

BÖLÜM 18

Diferansiye Tiroid Kanseri Radyoaktif İyot Tedavisinde Güncel Yaklaşım

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Özet

Radyoaktif I^{131} (RAI), bilinen ilk teranostik ajan olup yaklaşık 70 yıldır diferansiye tiroid kanseri (DTK)'nin tedavisinde sağ kalımı iyileştirmek, persistan hastalık veya rekürrense bağlı morbidite riskini azaltmak için kullanılmaktadır. Gelişen tanı yöntemleri ve değişen hastalık insidansı nedeniyle DTK tedavi yönetimi gibi radyoaktif iyot tedavisi (RAIT) prensipleri de güncellenmeye devam etmektedir. Dinamik risk belirleme süreci temelindeki güncel yaklaşımda, RAIT'in gerekli hastaya, gereken zamanda ve dozajda uygulanması öne çıkmaktadır. Buna paralel olarak Endokrinoloji ve Nükleer Tıp alanındaki birçok saygın kuruluş düşük riskli hasta grubunda RAIT uygulamasını kısıtlayacak önerilerde bulunmuştur. Ayrıca radyoprotektif bu yaklaşım; kümülatif dozu yüksek olan, yaygın metastatik hastalık nedeniyle yüksek dozaj RAIT endikasyonu olan veya radyasyona maruz kalma açısından hassas olan çocuk hastalarda kantitatif değerlendirme ve dozimetrik çalışmaları da hızlandırmıştır. Mevcut önerilerin hastalık prognozuna ve toplam sağ kalıma etkisini belirlemek için planlanan prospektif randomize klinik çalışmaların bazıları sonuçlanmış, büyük bir kısmı ise devam etmektedir. DTK'nın yavaş seyirli karakteri nedeniyle uzun yıllar takip gerekse de, klinik çalışmaların erken sonuçlarına göre, rekürrens riski yüksek ve/veya metastatik olan hasta grubu dışında RAIT seçici olarak uygulanmalıdır. Bu bölümde Endokrinoloji ve Nükleer Tıp alanındaki güncel kılavuzlar ışığında DTK'da RAIT için hasta seçimi, dozimetri ve tedavi prensipleri özetlenmektedir. Mevcut önerilere ait deneyimler ve veriye dayalı yeni kanıtlar sayesinde bu alanda yeni güncellemeler olması kaçınılmazdır. Bu nedenle, belirtilen prensipler kişiselleştirilmiş hasta takibinde öneri niteliğindedir.

Tedavi Sonrası Takip

RAIT sonrası 3-7 günde yapılan tüm vücut tarama sintigrafisi (TST) tedavi etkinliğini göstermenin yanında bakiye tiroid dokusunu ve metastazları belirlemede duyarlı bir incelemedir. Tanısal TVIT'ye göre daha yüksek dozaj ile yapıldığından daha önce tespit edilemeyen fonksiyonel tiroid dokuları saptanabilir ve teranostik yaklaşım ile RAIT'e yanıt vereceği öngörülebilir. Son yıllarda yaygınlaşan hibrid SPECT/CT gama kameraları sayesinde lezyonların üç boyutlu olarak görüntülenmesi ve anatomik lokalizasyonu da mümkün olmaktadır. RAIT etkinliğini değerlendirmek için serum Tg ve AntiTg düzeyleri, boyun US ve gerekli görülürse 6-8 ay sonra TVIT yapılır. Dinamik risk belirleme sürecinde tüm bu sonuçlara göre; mükemmel yanıt, biyokimyasal yetersiz yanıt, yapısal yetersiz yanıt ve belirsiz yanıt veren hastalara uygun takip protokolü oluşturulur. DTK yavaşı

ilerleyen ve nispeten yüksek rekürrens oranlı bir malignite olduğundan bu yüzden bu hastalarda ömür boyu takip önerilmektedir. (8,10,19,101).

RAIT sonrası persistan hastalıkta, rekürrens ve metastaz varlığında RAI tutulumu varsa tekrarlayan tedaviler yapılabilir. Genel olarak, yüksek evreli hastalık (evre III ve IV), 45-55 üstü yaş ve agresif histolojik alt tip içeren tümörlerde daha fazla doz gereklidir. RAI direnci gelişmedikçe tam remisyon sağlanana kadar tedaviye devam edilebilir. Maksimum güvenli kümülatif doz konusunda belirgin bir eşik değer belirlemek güçtür. 1Ci 'ye kadar tedaviye devam etmeyi destekleyen araştırmalar olsa >45 yaş hastalarda 600 mCi, <45 yaş hastalarda ise 800 mCi kümülatif doza ulaşan hastalarda RAIT'in riskleri ve hayat kalitesi üzerine etkisi dikkate alınarak diğer tedaviler (cerrahi eksizezyon, sistemik kemoterapi, kemoembolizasyon ve moleküler hedef tedavi gibi) açısından vaka bazında değerlendirme yapılmalıdır (51,56,97,102).

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