

Combination of western medicine and Chinese traditional patent medicine in treating a family case of COVID-19 in Wuhan

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Abstract In December 2019, an outbreak of novel coronavirus (2019-nCoV) occurred in Wuhan, Hubei Province, China. By February 14, 2020, it has led to 66 492 confirmed patients in China and high mortality up to ~2.96% (1123/37 914) in Wuhan. Here we report the first family case of coronavirus disease 2019 (COVID-19) confirmed in Wuhan and treated using the combination of western medicine and Chinese traditional patent medicine Shuanghuanglian oral liquid (SHL). This report describes the identification, diagnosis, clinical course, and management of three cases from a family, suggests the expected therapeutic effects of SHL on COVID-19, and warrants further clinical trials.

Keywords novel coronavirus (2019-nCoV); COVID-19; Chinese traditional patent medicine; Shuanghuanglian oral liquid

Introduction

Since December 31, 2019, a cluster of patients with pneumonia of unknown cause has been reported in Wuhan, China. This special pneumonia was associated with a novel coronavirus, 2019-nCoV, named initially by the World Health Organization (WHO) in January 2020 [1]. An outbreak of 2019-nCoV pneumonia (officially and internationally named as COVID-19 on February 11, 2020 by the WHO) occurred and spread to the entire China and multiple countries worldwide [2]. By February 14, the number of patients confirmed with COVID-19 reached 66 492, 8969 suspected patients, and 1523 deaths were identified in China [3]. By February 14, the number of confirmed patients and deaths were 37 914 and 1123, respectively, and the mortality of COVID-19 was 2.96% in Wuhan City, Hubei Province, which is considered as the origin of first cluster of patients [3]. The confusing and difficult thing for physicians is that they do not have specific drugs to either treat or prevent the aggravation and serious complications of COVID-19 for such patients [4]. Thus, Chinese herbs attract our attention, and investigations for different clinical trials are in progress. The present

report described a family case, including three cases who received western medicine and Chinese traditional patent medicine Shuanghuanglian oral liquid (SHL) treatments and achieved rapid recovery.

Case report

The family case includes parents and a daughter. The family lives in an apartment 3 km from the Huanan Seafood Wholesale Market in Wuhan, Hubei Province. These cases are a typical familial cluster where all of them had COVID-19 [5]. All the patients were informed about the clinical trial (ChiCTR2000029605) and signed informed consent.

Case 1 is a 51-year-old female. On January 17, 2020, the patient presented a feeling of general malaise and coldness. On January 19, she had fever with body temperature of 37.3 °C and experienced diarrhea and vomiting. She started to receive intravenous injection of cefotaxime in community clinic and took oral Jinyebaidu granules (another Chinese traditional patent medicine) and oseltamivir (75 mg, twice a day) for 4 days. She had persistent severe fever with body temperature from 37.6 °C to 38.3 °C for four days. On January 23, her chest computed tomography (CT) scan showed multiple patchy ground glass opacity and consolidation shadow in bilateral lung

and subpleural regions (Fig. 1A). She was highly suspected with COVID-19. Oral moxifloxacin and arbidol were prescribed, and she continued to take Jinyebaidu granules and oseltamivir. On January 24, her body temperature reached 39.9 °C and experienced severe fatigue, diarrhea, and breathlessness despite continuous treatments of the above drugs. She felt better after intravenous injection of immunoglobulin (IVIG, 5 g per day) and dexamethasone (5 mg, once to twice a day). However, the patient experienced recurrent fever (body temperature of 39 °C) and breathlessness, and her blood oxygen saturation fluctuated from 90% to 95% at the night of January 27. On January 28, the second chest CT scan indicated that her pneumonia aggravated for the past 5 days (Fig. 1B). A nasopharyngeal swab specimen was obtained and sent for detection of 2019-nCoV. Although the patient’s 2019-nCoV test was negative (Table 1), she

was diagnosed of COVID-19 in accordance with her symptoms and chest CT display. On the same day, she was confined into an isolation ward and started to take oral SHL (twice a day, 20 mL once). On the next day, SHL administration increased to three times a day (20 mL once) without using any other drugs. From January 29 to 31, the patient’s symptoms resolved with body temperature decreasing from 37.3 °C to 36.5 °C and without vomiting and diarrhea. The patient gradually felt strong except for slight cough. After February 6, the patient’s symptoms disappeared, and her third chest CT scan (Fig. 1C) showed significant absorption of bilateral ground glass opacity compared with the previous ones. The association of her symptoms with treatments is shown in Fig. 2, and the clinical laboratory results are shown in Table 2.

Cases 2 and 3 became simultaneously ill 9 days after they had close contacts with case 1 [5].

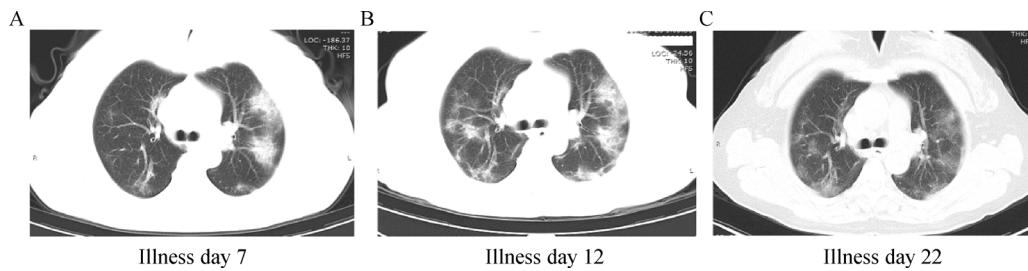


Fig. 1 Chest CT images of case 1. (A) CT imaging on January 23, 2020 shows ground glass opacity in both lungs on illness day 7. (B) Image taken on January 28, 2020, shows aggravation of pneumonia on illness day 12. (C) Image taken on February 6, 2020 shows the absorption of bilateral ground glass opacity after SHL treatment from January 28.

Table 1 Testing results of 2019-nCoV

Specimen	Case 1			Case 2			Case 3	
	Illness day 12	Illness day 20	Illness day 23	Illness day 3	Illness day 11	Illness day 14	Illness day 3	Illness day 11
Nasopharyngeal swab	Negative	Negative	NT	Negative	Negative	NT	Positive	Negative
Anal swab	NT	NT	Negative	NT	NT	Negative	NT	NT

NT, not tested.

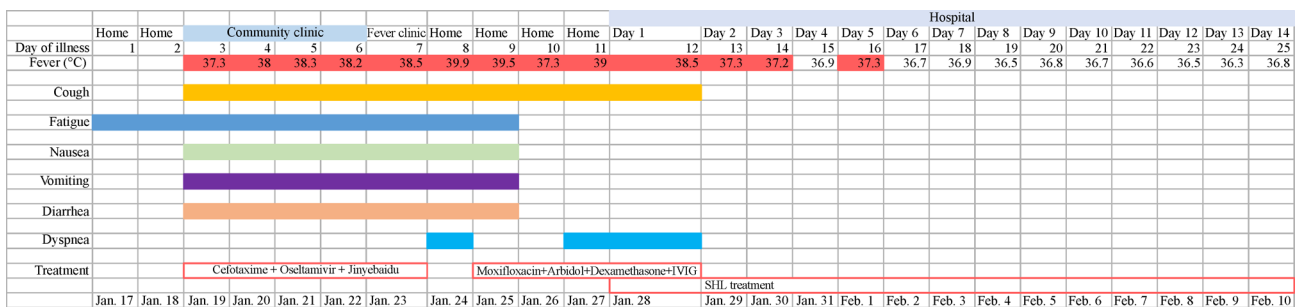


Fig. 2 Symptoms, maximum body temperatures, and treatment timeline in accordance with day of illness and day of hospitalization (January 17 to February 10).

Table 2 Clinical laboratory results of case 1

Measure	Reference range	Illness day 13	Illness day 24
		Hospital day 2	Hospital day 13
White cell count ($\times 10^9/L$)	3.50–9.50	5.93	5.46
Red cell count ($\times 10^9/L$)	3.80–5.10	4.13	4.06
Neutrophil count ($\times 10^9/L$)	1.90–6.30	3.04	3.27
Lymphocyte count ($\times 10^9/L$)	1.10–3.20	2.11	1.5
Eosinophil count ($\times 10^9/L$)	0.02–0.52	0.12	0.15
Platelet count ($\times 10^9/L$)	125.0–350.0	361	323
Hemoglobin (g/L)	115.0–150.0	113	111
Hematocrit (%)	35.0–45.0	35	35
Sodium (mmol/L)	136–214	140	140.4
Potassium (mmol/L)	3.50–5.10	4.22	4.8
Chloride (mmol/L)	99–110	104.6	102.4
Calcium (mmol/L)	2.15–2.50	2.06 ↓	2.19
Carbon dioxide (mmol/L)	22.0–29.0	21 ↓	25.1
Glucose (mmol/L)	4.11–6.05	–	5.01
Blood urea nitrogen ($\mu\text{mol/L}$)	2.6–7.5	3.1	3.6
Creatinine ($\mu\text{mol/L}$)	45–84	53	60
Total protein (g/L)	64–83	68.4	68.6
Albumin (g/L)	35–52	29.6 ↓	33.2 ↓
Total bilirubin ($\mu\text{mol/L}$)	≤ 21	6.4	4.5
Procalcitonin (ng/mL)	≤ 0.05	0.03	0.03
Alanine aminotransferase (U/L)	≤ 33	17	20
Aspartate aminotransferase (U/L)	≤ 32	16	19
Alkaline phosphatase (U/L)	35–105	59	83
Fibrinogen (g/L)	2.00–4.00	5.31 ↑	–
Lactate dehydrogenase (U/L)	135–214	248 ↑	–
Prothrombin time (s)	11.5–14.5	13.6	–
International normalized ratio	0.80–1.20	1.03	–
Creatine kinase (U/L)	≤ 170	35	27
C-reactive protein (mg/L)	≤ 3	57.8 ↑	–

↓ The patient's value was below normal. ↑ The patient's value was above normal.

Case 2 is a 27-year-old female nurse at the clinical trial center of Division of Cardiology of Tongji Hospital, Wuhan, daughter of case 1, who took care of her mother. On January 26, the patient presented mild weakness, diarrhea, and low fever, and she started to take oral Jinyebaidu granules, oseltamivir, moxifloxacin, and arbidol. On the next day, all her symptoms aggravated with body temperature reaching to 38.3 °C, frequent vomiting, and diarrhea for 5 times a day. At the night of January 27, her body temperature reached 39.5 °C with chest tightness and shortness of breath. Although she took nonsteroidal anti-inflammatory drug (loxoprofen), her fever remained high (39 °C). On January 28, her chest CT scan showed consolidation shadow in the left lung, and her 2019-nCoV test showed negative using nasopharyngeal swab specimen. Considering her contact history with her mother, she was also confined into an isolation ward and started to take SHL 20 mL once for three times a day without taking other

drugs. Her body temperature ranged from 37.5 °C to 38.5 °C during January 29 to 31 and decreased from 37.5 °C to 36.5 °C on February 1. All other symptoms resolved on February 2 with recovered appetite and spirit. Two repeated 2019-nCoV tests were negative (Table 1). On February 6, her second chest CT scan showed the absorption of the left lung shadow (Fig. 3). After her disease symptoms disappeared, the oral dose of SHL reduced to 10 mL once for three times a day. The patient's clinical laboratory results are shown in Table 3.

Case 3 is a 53-year-old male, husband of case 1 and father of case 2, who presented mild diarrhea, vomiting, and fever on January 26. On January 28, the patient was diagnosed of COVID-19 with positive 2019-nCoV test using nasopharyngeal swab specimen (Table 1) and chest CT scan showing patchy ground glass opacity in the right lower lung subpleural fields (Fig. 4). He had no fever, cough, and breathlessness. Thus, the patient started to

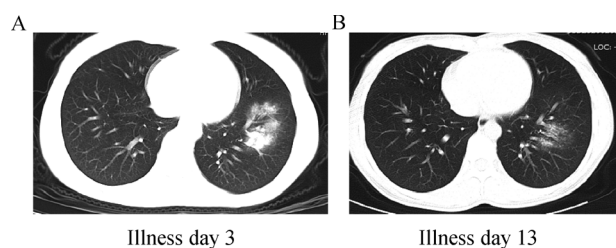


Fig. 3 Chest CT images of case 2. (A) CT imaging obtained from case 2 on January 28, 2020 shows consolidation shadow in the left lung on illness day 3. (B) Image taken on February 6, 2020 shows the absorption of consolidation shadow after SHL treatment from January 28.

isolate himself at home and took SHL (20 mL once, three times a day), moxifloxacin, and arbidol on January 28.

Since February 2, all his symptoms resolved with exception of light nausea, and he continued taking SHL with other drugs. On February 4, he felt slightly weak. On February 7, the patient fully recovered, and the repeated 2019-nCoV RNA test was negative (Table 1).

Discussion

This report described the first typical family case of COVID-19 treated using the Chinese traditional patent medicine SHL because of poor response to other treatments. The three patients were a close family, and the mother was the first victim with typical symptoms of severe viral pneumonia and confirmed as COVID-19 in accordance with the symptoms and chest CT scan. Cases 2 and 3 had close contact history with case 1, and 9 days later

Table 3 Clinical laboratory results of case 2

Measure	Reference range	Illness day 4	Illness day 15
		Hospital day 2	Hospital day 13
White cell count ($\times 10^9/L$)	3.50–9.50	3.63	5.79
Red cell count ($\times 10^9/L$)	3.80–5.10	4.5	4.28
Neutrophil count ($\times 10^9/L$)	1.90–6.30	2.12	3.2
Lymphocyte count ($\times 10^9/L$)	1.10–3.20	1.25	1.98
Eosinophil count ($\times 10^9/L$)	0.02–0.52	0.01	0.17
Platelet count ($\times 10^9/L$)	125.0–350.0	139	308
Hemoglobin (g/L)	115.0–150.0	125	117
Hematocrit (%)	35.0–45.0	37.9	36.2
Sodium (mmol/L)	136–214	135.3	143.5
Potassium (mmol/L)	3.50–5.10	3.69	4.27
Chloride (mmol/L)	99–110	97 ↓	106.3
Calcium (mmol/L)	2.15–2.50	2.25	2.27
Carbon dioxide (mmol/L)	22.0–29.0	22.3	25.2
Glucose (mmol/L)	4.11–6.05	4.31	4.82
Blood urea nitrogen ($\mu\text{mol/L}$)	2.6–7.5	2.7	3.4
Creatinine ($\mu\text{mol/L}$)	45–84	59	52
Total protein (g/L)	64–83	77.2	65.6
Albumin (g/L)	35–52	41.5	37.5
Total bilirubin ($\mu\text{mol/L}$)	≤ 21	6.6	4
Procalcitonin (ng/mL)	≤ 0.05	0.04	–
Alanine aminotransferase (U/L)	≤ 33	8	50 ↑
Aspartate aminotransferase (U/L)	≤ 32	21	27
Alkaline phosphatase (U/L)	35–105	52	22 ↓
Fibrinogen (g/L)	2.00–4.00	3.42	–
Lactate dehydrogenase (U/L)	135–214	170	186
Prothrombin time (s)	11.5–14.5	13.6	–
International normalized ratio	0.80–1.20	1.03	–
Creatine kinase (U/L)	≤ 170	52	27
C-reactive protein (mg/L)	≤ 3	9.4 ↑	–

↓ The patient's value was below normal. ↑ The patient's value was above normal.

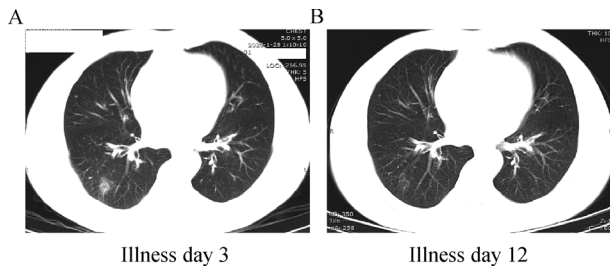


Fig. 4 Chest CT images of case 3. (A) CT imaging obtained from case 3 on January 28, 2020 shows ground glass opacity in the left lung on illness day 3. (B) Image taken on February 7, 2020 shows the absorption of ground glass opacity after SHL treatment from January 28.

they became ill with positive diagnosis of COVID-19. All their symptoms resolved after using the Chinese traditional patent medicine SHL and rapidly recovered without obvious adverse effects when the patients showed no response and their symptoms continued to aggravate after other treatments, including IVIG (5 g per day) and dexamethasone, antibiotics, and antiviral compounds.

SHL, a Chinese traditional patent medicine containing extracts of three Chinese herbs, namely, honeysuckle, forsythia, and *Scutellaria baicalensis*, is usually used to treat cold, sore throat, and cough with fever. SHL has been used in clinical practice for a long time because of its affordable cost and no serious adverse reaction. Recent news from preliminary study findings indicated that SHL can inhibit 2019-nCoV (http://www.cas.cn/yw/202001/t20200131_4733137.shtml, accessed on January 31, 2020). Considering that no specific drugs are recommended to treat COVID-19, we started our clinical trial (ChiCTR2000029605) to investigate whether SHL can treat this disease and the family case is a part of the clinical trial. These cases suggest that SHL might be effective for COVID-19 although subsequent clinical trials are needed. In this family case report, we described two patients who had poor response to other treatments but responded well to SHL therapy. Case 3 showed positive therapeutic effect although he simultaneously received arbidol. Early treatments may contribute to patients' outcome and several errors, including taking antibiotics, and combination of two antiviral drugs should be avoided. This report suggests that SHL treatment might be effective for COVID-19 and warrants subsequent clinical trials to obtain sufficient evidence for clinical recommendation.

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Compliance with ethics guidelines

Li Ni, Ling Zhou, Min Zhou, Jianping Zhao, and Dao Wen Wang declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article. Informed consent was obtained from all the patients in which their identifying information are included in this article. Other ethical board approval is not applicable in this case report.

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