A case of sclerotherapy for a giant lymphatic malformation in the neck

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Abstract: Lymphatic malformation is an abnormal development in the local area of the lymphatic system, often occurring in the head and neck of children within 2 years of age. Although this is a benign lesion, it rarely resolve spontaneously, enlarges and suppresses the vital organs around it, endangering life. We herein reported a case of a neonate with a large lymphatic malformation in the neck. The lesion was about 8.0 cm × 8.0 cm × 3.60 cm at birth. After multidisciplinary jointed diagnosis and treatment, sclerotherapy with pingyangmycin combined with triamcinolone acetonide or lauromacrogol foam remained very effective. The child was followed up until 1 year and 5 months, and the growth and development of the child were good. This case may demonstrate that large lymphatic malformations in the neck do not necessarily require surgical removal, and sclerotherapy can be effective.

Keywords: Lymphatic malformation; neck; sclerotherapy; pingyangmycin; multidisciplinary

Case presentation

A boy, born with a large lump in the left side of the neck, presented to our hospital 5 days later on January 29, 2018 (Figure 1). Physical examination revealed that the left neck mass was raised above the surrounding skin with an unclear boundary by naked eye. There was no pain upon pressure, and the skin temperature was normal. Pulsation was not obvious, and there was no rupture or infection. The child experienced difficulty in breathing and made a snoring sound while sleeping. Ultrasound examination revealed a cystic mass under the skin and multiple sections without obvious blood flow signals. The lymphatic malformations were approximately 8.0 cm × 8.0 cm × 3.6 cm in size (Figure 2). The child weighed 3 kg and was too young for surgery and the polycystic lesion located in the neck had the potential to affect breathing; hence, we considered sclerotherapy as a treatment option. The injectable drugs included lauromacrogol foam and triamcinolone acetonide. Lauromacrogol solution (1%) and air were quickly mixed in a 1:4 ratio to make foam using the Tessari method (8). Triamcinolone acetonide was diluted to 13.3 mg/mL with...
2 mL of 0.9% normal saline. A small amount of lauromacrogol foam and triamcinolone acetonide was injected in multiple areas using the double-needle method. Two 4.5# scalp needles were placed above and below a certain area to inject each sclerosing agent. The cystic fluid and a small amount of sclerosing agent was taken out. A faint yellow clear liquid was extracted from multiple points in the cyst. Cytological examination revealed the presence of many mature lymphocytes in the liquids; no pathological cells were found. Another two rounds of treatment, each treatments with an interval of 3 weeks, did not have a satisfactory effect and the sac fluid gradually became dark red.

Figure 1 The child during the treatment. (A) The child at the age of 5 days. (B) The child at the age of 3 months. (C) Injection of pingyangmycin 1 month later. (D) Injection of pingyangmycin 2 months later. (E) The child at the age of 1 year and 5 months.

Figure 2 Ultrasonography during treatment. (A) Ultrasonography on January 29, 2018. (B) Ultrasound examination after 2 months of pingyangmycin treatment. (C) Ultrasound examination of the child at the age of 1 year and 5 months.
By April 8, 2018, when the child was 2.5 months old, the size of the lesion has increased to 11.5 cm × 12.0 cm × 7.7 cm (Figure 1B) and the child's body weight was 4.5 kg. Pingyangmycin was added at the fourth treatment. Pingyangmycin was diluted with 4 mL of 0.9% normal saline and 2 mL multipoint injections were given. After a small dose of the sclerosing agent into cavity, it was washed with a small amount of 0.9% normal saline and was inserted from above and removed from below to reduce excess drug in the capsule and reduce side effects. For this procedure, 330 mL of cyst fluid was removed (Figure 3). Figure 4 shows the double-needle injection method used during the operation. After the operation, the child was transferred to the pediatric intensive medicine department at our hospital. The child's postoperative condition remained stable and no special treatment was given. The child was discharged after 3 days post-operation. During the 1-month treatment interval for pingyangmycin (9), and the child was closely followed up. During the follow up, the condition of the child was stable and the size of the lesion had significantly reduced. On April 22, 2018, the child was re-examined (Figure 1C) and the previous treatment procedure was repeated. After the operation, the child was again transferred to the pediatric intensive medicine department for 1 day. The child's postoperative condition was stable and was discharged without any special treatment. On May 29, 2018, the child was re-examined (Figure 1D), and the ultrasound examination showed a lesion of about 7.5 cm × 9.0 cm × 5.12 cm in size (Figure 2B). The treatment plan was continued as described above.

Subsequently, the child was closely followed up once a month, and the size of the lesion was gradually reducing. On October 15, 2018, a reexamination ultrasound revealed that size of the lesion was reduced to approximately 6.0 cm × 2.3 cm × 6.1 cm, and the child's condition was stable. Therefore, we decided not to intervene any further. On December 18, 2018, the child's parents complained of a sudden increase in the size of the lesion within 1 week and visited our hospital for treatment. Ultrasound results showed lymphatic malformations combined with hemorrhage. We continued the treatment that was previously effective. During the operation, 45 mL of dark red liquid was drained and the previous treatment plan was continued. On March 20, 2019, when the patient was 1 year and 2 months old, the lesion was stable (Figure 1E). Ultrasound results revealed that the size of the lesion was further reduced to approximately 5.6 cm × 3.4 cm × 5.3 cm (Figure 2C); however, the child was continued to be followed up.

**Discussion**

Head and neck lymphatic malformations account for 75% of all lymphatic malformations, and this may be due to the abundance of the lymphatic system in the head and neck (7). Under normal circumstances, lymphatic malformations grow alongside the growth of the child, but there are two worrying complications, infection and intralesional bleeding (1). In our case, the child also suffered from intralesional bleeding during the therapeutic process. Lymphatic malformations are regarded as benign lesions, but more attention should be paid to these in clinical practice (10). Even if the malformation is controlled, it still needs daily care and attention.

Traditionally, surgical resection was preferred treatment option (11); however, many researchers currently believe that sclerotherapy is a good alternative to surgical treatment (12). There are many types of sclerosing agents. Yura et al. were the first to use bleomycin injection for
the treatment of lymphatic malformations in 1977 (13). Recently, treatment with pingyangmycin has become popular in China. Pingyangmycin, also known as bleomycin A5, has shown good results (14). Furthermore, triamcinolone acetonide can also enhance the efficacy of pingyangmycin (15). Wang et al. showed that lauromacrogol foam is effective for treating head and neck lymphatic malformations (8). Moreover, the treatment of cervical lymphatic malformations requires a multidisciplinary team, involving the surgery, anesthesia, pediatric, and imaging departments. Even if the treatment is nearly complete, cervical lymphatic malformations cannot be taken lightly; participation of a multidisciplinary team provides safer and more effective treatment outcomes.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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References


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