Vertical transmission of dengue virus in the Yogyakarta airport area

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Abstract

Background: International Health Regulations controls international travel including human movement, disease vector, and imported items to prevent the spread of dengue, especially in seaports, airports, and border crossing posts. This study aimed to determine dengue Transovarial Transmission Index (TTI) and distribution of dengue virus in the areas around Adisucipto Airport of Yogyakarta, Indonesia.

Methods: The study was a descriptive analytic study with cross sectional design, conducted by mapping the spread of the dengue virus and identifying TTI in Adisucipto Airport. A total of 145 ovitraps were installed in both perimeter and buffer areas of the airport. Positive Ovitrap Index (OI), TTI, and serotype of dengue virus were examined. The TTI was identified using immunocytochemistry immunoperoxidase streptavidin biotin complex (IISBC) method in mosquito head squash preparations.

Results: OI in the buffer area was 32 (45.1%), whereas OI in the perimeter area was 24 (32.4%). The TTI in the buffer and perimeter areas were 21 (18.3%) and 11 (18.9%), respectively. The TTI was found greater in the Aedes aegypti population compared to the Aedes albopictus population, both in the perimeter area (20% versus 16.7%) and the buffer area (20.3% versus 16.1%). Dengue virus serotype-2 (DENV-2) and dengue virus serotype-3 (DENV-3) were predominantly found in Ae. aegypti and Ae. albopictus.

Conclusions: Buffer areas of Adisucipto Airport of Yogyakarta have higher risk as breeding sites for Aedes spp., predominantly DENV-2 and DENV-3 serotypes. High OI shows that the areas are likely to have higher risk of developing dengue outbreak.

Keywords: Transovarial Transmission Index, Airport, Dengue, Aedes spp.
to 159 cases per 1000 travelers during epidemics [3], although dengue transmission in the airport has not yet been reported until now.

The New Tokyo Narita International Airport Quarantine Post in Chiba Prefecture in year 2000 to 2002 had examined 233 passengers suspected of being infected with dengue virus: 1 case (4%) out of 26 cases identified in year 2000, 8 cases (12%) out of 69 cases identified in year 2001, and 22 cases (16%) out of 138 cases identified in year 2002 were confirmed as dengue infection [4]. Most of the passengers were infected after traveling from Southeast Asian and South Asian countries, one from African countries, one from Central American countries, one from Central and South American countries, and one from South American countries [4]. Fever screening in Taiwan Airport begun in July 2003 to June 2004 identified 40 dengue cases, among which 33 people (82.5%) were confirmed as viremic patients [5]. During 2007 to 2010, sentinel surveillance in Taiwan Airport showed that most of the dengue-infected travelers had just returned from endemic areas around the Southeast Asian region, namely Indonesia (21.0 to 35.1%), Thailand (5.0 to 13.0%), the Philippines (9.0 to 12.3%), Cambodia (4.1 to 8.0%), Malaysia (2.0 to 4.1%), Singapore (1.1 to 3.4%), India (0 to 1.1%), and only few travelers who had just returned from South America (0 to 0.7%) [6]. In September 2013, in Germany, a traveler who had just returned from Japan was confirmed as having type 2 dengue virus infection; hence, the German Health Authority performed strict monitoring towards travel history of the travelers in order to evaluate risk potential of travelers having dengue virus infection [7].

International travel such as human movement, disease vector, and contaminated items that potentially cause widespread disease are regulated by the WHO in the International Health Regulations 2005 Article 9 (nine). Each country is required to perform dengue risk assessment to prevent the spread of dengue between countries by strengthening surveillance and supervision at the entrance areas, i.e., seaports, airports, and border crossing posts [8].

According to the Law Act 1 Year 1962 regarding Marine Quarantine and Act 2 Year 1962 regarding Sky Quarantine, as well as the IHR Year 2005 Article 2, all of which state that seaports and airports are obliged to be free from disease vectors and are required to perform disease control and prevention suitable to the potential risk factors without interfering with the commercial traffic [8]. Perimeter ports including seaports, airports, and rural ports are expected to be free from both larval stage and adult stage of Aedes aegypti mosquitoes, whereas in the buffer areas the following values are considered essential: House Index (HI) of less than, or equal to 1%, Breteau Index (BI) of < 50, biting rate of < 2.5, and OI of < 15% [9].

This study was conducted to acquire information regarding the presence of Aedes spp. mosquitoes and the extent of transovarial transmission in the perimeter and buffer areas of Yogyakarta’s Adisucipto Airport. The study was also intended to determine dengue Transovarial Transmission Index (TTI) and distribution of dengue virus in areas in and around Adisucipto Airport, Yogyakarta, Indonesia.

**Methods**

This research is a descriptive analytic study with a cross sectional design. The study was conducted by mapping the spread of dengue virus and identifying TTI in Adisucipto Airport of Yogyakarta, Indonesia (07° 47’ 17” S and 110° 25’ 54” E). The study was done in December 2015 to May 2016, during which rainy season took place.

Dengue cases were collected based on data retrieved from local primary health center. All of dengue cases were diagnosed with dengue non-structural protein 1 (NS-1) antigen test. However, serotypes of infecting dengue virus were unknown. The study was conducted by installing 145 ovitraps in both the perimeter and buffer areas of the airport. We defined the zones as the following definitions: perimeter area is 100-m peripheral area from airport apron; buffer area is 400-m peripheral area from airport apron [9]. The study could only be performed at the northern and western area of the airport due to limited public access to the eastern and southern area.

Ovitrain Index (OI) is calculated from number of ovitraps with positive eggs divided by sum of ovitraps installed. Ovitraps were installed outdoors and indoors within 35–50 m distance. Positive OI and TTI of dengue virus were examined later on. After colonization of Aedes spp. eggs, TTI was identified using immunocytchemistry immunoperoxidase streptavidin biotin complex (IISBC) method in mosquito head squash preparations [10]. Rate of TTI was obtained by dividing the number of DENV-positive IISBC samples by total number of IISBC samples examined and was expressed as percentage [10]. The virus was identified through serotype examination by Lanciotti’s RT-PCR. Transovarial transmission in both Aedes aegypti and Aedes albopictus mosquitoes obtained from indoor and outdoor ovitraps were compared. The analysis of OI distribution was conducted using the software ArcGIS 9.2 (Esri, New York) to provide mapping and spatial reasoning to yield location-based data. The data were entered into nearest neighbor analysis, resulting in the distance between each feature centroid and its nearest neighbor’s centroid location. The z-score and p value are measures of statistical significance that are used in determining whether or not to reject the null hypothesis: features are randomly distributed.
The presence of high TTI value is considered an important mechanism for the maintenance of the virus in nature and may be associated in the occurrence of dengue epidemics and outbreak. High OI supports the potential of dengue outbreaks as well [11].

**Results**

Positive OI of *Aedes* spp. mosquitoes in the buffer area was 32 (45.1%), with outdoor ovitraps showing greater proportion (50%) compared to indoor ovitraps (37.9%). OI in the buffer area was 32 (45.1%), whereas OI in the perimeter area was 24 (32.4%) as shown in Table 1. The distribution of *Aedes* spp. mosquitoes based on positive OI in Yogyakarta’s Adisucipto Airport was found evenly distributed in the surrounding areas, with the highest number of positive OI found in the zone 4, the neighborhood 8 (Rukun Tetangga/RT 8) of village of Maguwoharjo (14 ovitraps or 9.6%), and the least number of positive OI found in zone 4 of perimeter area, the B Terminal (2 ovitraps or 1.4%) as shown in Fig. 1.

The nearest neighbor analysis shows that if the z-score is less than 1, the pattern exhibits clustering, whereas if the index is greater than 1, the trend is towards dispersion. The analysis of OI distribution resulted in z-score of −6.1 and p value of <0.001 that indicates the tendency of clustered OI distribution. The shortest distance of positive ovitraps was 3.6 m, while the farthest distance was 19 m with average distance of 13.2 m. The results indicate that the distance is still within the range of mosquito disease transmission.

Through identification by examination using immunohistochemistry (IHC) method, as many as 11 samples (18.9%) in the perimeter area and 21 samples (18.3%) in the buffer area of Yogyakarta’s Adisucipto Airport were found positive for transovarial transmission. The transovarial transmission was observed higher in percentage in *Aedes aegypti* mosquitoes, with 20% to 16.7% in the perimeter area and 20.3 to 16.1% in the buffer area (Table 2).

In *Aedes aegypti* mosquitoes, positive transovarial transmissions were more commonly detected in samples obtained from indoor-installed ovitraps: 23.5 to 17.4% in the perimeter area and 25 to 14.8% in the buffer area. On the other hand, positive transovarial transmissions in *Aedes albopictus* mosquitoes were all found in samples from outdoor-installed ovitraps, with 16.7% in the perimeter area and 16.1% in the buffer area. Unfortunately, we did not perform further analysis to identify statistical significance of this difference in findings between mosquito species. Distribution of dengue virus in the Adisucipto Airport of Yogyakarta was predominantly found in zone 4 of the buffer area, the RT 8 of village of Maguwoharjo (4 cases or 12.5%), and was least found in zone 2, the parking lot and common facility, and zone 3, RT 7 of village of Maguwoharjo, with both areas having only 1 case detected (3.1%). Dengue virus distribution tends to follow the distribution pattern of ovitraps. In both perimeter and buffer areas of the Adisucipto Airport, there were no dengue cases found. As described in Fig. 1, the shortest distance of dengue patients was 389 m, whereas the farthest distance was 1050 m.

The analysis of dengue virus distribution was conducted using the nearest neighbor analysis and resulted in z-score of −2.0 and p value of 0.04 that indicates the tendency of clustered distribution. The shortest distance of positive ovitraps was 3.1 m, while the farthest distance was 33 m with an average distance of 15.2 m.

Dengue virus serotype examination was performed in dengue-positive *Aedes* spp. mosquito-colonized samples and in positive transovarial transmission samples, all of which were coded in species and location. The serotype examination was conducted using Lanciotti’s RT-PCR method and resulted in the detection of DENV-2 and DENV-3 serotypes in *Aedes aegypti* and *Aedes albopictus* mosquitoes in Adisucipto Airport of Yogyakarta (Table 3). Data analysis on dengue virus serotype distribution in Yogyakarta’s Adisucipto Airport resulted in z-score of 1.4 and p value of 0.14 that indicates tendency of random dengue virus serotype distribution. The shortest distance of dengue virus serotype was 33 m, while the farthest distance was 173 m with average distance of 94.1 m as shown in Fig. 2.

**Discussion**

Ovitrap Index (OI) in the buffer area was higher compared to that in the perimeter area. The results showed that the buffer areas have more potential as breeding sites of *Aedes* spp. mosquitoes. The distribution of *Aedes* spp. mosquitoes based on positive OI in Yogyakarta’s Adisucipto Airport was found evenly distributed in the surrounding areas, with the highest number of positive OI found in the zone 4, the neighborhood 8 (Rukun Tetangga/RT 8) of village of Maguwoharjo. OI of <5% shows that the distribution of *Aedes* spp. mosquitoes is least spread, while OI of 5–20% shows that the *Aedes* spp. mosquitoes are moderately spread in the study area [12]. OI of <10% indicates that the area has no risk in the development of dengue outbreak, whereas OI of >10% indicates that the area has the potential for developing

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**Table 1** Distribution of ovitrap installation in Adisucipto Airport of Yogyakarta

| Location | Situation | (+) Eggs | OI (%) |
|----------|-----------|----------|--------|
|          | Indoor    | Outdoor  | Indoor | Outdoor |
| Perimeter area | 40 | 34 | 12 | 12 | 30 | 35.3 |
| Total number | 74 | 24 | 32.4 |
| Buffer area | 29 | 42 | 11 | 21 | 37.9 | 50 |
| Total number | 71 | 32 | 45.1 |
dengue outbreak [13]. In the area of Yogyakarta’s Adisucipto Airport, OI of 45.1% was found, which indicates potential for developing dengue outbreak.

The analysis of OI distribution resulted in z-score of −6.1 and p value of < 0.001 that indicates the tendency of clustered OI distribution. Through identification by using IHC method, the transovarial transmission was observed higher in percentage in *Aedes aegypti* mosquitoes in the perimeter area compared to that in the buffer area. The percentage alone presumes that *Aedes aegypti* mosquitoes may have a more dominant role in the transmission of dengue in the area of Adisucipto Airport of Yogyakarta. A similar study in Manado, Indonesia, also showed that the TTI in *Aedes aegypti* mosquitoes is significantly higher than that in *Aedes albopictus* mosquitoes [14]. Previous research established that *Aedes aegypti* mosquitoes have a major role in human dengue transmission [15]. A TTI rate of 20% allows the maintenance of highly stable vertical infection that persists for several generations [16].

Positive transovarial transmissions in *Aedes aegypti* mosquitoes were more commonly detected in samples obtained from indoor-installed ovitraps in the perimeter area, and the perimeter area had a higher TTI compared to the buffer area. This finding highlights the importance of vector control strategies in reducing the transmission of dengue in the area of Adisucipto Airport of Yogyakarta.

**Table 2** Analysis of transovarial transmission of *Aedes* spp. mosquitoes in Adisucipto Airport of Yogyakarta

| Location       | Immunohistochemistry (IHC) | Transovarial transmission | % transovarial transmission |
|----------------|-----------------------------|---------------------------|----------------------------|
|                | *Ae. aegypti* | *Ae. albopictus* | *Ae. aegypti* | *Ae. albopictus* | *Ae. aegypti* | *Ae. albopictus* |
| Perimeter area | 40 | 18 | 8 | 3 | 20 | 16.7 |
| Total number   | 58 | 11 | 12 | 9 | 20.3 | 16.1 |
| Buffer area    | 59 | 56 | 12 | 9 | 20.3 | 16.1 |
| Total number   | 115 | 21 | 21 | 21 | 21 | 21 |

Negative ovitraps and dengue patients (sufferers) distribution may also be observed.
area. The results suggest that *Aedes aegypti* mosquitoes tend to do oviposition indoor rather than outdoor. A study in three localities in Malaysia also showed that ovitraps were more likely to yield positive results if installed indoor [17].

Positive transovarial transmissions in *Aedes albopictus* mosquitoes were all found in samples from outdoor-installed ovitraps. Distribution of dengue virus in the Adisucipto Airport of Yogyakarta was predominantly found in zone 4 of the buffer area, the RT 8 of village of Maguwoharjo. Dengue virus distribution tends to follow the distribution pattern of ovitraps. In both perimeter and buffer areas of the Adisucipto Airport, there were no dengue cases found. This result may be due to lack of good host immunity, higher activities of humans in the airport so that the mosquitoes were not able to bite very often [18], and low number of *Aedes* spp. mosquitoes in the Adisucipto Airport: 378 mosquitoes in ± 560,000 m² perimeter and buffer areas. Other possible contributing factor is low TTI of dengue virus (18.9% in perimeter area and 18.3% in buffer area). These TTI values are considered low as the study was conducted in endemic area and in rainy season, during which dengue transmission is on its peak; therefore TTI values should exceed 30% [19]. As described in Fig. 1, the shortest distance of dengue patients was 389 m, whereas the farthest distance was 1050 m.

The analysis of dengue virus distribution was conducted using the nearest neighbor analysis and resulted in a z-score of −2.0 and p value of 0.04 that indicates the tendency of clustered distribution. The shortest distance of

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**Table 3** Serotypes of dengue virus in *Aedes* spp. mosquitoes in Adisucipto Airport of Yogyakarta

| Location | RT-PCR | DENV serotype |
|----------|--------|---------------|
| Perimeter area | | |
| 17a | 12 | DENV 2 |
| 54a | 2 | DENV 2 and 3 |
| Buffer area | | |
| 110a | 12 | DENV 2 and 3 |
| 119a | 5 | DENV 2 and 3 |
| 132a | 7 | DENV 2 and 3 |
| 134a | 4 | DENV 2 and 3 |
| 136a | 2 | DENV 2 |

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**Fig. 2** Distribution map of dengue virus and dengue cases in Adisucipto Airport Yogyakarta. Four zones in perimeter area are cargo log (zone 1), terminal A (zone 2), office (zone 3), and terminal B (zone 4), while four areas in buffer area are terminal B (zone 1), parking and public facilities (zone 2), neighborhood 7 (zone 3), and neighborhood 8 (zone 4). One spot was positive in perimeter area for DENV-2 and one spot positive for DENV-2 and DENV-3, both of which were also positive for *Ae. albopictus*. One spot was positive in buffer area for DENV-2 and two spots were positive for DENV-2 and DENV-3, all of which were also positive for *Ae. albopictus*. There were two spots positive in buffer area for DENV-2, DENV-3, and *Ae. aegypti*. ---
positive ovitraps was 3.1 m, while the farthest distance was 33 m with average distance of 15.2 m. Although no specific threshold values have been established for each arbovirus, absence of severe dengue cases in Thailand was noted when the density of *Ae. aegypti* eggs per ovitrap per week was less than two [20]. Moreover, despite using a different ovitrap, DENV transmission occurred in Taiwan when the density of eggs per house (two ovitraps per house) was around two [21]. In determining transmission of dengue virus, TTI is important in showing that dengue virus from infected female mosquitoes may be spread into the ovarii and transmitted to the next generation, whereas OI correlates with mosquito population and represents true mosquito infestations in the area [11, 22].

Dengue virus serotype examination was performed in dengue-positive *Aedes* spp. mosquito-colonized samples and in positive transovarial transmission samples, all of which were coded in species and location: DENV-2 and DENV-3. Similar results were obtained in Bantul, Indonesia, which showed the predominant serotypes of DENV-3. Studies conducted in positive transovarial transmission samples, all of which were coded in species and location: DENV-2 and DENV-3. Similar results were obtained in Bantul, Indonesia, which showed the predominant serotypes of DENV-3. Studies conducted in Taiwan [24] and Singapore [25] showed that DENV-2 was the predominant serotype found. In areas with high dengue endemicity, the predominant serotypes were DENV-2 and DENV-3, while in areas with low dengue endemicity the prevalent serotypes were DENV-1 and DENV-2 [21]. A similar study conducted in 2007 to 2009 in Brazil reported that the highest number of mosquito larvae found was *Aedes aegypti* (3417 mosquitoes or 91%), followed by *Aedes albopictus* (336 mosquitoes or 9%), with detected serotypes of DENV-3 *Aedes albopictus*, DENV-2 *Aedes aegypti*, and DENV-2 and DENV-3 *Aedes albopictus* [26].

The study has several limitations. First, the study was conducted only once during rainy season, which would have been better if performed simultaneously in other season as well. Second, study area was limited only to areas open for public access, not including those areas that belonged to the Air Force and not open for public.

**Conclusion**

The study shows that buffer areas of Adisucipto Airport of Yogyakarta have higher risk as breeding sites for *Aedes* spp. mosquitoes, predominantly DENV-2 and DENV-3 serotypes. OI of 45.1 indicates that the area has the potential for developing dengue outbreak.

**Abbreviations**

BII: Brieux Index; DENV-1: Dengue virus serotype-1; DENV-2: Dengue virus serotype-2; DENV-3: Dengue virus serotype-3; DENV-4: Dengue virus serotype-4; HI: House Index; IHR: International Health Regulations; iiSBC: Immunochemistry immunoperoxidase streptavidin biotin complex; NS-1: Non-structural protein; OI: Ovitrap Index; PHEIC: Public Health Emergency of International Concern; RT: Rulkun Tetangga (neighborhood); RT-PCR: Reverse transcriptase-polymerase chain reaction; TTI: Transovarial Transmission Index; WHO: World Health Organization

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**Availability of data and materials**

Data sharing is not applicable to this article as no datasets were generated or analyzed during the current study. The study was a part of AL’s thesis. The data used in this article are not published elsewhere.

**Authors’ contributions**

TBTS carried out the molecular genetic studies, conceived of the study, and participated in its design and coordination and in drafting the manuscript. AL carried out the field works and data analysis, while SDA participated in the surveillance. HKJ participated in the design of the study and performed the statistical analysis. BSW has provided assistance in designing and reviewing the map. All authors have read and approved the final manuscript.

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TBTS is a member of Indonesian Entomological Association and is active in research in mosquito-borne diseases. AL and SDA are surveys in Port Quarantine. HKJ is a professor in Public Health. BSW is an expert in map drawing.

**Ethics approval and consent to participate**

Ethical approval was obtained from the Medical and Health Research Ethics Committee (MHREC), Faculty of Medicine Universitas Gadjah Mada, recognized by FERCAP.

**Competing interests**

The authors declare that they have no competing interests.

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