Diagnosis of Pneumocystosis pneumonia with two staining methods using various specimens collected from animal model

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Citation

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Abstract

Correct diagnosis of pneumocystosis pneumonia depends on suitable sampling procedure and staining method. The aim of this study was comparison two staining methods “Giemsa and GMS” in three collected samples from studied animal model “oral swab, BAL and lung homogenate”.

Twenty two female rats of two month-ages (Sprague-Dawley) with 150-200 grams of weight were used. Methyl prednizolone acetate was used to stimulate pneumocystosis. Oral swab (OS), broncho-alveolar lavage (BAL) and lung homogenate (LH) were collected during studied weeks from control and test groups. All samples were stained with Giemsa and GMS (Gomori's Methenamine Silver). These specimens were classified in four stages based on 35 microscopic fields (100X) for counting of cyst numbers, tested by two staining methods for common specificity such as number of cysts, required time for observations, cost, and possibility of improper diagnosis.

All controls were negative in all specimens by both staining methods. Oral swabs were negative by both staining methods. The Lavage specimens were negative from weeks zero to five by Giemsa method but were positive from week two to the end by GMS method. Lung homogenate specimens were negative in weeks 0 and 2 by Giemsa but negative in week 0 by GMS. These samples were positive in later weeks in two staining methods.

Based on the analyzed results Giemsa and GMS had no enough sensitivity in oral swab specimens. The best sensitivity was obtained by GMS use of LH specimens.

INTRODUCTION

Direct observation of Pneumocystis cyst or trophozoite is required for diagnosis pneumocystosis in clinical samples by staining method. Gram-Wright, Papanicolaou, Toluidine blue and GMS has been used for identification of cysts or trophozoites in prepared specimens on tissue sections. Those staining procedures using immunofluorescence monoclonal antibody have increased sensitivity and specificity (1). Giemsa and GMS are the most applied methods to diagnosis of PCP (2), while Giemsa has been the most rapid and cheapest method. The organisms are diagnosed in all steps of their life cycle. Those cysts with and without organisms not only can differentiate by Giemsa but can also differentiate these cysts from yeast cells (3,10) although isolation of the organisms on cell line can be an alternative procedures (3).

GMS is a silver staining method that is usually used for the observation of fungi in tissue specimens. This method is the standard method for diagnosis of the pneumocystosis (11).

Proper diagnosis depends on applied staining method, kind of specimens and properly sampling from infected area. Specimens are provided by either non-invasive methods (such as sputum or stimulation procedure) or even invasive methods like broncho-alveolar lavage, lung homogenate and lung biopsy. Invasive methods have higher sensitivity (12). Those samples collected from HIV positive patients have also higher sensitivity because of high load of the organisms in these samples (13).

With looking through advantages and disadvantages these two methods, we want to evaluate the efficiency of the Giemsa and GMS in diagnosis of the PCP in animal model in different samples and applied conditions.
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MATERIALS AND METHODS
Animal model: Twenty two female rats of two month-ages (Sprague-Dawley) with 150-200 grams of weight was provided from Razi Vaccine & Serum Research Institute. 17 rats for tests that divided in six groups and 5 rats for control that one specified for each group except zero group (1).

Stimulation of pneumocystosis proliferation in rat lung: methyl prednizolone acetate (40mg/ml) made by Pharmacia & Upjohn provided from Belgium was used subcutaneously to stimulate pneumocystosis in all rat lung of test group to increase the Pneumocystis carinii (2).

Specimens: Oral swab (OS), broncho-alveolar lavage (BAL) and lung homogenate (LH) were collected at weeks zero (before injection), 2, 3, 4, 5, 6, and 7 from test and control groups. All samples were divided in 1 ml micro tubes and kept at -70°C.

PROCEDURE FOR COLLECTING SPECIMENS
Oral swabs: Oral specimens were obtained by sterile applicator from mouth area; above and beneath the tongue, under Bio-safety cabinet. Cotton head of the swabs were suspended inside 5 ml of sterile PBS in sterile capped tube. The tubes were vortexed for 1 min and swabs were taken out. Specimens were centrifuged for 15 min at 3000 RPM and top 4 ml of supernatant were taken out and the rest were transferred in sterile 1.5 ml tubes and kept at -70°C.

Broncho-alveolar lavage: Separated lung of the rats was put in sterile Petri-dish. 2.5 ml of Dithiothreitol and 2.5 ml PBS were injected through trachea inside the lung. Lavage fluid were then collected and incubated for 15 min at 37°C. 2.5 ml of 159 mMol (0.85%) Ammonium Chloride solution were added to the fluids and kept at laboratory temperature for 30 min. Specimens was centrifuged for 15 min at 3000 RPM. Top of supernatant were taken out and the rest 1 ml were transferred in sterile 1.5 ml tubes and kept at -70°C
till to be used (3).

Lung homogenate: Lung homogenate specimens were prepared from whole left lobe of the lung using sterile homogenizer containing 5 ml sterile PBS. Obtained fluids were transferred from sterile 3 layer of gauze filter. 2.5 ml of Dithiothreitol were added and incubated for 15 min at 37°C. 2.5 ml of ammonium chloride (0.85%, pH 6.00) were added to the fluids to lyses of the host and blood cells and kept at laboratory temperature for 30 min. These samples were centrifuged for 15 min at 3000 RPM. Top of supernatant were taken out and the rest 1 ml were transferred in sterile 1.5 ml tubes and kept at -70°C till to be used (3).

STAINING METHODS
Giemsa: Slide smears were prepared with 10 ul of specimens and stained with Giemsa after dry up the smears. Positive results are determined by observation of Pneumocystis cysts looking either of these clues:
- Eight circular bodies (4-8 µM)
- Purple banana shape (sporozoite) along each other
- Observation of trophozoite (2-4 µM) with blue nucleolus and pink cytoplasm that are usually in mass shape near each other or separated

Gomori’s Methenamine Silver: Slide smears were prepared with 10 ul of specimens and stained with CDC procedure (4). Positive results were determined by observation of brown cyst (4-8 µM) with specific clue of GMS staining that seems to green blue with two dark spot inside

Counting method for Pneumocystis cysts: Slide smears were prepared with 10 ul of OS, BAL and LH. It was spread in 1 cm2 and fixed with methanol and stained with GMS or Giemsa. 35 microscopic fields in the center of the smears were counted and average of the cyst number was registered for each sampling and staining method.

Scoring system for Giemsa and GMS staining method: Average counting of 35 microscopic fields with 100 magnification were classified less than 10 cysts=1+, 11-100 cysts 2+, 101-1000 cysts 3+ and for more than 1000 cysts 4+

RESULTS
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OBSERVED CYSTS BY GIEMSA STAINING METHOD

Oral swab specimens: No cysts were observed in all test and control groups during studied weeks after examining 35 microscopic fields (Figure No1).

Figure 1
Picture 1: Percentage of Positive cases in studied weeks and tested specimens with Giemsa staining method

![Figure 1](image1.png)

BAL specimens: Examined BAL specimens of control group in all weeks and test group collected in weeks zero, 2, 4, and 5 were negative while 25% of week six and 100% in week seven were positive (Figure No. 1).

Lung homogenate specimens: Specimens of test group collected in weeks zero and 2 and all control group specimens were remained negative. Those specimens collected from weeks 4, 5, and 7 were 100% positive while samples in week six were 75% positive (Figure No. 1).

Figure 4
Figure 4: Average of estimated cysts of tested samples in control and test animal groups during weeks by GMS method

![Figure 4](image2.png)

OBSERVED CYSTS BY GMS STAINING METHOD

Oral swab specimens: No cysts were observed in all test and control groups during studied weeks after examining 35 microscopic fields (Figure No 2).

BAL specimens: Examined BAL specimens of control group in all weeks and test group collected in weeks zero and 2 were negative while 66.6% of week four, 100% of week five, 75% of week six and 100% in week seven were positive (Figure No.2).

Lung homogenate specimens: Specimens of test group collected in weeks zero and all control group specimens were remained negative. That specimens collected from week 2 was 50% and weeks 4, 5, 6 and 7 were 100% positive (Figure No.2).

Figure 3
Figure 3: Average of estimated cysts of tested samples in control and test animal groups during weeks by Giemsa method

![Figure 3](image3.png)

Estimation of Pneumocystis cysts in all three collected samples by both methods: Oral swab samples have no cysts were observed in all groups during studied weeks. In BAL Samples 10 cysts in average were observed only in week seven by Giemsa while less than 10 cysts in weeks 4 and 5 and more than 100 cysts in weeks 6 and 7 by GMS method. Observed cysts were different in lung homogenate.
specimens in studied weeks by Giemsa and GMS method. In Giemsa method, no cysts were observed in weeks zero and 2, less than 10 cysts were estimated in weeks 4, 5 and 6. Estimated cysts in week seven were 10 to 100. In GMS method no cysts were observed just in week zero. Its number were less than 10 in week 2 and 10 to 100 in weeks 4, 5 and 100 to 1000 in week six and were more than 1000 in week seven (Figures No 3 and 4).

Comparison of specific criteria and scoring of two applied staining methods: considered criteria were compared with each other and methods were scored, the results are shown in (Table No.1).

**Figure 5**
Table 1: Comparison of the specificity of two applied staining method

<table>
<thead>
<tr>
<th></th>
<th>GMS</th>
<th>Giemsa</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sample Type</td>
<td>High</td>
<td>Low</td>
</tr>
<tr>
<td>Presence of stained organisms</td>
<td>Negative</td>
<td>Negative</td>
</tr>
<tr>
<td>Number of cysts</td>
<td>40</td>
<td>30</td>
</tr>
<tr>
<td>Yeast Cells</td>
<td>None</td>
<td>None</td>
</tr>
<tr>
<td>False Positive</td>
<td>No</td>
<td>No</td>
</tr>
<tr>
<td>False Negative</td>
<td>No</td>
<td>No</td>
</tr>
</tbody>
</table>

**Figure 6**
Table 2: Percentage of Scoring in diagnosis method of Giemsa and GMS

<table>
<thead>
<tr>
<th></th>
<th>Giemsa</th>
<th>GMS</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>-</strong></td>
<td>1.8</td>
<td>51.3</td>
</tr>
<tr>
<td><strong>+</strong></td>
<td>23.1</td>
<td>12.8</td>
</tr>
<tr>
<td><strong>++</strong></td>
<td>5.1</td>
<td>15.4</td>
</tr>
<tr>
<td><strong>+++</strong></td>
<td>0</td>
<td>15.4</td>
</tr>
<tr>
<td><strong>++++</strong></td>
<td>0</td>
<td>5.1</td>
</tr>
</tbody>
</table>

**SCORING RESULTS OF STAINING METHODS**

71.8% of tested specimens were negative and the rest 28.2% had just 1+ and 2+ grade by Giemsa. Negative rate of GMS was less than Giemsa. 51.3% of GMS had negative results while 15.4% and 5.1% had 3+ and 4+ grades (Table No. 2).

**DISCUSSION**

Three specimens (oral swab, lung homogenate and BAL) have been compared with each other in this study. Analyzed results revealed none of oral swab samples were positive even in the presence of the infection. These samples had the highest positive rate in lung homogenate and then in BAL specimens just in the early stage of the infection, while both had the same sensitivity in the last weeks.

It is frequently observed that microscopic examination of stained smears on specimens is standard method for diagnosis of PCP. Staining method is usually based on diagnosis of the cysts, because trophozoites are normally mistaken with counter-stain materials (13, 17). Giemsa method is the easiest, cheapest and the most rapid procedure, enable to differentiate those empty cysts from those containing organisms or even with yeast cells (5,6). Despite these advantages, diagnosis of the PCP is rather difficult and requires skilled microscopist because differentiation of the organisms from smears materials is not easy (17). Sensitivity of this method is also expected to be lower than GMS because of incapability of those empty cysts.

On the other hand, GMS method is usually used to observed fungi in tissue specimens. This method is specific for cyst wall and is applied as a gold standard method for diagnosis of pneumocystosis. Having long steps of staining and requirement of more than one hour time to perform are of its limitations. This method also can not diagnose trophozoite. There is possibility infection can not be identified in those mild infections or those cases with high ratio of trophozoite to cyst (17). Misdiagnosing is other limitation (1). Additionally, effect of anti-pneumocystosis drugs can not be determined because those empty cysts are not differentiated from the cysts containing trophozoites.

It is frequently observed HIV positive patients had pneumocystis cysts in their lung after treatment. Sporozoites of these cysts that have been destroyed by the drugs are responsible for these situations (18). We can differentiate pneumocystis cysts from other organisms because of two dark spots attached to each other inside the cysts, if GMS staining method to be performed in best conditions. We could decrease require necessary time for staining steps up to 30 min in this study.

It is reported Giemsa method has low sensitivity among studied staining procedures (13, 17, 19). Obtaining results in this study is in agreement with the other reports. Analyzed Results revealed GMS has higher sensitivity than Giemsa. Gurpreet has reported that routine staining method has enough sensitivity in those specimens containing high organisms such as biopsy, BAL specimens and taken specimens of those HIV positive patients (11, 20). Obviously, this routine method has high false positive results in HIV negative patients even in BAL specimens (15, 21).

The standard staining method is microscopic observation of Pneumocystis in specimens such as BAL, Stimulated sputum, or biopsy (22,23,24). Oral swab sampling is non-
invasive method that has been applied in animal model \( (32) \) with no positive results. Diagnosis of PCP is usually made by stimulated sputum sampling in HIV positive patients that is rarely positive in HIV negative patients with immunodeficiency problem. Correct diagnosis requires proper sampling such as BAL or lung biopsy. However, stimulated sputum sampling is more applicable in HIV positive patients because of higher load of organisms in this patients group \( (33,34) \). The sensitivity will be increase in those invasive samples. The quality of the result will be better in those invasive specimens that sampling is near to the colonized area \( (31,32) \). BAL sample is an invasive method that is occasionally facing with danger, although, it is a selective sample \( (32) \). Its sensitivity by staining methods is about 55-78 percent \( (11,13) \). False negative is even high in BAL sample in HIV negative patients \( (32) \). Besides false negative rate is high in non-invasive sampling methods and those patients that are in early stage of the disease \( (9) \). Open lung biopsy is the most invasive method that require for operating room. Use of this technique enables us to have the best specimens, although patients are facing with side effects \( (13) \).

CONCLUSION

Staining is necessary for diagnosis of pneumocystis if no other alternative method is available. Therefore, it is underlining that Giemsa procedure to be considered as the last alternative method. However, it is recommended to use other new diagnostic method such as PCR to decrease the risk of danger invasive sampling.

References

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