



The Bairn's Blain- Fibromatosis Colli



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Received: July 24, 2019; Published: August 28, 2020

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Preface

Fibromatosis colli is an exceptional, benign neoplasm of infancy constituted by spindle-shaped cells of sternocleidomastoid muscle. Fibromatosis colli emerges within specific sites such as distal or inferior segment of sternocleidomastoid muscle and is accompanied by diffuse enlargement of the muscle. Although nomenclated as “sternocleidomastoid tumour” or “sternocleidomastoid pseudo-tumour of infancy”, the designation is a misnomer, as the condition is non neoplastic although it may be denominated as a congenital fibrotic disorder. Fibromatosis colli was initially scripted by Hulbert in 1812 as “sternomastoid tumour torticollis” in German literature [1]. Thereafter, Chandler and Altenberg demonstrated a characteristic, hard, immobile, fusiform swelling within the sternocleidomastoid muscle, usually discerned within two weeks of birth, the magnitude of which subsequently enhances for two to four weeks until it attains the dimension of a “very large almond” [2]. Coined by Hulbert as “tumour torticollis” of sternocleidomastoid muscle, fibromatosis colli is a benign neoplasm associated with congenital fibrosis and manifests a swelling within lateral cervical or supraclavicular region [1]. The neoplasm ensues from fusiform expansion of sternocleidomastoid muscle concomitant with muscle contraction leading to torticollis and cervico-fascial asymmetry. Fibromatosis colli or sternocleidomastoid tumour is a self-limiting, fibroblastic lesion accompanied by congenital torticollis and history of birth trauma [3].

Keywords: Fibromatosis colli; Trauma; sternocleidomastoid; Birth injury; Tumour torticollis; Fibroblastic lesion; Congenital torticollis

Disease Characteristics

Of uncertain aetiology, possible birth trauma is suggested as a mechanism of tumour emergence although cogent mechanics with consequent muscle fibrosis remain unexplained. Several theories are posited in the emergence of fibromatosis colli such as foetal malposition, birth trauma, ischemic necrosis following vascular compression during birth, neonatal infection, and occurrence of endogenous factors [3,4].

A preferred premise is occurrence of birth trauma or micro trauma of sternocleidomastoid with engendered from difficult or assisted labour. Pertinent history of a laborious birth is discerned in an estimated 50% instances. Trauma with birth, prolonged difficult labour, breech presentation, forceps delivery and primiparous birth are implicated in genesis of the condition [3,4]. Another plausible hypothesis is obstruction of venous outflow within the muscle during delivery or intrauterine development. Birth injury engenders necrosis with subsequent fibrosis within muscle fibres, thereby inciting a secondary pressure within the muscle. The tumefaction occurs from an in-utero positioning of foetal head which initiates a selective injury to sternocleidomastoid muscle. With pertinent intramuscular injury, development of secondary compartment syndrome with

consequent pressure necrosis and fibrosis of the muscle ensues. Anomalous sternocleidomastoid muscle in association with fibrosis, secondary muscle contracture and manifestations of a compartment syndrome with ischemic lesions occurring due to intrauterine foetal malposition is theorized. Dual mechanism may be interlinked with foetal malposition terminating in laborious delivery [3,4]. The uncommon condition is discerned in around 0.4% of live births wherein prevalence ranges from 0.3% to 2% and majority of instances are detected prior to six months. However, roughly 50% neoplasms are detected beyond 6 months of neonatal life. A male predilection is observed [3,4]. An estimated 80% of the benign neoplasm gradually decrease and resolve completely. Concomitant congenital torticollis is documented in around 14% to 20% infants [3,4]. Right sternocleidomastoid muscle is incriminated in an estimated 60% to 75% instances whereas bilateral tumour occurrence is delineated in around 22% subjects [3,4]. Fibromatosis colli is generally discerned between 2 weeks to 4 weeks of neonatal life although majority of tumefaction are detected prior to six months. Mean age of disease representation is twenty-four days. Fibromatosis colli can be associated with congenital defects, particularly congenital muscular disorders as club foot or congenital dislocation of hip [3,4].

Clinical Elucidation

Trichilemmal carcinoma arises as a solitary, indolent neoplasm although multiple lesions can emerge upon non-exposed cutaneous surfaces. Trichilemmal carcinoma is a gradually progressive neoplasm cogitated within an elderly population, appearing as a pale, tan or reddish papule, indurated plaque or a nodule. The lesion is generally discerned upon scalp, forehead, face, neck, trunk, upper extremities, temporal or occipital region, nape or lower lip and demonstrates a varying magnitude betwixt 0.4 centimetre to 8 centimetres with a mean dimension of 2.8 centimetres. Majority (80%) of lesions demonstrate a clear, tumour-free margin. Prior to appropriate detection, the tumefaction can occur from 2 months to up to 50 years, followed by rapid tumour progression [3,4]. As trichilemmal carcinoma can be frequently misinterpreted as a benign lesion, a clinical Fibromatosis colli is accompanied by a characteristic clinical history and representation. Infant typically manifests a smooth, fusiform mass within the distal or inferior one third of sternocleidomastoid muscle and stiff neck with restricted neck mobility. Clinically, a lateral cervical swelling occurring during early childhood or within first month of life is indicative of fibromatosis colli. As the tumefaction is generally incorporated within distal one third of sternocleidomastoid muscle, the mobile mass can be palpated beneath superimposed cutaneous surface [5,6]. Fibromatosis colli typically demonstrates an intramuscular mass within one week to eight weeks which is accompanied by torticollis and a non tender, firm, fusiform swelling within the body of sternocleidomastoid muscle [5,6]. Classically, a palpable, solid tumefaction is observed within anterior neck segment superimposed upon the sternocleidomastoid muscle. Tumour nodule is usually absent at birth and commonly appears between second week to fourth week of neonatal life. Lateral cervical tumefaction induces a reduced lateral mobility of the head within first few weeks. The intramuscular mass demonstrates an initial growth phase and can increase for several weeks followed by a phase of stabilization, which lasts for a few months and finally retrogresses spontaneously within four months to eight months [6]. Tumour nodule is devoid of additional clinical anomalies. Although fibromatosis colli can induce respiratory failure and may necessitate a subsequent tracheostomy, the complication remains undocumented [4].

Histological Elucidation

On gross examination, a tan coloured, gritty nodule, confined to the sternocleidomastoid muscle is delineated. Characteristically, smears obtained by fine needle aspiration cytology are composed of bland appearing, plump or normal fibroblasts and proliferating fibroblasts with an admixture of degenerative, atrophic skeletal muscle cells, giant myocytes with several nuclei, scattered, bare, bland nuclei or cells with fragmented, wispy cytoplasm and parallel aggregates of fibroblasts. The background is clear and devoid of cellular or nuclear debris. Sheets of spindle-shaped fibroblasts, atrophic muscle fibres and regenerating, multinucleated muscle giant cells are intermingled with the fibroblastic component. On

microscopy, a collagen- rich, minimally cellular, fibrotic soft tissue nodule is exemplified which simulates a scar or conventional fibroma. The neoplasm is comprised of uniform, plump, fibroblastic, or myo-fibroblastic cells encompassed in a collagen-rich stroma with infiltration and entrapment of skeletal myocytes [5,6]. Cogent histological examination depicts replacement of muscle fibres and muscle mass with fascicles of fibrous tissue comprised of mature fibroblastic cells, thereby conferring fibrosis [6] (Figures 1-8).

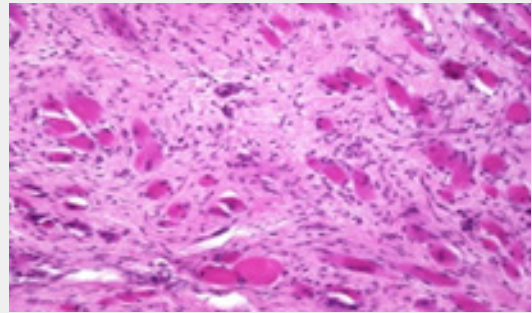


Figure 1: Fibromatosis colli depicting bundles of fibroblasts and myo-fibroblasts intermixed with a collagenous stroma and entrapped myocytes [9].

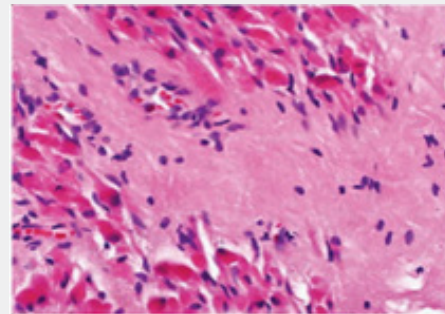


Figure 2: Fibromatosis colli demonstrating fascicles of fibroblasts and myo-fibroblasts within an abundant collagenous stroma and several mature myocytes [10].

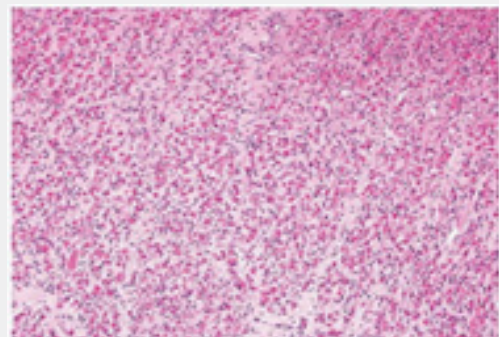


Figure 3: Fibromatosis colli depicting fascicles of fibroblasts and myo-fibroblasts admixed with a collagenous stroma and extravasation of numerous red blood cells [10].

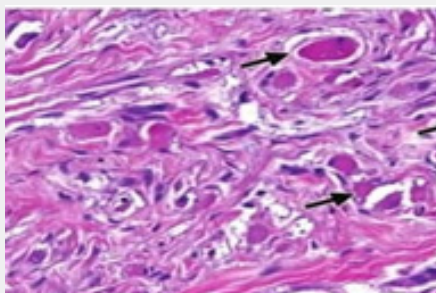


Figure 4: Fibromatosis colli delineating aggregates of fibroblasts and myo-fibroblasts commingled with several mature myocytes and circumscribing collagenous stroma [11].

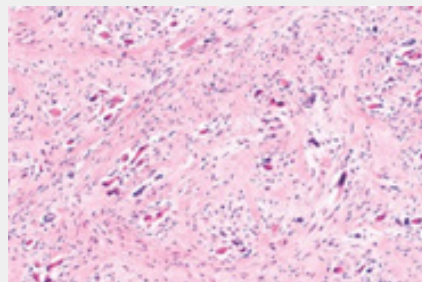


Figure 8: Fibromatosis colli enunciating fibroblasts, myo-fibroblasts and collagenous stroma with dispersed mature myocytes [14].

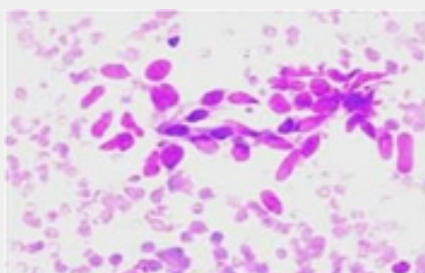


Figure 5: Fine needle aspiration cytology of fibromatosis colli exhibiting aggregates of spindle-shaped cells, plump fibroblasts, fibro-connective tissue fragments and a clear background with minimal red cell extravasation [12].

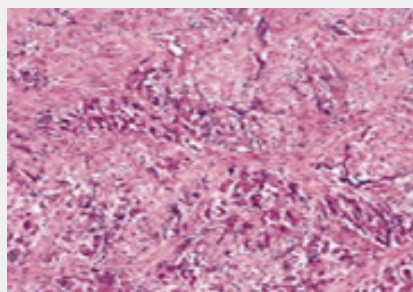


Figure 6: Fibromatosis colli depicting clusters of spindle-shaped cells, plump fibroblasts and commingled mature skeletal muscle fibres with collagen-rich stroma [13].

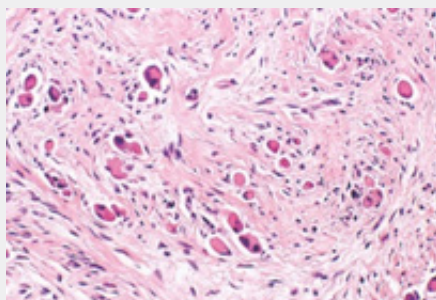


Figure 7: Fibromatosis colli exhibiting fibroblasts, myo-fibroblasts and an abundant collagenous stroma with commixture of enlarged myocytes [14].

Differential Diagnosis

Fibromatosis colli requires a segregation from congenital conditions such as enlarged cervical lymph nodes (possibly reactive with follicular hyperplasia), cervical abscess, bronchogenic cyst, dislocation of thyroid lobe, branchial cyst, thyroglossal duct cyst, inflammatory disorders as granulomatous, tubercular lymphadenitis. Neoplastic conditions as the benign haemangioma or cystic hygroma besides malignant neoplasia as neuroblastoma, rhabdomyosarcoma, lymphoma, teratoma and fibrosarcoma also necessitate a demarcation. Vascular and lymphatic malformations such as haemangioma or lymphangioma mandate evaluation besides bronchogenic cyst and thyroglossal duct cyst. Neck abscess is infrequent in neonates although can be discerned in older children or when accompanied by immunodeficiency [3,4].

Investigative Assay

On radiographic imaging, a uniform, isoechoic mass is detected, confined to the muscle. Fine needle aspiration cytology (FNAC) can be successfully adopted for diagnostic confirmation and segregation from associated congenital, inflammatory or neoplastic disorders, emerging at identical sites and within identical age group. The procedure is contemplated as a first line investigative modality for detecting infantile fibromatosis colli. It is a rapid, cost-effective, reliable procedure which can appropriately exclude adjunctive, benign or malignant supraclavicular or cervical nodules [3,4]. Extensive clinical examination can satisfactorily diagnose the condition. Ultrasonography is a preferential diagnostic tool on account of accessibility, cost effectiveness and lack of exposure to ionizing radiation. A competent ultrasound is beneficial and can eliminate associated aetiologies in instances where clinical aspect is atypical [7]. Ultrasonography delineates factors associated with congenital torticollis and lateral, cervical masses besides monitoring tumour development. Ultrasound scan of congenital fibromatosis colli typically displays a fusiform thickening of sternocleidomastoid muscle. Characteristically, fibromatosis colli demonstrates a fusiform or diffuse enlargement of sternocleidomastoid muscle with consequent muscle

shortening and head tilting towards the opposite side. Tumour echogenicity is variable. Colour Doppler can display an enhanced resistance waveform. Computerized tomography (CT) and magnetic resonance imaging (MRI) are advantageous in discerning the neoplasm and enunciate thickening of sternocleidomastoid muscle. Cross sectional computerized tomography (CT) or magnetic resonance imaging (MRI) can be occasionally employed to eliminate adjunctive conditions, especially with equivocal or atypical clinical features. Also, the modalities are beneficial for further characterization of disease and assessing extent of muscular involvement [7,8]. Computerized tomography (CT) typically demonstrates an enlarged sternocleidomastoid muscle which is iso-attenuating from normal, neighbouring musculature. Planes of adjacent adipose tissue are preserved. Calcification may be detected [8]. Magnetic resonance imaging (MRI) displays a well-defined mass with an enhanced signal intensity upon T1 weighted and T2 weighted imaging circumscribed by tissue with normal signal intensity and muscle [8].

Therapeutic Options

In the absence of adequate treatment, spontaneous retrogression of the neoplasm can ensue in 4 months or 6 months which can be facilitated or accelerated by appropriate physiotherapy. Surgical treatment is infrequent and adopted in beneath < 5% instances with antecedent diagnosis. Preliminarily discerned tumefaction can be appropriately managed with non-surgical techniques. Conservative management can be suitably adopted, contingent to appropriate diagnosis, thus circumventing superfluous surgical intervention [7,8]. Majority (90%) of instances are suitably alleviated with conservative management. Cogent therapies comprise of passive stretching of incriminated muscle and physiotherapy. An estimated 70% infants demonstrate complete resolution of the mass with normal cervico-fascial posture and mobility. Surgical intervention is beneficial in around

10% to 15% instances, especially in refractory neoplasms or incriminated children exceeding > one year, wherein tenotomy or tenomyotomy is a principally adopted procedure. Delayed diagnosis and therapeutic intervention, when occurring in children beyond > one year, is accompanied by inferior outcomes [7,8].

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9. Image 1 Courtesy: Webpathology.com
10. Image 2 and 3 Courtesy: Basic Medical Key.
11. Image 4 Courtesy: Research Gate
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13. Image 6 Courtesy: Pathology outlines.
14. Image 7 and 8 Courtesy: Paediatric and Orthopaedic Pathology.



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DOI: [10.19080/OAJS.2020.12.555829](https://doi.org/10.19080/OAJS.2020.12.555829)

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