

Lupus Nephritis and Pregnancy: Case Report

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Our aim is to report of a pregnant woman with lupus nephritis; A 33-year-old multigravida (gravida:5, parity:4) with an intrauterine pregnancy at 34 weeks of pregnant woman with 5 years of uncontrolled SLE.

Pregnancy can be successful in most women with lupus nephritis. Pregnancy in SLE should be planned and a management strategy should be agreed in full consultation with the patient, prior to conception. Women with SLE frequently need treatment throughout pregnancy. It is essential that the maternal disease is well controlled prior to, during and after pregnancy to ensure the best possible outcome for the mother and child.

Key Words: SLE, Pregnancy, Nephritis, Preterm Labor

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Introduction

Lupus is an chronic inflammatory and multisystem disease which damages the tissues and cells by autoimmunes and immune complexes.¹ By the years, understanding the mechanism and multidisciplinary approach to therapy of lupus have led to better pregnancy outcome.² The management of pregnancy in SLE should start before conception so as to optimize maternal health. The disease is not in itself a contra-indication to pregnancy, but the relations between SLE and pregnancy found high fetal and maternal risks notably, when pregnancy occurs in active SLE. SLE tends to flare during pregnancy and the puerperium. Maternal flares are associated with increased prematurity,³ and active nephritis has been shown to be an independent factor for fetal mortality.⁴ Nephritis is known to be one of the most serious complications of SLE and a strong predictor of poor outcome. Prophylactic steroids are a choice in management but There is no evidence that prophylactic steroids lower the frequency of flares, and there are significant adverse effects during pregnancy: premature rupture of membranes; infections; intra-uterine growth restriction; hypertension; gestational diabetes; osteoporosis; and avascular necrosis.⁵

The purpose of this case report is to inform a pregnant woman with uncontrolled lupus nephritis.

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Case Report

A 33-year-old multigravida (gravida:5, parite: 4) with an intrauterine pregnancy at 34 weeks of pregnant woman with 5 years of uncontrolled SLE. Her family reported that she has never used her therapy orderly. We noted during her physical examination that; she had photosensitivity, bilateral lower extremity edema, malar rash, discoid rash and artralgia. Her vital signs were temperature 36.8°C, blood pressure 190/120 mmHg, pulse 92 beats/min, respiratory rate 18/min. The examination of heart, chest, lymph node, neurological system were unremarkable. Initial laboratory values were WBC 7,85 K/UL, Hb 9.7 g/dl, Hct 26,9%, mean corpuscular volume 87,2 fL, platelet 244.000 K/UL, reticulocytes 3,2%. CRP 4,68 mg/dl. Serum blood urea nitrogen 29 mg/dl, creatine 0.92mg/dl, potassium 4.1 mEq/l, aspartate aminotransferase 23 U/l, alanin aminotransferase 15 U/l, total bilirubine 0.2 mg/dl, lactic dehydrogenase 226 U/l, albumin 0,97 g/dl, calcium 7.5 mg/dl, vitamin B12 199 pg/ml, folate 7 ng/ml. Protrombin time, activated partial protrombin time and fibrinogen levels were in normal limits. ANTI-SSA, ANTI Histones, ANTI Nucleosomes, ANTI ds DNA were positive and C3 and C4 levels were decreased. Anti-cardiolipin antibody-M, anti-cardiolipin antibody-G, ANTI-ENA, ANTI SCL 70, ANTI JO 1 were found negative. Urinalysis showed abundant erythrocytes and 5–10 leucocytes per high power field and 24 h urinalysis showed 9 g/day of proteinuria. Ultrasonography of abdomen was unremarkable. The patient was given 32mg/day of methylprednisolon by orally. Fetal heart tones were noted to be 136 bpm with reassuring variability. Uterine contractions were noted every 5 minutes and initial sterile vaginal exam indicated cervical dilation of 2 cm with 30% effacement. The fetus was vertex in presentation with a size consistent for stated gestational age and a normal amniotic fluid index. We

began orally nippedhipine as tocolytic therapy to the patient. Then we consulted the patient with nephrology. After an evaluation we decided that the patient is at the acute phase of lupus. Then we stopped tocolytic therapy and took the patient into delivery room. We gave oxytocin induction (10 units in 1000 cc %5 DRL and 0.01 u/min) to her and after 6 hours of delivery she gave a 2400 gr, 44 cm, 6-8 APGAR of fetus.

Discussion

A woman with SLE who wants to become pregnant faces a number of risks to her health and that of her unborn child. A severe disease flare may be potentially life threatening, while some of the drug therapies are teratogenic and fetotoxic. The risks involved may be minimized by appropriate timing of pregnancy and optimization of therapy prior to conception. After the development in the management and better understanding the mechanism of lupus nephritis, the pregnant women have successful maternal and fetal outcome.⁶ Pregnancy in SLE should be planned and a management strategy should be agreed in full consultation with the patient, prior to conception. Women with SLE frequently need treatment throughout pregnancy. Pregnancies in SLE have high risks of spontaneous abortion, premature delivery, and stillbirth,⁷ and also our case had premature delivery. Mok et al, reported that proteinuria is an important factor that causes fetal loss,⁸ our case also had proteinuria but she had a successful delivery and had a healthy baby. In other two of the studies, the predictors for bad maternal and fetal outcomes were, SLE activity at pregnancy onset, severity of maternal renal disease, the presence of hypertension or lupus anticoagulant,^{9,10} although our case had all of these predictors she didn't have bad maternal or fetal outcome. Although most authorities recommend continuation of immunosuppressive therapy during pregnancy in women with nephritis, it is not clear whether the dosage should be increased peripartum. While SLE is affecting the pregnancy, the pregnancy could worsen the SLE. We think that SLE is not a contraindication for pregnancy but the patients should be followed up by an obstetrician, pediatrician and nephrologist. And also the patients must be informed about the pregnancy progress in SLE.

Lupus Nefriti ve Gebelik: Olgu Sunumu

Amacımız gebe lupus nefriti olan olgunun sunulması; 33 yaşında multigravide (gravida :5, parite:4) , 34 haftalık intra-

uterine gebeliği olan ve 5 yıldır kontrolsüz SLE hastalığı olan bayan hasta.

Lupus nefriti olan kadınlarda gebelik başarılı olabilir. SLE'de gebelik ve gebelikte takip ve tedavi stratejisi, konsepsiyon öncesinde yapılmalıdır. SLE'li kadınlarda tedavi tüm gebelik boyunca devam etmelidir. En iyi maternal ve fetal sonuçlar için maternal hastalığın gebelik öncesinde, gebelik süresince ve sonrasında sıkı ve iyi kontrol altında olması gerekmektedir.

Anahtar Kelimeler: SLE, Gebelik, Nefrit, Preterm Eylem

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