

Bölüm 30

KAROTİS CISİM TÜMÖRLERİ

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GİRİŞ

Baş boyun tümörlerinin yaklaşık olarak % 0,5 ini oluşturan (1) karotis cisim tümörü ilk defa İsviçreli Albrecht von Haller tarafından 1743 yılında tanımlanmıştır (2). Yavaş büyümeye eğiliminde olan ve genellikle asemptomatik olan karotis cisim tümörleri nadir tümörler arasında sayılmaktadır. 1971 yılında yaklaşık 500 vaka sunulmuş olmasına karşın 1980 lerde bu sayı bin vakaya ulaşmıştır (3). 1950 yılında Monro karotis cisim tümörünün tanısını koymaktaki ilk adının bu tümörden şüphelenmek olduğunu söylemiştir (4). R.M. Mulligan 1950 yılında damar duvarındaki adventisyadan köken alan parasempatik sinir sistemiyle ilişkili olan kemoreseptör hücrelerin bir neoplazmi olduğunu tanımlamıştır. Bu tümörün ek-toderme yerleşik nöral krestten köken aldığıını bildirmiştir (2).

Nadir görülen bir tümör olması nedeniyle süregelen dönemde etiyoloji, klinik, biyolojik davranışları, hormonal etkileri ve malignite özelliklerini tam olarak değerlendirememiştir. Scudder 1903 yılında ilk başarılı carotis cisim tümör cerrahisi vakasını yayımlamıştır (5). Cerrahi olarak gelişen süreçte önemli yol alınsa da birçok konu tartışılmaya devam etmektedir.

EPİDEMİYOLOJİ VE ETİYOLOJİ

Baş ve boyunun paragangiomaları nadir görülen vasküler tümörlerdir. Nöral krestten köken alan ve sempatik zincir boyunca uzanan otonomik ganglionlarla ilişkili olan paraganrialardan oluşur(6,7).Baş boyun tümörlerinin yaklaşık %0,5 ini oluşturan karotis cisim tümörleri oldukça nadir tümörlerdir ve tüm vücut tümörlerinin %0,012 sini oluştururken insidansı 100000 de birdir(8). Sıklıkla 45-54

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