

Bölüm 6

GEBELİKTE ÜRİNER SİSTEM ENFEKSİYONLARI

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GİRİŞ

Üriner sistem enfeksiyonları (İYE), anemiden sonra hamile kadınlarda görülen en yaygın ikinci patolojidir. Ayrıca hamile kadınlarda en sık görülen enfeksiyondur⁽¹⁾. Üretranın erkeklerde göre daha kısa olması, anüsün vajinaya yakın olması ve cinsel aktivite ile patojen mikroorganizmaların daha kolay etkilemesi kadınlarda İYE nin daha sık görülmesine neden olur⁽²⁾. Üriner sistem enfeksiyonları, gebelikte maternal ve neonatal morbidite ve mortaliteyi arttıran en yaygın bakteriyel enfeksiyon olarak kabul edilmektedir⁽³⁾.

Hamilelikleri sırasında kadınlarda İYE prevalansı % 5-10 civarıdır. Hamile kadınların başvurularının % 5'inin sebebinin İYE olduğu tahmin edilmektedir. Üriner sistem enfeksiyonları tedavi edilmezse, üriner sistemin gebeliğe bağlı adaptif değişiklikleri sonucu sekonder piyelonefrit gelişme riskini artırabilir. Bu da erken doğum, düşük doğum ağırlığı, maternal sistemik enfeksiyon gibi komplikasyonlara neden olabilir⁽⁴⁾.

Gebelik sırasında idrar yolu enfeksiyonları, asemptomatik bakterüri (ASB), alt üriner sistem enfeksiyonları (sistit) veya üst üriner sistem enfeksiyonları (piyelonefrit) olarak sınıflandırılabilir.

1. GEBELİKTE ÜRİNER SİSTEMDE GÖRÜLEN DEĞİŞİKLİKLER

Hamilelikte üriner sistemde meydana gelen hormonal ve anatomik değişiklikler kadınları enfeksiyona daha duyarlı hale getirir⁽⁵⁾. Bu değişiklikler nedeniyle ASB ve sistit varlığında, piyelonefrit gelişme riskinin arttığı düşünülmektedir⁽⁶⁻⁷⁾. Gebelik sırasında böbreğin uzunluğu yaklaşık 1 cm ve glomerüler filtrasyon hızı % 30-50 artar⁽⁸⁾. Gebeliğin yedinci haftasından itibaren hidroüreteronefroz gözlenebilir⁽⁹⁾. Bu durum, üreterlerdeki azalmış peristaltizme, progesteronun kas gevşetici etkilerine ve uterusun giderek artan basisına bağlanabilir^(8,9,10,11). Ayrıca hamilelik ilerledikçe mesane üste ve öne doğru yer değiştirir, hipertrofi ve düz kas

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(⁵³). Nitrofurantoin idrarda terapötik seviyelere ulaşır, ancak dokuda yeterli düzeyde değildir. Bu nedenle piyelonefrit tedavisinde kullanılmamalıdır (¹⁰). Ancak alt üriner sistem enfeksiyonlarında nitrofurantoin, proteus, serratia veya pseudomonasa karşı aktivitesi olmamasına rağmen, E coli, Enterococcus faecalis, Klebsiella spp, Staphylococcus saprophyticus ve Staphylococcus aureus dahil birçok yaygın üropatojenlere karşı sınırlı direnç ve bakterisidal aktivite nedeniyle yararlıdır (^{54,55}). Yapılan bir çalışmada nitrofurantoin ile anoftalmi, mikroftalmi, atriyal septal kusurlar, yarık dudak ve damak gibi konjenital malformasyonlar arasında bir ilişki olduğu gösterilmiştir (⁵⁶). Klindamisin, Staphylococcus ve GBS'nin neden olduğu enfeksiyonların tedavisi için penisiline alerjisi olan hastalarda kullanılsa da Enterococcus ve çoğu aerobik gram-negatif bakteriye karşı etkisizdir (¹⁰). Klindamisin kullanan hastaların yaklaşık %10'unda ishal ve nadiren psödomembranöz kolit görülebilir, ancak yapılan çalışmalarda klindamisini teratojenite ile ilişkilendiren veri yoktur (^{29,30,54,57,58}). Aminoglikozidler gram negatif basillerin tedavisinde etkilidir. Genellikle ampicilin ile birlikte piyelonefrit tedavisinde önerilir (^{30,53}). Ancak hem anne hem de fetüste ototoksisite ve nefrotoksisite riski mevcuttur (^{29,30}). Gentamisin, nispeten üstün güvenlik profili ve düşük maliyeti nedeniyle gerekirse obstetrikte tercih edilen bir aminoglikoziddir (⁵⁴). Potansiyel risklere rağmen, gebelikte gentamisin uygulamasından sonra teratojenite, neonatal ototoksisite veya nefrotoksisite bildirilmemiştir ve gentamisin yaygın olarak kullanılmaktadır (^{30,57}).

SONUÇ

İdrar yolu enfeksiyonları gebelikte en sık görülen bakteriyel enfeksiyondur. Aynı zamanda, maternal ve neonatal morbidite ve mortalite riskini arttırmır. Bu nedenle ASB, sistit ve piyelonefrit tedavi edilmeli ve hastalar tedaviden sonra antimikrobiyal profilaksiye dikkat edilerek yakından takip edilmelidir.

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