IN VITRO STUDY OF URSOLIC ACID COMBINATION FIRST-LINE ANTITUBERCULOSIS DRUGS AGAINST DRUG-SENSITIVE AND DRUG-RESISTANT STRAINS OF MYCOBACTERIUM TUBERCULOSIS

DIAN AYU EKA PITALOKA*, ELIN YULINAH SUKANDAR

ABSTRACT

Objective: The resurgence of tuberculosis (TB) caused by Mycobacterium TB (MTB) is associated with the rapid spread of multidrug-resistant, therefore, the development of new antimycobacterial agents is necessary. The aim of this study was to evaluate the antimycobacterial activity of ursolic acid (UA) when it using alone and combination with TB drugs.

METHODS: MTB H37Rv strain, streptomycin-rifampicin resistant strain, and isoniazid-ethambutol resistant strain were evaluated by susceptibility test using a serial number of UA (25-150 µg/mL). Minimum inhibitory concentration (MIC) was read as minimum concentration of drugs that completely inhibit visible growth of organism. Activities of drug combination of UA with TB drug were determined in Lowenstein-Jensen media by calculating the fractional inhibitory concentration index.

RESULTS: The results showed that MIC of UA was 50 µg/mL against three different strains of MTB. The combination of UA and TB drugs displayed synergistic interaction, and no antagonism result from the combination was observed for strains of MTB.

Conclusion: These results indicate that UA may serve as a promising lead compound for future antimycobacterial drug development.

Keywords: Ursolic acid, Tuberculosis, Drug combination, Susceptibility test.

INTRODUCTION

Tuberculosis (TB) is the world’s second most common cause of death by infectious disease. It has been estimated that one-third of the world’s population is infected with Mycobacterium TB (MTB) and 10% of these people will become active patients during their lifetime [1]. The situation is getting worse day by day owing to the emergence of multidrug-resistant (MDR) TB strains and its association with human immunodeficiency virus or acquired immunodeficiency syndrome renders its control more difficult. In 2014, 400,000 people developed MDR TB in the world and 1.9% of TB cases with MDR-TB in Indonesia [2]. Thus, there is an urgent need to discover new anti-TB agents that are effective in TB treatment not only new synthetic drugs but also natural products are potential sources of new antimycobacterial products.

Ursolic acid (UA) is pentacyclic triterpenoid which considered as ubiquitous triterpenoids in the plant kingdom. It has low solubility in water and high lipophilicity [3]. This triterpene is quite interesting because it demonstrated variety biological effects such as anti-inflammatory, hepatoprotection, antihyperglycemia, and antitumor promotion [4]. Research conducted by López-García et al. [5] shown that UA causes a decrease in the growth and development of MTB in macrophages, stimulates the production of nitric oxide in the early phase, and stimulate to suppress transforming growth factor-alpha (TGF-α) and TGF-β. In addition, dichloromethane extract Duroia macrophylla which contained UA showed a significant reduction of activity in rats induced by MTB H37Rv sensitive strain and MDR strain [6]. Meanwhile, synergistic interaction of UA when it is used combination with the first-line anti-TB drugs is still unclear.

This study investigated the in vitro antimycobacterial activity of UA against three different strains of MTB (H37Rv sensitive strain, streptomycin-rifampicin (RMP) resistant strain, and isoniazid-ethambutol resistant strain) when is used as individual and combination with first-line anti-TB drugs. Minimum inhibitory concentration (MICs) values and fractional inhibitory concentration (FICIs) were used in order to assess the synergistic activity of UA.

METHODS

Reagents and antibiotics

UA was purchased from Bio Basic Canada Inc. Markham Ontario, Canada. It was dissolved in dimethyl sulfoxide (DMSO) (Merck) as a solvent. The antibiotics RMP, isoniazid (INH), streptomycin (STR), and ethambutol (EMB) were purchased from Sigma-Aldrich. INH, STR, and EMB were prepared in deionized water, and RMP was prepared in dimethylformamide (Merck). DMSO was used as an appropriate solvent control with the concentration of 0.5% v/v [7].

Microorganism

The antimycobacterial activity was evaluated using MTB H37Rv sensitive strain, streptomycin-RMP resistant strain, and isoniazid-ethambutol resistant strain. All bacterial strains were provided from Balai Pengembangan Laboratorium Kesehatan Provinsi Jawa Barat. Bacterial strains were maintained by subculture on Ogawa 3% favorable media. The antimycobacterial activities of UA and INH, RMP, STR, and EMB against three different type of MTB were tested by susceptibility test using proportion method in Lowenstein-Jensen (LJ) media. The inoculum was adjusted to 3×10⁴ cfu/mL by comparison with Mc. Farland no.1 turbidity standard and 100 µl bacterial suspension containing approximately 3×10⁴ cfu/mL were spotted onto LJ media in Mc Cartney tubes. Cultivation was done at 37°C for 8 weeks, and MICs were read as minimum concentrations of drugs completely inhibiting visible growth of the organism.

Activities of drug combinations

The effects of combinations of UA and anti-TB drugs (STR, RMP, INH, or EMB) were determined as described as determination of MICs. FICIs, 1/2
and \( \frac{1}{2} \) of MIC UA and anti-TB drugs were combined and tested in LJ media using susceptibility test. Interpretation data were achieved by calculating the FICIs as described by De Logu et al. [6]

\[
FICI = \frac{MICA_{combination}}{MICA_{alone}} + \frac{MIC_{combination}}{MIC_{alone}}
\]

Interpretation data of FICI of the results as follows: FICI ≤0.5, synergism; FICI 0.5–4, no interaction; FICI >4.0, antagonism.

**RESULTS**

**Antimycobacterial susceptibility in vitro**

The MICs of UA was determined according to susceptibility test using proportion method as described in methods. MIC of MTB in different strains is showed in Table 1. According to that data, it can be shown that UA can inhibit the growth of MTB in all different strains at concentration 50 µg/mL.

STR, INH, RMP, and EMB were used as drug control in determining MIC of UA and anti-TB drugs. The antimycobacterial susceptibility test results showed that H37Rv strain is sensitive to all anti-TB drugs used in this study. RMP-streptomycin resistant strain was found to be resistant to both RMP and STR, and isoniazid-ethambutol resistant strain was showed to be resistant to both INH and EMB.

**Combination testing**

The results of the combination of UA with STR, INH, RMP, EMB, or all first-line drugs of anti-TB are listed in Table 2. The MIC for anti-TB drugs used in this combination study are streptomycin 4 µg/mL, isoniazid 0.2 µg/mL, RMP 40 µg/mL, and ethambutol 2 µg/mL. The combination of UA and STR, INH, RMP, EMB, and all first-line drugs of TB showed synergistic activity with FICI values ≤0.5. No interaction was found when UA in combination with STR or RMP against RMP-streptomycin resistant strains but the synergistic effect was showed when UA combined with INH, EMB, or all first-line drugs. No interaction result was also found when UA combined with INH and EMB against isoniazid-ethambutol resistant strains, but the synergistic interaction was also showed as the result of the combination UA with STR, RMP, and all first-line drugs. The MICs of UA against drug-sensitive strain and drug-resistant strain are significantly decreased ranging from 6.25 to 25 µg/mL and also the MICs of anti-TB drugs, respectively. The FICI values were showed ranging from 0.25 to 0.5 and some of the combination found to have FICI values of 1. Noticeably, no antagonism result from the combination of UA with anti-TB drugs observed for any of the strains tested.

**DISCUSSION**

Until now, the treatment for this infection needs three to five different drugs simultaneously, depending on the patient category. These anti-TB drugs can be given as single-drug formulations or as fixed-dose combination formulations where two or more anti-TB drugs are present in fixed proportions in the same formulation [6]. Each of the anti-TB drugs has a major role in killing of MTB. For example, INH is the most widely used first-line anti-TB drugs. This bactericidal activity has critical early in therapy which rapidly reduces the sputum viable count because it is active mainly against the organisms growing aerobically in pulmonary cavities [9]. RMP is important in killing organisms that are metabolizing slowly, killing the persisters, and sterilizing the patient’s sputum [10]. EMB is the first-line drug that is used in combination with INH and RMP EMB is a bacteriostatic that interferes the biosynthesis of cell wall arabinogalactan [11]. Streptomycin is an alternative first-line anti-TB drug recommended by WHO. This aminocyclitol glycoside has been demonstrated to take place at the ribosomal level [12].

Treatment of drug resistant in MTB has become a major problem worldwide and more serious because of the resistant phenomena. The cell wall hydrophobicity of MTB naturally prevents diffusion and binding of many drugs. Moreover, this mycobacterium has many efflux pumps which contribute to expel the drugs rapidly before they enter to the site action [13]. Based on that phenomena, development of new antimycobacterial agent becomes necessary. UA is pentacyclic triterpenes that widely found in several plants and reported to have many activities. Several studies showed that pentacyclic triterpenes can be responsible for the antimycobacterial activity.

UA (3β-hydroxy-urs-12-en-28-oic acid) is an isomer from oleanolic acid (OA) (3β-hydroxy-olean-12-en-28-oic acid) which differs in the carbon skeleton [14]. Study conducted by Wójciak-Kosior [15] reported that these compounds have similar cell wall hydrophobicity of MTB naturally prevents diffusion and binding of many drugs. Moreover, this mycobacterium has many efflux pumps which contribute to expel the drugs rapidly before they enter to the site action [13]. Based on that phenomena, development of new antimycobacterial agent becomes necessary. UA is pentacyclic triterpenes that widely found in several plants and reported to have many activities. Several studies showed that pentacyclic triterpenes can be responsible for the antimycobacterial activity.

**Table 1: MICs of UA against drug-sensitive and drug-resistant strains of MtB**

<table>
<thead>
<tr>
<th>Properties</th>
<th>Strain of Mycobacterium tuberculosis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>MIC±standard deviation (µg/mL)</td>
</tr>
<tr>
<td></td>
<td>H37Rv sensitive</td>
</tr>
<tr>
<td>UA</td>
<td>50±0</td>
</tr>
</tbody>
</table>

SD: Standard deviation; in vitro MIC values are the means±standard deviation, n=3; MIC: Minimum inhibitory concentration, UA: Ursolic acid

**Table 2: Combination testing of UA and antituberculosis drugs against different strains of Mycobacterium tuberculosis**

<table>
<thead>
<tr>
<th>Drug combination</th>
<th>Status of strains</th>
<th>Combination MIC±standard deviation (µg/mL)</th>
<th>FICI</th>
<th>Conclusion</th>
</tr>
</thead>
<tbody>
<tr>
<td>UA/STR</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.5±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>RS&lt;sup&gt;S&lt;/sup&gt;</td>
<td>25/2±0</td>
<td>1</td>
<td>N</td>
</tr>
<tr>
<td>UA/INH</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.5±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>RS&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.025±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>HE&lt;sup&gt;H&lt;/sup&gt;</td>
<td>25/0.1±0</td>
<td>1</td>
<td>N</td>
</tr>
<tr>
<td>UA/RMP</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/5±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>RS&lt;sup&gt;S&lt;/sup&gt;</td>
<td>25/20±0</td>
<td>1</td>
<td>N</td>
</tr>
<tr>
<td></td>
<td>HE&lt;sup&gt;H&lt;/sup&gt;</td>
<td>12.5/10±0</td>
<td>0.5</td>
<td>Sy</td>
</tr>
<tr>
<td>UA/EMB</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>12.5/0.5±0</td>
<td>0.5</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>RS&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.25±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td></td>
<td>HE&lt;sup&gt;H&lt;/sup&gt;</td>
<td>25/1±0</td>
<td>1</td>
<td>N</td>
</tr>
<tr>
<td>UA/STR/INH</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.5/0.025/5/0.25±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
<tr>
<td>UA/STR/RMP</td>
<td>H37Rv&lt;sup&gt;S&lt;/sup&gt;</td>
<td>6.25/0.5/0.025/5/0.25±0</td>
<td>0.25</td>
<td>Sy</td>
</tr>
</tbody>
</table>

SD: Standard deviation; in vitro MIC values are the means±standard deviation, n=3; S: Sensitive, R: Resistant, RS: Rifampicin-streptomycin, HE: Isoniazid-ethambutol, N: No interaction, Sy: Synergistic, STR: Streptomycin, EMB: Ethambutol, INH: Isoniazid, RMP: Rifampicin, UA: Ursolic acid, MIC: Minimum inhibitory concentration

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RF values which make it difficult to separate in medicinal plants. In addition, UA and OA also have similar chemical structure that makes them have similar characteristics in pharmacology properties [16,17].

Many studies reported that UA and OA can inhibit the growth and survival of many bacteria, including pathogen one. Study literature data reported that OA, other pentacyclic triterpenes, has a synergistic effect when it combined with INH, RMP, or EMB [18]. Another study showed that UA and OA found to have synergistic activity. Intracellular activity of these two compounds can inhibit the growth of mycobacteria at low doses [16].

The synergistic effects from UA and anti-TB drugs indicated that the antimycobacterial combinations were more effective than the activity of the individual agents. The decrease of MIC value resulting from the combinations indicated the improved bactericidal potentials of UA and the anti-TB drugs as combined antimycobacterial agents. The high lipophilicity of pentacyclic triterpenes is probably the main factor that allows their penetration through the mycobacterial cell wall [18,19]. Acids from UA affected cell morphology and enhanced autolysis of the bacterial cells. UA also can inhibit peptidoglycan turnover and affected the profile of muropeptides after digestion of peptidoglycan with mutanolysin [20]. Another study showed that the mechanism of antimicrobial action of UA has been investigated and found to be based on the inhibition of DNA polymerase [21]. In addition, triterpenes UA was found relatively non-toxic [13].

CONCLUSION

From this study, we can see that combination of UA and first-line anti-TB drugs showed significant potential for the development of new antimycobacterial agent and reduction of drug resistance. Reducing the dose of STR, INH, RMP, and EMB as the result of combining them with UA hopefully can reduce the side effect caused by these anti-TB drugs. These results are preliminary but are of sufficient interest to warrant further study and indicate that UA is worthy of further investigation as a template for the development of novel antimycobacterial compounds. More study are needed on molecular basis of the synergistic interaction to understand the synergistic mechanism. For further study, we will evaluate the mechanism of action in docking molecular and in vivo testing of drug efficacy using genetic tools against MTB.

REFERENCES