Assessment of Immunostimulating effect of Carica papaya-based Immunostimulatory Complexes Against Streptococcus pneumoniae Infections

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ABSTRACT

Objectives: Streptococcus pneumoniae infection is a bacterial infection responsible for one-fifth of child mortality in Africa. The aim of this study was to formulate and evaluate Carica papaya-based Immunostimulatory complexes against Streptococcus pneumoniae infections. Methods: Leaves of the plant were harvested, air-dried, pulverized and the saponin content extracted using distilled water and but anole in a Soxhlet’s apparatus. The formulation of Immunostimulatory complexes was prepared using the ethanol injection method. Acute toxicity test and Immunological assays, namely: neutrophil adhesion, carbon clearance, hemagglutinating antibody titers were conducted using healthy laboratory mice. The results were compared to standard drug (levamisole). Results: The aqueous extract and butanol fraction yielded 7.45% and 0.38% saponin. At 5000mg/kg body weight of administered immunostimulatory complexes, no death or behavioral changes were recorded. Neutrophil adhesion showed no significant increase (20.8±4.34) as against the levamisole (20.7±3.31). The increased phagocytic index (0.089±0.017) showed better stimulation of reticuloendothelial system when compared to both controls (distilled water = 0.032 ± 0.001, levamisole = 0.071 ± 0.001). The lower Hemagglutination antibody titre value indicates the potency of the formulated ISCOM to elicit an immune response. The ISCOM dose (250 mg/kg body weight) significantly elevated hemagglutination antibody titre (7.81±1.60).

Conclusion: It can be inferred from this study that Carica papaya leaves has immunomodulatory capacity and has significant saponin yield which could potentially serve as a component of Immunostimulatory complexes formulation. Carica papaya-based Immunostimulatory complexes exhibit improved immunogenicity against Streptococcus pneumoniae infections.

Key words: Immunity, Adjuvant, ISCOMs, Streptococcus pneumoniae, Carica papaya, Saponin, ISCOMATRIX.

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INTRODUCTION

Papaya (Carica papaya) is a large succulent herbaceous plant that has medicinal and nutritional properties. It is considered a nutraceutical fruit because of its several medicinal qualities such as wound healing, anti-fungal and anti-bacterial activities. Phytochemical analysis reveals the presence of papain (enzyme), cardiac glycosides, lycopene, carotenoids, alkaloids, flavonoids and vitamins. The presence of saponin in the leaves confers cytotoxic effects on the plant. Papaya leaf green tea is taken to aid digestion and treat ailments like chronic indigestion, overweight and obesity and cardiovascular associated health problems. Streptococcus pneumoniae or pneumococcus, a Gram-positive cocci bacterium, is alpha-haemolytic under aerobic conditions or beta-haemolytic under anaerobic conditions. It is a facultative anaerobic member of the genus Streptococcus. Streptococcus pneumoniae is the most common cause of pneumonia and invasive and non-invasive mucosal diseases. Streptococcus pneumoniae infections are responsible for one-fifth of child mortality in Africa. The bacterium colonizes the nasopharynx and is responsible for bacteraemia, pneumonia, meningitis, otitis media. Although pneumococcal diseases can occur in all ages, they are prone to occur in elderly, infants, cancer patients with compromised immune system. The immune system is made up of a complex network of cells with the ability to protect the body from the harmful effect of pathogens and toxic substances. This pathogens or toxic substances include toxins, bacteria, fungi, parasites, prions, viruses. The immune cells are located in various parts of the body like the mucosa of the internal organs, eyes, skin, bone marrow and blood. They are specialised aggregate cells that ward off and fight pathogens. Adaptive and Innate immunity are components of the immune system that mobilizes broad collection of responses against pathogens and have the ability to distinguish body tissues from foreign agents. Phagocytic mechanism is employed by the immune system to combat pathogens by engulfment and digestion. Immune cells such as neutrophils, macrophages, mast cells and monocytes are responsible for phagocytosis. Immune boosters or immune stimulants are often used to improve resistance against infections in the body and can respond via innate and adaptive immune responses. Immune stimulants may be biological or synthetic in nature. In immunocompromised...
individuals, they tend to be curative as therapeutic agents while as prophylactic and boosters in healthy subjects.\textsuperscript{14}

The immunostimulating effect of vaccines as a means of combating infectious diseases have drastically reduced the rate of morbidity and mortality globally. Vaccines combat the effect of pathogens by stimulating protective long lasting immune responses.\textsuperscript{15} The need for improvement of the immunogenicity and efficacy of vaccine leads to the use of adjuvants. Adjuvants are immune boosters that respond to the effect of a particular antigen. Immunostimulating complexes (ISCOMs) are adjuvants that are made up of Saponin, Cholesterol and Phospholipid. ISCOMs are “adjuvant with multiple adjuvant properties” that are “open cage-like complexes typically with a diameter of about 40 nm”,\textsuperscript{16,17} and have been demonstrated to promote antibody responses and induce T helper cell as well as cytotoxic T lymphocyte responses.\textsuperscript{16,17,23}

They can function both as vaccine adjuvants and vaccine delivery system. ISCOMs are unique in their ability to provoke a wide range of protective B-cell and T-cell responses to protein antigens following either oral or parenteral vaccination. “The adjuvant properties of ISCOM-matrix fulfill the demands of a modern adjuvant: a formulation with dose-sparing capacity, improved quality of immune responses and an acceptable safety profile”.\textsuperscript{17}

The immense benefits of currently available pneumococcal vaccines, notwithstanding, the discovery and implementation of additional vaccine platforms is of uttermost importance due to the inherent shortcomings of those available pneumococcal vaccines. To improve on this limitation, \textit{Carica papaya}-based ISCOM-matrix with its unique ability to provoke long-lasting protective immune responses was formulated and evaluated for improved immunogenicity against \textit{Streptococcus pneumoniae} infections.

\textbf{MATERIALS AND METHODS}

\textbf{Plant material}

The \textit{Carica papaya} leaves were collected from Afikpo, Ebonyi State, Nigeria, in August 2016 and identified at the Department of Pharmacognosy, Faculty of Pharmaceutical Sciences; Agulu of Nnamdi Azikiwe University in Anambra State. The leaves were air dried in a cool dry place away from sun and pulverized to powder. Two (2) kg of the grounded \textit{C. papaya} leaves was soaked in ethanol at a ratio of 1:5 for 2 days and then filtered through a muslin cloth. The residue was discarded while the filtrate was concentrated through evaporation technique under reduced pressure at 45°C using a rotary evaporator.

\textbf{Extraction and fractionation to obtain saponin}

Extraction and fractionation method as described by Okigbo \textit{et al.}\textsuperscript{1} was used to extract the saponin from the \textit{C. papaya} leaves. The total yield of \textit{C. papaya} extract was calculated using the formula:

\[
\text{Extract yield (\%) = } \left( \frac{\text{Extracted Material}}{\text{Dried Material}} \right) \times 100
\]

\textbf{Experimental animals and test organisms}

The weight range of the mice (\textit{Mus musculus}) used for the study was 20-30g (male and female gender) and aged 5-6 weeks old. They were sourced from the Department of Zoology, University of Nigeria Nsukka and fed for a week under pathogen free condition before being used for experiments. The animals were handled according to established guidelines.\textsuperscript{18-20} while the ethical approval was obtained from the Ethics Committee of the Faculty of Pharmaceutical Sciences, Nnamdi Azikiwe University Awka Nigeria. Used animals were euthanized using isoflurane and buried after death.

Clinical isolates of \textit{Streptococcus pneumoniae} were obtained from stock samples in the Laboratory of the Department of Pharmaceutical Microbiology and Biotechnology, Faculty of Pharmaceutical Sciences, Nnamdi Azikiwe University Awka, Anambra State, Nigeria. The specimens were re-activated by incubating in nutrient broth at 37°C for 24 hr and sub-cultured onto sterile Blood Agar. The plates were incubated at 37°C with 5-10% CO₂ for 24-48 hr and observed for growth after which Gram staining and biochemical tests were carried out for confirmation.

\textbf{Preparation of ISCOMATRIX}

The ISCOMATRIX adjuvant was formulated using Ethanol injection method (Direct and Reverse) as reported previously.\textsuperscript{21,22}

\textbf{Acute toxicity study}

Hybrid methods of Lorke’s\textsuperscript{23} and Salgal \textit{et al.}\textsuperscript{24} were used. The method involves two phases consisting of a total of seventeen animals. The first phase consisted of nine animals divided into three groups of three animals each. Each group of animal were administered different doses of 10, 100 and 1000mg/kg of the test substance. The animals were afterwards observed for 24 hr to check behavioural characteristics and death. The second phase involved the use of eight mice which were distributed into four groups of two animals each. The animals were administered 2000, 3000, 4000 and 5000 mg/kg doses of test substance and then observed for 24 hr for behavioural changes and mortality.

\textbf{Immunological evaluation}

\textit{Neutrophil adhesion tests} Mallwarar \textit{et al.}\textsuperscript{25} method was used with slight modifications. The mice were grouped into six of five animals per group. Group 1 mice were administered distilled water only (10 ml), Group 2 were administered 100 mg/kg of levamisole which is the positive control, Group 3 were administered 250 mg/kg of test substance – the formulated ISCOM, Group 4 were administered 0.1ml of \textit{Streptococcus pneumoniae} suspension containing 0.5x10\textsuperscript{7}/cells only which serve as negative control, Group 5 were administered 0.1ml of \textit{Streptococcus pneumoniae} suspension containing 0.5x10\textsuperscript{7}/cells then followed by 250 mg/kg of ISCOM, Group 6 were administered 250 mg/kg then followed with 0.1ml of \textit{Streptococcus pneumoniae} suspension containing 0.5x10\textsuperscript{7}/cells. Blood samples were collected from all groups on the 14th day of treatment by puncturing the retro-orbital plexus under mild isoflurane anaesthesia. Blood was collected in pre-treated disodium EDTA vials and analysed for total leucocyte count. The blood sample drawn was diluted with Turk’s fluid in WBC pipette, in which red cells were lysed without affecting the leucocyte population. Leucocytes were counted using improved Neubauer’s counting chamber.

In differential leucocyte count (DLC), blood smear was prepared on a clean glass slide and stained with Leishman’s and field 1 stains. The differential populations of leucocytes were differentiated and identified based on the cell size, presence of granules, colour and shape of nucleus under the microscope using immersion oil. After initial counts, blood samples were inoculated with 80 mg/ml of nylon fibre for 15 min at 37°C. The incubated blood samples were analysed for Total Leucocyte Count (TLC) and DLC. The product of TLC and % neutrophil gives neutrophil index (NI) of blood sample. Percentage of neutrophil adhesion was calculated as

\[
\text{Neutrophil Adhesion (\%) = } \left( \frac{N_{LU}-N_{NI}}{N_{LU}} \right) \times 100
\]

Where \(N_{LU}\) is neutrophil index of untreated blood samples, \(N_{NI}\) is neutrophil index of treated blood samples.

\textit{Carbon clearance assay}: Carbon Clearance Assay was done according to the method described by Cheng \textit{et al.}\textsuperscript{26} Mice were divided into six
groups of five animals per group. Group 1 mice were administered distilled water only (10 ml), Group 2 were administered (100 mg/kg) of levamisole which is the positive control, Group 3 were administered (250 mg/kg) of test substance ISCOM, group 4 were administered 0.1 ml of Streptococcus pneumoniae suspension containing 0.5x10⁷/cells only which serve as negative control, Group 5 were administered 0.1 ml of Streptococcus pneumoniae suspension containing 0.5x10⁷/cells then followed by 250 mg/kg of ISCOM, Group 6 were administered 250 mg/kg then followed with 0.1 ml of Streptococcus pneumoniae suspension containing 0.5x10⁷/cells. On the 7th day of treatment, mice of all groups received an intravenous injection of 0.5 ml Indian ink solution (pre-warmed at 37°C). Blood samples were then collected from retro-orbital bleeding using glass capillaries at an interval of 3 and 10 min after injection of ink dispersion. 50 µl blood samples were added to 4 ml of 0.1 % sodium carbonate solution to lyse the erythrocytes and absorbance of these samples was measured at 675 nm using spectrophotometer, after 10 min. Rates of carbon clearance (K) and phagocytic index were calculated using the formula:

\[ K = \frac{\log OD_t - \log OD_0}{T_2 - T_1} \]  

\[ OD_t \] is the absorbance of blood at 3 min, \( OD_0 \) is the absorbance of blood at 10 min, \( T_2 \) is the last time point of blood collection and \( T_1 \) is the first time point of blood collection.

Rate of carbon clearance and phagocytic index of treated group mice was compared with the control group animals.

**Hemagglutinating antibody (HA) titre:** Hemagglutinating antibody titre was determined as reported by earlier researchers. The animals were challenged by injecting 0.1 ml of Streptococcus pneumoniae suspension containing 0.5x10⁷/cells intra-peritoneal on day 0. The test ISCOM was administered separately to all mice continuously for 7 days. Blood samples were collected in micro-centrifuge tubes from individual mice by retro-orbital puncture on 7th day. The blood samples were centrifuged at 2500 rpm for 10 min to obtain the serum. Antibody levels were determined by the hemagglutination technique. Equal volumes of individual serum samples of each group were pooled and two fold serial dilutions of samples made in 25 µl volume of normal saline in micro-titre plates with 25 µl of 1 % suspension of Streptococcus pneumoniae in saline. After mixing, the plates were then incubated at 37°C for 1 hr and examined for hemagglutination. The reciprocal of the highest dilution of the test serum showing agglutination was taken as the antibody titre.

**Statistical analysis**

Statistical analysis was performed using Graph pad Prism Software Version 5.0. All the results were expressed as Mean ± Standard Error Mean (SEM). Data were analysed using one-way Analysis of Variance (ANOVA) followed by Dennett Multiple comparison test. \( P \)-values < 0.05 were considered significant.

**RESULTS**

Percentage yields of Carica papaya extract

Table 1 shows percentage yields/composition of saponin crude aqueous extract and butanol fraction of C. papaya. Plants were extracted with aqueous or butanol and the composition of saponins in the extracts calculated as a percentage of the dried leaves.

**Isocomatrix formulation**

The ISCOM formulated using the rapid injection method had clear colloidal suspension with a foamy top layer in appearance and lightly yellow in colour, while ISCOM formulated using the reverse rapid injection method had a cloudy colloidal suspension with foamy top layer a yellow colouration.

**Acute toxicity study**

This was done to evaluate the toxicity and behavioural effects of the C. papaya-based ISCOMATRIX on animal models. The ISCOM administered at several doses of 10, 100, 1000, to 5000 mg/kg body weight in two phases of toxicity study showed no lethality after 7 days of treatment. This suggests that the LD₅₀ of the ISCOM is above 5000 mg/kg body weight. Table 2). In addition, no changes in behaviour were observed in the animals at the highest dose of 500mg/Kg body weight tested after 24 hr observation.

**Table 1: Percentage yields of crude aqueous extract and butanol fraction.**

<table>
<thead>
<tr>
<th>Solvents</th>
<th>Percentage yield (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Aqueous</td>
<td>7.45</td>
</tr>
<tr>
<td>Butanol</td>
<td>0.38</td>
</tr>
</tbody>
</table>

**Figure 1:** Index of phagocytic activity in the carbon clearance. (A=Distilled Water (10ml/kg); B= Levamisole (100mg/kg); C= ISCOM only (250mg/kg); D= Organism only (0.1ml/kg); E= Organism+ISCOM (0.1 ml/kg+250mg/kg) while F= ISCOM+Organism (250mg/kg+0.1ml/kg). Data are presented as means ± SEM (n=3). *p<0.05; ***p<0.01 compared to organism only.

**Figure 2:** Hemagglutinating Antibody Titre. (A=Distilled Water (10ml/kg); B= Levamisole (100mg/kg); C= ISCOM only (250mg/kg); D= Organism only (0.1ml/kg); E= Organism+ISCOM (0.1 ml/kg+250mg/kg) while F= ISCOM+Organism (250mg/kg+0.1ml/kg)).
Table 2: LD<sub>50</sub> of the formulated ISCOM.

<table>
<thead>
<tr>
<th>Phases</th>
<th>Treatment Groups</th>
<th>Number of Animals</th>
<th>Doses administered (mg/Kg)</th>
<th>Number of Deaths</th>
<th>Behavioural Changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>A</td>
<td>3</td>
<td>10</td>
<td>0/3</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td>B</td>
<td>3</td>
<td>100</td>
<td>0/3</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td>C</td>
<td>3</td>
<td>1,000</td>
<td>0/3</td>
<td>Normal</td>
</tr>
<tr>
<td>2</td>
<td>D</td>
<td>2</td>
<td>2,000</td>
<td>0/2</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td>E</td>
<td>2</td>
<td>3,000</td>
<td>0/2</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td>F</td>
<td>2</td>
<td>4,000</td>
<td>0/2</td>
<td>Normal</td>
</tr>
<tr>
<td></td>
<td>G</td>
<td>2</td>
<td>5,000</td>
<td>0/2</td>
<td>Normal</td>
</tr>
</tbody>
</table>

Table 3: The effect of the C. papaya-based ISCOM on neutrophil activation.

<table>
<thead>
<tr>
<th>Treatment Groups</th>
<th>Dose (mg/Kg)</th>
<th>Total Leucocyte Count (103/mm3)</th>
<th>Neutrophil %</th>
<th>Neutrophil Index</th>
<th>Neutrophil Adhesion (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Untreated Blood</td>
<td>Fiber Treated Blood</td>
<td>Untreated Blood</td>
<td>Fiber Treated Blood</td>
<td>Untreated Blood</td>
</tr>
<tr>
<td>Distilled Water</td>
<td>10</td>
<td>6.70±0.27</td>
<td>6.57±0.15</td>
<td>40.67±2.60</td>
<td>37.67±3.76</td>
</tr>
<tr>
<td>Levamisole (Positive Control)</td>
<td>100</td>
<td>8.70±0.25</td>
<td>8.60±0.25</td>
<td>51.67±1.45</td>
<td>41.33±0.67</td>
</tr>
<tr>
<td>ISCOM alone</td>
<td>250</td>
<td>9.70±0.21</td>
<td>9.10±0.21</td>
<td>52.33±2.03</td>
<td>44.00±1.16</td>
</tr>
<tr>
<td>Organism (Negative Control)</td>
<td>0.1</td>
<td>4.27±0.33</td>
<td>3.60±0.12</td>
<td>32.67±1.76</td>
<td>29.00±3.22</td>
</tr>
<tr>
<td>Organism then ISCOM</td>
<td>0.1 + 250</td>
<td>6.00±0.23</td>
<td>5.77±0.19</td>
<td>38.33±1.45</td>
<td>35.33±0.67</td>
</tr>
<tr>
<td>ISCOM then Organism</td>
<td>250 + 0.1</td>
<td>8.33±0.18</td>
<td>7.93±0.20</td>
<td>48.00±1.16</td>
<td>42.33±1.45</td>
</tr>
</tbody>
</table>

Immunological evaluation

Neutrophil adhesion test: Table 3 shows the effect of the C. papaya-based ISCOM on neutrophil activation. Treatment with ISCOM resulted in an increase in neutrophil adhesion compared to water treated control animals. However, neutrophil adhesion decreased significantly when the animals were treated with organism. This decrease was significantly reversed when the organism-treated animals were supplemented with ISCOM derived fraction at 250 mg/kg. Animals were prepared and described in the experimental section on immunological evaluation. The neutrophil index calculated from total leucocyte and neutrophils counts was used to determine the percent neutrophil adhesion. The neutrophil index was calculated for the treatment groups and analysed for statistical difference in GraphPad PRISM. Data are presented as means ±SEM obtained from 3 experiments (*p<0.05; or ** p<0.001) compared to distilled water.

Carbon clearance assay: To demonstrate the phagocytic activity using ISCOM, groups treated with formulated ISCOM showed concentration dependent phagocytic activity when compared with control groups. The phagocytic indices of 250mg/kg of ISCOM showed significant (p<0.01; p<0.001) increase as presented in Figure 1. The result showed that ISCOM exhibited more potent phagocytic activity when compared to the control group.

Hemagglutination antibody titre: To demonstrate antibody titre activity of ISCOM, administration of ISCOM produced significant (p< 0.01) humoral antibody titre as compared to control (Figure 2). Animals were treated as indicated in Figure 2. Data are presented as means ± SEM from 4 experiments (***p<0.001 while **p<0.0001) compared to organism only. The lower titre value indicates the potency of the formulated ISCOM to elicit an immune response. The ISCOM dose (250 mg/kg body weight) significantly elevated hemagglutination antibody titre (7.81±1.60).

DISCUSSION

In medical science, immune stimulation is an area wherein extensive studies have been conducted to devise methods to improve disease resistance as well as prevent or control immune disorders of host by optimum regulation of the immune system. Today, most infectious diseases of man are treated and controlled mainly by using broad-spectrum antibiotics and vaccines. However, the antibacterial agents are becoming increasingly ineffective due to rapid emergence of resistant microbial strains. So, there is urgent need for novel and improved alternative therapeutic and prophylactic strategies to manage several immune-related diseases. Immunostimulation is focused on manipulation of immune system to control the infections and other adverse health effects with precise regulation to avoid any complications while suppressive or potentiating efforts are made to benefit the human health. The main aim of this study was to explore the potential immunostimulatory effect of C. papaya-based ISCOM.

The use of ethnomedicinal plants as immunostimulatory agents has been more pronounced recently as plant extracts have widely been explored during last few decades in various parts of the world for their possible immunostimulatory properties. Many studies have demonstrated the isolation of potential bioactive molecules which have been patented and have been tested as herbal formulations. This study shows the percentage yield of aqueous extraction of saponin from C. papaya leaves as 7.45 %. The saponin content of C. papaya leaves have been established by several other studies as responsible for the bitter taste of the leaves. ISCOM was formulated using ethanol injection method due to its low percentage yield of aqueous extraction of saponin from C. papaya leaves as 7.45 %.
cost, simplicity, short time of preparation and high yield of homogenous ISCOM.21

In the acute toxicity study and at 5000 mg/kg body weight, no death was recorded and behavioural characteristics were normal. This supports the findings of Sahgal et al.24

The formulated C. papaya-based ISCOM had significant increase in phagocytic function of immune cells specifically neutrophils as compared to control suggesting immune enhancing effect. Pathogens are identified and destroyed by immune cells by the action of phagocytosis.31

Neutrophil adhesion to nylon fibre models the movement of polymorph nuclear lymphocytes through blood vessel walls and macrophages towards inflammation site, by the action of neutrophils in the neutrophil adhesion test.20,32,33 This implies stimulatory effect of neutrophil migration to inflammatory site. Formulated C. papaya-based ISCOM showed significant difference in the neutrophil adhesion (20.8±4.34) at a dose of 250 mg/kg for test group when compared to the negative control group (9.58±1.75). The adhesion of neutrophils to nylon fibre might be connected with up-regulation of β2 integrins as reported by Srikumar et al.28

Carbon clearance assay is used to evaluate effect on reticuloendothelial cell mediated phagocytosis.34 Macrophages engulf carbon particles in ink containing colloidal carbon, when the ink containing colloidal carbon is injected intravenously. The ISCOM when solely administered stimulated the reticuloendothelial system causing significant increase in the phagocytic index. ISCOM possesses macrophage stimulatory activity as evidenced by increased phagocytic index in carbon clearance test indicating stimulation of the reticuloendothelial system. The neutrophils are constituents of the white blood cell that normally identify, engulf and digest a bacterium according to earlier report.30

Antibody functions as effectors of humoral response by binding to antigen and neutralizing or enhancing its elimination via cross-linking to form clusters that are ingested by phagocytic cells. B cells interaction with antigen and proliferation and differentiation to antibody secreting plasma cells is characterized in humoral immunity.35,36,37 There will be agglutination if the serum contains antibody to the organism. The increase antibody percentage stimulation value by the ISCOM only indicates the responsive effect of macrophages, T and B lymphocyte subset involved in the synthesis of antibodies when compared to the standard.

CONCLUSION

C. papaya leaves yielded saponin which served as a component of ISCOM formulation to induce both cell and antibody mediated immune responses. C. papaya-based ISCOM may be useful as an immune enhance against Streptococcus pneumoniae infections. It can also be inferred that C. papaya-based ISCOM has therapeutic and immunomodulatory activities.

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CONFLICT OF INTEREST

The authors declare no conflict of interest.

ABBREVIATIONS


REFERENCES


