## REVIEW

1457  Nonalcoholic fatty liver disease shows significant sex dimorphism  
*Chen XY, Wang C, Huang YZ, Zhang LL*

## MINIREVIEWS

1473  Management of procedural pain in the intensive care unit  

## ORIGINAL ARTICLE

### Clinical and Translational Research

1485  Effect of prior malignancy on the prognosis of gastric cancer and somatic mutation  
*Yin X, He XK, Wu LY, Yan SX*

### Retrospective Cohort Study

1498  Elemene-containing hyperthermic intraperitoneal chemotherapy combined with chemotherapy for elderly patients with peritoneal metastatic advanced gastric cancer  
*Chen ZX, Li J, Liu WB, Zhang SR, Sun H*

### Retrospective Study

1508  Timing theory continuous nursing, resistance training: Rehabilitation and mental health of caregivers and stroke patients with traumatic fractures  
*Shen YL, Zhang ZQ, Zhu LJ, Liu JH*

1517  Effect of precise nursing service mode on postoperative urinary incontinence prevention in patients with prostate disease  
*Zheng XC, Luo TT, Cao DD, Cai WZ*

1527  Significance of serum glucagon-like peptide-1 and matrix Gla protein levels in patients with diabetes and osteoporosis  
*Xie FF, Zhang YF, Hu YF, Xie YY, Wang XY, Wang SZ, Xie BQ*

1536  Castleman disease and TAFRO syndrome: To improve the diagnostic consciousness is the key  
*Zhou QY*

### Observational Study

1548  Correlation of myopia onset and progression with corneal biomechanical parameters in children  
*Lu LL, Hu XJ, Yang Y, Xu S, Yang SY, Zhang CY, Zhao QY*
<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
</tr>
</thead>
<tbody>
<tr>
<td>1572</td>
<td>Giant nodular fasciitis originating from the humeral periosteum: A case report</td>
<td>Yu SL, Sun PL, Li J, Jia M, Gao HW</td>
</tr>
<tr>
<td>1580</td>
<td>Tumor-related cytokine release syndrome in a treatment-naïve patient with lung adenocarcinoma: A case report</td>
<td>Deng PB, Jiang J, Hu CP, Cao LM, Li M</td>
</tr>
<tr>
<td>1586</td>
<td>Submucosal protuberance caused by a fish bone in the absence of preoperative positive signs: A case report</td>
<td>Du WW, Huang T, Yang GD, Zhang J, Chen J, Wang YB</td>
</tr>
<tr>
<td>1592</td>
<td>Misdiagnosis of unroofed coronary sinus syndrome as an ostium primum atrial septal defect by echocardiography: A case report</td>
<td>Chen JL, Yu CG, Wang DJ, Chen HB</td>
</tr>
<tr>
<td>1602</td>
<td>Treatment of extracranial internal carotid artery dissecting aneurysm with SUPERA stent implantation: Two case reports</td>
<td>Qiu MJ, Zhang BR, Song SJ</td>
</tr>
<tr>
<td>1609</td>
<td>Combination of atezolizumab and chidamide to maintain long-term remission in refractory metastatic extranodal natural killer/T-cell lymphoma: A case report</td>
<td>Wang J, Gao YS, Xu K, Li XD</td>
</tr>
<tr>
<td>1623</td>
<td>Primary orbital monophasic synovial sarcoma with calcification: A case report</td>
<td>Ren MY, Li J, Li RM, Wu YX, Han RJ, Zhang C</td>
</tr>
<tr>
<td>1639</td>
<td>Disseminated peritoneal leiomyomatosis with malignant transformation involving right ureter: A case report</td>
<td>Wen CY, Lee HS, Lin JT, Yu CC</td>
</tr>
</tbody>
</table>
Contents

Arthroscopic surgery for synovial chondroma of the subacromial bursa with non-traumatic shoulder subluxation complications: Two case reports
Tang XF, Qin YG, Shen XY, Chen B, Li YZ

Wilkie's syndrome as a cause of anxiety-depressive disorder: A case report and review of literature

Gastric schwannoma misdiagnosed as gastrointestinal stromal tumor by ultrasonography before surgery: A case report
Li QQ, Liu D

Giant retroperitoneal lipoma presenting with abdominal distention: A case report and review of the literature
Chen ZY, Chen XJ, Yu Q, Fan QB

Pneumothorax during retroperitoneal laparoscopic partial nephrectomy in a lupus nephritis patient: A case report
Zhao Y, Xue XQ, Xia D, Xu WF, Liu GH, Xie Y, Ji ZG

Bulbar conjunctival vascular lesion combined with spontaneous retrobulbar hematoma: A case report
Lei JY, Wang H

Hepatitis B virus in cerebrospinal fluid of a patient with purulent bacterial meningitis detected by multiplex-PCR: A case report
Gao DQ, Hu YQ, Wang X, Zhang YZ

Aseptic abscess in the abdominal wall accompanied by monoclonal gammopathy simulating the local recurrence of rectal cancer: A case report
Yu Y, Feng YD, Zhang C, Li R, Tian DA, Huang HJ

Tacrolimus treatment for relapsing-remitting chronic inflammatory demyelinating polyradiculoneuropathy: Two case reports
Zhu WJ, Da YW, Chen H, Xu M, Lu Y, Di L, Duo JY

Vedolizumab-associated diffuse interstitial lung disease in patients with ulcerative colitis: A case report
Zhang J, Liu MH, Gao X, Dong C, Li YX

Unusual magnetic resonance imaging findings of brain and leptomeningeal metastasis in lung adenocarcinoma: A case report
Li N, Wang YJ, Zhu FM, Deng ST

Diffuse invasive signet ring cell carcinoma in total colorectum caused by ulcerative colitis: A case report and review of literature
Zhang Z, Yu PF, Gu GL, Zhang YH, Wang YM, Dong ZW, Yang HR

Neurothekeoma located in the hallux and axilla: Two case reports
Huang WY, Zhang YQ, Yang XH
| 1747 | Subclavian artery stenting via bilateral radial artery access: Four case reports 

Qiu T, Fu SQ, Deng XY, Chen M, Dai XY |
ABOUT COVER
Editorial Board Member of World Journal of Clinical Cases, Prashanth Panta, MDS, Reader (Associate Professor), Department of Oral Medicine and Radiology, Malla Reddy Institute of Dental Sciences, Suraram 500055, Telangana, India. maithreya.prashanth@gmail.com

AIMS AND SCOPE
The primary aim of World Journal of Clinical Cases (WJCC, World J Clin Cases) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

INDEXING/ABSTRACTING
The WJCC is now indexed in Science Citation Index Expanded (also known as SciSearch®), Journal Citation Reports/Science Edition, Scopus, PubMed, and PubMed Central. The 2021 Edition of Journal Citation Reports® cites the 2020 impact factor (IF) for WJCC as 1.337; IF without journal self cites: 1.301; 5-year IF: 1.742; Journal Citation Indicator: 0.33; Ranking: 119 among 169 journals in medicine, general and internal; and Quartile category: Q3. The WJCC’s CiteScore for 2020 is 0.8 and Scopus CiteScore rank 2020: General Medicine is 493/793.

RESPONSIBLE EDITORS FOR THIS ISSUE
Production Editor: Lin-Yu Tong Wang; Production Department Director: Xiang Li; Editorial Office Director: Jin-Lei Wang.

NAME OF JOURNAL
World Journal of Clinical Cases

ISSN
ISSN 2307-8960 (online)

LAUNCH DATE
April 16, 2013

FREQUENCY
Thrice Monthly

EDITORS-IN-CHIEF
Bao-Gan Peng, Jerzy Tadeusz Chudek, George Kontogeorgos, Maurizio Serati, Ja Hyeon Ku

EDITORIAL BOARD MEMBERS
https://www.wjgnet.com/2307-8960/editorialboard.htm

PUBLICATION DATE
February 16, 2022

COPYRIGHT
© 2022 Baishideng Publishing Group Inc

INSTRUCTIONS TO AUTHORS
https://www.wjgnet.com/bpg/gerinfo/204

GUIDELINES FOR ETHICS DOCUMENTS
https://www.wjgnet.com/bpg/GerInfo/287

GUIDELINES FOR NON-NATIVE SPEAKERS OF ENGLISH
https://www.wjgnet.com/bpg/gerinfo/240

PUBLICATION ETHICS
https://www.wjgnet.com/bpg/GerInfo/288

PUBLICATION MISCONDUCT
https://www.wjgnet.com/bpg/gerinfo/208

ARTICLE PROCESSING CHARGE
https://www.wjgnet.com/bpg/gerinfo/242

STEPS FOR SUBMITTING MANUSCRIPTS
https://www.wjgnet.com/bpg/gerinfo/239

ONLINE SUBMISSION
https://www.ffpublishing.com
Hemangioma in the lower labial vestibule of an eleven-year-old girl: A case report

Ashwag Yagoub Aloyouny, Afrah Jaber Alfaifi, Shahad Mohammed Aladhyani, Ahad Ali Alshalan, Hadeel Mohammed Alfayadh, Hend Mahmoud Salem

ORCID number: Ashwag Yagoub Aloyouny 0000-0001-6759-2846; Afrah Jaber Alfaifi 0000-0002-9154-3806; Shahad Mohammed Aladhyani 0000-0003-1339-2747; Ahad Ali Alshalan 0000-0001-9934-4070; Hadeel Mohammed Alfayadh 0000-0003-1690-8470; Hend Mahmoud Salem 0000-0003-1126-80056.

Author contributions: Aloyouny AY contributed to data collection, reviewed the literature, interpreted the data, manuscript drafting and revision; Salem H prepared the specimen in the pathology laboratory and analysed it under the microscope; Alfaifi AJ, Aladhyani SM, Alshalan AA, and Alfayadh HM contributed to data collection, and manuscript drafting.

Informed consent statement: Informed written consent was obtained from the patient for publication of this case report and accompanying images.

Conflict-of-interest statement: The authors declare that they have no conflicts of interest.

CARE Checklist (2016) statement: The authors have read the CARE Checklist (2016) and have prepared and revised the manuscript accordingly.

Ashwag Yagoub Aloyouny, Afrah Jaber Alfaifi, Hend Mahmoud Salem, Department of Basic Dental Science, Princess Nourah Bint Abdulrahman University, Riyadh 11671, Saudi Arabia

Shahad Mohammed Aladhyani, Oral Medicine Special Care Dentistry, Prince Sultan Military Medical City, Riyadh 12211, Saudi Arabia

Ahad Ali Alshalan, Hadeel Mohammed Alfayadh, Department of Dentistry, Ministry of Health, Riyadh 11176, Saudi Arabia

Corresponding author: Ashwag Yagoub Aloyouny, DDS, Doctor, Department of Basic Dental Science, Princess Nourah Bint Abdulrahman University, King Khalid International Rd, Riyadh 11671, Saudi Arabia. ayaloyouny@pnu.edu.sa

Abstract

BACKGROUND
Hemangioma is a vascular benign tumour of endothelial origin. It appears commonly in the first decade of life with increases incidence in females. Hemangioma is not common to happen in the oral cavity and it is extremely rare to appear in the labial vestibule.

CASE SUMMARY
We present a case of an 11-year-old girl who complained of a painful, slowly growing mass which was consistent with the capillary hemangioma in the left mandibular vestibule. Vascular tumor such as hemangioma in the mandibular vestibule is extremely rare; hence, the clinical definitive diagnosis is very challenging. Therefore, radiographic imaging and histopathologic analysis are crucial to reach to the final diagnosis for proper management.

CONCLUSION
Comprehensive clinical evaluation, proper diagnostic imaging and microscopic analysis of the mass establish a precise diagnosis of the hemangioma for better management.

Key Words: Capillary hemangioma; Vascular malformation; Labial vestibule; Childhood; Case report

©The Author(s) 2021. Published by Baishideng Publishing Group Inc. All rights reserved.
Core Tip: Although hemangioma rarely occurs in the oral cavity, it should be considered in the diagnosis of a red-bluish isolated mass. In this case report, the patient presented with a painful, slowly growing mass in the left labial vestibule which resulted in asymmetry and swelling of the lower lip. The final diagnosis of the mass was consistent with capillary hemangioma in the mandibular vestibule. Early detection and treatment of oral masses is essential to avoid any complications.

INTRODUCTION

Hemangioma commonly appears early in life with increases incidence in females more than males. It usually gets smaller with time until it completely disappears. Mulliken et al. [1982] published the novel classification of vascular lesion and the International Society for the Study of Vascular Anomalies (2018) has provided updated guidelines for vascular anomalies classification. Accordingly, the vascular anomalies have been classified into vascular malformations and vascular tumors. Vascular tumors include non-harmful, locally destructive, and malignant lesions, whereas vascular malformations include simple and combined malformations.

Hemangiomas are vascular benign tumours of endothelial origin, presenting clinically with varying sizes and shapes. In some cases, hemangioma could cause functional disability and disfiguring appearance, which may lead to psychological issues. Histologically, hemangioma has another classification, into either cavernous or capillary types.

Hemangioma is uncommon to occur in the oral cavity and it is extremely rare to appear in the labial vestibule. To our knowledge and based on the review of English literature (PubMed), this case is the first report of capillary hemangioma in the mandibular vestibule. In this report, we present an 11-year-old girl patient complaining of a slowly growing mass that caused labial swelling and asymmetry. The diagnosis was consistent with capillary hemangioma in the mandibular vestibule.

CASE PRESENTATION

Chief complaints

An 11-year-old healthy girl was referred to the oral and maxillofacial surgery clinic for evaluation of labial asymmetry and swelling.

History of present illness

The patient had a two-month-history of a slowly growing lesion in the left side of the lower labial vestibule accompanied with persistent mild pain. The patient and her parents reported no history of trauma at the site of the mass.

History of past illness

The patient was healthy and did not undergo any surgeries.

Personal and family history

The parents revealed no significant family history and no genetic abnormalities.

Physical examination

Physical and systemic examination: The patient was healthy and had only taken Tylenol 15 mg as needed for fever.

Extraoral examination: Extraoral examination showed lower lip asymmetry and swelling in the left side.
**Intraoral examination:** Intraoral examination showed a 2.0-1.5 cm, solitary, fluctuant, bluish, smooth, palpable submucosal mass, rubbery in consistency, tender, and blanch on pressure (positive diascopy test). The submucosal mass located in the left mandibular vestibule opposite to tooth number 32 and 33 (23 and 33, according to the FDI World Dental Federation Notation) (Figure 1).

**Imaging examinations**
A panoramic radiograph showed normal structures with no significant pathologic findings. Additionally, Color-Doppler-ultrasound was performed to confirm the nature of the lesion. The imaging interpretation revealed a slow-flow vascular lesion in the left lower vestibule and attached to the lower orbicularis oris muscle.

**FINAL DIAGNOSIS**
The final diagnosis was based on histopathological analysis. The histopathological diagnosis was consistent with capillary hemangioma in the labial vestibule (Figure 2).

**TREATMENT**
Complete surgical removal of the lesion to reduce the risk of the recurrence.

**OUTCOME AND FOLLOW-UP**
At two-, four- and eight-week-follow up, the site of the surgery healed well with no sign of bleeding, infection, and swelling. At one- and three-year- follow up, there was no recurrence of the lesion or complications noted. Additionally, the patient was in a good health.

**DISCUSSION**
In 1982, vascular anomalies were categorised into two main categories: Vascular malformations and vascular tumors. Hemangiomas are true neoplasms represented by increased rate and proliferation of endothelial cell turnover. On the other hand, vascular malformations are localised abnormality and disorganisation of the blood vessel caused by defects in vascular development [1-3]. Simple vascular malformations are classified histologically, based on the vessel size, into capillary, venous, lymphatics, arteriovenous fistula, and arteriovenous malformations. Vascular lesions are further categorised into non-harmful, locally destructive, and malignant lesions [4]. Namely, hemangioma is a neoplasm of endothelial origin which is commonly found in the early years of life and then the neoplasm regresses gradually with age [5]. Intraoral and intramuscular hemangiomas are rare, dissimilar to cutaneous and subcutaneous hemangiomas. Oral hemangiomas could occur in more than 6% of infants and have high prevalence in female presenting 3:1 (female: male). Infants are more likely to develop oral hemangiomas if they fall in one of the following conditions; infants who are born to older mothers, twins or triplets, premature, or have low birth weight [6]. Hemangioma is a common vascular benign tumor which falls under the category of benign vascular tumors and it is further divided into capillary and cavernous hemangioma [2].

Capillary hemangioma is a common lesion, but it rarely occurs in the oral cavity. According to Matsumoto et al [7], 45.2% of capillary hemangiomas occur on buccal mucosa, 35.5% on the tongue, and only a small percentage occur in the lip, gingiva and palate. Capillary hemangiomas are firm in consistency and have a limited history of symptoms. Although the exact cause of oral hemangioma is not fully understood, hormonal changes, embolic phenomenon and genetic mutations are believed to play an important role in the tumor development [8].

Hemangiomas are hypothesised to develop because of both angiogenesis and vasculogenesis through three different stages, as follows: Endothelial cell proliferation stage, rapid growth stage and spontaneous disappearance. Endothelial cell proliferation is stimulated by many factors, such as basic fibroblast growth factor, vascular...
endothelial growth factor and transforming growth factor-beta. Then, the quantity of endothelial cells is sustained, and each cell increases in size, leading to comprehensive enlargement of the structure size. At the end, spontaneous involution occurs when the endothelial cells are replaced by connective tissue, adipose, and fibroblast, and the number of small vessels decrease in quantity[9].

Hemangioma could be classified clinically as congenital or infantile (previously named strawberry or juvenile). Congenital hemangioma presents at birth and does not demonstrate proliferation stage. In contrast, infantile hemangioma may develop at the first months of the infant life and show a proliferative phase during the period of six to twelve months; then, most cases spontaneously regress between the age of six to nine years. High percentage of hemangiomas disappear completely in childhood, with < 20% carrying on to puberty[10,11]. Oral hemangioma presents as a solitary, soft, fluctuant, compressible, smooth, red, or bluish submucosal mass. Significant variations may present based on the depth and site of the mass. Superficial masses are easy to visualise and may present as pedunculated, sessile, or lobulated and reddish in colour. In contrast to deeper masses, they appear as a dark blue discolouration recognisable from surrounding normal colour mucosa. It also reveals tenderness on palpation and blanch on compression with glass slide (positive diascopy test)[12]. In this case, the differential diagnosis of the tumor was written down as vascular anomalies, including hemangioma, and vascular malformation, including venous, capillary, lymphatic and arterial malformations. Salivary gland tumor, mucocele and angioleiomyoma were also considered.
It is worth mentioning that vascular malformations, salivary gland tumor, mucocele and angioleiomyoma were all excluded because the lesion showed a slow-flow vascular lesion by using Colour-Doppler-ultrasound, which is highly consistent with hemangioma.

Histological analysis of vascular anomalies, including capillary hemangioma, is still the most acceptable and accurate method of diagnosis[3]. Microscopic description of capillary hemangioma illustrates several dilated capillaries lined by endothelial cells, filled with blood, and surrounded by inflammatory infiltrate.

Hemangioma is mostly characterised by its benign feature and has high tendency of involution by itself over time. However, sometimes hemangioma requires intervention, especially in case of impairment in breathing, swallowing and speech. The first line of evaluation and diagnosis would be by Color-Doppler ultrasound imaging. This imaging modality is non-invasive, cost-effective and has no risk of radiation. If intraosseous lesion is anticipated, other imaging modalities could be useful for the diagnosis, such as a contrast-enhanced magnetic resonance imaging (MRI), computed tomography (CT), and angiography[13]. A contrast-enhanced MRI and CT imaging identify the shape, size, and calcification of the tumor. The Color-Doppler ultrasound imaging modality was the suitable choice for the patient due to many factors, such as financial issues and the age of the patient[14].

Choosing a suitable method for managing hemangioma is based on multiple factors such as the aesthetic consideration, clinical nature, size, site, growth rate, accessibility, extent of the tumor, and age of the patient. Hemangioma could be managed by different ways; for instance, surgical excision of the tumor, embolization, electro-surgery, cryosurgery, laser, steroid injection, or sclerosing materials. In case of small oral hemangioma, the commonly used method is the total surgical excision of the whole mass to decrease the potential risk of recurrence[15]. However, if the lesion is large and located in a significant part of the mouth, such as the tongue, in this case the surgical excision of the lesion would not be preferred, so as to avoid post-surgical complications in swallowing and speech. Sclerotherapy is recommended to manage large hemangiomas in the oral cavity in which 3% sodium tetradecyl sulfate or ethanolamine olate is injected into the main vessels of the lesion to destroy the endothelial cells, leading to lesion destruction. Precautionary measures should be taken to avoid bleeding during the surgical procedure or afterward during recovery phase. In the present case, the tumor was excised surgically with a thin rim of the attached orbicularis oris muscle to decrease the risk of recurrence[16,17].

CONCLUSION

Although hemangioma rarely appears in the oral cavity, it should be considered in the diagnosis of a red-blush isolated mass. Comprehensive clinical evaluation, proper diagnostic imaging and microscopic analysis of the mass establish a precise diagnosis of the hemangioma for better treatment.

REFERENCES

3 Larsen AK, Damsgaard TE, Hedelund L. [Classification of vascular anomalies]. Ugeskr Laeger 2018; 180 [PMID: 30187855]


