Chemical pleurodesis and somatostatin in treating spontaneous chylothorax in pediatric patients: a retrospective analysis and review of the literature

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Background: Chylothorax is a rare disease, defined as an abnormal accumulation of chylous lymphatic effusion in thoracic cavity, with a high mortality rate in pediatric patients. At present, there are few studies on the treatment of pediatric chylothorax, and conservative treatments like somatostatin (SST) and pleurodesis are performed empirically. SST has been used for treating pediatric chylothorax over 20 years, and povidone-iodine chemical pleurodesis (PICP) is adopted in recent years with a high cure rate, but both the effect are still uncertain. The safety and efficacy of SST and PICP in treating pediatric chylothorax was compared in this study.

Methods: From January 2009 to May 2020, 27 pediatric chylothorax patients who accepted life support, SST and PICP treatment were retrospectively reviewed in this study. Their medical history, clinical manifestations, pleural effusion examinations, treatment methods, effects, and complications were screened. PICP and SST was performed in 19 and 8 children, respectively. Each patient was followed up after discharge from the hospital.

Results: After admission, chest X-ray and ultrasound diagnosed pleural effusion in 27 patients (18 males and 9 females), including 13 and 6 cases on the left and right side, and 8 cases on both sides. In the pleural effusion, the mean leukocyte count was \((9,826 \pm 9,482) \times 10^6\)/L, the mean lymphocyte ratio was \((84.82 \pm 6.58)\)% , the mean triglyceride content was 7.11±6.63 mmol/L. In PICP and SST group, the mean length of stay was 38.42±19.42 and 35.13±12.72 d (P=0.664), the mean time of thoracic drainage was 24.58±16.34 and 19.63±8.88 d (P=0.440), the mean time of parenteral nutrition was 27.16±18.29 and 25.25±13.52 d (P=0.793), respectively. PICP was performed for 3.16±2.27 times, and SST was used for 14.75±9.08 d with the dosage of 0.5 to 10 μg/kg/hour. After the treatment, all patients in PICP group had mild chest pain and low-grade fever, and 12 cases were observed with transient decrease of oxygen saturation. Eight patients in SST group had nausea and vomit, and 1 child had diarrhea simultaneously. Both were cured by symptomatic treatment, thoracic drainage and discontinuing using SST. Three children were diagnosed with refractory chylothorax and cured by PICP after SST. Pleural effusion in 27 patients was finally absorbed. All of them achieved a clinical cure successfully, and there was no recurrence during discharge follow-up.

Conclusions: PICP has a high success rate and few adverse reactions in treating spontaneous chylothorax in pediatric patients, expected to be a new treatment option for pediatric chylothorax after SST.

Keywords: Chylothorax; pleural effusion; pleurodesis; povidone-iodine (PI); somatostatin (SST)
Introduction

Chylothorax is a rare disease resulting from the abnormal accumulation of lymphatic fluid in the thoracic cavity, and it is one of the most common causes of pleural effusion in young children. Although congenital chylothorax demonstrates prevalence in pediatric patients, occurring with an incidence of 1:6,000 to 1:24,000 live births, the mortality associated with this disease could reach 60%, uniquely combining with premature birth, low birth weight, fetal edema, lung hypoplasia, hypoproteinemia, or continuous chylous exudation (1-4). In this study, the author classified congenital chylothorax and idiopathic chylothorax (in 5–10% of pediatric patients) as spontaneous chylothorax for the cause of both types of chylothorax is not found.

Chylothorax has been clinically reported for nearly a century, but due to its low incidence and few case reports, there are still insufficient guidelines and clinical pathways on its medical management. From the earlier experience, conservative treatment like multiple organ function support, thoracic drainage, somatostatin (SST), and pleurodesis are the first line treatment. Few children will finally be suggested to accept surgery, including ligation of the thoracic duct when conservative treatment fails (5).

SST has been used in treating chylothorax for twenty years, but due to the few reports on chylothorax, its curative effect and medication regimen are still unclear. Povidone-iodine chemical pleurodesis (PICP) is a minimally invasive treatment used in treating pneumothorax or refractory pleural effusion with a high success rate and few complications, so it is expected to be the alternative method in treating pediatric chylothorax in recent years. With this background, we had performed a retrospective analysis to compare the efficacy and safety of PICP and SST in treating spontaneous chylothorax in pediatric patients and review our experience simultaneously. We presented the following article following the STROBE reporting checklist (available at http://dx.doi.org/10.21037/tp-20-199).

Methods

Patients

Pediatric patients who were younger than 8-year-old and diagnosed with spontaneous chylothorax by chest X-ray, ultrasound, and cytological and biochemical analysis in the pleural effusion between January 2009 and May 2020, with no history of trauma, thoracic surgery other than thoracentesis and thoracic drainage, and other congenital and genetic diseases, were enrolled in this study.

After screening, 27 patients were enrolled and divided into the SST group and PICP group according to the therapeutic regimen. The data of their general condition, pleural effusion examination results, diagnosis, treatment, and adverse reactions were retrospectively analyzed and compared.

Written informed consent was obtained before the treatment. The ethical committee approved this study of Guangdong Second Provincial General Hospital (No. 20200615-YXXXYJ-LW-01-01). The research complied with the Declaration of Helsinki (as revised in 2013).

Diagnosis of chylothorax

The diagnosis of chylothorax in all the 27 children was confirmed by thoracic drainage after admission, which showed a milky or yellow turbid fluid on gross examination and positive chyle test when the patients accepted normal feeding, with increased levels of triglycerides ≥1.24 mmol/L, a total leukocyte count ≥1,000 cells/μL (1.0×10⁶/L) or at least 80% lymphocytes, and sterile (6,7).

Basic support treatment

After admission, chest X-ray and ultrasound were performed as soon as possible to confirm the affected side of the thoracic cavity in all the patients, and a chest tube was put into the thoracic cavity to drain out the pleural effusion. Chyle testing, cytological and biochemical examination were checked on the effusion, and blood routine, serum electrolytes, lipid test, C-reactive protein, liver, kidney, and thyroid function test, and electrocardiography were also performed.

After the tests were finished, patients were then given electrocardiographic monitoring and oxygen treatment. Nutrition support therapy was carried out combining the way of internal and parental nutrition according to the daily need for fluid and energy from the patients’ age and weight. Both enteral and parenteral nutrition were adopted in the treatment. Patients were only allowed to eat carbohydrates, electrolytes, vitamins, vegetables, and fruits; we only allowed a limited number of milk-fat products (1/4 to 1/3 of the daily demand) and liquids (1/3 of the daily demand). The rest of the calories were supplied with the parental nutrition solution comprised of glucose, amino acids, fat emulsions, and vitamins. Human immunoglobulin and albumin were given according to the results of blood tests. Cephalosporin
antibiotics were used to treat pneumonia. Blood routine and serum biochemical examination, liver and kidney function tests, chest X-ray, or ultrasound are rechecked routinely.

**PICP group**

After admission, the patient’s food intake was controlled to reduce the amount of chyle exudation. PICP was possible when the daily amount of pleural effusion decreased and supported in a low volume. A chest X-ray or ultrasound was performed in each patient before the treatment to confirm that the affected side had sufficient lung recruitment. Then, a small amount of normal saline was used to clean the chest tube to ensure smooth drainage. A modified PICP treatment method was adopted for treatment (8), that is, to increase the amount of povidone-iodine (PI) sclerosant, reduce the concentration of PI, and reduce the stay time of the sclerosant in the thoracic cavity. The vital signs of the patients were watched closely, and the sclerosant was then injected slowly through the chest tube when the signs became stable, and the children were calm. Depending on the reaction of the child’s response, the sclerosant was wholly or partially injected. If the child had apparent cyanosis, difficulty breathing, and a noticeable change in the vital signs like a rapid drop in the blood oxygen saturation, the injection was stopped immediately, sclerosant should be drained out, and rescue treatment should be performed if it was necessary. After the treatment, their parents changed their position appropriately, and the chest tube was connected to the drainage bag for continuous drainage. A chest X-ray or ultrasound was rechecked within 12–24 hours to confirm the condition of the thoracic cavity. If the pleural effusion was significantly reduced or disappeared, the chest tube was closed for 1–2 weeks, and chest x-rays or ultrasonograms was regularly rechecked to ensure no recurrence of the effusion. If the amount of pleural effusion was increased after reexamination, the chest tube was opened for drainage again, and PICP treatment could be considered when the patients had a steady general condition.

**SST group**

Eight patients were treated with SST simultaneously during the primary support treatment. Intravenous injection used SST with an intravenous infusion pump, with a starting dosage of 0.5 μg/kg/hour. If the child’s vital signs were stable, and no visible adverse reactions occurred, the dosage of SST could be gradually increased to 10 μg/kg/hour maximally. When the daily amount of pleural effusion decreased significantly and did not increase for about three days, the dosage could be gradually decreased until SST was stopped. During the treatment, the electrolytes, blood glucose, liver, kidney, and thyroid function should be re-examined regularly.

**Clinical cure standards**

After the treatment of PICP or SST, pleural effusion was absorbed, and no recurrence of effusion happened during the drainage tube was clamped for 1 to 2 weeks. No severe adverse reactions occurred during hospitalization (e.g., organ dysfunction). The results of the blood test and thyroid function were typical.

**Treatment and follow-up after discharge**

All patients were asked to keep on a low-fat diet after clinical cure, and their general condition was observed by their legal guardians and reported to us if it was abnormal in weight, mental state, daily activities, and any uncomfortable symptoms. Chest X-rays or ultrasound was rechecked for each patient on one month and three months after discharge. And they could have a regular diet if the pleural effusion had not recurred. The follow-up period was 1–10 years.

**Statistical analysis**

Data are presented as the minimum value to maximum value (in Tables 1,2) and mean ± standard deviation for categorical variables. Group comparisons performed on the mean length of stay, the mean total time of thoracic drainage, and parenteral nutrition were acquired using independent-samples t-test. All analyses were performed using IBM SPSS Statistics software (Version 22.0. IBM Corporation, NY, USA), with statistical significance set at P values less than 0.05.

**Results**

**Patient clinical characteristics**

Twenty-seven patients (18 males and 9 females) were enrolled in this study, aged between 19 hours and seven years old. All of them were admitted to the hospital due
Patients for 3.16±2.27 times. The sclerosant was used when the volume ranged from 20–50 mL during PICP, depending on the patient's general condition. After the treatment, all the children had symptoms of chest pain, including crying and restlessness, which could be relieved by feeding, appeasement, and other ways to divert their attention. Moreover, low-grade fever was recorded in all cases, with the highest temperature of 37.8 °C, and cured by physical cooling and ibuprofen suspension. Twelve cases displayed short of breath, with the blood oxygen saturation decreasing minimally to 90% shortly after the injection of sclerosant, which returned to normal after the drainage of sclerosant. During the treatment, 19 children showed good general condition and physical activities. Their pleural effusion was finally absorbed without recurrence confirmed by the chest X-ray or ultrasound.

Eight patients in the SST group accepted SST for 14.75±9.08 d, with a dosage of 0.5–10 μg/kg/hour. During the treatment, all the patients had nausea and vomit, and 1 of them had diarrhea simultaneously. Symptomatic treatment, reducing the dosage and stopping using SST relieved the above symptoms. After the treatment, five patients reached the standards of clinical cure. Three patients were diagnosed with refractory chylothorax by continuously recurrent pleural effusion after treated by SST for 12–31 days, with a maximum dosage of 10 μg/kg/hour. Daily drainage volume in these child had decreased from 24–29 to 8–13 mL/kg/d after the treatment, but the pleural effusion still recurred and could not be absorbed completely. After the symptoms of gastrointestinal dysfunction found in the

### Table 1 General condition, diagnosis, and pleural effusion results of patients in PICP group and SST group

<table>
<thead>
<tr>
<th>Group details</th>
<th>PICP group</th>
<th>SST group</th>
</tr>
</thead>
<tbody>
<tr>
<td>n (cases)</td>
<td>19</td>
<td>8</td>
</tr>
<tr>
<td>Age (h/d/m/y)</td>
<td>21 d–7 y</td>
<td>19 h–3 y</td>
</tr>
<tr>
<td>Gender (M:F, cases)</td>
<td>12:7</td>
<td>6:2</td>
</tr>
<tr>
<td>Side of chylothorax (cases)</td>
<td>Left 8, right 5, bilateral 6</td>
<td>Left 5, right 1, bilateral 2</td>
</tr>
<tr>
<td>The appearance of pleural effusion (cases)</td>
<td>Milky white 7, yellow 10, milky red 2</td>
<td>Milky white 2, yellow 6</td>
</tr>
<tr>
<td>Results of pleural effusion</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Leukocyte (×10⁶/L)</td>
<td>1,016–36,456 (10,183±10,331)</td>
<td>2,771–21,372 (8,697±6,737)</td>
</tr>
<tr>
<td>Lymphocyte (%)</td>
<td>72.1–93.2 (83.85±6.03)</td>
<td>77.3–97.7 (87.97±7.89)</td>
</tr>
<tr>
<td>Triglyceride (mmol/L)</td>
<td>1.35–19.23 (7.71±6.84)</td>
<td>1.32–19.81 (5.69±6.28)</td>
</tr>
<tr>
<td>Albumin (g/L)</td>
<td>6.90–35.2 (20.93±8.07)</td>
<td>10.10–36.30 (22.53±8.89)</td>
</tr>
</tbody>
</table>

PICP, povidone-iodine chemical pleurodesis; SST, somatostatin; h/d/m/y, hour/day/month/year; M:F, male:female.

After treatment of necessary support, PICP and SST

All patients had stable vital signs during the necessary support treatment. The symptoms of dyspnea were alleviated after the effusion was drained out, and the amount of pleural effusion gradually decreased and maintained in a low volume after diet control (about 1/3 of the daily amount measured at the time of admission). Cephalosporin antibiotics cured pneumonia diagnosed in 6 children.

PICP treatment was successfully performed in all
Table 2 Results of the treatment and statistical analysis in PICP group and SST group

<table>
<thead>
<tr>
<th>Outcomes</th>
<th>PICP group</th>
<th>SST group</th>
<th>P values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Length of hospitalization (d)</td>
<td>15–92 (38.42±19.42)</td>
<td>15–51 (35.13±12.72)</td>
<td>0.664</td>
</tr>
<tr>
<td>Time of thoracic drainage (d)</td>
<td>6–61 (24.58±16.76)</td>
<td>8–38 (19.63±8.88)</td>
<td>0.440</td>
</tr>
<tr>
<td>Time of parenteral nutrition (d)</td>
<td>3–69 (27.16±18.29)</td>
<td>11–43 (25.25±13.52)</td>
<td>0.793</td>
</tr>
</tbody>
</table>

PICP, povidone-iodine chemical pleurodesis; SST, somatostatin.

Figure 1 Appearance of pleural effusion before and after diet control in a part of the patients. The different appearance of pleural effusion, like milky white (A), milky red (B) had been found in a part of the patients when they were admitted to the hospital. After diet control, the pleural effusion gradually became yellowish and slightly turbid (C).

patients, SST was gradually stopped to avoid deterioration of the above adverse reactions. Considering a poor cardiopulmonary function and immunity and an elevated risk of anesthesia that these low-age patients had, PICP was adopted for treatment after SST instead of surgery. After 2–4 times of PICP treatment, the patients’ pleural effusion was gradually absorbed. Eight patients were clinically cured finally, and chylothorax did not recur in the follow-up period.

The statistical analysis results in the two groups were shown in Table 2. The chest X-ray results of each patient in the PICP group and SST group before and after treatment were shown in Figure 2, respectively. The pleural effusion was absorbed after a clinical cure.

Discussion

Spontaneous chylothorax is a common cause of pleural effusion in young children, and its etiology is unclear, usually considered associated with the abnormal growth of lymphatic vessels in the thorax (9). Because the organ function of young children is not yet mature and the tolerance of their body is relatively weak, the condition of chylothorax in pediatric patients is often more complicated than adult patients, so an appropriate distinction needs to be made in the process of clinical diagnosis and treatment. Children with spontaneous chylothorax are usually faced with a higher possibility of premature delivery with low birth weight. They often need to be treated by mechanical ventilation, thoracentesis, and thoracic drainage immediately for the reason of anoxia causing by atelectasis and acute respiratory distress syndrome. Continuously thoracic drainage is more likely to result in malnutrition, electrolytes disorder, and decreasing in immunity due to the loss of lipids, albumin, globulin, vitamins, and electrolytes as well...
as lymphocytes and antibodies through the effusion (10). It has a high possibility to cause septicemia and nosocomial infection during the treatment. Therefore, life support must be adopted at first to support a stable general condition for pediatric patients with chylothorax.

As the incidence of chylothorax is low, and there have been few related studies, the treatment efficacy has not been sufficiently demonstrated. It is challenging and unprecise to perform surgeries like lymphangiography, embolization, or ligation of the thoracic duct on pediatric chylothorax patients because their chest wall lymphatics are small and have many collateral vessels. Moreover, pediatric patients with chylothorax often have a poor cardiopulmonary function, with an elevated risk of anesthesia. According to most of the previous studies, conservative treatment (e.g., necessary life support, thoracic drainage, SST, and chemical pleurodesis) is the first choice for spontaneous chylothorax in pediatric patients at present (5). Thus, it is

**Figure 2** Comparison of chest X-ray of 1 child in PICP group and SST group respectively before and after clinical cure. A large and medium amount of pleural effusion in the right pleural cavity was found in one case in PICP group (A) and SST group (C) at admission. After the treatments, the effusion in two patients was absorbed respectively (B,D). Intestinal flatulence was a performance of gastrointestinal dysfunction found by X-ray in the case after SST treatment (D). PICP, povidone-iodine chemical pleurodesis; SST, somatostatin.
highly relevant to explore a safe and effective treatment to promote the absorption of pleural effusion, the self-healing of pleural lymphatic fistula, and the reconstruction of thoracic lymphatic circulation.

SST (or octreotide) is a type of regulatory inhibitory polypeptides, which can act on receptors in splanchnic vessels to decrease intestinal blood flow, gastrointestinal motility, intestinal fat absorption and inhibit the release of gastrin, vasoactive intestinal peptide, secretin, motilin, and pancreatic polypeptide. Therefore, it can decrease the exudation of chylous lymphatic effusion in the thoracic cavity and promote the self-healing of pleural lymphatic fistula. In 1998, Rimensberger et al. (11) firstly used SST to cure an infant with chylothorax. Nevertheless, due to insufficient clinical experience and related researches, the dosage and cure rate of SST in treating spontaneous chylothorax vary widely in different case reports and series in early stage (11). In 2010 and 2018, Das et al. (12) and Bellini et al. (13) had performed a meta-analysis and systematic review of the efficacy of octreotide in treating spontaneous chylothorax, with a success rate of 70% and 53%, respectively. Adverse reactions reported in the study included nausea, bloating, diarrhea or constipation, hyperglycemia, skin flushing, liver dysfunction, necrotizing enterocolitis, and transient hypothyroidism (14,15). Therefore, avoiding adverse reactions is a critical point for SST treatment. A starting dosage of 0.5–1.0 μg/kg/hour with an increment of 1–2 μg/kg/hour to a maximum of 10 μg/kg/hour, continuing using for 2–5 weeks was recommended referring to the treatment experience in relevant studies in recent years (7,16,17). SST’s treatment in the dosage mentioned above had been observed with better curative efficacy and safety. Besides, SST had been used either intravenously in a high dosage (20 μg/kg/hour and even more) or subcutaneously in an extended course for over five months also had been reported clinically, and no apparent adverse reaction was reported (15,18).

In conclusion, a larger sample of clinical trials still must demonstrate the appropriate dosage and course of treatment with SST in the treatment of spontaneous chylothorax in pediatric patients.

PICP is a minimally invasive treatment mainly used in conservative therapy for pneumothorax and refractory pleural effusion in adult patients (19), with a highly successful rate of up to 80–90% (20,21). It was firstly reported in the treatment of pediatric chylothorax in 2003 (22) and gradually adopted in recent years with its highly successful rate and excellent curative effect illustrated in some case reports (23–25). During the treatment, PI sclerosant was injected into the thoracic cavity to induce aseptic pleurisy and create adhesions between parietal and visceral pleura, and the thoracic cavity was closed, therefore; this was done to prevent chylous effusion exudates into the thoracic cavity. However, it was reported that the incidence of chest pain after PICP could reach 31–55% (20,21) and side effects caused by sclerosant like allergic reactions, thyrotoxicosis, and nephrotoxicity may occur. While other studies pointed out that the above side effects were usually rare, as long as the concentration and dosage of the PI sclerosant were well controlled to avoid being absorbed excessively by the pleura during the treatment process (26). Therefore, the proper use of PI sclerosant also needs further studies.

In this study, the efficacy of PICP and SST in treating spontaneous chylothorax in pediatric patients was compared. Statistics results showed no significant differences in the mean length of stay (P=0.664), the mean time of thoracic drainage (P=0.440), and the mean time of parenteral nutrition (P=0.793) between PICP group and SST group, which indicated two methods had a specific curative effect. However, when it comes to the adverse reactions and side effects, the symptoms displayed by the patients between PICP group and SST group were different in our study. Short-term of low-grade fever, and mild chest pain were the main adverse reactions found in 19 patients who accepted PICP treatment and could be relieved by the ways to divert the children’s attention and physical and drug treatment. Also, anhelation and decreased blood oxygen saturation recorded in 12 cases after PICP treatment were alleviated and returned to normal after the sclerosant was drained out. This condition was identified as an artificial compression on the affected lung caused by the sclerosant during the PICP process, but not an adverse reaction in fact, which could be solved by eliminating the compression after drainage of the effusion. In the SST group, gastrointestinal dysfunction was the main adverse reactions in our study. All the eight children had been found nausea and vomiting, and one patient had mild diarrhea simultaneously after SST treatment. Compared with chest pain and fever, it is more challenging treating gastrointestinal dysfunction in pediatric patients, not to mention the need to use SST to treat chylothorax. Finally, symptomatic treatment cured these eight children, reducing the dosage or stopping using SST. To avoid PI and SST related adverse reactions, we had adopted a modified PICP treatment and controlled the...
dosage of SST within 10 μg/kg/hour. Twenty-seven patients were discharged after meeting the clinical cure standards, and no complications and recurrence of chylothorax were reported during the follow-up period.

During the treatment, 3 cases of refractory chylothorax were found in the SST group. The daily volume of pleural effusion, the duration of conservative treatment, and the patient’s condition should be carefully considered for the diagnosis of refractory chylothorax. In some earlier studies, refractory chylothorax was defined in the case of drainage volume more than 10 mL/kg/day after 2–4 weeks of conservative treatment (27,28). Moreover, no significant improvement in drainage volume over ten days, or drainage volume of pleural effusion over 30 mL/kg/d, was considered meeting the surgical indications by some researchers (29).

Minimally invasive treatment like chemical pleurodesis with PI and other sclerosants (27,28) and surgeries like thoracic duct ligation and (29,30) diaphragmatic fenestration (31) had been reported before. In this study, three children were diagnosed with refractory chylothorax for failing to be cured with SST used intravenously for 12–31 days. Their daily drainage volume had decreased from 24–29 to 8–13 mL/kg/d after treating by SST, but their pleural effusion could not be absorbed completely. Surgery was not adopted considering the elevated risk of anesthesia, and PICP treatment finally cured them. However, up to now, there are still sporadic reports of refractory chylothorax in pediatric patients. More research is needed to prove the treatment protocols of refractory chylothorax.

Our study has limitations. With the limited sample size, the results and conclusion in our paper could not represent the efficacy of the two treatment methods, and publication bias is prone to appear in the SST group. Moreover, research approaches like random control and prospective observation did not be applied in this study. The theory of PICP and SST treatment is also different. PICP works in closing the pleural cavity and induces the reconstruction of the lymphatic circulation by causing pleurisy (32,33), and SST mainly promotes the self-healing of the lymphatic fistula by reducing the generation of lymph fluid and the pressure of lymphatic vessels. With the results and theory, we think that PICP is safer, more direct, and comprehensive, with slight adverse reactions, expected to become a new possibility for minimally invasive treatment of chylothorax in pediatric patients. As for refractory chylothorax cases, the two methods may be combined to explore a comprehensive treatment method. More randomized controlled trials with larger samples are needed for further study to prove the curative efficacy of PICP and SST treatment.

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Footnote
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Data Sharing Statement: Available at http://dx.doi.org/10.21037/tp-20-199

Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at http://dx.doi.org/10.21037/tp-20-199). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The research complied with the Declaration of Helsinki (as revised in 2013) and was approved by the ethical committee of Guangdong Second Provincial General Hospital (No. 20200615-YXKXYJ-LW-01-01). Written informed consent was obtained before the treatment.

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