Nilai Akhir

LEMBAR HASIL PENILAIAN SEJAWAT SEBIDANG ATAU PEER REVIEW KARYA ILMIAH: JURNAL ILMIAH

Judul Artikel Ilmiah Penulis Artikel Ilmiah : Pneumorrhachis and Hyponatremia After a Neck Hack - A Case Report

: 5 orang

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Status Pengusul Identitas Jurnal Ilmiah

: International Journal of Surgery Case Reports : a. Nama Jurnal

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h. Terindeks di

: SQOPUS (Q3) SJR 0,232

i. Link turnitin

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d. Kelengkapan unsur dan kualitas

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		40			·	
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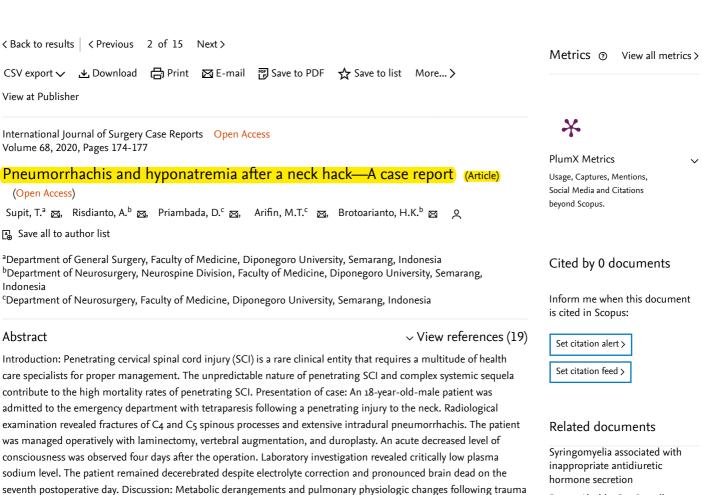
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are lethal complications. Hyponatremic encephalopathy and disrupted pulmonary function caused by high cervical

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report presents a rare clinical entity along with its' complications. Prompt clinical stabilization, strict biochemical

monitoring, and multidisciplinary care from health care specialists are mandatory for SCI patients. © 2020 The

Prominence percentile: 74.884

Author keywords

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 Cervical
 Penetrating
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EMTREE drug terms: (antibiotic agent) (sodium)

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Barros Alcalde, P., González Quintela, A., Pena Seijo, M. (2014) BMJ Case Reports

A case of hyponatremia after cervical spinal cord injury

Kageyama, K. , Suda, T. *(2011) Endocrine Journal*

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Treatment of an intraarticular comminuted fracture of the base of the proximal phalanx in a ring finger using the Ichi-Fixator external fixator system: A case report



Akira Hara a,b,*, Minoru Yokoyama b, Satoshi Ichihara a, Yuichiro Maruyama a

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Keywords: Metacarpophalangeal joint External fixator Intraarticular fracture Comminuted fracture

ABSTRACT

INTRODUCTION: Comminuted fractures involving the articular surface of the base of the proximal phalanx are relatively rare. We treated a patient with this type of fracture by open reduction and internal fixation with a locked-wire-type external fixator (Ichi-Fixator System).

PRESENTATION OF CASE: A 45-year-old man was injured because his ring finger was kicked during a Futsal game. Radiographs and computed tomography revealed a comminuted intraarticular fracture of the proximal phalanx of this ring finger. We treated the fracture with open reduction and K-wires and external fixation. We removed the K-wire and external fixator 5 weeks postoperatively and initiated range of motion exercises. Five months postoperatively, his finger motion was fully recovered without restriction.

DISCUSSION: Comminuted intraarticular fractures of the base of the proximal phalanx are usually treated with plating. Complications such as interference with excursion of the central slip and lateral bands, extensor tendon rupture, and plate prominence have been reported in these fractures. In our patient, the Ichi-Fixator System was useful as a distraction apparatus for metacarpophalangeal joint fixation.

CONCLUSION: A comminuted intra-articular fracture of the base of the proximal phalanx was treated successfully using the Ichi-Fixator system.

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1. Introduction

Fractures of the lateral volar base of the proximal phalanx are common injuries and usually represent with collateral ligament avulsion injuries. In contrast, comminuted fractures involving the articular surface of the base of the proximal phalanx are relatively rare and usually represent as a volar base fracture with a central depression of the articular surface. Failure to reduce and secure the fracture leads to persistent subluxation, articular incongruity, and post-traumatic arthritis. These fractures are generally approached through a dorsal extensor-tendon-splitting incision [1] or volar A1 pulley approach to visualize the articular surface [2]. Most commonly, a significantly-sized palmar–ulnar or palmar–radial fragment exists, and fixation can be accomplished with a minicondylar plate or K-wires. We experienced a patient

with a comminuted intraarticular fracture of the base of the proximal phalanx, which we treated by open reduction and internal fixation with a locked-wire-type external fixator (Ichi-Fixator System (IFS); Neo-medical, Saitama, Japan) [3,4]. The work has been reported in line with the SCARE criteria [5].

2. Case report

A 45-year-old man presented to our hospital because of a ring finger injury. One day earlier, he played Futsal as a goalkeeper, when another player accidentally kicked the patient's ring finger while the patient was saving the ball. At presentation, his ring finger was swollen without a wound. Plain radiographs showed a comminuted intraarticular fracture of the base of his ring finger proximal phalanx (Fig. 1). Computed tomography revealed a comminuted fracture with articular depression of a fragment of the proximal phalangeal base (Fig. 2). We explained that closed reduction and conservative treatment would be failed. The patient chose operative treatment. Surgery was performed 10 days after his first visit. We approached the fracture site via dorsal extensor-tendon-

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The infected hematometra in a rudimentary noncommunicating horn misdiagnosed as pelvic mass: A case report



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ABSTRACT

INTRODUCTION: The rudimentary noncommunicating horn with a functional endometrial cavity is rare and often challenging to diagnose because of the variety in clinical features. We present a case of a patient for whom the diagnosis of a uterine horn was missed during the prior cesarean section, which later successfully treated with robotic-assisted laparoscopic removal of a rudimentary noncommunicating horn of uterus and ipsilateral tube.

PRESENTATION OF CASE: A 20-year old woman, gravida 3 para 2, presented with a complaint of acute and severe pelvic pain with fever. Multiple imaging modalities of pelvis and abdomen showed an 8 cm right-sided pelvic mass with a tubular structure adjacent to the uterus. The pelvic inflammatory disease was diagnosed and treated with intravenous antibiotics. After reviewing multiple radiology images, Müllerian anomaly was suspected, and the rudimentary horn with the fallopian tube was confirmed via diagnostic hysteroscopy and laparoscopy. Subsequently, robotic-assisted laparoscopic removal of the right horn with the fallopian tube was performed.

DISCUSSION: Assessment of a rudimentary noncommunicating horn with unicornuate uterus can be achieved by several radiology methods, including computed tomography, magnetic resonance imaging, two and 3-dimensional ultrasonography, hysterosalpingogram, and sonohysterography. In addition, evaluation of concomitant skeletal and renal anomalies is essential in enhancing diagnostic accuracy. In our case, the Müllerian anomaly with delayed onset complications was diagnosed by multiple imaging studies and treated successfully.

CONCLUSION: The early and correct diagnosis of the Müllerian anomaly remains difficult but essential as misdiagnosis can be associated with serious complications in patients.

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1. Introduction

Congenital abnormalities of the Müllerian ducts are relatively common and contribute to complications in obstetrics and gynecology. A unicornuate uterus is caused by a failure of development in one Müllerian duct and frequently present with a rudimentary horn [1]. Most rudimentary horns are asymptomatic when the endometrium is both non-functional and noncommunicating [2]. However, rudimentary horns with a functional endometrial cavity can cause chronic pain, dysmenorrhea, and endometriosis, and they are often difficult to diagnose by noninvasive means [3]. Therefore, proper preoperative diagnosis and surgical treatment are essential to prevent serious complications in the future as well as to provide symptomatic relief to patients. We present the case of a patient for whom the diagnosis of a uterine horn was missed during the

prior cesarean section, and we successfully treated with roboticassisted laparoscopic removal of a rudimentary noncommunicating horn and ipsilateral tube causing severe pelvic pain. The case report described here is in line with the SCARE criteria [4].

2. Presentation of case

A 20-year-old woman, gravida 3 para 2, presented to the emergency department with acute and severe pelvic pain with fever $38.3\,^{\circ}$ C ($101\,^{\circ}$ F). Her gynecologic and obstetrical history consisted of one cesarean section due to preterm prelabour rupture of membranes and increased fetal risk at 35 week-gestation in 2011. Her second pregnancy in 2013 was a successful vaginal birth after the cesarean section at 37 week-gestation. Then, the patient was treated for chlamydia infection about three months after her last delivery. She endorsed cyclic pelvic pain, particularly at the time of her regular menses. A patient denied any significant medical and family history.

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