Results of Surgical treatment for Clavicle Fractures in Vietnamese Adults

Published On: November 24, 2017 | Pages: 023 - 025

Author(s): Dung Trung Tran*, Khanh Trinh Le, Ban Hoang Van, Tung Pham Son, Minh Ho Ngoc, Hanh Tran Thi My and Dat Tran Tien

Objectives: To describe the pattern of clavicle fractures and to evaluate the results of surgical treatment for clavicle fractures. ...

Nuances of Microsurgical and Endoscope Assisted Surgical Techniques to the Cranio-Vertebral Junction: Review of the Literature

Published On: February 06, 2017 | Pages: 001 - 008

Author(s): Visocchi Massimiliano*, Signorelli Francesco, Iacopino Gerardo and Barbagallo Giuseppe,

Purpose: An update of the technical nuances of microsurgical - endoscopic assisted approaches to the craniocervical junction (transnasal, transoral and transcervical) if provided from the literature in order to better contribute to identify the best strategy. ...

Congenital pseudarthrosis of the clavicle in a 7-year-old girl: A case report
Congenital pseudarthrosis of the clavicle is a rare clinical entity. The literature reports to date only 200 clinical cases in the world. ...

Familial Mediterranean Fever (FMF) is an autoimmune disease caused by mutations in the MEFV gene and characterized by recurrent fever, polyserositis and arthritis. It is transmitted in an autosomal recessive pattern. ...

Shoulder dislocations are the most common joint dislocations, and only 2% of these are seen as posterior shoulder dislocations. The floating elbow was first described in children, after that shown in adults. Floating elbow cases are very rare, and usually seen with high-energy trauma. Classical definition is the coexistence of the humeral diaphyseal and forearm ...

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Abstract View | Full Article View | DOI: 10.17352/ojor.000010

Abstract View | Full Article View | DOI: 10.17352/ojor.000009
Camptodactyly Arthropathy CoxaVara Pericarditis Syndrome: Early diagnosis prevents unnecessary and harmful treatment

Published On: May 29, 2017 | Pages: 012 - 014

Author(s): Sheren Esam Maher*, Mohamed Hashem M Mahgoob and Nadia El Ameen

Camptodactyly-arthropathy-coxa vara-pericarditis (CACP) syndrome is a genetic disorder caused by mutation in the Proteoglycan PRG4 gene on chromosome 1. The syndrome is characterized by congenital or early onset camptodactyly and childhood-onset of non-inflammatory arthropathy, coxa vara deformity, or other dysplasia associated with progressive hip disease and non-i ...

Muscle Vanishing in Poliomyelitis Manifested on F-18 FDG PET/CT: An Interesting Imaging Finding

Published On: March 27, 2017 | Pages: 009 - 011

Author(s): Rong-Hsin Yang and Yum-Kung Chu*

Muscle atrophy is the loss of muscles bulk; it can be a partial or complete wasting away of muscle. Herein, we describe a 56-year-old man with diffuse large B-cell lymphoma underwent F-18 FDG PET for postchemotherapy evaluation. ...