Fibromatosis colli or pseudotumor of infancy of the sternocleidomastoid muscle is a rare benign cause of neck swelling or mass in neonates and infants. Though the exact etiology is not known, it is most likely due to birth trauma or malposition in uterus. It is one of the cause of congenital torticollis and usually recognized by mother as neck swelling with or without restricted neck movement. Ultrasonography of neck is the imaging modality of choice and sometimes CT scan or MRI scan may be required to further characterize the disease and extent of involvement. Real time ultrasonography demonstrate synchronous motion of the mass with the sternocleidomastoid muscle, thus confirming the diagnosis.

We present a case report of fibromatosis colli in a 1 month old neonate diagnosed using ultrasonography. In addition to this brief discussion was made on literature review, clinical and radiological findings and differentials.

2. Case Report

A one month old neonate was referred to the radiology department for ultrasonography of a neck swelling on the right side which was incidentally noticed by the mother 10 days ago. On examination the neck swelling was firm, freely mobile, no warm on touch and approximately 1.5-2 cm soft tissue mass was felt on the right side of neck (figure 1). Neonate was afebrile. There was restriction of neck movements on the affected side and chin is directed away. Past history revealed prolonged labour with breech presentation; however no assisted delivery was noted. Baby did not cry immediately after birth and was admitted for three days in neonatal intensive care unit. Later baby started feeding well and was immunized till date. Ultrasonography of neck showed ill-defined fusiform thickening of right distal sternocleidomastoid muscle with heterogeneous echotexture (figure 2). The fibrillar structures of muscle fibers are well maintained. On color doppler study, minimal peripheral vascularity with high resistance flow is noted (figure 3).
heads of the muscle and seen more commonly in male child. About 14-20 % cases of congenital torticollis are due to contraction of the sternocleidomastoid muscle. The exact cause is unclear; it is likely related to birth trauma greater than 90% of cases associated with a difficult or forceps delivery. Some cases are due to abnormal utero fetal head position, which causes selective injury and leads to development of a secondary compartment syndrome, pressure necrosis and fibrosis within the sternocleidomastoid muscle. No cervical lymphadenopathy, vascular invasion or bony involvements are seen as compared other neck masses.

Ultrasonography (USG) is the best imaging modality for diagnosis due to its relative low cost, its lack of radiation and the proximity of the lesion to the skin. USG findings show fusiform thickening or diffusely enlarged sternocleidomastoid muscle with variable echogenicity. Color doppler interrogation may reveal a high resistance waveform. The enlarged area often moves synchronously with the rest of the sternocleidomastoid muscle (SCM) on real time sonography. Almost all the cases can be identified by ultrasonography as compared to CT or MRI. Computed tomography (CT) show a homogeneous diffusely enlarged sternocleidomastoid muscle with surrounding well preserved fat planes. At times calcification may be present. Magnetic resonance imaging (MRI) shows decreased signal intensity of the mass on T2W images as compared to gradient-recalled T1W images, because of the presence of fibrous tissue. The extent of involved muscle is better delineated with MRI than with USG. Radiographs are usually normal and sometimes used to exclude congenital bony abnormality. Histological features include bland-appearing fibroblasts, myofibroblasts and atrophic skeletal muscle along with muscle giant cells and bare nuclei.

Fibromatosis colli is self-limiting condition and usually resolves within 4-8 months and requires physiotherapy and neck stretching exercises. Approximately two-third cases regress by the age of 2 years. The differential diagnoses of solid masses in the neck include neuroblastoma, rhabdomyosarcoma, lymphoma and other sarcomas. In addition to neck mass, may have enlarged cervical lymph nodes, vascular encasement, or invasion of surrounding structures. Cystic mass in the neck near the midline includes cystic hygroma, branchial cleft cyst, dermoid cyst, teratoma or thyroglossal duct cyst are considered in the differentials.

4. Conclusion

Our case highlights, fibromatosis colli or pseudotumor of infancy of the sternocleidomastoid muscle is a rare benign cause of neck swelling or mass in neonate and infants. The radiologist must be aware of its imaging features in order to differentiate it from other neck masses. This entity is a self-limiting condition and resolves over a period of time. If diagnosed correctly can be managed conservatively and prevent unnecessary diagnostic and therapeutic maneuvers.

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Figure 3: Longitudinal USG image of normal sternocleidomastoid muscle (left).

Figure 4: Transverse USG image of the neck shows oval soft tissue mass at distal right sternocleidomastoid muscle.

Figure 5: Transverse USG image with normal bulk in the left sternocleidomastoid muscle.

3. Discussion

Fibromatosis colli is a rare benign condition in neonates and infants with neck swelling. Most cases show no abnormality at birth but manifest between the 2nd and 4th weeks of life as a firm soft-tissue mass in the lower one-third of the sternocleidomastoid muscle. It is usually unilateral, slightly more common on the right side while bilateral involvement is rare. It affects both the sternal and clavicular
was received in this work.

References