### REVIEW

**Liver replacement therapy with extracorporeal blood purification techniques: current knowledge and future directions**


**Prediction models for recurrence in patients with small bowel bleeding**

Kim JH, Nam SJ

**Investigation of possible relationship between atopic dermatitis and salivary biomarkers, stress, and sleep disorders**

Estefan J, Ferreira DC, Cavalcante FS, dos Santos KRN, Ribeiro M

**Value of clinical applications of differential pressure and relative pressure imaging in the left ventricle**

Zheng AS, Yu HX

**Low-dose immunotherapy as a potentiator to increase the response with neo-adjuvant chemotherapy in oral cancers**

Rathinasamy N, Muthu S, Krishnan A

**Kidney disease in patients with chronic liver disease: Does sex matter?**

Cooper KM, Colletta A, Moulton K, Ralto KM, Devani D

### MINIREVIEWS

**Case Control Study**

**Elabela is a reliable biomarker for predicting early onset preeclampsia: A comparative study**

Amer Ali E, Nori W, Salman AF, Al-Rawi TSS, Hameed BH, Al-Ani RM

**Retrospective Cohort Study**

**Acute-on-chronic liver failure is independently associated with higher mortality for cirrhotic patients with acute esophageal variceal hemorrhage: Retrospective cohort study**

Terres AZ, Balbinot RS, Muscope ALF, Longen ML, Schena B, Cini BT, Rost Jr GL, Balensiefer JIL, Eberhardt LZ, Balbinot RA, Balbinot SS, Soldera J

**Retrospective Study**

**Elastic fiber degradation in the development of pediatric granuloma annulare: Report of 39 cases**

Zhang DY, Zhang L, Yang QY, Xie YC, Jiang HC, Li JZ, Shu H
### Contents

**Thrice Monthly Volume 11 Number 17 June 16, 2023**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>4026</td>
<td>Anti-bacterial mechanism of baicalin-tobramycin combination on carbapenem-resistant <em>Pseudomonas aeruginosa</em></td>
<td>Jin LM, Shen H, Che XY, Jin Y, Yuan CM, Zhang NH</td>
</tr>
<tr>
<td>4035</td>
<td>Acknowledging the use of botanicals to treat diabetic foot ulcer during the 21st century: A systematic review</td>
<td>Narzary I, Swarnakar A, Kalita M, Middha SK, Usha T, Babu D, Mochahary B, Brahma S, Basumatary J, Goyal AK</td>
</tr>
<tr>
<td>4059</td>
<td>Pregabalin induced balance disorder, asthenia, edema, and constipation in an elderly adult: A case report</td>
<td>Ma LP, Wen C, Zhao TX, Jiang XM, Gu J</td>
</tr>
<tr>
<td>4072</td>
<td>Hemophagocytic lymphohistiocytosis after autologous stem cell transplantation in angioimmunoblastic T-cell lymphoma: A case report</td>
<td>Zhang ZR, Dou AX, Liu Y, Zhu HB, Jia HP, Kong QH, Sun LK, Qin AQ</td>
</tr>
<tr>
<td>4079</td>
<td>Successful reconstruction of an ankle defect with free tissue transfer in a hemophilia A patient with repetitive hemorrhosis: A case report</td>
<td>Lee DY, Lim S, Eo S, Yoon JS</td>
</tr>
<tr>
<td>4084</td>
<td>Primary pelvic <em>Echinococcus granulosus</em> infection: A case report</td>
<td>Abulaiti Y, Kadi A, Tayib B, Tuergan T, Shalayiadang P, Abulizi A, Ahan A</td>
</tr>
<tr>
<td>4090</td>
<td>Epstein-Barr virus-induced infection-associated hemophagocytic lymphohistiocytosis with acute liver injury: A case report</td>
<td>Sun FY, Ouyang BQ, Li XX, Zhang T, Feng WT, Han YG</td>
</tr>
<tr>
<td>4098</td>
<td>Cardiac arrest secondary to pulmonary embolism treated with extracorporeal cardiopulmonary resuscitation: Six case reports</td>
<td>Qiu MS, Deng YJ, Yang X, Shao HQ</td>
</tr>
<tr>
<td>4105</td>
<td>Flared inflammatory episode transforms advanced myelodysplastic syndrome into aplastic pancytopenia: A case report and literature review</td>
<td>Ju B, Xiu NN, Xu J, Yang XD, Sun XY, Zhao XC</td>
</tr>
<tr>
<td>4123</td>
<td>Chest wall osteochondroma resection with biologic acellular bovine dermal mesh reconstruction in pediatric hereditary multiple exostoses: A case report and review of literature</td>
<td>Alshehri A</td>
</tr>
</tbody>
</table>
## Contents

**Thrice Monthly Volume 11 Number 17 June 16, 2023**

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>4133</td>
<td>Massive pulmonary embolism in Klippel-Trenaunay syndrome after leg raising: A case report</td>
<td>Lo CY, Chen KB, Chen LK, Chiu CS</td>
</tr>
<tr>
<td>4142</td>
<td>Improved super-elastic Ti-Ni alloy wire intrusion arch for skeletal class II malocclusion combined with deep overbite: A case report</td>
<td>Yang CY, Lin CC, Wang LJ, Chen YH, Yu JH</td>
</tr>
<tr>
<td>4159</td>
<td>Endoscopic and surgical treatment of jejunal gallstone ileus caused by cholecystoduodenal fistula: A case report</td>
<td>Fan WJ, Liu M, Feng XX</td>
</tr>
<tr>
<td>4168</td>
<td>Application of advanced platelet-rich fibrin for through-and-through bony defect during endodontic surgery: Three case reports and review of the literature</td>
<td>Algahtani FN, Almohareb R, Aljamie M, Alkhunaini N, ALHarthi SS, Barakat R</td>
</tr>
<tr>
<td>4179</td>
<td>Facial Merkel cell carcinoma in a patient with diabetes and hepatitis B: A case report</td>
<td>Ren MY, Shi YJ, Lu W, Fan SS, Tao XH, Ding Y</td>
</tr>
<tr>
<td>4187</td>
<td>Pregnancy and lactation-associated osteoporosis with pyogenic spondylitis: A case report</td>
<td>Zhai K, Wang L, Wu AF, Qian Y, Huang WM</td>
</tr>
</tbody>
</table>
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Massive pulmonary embolism in Klippel-Trenaunay syndrome after leg raising: A case report

Chih-Yu Lo, Kuen-Bao Chen, Li-Kuei Chen, Chiuan-Shiou Chiou

Abstract

BACKGROUND
Klippel-Trenaunay syndrome (KTS) is a rare congenital disorder characterized by a combination of capillary malformations, soft-tissue or bone hypertrophy, and varicose veins or venous malformations. The syndrome predisposes patients to hypercoagulable states, including venous thromboembolism and pulmonary embolism (PE).

CASE SUMMARY
A 12-year-old girl with KTS was scheduled excision of verrucous hyperkeratosis in the left foot and posterior aspect of the left leg and left thigh and excision of a cutaneous hemangioma in the right buttock. After induction, the surgeon elevated the patient’s leg for sterilization, whereupon she experienced a massive PE and refractory cardiac arrest. Extracorporeal membrane oxygenation (ECMO) was performed after prolonged resuscitation, and she had a return of spontaneous circulation. After this episode, the patient was discharged without any neurologic complications.

CONCLUSION
The mechanism of PE, a lethal disease, involves a preexisting deep vein thrombosis that is mechanically dislodged by compression or changing positions and travels to the pulmonary artery. Therefore, patients predisposed to PE should be prescribed prophylactic anticoagulants. If the patient has unstable vital signs, resuscitation should be started immediately, and extracorporeal cardiopulmonary resuscitation should be considered in settings with existing ECMO protocols, expertise, and equipment. Awareness of PE in patients with KTS while leg raising for sterilization is critical.
INTRODUCTION

Klippel-Trenaunay syndrome (KTS) is a rare congenital disorder and diagnosis of KTS is established when two of the triad features, viz. capillary malformation (port wine stains), hypertrophy of soft tissue or bones and varicose veins, are present. Vascular malformations involve the lower limbs, gastrointestinal tract, and genitourinary tract. The incidence has been estimated at two to five per 100000[1]. KTS predisposes patients to hypercoagulable states, including venous thromboembolism (VTE) and pulmonary embolism (PE). Dislodgement of the clots from pre-existing thrombosed veins upon movements and subsequent migration to pulmonary circulation can lead to massive pulmonary embolism. There have been no reports of severe pulmonary embolisms occurring after leg sterilization, except in cases where a massage or ultrasound examination was being performed.

CASE PRESENTATION

Chief complaints

A 12-year-old girl with KTS (height, 124 cm; weight, 20 kg) was referred to our hospital for recurrent left lower limb lymphangioma since the age of eight years.

History of present illness

She presented with clitoral hypertrophy, hemihypertrophy in the left lower limb along with varicose veins and multiple lipomas, and gastrointestinal bleeding. She was admitted for excision of verrucous hyperkeratosis on the left foot and posterior aspect of the left leg and thigh and excision of a cutaneous hemangioma on the right buttock.

Before induction, the patient was irritably crying in her father's arms at the time. The patient’s vital signs were as follows: Heart rate, 111 beats/min; blood pressure, 127/75 mmHg; saturation, 100%; respiratory rate, 18 breaths/min; and body temperature, 36.2°C. After preoxygenation, lidocaine (20 mg), propofol (50 mg), and cisatracurium (4 mg) were administered, and a 5.5 mm cuffed endotracheal tube was inserted. Monitoring was established using standard monitors and an intra-arterial line. Anesthesia was maintained with 45% oxygen and 2% sevoflurane.

After induction, the vital signs were as follows: Saturation, 100%; blood pressure, 97/61 mmHg; heart rate, 113 beats/min; respiratory rate, 17 breaths/min; tidal volume, 200 mL (controlled by ventilator); and body temperature, 36.2°C. Ten minutes after induction, her baseline blood gas test showed pCO2, 26 mmHg; pO2, 216.2 mmHg; HCO3 -, 19.0 mmol/L; and Hb, 8.4. Sixty minutes after induction, the surgeon elevated the patient’s leg for sterilization. Her end-tidal CO2 dropped from 35 to 7 mmHg within a few minutes. She had hypotension and bradycardia; therefore, atropine and epinephrine were sequentially administered intravenously. The ABG analysis was repeated, and testing revealed acidosis (pH, 7.263; pCO2, 42.4 mmHg; pO2, 69.4 mmHg; HCO3 -, 19.4 mmol/L; lactate, 1.2 mmol/L; and Ca2+, 1.04 mmol/L). Sixty-three minutes after induction, the patient have a cardiac arrest and cardiopulmonary cerebral resuscitation (CPCR) was started. Seventy-eight minutes after induction, a return of spontaneous circulation occurred, and transesophageal echocardiography (TEE) was performed. TEE revealed severe right ventricular distension and little blood flow to the right and left pulmonary arteries. Eighty-six minutes after induction, the patient had a second cardiac arrest and CPCR was...
started again. Ninety minutes after induction, a right femoral 5 French-size two-way central line was placed at 13 cm. A cardiac surgeon was consulted who conducted ECMO 106 min after induction. Thirty minutes after ECMO, a normal sinus rhythm returned. CPR was performed for approximately 70 min. The cardiac surgeon started target temperature management (TTM) of 32°C to protect the patient’s brain. Due to the incident of severe PE, the surgery was abandoned. Before getting transferred to the pediatric intensive care unit (PICU), her vital signs were as follows: Heart rate, 112 beats/min; blood pressure, 119/100 mmHg; saturation, 100%; respiratory rate, 17 breaths/min; and body temperature, 31.2°C (Figure 1).

Initially, in the PICU, her bilateral pupil was dilated approximately 9 mm with no light reflex. She was under sedation with a midazolam pump, as well as a dopamine (10 mcg/kg/min) and heparin pump. Computed tomography (CT) performed three hours after the surgery revealed PEs in the left truncus anterior, right interlobar, bilateral lower lobar, and segmental pulmonary arteries (Figure 2). Laboratory data showed elevated D-dimer (7939.76 ng/mL) and decreased fibrinogen (203.3 mg/dL) levels. Six hours later, her body temperature had risen from 31.2°C to 35°C, and her bilateral pupil size was 5 mm with a light reflex. A cardiologist was consulted for thrombolysis six hours after the surgery. Ultrasound-facilitated catheter thrombolysis with the Ekosonic Endovascular System (EKOS) was performed via the right femoral vein to the right upper pulmonary artery and left lower pulmonary artery for 48 h from the morning of the first post-incident day (Figure 3). On the third post-incident day, a post-procedure CT showed complete resolution of the PEs. The patient was weaned from ECMO on the fifth postoperative day, extubated on the eighth post-incident day, and prophylaxis treatment with enoxaparin was provided. Two months later, the patient was discharged without any neurologic complications (Figure 4). The patient had received the surgery 3 months later after the episode.

**History of past illness**
Four years ago, she had undergone extensive resection of the vascular malformations in the left lower limb, including amputation of the lateral fourth toe on her left foot (Figure 5). There was no record of difficulty in intubation.

**Personal and family history**
There was no specific personal or family history.

**Physical examination**
Physical examination presented with clitoral hypertrophy, hemihypertrophy in the left lower limb.

**Laboratory examinations**
Preoperative laboratory data showed anemia (hemoglobulin: 9.2 g/dL, mean corpuscular volume: 66.9 fL) no thrombocytopenia or coagulopathy (platelet count, 379000/μL; prothrombin time, 11.1 s; international normalized ratio, 1.07; and partial thromboplastin time, 27.9 s), liver and renal function were within normal limit, C-reactive protein: 7.43 mg/L.

**Imaging examinations**
The preoperative CT revealed angiomatosis extending from the left pelvis to the left foot, pelvic hemangiomas involving the rectum, a splenic lymphangioma, and engorged epidural vessels on the left side of the sacral canal. There is no obvious deep vein thrombosis (DVT) on the preoperative CT.

**FINAL DIAGNOSIS**
Klippel-Trenaunay syndrome complicated by refractory pulmonary embolism.

**TREATMENT**
The patient underwent ECMO and received ultrasound-facilitated catheter thrombolysis with the EKOS.

**OUTCOME AND FOLLOW-UP**
The patient was discharged without any neurologic complications.
DISCUSSION

KTS was first described by two French doctors, Klippel and Trenaunay (1900). KTS is a congenital vascular disorder characterized by a triad of main symptoms affecting one or more limbs, namely cutaneous hemangiomas, varicose veins, and bone and soft tissue hypertrophy. Approximately 95% of patients have lower limb malformations[2]. If capillary malformations are sufficiently large, the cutaneous lesions may sequester platelets[3]. In addition, large varicose veins may cause localized intravascular coagulation, and many patients have hematologic evidence of coagulopathy, defined by elevated D-dimer and soluble fibrin complex levels, decreased fibrinogen levels, and elevated prothrombin times, with normal to moderately low platelet counts[4]. Therefore, patients with KTS have a known significantly increased risk for PE[5].

PE while leg raising for sterilization before surgery is rare. The mechanism of PE is that a preexisting deep vein thrombosis is mechanically dislodged by compression or changing positions and travels to the pulmonary artery. Patients can be stratified before surgery based on the clinical prediction scoring system, the Modified Well’s criteria (Table 1) recommended by the European Society of Cardiology and the American College of Chest Physicians (ACCP), which standardizes the diagnosis and management of acute PEs. Our patient’s score of the Modified Well’s criteria is 1.5. However, the Modified Wells’
Table 1 The Clinical prediction scoring system

<table>
<thead>
<tr>
<th>Modified well's criteria</th>
<th>Points</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical symptoms of DVT</td>
<td>3</td>
</tr>
<tr>
<td>PE the most likely diagnosis</td>
<td>3</td>
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<tr>
<td>Heart rate &gt; 100/min</td>
<td>1.5</td>
</tr>
<tr>
<td>Immobilization or surgery with 4 weeks</td>
<td>1.5</td>
</tr>
<tr>
<td>Previous DVT or PE</td>
<td>1.5</td>
</tr>
<tr>
<td>Hemoptysis</td>
<td>1</td>
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<tr>
<td>Malignancy</td>
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</tbody>
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Figure 3 Catheter thrombolysis. Ekosonic Endovascular System-directed thrombolysis was advanced to, A: The left truncus; B: The right upper pulmonary artery.

Figure 4 The post-incident timeline for the patient. CT: Computed tomography; ECMO: Extracorporeal membrane oxygenation; PE: Pulmonary embolism.

criteria may not be applicable to patients with KTS, as the score may not accurately reflect the patient's high risk for pulmonary embolism[6]. Patients with KTS are at high risk and require a series of examinations, such as ultrasonography, CT, or magnetic resonance imaging. In some case reports[7-9], the thrombi can dislodge from the vein by leg massage, probe compression, and application of sterile elastic exsanguination tourniquets. If thromboembolism is suspected, prevention using anticoagulants and inferior vena cava (IVC) filters might be indicated. However, prophylactic anticoagulants are not prescribed in some Asian regions, and studies have reported a lower incidence of VTE in the Asian population than in Caucasians[10,11]. Although data on pediatric thromboembolism prophylaxis are limited, a systemic review demonstrated that low-molecular-weight heparin is safe and effective in
children[12]. The IVC filter is ineffective in KTS because of an anomalous venous connection between the lower extremities and the IVC, effectively bypassing the filter[13]. Considering the risk of lower limb thrombophlebitis associated with venous anomalies, femoral cannulation is better avoided[14].

If PE is suspected intraoperatively, supportive care should be started in unstable patients, including ventilation with 100% O₂, use of positive end-expiratory pressure, resuscitation of the circulating fluid volume, and administration of inotropic drugs as early as possible. In addition, invasive monitors should be placed for diagnosis and management, including arterial and central venous lines, transthoracic echocardiography, and trans-esophageal echocardiogram, which is generally considered the primary diagnostic technique for identifying intraoperative PE because of its high safety, availability, and utility in the operating room and its lack of interference with resuscitation efforts[15]. If vital signs are profoundly unstable, cardiopulmonary cerebral resuscitation should be performed, and a cardiac surgeon should be consulted for cardiopulmonary bypass or venoarterial ECMO. Extracorporeal life support is an effective therapy for unstable patients, offering acceptable outcomes; however, these studies lacked pediatric patients. The American Heart Association guidelines, updated for pediatric advanced life support in 2019[16], recommend ECPR for pediatric patients with cardiac diagnoses and an in-hospital cardiac arrest in settings with existing ECMO protocols, expertise, and equipment. For infants and children who remain comatose after out-of-hospital or in-hospital cardiac arrest, either TTM from 32°C to 34°C followed by TTM of 36°C to 37.5°C or TTM of 36°C to 37.5°C can be used. There were no significant differences in the outcomes between the two TTM groups in the Therapeutic Hypothermia after Pediatric Cardiac Arrest trials[17].

Postoperatively, the patient should be admitted to the intensive care unit, and CT pulmonary angiography should be performed. If there is no major bleeding, anticoagulant treatment should be started as soon as possible, and experts should be consulted for systemic or catheter-directed thrombolysis.

In current standard therapy, if the patient has no major active bleeding, the ACCP recommends immediate initiation of parenteral anticoagulation using low-molecular-weight heparin and fondaparinux, as they are superior to intravenous unfractionated heparin, owing to a lower risk of adverse bleeding events and the absence of need for serial laboratory monitoring. However, there are no guidelines or group consensus for therapies for intraoperative PE. Therefore, the authors organized a reference algorithm for intraoperative PE management (Figure 6).

Despite the lack of established strategies for anesthetic management of KTS patients undergoing surgery, some preoperative evaluations have been recommended. Firstly, anesthesiologists should consider the potential difficulties in managing the airway due to soft tissue hypertrophy in the mouth, hypopharynx, and facial anomalies commonly observed in KTS patients. Secondly, neuraxial anesthesia should be avoided as it is contraindicated due to the possibility of neurovascular malformations in the spinal cord and surrounding structures. It must be noted that central regional blockade can be performed safely in KTS patients, provided that vascular malformations in the central nervous system have been ruled out through computed tomography/magnetic resonance imaging and that there are no cutaneous lesions at the site of needle insertion. Thirdly, the risk of excessive intraoperative blood loss should always be taken into consideration, even in minor surgeries, due to the presence of widespread varicosities and venous malformations. Fourthly, KTS patients have a relatively high risk of developing venous thrombosis and pulmonary thromboembolism. Chronic coagulopathy such as disseminated intravascular coagulation can also occur in these patients. Finally, intracerebral aneurysm is a potential complication in KTS patients, which can rupture during the perioperative period[14,18,19].
CONCLUSION

KTS is a congenital vascular disorder that primarily affects the lower limbs and is characterized by cutaneous hemangiomas, varicose veins, and bone and soft tissue hypertrophy. Patients with KTS are at high risk for PE. All patients having KTS should be evaluated for lower limb circulation and pre-existing DVT pre-operatively. Also, all patients with KTS should receive DVT prophylaxis at least 8 h prior to surgery irrespective of the age. The Modified Wells' criteria may not be applicable to patients with KTS. Care should be taken to monitor for PE in patients with KTS while leg raising for sterilization. Intraoperative PE is lethal, and the management of hemodynamically unstable patients requires efficient CPR and early mechanical support. Anesthesiologists should consider potential difficulties in managing the airway and avoid neuraxial anesthesia. Central regional blockade can be performed safely in KTS patients if vascular malformations in the central nervous system have been ruled out.

FOOTNOTES

Author contributions: Lo CY and Chen KB are the first authors; Lo CY and Chen KB conceptualized and drafted the initial manuscript, and reviewed and revised the manuscript; Chiou CS and Chen LK conceptualized and coordinated, reviewed, and revised the manuscript; All authors approved the final manuscript as submitted and agree to be accountable for all aspects of the work.
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