Seroprevalence of mumps before the introduction of mumps-containing vaccine in Lao PDR: results from a nationwide cross-sectional population-based survey

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Abstract
Objective: Mumps-containing vaccine is currently not a component of the national immunization schedule in Lao People’s Democratic Republic (PDR). Mumps itself is not a notifiable disease in the country and the seroprevalence of anti-mumps immunoglobulin G (IgG) in the general population is unknown. In this study, anti-mumps IgG was measured in 2058 blood samples to evaluate population immunity in the country.

Results: The seroprevalence of anti-mumps IgG showed a gradual increase with increasing age, starting at 10.6% (95% CI 7.4–13.7) in participants aged 1–2 years, and almost plateaued at about 75% in individuals older than 11–12 years, though it still tended toward a small increase up to 89.6% (95% CI 86.6–92.6) in participants aged 40 years or older. Compared with the results of previous studies, this increase with increasing age is less marked and the plateau of anti-mumps seroprevalence is lower. We attribute this result mainly to the lower population density in Lao PDR.

Keywords: Seroprevalence, Mumps, Pre-vaccination era, Nationwide cross-sectional population-based survey

Introduction
Mumps is a common childhood viral infectious disease that is preventable by vaccination. Live mumps vaccines are available as monovalent (mumps only), bivalent measles-mumps (MM), and trivalent measles-mumps-rubella (MMR) vaccines, and the effectiveness of at least one dose of MMR in preventing clinical mumps in children is estimated to be around 70–80% [1]. In total, 122 countries have introduced one of the mumps-containing vaccines nationwide as of 2017 [2].

The majority of studies on anti-mumps antibody seroprevalence in the pre-vaccination era were conducted either in European countries before 1990 [7–10] or outside Europe [11–13]. However, the participants were not randomly selected but involved specific populations or were randomly selected but did not involve a nationwide population. To our knowledge, no studies have investigated the nationwide seroprevalence of anti-mumps antibody in a pre-vaccination era.

Lao People’s Democratic Republic (PDR) is a small developing country in Southeast Asia. It is landlocked and mountainous, and its road infrastructure is underdeveloped. The expanded program on immunization (EPI) was started in Lao PDR in 1984 but does not currently include the mumps-containing vaccine. Sporadic outbreaks of mumps have been reported, but mumps is not...
a notifiable disease as yet. A seroprevalence study of anti-mumps antibody has been conducted for specific populations in some areas [14], but nationwide age-specific seroprevalence in the general population is unknown.

In 2011, Lao PDR conducted a nationwide supplemental immunization activity (SIA) using the measles and rubella (MR) combined vaccine targeted at children aged 9 months to 19 years. In 2014, a nationwide multistage random cluster sampling survey was conducted to evaluate the SIA by determining anti-measles and anti-rubella IgG seroprevalence among children and adults [15].

This study sought to estimate the age-specific seroprevalence of anti-mumps IgG in the nationwide general population of Lao PDR, where mumps antigen-containing vaccine is currently not included in the national immunization schedule, and analyzed the same blood samples from the 2014 survey.

Main text

Materials and methods

Study population and sampling

In this study, we used the same blood samples remaining from a nationwide multistage cluster sampling survey that we conducted in 2014 to measure anti-measles and anti-rubella IgG and evaluate the effectiveness of the SIA in 2011 [15]. The 2014 study used three-stage random cluster sampling with probability proportionate to size sampling based on the 2005 population census conducted by the Department of Statistics, Lao PDR. In total, 26 of 143 districts and 2 villages from the respective selected districts were selected, then 42 participants from each of the selected villages were selected. Participants aged 3–4 years were excluded because of indeterminate immunization history and date of birth due to inconsistencies between calendar and traditional ages that often occur in rural areas [16, 17]. From 2184 expected participants, blood samples were obtained from 2153 (97.8% of the required sample size; males 44.8%; mean age 23.2 years, age range 1–81 years). Blood sampling (97.8% of the required sample size; males 44.8%; mean age 23.2 years, age range 1–81 years). Blood sampling was performed using dried blood spots from finger prick blood spots onto Whatman® 903 Protein Saver filter paper (Whatman, Maidstone, Kent, UK) and transported to the National Institute of Infectious Diseases, Japan, within a few weeks [18, 19].

Anti-mumps IgG measurement

In the 2014 study, anti-measles IgG and anti-rubella IgG levels were evaluated for 2135 samples after exclusion of 18 samples due to missing vital information from 2153 samples collected between January and February. The remaining samples were stored at 4 °C as dried blood spots collected on the filter paper. Of these 2135 samples, 2058 had enough dried blood spots to measure anti-mumps IgG. Blood samples were extracted from the dried blood spots from May to July 2015 and stored at −80 °C. Measurements were made from April to July 2017 with a commercially available enzyme-linked immunosorbent assay (ELISA) kit (Enzygnost® Anti-Parotitis Virus/IgG, Siemens Healthcare Diagnostics, Eschborn, Germany) according to the manufacturer’s instructions. Anti-mumps IgG levels were determined as positive (ΔA > 0.200), equivocal (0.100 ≤ ΔA ≤ 0.200), or negative (ΔA < 0.100) according to the manufacturer’s instructions. Only the samples determined as positive were regarded as ‘positive’ for analysis.

Data entry and statistical analysis

All collected data were double-entered and cleaned in Microsoft Excel 2016 spreadsheet. Statistical analysis was performed and IgG seroprevalence was calculated using STATA versions 13 and 14 (Stata Corp., College Station, TX).

Results

The seroprevalence of anti-mumps IgG for participants aged 1–2 years was the lowest among all age groups, at 10.6% (95% CI 7.4–13.7), and showed a gradual increase with increasing age (Fig. 1). That for participants aged 5–6 years, 7–8 years, and 9–10 years was 33.3% (95% CI 21.1–45.6), 44.3% (95% CI 32.4–56.2), and 48.4% (95% CI 35.6–61.2), respectively. Anti-mumps IgG seroprevalence reached 75.4% (95% CI 64.3–86.5) for those aged 11–12 years and almost plateaued; however, it still tended toward a small increase with increasing age in those older than 13–14 years though it fluctuated in the 70–80% range until those for aged 30–34 years. The seroprevalence of anti-mumps IgG for participants aged 35–39 years was 83.6% (95% CI 77.1–90.1) and reached 89.6% (95% CI 86.6–92.6) for those aged 40 years or older.

Discussion

Previous studies on age-specific seroprevalence of anti-mumps IgG before the introduction of mumps-containing vaccines have been conducted in Europe [7–10] and other countries [11–13], but ours is the first nationwide study to be conducted in Lao PDR. Our study revealed a rather moderate degree of increase in IgG seroprevalence by age until age 11–12 years when it almost plateaued at about 75%. The seroprevalence tended toward a small increase after age 13–14 years and reached almost 90% in age 40 years or older. Another study in Lao PDR, which targeted elementary to high school students in four provinces, showed findings similar to our results, such as 50.0% in age group 5–9 years, 65.6% in age group 10–14 years, and 66.9% in age group 15–19 years [14].
In contrast, previous studies in other countries showed a sharp increase in IgG seroprevalence and a higher plateau, where the increase started during infancy and rose sharply to about 80% before age 10 years and then gradually increased after around age 10 to reach almost 90% before age 30. The seroprevalence tended toward a small increase fluctuating in the 90–100% range after age 30 [7, 9–13]. To suggest an explanation for our results in Lao PDR, a comparison with these earlier studies is helpful. In Wagenvoort et al.’s seroprevalence study of mumps immunity before introduction of mumps-containing vaccine in the Netherlands, the estimated population density of the study area was more than the national mean population density of 409 persons per km² [8], and in Liu et al.’s study in China, the population density was 767 persons per km² [13]. Lao PDR had a reported population density of 27 persons per km² in 2015 [20], which is much lower than that in previous studies. Low population density implies less human-to-human contact and this may be the reason for the less marked increase in anti-mumps IgG seroprevalence levels and lower plateau between teens and thirties in Lao PDR. Arroyo et al.’s study among unvaccinated children in Spain revealed that rural environment, not attending school, and lack of brothers were associated with significantly lower levels of seroprevalence in the 3- to 5-year-old age group [9]. However, to our knowledge, few studies have investigated the direct relationship between anti-mumps IgG seroprevalence levels and population density before introduction of mumps-containing vaccine. Further study is necessary to investigate the direct relationship between seroprevalence levels and population density and other factors related to human-to-human contact such as school attendance and number of siblings.

The results of the present study show that many adolescents and adults are susceptible to mumps in Lao PDR, suggesting that they might be infected if a mumps outbreak were to occur. Complications of mumps among adolescents and adults are commonly seen. Orchitis occurs in about 35% of postpubertal men and oophoritis is observed in about 5% of adult women [21]. Our results provide essential information that can be utilized for planning the introduction of the MMR vaccine in the future. In addition, however, it is necessary to add mumps to the surveillance system because it is currently not a notifiable disease. Further examination of data on the incidence of mumps by age group is also needed to help estimate the burden of the disease [22].

In conclusion, this is the first study to describe the age-specific seroprevalence of anti-mumps IgG in a nationwide general population in Lao PDR and, to our knowledge, it is the first study worldwide to describe this seroprevalence in such a population before the introduction of mumps-containing vaccine into a national immunization schedule. A study to investigate age-specific seroprevalence of anti-mumps IgG offers valuable information for planning the introduction of a mumps-containing vaccine such as MMR.
Limitations
Firstly, we designed sampling based on the 2005 population census which was rather old though it was the latest data officially available of the time. The population growth or mobility after 2005 may affect representation of the whole population of the country at the time of the survey.

Secondly, measurement of anti-mumps IgG levels were performed about three and half years after sampling. The period of the storage at 4 °C about 15 months as dried blood spots before extraction of blood samples could affect the levels of IgG. However, the previous study reported that dried blood spots stored at 4 °C up to 17 months less likely affected sensitivity and specificity for measurement of the antibody [23].

Thirdly, numerous ELISA kits are commercially available and have been used in previous studies investigating anti-mumps IgG seroprevalence [3–6, 13]. However, the cutoff value for anti-mumps IgG levels measured using these kits is not standardized. In the present study, we followed the manufacturer’s instructions to determine the IgG levels as positive, but other studies used different quantitative ELISA values with the same Enzygnost® ELISA kit (Siemens Healthcare Diagnostics) [6]. Therefore, direct comparison of age-specific mumps IgG seroprevalence is not conclusive. To better assess the seroepidemiological data, cutoff values and methods for comparison among different test kits should be standardized.

Abbreviations
CI: confidence interval; ELISA: enzyme-linked immunosorbent assay; EPI: expanded program on immunization; IgG: immunoglobulin G; MM: measles-mumps; MMR: measles-mumps-rubella; MR: measles-rubella; SIA: supplemental immunization activity.

Authors’ contributions
MH, TI, MT, MK, KKomas, and YM conceptualized and designed the study. TK, SM, KF, and CP collected samples and data, and MH and AX supervised the survey. MT, MK, KKomas, YM, KM, and MN performed laboratory work, and KKomas and HO analyzed the data. HO prepared the draft manuscript and MH, KKomas, SM, TK, MT, MK, KKomas, and YM critically revised it. All authors read and approved the final manuscript.

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Competing interests
The authors declare that they have no competing interests.

Availability of data and materials
The datasets analyzed in the present study are available from the corresponding author on reasonable request.

Consent for publication
Not applicable.

Ethics approval and consent to participate
The study protocol for analysis of anti-mumps IgG levels was approved by the ethics committees of the National Center for Global Health and Medicine Japan (NCIM-G-002095-00) and the National Institute of Infectious Diseases, Japan (NIID-714). The 2014 study protocol was approved by the ethics committees of the National Center for Global Health and Medicine, Japan (NCIM-G-001459-00) and the National Institute of Infectious Diseases, Japan (NIID-494). Both protocols were submitted to the ethics committee of the Ministry of Health, Lao PDR for approval in a combined submission in Lao PDR and were approved at the same time (025-NECHR). All selected participants during the 2014 survey had provided written informed consent. For participants aged under 15 years, consent was obtained from their parents or legal guardians.

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References
1. Demicheli V, Rivetti A, Debalini MG, Di Pietrantonj C. Vaccines for measles, mumps and rubella in children. Cochrane Database Syst Rev 2012;2:CD004407. https://doi.org/10.1002/14651858.cd004407.pub3.
2. WHO. Global immunization coverage 2016. http://www.who.int/en/news-room/factsheets/detail/immunization-coverage Accessed 24 Jan 2019.
3. Muhsen K, Shohat T, Aboudy Y, Mendelson E, Algor N, Anis E, et al. Sero-prevalence of mumps antibodies in subpopulations subsequenty affected by a large scale mumps epidemic in Israel. Vaccine. 2011;29(22):3878–82. https://doi.org/10.1016/j.vaccine.2011.03.047 (Epub 2011 Mar 29).
4. Poethko-Müller C, Mankertz A. Seroprevalence of measles-, mumps- and rubella-specific IgG antibodies in German children and adolescents and predictors for seronegativity. PLoS ONE. 2012;7(8):42867. https://doi.org/10.1371/journal.pone.0042867 (Epub 2012 Aug 6).
5. Plans P. New preventive strategy to eliminate measles, mumps and rubella from Europe based on the serological assessment of herd immunity levels in the population. Eur J Clin Microbiol Infect Dis. 2013;32(7):961–6. https://doi.org/10.1007/s10096-013-1836-6 (Epub 2013 Feb 16).
6. Leundan E, Maertens K, Wautier M, Hutse V, Theeten H. Susceptibility to measles, mumps, and rubella in 5-year-old children in Flanders, Belgium. Eur J Pediatr. 2015;174(7):925–32. https://doi.org/10.1007/s00431-014-2481-5 (Epub 2015 Jan 13).
7. Edmunds WJ, Gay NJ, Kretzschmar M, Pebody RG, Wachmann H, ESEN Project. European Sero-epidemiology Network. The pre-vaccination epidemiology of measles, mumps and rubella in Europe: implications for modelling studies. Epidemiol Infect. 2000;125(3):635–50.
8. Wagenvoort JH, Harmsen M, Boutahar-Trouw BJ, Kraijeeveld CA, Winkler KC. Epidemiology of mumps in the Netherlands. J Hyg. 1980;85(3):313–26.
9. Arroyo M, Aller JM, Mateos ML, Carasco JL, Ballesteros F, Lardinos R. Natural immunity to measles, rubella and mumps among Spanish children in the pre-vaccination era. Int J Epidemiol. 1985;14(1):95–100.
10. Nokes DJ, Wright J, Morgan-Capner P, Anderson RM. Serological study of the epidemiology of mumps virus infection in north-west England. Epidemiol Infect. 1990;105(1):175–95.
11. Cox MJ, Anderson RM, Bundy DA, Nokes DJ, Didier JM, Simmons I, et al. Seroepidemiological study of the transmission of the mumps virus in St. Lucia, West Indies. Epidemiol Infect. 1989;102(1):147–60.
12. Akst S, Egemen A, Ozacar T, Kurugol Z. Mumps seroprevalence in an unvaccinated population in Izmir, Turkey. Acta Paediatr. 2000;89(3):370–1.
13. Liu Y, Hu Y, Deng X, Wang Z, Lu P, Ma F, et al. Seroepidemiology of mumps in the general population of Jiangsu province, China after introduction of a one-dose measles-mumps-rubella vaccine. Sci Rep. 2015;5:14660. https://doi.org/10.1038/srep14660.
14. Hübchen JM, Wilvong K, Souvannaso C, Black AP, Lütteke N, Samountry B, et al. High prevalence of mumps in Lao People’s Democratic Republic. Clin Microbiol Infect. 2014;20(10):O664–71. https://doi.org/10.1111/1469-0691.12586 (Epub 2014 Mar 6).
15. Hachiya M, Miyano S, Morii Y, Vunmnyey E, Keungsaneth P, Vongphrachanh P, et al. Evaluation of nationwide supplementary immunization in Lao People’s Democratic Republic: population-based seroprevalence survey of anti-measles and anti-rubella IgG in children and adults, mathematical modelling and a stability testing of the vaccine. PLoS ONE. 2018;13(3):e0194931. https://doi.org/10.1371/journal.pone.0194931 (eCollection 2018).
16. Xeuatvongsa A, Komada K, Kitamura T, Vongphrachanh P, Pathammavong C, Phounphenghak K, et al. Chronic hepatitis B prevalence among children and mothers: results from a nationwide, population-based survey in Lao People’s Democratic Republic. PLoS ONE. 2014;9(2):e88829. https://doi.org/10.1371/journal.pone.0088829 (eCollection 2014).
17. Komada K, Sugiyama M, Vongphrachanh P, Xeuatvongsa A, Khampaphongphane B, Kitamura T, et al. Sero-prevalence of chronic hepatitis B, as determined from dried blood spots, among children and their mothers in central Lao People’s Democratic Republic: a multistage, stratified cluster sampling survey. Int J Infect Dis. 2015;36:21–6. https://doi.org/10.1016/j.ijid.2015.04.020 (Epub 2015 May 6).
18. Uzcainan A, Lubega J, Nanunya M, Mercader S, Rota P, Bellini W, et al. Dried blood spots on filter paper as an alternative specimen for measles diagnostics: detection of measles immunoglobulin M antibody by a commercial enzyme immunoassay. J Infect Dis J Infect Dis. 2011;204(Suppl 1):S564–9. https://doi.org/10.1093/infdis/jir088.
19. Riddell MA, Byrnes GB, Leydon JA, Kelly HA. Dried venous blood samples for the detection and quantification of measles IgG using a commercial enzyme immunoassay. Bull World Health Organ. 2003;81(10):701–7 (Epub 2003 Nov 25).
20. Lao Statistics Bureau. Results of Population and Housing Census 2015. https://lao.unfpa.org/sites/default/files/pub-pdf/PHC-ENG-FNAL-WEB_0.pdf Accessed 24 Jan 2019.
21. Cohen J, Powderly WG. Infectious Diseases Second Edition. 2004, p.1900.
22. Galazka AM, Robertson SE, Kraigher A. Mumps and mumps vaccine: a global review. Bull World Health Organ. 1999;77(1):13–14.
23. Riddell MA, Leydon JA, Catton MG, Kelly HA. Detection of measles virus-specific immunoglobulin M in dried venous blood samples by using a commercial enzyme immunoassay. J Clin Microbiol. 2002;40(1):5–9.