Hepatoprotective effect of Phytosome Curcumin against paracetamol-induced liver toxicity in mice

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Curcuma longa, which contains curcumin as a major constituent, has been shown many pharmacological effects, but it is limited using in clinical due to low bioavailability. In this study, we developed a phytosome curcumin formulation and evaluated the hepatoprotective effect of phytosome curcumin on paracetamol induced liver damage in mice. Phytosome curcumin (equivalent to curcumin 100 and 200 mg/kg body weight) and curcumin (200 mg/kg body weight) were given by gastrically and toxicity was induced by paracetamol (500 mg/kg) during 7 days. On the final day animals were sacrificed and liver function markers (ALT, AST), hepatic antioxidants (SOD, CAT and GPx) and lipid peroxidation in liver homogenate were estimated. Our data showed that phytosome has stronger hepatoprotective effect compared to curcumin-free. Administration of phytosome curcumin effectively suppressed paracetamol-induced liver injury evidenced by a reduction of lipid peroxidation level, and elevated enzymatic antioxidant activities of superoxide dismutase, catalase, glutathione peroxidase in mice liver tissue. Our study suggests that phytosome curcumin has strong antioxidant activity and potential hepatoprotective effects.


INTRODUCTION

Curcumin is a polyphenol extracted from the rhizomes of Curcuma longa. It has a yellow color, and traditional used in many foods. Curcumin has been shown many beneficial biological activities including antioxidant, anti-inflammatory, anticoagulant, antitumor and hepatoprotective activities (Anand et al., 2008). However, curcumin has a limited to use as a drug to treat the disease because of its poor solubility in water and its low oral bioavailability (Prasad, Tyagi, Aggarwal, 2014; Siviero et al., 2015). Therefore, it is needed to develop a new preparation of curcumin, to enhance the absorption and pharmacological activity.

Some intents to improve the bioavailability of curcumin, such as the using of nanoparticles, liposomes, structural analogues and phospholipid complexes (Gupta, Patchva, Aggarwal, 2013). Phytosome is a technology, which incorporate natural product compound or extract into phospholipids to produce lipid compatible molecular complex and then increase their absorption and bioavailability. Marczylo have prepared a formulation of curcumin with phosphatidylecholine (phytosome curcumin) to increase its oral bioavailability. The author showed that the maximum plasma concentration and area under the plasma concentration time curve values for phytosome curcumin after administration were fivefold higher than the equivalent values seen after curcumin free in rats plasma (Marczylo et al., 2007). In other study, the bioavailability of phytosome curcumin was investigated in human clinical trial. Phytosome curcumin showed the total curcuminoid absorption was 29-fold higher than for its corresponding curcuminoid free. Interestingly, the main plasma curcuminoid after administration of phytosome curcumin was demethoxycurcumin, but not curcumin (Cuomo et al., 2011).

Many study has demonstrated that the main causes of the damage liver is linked to reactive oxygen species (ROS) (Jaeschke, Ramachandran 2011). ROS can attack to polyunsaturated fatty acids in cellular membranes,
the proteins groups and DNA bases. The cells had to
develop efficiency defense systems to prevent damages
with their own antioxidant enzyme system. Paracetamol
has been widely used in model of hepatotoxicity in
mice (McGill et al., 2012). The biochemical changed
with paracetamol toxicity seems to be a significantly
increasing in serum alanine aminotransferase (ALT) and
aspartate aminotransferase (AST) levels and decreasing
the antioxidant enzyme endogenous in liver (Hinson,
Roberts, James, 2010).

In our recent publication, we have prepared saponin-
phospholipids complex to increase the pharmacological
effect of saponin having poor oral absorption (Kim et
al., 2016). Continue studying to improve bioavailability
of natural product, in this paper we aimed to prepare
phytosome curcumin and determine its physicochemical
characteristics and evaluate its hepatoprotective effect in
paracetamol-induced mice, compare with curcumin free.

MATERIAL AND METHODS

Preparation of phytosome curcumin

Phytosome curcumin was prepared by reaction
between curcumin and phosphatidylcholine at different
molar ratios: 1:1; 1:2; 1:4, each ratio was repeated three
times. Weigh exactly 2.04 g curcumin powder and 4.35
g phosphatidylcholine and put to 100 mL round bottom
flask, then added 30 mL of dichloromethane. The mixture
was refluxed at 40 °C with magnetic stirring for 2 hours.
Then solution was evaporated to remove dichloromethane
and added 50 mL n-hexane. The obtained complex was
precipitated, filtered and dried under vacuum to obtain the
phytosome complex.

Morphology and structure of phytosome
curcumin

Using the method of negative staining transmission
electron microscopy (TEM) scanning electron microscope
(SEM).

Determination of curcumin content in the
phytosome curcumin

Standard curve of curcumin concentration
Approximately 100 mg curcumin powder was
dissolved in methanol in 25 mL volumetric flask (4 mg/
ml). This solution was diluted 1000 – 10000x (4 µg/mL-
0.4 µg/mL) and filtered through membrane 0.45 µm and
prepared a standard curve of curcumin by using a HPLC
method. The mobile phase used was water and acetonitrile
gradient as following:

<table>
<thead>
<tr>
<th>Time (min)</th>
<th>% water</th>
<th>% acetonitrile</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.00</td>
<td>70.0</td>
<td>30.0</td>
</tr>
<tr>
<td>20.00</td>
<td>50.0</td>
<td>50.0</td>
</tr>
<tr>
<td>23.00</td>
<td>70.0</td>
<td>30.0</td>
</tr>
</tbody>
</table>

Detector: UV –VIS, 425 nm.
Flow rate: 1.0 mL/ min
Injection volume: 10 µL
Column: Agilent ZORBAX Eclipse Plus 95Å C18, 4.6 x
100 mm, diameter 3.5 µm.

Yield of process

Weight exactly amount of phytosome, curcumin,
disperse in water with 1:100 (w/v). Centrifugate at 8000
rpm during 30 min. Filter, take the solid and dissolve in
methanol. Curcumin’s was concentration was determined by using the HPLC method
as described above.

Infrared (IR) spectroscopy

Infrared spectrum of curcumin, phosphatidylcholine
and phytosome was measured by Fourier transform
infrared spectroscopy (FTIR).

Differential scanning calorimetry (DSC)

The samples were sealed in the aluminum crimp

cell and heated at the speed of 10 °C/min from 0 to 800 °C
in nitrogen atmosphere (60 mL/min). The peak transition
onset temperature of the obtained complexes were
determined and compared with the help of a Mettler DSC
30S (Mettler Toledo, US) (Shyam, Kumar, 2012).
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1H-NMR Spectroscopy

1H-NMR (500 MHz, CDCl3, δ in ppm) spectroscopy of curcumin, phosphatidylcholine, and phytosome curcumin was carried out by using Brucker Advance DRX-500, BruckerBioSciences Corporation, Billerica, MA, USA) at 500 MHz (Maiti et al., 2007).

Particle Size and Zeta Potential analysis

The particle size and zeta potential of phytosome curcumin were determined at 25°C using photon correlation spectroscopy (ZetaSizer Nano-ZS90, Malvern, UK). The analysis consisted of 100 mg of phytosome curcumin powders were dispersed in about 15 mL of double-distilled water before analysis.

Solubility studies

Solubility of curcumin, phytosome curcumin and physical mixture of curcumin and phosphatidylcholine were evaluated by adding excess of the samples to 5 mL of water in glass container at room temperature. The liquids were shaken for 24 h and centrifuged at 5000 rpm for 10 min. The supernatant was filtered by membrane 0.45 μm. Dilute 1 mL with methanol to 10 mL and their curcumin’s concentration were measured by HPLC method (Maiti et al., 2007).

Animals

Fifty (25–30 g) Swiss mice were used to study the hepatoprotective activity of the phytosome curcumin. The animals were kept at 27 ± 2 °C, relative humidity 44–56% and light and dark cycles of 12 h, for 5 days before and during the experiments. Animals were provided with standard diet and water ad libitum. Mice were randomly divided into five groups, each group consisting of ten mice.

Group I received a single daily dose of 1 mL/kg of saline orally (NC group).

Group II was given a single daily dosing of paracetamol (500 mg/kg) orally (PAR group).

Group III was given orally a single daily dose of both 500 mg/kg paracetamol and 200 mg/kg of Curcumin (CUR group).

Group IV was given orally a single daily dose of both 500 mg/kg paracetamol and amount phytosome curcumin equivalent to curcumin 100 mg/kg b.w (Phyt 100 group).

Group V was given orally a single daily dose of both 500 mg/kg paracetamol and amount phytosome curcumin equivalent to curcumin 200 mg/kg b.w (Phyt 200 group).

Phytosome curcumin was administered three hours after the administration of paracetamol. The treatments were continued for seven days and on the final day of the experiment blood were taken from carotid artery of all animals and then mice were sacrificed by cervical dislocation. All liver tissues were dissected, washed in 0.9% NaCl and frozen rapidly at -80 °C. Frozen tissues defrosted, weighted and homogenized in ice-cold lysis buffer, containing 50 mM Tris-HCl (pH 7.5), 8 mM MgCl2, 5 mM ethylene glycol bis (2-aminoethyl ether)-N,N,N’,N’-tetraacetic acid (EGTA), 0.5 mM EDTA, 0.01 mg/mL leupeptin, 0.01 mg/mL pepstatin, 0.01 mg/mL aprotinin, 1 mM phenylmethylsulfonyl fluoride (PMSF) and 250 mM NaCl. Homogenates were then centrifuged at 12000 × g for 10 min at 4 °C. The supernatants was collected and stored until use at -80 °C. Protein concentration was determined by Bradford’s method (Noble, Bailey, 2009).

Hepatotoxicity

Serum levels of ALT and AST as markers of hepatic function were measured by using a ALT Activity Assay Kit and AST Activity Assay Kit (Sigma-Aldrich, Singapore) according to the manufacturer’s instructions.

Lipid peroxidation assay

Lipid peroxidation assay was performed by determining the reaction of malonaldehyde with two molecules of 1-methyl-2-phenylindole at 45°C as described previously (Thanh et al., 2016; Thanh et al., 2015). The reaction mixture consisted of 0.64 mL of 10.3 mM 1-methyl-2-phenylindole, 0.2 mL of sample and 10 µL of 2 µg/mL butylated hydroxytoluene. After vigorously mixing, 0.15 mL of 37% v/v HCl was added. The mixture was incubated at 45°C for 45 min and centrifuged at 10,000 g for 10 min. Cleared supernatant absorbance was recorded at 586 nm. A calibration curve prepared from 1,1,3,3- tetramethoxypropane (Sigma-Aldrich, Singapore) was used for calculation. Peroxidized lipids were expressed as nmol MDA equivalents/mg protein.

Superoxide dismutase (SOD) activity determination

SOD activity was determined as described previously (Thanh et al., 2015). This method is based on the capacity of SOD to inhibit the autoxidation of pyrogallol. Each assay was measured in triplicate.
Catalase (CAT) activity determination

CAT activity was measured in triplicate by monitoring the disappearance of H$_2$O$_2$ at 240 nm as described previously (Thanh et al., 2015). Each assay was measured in triplicate.

Glutathione peroxidase (GPx) activity determination

Glutathione peroxidase (GPx) activity was measured with a coupled enzyme assay described previously (Thanh et al., 2015). Each assay was measured in triplicate.

Statistical analysis

All data are expressed as the mean ± standard deviation (SD). One-way analysis of variance (ANOVA) was used to determine significance among groups. Statistical significance was set at p<0.05.

RESULTS

Physical properties of phytosome curcumin

The resultant complexes after dried under vacuum were kept in petri disk (Figure 1). All phytosomes have orange–yellow powders, but the phytosome with the ratio curcumin and phosphatidylcholine 1:1 were softer and smoother while phytosome with ratio curcumin and phosphatidylcholine 1:2 and 1:4 seemed to be more sticky and clammy. That may be explained by the more presence of phosphatidylcholine in the samples.

Morphology of phytosome curcumin

SEM and TEM images demonstrate the formation of particles. As shown in Figure 2 (A, B) the particles observed under SEM showed spherical-shaped particles. The TEM showed that particles phytosome curcumin are uniforms and homogeneous.

FIGURE 1 - Physical properties of curcumin phytosome at different ratios.

FIGURE 2 - Phytosome curcumin was taken by SEM (A) and TEM (B) of phytosome curcumin: phosphatidylcholine ratio is 1:1.

Yield of phytosome preparation process

Yield of phytosome preparation process was presented in Table I.
Table I showed that total yield of the preparation process increased following the rise of phosphatidylcholine amount added in the complexes. With the curcumin: phosphatidylcholine ratio is 1:2 and 1:4, the yield can be about 94%. However, this yield is only about 81% when the ratio is 1:1. This can be explained by adding the large amount of phosphatidylcholine, the ability of interaction between curcumin and phosphatidylcholine was higher, lead to form more phytosome curcumin complex.

**Determinant of curcumin content in the complex**

The curcumin content in the phytosome was shown in Table II.

**TABLE I - Total yield of phytosome preparation process**

<table>
<thead>
<tr>
<th>Curcumin: phosphatidylcholine ratio</th>
<th>Total yield (w/w, %)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1:1</td>
<td>87.74 ± 1.73</td>
</tr>
<tr>
<td>1:2</td>
<td>91.61 ± 2.24</td>
</tr>
<tr>
<td>1:4</td>
<td>94.64 ± 2.64</td>
</tr>
</tbody>
</table>

The curcumin content in the complex was determined by HPLC method. A HPLC chromatographical of phytosome curcumin with curcumin: phosphatidylcholine ratio (mol:mol) 1:1 is shown in Figure 3. The percentages of curcumin in the phytosome in sample at ratio 1:1 are higher than curcumin in other ratio. The amount of curcumin reached 25.64% in the complexes with curcumin: phospholipis ratio 1:1. However, curcumin content reached 12.61 % and 8.97 % in the complexes with 1:2 and 1:3 ratios, respectively.

**Particles size and zeta potential of phytosome curcumin**

The distribution of the size of phytosome curcumin was shown in Figure 4. The average size of phytosome curcumin was 131.8 nm and PDI was 0.191. The zeta potential of PEG-CUR was -44.5 mV. One important parameter of nanoparticles is the polydispersity index (PDI), which is measured the particle size distribution. If PDI is smaller than 0.1, the particles are typically referred to as “monodisperse” (Moreira, Gaspar, Allen, 2001; Pereira-Lachataignerais et al., 2006; Pham et al., 2015). Our particles of phytosome curcumin are shown to be

**TABLE II - Curcumin content in the phytosome**

<table>
<thead>
<tr>
<th>Sample (curcumin:phosphatidylcholine ratio)</th>
<th>1:1</th>
<th>1:2</th>
<th>1:4</th>
</tr>
</thead>
<tbody>
<tr>
<td>Curcumin content (%)</td>
<td>25.64 ± 2.01</td>
<td>12.61 ± 1.76</td>
<td>8.97 ± 1.25</td>
</tr>
</tbody>
</table>

**FIGURE 3** - Representative chromatograms HPLC chromatographical of Phytosome curcumin with curcumin: phosphatidylcholine ratio (mol:mol) is 1:1.
quite uniforms with PDI was 0.191 and have an average size of 131.8 nm.

The index used to evaluate the stability of particle system is zeta potential value. If the particles possess the absolute value of zeta potential greater than 30 mV, then the particle system is highly stable and able to prevent the aggregation of particles. If zeta potential values are in the range of 20-30 mV, the particle system is relatively stable (Eloy et al., 2014; Pham et al., 2015). Our PEG-CUR has zeta potential values of -44.5 mV, it can be considered they are a relatively stable system.

**Infrared (IR) spectroscopy**

IR spectra of curcumin, phosphatidylcholine and phytosome are shown in Figure 5. The IR spectra of curcumin showed two peaks at λ 1625.99 and 1600.92 cm⁻¹ which represent for C=C and C=O bonds, but disappear

![FIGURE 4](image-url) - Distribution of PEG-CUR size (A) and zeta potential (B).
FIGURE 5 - Infrared spectrum of curcumin, phosphatidylcholine and phytosome curcumin.
in phytosome’s IR spectra. The peak at \( \lambda \) 1734.01 cm\(^{-1}\) presents in both IR spectra of phosphatidylcholine and phytosome curcumin. Moreover, the peak at \( \lambda \) 3506.59 cm\(^{-1}\) of OH group in curcumin structure and two peaks at \( \lambda \) 2916.37 and 2848.86 cm\(^{-1}\) in phosphatidylcholine structure are also presented in phytosome curcumin’s IR spectrum. Interestingly, there is a new peak at \( \lambda \) 3745.76 cm\(^{-1}\) and some fluctuations appearing from 3200 to 4000 cm\(^{-1}\) in phytosome’s IR spectrum. This confirms the presence of hydrogen bonds between curcumin and phosphatidylcholine. Therefore, it is suggested that in the phytosome, curcumin bounds to the polar head of phosphatidylcholine while the non–polar part of phosphatidylcholine still are freely and envelops the polar part containing curcumin molecules.

**Differential scanning calorimetry (DSC)**

Figure 6 reveals the DSC thermographs of curcumin (A), phosphatidylcholine (B) and phytosome curcumin (C). DSC spectrum of curcumin showed the onset temperature of the melting processes is 155.8172°C corresponds to peak at 185.0430 °C which is the curcumin’s melting temperature. Phosphatidylcholine has a melting peak at 149.5276 °C corresponds to the temperature when hydrocarbon tails of phosphatidylcholine transform from gel state to liquid crystalline. DSC of phytosome curcumin do not present the endothermic peak of both curcumin and phosphatidylcholine. The melting peak of curcumin and phosphatidylcholine were completely disappeared. Instead, it showed new melting peaks with lower endothermic effect of phosphatidylcholine (onset temperature 131.4244 °C and melting peak 116.6517 °C). This proves that curcumin reacts with phosphatidylcholine to form the chemical bonds between the OH in phenol group of curcumin structure with phosphatidylcholine’s polar head.

**\(^1\)H NMR spectroscopy**

The proton NMR spectrum of curcumin, phosphatidylcholine and phytosome curcumin was represented in Figure 7. In \(^1\)H-NMR of phytosome curcumin signals at \( \delta \) 0.876, 1.296 and 2.764 ppm showed that the signals of protons of methyl, methylene group of aliphatic side chain and methylene protons linked to –C(=O)–C group, respectively. They are characteristics of nonpolar portion of phosphatidylcholine molecule. The signal of protons at \( \delta \) 5.334 ppm is due to signals of second methylene group proton near N atom (–CH2CH2N) of choline and signals at \( \delta \) 3.362 ppm is due to protons of methyl group attached to N–atom of choline which represented a signal broadening due to their involvement in phytosome curcumin (Sikarwar et al., 2008). The

![Figure 6](image-url)
protons signals at 5.779-5.983; 6.459-6.491; 6.926-6.942; 7.049-7.052; 7.112-7.132 and 7.572-7.604 [m, ArH] are due to aromatic ring of curcumin. These data suggested that phenyl group of curcumin was complexed with choline part of phosphatidylcholine.

**Solubility studies**

Solubility of curcumin, phytosome and physical mixture of curcumin and phosphatidylcholine in water was shown in Table III.

Curcumin, a hydrophobic molecule, is practically insoluble in water, especially at pH 1.2. Therefore, curcumin has very low bioavailability; this is a barrier of using curcumin in clinical application. When pH was increased, curcumin’s solubility tends to increase but still is very low. Solubility of curcumin in n-octanol is also very low. Therefore, bioavailability of curcumin by oral via or derma via is extremely low. Phytosome curcumin can significantly improve solubility of curcumin in solution at different pH and also in n-octanol. Thereby phytosome curcumin can increase the partition coefficient O/W, lead curcumin easily to diffuse into the cell membrane, easily transferred from the aqueous phase to the lipid phase, then increasing its bioavailability.

**Biological activity**

Enzyme such as AST and ALT are main liver transaminases have been used for the assessment of liver damage (Howell et al., 2014).

Serum ALT and AST activities were significantly increased in PAR group as compared with control group (Table IV). When mice were treated with phytosome, the ALT and AST activities were significantly decreased as compared to the PAR group. These enzymes activities tended to decrease in Cur 200 group compared with PAR group.

**Lipid peroxidation**

The levels of lipid peroxidation product (MDA) from the liver tissues in the studied groups were shown in Table V and Figure 8. An increased in MDA level was observed significantly in the PAR group when compared with control group (p<0.05). A decreased in MDA levels were observed significantly in group of Phyt 100 and Phyt 200 (p<0.05). The level of MDA tended to decrease in Cur 200 group compared with PAR group.

**Antioxidant enzyme analysis**

We also determined the activities of several antioxidant enzymes including CAT, SOD and GPx in the liver tissue of control and experimental animals. The results were reported in Table V and Figures 9, 10 and 11. Mice in PAR group showed a significant lower in the activities of these antioxidants compared with control group (p<0.05). Mice in groups Phyt 100 and Phyt 200 exhibited significant increasing in the levels of CAT,
TABLE III - Solubility of curcumin, phytosome and mixture of curcumin and phosphatidylcholine in different medium

<table>
<thead>
<tr>
<th>Sample</th>
<th>Solubility in water (μg/mL)</th>
<th>Solubility in solution HCL 0.1 N (μg/mL)</th>
<th>Solubility in buffer solution phosphate pH 4.5 (μg/mL)</th>
<th>Solubility in buffer solution phosphate pH 6.8 (μg/mL)</th>
<th>Solubility in n-octanol (mg/mL)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Curcumin</td>
<td>8.2 ± 3.1</td>
<td>3.1 ± 2.3</td>
<td>13.3 ± 1.4</td>
<td>21.7 ± 1.4</td>
<td>4.1 ± 3.7</td>
</tr>
<tr>
<td>Mixture of curcumin and phosphatidylcholine (ratio 1:1)</td>
<td>17.7 ± 2.4</td>
<td>10.1 ± 2.4</td>
<td>19.4 ± 2.8</td>
<td>30.1 ± 5.1</td>
<td>8.4 ± 1.6</td>
</tr>
<tr>
<td>Phytosome curcumin (ratio 1:1)</td>
<td>38.7 ± 1.4</td>
<td>20.7 ± 3.5</td>
<td>32.7 ± 4.3</td>
<td>60.1 ± 1.1</td>
<td>44.1 ± 1.8</td>
</tr>
</tbody>
</table>

TABLE IV - Effect of phytosome curcumin on liver marker enzymes

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>PAR</th>
<th>Cur 200</th>
<th>Phyt 100</th>
<th>Phyt 200</th>
</tr>
</thead>
<tbody>
<tr>
<td>AST (IU/L)</td>
<td>38.24 ± 4.23</td>
<td>101.28 ± 11.25*</td>
<td>81.28 ± 12.27</td>
<td>47.25 ± 5.24#</td>
<td>41.25 ± 5.32#</td>
</tr>
<tr>
<td>ALT (IU/L)</td>
<td>31.12 ± 5.21</td>
<td>97.67 ± 11.84*</td>
<td>68.58 ± 14.45</td>
<td>42.15 ± 6.38#</td>
<td>37.17 ± 3.25#</td>
</tr>
</tbody>
</table>

TABLE V - The effect of administration of phytosome curcumin on lipid peroxidation and antioxidant enzymes in mice liver tissue

<table>
<thead>
<tr>
<th>Parameters</th>
<th>Control</th>
<th>PAR</th>
<th>Cur 200</th>
<th>Phyt 100</th>
<th>Phyt 200</th>
</tr>
</thead>
<tbody>
<tr>
<td>MDA</td>
<td>0.52 ± 0.14</td>
<td>1.89 ± 0.15*</td>
<td>1.61 ± 0.13</td>
<td>1.42 ± 0.12#</td>
<td>1.28 ± 0.11#</td>
</tr>
<tr>
<td>SOD</td>
<td>0.573 ± 0.09</td>
<td>0.187 ± 0.08*</td>
<td>0.210 ± 0.07</td>
<td>0.282 ± 0.10#</td>
<td>0.362 ± 0.12#</td>
</tr>
<tr>
<td>CAT</td>
<td>298.87 ± 39.15</td>
<td>89.8 ± 13.19*</td>
<td>129.8 ± 14.21</td>
<td>165.15 ± 21.12#</td>
<td>196.12 ± 23.15#</td>
</tr>
<tr>
<td>GPx</td>
<td>25.35 ± 3.27</td>
<td>10.19 ± 2.32*</td>
<td>11.23 ± 3.18</td>
<td>13.68 ± 2.72</td>
<td>17.86 ± 4.54#</td>
</tr>
</tbody>
</table>

Value represents as mean ± S.D. *p < 0.05, significant difference compared with control group, #p < 0.05, significant difference compared with paracetamol group (n = 10).

SOD and GPx compared with PAR group (p < 0.05). These enzymes activities tended to increase in Cur 200 group compared with PAR group.

DISCUSSION

Preparing new formulation for curcumin delivery is important because of many beneficial effect of curcumin. Curcumin has poor absorption because of its low solubility (Maiti et al., 2007). Phospholipids are now using in many application of drug delivery technology. The advantage of phospholipids is increasing solubilizing property of many natural products. In this study, we have developed successfully phytosome curcumin which can improve the bioavailability of curcumin. The physicochemical data showed that curcumin formed a complex with phosphatidylcholine by hydrogen bonds. We also showed that the phytosome has increased the solubility of curcumin in different medium.

Paracetamol have been used in many study for induce liver damage (Faraghaly, Hussein 2010). Liver is the main organ in body, which metabolizes chemicals and drug. It has been well known that enzyme AST and ALT are markers of hepatocyte damage and the high level of AST and ALT is an important marker for liver injury (McGill, Jaeschke, 2013). In our study, paracetamol significant induced hepatic damage in mice by increasing the level of AST and ALT. Our data are agreed with Nithianantham et al. (2011) they have showed that paracetamol significant increased the ALT, AST, and bilirubin levels in mice.

Phytosome curcumin protects the mice from PAR-induced acute liver injury in vivo. After administration of paracetamol, serum ALT and AST levels in mice were significantly greater than those in control group, and phytosome curcumin could reduce those levels. Our results indicate that phytosome curcumin protects hepatocytes in vivo from damage induced by paracetamol administration. Our data are in line with previous report that curcumin can protect damages in liver caused by paracetamol (Kheradpezhouh et al., 2010).

When reactive oxidative stress attack polyunsaturated...
fatty acids and disrupt the cell membrane. It leads to oxidative lipid and forms MDA, a product of lipid peroxidation. Increasing production of liver MDA observed in our experiments by PAR are in agreement with previous study which reported that PAR increased extracellular MDA level (Boonruamkaew, Chonpathompikunlert et al., 2016). In addition, we have shown phytosome curcumin may diminish the level of MDA in liver tissues. Our data are in line with study of Hatem et al. which showed that hepatic lipid peroxidation level was suppressed by administration of curcumin to paracetamol-treated rats (Farghaly, Hussein 2010).

The organism can develop a mechanism, such as the antioxidant system SOD, CAT and GPx to decrease the damage cause by ROS. SOD, CAT and GPx are...
endogenous enzymatic against ROS. SOD is antioxidant enzyme that converts superoxide anion \(O_2^-\) to \(H_2O_2\). CAT converts \(H_2O_2\) to water and \(O_2\). GPx catalyzes the reduction of \(H_2O_2\) and other peroxides by coupling reduced glutathione (Madrigal-Santillan et al., 2014; Thanh et al., 2015). Our data have showed the activities of these enzymes in PAR group were declined. Interestingly when mice were treated with phytosome curcumin, these enzymes can be reversed significantly.

The hepatoprotective effect of phytosome was significantly higher than curcumin free at the same dose. Free curcumin at the dose of 200 mg/kg only slightly reduced the damage conditions in mice induced by paracetamol. The phytosome at dose equivalent to 100 mg/kg of curcumin showed higher restored the damage of mice liver compared with curcumin free at double dose (200 mg/kg). The increasing hepatoprotective efficacy of phytosome curcumin may be explained by increasing bioavailability of the curcumin.

In summary, this study demonstrates that phytosome curcumin had a strong protective effect against paracetamol-induced acute hepatic damage in mice. The hepatoprotective effect of phytosome curcumin may be explained by increasing levels of antioxidant enzymes and decreasing the lipid peroxidation and liver enzyme on paracetamol-induced damage in mice.

CONFLICTS OF INTERESTS

The authors declare no conflict of interest.

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