A Case-control Study of Vitamin A Level in Hydatidiform Mole

Andrijono*, Kukung Kurnia*, Nur Askin**

INTRODUCTION

Hydatidiform mole is one of the problems in women's reproductive health. This is because this disease is suffered mostly by women of young age and without children who still need their reproductive functions, and its incidence is high enough, i.e. one of 60 pregnancies or 16%.1,2 One of the most dangerous hydatidiform mole complications is the occurrence of malignancy degeneration which results in the failure of reproductive functions, or even death.3,4

One of the efforts to promote women's reproductive health is through the prevention of the occurrence of hydatidiform mole by undertaking the studies on the etiology of this disease and its risk factors. The decrease of hydatidiform mole incidence will be a great significance in the efforts to promote women's reproductive health.

Hydatidiform mole is an abnormal pregnancy indicated by degeneration of hydropic chorionic villi, avascular chorionic villi and proliferation of trophoblast
The cause of these changes has not yet been identified. However, it has been understood that the changes in cell differentiation and cell proliferation are caused by the disorder of intracellular metabolism. One of the metabolism chains is the function of vitamin A. Therefore, it will be interesting to study whether there is a correlation between vitamin A and the incidence of hydatidiform mole. This study is performed with the purpose of attempting to give the tentative answer to that problem.

**MATERIALS AND METHODS**

The study was designed as a case control study. The cases were the patients with hydatidiform mole and the controls were the pregnant women. Two determining factors were age and parity. These two factors were used as matched parity with an interval of one and ages with interval of four. All the patients with hydatidiform mole which met the criteria of the study were included in the study, and these matched factors served as the control. The patients with hydatidiform mole included in this study were those evacuated at Dr. Cipto Mangunkusumo Hospital, neither receiving blood transfusion yet nor the treatment with vitamin A > 5000 IU. The measurement of vitamin A performed was the level of fasting retinol, and it was carried out at the Laboratory Research Center for Immuonoendocrinology Jakarta, according to Neeld and Pearson's method.

**RESULTS**

During the period of a year study (June 1990-May 1991), a number of 30 cases of hydatidiform mole were qualified for the study.

The average cases were 25 years (± SD: 5.73 ) with the youngest age being 18 years and the oldest 40 years. The average age of the control group was 26,43 years (± SD: 4.28 ), the youngest age being 14 years and the oldest 38 years.

For parity, 70% of the cases and 76% of control group had parity of 1 or 0, the rest had parity more than 1. The average education length of case group was 7.96 years (± SD: 3.64 ) while the control group being 10.7 years (± SD: 3.78 ). Statistically, all these factors did not differ significantly.
The results showed no influence of age, parity and education with regard to vitamin A level. These three factors in our study did not seem to affect the study on vitamin A in both the case group and the control group. Thus, these results differed from several studies which suggested that the above three factors play a role in affecting vitamin A level.

### Vitamin A deficiency

The assessment of risks for the occurrence of hydatidiform mole in vitamin A deficiency showed OR = 2.51.

#### Table 4. The average vitamin A level according to the level of education in both groups

<table>
<thead>
<tr>
<th>Education (year)</th>
<th>Case</th>
<th>Control</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>n</td>
<td>x ± SD</td>
</tr>
<tr>
<td>0</td>
<td>1</td>
<td>13,40 ± 0.00</td>
</tr>
<tr>
<td>1-6</td>
<td>15</td>
<td>10.99 ± 4.38</td>
</tr>
<tr>
<td>7-12</td>
<td>13</td>
<td>9.24 ± 2.95</td>
</tr>
<tr>
<td>12+</td>
<td>1</td>
<td>17.10 ± 0.00</td>
</tr>
</tbody>
</table>

*p > 0.05  p > 0.05*

Although there was no statistical difference, it was evident that the percentage of vitamin A deficiency in the patients with hydatidiform mole (43.33%) was higher than that of control group (23.33%). In the distribution of age and parity, the majority of age and parity in the cases of hydatidiform mole were the cases of less than 24 years of age, and parity of 1. This finding was in according with the literature suggesting that the highest incidence of hydatidiform mole occurred in the young women with low parity. It will be of interest to study such tendency separately.

#### Analysis of the women in the age group less than 24 years.

#### Table 6. Vitamin A deficiency in the case and control group with age less than 24 years old

<table>
<thead>
<tr>
<th>Group</th>
<th>Vit. A Def. (+)</th>
<th>Vit. A Def. (-)</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case (mole)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control (pregnant)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*p = 0.038  OR = 6.29*

These data suggest high risks, i.e. 6.29 times in the women group of ≤ 24 years of age with vitamin A deficiency for developing hydatidiform mole.

#### Table 7. Vitamin A deficiency in the case and control group with parity of 1 or less

<table>
<thead>
<tr>
<th>Group</th>
<th>Vit. A Def. (+)</th>
<th>Vit. A Def. (-)</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case (mole)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control (pregnant)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*p > 0.05  OR = 3.38*

On the other hand, analysis of the patients with parity of 1 showed no significant difference (OR = 3.38). It will be of interest, therefore, to perform a separate study of the cases with nil parity.

#### Table 8. Vitamin A deficiency in the case and control group with nil parity and age less than 24 years old

<table>
<thead>
<tr>
<th>Group</th>
<th>Vit. A Def. (+)</th>
<th>Vit. A Def. (-)</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Control</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*p < 0.05  OR = 7*

It is obvious that there is a significant different between the case and control groups with nil parity, age less than 24 years old and vitamin A deficiency. The risk
of hydatidiform mole in the women of less than 24 years with nil parity and suffering from vitamin A deficiency is 7 times higher than in the control group.

Histologic Classification

Several investigators did not find the influence of histologic classification on the likelihood of the post-hydadiform mole malignancy. However, this histologic classification did not provide a brief description of the essential changes in trophoblast cells that occurred, i.e. the more severe and the more numerous cases were classified as the more severe cases. Histologic classification of hydatidiform mole commonly referred to is the classification according to Hertig-Manse1.11

Table 9. The relationship between histopathologic classification and the average vitamin A level

<table>
<thead>
<tr>
<th>Histopathologic classification</th>
<th>Vitamin A level</th>
<th>n</th>
<th>x</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>I</td>
<td></td>
<td>6</td>
<td>13.12</td>
<td>4.56</td>
</tr>
<tr>
<td>II</td>
<td></td>
<td>20</td>
<td>9.90</td>
<td>3.32</td>
</tr>
<tr>
<td>III</td>
<td></td>
<td>4</td>
<td>9.70</td>
<td>5.11</td>
</tr>
</tbody>
</table>

\[ F = 1.7, \quad df = 2.27, \quad p > 0.05 \]

Statistically, there was no significant difference between histologic level and vitamin A level. However, it was evident that the higher the histologic changes, the lower the vitamin A level.

DISCUSSION

One of the diagnostic parameters of hydatidiform mole is the presence of trophoblast cell proliferation. The direct cause of this proliferation is not known yet. The in vivo study showed that vitamin A has the effect of inhibiting or controlling cell proliferation and enhancing cell differentiation.7,8,9 How vitamin A performs that role is unclear yet. Several mechanisms that may be affected include protein system of kinase C, cascade system, and several other biological effects, such as enzyme synthesis, genomic expression, extracellular effects, immunologic activity and protein kinase C cascade.12,13,14,15 It is of interest, therefore, to study whether vitamin A also plays a role against the proliferation of trophoblast cells.

Started by clinical study, i.e. control case study to identify the correlation of the occurrence of hydatidiform mole and vitamin A level, the analysis of vitamin A level showed that the majority of the cases and controls had vitamin A level below the normal (20-40 µg/dl). These results suggest that most pregnant women in the study suffered from vitamin A deficiency. This situation calls for special attention on the impacts it may have on pregnancy and how to deal with them. The level of vitamin A deficiency is mild if blood retinol level is 10-20 µg/dl, and the deficiency is severe if the retinol level is <10 µg/dl. However, the average results of vitamin A level in the cases of hydatidiform mole and in the control group were statistically different. Analysis of age, parity, and education factors did not seem to show different result. Education factor is suspected to play a major role because there is a close correlation between education and socioeconomic conditions. Furthermore, the socioeconomic conditions play a major role in the quality of nutritional food. This correlation suggests a probable role of vitamin A deficiency in the incidence of hydatidiform mole.

One of the hypotheses explaining the occurrence of hydatidiform mole is the "blighted-ovum" theory. The genetic study revealed that the genotype of hydatidiform mole was of the father's gene, without the mother's.16 This, hydatidiform mole originates from fertilization without mothers factor, i.e. the ovum whose gene factor is not fertilized by the father's sperm.15 However, this situation still requires other factors because not all the blighted ovum develops into hydatidiform mole. This other factor is yet to be identified. And the factor that we attempted to study is vitamin A deficiency.

Epidemiological study of hydatidiform mole revealed that hydatidiform mole was generally developed by pregnant women of young age without children. This risk factor also constitutes the problem that is yet to be answered. It is of interest, therefore, to perform a study on the factors of young age, low parity or nil parity, and vitamin A deficiency. The interesting data of our study are the data of hydatidiform mole patients of ≤24 years of age suffering from severe vitamin A deficiency. These data suggest a significant difference between the case and the control group. The risk for developing hydatidiform mole increases 6.29 times if the woman is under 24 years, pregnant, and suffering from severe vitamin A deficiency.
Similarly, in the cases of ≤ 24 years with nil parity, the data found revealed the risk of 7 times higher than in the control group. The magnitude of risk for the occurrence of hydatidiform mole by the accumulation of these factors underlines that the etiology of hydatidiform mole is multifactors.

Thus, the prevention for the occurrence of hydatidiform mole should be attempted through various aspects. Nevertheless, as it is a natural phenomenon that the first pregnancy occurs at young age, the risk factors that play a role, such as vitamin A deficiency. The severity of histologic changes can be seen from the compared data of histologic level and the average vitamin A level. The lower vitamin A level reflects the more severe histologic changes of hydatidiform mole cells.

Several studies of histologic levels of hydatidiform mole revealed that this histologic level did not affect the risks for the occurrence of malignancy degeneration (Genest DR et al. personal communication). Thus, the results of this study are sufficient to show that the women of ≤ 24 years with nil parity and suffering from severe vitamin A deficiency have the risk for developing hydatidiform mole seven times higher. Vitamin A level in the patients with hydatidiform mole compared with the pregnant woman control and histologic changes, which occurred more severely with the lower vitamin A level, suggest a strong likelihood of the role of vitamin A level deficiency in the occurrence of hydatidiform mole.

CONCLUSION

There is a significant difference of vitamin A level in the patients with hydatidiform mole and the control group. The risk for developing hydatidiform mole in the women less than 24 years of age with nil parity and suffering from severe vitamin A deficiency is 7 times higher.

REFERENCES