June 2023

When Your Teammate Kneeds You

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Laser-Assisted Delivery of Trichloroacetic Acid for Successful Treatment of Persistent Facial Hyperpigmentation

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**Mentor:** Ronald J. Sulewski  
**Program:** Dermatology  
**Type:** Case Report  

**Background:** Hyperpigmentation is a common cosmetic dermatology concern that can be difficult to treat. Standard treatment options include various skin lightening topical and oral medications, as well as in-office options such as chemical peels and laser treatments. While laser-assisted drug delivery is an emerging treatment in the field of dermatology, there are no reports in the literature of laser-assisted delivery of trichloroacetic acid (TCA), a chemical peel commonly used for hyperpigmentation.

**Case:** A 35-year-old female, with previous use of injectables (botulinum toxin and dermal fillers), presented with dynamic rhytids and hyperpigmentation. The patient reported using sunscreen and antioxidant serum (SkinCeuticals CE ferulic) daily. On physical examination, there were scattered brown macules on the face and a brown patch on the left upper lip. The patient initially underwent treatment with a series of four single pass TCA 15% peels to the face in 1-month intervals, with an additional pass of TCA 25% to the upper lip and cheeks in the latter two sessions. The patient reported improved hyperpigmentation, but with a persistent brown patch on the upper lip.

Eight months later, the patient underwent laser-assisted delivery of TCA 15% during full face treatment with Fraxel Dual for the patch on the left upper lip [Settings: wavelength 1927 nm, 10 mJ, treatment level 4 (35%), 8 passes, 1.43KJ]. The treatment resulted in significant improvement of the hyperpigmented patch on the upper lip (Figure 1).

**Conclusion:** In this case, laser assisted delivery was successfully utilized for better penetration of TCA and resulting improvement in facial hyperpigmentation.

[Poster Image: Before and After Laser-Assisted Delivery of TCA]

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Not your Typical Lateral Ankle Sprain

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**Mentor:** Jason Meredith  
**Program:** Family Medicine  
**Type:** Case Report  

**Background:** Extensor digitorum brevis (EDB) avulsions present with anterior tenderness below the anterior talofibular ligament after inversion ankle injury. This injury can be misdiagnosed as normal anatomy, sprain, or another type of fracture. Misleading anatomy includes the os peroneum and sesamoid bone. Fractures that can be misdirecting include comminuted fracture of the cuboid, anterior process of calcaneus, and avulsion of calcaneofibular ligament (CFL).

Case: A 39-year-old male presents to sports medicine clinic for evaluation of right ankle injury. One week prior, he had been hiking in Colorado and had multiple inversion ankle injuries which led to swelling but denied limitations to activity. On his return trip he suffered another inversion injury going downstairs resulting in significant swelling and difficulty ambulating secondary to pain. Patient saw his primary care provider and Xray of right ankle revealing multiple fractures of unknown acuity. CT w/o contrast revealed acute to subacute mildly displaced avulsion fractures of the dorsal talus and anterolateral calcaneus. Given the location of his symptoms, our final diagnosis was an extensor digitorum brevis avulsion fracture. Patient utilized CAM boot for 4 weeks followed by one-week progression out of CAM boot. Patient reported improvement in pain and ability to bear weight. He utilized an ASO ankle brace with activity and started physical therapy for proprioception retraining.

**Conclusion:** EBD avulsion is an atypical cause of lateral ankle pain and should be on the differential for inversion ankle injuries. Multiplanar imaging can assist in its diagnosis if plain films are nonconfirmatory.

[Poster Image: Ankle Injury Image]
Delayed Malignant Cerebellar Posterior Reversible Encephalopathy Syndrome secondary to CLAG-M and Venetoclax regimen

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Mentor: Subin Mathew
Program: Neurological Sciences
Type: Case Report

**Background:** Isolated cerebellar malignant posterior reversible encephalopathy syndrome (PRES) is rare. We report a case of delayed malignant cerebellar PRES triggered by cladribine, cytarabine, granulocyte colony-stimulating factor, mitoxantrone (CLAG-M), and venetoclax regimen. PRES is a heterogeneous syndrome caused by multiple factors. Cerebellar PRES is commonly seen with autoimmune disease and rarely with chemotherapy.

**Case:** A 73-year-old male presented with a history of relapse after allogeneic stem cell transplantation for refractory myelodysplastic syndrome on maintenance tacrolimus. He was admitted for salvage therapy with (CLAG-M) + venetoclax regimen. On day 16 post salvage therapy the patient developed acute onset altered mental status and headache followed by fever and hypotension. Brain MRI revealed posterior bilateral cerebellar, vermian edema with tiny petechial hemorrhages, ventriculomegaly, trans ependymal edema, and mass effect on fourth ventricular effacement. Despite external ventricular drain placement, hyperosmolar therapy, and steroids, on day 19 the patient suffered a devastating neurological injury and the family opted for hospice care. Off note, maintenance tacrolimus was held prior to admission and was started 4 days prior to the onset of neurological symptoms with subtherapeutic levels.

**Conclusion:** The potential role of (CLAG-M) and venetoclax regimen in the pathogenesis of PRES needs to be considered. Cerebellar PRES is rare and can be complicated by early hydrocephalus and 4th ventricle compression. Although PRES is usually benign, prompt recognition of indicators of the malignant phenotype and aggressive management is crucial.

https://doi.org/10.32873/unmc.dc.gmerj.5.1.069

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Intravenous Alteplase in CADASIL

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Mentor: Marco Gonzalez-Castellon
Program: Neurology
Type: Case Report

**Background:** Cerebral autosomal dominant arteriopathy with subcortical infarcts and leukoencephalopathy (CADASIL) is a genetic small vessel vasculopathy caused by mutations in the NOTCH3 gene that result in vasculopathic changes in cerebral small penetrating arteries, arterioles, and capillaries. Stroke is an important cause of morbidity and mortality. Theoretical concerns have been raised about the efficacy and safety of intravenous alteplase (IV tPA) in the treatment of small vessel stroke in these patients. However, there is little clinical evidence to validate or discredit these concerns.

**Case:** A 40-year-old female with a diagnosis of CADASIL presented to our emergency department for acute onset non-fluent aphasia and significant worsening of baseline left-sided hemiparesis. National Institutes of Health Stroke Score (NIHSS) was 12. Computed tomography (CT) did not show any signs of intracranial hemorrhage or large vessel occlusion. The patient presented within the window for IV tPA treatment, and no contraindications were identified. IV tPA was administered with no immediate complications. Approximately 12 hours later, the patient had a brief episode of left-sided head deviation and prominent nystagmus concerning for focal seizure. Magnetic resonance imaging (MRI) showed a 3.2cm right frontal lobe intraparenchymal hemorrhage (Figure 1) and small acute infarcts in the right posterior internal capsule and left postcentral gyrus.

**Conclusion:** CADASIL carries a risk of bleeding, and the mechanism by which it causes small vessel stroke may be unrelated to thrombosis. Further study is necessary to determine if and/or which patients with CADASIL may benefit from IV tPA.

https://doi.org/10.32873/unmc.dc.gmerj.5.1.070

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*Names in bold type indicate presenting author.

**Figure 1:** Axial MRI gradient echo (GRE) sequence showing right frontal lobe intraparenchymal hemorrhage and surrounding microhemorrhages
Atrial Flutter due to Amiodarone-Induced Thyrotoxicosis Months after Discontinuation
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Mentor: Dalton Nelsen
Program: Emergency Medicine
Type: Case Report

Background: A 56-year-old man presenting with one month of recurrent episodes of atrial flutter and seizure-like activity. He had no history of alcohol or illicit drug use, no recent fall or trauma event, and no headaches or recent infections.

Case: On presentation, he was afebrile and blood pressure was 151/85. He had proximal muscle weakness and bilateral lower extremity edema on exam. Additional examination findings were unremarkable. Initial lab work revealed blood glucose 294, potassium 2.3, and calcium 7.3 (ionized calcium 1.07). Eight AM cortisol was 101, ACTH was 437, and DHEA-S was 180. PET scan revealed bilateral adrenal hyperplasia without any extra adrenal masses. MRI of the pituitary revealed a 1-2 mm hypodensity, and inferior petrosal sinus sampling was performed. He subsequently underwent bilateral adrenalectomy and was placed on a hydrocortisone plus fludrocortisone steroid taper.

Conclusion: The adrenal glands secrete glucocorticoids, mineralocorticoids, and sex hormones, which are regulated by the hypothalamic-pituitary-adrenal (HPA) axis. Increased plasma ACTH concentrations cause bilateral adrenocortical hyperplasia. Overactive adrenal glands resulting in hypercortisolism can lead to elevated blood glucose levels due to increased glucocorticoid activity. Further, increased mineralocorticoid activity can cause hypokalemia and hypertension from the effects of abnormal aldosterone secretion. This patient was diagnosed with Cushing’s disease secondary to a pituitary tumor. The treatment of choice for Cushing’s disease (ACTH-producing pituitary tumor) is transphenoidal microadenomectomy when a microadenoma can be identified. Pituitary irradiation can also be considered. Ultimately, bilateral total adrenalectomy with lifelong daily glucocorticoid (eg, hydrocortisone) and mineralocorticoid (eg, fludrocortisone) replacement therapy is the final definitive cure.

https://doi.org/10.32873/unmc.dc.gmerj.5.1.071

Rabbit Allergy and Kidney Transplant: What’s the connection?
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Mentor: Scott Westphal
Program: Internal Medicine
Type: Case Report

Background: An anti-thymocyte globulin (ATG, Thymoglobulin®) is a polyclonal, anti-human thymocyte antibody derived from rabbits, used for lymphocyte depleting induction immunosuppression in solid organ transplantation. Despite its routine use in transplantation, reports of life-threatening anaphylaxis are extremely rare.

Case: A 37-year-old woman presented for a kidney transplant. She received midazolam, propofol, lidocaine, rocuronium, fentanyl for anesthesia, preoperative cefazolin, followed by antithymocyte globulin. Shortly thereafter she developed shock and ultimately cardiac arrest. With CPR and aggressive hemodynamic support, she achieved return of spontaneous circulation and did eventually recover, although her transplant surgery was aborted. Serum tryptase was elevated (consistent with mast cell degranulation). Subsequent allergy skin prick testing was negative, except ATG which was positive at a 1:1,000 dilution, and elevated serum antigen...
When Your Teammate Kneeds You

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Mentors: Jason Meredith, Ross Mathiasen
Program: Primary Care Sports Medicine
Type: Case Report

Background: A 21-year-old male collegiate basketball athlete initially presented to the athletic training room for evaluation of left groin pain. He reported being kneed in the left thigh near the time of onset 5 days ago. He had consistent soreness since onset. Notably, he recently had surgical fixation of a Jones fracture resulting from stress injury. These new symptoms occurred during his return to play progression after surgery.

Case: Exam consisted of palpatory tenderness near the left inguinal canal and LLQ. He also experienced pain with a sit-up and straight leg. An MRI was ordered due to concern of a possible sports hernia. MRI showed a left pubic body stress fracture. A metabolic workup was pursued which demonstrated low vitamin D levels. Calcium and vitamin D were supplemented, and he stopped all pain provoking activities for 6 weeks. During his subsequent return to play progression, he developed right groin pain. Repeat pelvis MRI now showed a right pubic body stress fracture and unchanged left pubic body stress fracture. He was subsequently removed again from activity. Vitamin D and calcium supplementation were continued during this time.

Conclusion: Fourteen weeks after the repeat MRI, he was symptom free with no tenderness to palpation. Vitamin D level was rechecked. They were adequate at 44 ng/ML. He slowly re-started his return to play progression, adding impact activities over the following 6 weeks. During this time an unremarkable gait analysis was performed. There was no return of symptoms and he has returned to full activity.

https://doi.org/10.32873/unmc.dc.gmerj.5.1.074

Not your TIPICal Neck Pain: Case of Transient Perivascular Inflammation of the Carotid Artery (TIPIC) Syndrome

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Mentor: Jonathan Muldermans
Program: Family Medicine
Type: Case Report

Background: We present a case of a rare cause of neck pain called Transient Perivascular Inflammation of the Carotid artery (TIPIC) syndrome. TIPIC is an idiopathic self-resolving process identified by clinical and radiographic criteria.

Case: A 41-year-old female with history of smoking presented to clinic with acute right sided neck pain, but otherwise feeling well. Vitals were normal. On exam she had tenderness overlying the right carotid artery. CT neck revealed mild perivascular thickening around the right common carotid artery extending through the carotid bulb, without associated occlusion or high-grade stenosis. CBC, ESR, CRP were normal. U.S. demonstrated mildly increased velocity of right carotid, without evidence of intimal thickening or plaque. These imaging findings are consistent with TIPIC syndrome. Patient’s neck pain resolved over next 2 weeks. Follow-up ultrasound 6 months later demonstrated resolution of perivascular findings.

Conclusion: TIPIC is important to include in patients with acute unilateral neck pain and carotid tenderness. TIPIC was newly identified and defined in 2017 with four proposed major criteria: acute pain overlying carotid artery, perivascular inflammation on imaging, exclusion of another diagnosis, and improvement within 14 days. Mild elevations in inflammatory markers can be seen. Treatment is supportive with NSAIDs. It is self-resolving process within 14 days. It is likely under-recognized due to its transient nature. Among various other risk factors, TIPIC has been seen in patients with tobacco use as well as recent viral illness.

https://doi.org/10.32873/unmc.dc.gmerj.5.1.075