BMJ Open  Serological investigation to identify risk factors for post-flood infectious diseases: a longitudinal survey among people displaced by Typhoon Morakot in Taiwan

Chun-Yu Lin, Tun-Chieh Chen, Chia-Yen Dai, Ming-Lung Yu, Po-Liang Lu, Jeng-Hsien Yen, Yen-Hsu Chen

ABSTRACT

Objectives: After Typhoon Morakot struck Taiwan in 2009, thousands of Taiwanese citizens were displaced to shelters for several weeks. This study aimed to investigate serological status in both groups and identify risk factors associated with seroconversion of infectious diseases.

Design: A longitudinal survey.

Setting: All experimental and clinical investigations were performed in a tertiary teaching hospital.

Participants: A total of 288 displaced persons (96 males and 192 females) were recruited and complete follow-up data through two rounds of sampling were collected. The average age was 58.42 years (range 31–87 years).

Interventions: First, serum specimens were collected between December 2009 and January 2010, 4–5 months after the typhoon. The second round of specimen collection was carried out after 6 months.

Primary and secondary outcome measures: The primary outcome measured was serological status of vaccine-preventable droplet-borne infectious diseases (measles, mumps, rubella) and water-borne diseases (amoebiasis and leptospirosis). The secondary outcome was identification of risk factors for seroconversion using univariate and multivariate analyses.

Results: Complete data were available for all 288 displaced persons (114 from the shelter group; 174 from the community group). Seroconversion of Entamoeba histolytica was observed in 128 (44.4%) participants, with a significantly higher rate in the shelter group than in the community group (56.1% vs 36.8%; p=0.001). There were 10 cases of rubella seroconversion. After adjusting for medical history, hypertension and hyperlipidaemia, shelter stay was associated with higher risk for seroconversion (OR=2.055, 95% CI 1.251 to 3.374; p=0.004). Amoebiasis was more evident in the shelter group, although the manifestations were mild.

Conclusions: Our results suggested that (1) a clean water supply is essential postdisaster, especially in crowded shelters, and (2) vaccination programmes should be extended to populations at higher risk for post-disaster displacement or to those with weakened immune status.

INTRODUCTION

Climate change, resulting from an imbalance between radiation entering and exiting the atmosphere, is thought to alter the hydrological cycle. In the past few decades, climate change and global warming have been suggested as the triggers for increased severity of cyclones, flooding and major hurricane-related damage, even in temperate regions. Flooding, which is the most common type of disaster globally, has significantly impacted places such as Hunan, China, where river flooding has caused an increase in mortality rates as well as potential years of lost life.

Natural disasters such as typhoons, cyclones, earthquakes, tsunamis or flooding are also associated with an increased incidence of vector-borne and water-borne infectious diseases. Indeed, a number of reports have suggested that large-scale cyclones with extensive flooding impose a burden on the public health systems of affected countries.
health system due to outbreaks of infectious diseases such as cholera, typhoid fever, acute hepatitis A, leptospirosis, melioidosis, measles, dengue fever, malaria and amoebiasis. An important contributory factor for such disease outbreaks may be the displacement of large numbers of people from their homes into overcrowded shelters where supplies may be limited. The availability and accessibility of medical services after such natural disasters is also a major concern. Epidemiological studies have reported that the increased incidence of infectious diseases after such disasters is associated with environmental changes, crowded shelter conditions and an inadequate or dirty water supply. Risk behaviours linked to sexually transmitted diseases also may increase after a disaster as a result of the unaccompanied status of primary sex partners.

Typhoon Morakot struck southern Taiwan between 7 and 9 August 2009, with massive rainfall (>1000 mm) and flooding, resulting in 673 deaths and property losses of more than 19 billion New Taiwan Dollars. In contrast to tsunamis, where flooding occurs along coastal areas, the major rainfall and flooding during Typhoon Morakot occurred along coastal areas as well as inland areas. Southern Taiwan had the most severe damage in mountainous and coastal areas. Thousands of Taiwanese citizens were displaced from their original residences in coastal, plains and mountainous areas, and moved to shelters or to urban communities. This movement made the collection and accuracy of medical records unreliable. Owing to these challenges, no longitudinal population-based surveys were conducted either during or after Typhoon Morakot to examine the occurrence of infectious diseases among displaced people. It is also not clear which post-disaster shelter policies should be examined with caution when drafting guidelines for prevention of infectious diseases.

In order for government agencies to prepare more effectively for post-disaster management, it is important: (1) to understand differences between disaster-displaced persons staying in shelters and those remaining in the community, and (2) to collect adequate and accurate population-based serological epidemiological evidence of infectious diseases after such disasters.

In this longitudinal study, we performed serological testing to track communicable diseases in two groups of people displaced from their homes: those who were moved to shelters, and those who remained in the community. The major aims of this study were: (1) to investigate the serological status of vaccine-preventable droplet-borne and water-borne infectious diseases in displaced persons staying in shelters, and those in the community, and (2) to identify risk factors associated with seroconversion of infectious diseases.

**MATERIALS AND METHODS**

This longitudinal epidemiological survey of 288 adult flood-displaced persons was conducted between December 2009 and July 2010, during the post-flood period 4–11 months after Typhoon Morakot in Taiwan. People living in the mountainous areas who were moved for longer than 2 weeks into shelters equipped with beds and a water supply were categorised as the shelter group. Those who had lived in a shelter for less than 2 weeks were excluded from further analysis. In contrast, storm-displaced persons who did not move into shelters, but went to live with their families in urban areas, were categorised as the community group. All 288 cases were from the Kaohsiung County. Hepatitis A vaccination was provided to those people who lived in the mountainous areas for several years before this typhoon. The Institutional Review Board of Kaohsiung Medical University Hospital reviewed and approved the study protocol, and all participants provided written informed consent. People younger than 20 years old were excluded from the study.

**Study design**

We sampled the first serum specimens between December 2009 and January 2010, 4–5 months after the typhoon. We performed second-round testing after 6 months. At the first and second sampling, each participant was asked to complete a structured questionnaire, either individually or with the help of trained staff during a face-to-face interview. The first questionnaire (for first sampling) included epidemiological data, underlying diseases and severity of trauma during the typhoon (family member’s death or major trauma). The second questionnaire (for second sampling) included duration of shelter stay before returning to one’s original residence.

**First-round sampling**

The first-round specimens were analysed for biochemistry data and nutritional status. Serological tests included IgG levels for vaccine-preventable diseases (measles, mumps and rubella, which all require mandatory notification), *E. histolytica* serological titres (IgG using IHA (indirect haemagglutination)) and IgM/IgG for leptospirosis. We also measured IgG levels of herpes simplex virus (HSV)-1, HSV-2 and Venereal Disease Research Laboratory, for syphilis. Amoebiasis (IHA), HSV-1 and HSV-2 tests were performed by the College of American Pathologists (CAP) Accreditation Programmes Laboratory. All other tests were performed in our own laboratory, which is a Biosafety level-2 (BSL-2) certified lab.

**Table 1** lists the details of the serological test kits used in the present study. Patients with positive syphilis screening results were contacted between first-round and second-round specimen collection, their serum was reanalysed and they were provided adequate treatment. No scheduled vaccination programme was provided to the shelter group or to the community group between the two samplings, except for vaccination for the 2009 pandemic H1N1 influenza strain.
**Second-round sampling**

Six months after the first sampling, a second-round serum sampling was conducted to evaluate for measles, mumps, rubella and *E. histolytica*, using the same methods. If a participant had negative results for measles, mumps and/or rubella in the first round, and positive results in the second, we defined this as an IgG seroconversion. (If a participant had positive results for measles, mumps and/or rubella in the first-round already, he or she did not receive recheck of measles, mumps and/or rubella IgG in the second-round). The cut-off value of positive results of the IHA test for amoebiasis was previously established as ≥1:128.\(^{19,20}\) However, there is evidence that this cut-off value may miss some cases with parasitologically positive amoebiasis.\(^{21,22}\) Therefore, with two blood samples collected at an interval of at least 16 weeks, we defined IgG seroconversion for amoebiasis as ≥1:64 at the second sampling as well as a fourfold titre increase when compared to first sampling results, which indicates recent infection.\(^{23}\)

**Follow-up telephone interview**

Telephone interviews were conducted for all displaced persons with IHA seroconversion within 1 month after the second sampling. The structured questionnaire that was used mainly inquired about gastrointestinal symptoms and signs of amoebiasis, including diarrhoea, abdominal pain and dysentery.

**Statistical analysis**

Data were summarised as n (%) for categorical variables, mean±SD for continuous variables and median (IQR: Q1, Q3) for continuous variables with non-normal distribution. Between-group differences in categorical data were compared by Pearson \(\chi^2\) test or Fisher’s exact test if case numbers were less than five. Between-group differences in continuous data were compared by two-sample t test and by Mann-Whitney U test if data were without normal distribution. Associations between seroconversion among shelter stay, participants’ demographics, medical history and laboratory data were further identified. Variables significantly associated with seroconversion were placed into univariate and multivariate logistic regression analysis. Results were presented as ORs and corresponding 95% CIs of OR (95% CI) and p values. All statistical assessments were two-tailed and \(p<0.05\) represented statistical significance. All analyses were conducted using SPSS statistical software V.18.0 (SPSS Inc, Chicago, Illinois, USA).

**RESULTS**

**Demographic characteristics and laboratory data**

A total of 288 participants (96 males/192 females) with complete follow-up data through two rounds of sampling were enrolled into evaluation. The average age was 58.42 years (SD=12.73) with a range of 31 to 87 years. Table 2 summarises the demographics, medical history and laboratory data of participants in the shelter group, and those in the community group. There was no significant difference in the distribution of demographics and medical histories between the two groups (all \(p>0.05\)). Although there was a difference in age between the shelter and community groups, this difference was of borderline significance (median (IQR): 55 (48, 66) vs 57 (50, 70), \(p=0.049\)). The second round of testing showed that participants placed in shelters had significantly higher levels of serum glutamic pyruvate transaminase (GPT values; serum GPT >40 IU/L) as well as significantly lower total cholesterol values than those in the community (serum GPT: 28.9% vs 13.2%, \(p=0.001\); total cholesterol: 188.65±38.33 mg/dL vs 199.15 ±42.40 mg/dL, \(p=0.034\); table 2). Comparison of first-round and second-round data showed that participants in shelters had significantly higher levels of serum GPT, but significantly lower levels of serum albumin and total cholesterol than participants in the community (all \(p<0.05\)).

**First-round serology results**

Table 3 summarises the first round of serology tests. First round testing after the typhoon showed that participants in shelters had a significantly higher rate of mumps, but a significantly lower rate of HSV-2, compared to the community group (mumps: 81.6% vs 69.5%, \(p=0.022\); HSV-2: 27.2% vs 40.2%, \(p=0.023\)). All 288 participants in the first round examination were subjected to a second round of testing after 6 months. There were no

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**Table 1  Serological test kits used in the present study**

| Disease       | Products                                | Manufacturer   | City          | Country   |
|---------------|-----------------------------------------|----------------|---------------|-----------|
| Measles       | Measles virus IgG–ELISA                | NovaLisa       | Dietzenbach   | Germany   |
| Mumps         | Mumps virus IgG–ELISA                  | NovaLisa       | Dietzenbach   | Germany   |
| Rubella       | Rubella virus IgG–ELISA                | Behring Diagnostics | Marburg    | Germany   |
| Amoebiasis (IHA) | Cellognost–Amoebias                   | CTK Biotech, Inc | San Diego, CA | USA       |
| Leptospirosis | OnSite Leptospira IgG/IgM Combo Rapid Test | Euroimmun      | Lübeck        | Germany   |
| HSV-1         | Anti-HSV-1 ELISA (IgG)                 | Euroimmun      | Lübeck        | Germany   |
| HSV-2         | Anti-HSV-2 ELISA (IgG)                 | Human          | Wiesbaden     | Germany   |
| VDRL of syphilis | Syphilis RPR Test                     |                |               |           |

HSV, herpes simplex virus; RPR, rapid plasma reagin; VDRL, Venereal Disease Research Laboratory.
significant differences in the incidence of vaccine-preventable diseases or in water-borne diseases between participants in shelters and in community (all p>0.05). There was one case of measles seroconversion, 3 cases of mumps seroconversion and 10 cases of rubella seroconversion, in all. Of the 288 participants, 128 participants (44.4%) showed amoebiasis seroconversion by the IHA test. The rate of amoebiasis seroconversion was higher among participants who had stayed in shelters 2 weeks or more compared to those who remained in the community (56.1% vs 36.8%; p=0.001; table 3). Associations between amoebiasis seroconversion and shelter stay, participants' demographics, medical history and laboratory data, are summarised in table 4. Amebiasis seroconversion was significantly associated with shelter stay and medical history, including hypertension and hyperlipidaemia (all p values<0.05; table 4). The length of stay in the shelter was positively and significantly associated with the rate of seroconversion (figure 1).

Table 5 represents the results of binary logistic regression analysis. Variables that were significantly associated with amoebiasis seroconversion (table 4) were placed into univariate and multivariate logistic regression analysis. Univariate analysis showed that shelter stay and hypertension were significantly associated with amoebiasis seroconversion. Participants with hyperlipidaemia may be at lower risk for seroconversion (all p<0.05). After adjusting for medical history, hypertension and hyperlipidaemia, multivariate logistic regression analysis showed that participants who stayed in shelters had a higher

![Table 2](https://www.bmjopen.org/content/5/5/e007008)

Table 2: Participants' demographics, medical history and laboratory data by group (N=288)

| Variables                              | Total (n=288) | Shelter group (n=114) | Community group (n=174) | p Value |
|----------------------------------------|--------------|-----------------------|-------------------------|---------|
| Demographics                           |              |                       |                         |         |
| Gender, male (%)                       | 96 (33.3)    | 37 (32.5)             | 59 (33.9)               | 0.798   |
| Age ≥65 years, n (%)                   | 94 (32.6)    | 30 (26.3)             | 64 (36.8)               | 0.064   |
| BMI                                    | 28.20±4.92   | 28.60±4.67            | 27.94±5.08              | 0.268   |
| Ratio of waist and hip                 | 0.90±0.07    | 0.89±0.07             | 0.90±0.07               | 0.266   |
| Education                              |              |                       |                         |         |
| Junior high school or lower            | 159 (55.2%)  | 64 (56.1%)            | 95 (54.6%)              | 0.913   |
| High school and college                | 54 (18.8%)   | 20 (17.5%)            | 34 (19.5%)              |         |
| Without formal education and unknown   | 75 (26.0%)   | 30 (26.3%)            | 45 (25.9%)              |         |
| Single‡, n (%)                         | 121 (42.0)   | 46 (40.4)             | 75 (43.1)               | 0.643   |
| Family with major trauma               | 10 (3.5)     | 6 (5.3)               | 4 (2.3)                 | 0.201   |
| Medical history                        |              |                       |                         |         |
| Hypertension                           | 127 (46.2)   | 53 (47.3)             | 74 (45.4)               | 0.753   |
| Diabetes mellitus                      | 45 (16.4)    | 20 (17.9)             | 25 (15.3)               | 0.579   |
| Hyperlipidaemia                        | 177 (61.5)   | 20 (17.9)             | 111 (63.8)              | 0.315   |
| Hyperuricaemia                         | 91 (31.6)    | 33 (28.9)             | 58 (33.3)               | 0.434   |
| Ischaemic heart disease                | 10 (3.6)     | 5 (4.5)               | 5 (3.1)                 | 0.543   |
| Gouty arthritis                        | 35 (12.7)    | 19 (17.0)             | 16 (9.8)                | 0.081   |
| Cerebrovascular disease                | 6 (2.2)      | 1 (0.9)               | 5 (3.1)                 | 0.406   |
| Laboratory data                        |              |                       |                         |         |
| After typhoon (first round)            |              |                       |                         |         |
| SERUM GPT >40 IU/L                     | 49 (17.0)    | 23 (20.2)             | 26 (14.9)               | 0.248   |
| Creatinine >1.5 mg/dL                  | 22 (7.6)     | 8 (7.0)               | 14 (8.0)                | 0.748   |
| Serum albumin, mg/dL                   | 4.40 (4.30, 4.60) | 4.40 (4.30, 4.60) | 4.40 (4.20, 4.60) | 0.227   |
| Total cholesterol, mg/dL               | 203.98±46.34 | 197.85±37.00          | 208.00±51.26            | 0.084   |
| Triglyceride, mg/dL                    | 150 (92, 233.25) | 142 (89.75, 219.25) | 155.50 (92.00, 242.75) | 0.111   |
| Follow-up (second round)               |              |                       |                         |         |
| SERUM GPT >40 IU/L                     | 56 (19.4)    | 33 (28.9)‡            | 23 (13.2)†              | 0.001†  |
| Creatinine >1.5 mg/dL                  | 20 (6.9)     | 6 (5.3)               | 14 (8.0)§               | 0.364   |
| Serum albumin, mg/dL                   | 4.30 (4.20, 4.50)† | 4.30 (4.20, 4.50)‡ | 4.30 (4.18, 4.50)§ | 0.447   |
| Total cholesterol, mg/dL               | 194.99±41.09† | 188.65±38.33‡         | 199.15±42.40§           | 0.034§  |
| Triglyceride, mg/dL                    | 150 (102, 257.50)† | 142.50 (98.25, 208.75) | 156.50 (102, 275.75)§ | 0.240   |

Data are summarised as n (%) for categorical variables, means±SD for continuous variables and median (IQR: Q1, Q3) for continuous variables without normal distribution.

Differences between shelter and community groups in categorical data were compared using Pearson χ² test or Fisher's exact test if there were any cases numbering less than five; for continuous data, differences between shelter group and community group were compared using two-sample t test, and Mann-Whitney U test, in data without normal distribution.

* p<0.05, indicates significant differences between shelter and community groups.
†‡§ p<0.05, indicates significant differences when comparing data after typhoon in †total, ‡shelter group and §community group, separately.
¶Including divorced or unmarried participants, or widows or widowers.

BMI, body mass index.
risk for seroconversion than those in community. (OR=2.055, 95% CI 1.251 to 3.374; p=0.004; table 5).

Clinical manifestation of amoebiasis seroconversion
We retrospectively interviewed 60 seroconverted participants in order to evaluate disease severity. Only 3 (5%) of these patients reported acute diarrhoea (less than 14 days) without dysentery.

DISCUSSION
This longitudinal study, which was conducted after Typhoon Morakot and the accompanying flooding, showed that E. histolytica seroconversion rates were higher in persons displaced to shelters compared to those remaining in the community, although disease manifestation was mild or asymptomatic. There were a few instances of vaccine-preventable diseases such as measles, mumps and rubella, among displaced participants in the shelter group as well as among those in the community group. These diseases likely occurred in unvaccinated participants or in those with weakened immune systems. The risk of amoebiasis seroconversion was higher in participants displaced to shelters who had hypertension and lower in participants with hyperlipidaemia. However, after adjusting for medical history, hypertension and hyperlipidaemia, multivariate logistic regression analysis showed that participants who stayed in shelters had a significantly higher risk for amoebiasis seroconversion than those in the community.

Recent data suggested a relationship between massive precipitation and climate-related diseases in Taiwan (ie, enterovirus infection, Japanese encephalitis, scrub typhus, dengue fever and, to some extent, melioidosis and leptospirosis).24 However, the relationship between parasitic diseases and extreme rainfall remains unclear. Although Taiwan is considered a low-endemic area for amoebiasis,20 and notification of cases to the Center for Disease Control (CDC) is mandatory, our present study showed that amoebiasis is not uncommon in Taiwan. The annual number of cases reported to the CDC in Taiwan ranged from only 146 to 264 between 2007 and 2011,25 and studies suggested that there are many asymptomatic infections.26

Unlike outbreak reports, this longitudinal, paired serological epidemiological investigation relied on evidence...
Table 4  Associations between accommodations, participants’ demographics, medical history, laboratory data and seroconversion (N=288)

| Variables                                      | Seroconversion (n=128) | Without seroconversion (n=160) | p Value |
|-----------------------------------------------|------------------------|-------------------------------|---------|
| Accommodations                                |                        |                               |         |
| Community                                     | 64 (36.8)              | 110 (63.2)                    | 0.001*  |
| Shelter                                       | 64 (56.1)              | 50 (43.9)                     |         |
| Demographics                                  |                        |                               |         |
| Gender                                        |                        |                               |         |
| Males                                         | 43 (44.8)              | 53 (55.2)                     | 0.933   |
| Females                                       | 85 (44.3)              | 107 (55.7)                    |         |
| Age ≥65 years                                  |                        |                               |         |
| No                                            | 86 (44.3)              | 108 (55.7)                    | 0.955   |
| Yes                                           | 42 (44.7)              | 52 (55.3)                     |         |
| Age, yrs                                       | 59.02±12.33            | 57.94±13.05                   | 0.475   |
| BMI, kg/m²                                     | 28.29±4.81             | 28.13±5.02                    | 0.788   |
| Ratio of waist and hip                         | 0.90±0.07              | 0.89±0.07                     | 0.491   |
| Education                                     |                        |                               |         |
| Junior high school or lower                   | 69 (43.4)              | 90 (56.6)                     |         |
| High school and college                       | 25 (46.3)              | 29 (53.7)                     |         |
| Without formal education and unknown          | 34 (45.3)              | 41 (54.7)                     |         |
| Single†, n (%)                                |                        |                               |         |
| No                                            | 78 (46.7)              | 89 (53.3)                     | 0.364   |
| Yes                                           | 50 (41.3)              | 71 (58.7)                     |         |
| Family with major trauma                      |                        |                               |         |
| No                                            | 122 (43.9)             | 156 (56.1)                    | 0.348   |
| Yes                                           | 6 (60.0)               | 4 (40.0)                      |         |
| Medical history                               |                        |                               |         |
| Hypertension                                  | 66 (52.0)              | 61 (48.0)                     | 0.044*  |
| Diabetes mellitus                             | 24 (53.3)              | 21 (46.7)                     | 0.246   |
| Hyperlipidaemia                               | 70 (39.5)              | 107 (60.5)                    | 0.035*  |
| Hyperuricaemia                                | 44 (48.4)              | 47 (51.6)                     | 0.436   |
| Ischaemic heart disease                       | 3 (30.0)               | 7 (70.0)                      | 0.246   |
| Gouty arthritis                               | 16 (45.7)              | 19 (54.3)                     | 0.974   |
| Cerebrovascular disease                       | 3 (50.0)               | 3 (50.0)                      | 1.000   |
| Laboratory data                               |                        |                               |         |
| Pretest                                       |                        |                               |         |
| Serum GPT >40 IU/L                            | 23 (46.9)              | 26 (53.1)                     | 0.700   |
| Creatinine >1.5 mg/dL                         | 7 (31.8)               | 15 (68.2)                     | 0.215   |
| Serum albumin, mg/dL                          | 4.40 (4.30, 4.60)      | 4.40 (4.30, 4.60)             | 0.302   |
| Total cholesterol, mg/dL                      | 200.16±36.32           | 207.04±52.93                  | 0.194   |
| Triglyceride, mg/dL                           | 148.50 (84.00, 202.50) | 160.50 (95.30, 261.00)        | 0.167   |
| Follow-up                                     |                        |                               |         |
| Serum GPT >40 IU/L                            | 29 (51.8)              | 27 (48.2)                     | 0.218   |
| Creatinine >1.5 mg/dL                         | 7 (35.0)               | 13 (65.0)                     | 0.378   |
| Serum albumin, mg/dL                          | 4.30 (4.10, 4.40)      | 4.30 (4.20, 4.50)             | 0.087   |
| Total cholesterol, mg/dL                      | 191.74±39.90           | 197.59±41.96                  | 0.230   |
| Triglyceride, mg/dL                           | 193.00 (162.00, 214.75) | 156.50 (106.00, 261.00)       | 0.248   |

Data are summarised as n (%) for categorical variables, mean±SD for continuous variables, and median (IQR: Q1, Q3) for continuous variables without normal distribution.

Differences between participants with and without seroconversion in categorical data were compared using Pearson χ² test or Fisher’s exact test if there were any cells numbering less than five; for continuous data, differences between participants with and without seroconversion were compared using two-sample t test and Mann-Whitney U test in data without normal distribution.

*p<0.05, indicates significant differences between participants with and without seroconversion.

†Including divorced or unmarried participants, or widows or widowers.

BMI, body mass index.

of seroconversion to evaluate the incidence of infectious diseases among different populations of disaster-displaced persons. We found that prolonged shelter stay was associated with higher *E. histolytica* antibody seroconversion rates after Typhoon Morakot, even though those who showed seroconversion presented only mild symptoms or were asymptomatic. Our post-disaster serological data were consistent with a study of
clinical laboratory methods of *E. histolytica* detection, which noted the asymptomatic nature of amoebiasis.\(^23\)

Our data strongly suggested that contaminated water supply in crowded shelters could play a major role in amoebiasis outbreaks seen after natural disasters. We suggest that a clean drinking water supply is crucial during post-disaster care and management, especially for displaced persons admitted to shelters. The presence of *E. histolytica* seroconversion among disaster-displaced people living in the community may be due to disaster-damaged public drinking water systems whereby some people living with their families in urban areas may also become infected.

Although an outbreak of leptospirosis was previously reported in Pingtung County (a non-mountainous area) after Typhoon Morakot, and melioidosis was also confirmed in areas of heavy precipitation after the typhoon,\(^12\) there were no reports of leptospirosis outbreaks made to the CDC from the mountainous areas during our study period. This discrepancy suggested that rainfall levels and cumulative flooding may have a greater impact on leptospirosis and melioidosis outbreaks compared to typhoon strength.

Our data showed that a greater percentage of displaced persons admitted to shelters had family members who suffered a major trauma resulting in serious injury or death. On the other hand, the higher percentage of older displaced persons in the community group may be due to the fact that family members who originally resided in urban areas before Typhoon Morakot (not included in our cohort) were more likely to provide shelter to their elderly relatives. Different environmental circumstances as well as the different vaccine coverage rates among various age groups may explain the different seropositivity rates for mumps between younger adults in the two groups. However, among people aged ≥50 years in both groups, the baseline seropositivity rates for mumps were not different (74% in the shelter group vs 67.9% in the community group; p=0.150; data not shown).

Our present data showed some cases of seroconversion for vaccine-preventable diseases such as measles, mumps and, especially, rubella, among displaced persons during the study period. Although there were no significant differences in the seroconversion rates for these three diseases between the shelter and community groups, evidence of seroconversion among displaced persons suggested that there was a risk of outbreaks under these conditions. It is important to note that vaccine coverage rates are high in Taiwan, and there were no outbreaks during our study period, unlike the concurrent measles and rubella outbreaks in camps hosting Liberian displaced persons in Côte d’Ivoire in 2004.\(^27\) However, our data suggested that there was a risk of vaccine-preventable diseases after the disaster, regardless of the type of accommodations. Outbreaks of infectious diseases such as rubella in crowded shelters after natural disasters can be effectively prevented by extending vaccination programmes to populations who have a higher risk of post-disaster displacement and who are without antibody protection.

There is currently little evidence to provide guidance on whether staying in shelters is a more advantageous post-disaster accommodation policy in terms of preventing disease transmission among large numbers of displaced people, compared to remaining in the community. To our knowledge, this is the first study investigating differences in seroconversion rates between disaster-displaced persons staying in different types of accommodations. Our present data are consistent with previous results showing that effective post-disaster management and mitigation strategies are necessary to address the needs of flood-displaced persons, and to reduce the risk of communicable diseases,\(^4\)\(^,\)\(^8\) especially

![Figure 1](image-url)  
**Figure 1** Amoebiasis indirect haemagglutination (IHA) seroconversion rate in groups with different lengths of stay in the shelters.

### Table 5

| Variables† | Univariate OR (95% CI) | p Value | Multivariate OR (95% CI) | p Value |
|------------|------------------------|---------|--------------------------|---------|
| Shelter admission | | | | |
| Shelter versus community | 2.200 (1.359 to 3.561) | 0.001* | 2.055 (1.251 to 3.374) | 0.004* |
| Medical history | | | | |
| Hypertension | 1.632 (1.011 to 2.635) | 0.045* | 1.576 (0.964 to 2.575) | 0.070 |
| Hyperlipidaemia | 0.598 (0.370 to 0.965) | 0.035* | 0.642 (0.389 to 1.060) | 0.083 |

Results are presented as OR and corresponding 95% CIs of OR (95% CI) and p values.

*Indicates significance (p<0.05).

†Variables with significant association with seroconversion were selected into models.
in vulnerable groups such as younger children and older adults with weaker immune systems. This study has several limitations. First, only 46.9% of the participants with *E. histolytica* IHA seroconversion were retrospectively interviewed. Bias in participants’ ability to recall clinical symptoms or signs was also a limitation. Second, we did not directly examine the drinking water in the shelters; thus, a direct causal relationship between drinking water and *E. histolytica* seroconversion could not be established even though the risk of seroconversion was significantly higher in the shelter group compared to the community group. Another limitation was that although we recognise the importance of covering all infectious diseases in post-crisis situations, we could only investigate diseases based on local interests in this study. Finally, the sample size for this longitudinal study was relatively small. It is important to validate our findings with a larger sample size.

In conclusion, after Typhoon Morakot, shelter stay was associated with higher risk for serconversion of *E. histolytica*. Although displaced persons staying in shelters had higher seroconversion rates than those remaining in the community, manifestations were mild. Our results provide useful information for the pre-emptive preparation and response to relatively predictable typhoons and flooding, and suggested that a clean water supply is essential after a disaster, especially in relatively crowded shelters. Our finding of measles, mumps and rubella seroconversion among displaced participants, strongly suggested that prevention of large outbreaks of infectious diseases following natural disasters could be prevented by extending vaccination programmes to populations at higher risk for post-disaster displacement and to those with weakened immune systems.

**Author affiliations**

1 Division of Infectious Diseases, Department of Internal Medicine, Kaohsiung Medical University Hospital, Kaohsiung, Taiwan
2 Graduate Institute of Medicine, Tropical Medicine Research Center, College of Medicine, Kaohsiung Medical University, Kaohsiung, Taiwan
3 Hepatobiliary Division, Department of Internal Medicine, Kaohsiung Medical University Hospital, Kaohsiung Medical University, Kaohsiung, Taiwan

**Contributors**

C-YL, C-YD and Y-HC conceived and designed the experiments. C-YL, T-CC and P-LL analysed the data. C-YD, M-LY and J-HY contributed reagents/materials/analysis tools. C-YL, T-CC and Y-HC wrote the paper.

**Funding**

This study was supported by a grant to Y-TW, C-YD and Y-HC from the National Science Council, Taiwan (NSC 98-2321-B-037-063).

**Competing interests**

None declared.

**Patient consent**

Obtained.

**Ethics approval**

The institutional review board of Kaohsiung Medical University Hospital reviewed and approved the study protocol, and all participants provided written informed consent. People younger than age 20 years were excluded from this study.

**Provenance and peer review**

Not commissioned; externally peer reviewed.

**Data sharing statement**

No additional data are available.

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