**Characteristics and Factors Associated with Death among Patients Hospitalized for Severe Fever with Thrombocytopenia Syndrome, South Korea, 2013**

Jaeseung Shin, Donghyok Kwon, Seung-Ki Youn, Ji-Hyuk Park

In South Korea, nationwide surveillance for severe fever with thrombocytopenia syndrome (SFTS) began during 2013. Among 301 surveillance cases, 35 hospitalized case-patients in 25 areas were confirmed by using virologic testing, and 16 (46%) case-patients subsequently died. The SFTS cases occurred during May–November and peaked during June (9 cases, 26%). The incidence of SFTS was higher in the southern regions of South Korea. Age and neurologic symptoms, including decreased level of consciousness and slurred speech, were heavily associated with death; neurologic symptoms during the first week after disease onset were also associated with death. Although melena was common among patients who died, no other hemorrhagic manifestations were substantively more common among those who died. No effective treatments, including ribavirin, were identified. Expansion of SFTS surveillance to include the outpatient sector and development of an antibody test would enhance completeness of SFTS detection in South Korea.

Severe fever with thrombocytopenia syndrome (SFTS) is an emerging infectious disease that is caused by a novel phlebovirus in the Bunyaviridae family. This virus has been named severe fever with thrombocytopenia syndrome virus (SFTSV). The disease was first reported during 2009 in China, where it is most prevalent in the Hunan, Hubei, and Shandong provinces, which are located at a similar latitude to that of South Korea and Japan (1). SFTS is mainly transmitted to humans by SFTSV-infected ticks, most frequently Haemaphysalis longicornis (2). However, person-to-person transmission by direct contact with infected blood or body fluid has also been reported (3,4).

The clinical symptoms of SFTS include fever; gastrointestinal (GI) symptoms (e.g., diarrhea and vomiting); leukocytopenia and thrombocytopenia; bleeding tendency; and neurologic symptoms. The incubation period for SFTS is 1–2 weeks, and illness progresses through 3 stages: fever, multiorgan dysfunction, and convalescence. No effective treatment for SFTS has been established (5).

In South Korea, the laboratory diagnostic system for SFTS was established during March 2013, and the presence of SFTSV among H. longicornis ticks was confirmed by using samples from ticks that were collected during 2011–2012 (6). Based on the nationwide surveillance reports for SFTS and its designation as a notifiable infectious disease, the first SFTS case was retrospectively confirmed in May 2013, after the patient’s death in 2012 (7). However, because of the novel nature of this system, the clinical and demographic characteristics of patients in South Korea infected with SFTS are not well understood. Therefore, this study evaluated the characteristics and factors that were associated with SFTS-related fatalities in South Korea, as reported during 2013.

**Methods**

**Surveillance System and Case Definition**

A passive hospital-based surveillance system for SFTS was initiated nationwide in South Korea during March 2013. Physicians were advised to request SFTSV testing for patients with known clinical manifestations of SFTS, including fever (body temperature $\geq 38.0^\circ$C), thrombocytopenia, leukocytopenia, and GI symptoms. Each patient’s enrollment for SFTSV testing was dependent on the physician’s clinical suspicion of SFTS. Serum samples from 301 patients admitted on the basis of physician referrals at 125 hospitals throughout Korea were collected and tested. Case-patients were defined as patients who had clinical symptoms and were confirmed to have SFTSV by virologic testing at the Korea Centers for Disease Control and Prevention.
Severe Fever with Thrombocytopenia Syndrome (KCDC). This study did not require an institutional ethics review because it was conducted under the Infectious Disease Control and Prevention Act in South Korea.

**Laboratory Testing**

All acute-phase serum samples were tested to detect the SFTSV medium segment gene by one-step reverse transcription PCR, by using DiaStar 2× OneStep reverse transcription PCR Pre-Mix Kit (SolGent, Daejeon, South Korea), as described (8). The PCR primers were MF3 (5′-GATGAGATGGTCCATGCTGATTCT-3′) and MR2 (5′-CTCATGGGGTGGAATGTCCTCAC-3′). The PCR conditions were an initial step of 30 min at 50°C for reverse transcription; 15 min at 95°C for denaturation; 35 cycles of 20 s at 95°C, 40 s at 58°C, and 30 s at 72°C; and a final extension step of 5 min at 72°C.

**Epidemiologic Investigation**

We also performed an epidemiologic investigation of patients in whom SFTS was suspected immediately after the SFTSV testing was requested by their physicians. KCDC Epidemic Intelligence Service officers interviewed the patients and their physicians using a standardized questionnaire that evaluated demographic characteristics, exposure history, clinical symptoms, and laboratory results. Patients were also questioned regarding their exposure history within the month before their onset of symptoms. Multiple activities were documented, including agricultural and forestry work, mountain climbing, and visits to a family grave. The date of the tick bite was self-reported when the patient was aware of a bite or if the patient participated in only 1 exposure-related activity over a short period of time. In addition, after the death or discharge of confirmed case-patients during September 2013–January 2014, the clinical course and prognosis were investigated by reviewing medical records. The date for clinical suspicion of SFTS was recorded as the date reported to the KCDC.

**Statistical Analysis**

We obtained median regional population data during 2013 from Statistics Korea (http://www.kostat.go.kr) to calculate the regional incidences during 2013. These incidences were then overlaid on a map of South Korea by using bio-geographic information system software (DIVA-GIS 7.5; http://www.diva-gis.org). We used the Fisher exact test or the Mann-Whitney U test to compare the prognoses of patients with SFTS. All statistical analyses were performed by using SAS software version 9.2 (SAS Institute, Cary, NC, USA), and statistical significance was set at p<0.05.

**Results**

During 2013, a total of 36 hospitalized patients were confirmed to have SFTS. One case was excluded because the patient’s disease onset occurred during 2012. Thus, 35 cases from 20 hospitals were included in our analysis.

**Demographic Characteristics**

Symptom onset occurred during May–November 2013; a peak of symptoms among 9 (26%) patients occurred in June. Similar patterns were observed among the SFTS case-patients in our study (Figure 1, panel A) and the number of *H. longicornis* ticks that were collected each month by Park et al. during 2011–2012 (6) (Figure 1, panel B). Peaks of symptoms among patients occurred in June 2013, and peaks of the number of ticks collected occurred in May 2011 and May 2012.

The overall incidence of SFTS during 2013 was 0.7 cases/1 million persons. Geographic differences were documented in the incidences in 25 cities; higher incidences...
were observed in the southern regions, including Jeju Province (Figure 2).

The case-patient group comprised 18 women (51%) and 17 men (49%). The median age was 69 (range 28–84) years, and age was significantly higher among those who died (range 62–82 years; p = 0.026). Twenty-six (74%) case-patients resided in rural areas, and 25 (71%) were farmers. A trace of a tick bite was found on 11 (31%) case-patients, and 4 (13%) had recognized the tick bite before their hospital admission. The most common exposure-related activity was agricultural work (n = 20, 61%), followed by forestry work (n = 7, 21%), mountain climbing (n = 6, 18%), and visiting a family member’s grave (n = 3, 9%). The agricultural work was performed in dry fields by 16 (80%) of 20 case-patients and forestry work was done in orchards by 5 (71%) of 7 case-patients (Table 1).

The incubation period was available for 8 patients; the median was 9 (range 5–16) days. Median time from symptom onset to hospital admission was 4 (range 1–9) days and from symptom onset to clinical suspicion was 7 (range 3–22) days. We found no substantive differences in time from symptom onset to admission and to clinical suspicion between the group of persons who died and those who recovered. Twenty-four (69%) of the 35 patients with SFTS were admitted to intensive care units, and 16 (46%) died. The median survival time among the 16 was 10.5 (range 4–32) days; 11 (69%) patients died within 2 weeks after symptom onset. Among patients who survived, median time from symptom onset to discharge was 18 (range 10–49) days (Table 1).

The cohort included 1 family cluster, which consisted of an uncle and nephew. They did not live in the same village and did not have contact with each other after symptom onset. However, within 2 weeks before symptom onset, both had visited the same family member’s grave on a mountain and had mowed the grass.

**Clinical Characteristics and Prognoses**

The clinical characteristics of the 35 SFTS case-patients during the course of the illness are shown in Table 2. All patients experienced fever or chills, and other common symptoms included GI symptoms (n = 33, 94%), neurologic symptoms (n = 27, 77%), fatigue (n = 26, 74%), myalgia (n = 19, 54%), and hemorrhagic manifestations (n = 12, 34%). Among the 32 patients with medical records for the first 7 days after disease onset, all patients had a fever within 7 days of symptom onset. A high fever (≥39°C) was documented for 7 (22%) patients, although no significant difference was observed...
related to occurrence of fever among those who died and those who recovered (p = 0.195). Lymphadenopathy was identified by physical examination on admission on the neck, face, left thigh, left axilla, or left inguinal area of 5 (14%) patients.

The most common GI symptoms among all patients were diarrhea (n = 22, 63%), anorexia (n = 21, 60%), nausea (n = 16, 46%), and vomiting (n = 15, 43%). Both diarrhea and vomiting were reported for 26 (74%) patients; most (25/26, 96%) experienced diarrhea or vomiting within 1 week after symptom onset, and 18 (51%) patients had diarrhea or vomiting at the time of admission. However, no substantive differences in the occurrence of GI symptoms were observed for those who died compared with those who recovered.

Decreased level of consciousness (n = 26, 74%) was the most frequent neurologic symptom, followed by slurred speech (n = 9, 26%) and convulsion (n = 8, 23%). Neurologic symptoms occurred at a median of 6 days after symptom onset (range 2–10 days). On admission, 3 patients (9%) had neurologic symptoms. The group of case-patients who died had a significantly higher number of members who had neurologic symptoms (p = 0.047) and exhibited significantly more frequent neurologic symptoms that occurred within 7 days (p = 0.002). Decreased level of consciousness (p = 0.050) and slurred speech (p = 0.022) were significantly more common among those who died, although the frequency of convulsion was similar for both groups (p = 0.105).

The only hemorrhagic manifestation that occurred with a significant difference among those who died and those who recovered was melena (p = 0.035). None of the treatments of SFTS appeared to be effective, including the use of ribavirin, intravenous immunoglobulin, plasmapheresis, or continuous renal replacement therapy. Fungal pneumonia (n = 2, 6%) and cerebral hemorrhage (n = 2, 6%) were observed as complications among those who died.

**Laboratory Features and Prognoses**

The laboratory data from the 32 patients with available medical records for the first 7 days after symptom onset are shown in Table 3. All patients had thrombocytopenia (platelets <150 × 10^9/L) and leukocytopenia (leukocytes <4 × 10^9 cells/L) on admission. The median minimum platelet count was marginally lower among those who died (34 × 10^9/L), compared with that among those who recovered (47.5 × 10^9/L; p = 0.054). The maximum serum
TABLE 2. Clinical characteristics of hospitalized case-patients with confirmed severe fever with thrombocytopenia syndrome, by outcome, South Korea, 2013

| Characteristics                                      | No. (%) case-patients | p value†  |
|------------------------------------------------------|-----------------------|-----------|
|                                                      | Total, n = 35         | Died, n = 16 | Recovered, n = 19 |
| Fever/chills                                         | 35 (100)              | 16 (100)   | 19 (100)         |
| Fever, n = 32                                        |                       |           |                  |
| ≤39.0°C                                              | 25 (78)               | 9 (64)    | 16 (89)         | 0.195 |
| >39.0°C                                              | 7 (22)                | 5 (36)    | 2 (11)          | NA    |
| Fatigue                                              | 26 (74)               | 11 (69)   | 15 (79)         | 0.700 |
| Myalgia                                              | 19 (54)               | 9 (56)    | 10 (53)         | 1.000 |
| Headache                                             | 12 (34)               | 4 (25)    | 8 (42)          | 0.476 |
| Cough                                                | 7 (20)                | 3 (19)    | 4 (21)          | 1.000 |
| Sputum                                               | 9 (26)                | 3 (19)    | 6 (32)          | 0.460 |
| Lymphadenopathy on physical examination              | 5 (14)                | 1 (6)     | 4 (21)          | 0.347 |
| Lymph node enlargement found by CT, n = 32           | 20 (63)               | 7 (54)    | 13 (68)         | 0.473 |
| Gastrointestinal symptoms                            |                       |           |                  |
| Overall                                              | 33 (94)               | 15 (94)   | 18 (95)         | 1.000 |
| Anorexia                                             | 21 (60)               | 8 (50)    | 13 (68)         | 0.317 |
| Nausea                                               | 16 (46)               | 5 (31)    | 11 (58)         | 0.176 |
| Abdominal pain                                       | 9 (26)                | 6 (38)    | 3 (16)          | 0.245 |
| Diarrhea                                             | 22 (63)               | 12 (75)   | 10 (53)         | 0.293 |
| Vomiting                                             | 15 (43)               | 6 (38)    | 9 (47)          | 0.734 |
| Diarrhea/vomiting                                    | 26 (74)               | 13 (81)   | 13 (68)         | 0.460 |
| Neurologic symptoms                                  |                       |           |                  |
| Overall                                              | 27 (77)               | 15 (94)   | 12 (63)         | 0.047 |
| Within 7 d after disease onset                       | 20 (57)               | 14 (88)   | 6 (32)          | 0.002 |
| Slurred speech                                       | 9 (26)                | 7 (44)    | 2 (11)          | 0.050 |
| Decreased level of consciousness                     | 26 (74)               | 15 (94)   | 11 (58)         | 0.022 |
| Convulsions                                          | 8 (23)                | 6 (38)    | 2 (11)          | 0.105 |
| Hemorrhagic manifestations                            |                       |           |                  |
| Overall                                              | 12 (34)               | 8 (50)    | 4 (21)          | 0.090 |
| Gross hematuria                                       | 2 (6)                 | 1 (6)     | 1 (5)           | 1.000 |
| Petechiae                                            | 3 (8)                 | 1 (6)     | 2 (11)          | 1.000 |
| Gingival bleeding                                    | 5 (14)                | 3 (19)    | 2 (11)          | 0.642 |
| Hematemesis                                          | 2 (6)                 | 1 (6)     | 1 (5)           | 1.000 |
| Hematochezia                                         | 1 (3)                 | 1 (6)     | 0 (0)           | 0.457 |
| Melena                                               | 4 (11)                | 4 (25)    | 0 (0)           | 0.035 |
| Treatment                                            |                       |           |                  |
| Ribavirin                                            | 9 (26)                | 6 (38)    | 3 (16)          | 0.245 |
| IVIG                                                 | 7 (20)                | 5 (31)    | 2 (11)          | 0.208 |
| Plasmapheresis                                       | 7 (20)                | 4 (25)    | 3 (16)          | 0.877 |
| CRRT                                                 | 10 (29)               | 9 (56)    | 1 (5)           | 0.002 |

*NA, not applicable; CT, computed tomography; IVIG, intravenous immunoglobulin; CRRT, continuous renal replacement therapy.
†Fisher exact test was used to compare groups.

The higher incidence of SFTS in the southern part of South Korea was particularly notable for Jeju Province, which is the largest island and the most southern part of South Korea. The prevalence of *H. longicornis* ticks in Jeju Province is among the highest in South Korea (9,10); high temperatures are conducive to the survival and breeding of this species (11), and Jeju Province has the highest average temperature in South Korea, related to its low latitude (12). Furthermore, a study of ticks that were collected during 2011–2012 in South Korea reported that the minimum infection rates for SFTSV in *H. longicornis* ticks were higher in the southern part of the country (6).

Although the SFTS case-fatality rate in South Korea (46%) was higher than those that have been reported by using SFTS data from China (6%–30%) (1,13), it was similar to the rate (55%) that was reported in a retrospective tracing study in Japan (14). However, after the introduction of the China surveillance system in 2009, a decreasing trend in...
the case-fatality rate in that country has been observed \((13)\), which may be supported by the increased capability to detect mild SFTS cases. Age was associated with a prognosis of death in our study, and other studies of hospitalized patients in China with SFTS have reported similar findings \((15-17)\). However, in our study, the median ages for those who died versus those who survived (73.5 years and 61 years, respectively) tended to be higher than those among the patients in China \((62.1-74\) years and \(52.9-60\) years, respectively) \((15-17)\). This older age might partially explain why we observed a higher case-fatality rate compared with previous reports. However, we did not find any person-to-person clusters of infection, and found only 1 family cluster, although these patterns have been reported in China \((3,18)\).

The median time from symptom onset to admission was 4 days and from onset to clinical suspicion was 7 days. This finding indicates that there is a delay in diagnosis and appropriate care, which is likely related to lack of clinical experience with SFTS among physicians in South Korea. Nevertheless, despite the lack of a statistically significant relationship between this time period and death, a delayed diagnosis of SFTS could affect the prognosis of the patients.

The fever stage of SFTS occurs during the first week after disease onset and is characterized by the sudden onset of fever and GI symptoms \((19)\); our findings were similar. However, we found that neurologic symptoms were associated with a greater likelihood of death, and similar findings have been reported in previous studies of hospitalized patients in China \((19,20)\). Moreover, we observed a rapid progression and increased frequency of neurologic symptoms during the fever stage among the patients who died, suggesting that early neurologic symptoms portend a fatal outcome.

Hemorrhagic manifestations have been linked to fatal outcomes among SFTS patients in China \((19,20)\). However, we did not observe a similar association, with the exception of melena. Differences between the health care systems in South Korea and China, especially regarding the use of appropriate platelet transfusions, may partially explain the differences regarding hemorrhagic manifestations.

The lowest platelet counts during the first week were not substantively different between those who recovered and those who died. However, one study has reported that low platelet counts on admission or during the entire course were associated with SFTS severity \((20)\). Therefore, it is possible that our relatively small number of cases may obscure the association of the lowest platelet counts with death among patients in South Korea.

According to 2 studies in China, elevated LDH levels on admission or during the entire course of illness were significantly associated with a fatal outcome \((15,20)\). However, we did not observe a significant association between the highest LDH levels during the first week or at the time of admission and death \(\text{data not shown}\). We did observe significantly higher ALP levels during the first week among those who died, although a study of ALP levels on admission reported similar levels between those who died and those who recovered on the basis of multivariate analysis findings \((15)\). Additional studies are needed to evaluate the

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**Table 3. Laboratory features of hospitalized case-patients with confirmed severe fever with thrombocytopenia syndrome during the first week after onset, by outcome, South Korea, 2013**

| Laboratory tests                  | Median (range)                  | p value† |
|-----------------------------------|---------------------------------|----------|
| Total, n = 32                     | Total, n = 14                   | Recovered, n = 18 |
| Platelet count, \(\times 10^9/\mu L\)†‡ | 38 (15–113)                     | 47.5 (29–107) | 0.054 |
| Leukocyte count, \(\times 10^3\) cells/µL†‡ | 1.5 (0.7–3.0)                  | 1.5 (0.7–2.7) | 0.925 |
| ANC, \(\times 10^9\) cells/L†‡        | 969 (125–3,292)                | 920 (125–3,292) | 0.637 |
| Hemoglobin, g/dL†‡                | 12.9 (9.2–16.4)                | 13.5 (9.8–16.4) | 0.193 |
| aPTT, s, n = 30                   | 54 (35–97)                     | 54 (36–73) | 0.400 |
| AST, U/L, n = 31                  | 242 (63–4,567)                | 223 (63–2,145) | 0.109 |
| ALT, U/L, n = 31                  | 77 (27–1,432)                 | 63 (27–477) | 0.186 |
| Total bilirubin, mg/dL, n = 31    | 0.5 (0.2–4.0)                  | 0.5 (0.2–1.3) | 0.109 |
| Amylase, U/L, n = 26              | 84 (40–333)                    | 90 (40–333) | 0.540 |
| Lipase, U/L, n = 20               | 98 (38–682)                    | 117 (40–682) | 0.370 |
| CK, U/L, n = 21                   | 570 (67–4,362)                | 676 (67–2,760) | 0.651 |
| CK-MB, U/L, n = 18                | 6.9 (0.3–300.0)                | 7.6 (0.3–35.0) | 0.762 |
| LDH, U/L, n = 28                  | 908 (279–4,564)               | 893 (348–3,920) | 0.260 |
| ALP, U/L, n = 31                  | 119 (44–1,586)                | 79 (44–510) | 0.017 |

†ANC, absolute neutrophil count; aPTT, activated partial thromboplastin time; AST, aspartate aminotransferase; ALT, alanine aminotransferase; CK, creatinine kinase; CK-MB, creatinine kinase myocardial b fraction; LDH, lactate dehydrogenase; ALP, alkaline phosphatase.
‡Mann-Whitney U test was used to compare groups.
‡Minimum value.
relationships between a prognosis of death and high levels of LDH or ALP.

Our study has several limitations. First, mild cases of SFTS might have been missed, given that all of our case-patients were hospitalized. Thus, the true case-fatality rate in South Korea is likely lower than our reported value, related to case ascertainment bias. Second, because of the limited number of SFTS cases during 2013, we could not perform multivariate analysis for the prognosis. Therefore, our findings should be interpreted with caution. Third, because of the case series design, we could not analyze the risk factors for SFTS infection.

In conclusion, the clinical symptoms of SFTS in South Korea appear similar to those experienced by hospitalized SFTS patients in China. Older age and early neurologic symptoms were associated with a fatal outcome in studies in both countries. Expansion of SFTS surveillance into the outpatient sector, along with the development and incorporation of an SFTSV antibody test into the case detection algorithm, would detect milder cases and enhance completeness of SFTS case detection in South Korea.

Dr. Shin is an Epidemic Intelligence Service officer, currently working in the Division of Epidemic Intelligence Service, Korea Centers for Disease Control and Prevention. His research interests include the epidemiology of infectious diseases.

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