

RADYOLOJİ BAŞUCU SERİSİ

Nöroradyoloji

EDİTÖRLER
Mehmet Ali Gedik
Fuldem Mutlu

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1

KRANİAL TRAVMA

Mustafa HIZAL¹

Vaka 1: Kalvaryal kırık

Vaka 2: Travmatik epidural hematoma ve kalvaryal kırık

Vaka 3: Travmatik subdural hematoma ve difüz aksonal hasar

Vaka 4: Subaraknoid ve subdural kanama ve kalvarial kırık

Vaka 5: Travmatik serebral kontüzyon

Vaka 6: Parankimal ve subdural hematoma, subaraknoid kanama

Vaka 7: Difüz Aksonal Hasar

Vaka 8: Düşük hızlı penetran kafa travması

Vaka 9: Dural sinüs trombozu, serebellar venöz enfarkt, parankimal hematoma

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BÖLÜM

2

KONJENİTAL SEREBRAL MALFORMASYONLAR

Elif GÖZGEÇ1

Vaka 1: Oksipital sefalosel

Vaka 2: Chiari tip 1 malformasyon

Vaka 3: Chiari tip 2 malformasyon

Vaka 4: Alobar holoprozensefali

Vaka 5: Dandy Walker Malformasyonu

Vaka 6: Araknoid Kist

Vaka 7: Açık dudak şizensefali

Vaka 8: Klasik Lizenfali

Vaka 9: Korpus kallozum agenezisi

Vaka 10: Subependimal Heterotopi

Vaka 11: Baziler İnvajinasyon

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- McRae hattı: Foramen magnumun anterior hattının orta noktası (basion) ile posterior hattının orta noktasını (opistion) birleştiren çizgidir. Normalde odontoid proçes bu hattın altında kalır.
- Chamberlain hattı: Sert damak arka kenarından opistiona çizilen hattır. Odontoid proçesin bu hattı 5 mm'den fazla geçmesi tanı koydurur.
- McGregor hattı: Sert damak arka kenarından oksipital kemik en alt noktasına çizilen hattır. Odontoid proçesin bu hattı 7 mm'den fazla geçmesi tanı koydurur (20-22).

Tuzaklar

Platibazi kafatası tabanının anormal düzleşmesidir. Beraberinde baziler invajinasyon yoksa asemptomatiktir (3).

Tedavi ve Yaklaşım

Semptomatik olanlarda cerrahi tedavi uygulanır.

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BÖLÜM

3

NÖROKUTANÖZ SENDROMLAR

İlyas DÜNDAR¹

Vaka 1: Nörofibromatozis Tip 1 (Beyin Tutulumu + Nörofibromlar)

Vaka 2: Nörofibromatozis Tip 1 (Beyin Tutulumu) + Ewing Sarkom

Vaka 3: Nörofibromatozis Tip 2 (Meningiom + Sağ Vestibüler Schwannom + Optik Gliom)

Vaka 4: Tüberoskleroz (Beyin Tutulumu)

Vaka 5: Tüberoskleroz (Beyin Tutulumu) + Pankreas Orta Dereceli Nöroendokrin Tümör (Distal Pankreotektomi)

Vaka 6: Sturge-Weber Sendromu

Vaka 7: Sturge Weber Sendromu

Vaka 8: Von Hippel-Lindau Sendromu (Hemanjioblastom + Pankreatik Kist + Sol Böbrek RCC)

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Tuzaklar

MRG'de kistik lezyon ve kontrast tutan mural nodül ile tanınan serebellar hemanjiyoblastom VHL sendromuna eşlik edebildiği gibi sporadik de olabilir. Bu yüzden serebellar hemanjiyoblastom tanısı konan hastalarda diğer sistemlerin de taranması gerekir.

Tedavi ve yaklaşım

Kraniyospinal lezyonlar cerrahi olarak güvenli bir şekilde rezeke edilir ve sıklıkla küratiftir. SSS hemanjiyoblastomları farklı bölgelerde ve farklı zamanlarda büyüebildiğinden, hastalar semptomatik olana kadar cerrahi yaklaşım yapılmaz. Asemptomatik lezyonlar yıllık görüntüleme ile izlenir. Büyük boyutlu tümörlerde tümör vaskülaritesini kontrol etmek için ameliyat öncesi arteriyel embolizasyon yapılmalıdır (22). Postoperatif dönemde SSS MRG ile yakın takip edilmeli ve diğer sistemler de tutulum açısından araştırılmalıdır.

Anahtar Kelimeler

nörokütanöz sendrom, nörofibromatozis, von recklinghausen hastalığı, cafe' la lait lekeleri, legius sendromu, optik yol gliomları, nörofibrom, pleksiform nörofibrom, iris hamartomu, lisch nodülleri, tanımlanamayan parlak objeler, fokal sinyal alanları, vestibüler schwannom, menenjiom, ependimom, tüberoskleroz, bourneville hastalığı, kortikal tüber, subependimal nodül, subependimal dev hücreli astrositom, nöroendokrin tümör, sturge-weber sendromu, ensefalotrigeminal anjiyomatozis, serebral hemiatrofi, giral kalsifikasyonlar, oksipital kalsifikasyon, epilepsi, porto-şarap lekesi, leptomeningeal anjiyom, kalvaryal hipertrofi, CEC sendromu, PHACE sendromu, von hippel-lindau sendromu, serebellar hemanjiyoblastom, mural nodül, feokromositoma, paraganglioma, pankreatik kist, renal hücreli karsinom, MRG, BT

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BÖLÜM

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VASKÜLER OKLÜZİV HASTALIKLAR

Fatih KILINÇ¹

Vaka 1: Akut iskemik inme, MCA oklüzyonu

Vaka 2: Baziler arter oklüzyonu, vertebrobaziler inme

Vaka 3: Ekstrakranial ICA darlığı, hemodinamik enfarkt

Vaka 4: SAK sonrası vazospazm ve serebral iskemi

Vaka 5: Serebral venöz trombüs

Vaka 6: Santral sinir sistemi vaskülit

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Tuzaklar

Görüntüleme yöntemleri vaskülit tanısını destekler ancak kesin tanı koydurmaz.

Tedavi ve yaklaşım

Tedavide yüksek doz kortikosteroid ve gereklilik halinde immünsüpresif ilaçlar kullanılır.

Anahtar Kelimeler

Akut iskemik inme, vertebrobaziler inme, penumbra, iskemik kor, borderzone enfarkt, NIHSS, ASPECTS, vazospazm, gecikmiş serebral iskemik, MCA oklüzyonu, dens MCA, baziler arter oklüzyonu, serebral venöz tromboz, empty delta işareti, semptomatik ICA darlığı, bilateral talamik ödem, NASCET, trombolitik, trombektomi, IV tPA, vaskülit

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- American Society of Neuroradiology, Congress of Neurological Surgeons, Society of Atherosclerosis Imaging and Prevention, Society for Cardiovascular Angiography and Interventions, Society of Interventional Radiology, Society of NeuroInterventional Surgery, Society for Vascular Medicine, and Society for Vascular Surgery [published correction appears in *Circulation*. 2011 Jul 26;124(4):e146. Dosage error in article text] [published correction appears in *Circulation*. 2012 Jul 10;126(2):e26]. *Circulation*. 2011;124(4):e54-e130. doi:10.1161/CIR.0b013e31820d8c98
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BÖLÜM

5

NON OKLÜZİV VASKÜLER PATOLOJİLER

Seray Akçalar¹

Vaka 1: Benign Perimezensefalik Subaraknoid Kanama

Vaka 2: Hipertansif lobar hematom

Vaka 3: Epidural hematom ve travmatik SAK

Vaka 4: Akut ve subakut subdural hematom

Vaka 5: Serebral arteriyovenöz malformasyon

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BÖLÜM



DURAL VE LEPTOMENİNGEAL TUTULUM İLE SEYREDEN HASTALIKLAR

Ezra Çetinkaya¹

Vaka 1: EBV Meningoensefalit

Vaka 2: Menenjiom

Vaka 3: Primer Diffüz Leptomeningeal Gliomatozis(PDLG)

Vaka 4: Akut iskemik inme

Vaka 5: Sarkoidoz

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yönelik girişimlere de ihtiyaç duyulabilir(52,57). Hasta takibi her hastaya özgü değişmeliyken, göz muayenesi, kontrastlı Beyin MRG ya da PET görüntüleme fayda sağlar(59).

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BÖLÜM



HİDROSEFALİ VE DİĞER BOS BOZUKLUKLARI

Temel Fatih YILMAZ¹

Vaka 1: Koroidal pleksus papillomu

Vaka 2: Normal basınçlı hidrosefali

Vaka 3: Aquaduktal web

Vaka 4: Chiari malformasyonuna sekonder hidrosefali

Vaka 5: Rinore, BOS kaçağı

Vaka 6: Komünikan hidrosefali

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Tanı

- Kommünikan hidrocefali

Ayırıcı Tanı

Atrofik hidrocefali

Normal basınçlı hidrocefali

Önemli Noktalar

Etiyolojik faktörler arasında subaraknoid kanama, menenjit, normal basınçlı hidrocefali ve leptomeningeal tutulum gösterilebilir. Beyin omurilik sıvısındaki azalmış rezorpsiyona sekonder olduğu düşünülmektedir (9).

Tuzaklar

Komünikan hidrocefaliyi parankimal hacim kaybı nedeniyle ventriküllerin ex vacuo dilatasyonu olan hastalarda görülen ventriküler genişlemeden ayırt etmek zor olabilir. Üçüncü ventrikül ve lateral ventriküllerin temporal hornlarındaki genişleme tanı için önemlidir çünkü ex vacuo dilatasyonu olan hastalarda bu bölgeler nispeten korunur (10).

Tedavi ve yaklaşım

Komünikan hidrocefali tedavisinde endoskopik yöntemlerin aksine genellikle ventriküloperitoneal şant prosedürleri kullanılmaktadır.

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BÖLÜM

8

EPİLEPSİ

Berrak Barutcu ASFUROĞLU¹
Umut ASFUROĞLU²

Vaka 1: Disembriyoplastik Nöroendokrin Tümör (DNET)

Vaka 2: Tüberoskleroz

Vaka 3: Fokal kortikal displazi (FKD) Tip IIb

Vaka 4: Status Epileptikus (SE)

Vaka 5: Sol mezial temporal skleroz (MTS)

Vaka 6: Sturge Weber Sendromu (SWS)

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BÖLÜM



DEMİYELİNİZAN VE DİSMİYELİNİZAN HASTALIKLAR

Zakir Sakcı¹

Vaka 1: Multipl Skleroz

Vaka 2: Tümefaktif Multipl Skleroz, Servikal tutulum

Vaka 3: Akut Dissemine Ensefalomyelit (ADEM)

Vaka 4: Nöromiyelitis optika, Optik nörit

Vaka 5: Canavan Hastalığı

Vaka 6: Alexander Hastalığı

Vaka 7: X'e bağlı Adrenolökodistrofi

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yon görülür. T2 ağırlıklı MRG'de orta derece hiperintens olup kontrast tutmaz. ALD'de proton MR spektroskopisi NAA pikinde azalma, kolin pikinde artma miyoinozitol pikinde azalma ve laktat pikinde artma görülür (18–20).

Tuzaklar

Neonatal hipoglisemide korpus kallozum splenium, peritrigonal beyaz cevher ve oksipital korteks tutulabilir. Postkontrast görüntülerde kontrastlanma izlenmez. Aleksander hastalığında peritrigonal beyaz cevherden ziyade frontal periventriküler beyaz cevher tutulur. Postkontrast serilerde kontrastlanma görülebilir.

Tedavi ve yaklaşım

Lorenzo'nun yağı semptomların gelişmesini geciktirebilir. Kemik iliği transplantasyonu ve kök hücre tedavisi denenebilir. Tedavi sonrası metabolik stabilizasyon ve kısmen geri dönüş görülebilir (20).

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BÖLÜM

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METABOLİK VE TOKSİK HASTALIKLAR

Saliha ÇIRACI¹

Vaka 1: L-2-Hidroksiglutarik Asidüri

Vaka 2: Adrenolökodistrofi

Vaka 3: Glutarik asidüri tip 1

Vaka 4: Canavan hastalığı

Vaka 5: Metakromatik lökodistrofi (MLD)

Vaka 6: Krabbe hastalığı

Vaka 7: Van Der Knaap Hastalığı

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Ayrırcı Tanı

Alexander hastalığı, Canavan hastalığı ve Glutarik asidüri, ayrırcı tanıda düşünülebilir. Bu hastalıklarda MRG'de subkortikal kist yoktur ve hepsinde bazal gangliyon tutulumu görülür (13).

Önemli Noktalar

Van Der Knaap hastalığı megalensefalik lökoensefalopati ve subkortikal kistlerle karakterize, MLC 1 geni mutasyonuna bağlı kalıtsal otozomal resesif bozukluktur. Klinik olarak makrosefali, motor gelişim geriliği ve nöbetlerle karakterizedir. İlerleyen dönemlerde kademeli ataksi ve piramidal bulgular eklenir. Zihinsel kapasiteler genellikle korunur, ancak hafif bozulma olabilir. Tanı için klinik özellikler ve MRG bulgularının kombinasyonu gereklidir (13). Hastaların çoğu erken ergenlikte tekerlekli sandalyeye bağımlı hale gelir. 2. ya da 3. dekada ölüm görülür (14).

Tipik MRG bulguları 6. aydan itibaren görülebilir. Periventriküler bölgenin korunduğu beyaz cevher tutulumu izlenir, gri cevher tutulmaz. Temporal ve frontoparietal bölgede bilateral subkortikal kistler görülür (14).

Tuzaklar

Beyaz cevher tutulum şiddeti ile hastalığın klinik seyri arasında uyumsuzluk bulunur. Klinik bulgular yavaş seyirlidir (14).

Tedavi ve yaklaşım

Van Der Knaap hastalığının kesin bir tedavisi yoktur. Nöbetleri önlemek için antiepileptikler ve motor bozukluğa yönelik fizyoterapi uygulanır (14).

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NÖRODEJENERATİF HASTALIKLAR

Yusuf CAN¹

Vaka 1: Alzheimer Hastalığı (AH)

Vaka 2: Multipl beyaz cevher iskemileri, kronik laküner iskemi odakları
ve bununla ilişkili vasküler demans

Vaka 3: Huntington hastalığı

Vaka 4: Creutzfeldt-Jacob hastalığı

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En yaygın görüntüleme bulgusu tipik olarak frontal, temporal ve parietal loblarda hızlı ilerleyici atrofi görülmesidir. Nadir tiplerde oksipital (Heidenhain varyantı) ve serebellar (Oppenheimer-Brownell varyantı) hemisferlerde atrofi görülebilir (19, 20). Bunun yanında diğer spesifik görüntüleme bulgularından ilki, talamik pulvinarlarda difüzyon kısıtlanması, ikincisi talamik pulvinarlar ve dorsomedial talamuslarda T2 sinyal artışına bağlı 'hokey sopası' görünümünün olmasıdır (21). Diğer nadir bulgular ise serebral kortekslerde yamalı giriform T2 hiperintensitesi ve globus palliduslarda T1 sinyal artışı olmasıdır (22).

FDG PET/BT'de klasik bir görünüm tanımlanmamıştır.

Amiloid PET incelemede amiloid depozisyonu görülmez (16).

Tuzaklar

FDG PET/BT görüntüleme bulgusu olmasa da bazı olgularda Kortikobazal dejenerasyon ve Lewy cisimcikli demans yanlış tanısına neden olabilecek fokal hipometabolizma görülebilir. Bu olgularda klinik ve görüntüleme hızı progresyon olması CJH lehine tanı koydurur (16).

Tedavi ve Yaklaşım

Günümüzde küratif tedavi yöntemi yoktur. Olguların çoğunda ortalama 7 aylık yaşam beklentisi olan ölümcül bir hastalıktır.

Anahtar Kelimeler

Demans, nörodejeneratif, beyin, MRG, amiloid görüntüleme, FDG PET/BT, medial temporal lob atrofisi, Alzheimer, lewy cisimciği, vasküler demans, huntington, Creutzfeldt-Jacob hastalığı, prion hastalığı,

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BAŞ BOYUN ENFEKSİYON VE İNFLAMASYONLARI

Cansu ÖZTÜRK¹

Vaka 1: Retrofarengeal apse

Vaka 2: Süpüratif lenfadenit (intranodal apse), retrofarengeal ödem

Vaka 3: Mastoidit+serebellar apse

Vaka 4: Peritonsiller apse

Vaka 5: Tüberküloz (tbc) menenjit+tüberkülom

Vaka 6: Sol parotis gland; akut süpüratif parotitis

Vaka 7: Sağ submandibuler gland; sialolitiazis+sialoadenit

Vaka 8: Ludwig angina+boyun sağ yarı flegmon+apse

Vaka 9: Lakrimal gland psödötümör/idiopatik orbital enflamasyonu

Vaka 10: Akut serebellit

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Tuzaklar

Akut serebellit olgularının ilk başvuruları acil servislere olmaktadır, bu nedenle karşımıza öncelikle beyin BT ile gelmeleri mümkündür. Her ne kadar beyin BT'de ışın sertleşme artefaktları nedeniyle posterior fossa değerlendirilmesi güç olsa da serebellar hipodansite varlığı uyarıcı olmalıdır.

Tedavi ve yaklaşım

Çoğunlukla kendi kendini sınırlayan bir durumdur, tedavi için konsensusa varılmış bir protokol yoktur, etiyojoloji ve kafa içi basınç artışına yönelik tedaviler uygulanır. Bu amaçla; steroid, mannitol ve antiviral/antibiyotik tedavisi verilebilir. Komplike olgularda (hidrosefali, tonsiller herniasyon vb.) cerrahi dekompresyon ve drenaj gerekebilir.

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EKSTRAAKSİYEL TÜMÖRLER

*Aydın ASLAN¹
Elif GÜNAYDIN¹*

Vaka 1: Menenjiom

Vaka 2: Dural metastaz

Vaka 3: Vestibüler schwannom

Vaka 4: Trigeminal schwannom

Vaka 5: Hemanjioperisitom

Vaka 6: Epidermoid

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Tedavi ve yaklaşım

Karakteristik özellikleri rezeksiyonunu zorlaştırmaktadır. Total rezeksiyon postoperatif aseptik menenjit, hidrosefali ve tümör rekürrensi riskini etmesine rağmen agresif cerrahi kraniyal sinir ya da iskemik defisitler ile ilişkili olabilir.

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BÖLÜM

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İNTRAAKSIYEL SUPRATENTORIAL TÜMÖRLER

Ali Murat KOÇ¹

Vaka 1: Glioblastom, NSO, DSÖ Evre 4

Vaka 2: Adenokarsinom metastazı

Vaka 3: Difüz Büyük B Hücreli Lenfoma (Primer Santral Sinir Sistemi Lenfoması)

Vaka 4: Oligodendrogliom, NOS, DSÖ Evre II

Vaka 5: Disembriyoplastik Nöroepitelyal Tümör (DNET) ve Fokal kortikal displazi (FCD)

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Tedavi ve yaklaşım

DNET tedavisinin temel amacı epileptik ataklar ve bunların neden olduğu hasarı önlemektir. Tedavide seçenek cerrahi eksizyon olup, cerrahi tedavi kararı hastanın kliniğine göre verilir. Cerrahi yapılması planlanan olgularda, eşlik edebilecek kortikal displazilerin de değerlendirilmesi tedavi sonrasında ataksız bir yaşam için önem arz etmektedir.(15)

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BÖLÜM

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İNTRAAKSİYEL İNFRATENTORİAL BÖLGE TÜMÖRLERİ

Ezel YALTIRIK BİLGİN¹

Vaka 1: Medulloblastom.

Vaka 2: Piloitik astrositom

Vaka 3: Hemanjioblastom

Vaka 4: Ependimom

Vaka 5: Orta hat gliomu

Vaka 6: Akciğer küçük hücreli karsinomu, soliter beyin metastazı

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TEDAVİYE İKİNCİL DEĞİŞİKLİKLER

Okan DİLEK1

Vaka 1: Radyasyon Lökoensefalopatisi

Vaka 2: Radyasyon Nekrozu

Vaka 3: Psödoprogresyon

Vaka 4: Korpus Kallozum Splenium Toksitesi

Vaka 5: Ovaryan Stimülüs Tedavisine Sekonder Tromboembolik iskemi

Tedavi ve yaklaşım

OHSS sekonder enfarkt etiyolojisine göre tedavi planlaması yapılır. Venöz tromboza bağlı bir durum varsa antikoagulan tedavi hemen başlanılmıdır. Arteriyel kaynaklı enfarkt var ise trombolitik tedavi başlanılmıdır.

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BÖLÜM

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SİNONAZAL KAVİTE HASTALIKLARI

*İsa ÇAM¹
Ural KOÇ²*

Vaka 1: Olfaktör Nöroblastom (Estezionöroblastom)

Vaka 2: Nazoetmoidal ensefalosel

Vaka 3: Paranasal-orbital osteom

Vaka 4: Maksiller sinüs adenokarsinomu

Vaka 5: Akut invaziv fungal sinüzit

Vaka 6: Sinonazal Diffüz Büyük B hücreli lenfoma

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BÖLÜM

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PİTÜİTER BEZ PATOLOJİLER

Ali Can YALÇIN¹

Vaka 1: Hipofizer mikroadenom

Vaka 2: Hipofizer makroadenom

Vaka 3: Hipofizer hiperplazi

Vaka 4: Hipofizer apopleksi

Vaka 5: Rathke kleft kisti

Vaka 6: Empty sella

Vaka 7: Kavernöz sinüs menenjiomu

Vaka 8: Kraniofarenjiom

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BÖLÜM

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ORBİTA VE GÖRME YOLLARI PATOLOJİLERİ

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Anıl KAYA²*

Vaka 1: Primer Uveal Melanoma

Vaka 2: Preseptal orbital ve periorbital selülit

Vaka 3: Postseptal orbital selülit

Vaka 4: Optik gliom

Vaka 5: Fitizis Bulbi

Vaka 6: Dakriyosistitis

Vaka 7: Fibröz Displazi

Vaka 8: Retinoblastom

Vaka 9: Tiroid Oftalmopati

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Kaslarda izlenen volüm artışının derecesi hastalığın tutulumu ve optik sinir sıkışmasının derecesi ile ilişkilidir(17). Patoloji nedeniyle orbital apeks düzeyinde sinir sıkışması optik nöropati ile sonuçlanır. MRG inceleme daha yüksek yumuşak doku kontrastı ve çoklu planların olması nedeniyle BT' de tanımlanan bulguları daha demonstratif şekilde ortaya koyar. T1 ağırlıklı görüntülerde tutulum diğer kas dokularla izointens, yağ infiltrasyonu derecesine bağlı bazen yağ baskısız sekanslarda yüksek sinyalli izlenmektedir. T2 ağırlıklı görüntülerde enflamasyona sekonder ekstraoküler kaslarda sinyal artışı izlenir. Postkontrast görüntülerde ise kontrast madde artışı gözlenir.

Tuzaklar

Radyolojik ayırıcı tanısında, ekstraoküler kasların tendonlarının tutulmaması ve simetrik tutulum olması nedeniyle diğer orbital tutulumlu patolojilerden kolayca ayırımı yapılabilir.

Tedavi ve Yaklaşım

Hastalık genellikle 2-5 yılda kendini sınırlar. Tedavi genellikle, kornea ülseri, kozmetik sorunlar ve optik sinir kompresyonu için yapılır. Tedavide hasta durumuna göre steroid ile medikal tedavi, radyoterapi veya cerrahi dekompresyon yapılır.

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BÖLÜM

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TEMPORAL KEMİK VE KULAK PATOLOJİLERİ

Fuldem MUTLU¹

Vaka 1: Akut Otitis Media

Vaka 2: Kolesteatom

Vaka 3: Malign Otitis Eksterna (Nekrotizan Otitis Eksterna)

Vaka 4: Eksternal Akustik Duktusta Yabancı Cisim

Vaka 5: Temporal Kemik Transvers Fraktürü-Fasial Sinir Hasarı

Vaka 6: Vestibüler Schwannoma

Vaka 7: Superior Semisirküler Kanal Dehissansı

Vaka 8: Genişlemiş Vestibüler Akuadukt Sendromu

Vaka 9: Mastoidektomi Kavitesinde Transplante Yağ Doku

Vaka 10: Juguler Foramen Menenjiomu

Schwannomlar ile karşılaştırıldığında daha düşük T2 sinyal intensitesi gösterirler ve prekontrast BT imajlarda daha yüksek attenuasyondadırlar. Kalsifikasyon olması ve diploik mesafede infiltrasyon görülmesi menenjiomlar için güvenilir bulgulardır. Paraganglioma kemik destrüksiyonu, sinir kılıfı tümörleri foramen-de skalloping yaparken menenjiomlar karakteristik hiperosteoze yaparlar (33).

Tedavi ve yaklaşım

Multidisipliner yaklaşımla cerrahi olarak çıkartılır.

Anahtar Kelimeler

Otitis media, Kolesteatom, Hemotimpanium, Kulak ağrısı, Kulak akıntısı, Radyoloji, YRBT, Otomastoidit, Sekretuar otit, BT, MR, Serumen, Difüzyon MR, Nonechoplanar Single Shot Turbo Spin Echo (SSTSE), Ossiküler erozyon, Malign otitis eksterna, Nekrotizan otitis eksterna, Yabancı cisim, İletim tipi işitme kaybı, Sensörinöral, Eksternal akustik kanal, Temporal kemik, Longitudinal fraktür, Transvers fraktür, Fasial kanal, Otik kapsül, Travma, Petröz apeks, Psödofraktür, Pöschl, Stenvers, Ossiküler dislokasyon, Fasial sinir hasarı, BT Angiografi, MR Angiografi, Vestibüler schwannoma, MR sisternografi, Superior semisirküler kanal dehissansı, Tullio fenomeni, Progresif işitme kaybı, Genişlemiş vestibüler aquadukt, Postoperatif görüntüleme, Transplante yağ, Rezidüel/Rekürren kolesteatom, Non EPI/DW, Paraganglioma, Juguler foramen, Menenjiom, Sentrifugal büyüme, En plaque, Mastoidektomi, Timpanik membran, ADC, Serebellopontin bölge, İntrameatal, Ekstrameatal, Tegmen timpani, Enfeksiyon, Labirintis ossifikans

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SUPRAHYOİD BOYUN PATOLOJİLERİ

Özgür ÇAKIR¹

Vaka 1: Pleomorfik adenom

Vaka 2: Sağ parafarengeal alandan intrakranial alana uzanan apse

Vaka 3: Glomus karotikum

Vaka 4: Odontojenik miksonoma

Vaka 5: Submandibuler gland sialolitiazis

Vaka 6: Brankial kleft kleft kisti

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BÖLÜM

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KONJENİTAL SPİNAL ANOMALİLER

*Büşra ŞEKER¹
Maksude Esra Kadiođlu*

Vaka 1: Dorsal dermal sinüs

Vaka 2: Nöroenterik kist

Vaka 3: Diastometamiyeli Tip 1

Vaka 4: Anterior sakral meningesel

Vaka 5: Kaudal regresyon sendromu Tip 2

Vaka 6: Tip 3 Sakrokoksigeal teratom

Vaka 7: Ventrikülis terminalis

Vaka 8: Gergin kord sendromu, terminal lipom ve epidermoid kist

Vaka 9: Lipomiyelomeningesel

Vaka 10: Vertebra formasyon kusuru, hemivertebra, kelebek vertebra

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Önemli Noktalar

Vertebra formasyon kusuru ya da vertebral segmentasyon ve füzyon anomalisi vertebral formasyonun parsiyel veya komplet yetersizliği sonucu oluşmaktadır. Parsiyel formasyon yetersizliği sonucu kama şeklinde vertebra, komplet formasyon yetersizliği sonucu ise vertebra aplazisi, hemivertebra, kelebek vertebra anomalileri oluşmaktadır (12). En sık torakolomber bölgede görülmektedir. Deforme vertebra korpuslarını içeren keskin açılı, tek eğrili veya fokal skolyoz izlenmektedir. Görüntüleme radyografi vertebral anomaliler ve eşlik eden skolyozu (dengeli-dengesiz) değerlendirmede faydalıdır. BT'de sagittal ve koronal vertebral yarıklarının, hemivertebra ve kelebek vertebranın aksiyel planda değerlendirilmesi güç olabileceği için sagittal ve koronal reformatların oluşturulması faydalı olacaktır. Posterior elemanların disrafizmi ise en iyi aksiyel planda değerlendirilmektedir. Üç boyutlu BT reformat görüntüleri ise cerrahi planlama için faydalıdır. MR ise vertebra anomalileri yanı sıra eşlik eden spinal kord anomalilerinin değerlendirilmesinde faydalıdır. Vertebra korpus anomaliler agenezi, hemivertebra (dorsal-ventral), koronal kleft, kelebek vertebra, blok vertebra ve hipoplastik vertebra olarak sınıflandırılabilir. Klinik olarak hastalar asemptomatik olabilir ya da g bağlı semptomatik hale gelebilmektedirler (4).

Tuzaklar

Eşlik edebilecek sendromlar (VATER/VACTERL, Aicardi sendromu, kleidokranial dizostozis, kloakal ekstrofi, Goldenhar sendromu, Gorlin sendromu, Jarcho-Levin sendromu, Juberg-Hayward sendromu, Klipper Feil sendromu, OEIS kompleksi, Alagille sendromu) ve bunların görüntüleme bulguları raporlamada dikkate alınmalıdır (4).

Tedavi ve yaklaşım:

Hafif vakalarda hastalar takip edilmektedir. Orta ve ağır vakalarda ise kifoskolyoz gelişimini durdurmak için cerrahi füzyon yapılmaktadır (4).

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BÖLÜM

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SPİNAL BÖLGE ENFLAMMATUAR ve ENFEKSİYÖZ DURUMLAR

Fatma CAN¹

Vaka 1: Transvers miyelit

Vaka 2: Multipl Skleroz

Vaka 3: Guillan-Barre Sendromu

Vaka 4: Araknoidit (Tip2) (adeziv araknoidit)

Vaka 5: Enfeksiyöz diskrit

Vaka 6: Epidural Apse

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alanda kitle etkisi ve homojen kontrastlanma gösterirler. Kitle etkisi apseye göre daha azdır (19).

Tedavi ve yaklaşım

Tedavi acil cerrahi drenaj ve dekompresyonu içerir. Etken patojen izole edile- ne kadar geniş spektrumlu antibiyotikler kullanılabilir (18).

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Eray ATLI¹

- Vaka 1:** İntervertebral disk dejenerasyonu, yükseklik kaybı ve disk dehidrasyonu, anüler fissür
- Vaka 2:** Posteriyör santral, sağ paramediyal-subartiküler ve sol ekstraforaminal (far lateral) protrüzyon.
- Vaka 3:** Foraminal ekstrüzyon ve sekestrasyon, sinir köküne foramende bası.
- Vaka 4:** İntervertebral diskte taşma, schmorl nodülü, intervertebral disklerde dehidrasyon ve yükseklik kaybı.
- Vaka 5:** Osteofit, disk osteofit kompleksleri, skleroz (Modik tip III değişiklik), schmorl nodülü, dejeneratif değişiklikler, santral kanalda darlık, longitudinal ligamanda ossifikasyon (diffüz idiyopatik iskelet hiperostozis, DISH).
- Vaka 6:** Modik tip I değişikliği.
- Vaka 7:** Modik tip I ve II değişiklik.
- Vaka 8:** Modik tip III değişikliği ve Dejeneratif değişiklikler
- Vaka 9:** Faset eklemden artropati, foraminal daralma, sağ faset eklem ile ilişkili kist.
- Vaka 10:** Servikal unkovertebral dejeneratif artropati, disk aralığında daralma, faset artropati, anterolistezis.
- Vaka 11:** Dejenerasyon zemininde edinilmiş spinal darlık
- Vaka 12:** Anterolistezis, dejenerasyon, osteofit, pars interartikularis defekti (spondilolizis), lomber lordozda düzleşme, retrolistezis, anterolistezis,
- Vaka 13:** Anterolistezis, osteofit, pars interartikularis defekti (spondilolizis), anterolistezis

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Anahtar Kelimeler

İntervertebral disk dejenerasyonu, intervertebral disk dehidrasyonu, anüler fissür, disk herniyasyonu, protrüzyon, ekstrüzyon, disk taşması(bulging), dejeneratif omurga hastalığı, osteofit, disk osteofit kompleksi, stabilite, instabilite, endplate ve subkondral kemik iliği sinyal değişiklikleri, Modik tip I, vertebra korpusunda subkondral enflamasyon, Modik tip II, vertebra korpusunda subkondral yağlanma, Modik tip III, vertebra korpusunda subkondral skleroz, faset artropatisi, faset kisti, faset eklemde osseöz hipertrofi, unkovertebral eklem dejenerasyonu, foraminal stenoz, edinilmiş spinal kanal darlığı, spondilolistezis, anterolistezis, retrolistezis, spondilozis, pars interartikularis defekti

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