COVID-19 and Myositis: CASE REPORTS

BS. NGUYỄN HÒ TRÚC LINH

BS. TRẦN CHÍ ĐỦ

BS. NGUYỄN THÀNH ĐĂNG

BS. LÊ ĐÌNH VĨNH PHÚC

PATHOLOGY:

Findings

19 is associated with a viral myositis attributable to direct myocyte invasion or induction o unity. COVID-19-induced myositis may be varied in presentation, from typical dermatomy omyolysis, and a paraspinal affliction with back pain. It may or may not present with acute tial elevations of enzyme markers such as creatine kinase (CK). Virus-mediated muscle ation is attributed to ACE2 (angiotensin-converting enzyme) receptor-mediated direct entry of muscle fibers, leading on to innate and adaptive immune activation. A greater recogniti similarity between anti-MDA5-positive myositis with COVID-19 has thrown researchers in exploration — finding common etiopathogenic basis as well as therapeutic strategies. For p blished myositis, chronic care was disrupted during the pandemic with several logistic cha ment dilemmas leading to high flare rates. Teleconsultation bridged the gap while ushering tient-led care with the digital transition to tools of remote disease assessment.

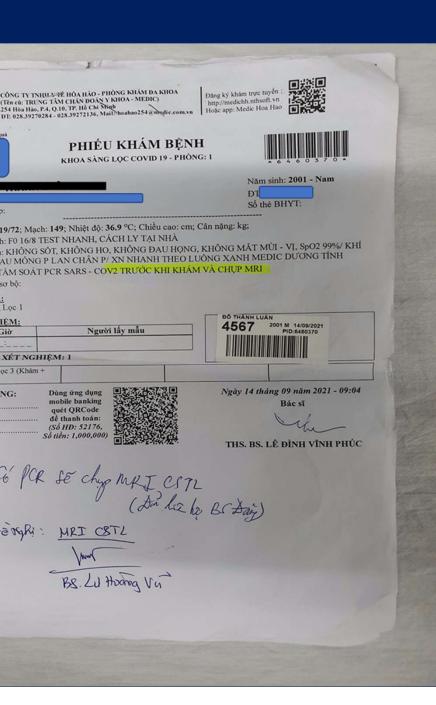
Alongside the typical respiratory manifestations like cough, fever, and sore throat, proximal muscle weakness is increasingly being reported as a manifestation causing significant morbidity in occasional COVID-19 patients. Myalgia is reported extensively in the literature as a common musculoskeletal manifestation of COVID-19 infection, presenting in nearly half of all COVID-19-infected patients. COVID-19-related myositis and consequently rhabdomyolysis are other reported manifestations, albeit relatively rare [7, 8]. By July 2020, a single case of COVID-19-related myositis had been reported in the literature [9]. Since then, several case reports and series reporting a virus-induced myositis attributed to COVID-19 disease have been published. The muscle involvement may vary from an asymptomatic elevation of CK to severe rhabdomyolysis.

Acute Viral Myositis

Nearly 23 patient cases of myositis attributable to COVID-19 have been described so far. COVID-19-induced myositis may vary in presentation, ranging from frank muscle weakness to typical dermatomyositis replete with classic rashes, or mere back pain with muscle disease on MRI. Most patients test positive for COVID-19 on initial presentation with most reported being males aged 33–87. Noteworthily, COVID-19 may or may not present with acute exponential elevations of enzyme markers such as CK, and muscle enzymes may not necessarily have a direct bearing on prognosis.

Rhabdomyolysis

Rhabdomyolysis is one of the rare and severe complications of COVID-19 infection which can be an initial presentation in some cases [10, 11]. In such cases, patients present with typical COVID-19 symptoms such as fever, cough, myalgia, and shortness of breath as well as manifest acute lower limb—dominant symmetric muscle weakness and subsequently go on to develop rhabdomyolysis associated with elevated CK levels. A study reported a peak CK value as high as 33,000 U/L [12•]. In this setting, they may present with frank muscle weakness, which is profound, proximal, lower limb—dominant, acute, and symmetric. At times, the patients are critically ill and requiring ventilatory support. The only manifestation of rhabdomyolysis in these may be myoglobinuria (dark urine) and acute kidney injury (AKI) needing hemodialysis with raised CK >5000 IU/L [13•, 14•, 15•]. Rhabdomyolysis can be fatal, with casualties reported in nearly 45% (4 of 9 reported) of those with this presentation over a short follow-up duration (Table 1). Among all reported cases (n = 23) of COVID-19-related myositis so far, 21.7% (n = 5) succumbed to the illness of which 80% (n=4) had rhabdomyolysis.



> 1ST CASE:

- A 20-year-old male patient, visited Hoa Hao on Sep 14th
- CHIEF COMPLAINT:
- On August 16th: The patient got covid from a housemate due to a quick test at home. Selfand treat at home with mild symptoms.
- On September 3rd, the patient had right bac spreading to the right buttock, went to a privious doctor and was given medicine but the symp did not decrease.
- On September 10th, the patient went to hose for traumatology and orthopaedics, did the following tests and treatment, but the symptodid not decrease.
- On September 14, the patient was assigned Medic center for an MRI.



BV CHẨN THƯƠNG CHÍNH HÌNH 929 TRẦN HƯNG ĐẠO-P1-Q5 Tel: 028-39235821

Số bệnh án: CH210901785 Số bệnh phẩm: 100921-5862

1/2

PHIẾU KẾT QUẢ XÉT NGHIỆM

Họ tên:		Năm sinh: 20	001 Giới tinh: Nam
Địa chỉ:			
BS chỉ định: Chẩn đoán:		Khoa: Viện Phí	Ngoài Giờ
Ngày giờ nhận mẫu: 10/09/2021	10:31:36		: 10/09/2021 11:04:18 h trang mẫu: Đạt

Yêu cầu xét nghiệm	Kết quả	Giá trị tham chiếu	Đơn vị	Tên máy	Mã PPXN
		Huyết học			
Tổng phân tích tế bào ma	áu ngoại vi bầ	ing máy đểm laser			
WBC	8.3	(5 - 10)	X10^9/L	Sysmex XN100	QTKT.HH.01
NEU%	72.5	(41 - 74)	%	Sysmex XN100	QTKT.HH.01
LYM%	16.5	(21 - 50)	%	Sysmex XN100	QTKT.HH.01
MONO%	9.4	(3 - 9)	%	Sysmex XN100	QTKT.HH.01
EOS%	1.6	(0 - 8.4)	%	Sysmex XN100	QTKT.HH.01
BASO%	0.0	(0 -1.5)	%	Sysmex XN100	QTKT.HH.01
NEU#	6.0		X10^9/L	Sysmex XN100	QTKT.HH.01
LYM#	1.4		X10^9/L	Sysmex XN100	QTKT.HH.01
MONO#	0.8		X10^9/L	Sysmex XN100	
EOS#	0.1		X10^9/L	Sysmex XN100	QTKT.HH.01
BASO#	0.0		X10^9/L	Sysmex XN100	QTKT.HH.01
RBC	4.88	Nữ(3.9-5.4);Nam(4.3-5.8)	X10^12/L	Sysmex XN100	QTKT.HH.01
HGB	13.4	Nữ(12.5-14.2)Nam(14-16)	g/dL	Sysmex XN100	QTKT.HH.01
HCT	41.7	Nữ(35-47);Nam(38-50)	%	Sysmex XN100	
MCV	85.5	Nữ(83-91);Nam(84-92)	fL	Sysmex XN100	
MCH	27.5	Nữ(27-31);Nam(28-32)	Pg	Sysmex XN100	
MCHC	32.1	Nữ(32-35);Nam(32-36)	g/dL	Sysmex XN100	
RDW	13.4	(9 - 15)	%	Sysmex XN100	
PLT	245.0	(150 - 400)	X10^9/L	Sysmex XN100	
MPV	9.2	(6.5-11)	fL	Sysmex XN100	
Máu lắng (máy tự động)		, , ,	9-27	pyomex xivio	Q I I I I I I I I I
ESR	120	(2 - 30)	mm/h	Roller 20	QTKT.HH.03
		Sinh hoá	10111011	Troiler 20	QTICT.TITL.00
Glucose	6.25	(3.9 - 6.4)	mmol/L	AU680	QTKT.HS.01
Creatinine	87	Nữ(44-100);Nam(62-120)	µmol/L	AU680	QTKT.HS.03
Jric Acid	385	Nữ(150-360);Nam(180-420)	µmol/L	AU680	QTKT.HS.03
AST (SGOT)	35.6	Nữ (< 31); Nam (< 37)	U/L	AU680	QTKT.HS.04
ALT (SGPT)	69.1	Nữ (< 31);Nam (< 40)	U/L	AU680	QTKT.HS.06
		Miễn dịch	U/L	1 40000	Q1K1.H5.06
RF	2.2	< 10	UI/mI	AU680	OTKTUS
CRP	134.2	< 6	mg/l	AU680 AU680	QTKT.HS.21
Shi chú:		1	mg/i	A0080	QTKT.HS.20

Thời gian trả kết quả:

Địa chỉ: Ấp Bình Châu Xã Tuyên Bình, Huyện Vĩnh Hưng, Long An Chẩn đoán: viêm khớp cúng chậu Phải Hướng xử trí: Xương Chậu thẳng(Chụp Xquang số hóa 1 phim)(1); Thuốc: 1, Celecoxib (Celofirm 200) Ngày uống 2 lần, mỗi lần 1 Viên(sau ăn) 28 Viên 2. Pregabalin 75mg (Prezel 75) 28 Viên Ngày uống 2 lần, mỗi lần 1 Viên(sau ăn) 3. Cefdinir 300mg (Topdinir 300) 28 Viên Ngày uống 2 lần, mỗi lần 1 Viên(sau ăn) 4. Tramadol 37.5mg+paracetamol 325mg (Paratramol

Ngày uống 2 lần, mỗi lần 1 viên(sau ăn) 5. Bromelain 50 F.I.P units (Bromanase) Ngày uống 2 lần, mỗi lần 1 viên(sau ăn)

viên

28 viên

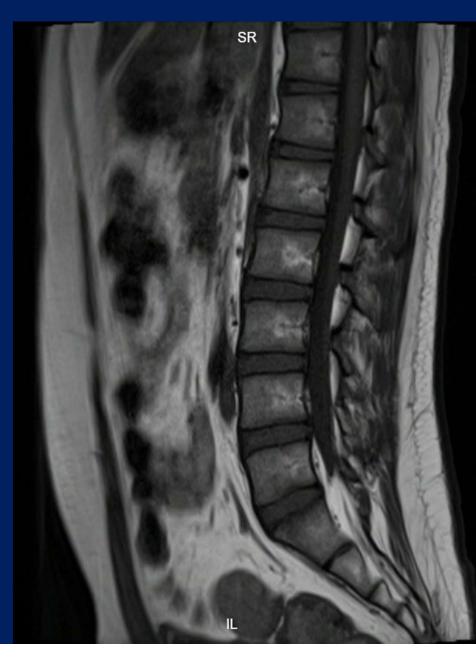
 ^{- 60} Phút cho các xét nghiệm thường qui (Sinh hóa, Huyết học...)
- 120 Phút cho các xét nghiệm thường qui + miễn dịch.

Dữ liệu xét nghiệm: Ngày 14/09/2021 (Đã đủ kết quả) TÊN XÉT NGHIỆM KÉT QUẢ KHOẢNG THAM CHIẾU MÃ QT I. MIEN DICH - IMMUNOLOGY ÂM TÍNH XN nhanh kháng nguyên SARS CoV-2 Dương tính test nhanh Chở kết quả khẳng định RT (Dịch ty hầu) PCR II. SINH HỌC PHẨN TỬ MOLLCULAR BIOLUGY Realtime PCR SARS-CoV-2 (mau don) Dwong tinh H Am TINH QTAD351 Ct 30 Đóng

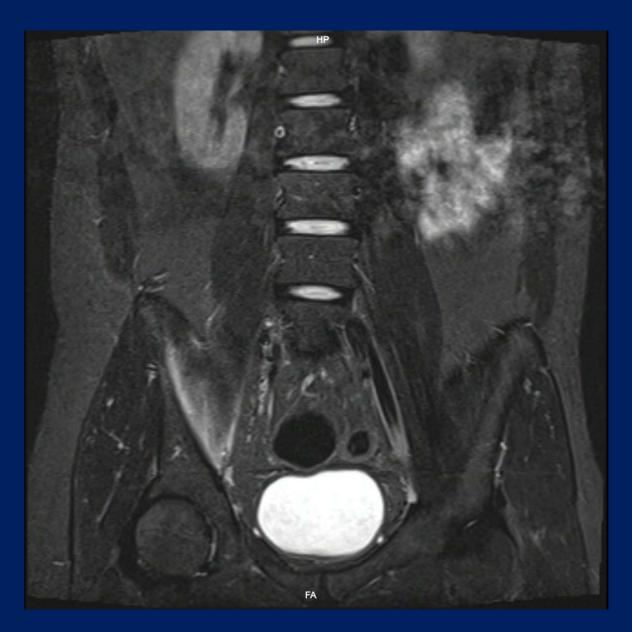
T2WI SAGITTAL

T1WI SAGITTAL

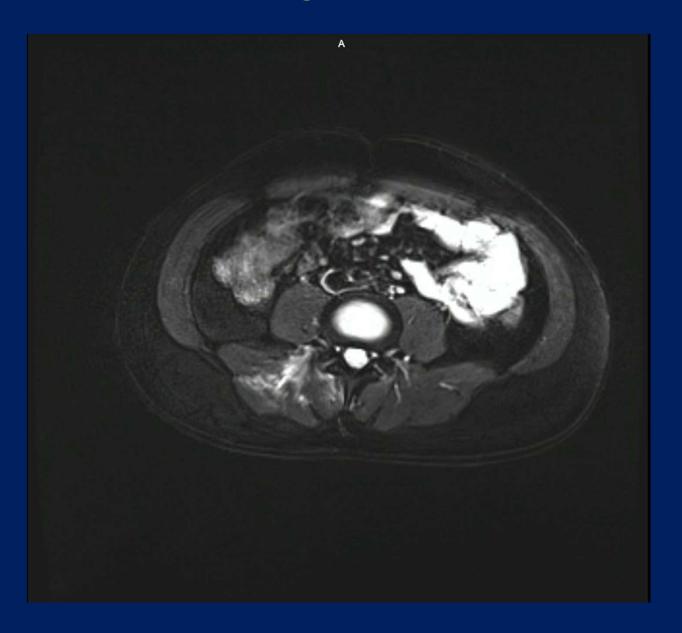




T2WI FATSAT CORONAL



T2WI FATSAT AXIAL





Bệnh viện : MEDIC Khoa : PK

LÝ DO KHÁM : đau lưng

Máy : SIEMEN ESSENZA 2

Vùng : MRI Cột Sống Thất Lưng Không tiêm tương phản

Kết quả : Cột sống thắt lưng được khảo sát qua các mặt cắt theo những chuỗi xung:

Sagittal T1WI.

Sagittal, Axial, Coronal T2WI.

Không tiêm thuốc tương phản.

*** MÔ TẢ:

- Không giảm tín hiệu đĩa đệm trên T2WI. Không giảm chiều cao thân sống và đĩa đệm.

Không ghi nhận lồi hay thóat vị đĩa đệm.

Đường kính trước sau đo ngang chân cung không hẹp.

Không hẹp lỗ liên hợp hai bên.

- Không dầy hay vôi hóa dây chẳng vàng và dọc sau.

Tùy sống ngực thấp, chóp cùng và các rễ chùm đuôi ngựa bình thường.

Không thấy u trong túi cùng màng cứng.

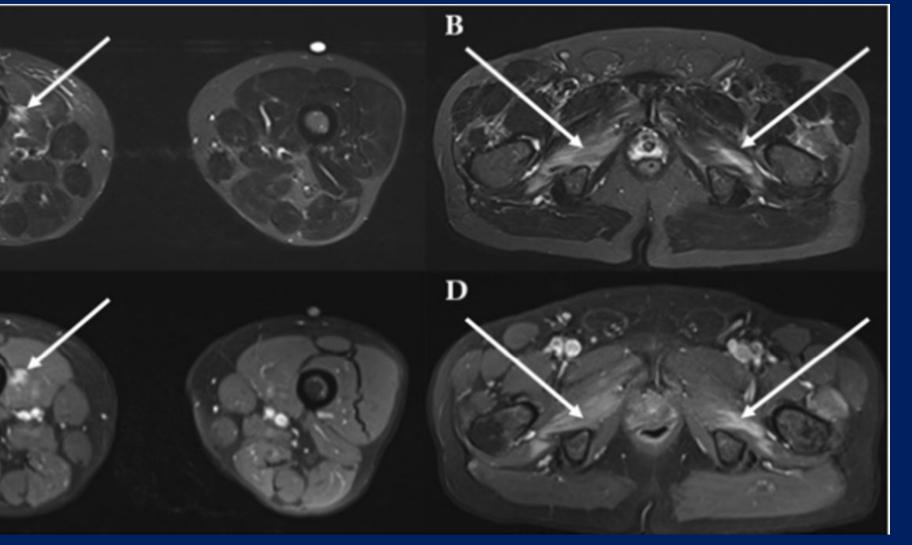
- Viêm khớp cùng chậu phải và cơ cạnh sống vùng thắt lưng phải.

*** KÉT LUÂN:

Không thấy thương tổn thân sống đĩa đệm cột sống thắt lưng.

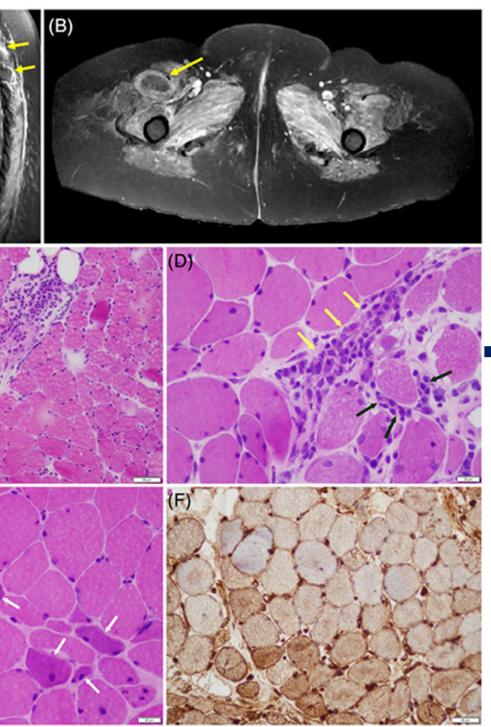
Viêm khớp cùng chậu phải và cơ cạnh sống vùng thắt lưng phải.

Tp. Hồ Chí Minh, ngày 14/09/2021 17:16



Symptoms suddenly or with diffuse and proximilimb weakness, him to fall. Out at the hosp patient was and did not any upper airway symptoms

and thigh MRI. (A) Thigh MRI in T2 STIR sequence showing oedema of the right vastus medialis (arrayic MRI in T2 STIR (short TI inversion recovery) sequence showing bilateral oedema of external obtus (arrows). (C and D) T1 sequences revealing enhancement of muscle lesions after gadolinium infus



A 58-year-old woman presented with cough, dyspnea, and myalgia. Vital signs of and her physical examination was unremarkable. Initial polymerase chain reactesting for SARS-CoV-2 was negative and the patient was discharged home. She weeks later with more severe dyspnea, cough, dysarthria, dysphagia, odynophysevere generalized weakness with inability to ambulate. She had no sensory sybowel or bladder dysfunction.

Physical examination was significant for tachycardia at 110 beats/min and oxygof 88% on room air. She had bilateral ptosis, facial weakness, hypernasal dysar profound symmetric proximal limb weakness. Reflexes were symmetrically din Repeat SARS-CoV-2 PCR test was positive.

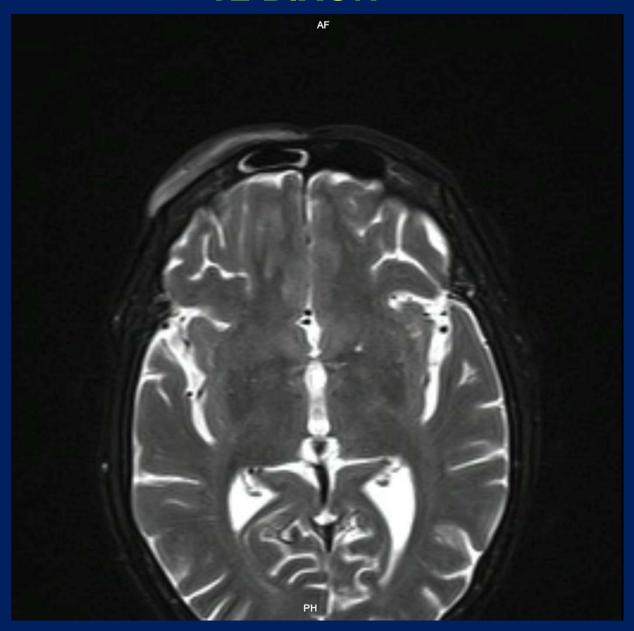
MRI of the entire neuroaxis showed no central or peripheral nervous system in but it did demonstrate diffuse muscle edema and enhancement, with a region myonecrosis (Figure <u>1A,B</u>).

A and B, Sagittal T1WIs of contrast-enhanced fat-suppressed MRI of the paraspinal and this demonstrate extensive edema and enhancement (yellow arrows in A) consistent with inflar myopathy; central nonenhancement in the vastus medialis (yellow arrow in B) is consistent myonecrosis. C-F, Biopsy of the left quadriceps muscle. Hematoxylin-and-eosin cryostat seed demonstrate multifocal, predominantly perimysial perivascular lymphocytic inflammation (arrows in D), with focal endomysial extension (black arrows in D). Multiple regenerating my arrows in E) are recognized by their mild sarcoplasmic basophilia and enlargement of visible upregulation of HLA class ABC on myofiber surfaces, and sarcoplasmic staining by immunican be identified by the brown staining of myofibers, most consistent with an inflammatory Scale bar = 50 µm. HLA, human leukocyte antigen; T1WI, T1-weighted image

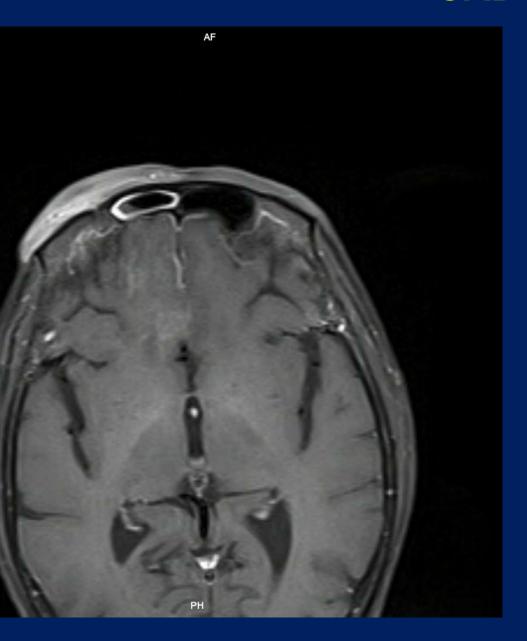
> 2ND CASE:

- CHIEF COMPLAINT: The patient had pain in the right forehead and eye for abt 2 days, went to Cho Ray hospital, was diagnosed with sinusitis and given medicine.
- Almost 1 week later, the patient's right eye was red, swollen and painful, with a slight fever, the patient went to Eye Hospital and was assigned to have an MRI at Medic center
- MEDICAL HISTORY:
 - Had Covid self-treatment at home with oxygen support, within 22 days, recovered more than 1 week ago.
 - Diabetes, Hepatitis for more than 10 years.

T2 DIXON



GADOVIST





LÝ DO KHÁM

viêm tổ chức hốc mắtSIEMEN ESSENZA 2

Máy Vùng

: MRI SO NÃO _ HỐC MẮT

Không, sau đó tiêm tương phản

Kết quả

: Mắt và sọ não được khảo sát cộng hưởng từ có tiêm thuốc tương phản với thông số kỹ thuật: Axial fat NE+CE-T1GRE, T2WI; sagittal T1WI; coronal CE-T1WI; axial Flair T2WI

XOANG

Dày niêm mạc xoang trán, hàm phải, sàng, bướm hai bên. Vòm hầu và các khoang cạnh hầu bên phải trái đều trống. Các tế bào chủm hai bên không viêm. Không tín hiệu bất thường xoang hang MẮT

Thương tổn dạng thâm nhiễm lan tỏa mô dưới da vùng trán hai bên - thái dương phải, giác mạc mắt p cơ thẳng ngoài và cơ thẳng trên phải. Dày màng não trán hai bên.

Nhãn cầu trái có kích thước, hình dạng và tín hiệu bình thường trên các chuỗi xung khảo sát, không t bong võng mạc.

Không thấy tín hiệu viêm thần kinh thị.

SO NÃO

Tẩng trên lều và dưới lều không thấy khối choán chỗ, không xuất huyết hoặc tụ máu cũ. Không thấy b lý chất trắng. Các não thất bình thường. Rãnh vỏ não bình thường. Không thương tổn tuyến yên, khôr thương tổn vùng giao thoa thị giác.

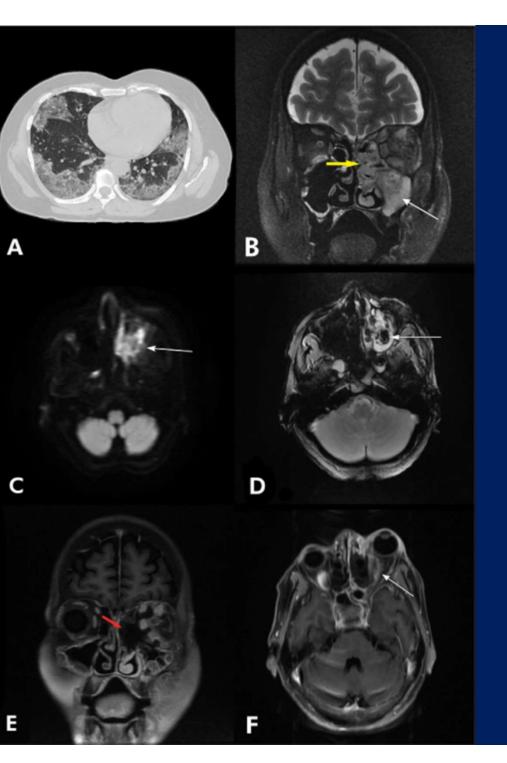
*** KẾT LUÂN:

Thương tổn dạng thâm nhiễm lan tỏa mô dưới da vùng trán hai bên - thái dương phải, giác mạc mắt p cơ thẳng ngoài và cơ thẳng trên phải nghĩ do Viêm. Theo dõi viêm màng não trán hai bên.

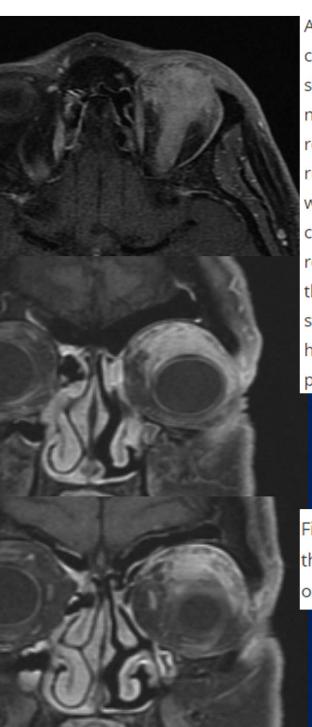
Không khối choán chỗ nội sọ. Không thấy thương tổn nhu mô não.

Viêm đa xoang.

Tp. Hồ Chí Minh, ngày 14/09/2021 10:49 (Bác sĩ đã ký)



a Axial HRCT thorax image showing ground opacities in subpleural regions of bilateral parenchyma with "crazy paving appearance." b C T2 FS image showing mucosal thickening and collect the left maxillary sinus (white arrow), left eth sinus, and left middle nasal turbinate (yellow arrov inflamed extraocular muscles. c Axial DWI image sl restricted diffusion in the left maxillary sinus a middle nasal turbinate. d Axial GRE image showing blooming in the left maxillary sinus arrow). e Coronal T1 post-contrast image showing non-enhancing soft tissue in left middle nasal tur and within the left maxillary antrum ("black tur sign"). f Axial T1 post-contrast image sl enhancement and inflammation involving extra muscles of left orbit causing proptosis



A 44-year-old male tested positive for SARS-CoV19 after experiencing back pain and fever to 100 complained of sinus congestion but had no purulent nasal discharge. His past medical history was significant for an idiopathic facial palsy which occurred spontaneously, resolved completely, and no additional workup or neuroimaging and a neurogenic tumor removed from his spine in 1995 recurrence. Two days after his COVID-19 diagnosis, the patient noted swelling of the left eyelids a rubbing while sitting in a hot tub. Over the next 24 hours, he developed proptosis and periocular with discomfort. He was treated in an outpatient urgent care clinic with a 3 day course of oral corticosteroids with improvement in his symptoms. However, upon steroid cessation, his eyelid serior recurred and he was started on a course of oral cefazolin. His swelling progressed to involve his the next day the patient was admitted to an outside hospital for intravenous vancomycin and an swelling did not progress and he was discharged several days later on oral antibiotics. He returns hospital five days after discharge with worsening symptoms. Intravenous antibiotics were restart patient was transferred to our facility.

Figure 1. Axial and coronal, post contrast T1 weighted magnetic resonance images show enhalthe superior orbit involving the left superior rectus/levator complex and extending into the sorbital soft tissue

CONCLUSION:

Much is still unknown about how COVID-19 will affect people over ime, but research is ongoing. Researchers recommend that doctors losely monitor people who have had COVID-19 to see how their organs are functioning after recovery.

Although MRI is not a first-line diagnostic tool for COVID-19 patients, still it has certain values in following up, evaluating lisease progression, detecting complications due to COVID-19

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https://onlinelibrary.wiley.com/doi/10.1002/mus.27003

https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8254439/

https://www.tandfonline.com/doi/full/10.1080/01676830.2021.1962366

https://www.mayoclinic.org/diseases-conditions/coronavirus/indepth/coronavirus-long-term-effects/art-20490351

THANKS FOR YOUR LISTENING!