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<td>猿マラリアを疑わしめる多重感染の著しい輸入マラリア例</td>
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<td>著者</td>
<td>塚本 増久</td>
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<td>雑誌名</td>
<td>熱帯医学</td>
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An Imported Human Malarial Case Characterized by Severe Multiple Infections of the Red Blood Cells

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Abstract: Thin blood smears prepared from a Japanese student showed malaria parasites of which characteristics were similar to those of *Plasmodium vivax* except for their heavy multiple infections of the host red blood cell (RBC). Immediately after returning to Japan from a trip to Afghanistan, the patient showed severe symptoms with an irregular pattern of fever. However, examination of the blood smear taken on admission indicated that the parasitaemia was only 0.82%. More than 30% of the infected RBC were invaded by 2 or more young trophozoites and up to 7 ring forms in a single RBC were photographed for illustration, whereas 8 rings/RBC had once been observed. All blood forms of the *vivax*-type parasite were detected in the host RBC with or without stippling. The infected RBC showed more or less enlargement in accordance with the growth of the parasites. The possibility of mixed infection with *P. falciparum* and *P. vivax* was rejected on the basis of the features of the infected host cells and the absence of elongated gametocytes. Merozoite numbers ranged from 12 to 19 with an average of 15.4, thus the diagnosis of *P. malariae* or *P. ovale* was also excluded. In some simian malaria parasites, it is known that such severe multiple infections are characteristic and that the morphology of blood stages is practically very difficult to distinguish from that of *P. vivax*. Therefore some possibilities of natural infection with a simian malaria parasite have also been discussed.

Imported cases of human malaria in Japan are increasing in number each year (Nakabayashi et al., 1976). The writer has recently had an opportunity to examine thin blood smears taken from a university student, who was admitted to the Tohoku University Hospital, Sendai, Japan, as a patient suspected of having malaria almost immediately after coming home from a trip to Afghanistan. No detailed clinical information was available except that he showed an irregular pattern of high fever before and after admission to the hospital. At a glance of the thin blood smear under the microscope, severe multiple infections of the host erythrocytes (RBC)* were noticed in addition to the usual *vivax*-like malaria infection. This could not be easily ascribed to any parasite species of the typical human malaria. Among the usual human

* Abbreviation used: RBC for erythrocyte(s) or red blood cell(s)
malaria parasites, multiple infections of parasites in a single RBC are rather common in the case of *falciparum* malaria, and up to 8 rings were reported within a single RBC by Springall (1943). Even in the case of *vivax* malaria double or triple infections of the host cell occurs but severe cases of multiple infections are uncommon in *vivax* malaria. It is, therefore, very intriguing to diagnose the parasite species in this particular case. The present paper describes the results of the examination of blood smears of such a curious cases of human malaria.

**MATERIALS AND METHODS**

Two thin blood smears taken from the young male student mentioned above were sent, after Giemsa staining, to be examined. One (Slide A) was said to have been prepared on admission to the hospital on September 20, 1976, and hence prior to initiating chemotherapy with antimalarial drugs; and another (Slide B) was said to have been prepared a few days later during the chemotherapy. These blood smears were examined using an oil-immersion optical system, and the parasitaemia was counted as usual.

**RESULTS**

1. **Parasitaemia**

   On the basis of trial counts of 15 microscopic fields on Slide A, 39 RBC were infected with malaria parasites among a total of 4736 RBC counted; thus the parasitaemia was 0.82% on admission. By a similar examination of Slide B, a greatly reduced parasitaemia (0.0027%) was found in the patient a few days after drug treatment had begun.

2. **Erythrocytic Forms of the Parasites**

   Several different stages of the erythrocytic forms were observed. Only the infected RBC were then examined and Table 1 summarizes the results of such differential counts of the parasites. Among a total of 1,280 infected cells counted, about 800 RBC contained the young trophozoite (ring form) and about 300 RBC were infected with older trophozoites. The general appearance of the malaria parasites of different stages are briefly described below.

   **Young Trophozoites** (Figs. 1-4): Typical ring forms are less amoeboid with a well developed vacuole and a diameter about one-fourth of the infected RBC. The nucleus is round or oval and usually single, but double chromatin dots of either equal or unequal size are often found; a small accessory dot is also observed frequently (Fig. 4).

   **Old Trophozoites** (Figs. 5-11): Trophozoites become much more amoeboid and their features resemble those of *P. vivax* in size and shape. The older trophozoites usually occupy from two-thirds to three-fourths of the enlarged host cell. No band forms have been detected. The fine brownish black granules of malaria pigment become visible within the markedly irregular-shaped cytoplasm.

   **Immature Schizonts** (Figs. 12-14): The nucleus is by now divided into 4–8 fragments of
Table 1. Differential counting of malaria parasites on the blood smear prepared from the patient on admission (Slide A)*

<table>
<thead>
<tr>
<th>Parasite stage</th>
<th>No. of infected RBC examined</th>
<th>Percentage</th>
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</thead>
<tbody>
<tr>
<td>Young trophozoites</td>
<td>797</td>
<td>62.3</td>
</tr>
<tr>
<td>Old trophozoites</td>
<td>308</td>
<td>24.1</td>
</tr>
<tr>
<td>Immature schizonts</td>
<td>92</td>
<td>7.2</td>
</tr>
<tr>
<td>Mature schizonts</td>
<td>28</td>
<td>2.2</td>
</tr>
<tr>
<td>Merozoite group</td>
<td>4</td>
<td>0.3</td>
</tr>
<tr>
<td>Gametocytes</td>
<td>51</td>
<td>4.0</td>
</tr>
<tr>
<td><strong>Total</strong></td>
<td><strong>1,280</strong></td>
<td><strong>(100%)</strong></td>
</tr>
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* Counting was based on the examinations for 500 microscopic fields corresponding to about $1.56 \times 10^5$ RBC.

irregular shape, and the pigment granules are still not so evident. The immature schizonts develop largely within the RBC.

*Mature Schizonts* (Figs. 15–17): The schizont occupies large parts of the whole RBC and malaria pigments gather together to form a large mass at the center or masses at some peripheral portions. Nuclei are isolated from each other and hence the number of merozoites (or at least well separated chromatin dots) can be counted individually. The counts ranged from 12 to 19 with an average of 15.4 merozoites:

<table>
<thead>
<tr>
<th>Merozoites/RBC</th>
<th>12</th>
<th>13</th>
<th>14</th>
<th>15</th>
<th>16</th>
<th>17</th>
<th>18</th>
<th>19</th>
</tr>
</thead>
<tbody>
<tr>
<td>Counts</td>
<td>2</td>
<td>2</td>
<td>7</td>
<td>5</td>
<td>8</td>
<td>4</td>
<td>3</td>
<td>1</td>
</tr>
</tbody>
</table>

*Gametocytes* (Figs. 18–20, 32): Matured or nearly matured gametocytes are not crescent in shape, thus this excludes the possibility of *P. falciparum*. The cytoplasm of the gametocytes nearly fills the enlarged RBC, and dark pigment granules of different sizes are scattered in the cytoplasm.

3. Multiple Infections

One of the most remarkable characteristics of the present case was the tendency to heavy multiple infections of a single host RBC. Then only the infected RBC were assorted into several groups on the bases of the number of parasite(s) within each RBC and the different stages of the blood forms. Distribution of blood forms of the parasites in such differential counts for Slide A is summarized in Table 2. On admission more than 60% of the parasitized RBC were infected with young trophozoites (ring forms). Among a total of 1,280 infected cells counted, at least 18.8% were invaded by 2 or more parasites, or in other words, about 30% of the RBC invaded by the ring forms were found to have multiple infections. Double or triple infections were common and easily noticed at a glance (Figs. 4, 5, 21, 29, 30). Although the highest case of multiple infections, 8 rings in a single RBC was observed when a photographic attachment was not in place during the examinations, two cases of 7 rings per RBC have been photographed later (Figs. 27, 28). In the cases of multiple infections, mostly
Table 2. Multiple infections of the host RBC with young trophozoites of vivax-like parasites from the Slide A

<table>
<thead>
<tr>
<th>No. of ring forms in a single RBC</th>
<th>No. of infected RBC examined</th>
<th>Percentage</th>
<th>Rate of multiple infections</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>557</td>
<td>69.9</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>139</td>
<td>17.4</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>51</td>
<td>6.4</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>29</td>
<td>3.6</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>17</td>
<td>2.1</td>
<td>30.1%</td>
</tr>
<tr>
<td>6</td>
<td>3</td>
<td>0.4</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>1</td>
<td>0.1</td>
<td></td>
</tr>
<tr>
<td>Total</td>
<td>797</td>
<td>(100%)</td>
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young trophozoites of about the same stage were found within a single RBC, but in some cases combinations of the different stages of parasites were also observed, for example, 2 growing trophozoites (Fig. 30), several young and old trophozoites (Fig. 31), one gametocyte and 1–3 young trophozoites (Figs. 19, 32), 2 gametocytes (not shown in figures), and so on. In such cases, parasites were described as old trophozoites or gametocytes in Table 1. Thus the actual rate of multiple infections is higher than that shown in Table 2 where multiple infections were recorded on the basis of counts for only young trophozoites.

4. Stippling and Enlargement of the Infected Host RBC

The RBC infected with a young ring form usually does not show any stippling and the size of the RBC is also not enlarged (Figs. 1, 3, 4). In some cases, however, a markedly enlarged RBC harboured only a single young trophozoite without any stippling (not shown in figures). The RBC infected by several young or old trophozoites becomes enlarged with or without stippling. In the present case stippling is not so remarkable, and, where present, its colour is bluish gray instead of pink, the colour typical of Schüffner's dots. This might be due, however, to staining conditions because at most hospitals in Japan blood smears are routinely stained under slightly acidic conditions, such as pH 6.5–6.8. Trial counts of the infected RBC demonstrated that there was no difference in distribution of the parasites of various blood stages between the stippled RBC group and the unstippled RBC group. In other words, it is impossible to divide malaria parasites into 2 groups on the basis of the presence or the absence of Schüffner's dots in this case. Thus it is unlikely that there was a mixed infection of distinct parasite species. A single infection with the malaria parasite, especially with an old trophozoite, found in an enlarged and stippled RBC shows characteristics rather

Figs. 1–3: Trophozoites in a host RBC with or without stippling; Fig. 4: Trophozoites with an accessory chromatin dot; Fig. 5: Growing trophozoites in a well-stippled and elongated RBC; Figs. 6—11: Old trophozoites in an enlarged RBC with or without stippling; Figs. 12–14: Immature schizonts; Figs. 15–17: Mature schizonts; Figs. 18–20: Gametocytes.
PLATE I. — Various blood stages of *vivax*-like malaria parasites.
typical of *P. vivax* infection. In some cases, the infected RBC showed the highly deformed shape similar to *P. ovale* infection (Figs. 5, 12, 15, 29).

5. Examination of Slide B

All the observations described in previous sections were based on the examination of Slide A, but parasitological informations obtained from the examination of Slide B have been markedly different from above because of the consequence of the chemotherapy with antimalarials. A total of 29 parasites could be detected after extensive examination of 387 microscopic fields corresponding to about $1.07 \times 10^5$ RBC. The parasitaemia calculated was thus 0.0027%. The parasites detected were: 2 ring forms, 9 old trophozoites, 3 immature schizonts, and 15 gametocytes. The morphology of all the parasites except gametocytes appeared to be greatly damaged presumably by the drug treatment, and no multiple infections were detected. Both microgametocytes and macrogametocytes appeared to be intact morphologically, and again no crescent-typed gametocytes were observed.

**DISCUSSION**

The phenomenon of multiple infections of the host RBC is not rare in human malaria. Especially, in *P. falciparum*, it is rather frequent. As an extreme case, up to 8 young trophozoites have been reported within a single RBC by Springall (1943) in a fatal case where the parasitaemia was more than 44% and the rate of multiple infections was 37.0%. Field and Shute (1956) also showed an example of multiple infections in a fatal case where the parasitaemia was more than 38.3% just before death, and the rate of multiple infections was 23.6% with a maximum of 7 rings in a cell. In the present case of human malaria, however, the possibility of mixed infections with *P. falciparum* and another species is not acceptable by the evidence mentioned above: *i.e.*, stippling and enlargement of the infected host cells and the shape of gametocytes.

In the case of *P. vivax*, double or triple invasions of the host RBC are also not uncommon whereas the frequency of multiple infections and the number of parasites found within single cells are usually not so high as in cases of *P. falciparum*. In an extreme case of even *P. vivax*, however, Field (1942) recorded exceptionally heavy infections where one RBC appeared to have 8 young trophozoites. The rate of multiple infections is calculated to be 37.6% from his data. As cited by Field and Shute (1956), similar severe cases of multiple infections with up to 6 rings/RBC were also recorded by Chalmers and Archibald (1920), Rodenhuis.

Figs. 21–22: Trophozoites with 2 chromatin dots or nearly divided nucleus; Fig. 23: 4 ring forms in a host cell with stippling; Figs. 24–26: 5 ring forms in a host RBC; Figs. 27–28: 7 ring forms in a single RBC; Fig. 29: Deformed RBC with young or old trophozoites; Fig. 30: Double infection of RBC with 2 growing trophozoites; Fig. 31: An enlarged RBC invaded by 3 old trophozoites and 3 young trophozoites; Fig. 32: A single RBC occupied by a large gametocyte and 3 young trophozoites.
PLATE II. — Multiple infections of a single host RBC.
Das Gupta (1939) and Young and Eyles (1949). In the case of P. ovale, double or triple infections of single RBC are more common than in cases of P. vivax but any heavy multiple infection of P. ovale has not been reported so far. On the other hand, multiple infection of the RBC is extremely rare in the case of P. malariae.

In contrast to these human malaria parasites, the phenomena of multiple infections have been reported to be more frequent in several simian malaria species. For example, in P. cynomolgi bastianellii the presence of up to 6 ring forms has been described by Garnham (1966), and in P. cynomolgi cyclopis up to 7 rings have been recorded by Inoki et al. (1942), whereas in the original P. cynomolgi cynomolgi such heavy multiple infections have not been reported (with up to 4 parasites). P. eylesi is also one of the simian malaria species characterized by a high frequency of multiple infections and the average rate of multiple infection reported was 29.45% by Warren et al. (1965).

In the present case of vivax-like malaria, the writer’s first impression during the microscopic examination of the thin blood smears was so curious that the possibility of simian malaria parasites came to mind. Garnham (1966) has pointed out that parasites of simian origin might explain some of the abnormal or aberrant forms reported as P. vivax. Indeed, the human is more or less susceptible to the infection of certain simian malaria parasites (Garnham, 1966; Coatney et al., 1971). So far the malaria parasites of simian origin distributing in south-east and central south Asia which are known to be infectious to humans whether naturally, accidentally or experimentally, are: P. knowlesi (Knowles and Das Gupta, 1932; Chin et al., 1965; Yap et al., 1971; etc.), P. cynomolgi (Schmidt et al., 1961; Coatney et al., 1961; Rennet and Warren, 1965), P. cynomolgi bastianellii (Eyles et al., 1960; Garnham et al., 1962; etc.), P. eylesi (Coatney et al., 1971), and P. inui (Das Gupta, 1938). It is, therefore, not unlikely that the malaria parasite of the present human case is of some simian origin.

P. vivax and P. cynomolgi bastianellii resemble each other and practically are not easy to distinguish by means of examination of a blood film alone whereas they can be differentiated by knowing other stages of the life cycle and their infectivity to the rhesus monkey. Because of the marked resemblance of P. cynomolgi bastianellii to the aberrant forms of P. vivax reported by Field (1942), Sandosham et al. (1962) have stated in their record that “a possible explanation may be that Field was in fact dealing with two cases of naturally acquired human infections of P. c. bastianellii of monkey origin.” According to Garnham et al. (1962), too, “the first important clue in the differential diagnosis of bastianellii and vivax malaria” is that “the human infections (of P. c. bastianellii) are characterized by the association of relative severity of symptoms with very low parasitaemia.” These characteristics are also applicable to the present situation, although the present parasitaemia of 0.82% is still higher than the very low level of their concept. A diagnostic method more conclusive than morphology is to test infectivity to the rhesus monkey: i.e., P. vivax cannot invade the rhesus monkey whereas P. c. bastianellii can easily do so. However, when the present writer could examine the blood smears, the patient was already discharged from the hospital after curative therapy. Hence a desirable subinoculation of the blood samples into rhesus monkeys and subsequent
confirmation of whether the parasite was an aberrant form of the human \textit{vivax} malaria or one of the rare cases of natural infection with a parasite of simian origin had become impossible. The likelihood of infection with \textit{P. eylesi} may be excluded by its higher number of merozoites (or at least divided chromatin dots) than that found in the present case. Another possibility of a mixed infection with \textit{P. vivax} and a similar simian malarial parasite still remains obscure. At present, therefore, the writer must suspend making a definite diagnosis of the parasite species.

\textbf{ACKNOWLEDGMENTS}

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\textbf{REFERENCES}


* Not seen, cited by Field & Shute (1956) ; Garnham (1966) ; Coatney et al. (1971)