variability in ASA release between batches, especially at 10, 20, and 30 minutes in buffer medium (the standard deviations were more than 20%) (see Table 1), which might indicate inconsistency in the manufacturing process.

Cardio Aspirin®, Aptor®, and Astika® are 100 mg enteric-coated tablets, and have shown low variabilities of ASA release between batches (Table 1). Aptor® and Astika® had a lag time of about 10 minutes, while Cardio Aspirin® had a lag time of about 20 minutes, before ASA was fastly released afterwards. These facts indicated that the tablets of Aptor® and Astika® required 10 minutes, while Cardio Aspirin® tablet required 20 minutes to dissolve at the near neutral pH of the small intestine. However, Astika® showed a high variability in producing salicylic acid compared to Aptor® and Cardio
Aspirin® at the end of buffer stage (see Table 3), which indicated that it was more unstable in intestinal pH compared to Aptor® and Cardio Aspirin®. Therefore, among these 3 enteric-coated tablets that contain 100 mg ASA, Cardio Aspirin® and Aptor® were the most stable towards deacetylation, and produced the lowest amount of salicylic acid at the end of buffer stage (90 minutes), as shown in Table 3.

Our data also showed that Thrombo Aspilet® had both enteric-coated and sustained-release properties. This sustained-release ASA preparation produced more salicylic acid (see Table 3), may be because it released ASA steadily but in small amounts, and therefore more prone to deacetylation. This instability towards deacetylation was shown in the high variability in producing salicylic acid at the end of the dissolution test, as shown in Table 3. As already mentioned, Thrombo Aspilet® also showed a high variability between batches in ASA release at 90 minutes in buffer stage (see Table 2). These results were in accordance with the study of Dooley et al., which showed that the slow-release 75 mg ASA was worse in terms of variation in platelet aggregation and serum TXB₂ levels in healthy volunteers. Consequently, sustained-release critically low doses of ASA may result in subtherapeutic effects. 9

Enteric-coated ASA may cause rectal bleeding, ulceration of ascending colon, collagenous colitis, and diverticular bleeding from a duration of use ranging from 18 days to 15 years. 10 However, enteric-coated ASA 100 mg/day caused significantly less gastroduodenal damage than the same dose of plain ASA. 4 This is due to enteric-coated ASA that released ASA mainly in the intestine, therefore decreased ulcerogenicity in the upper gastroduodenum and shifted the site of damage to the more distal intestine. 10 Moreover, the sustained-release preparations, due to the sustained release of the drug and its longer presence within the intestinal tract, induced more local damage in the distal intestinal wall in addition to the systemic intestinal effect.10-12 This fact was due to a significant increase in the permeability of the lower intestine produced by the sustained-release formulations, but not of the gastroduodenum, while the immediate-release products significantly increased the permeability at the gastroduodenal level. 11

Nevertheless, the gastrointestinal blood loss with enteric-coated ASA was less compared to that with plain ASA, but more compared with control, 13 which implies that the gastric contribution to gastrointestinal ASA induced blood loss may be more important than the contribution of the small intestine. 13 This reduced GI blood loss with enteric-coated ASA was accompanied by decreased gastroduodenal mucosal damage compared with plain ASA, as measured endoscopically and symptomatically. 14

In short, sustained-release low-dose ASA showed high variability in ASA release between batches and in producing salicylic acid (deacetylation) at the end of the dissolution test, and therefore may be less effective in inhibiting platelet aggregation. It may also cause more local damage in the distal intestine.

In conclusion, all of the investigated low-dose ASA products in this study are enteric-coated formulations and are marketed in Indonesia. Thrombo Aspilet® besides an enteric-coated is also a sustained-release product. Cardio Aspirin®, followed by Aptor®, has the right dose for low-dose enteric-coated preparation (100 mg), produces consistent ASA release between batches, and the most stable towards deacetylation (antiplatelet inactivation).

Acknowledgment

We thank PT Bayer Indonesia for funding the study.

REFERENCES