Antenatal follow-up, anesthesia management and perinatal outcomes in pregnancy with renal transplant

Renal transplantlı gebelerin antenatal takibi, anestezi yönetimi ve perinatal sonuçları

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Abstract

Objective: Due to the recent increase in the successful pregnancies after renal transplant, the number of renal transplant recipients having vaginal or cesarean delivery possibly associated with high maternal, fetal and/or neonatal risk requiring team approach increased. We aimed to evaluate antenatal follow-up, perinatal outcomes, and anesthesia management in pregnancies with renal transplantation and to compare them with the current literature.

Materials and Methods: After ethics committee approval, renal transplant recipients who gave birth in our hospital between January 2010 and December 2019 were documented in this retrospective study. Demographic characteristics, comorbidities, antenatal follow-up, anesthesia management, and maternal, fetal, and neonatal outcomes were presented.

Results: A total of 20 pregnant women who underwent renal transplant were identified. The mean age of the parturients was 31±5 years. The median interval from transplantation to conception was 8.15±4.8 years. Antenatal mean serum creatinine level and proteinuria were 1.48±1.39 mg/dL and 1.397±1.316 mg/dL, respectively. No allograft rejection was recorded. Comorbidities including hypertension (n=12), preeclampsia (n=6), and preterm delivery (n=10) were noted. The median gestational age was 35±3 weeks and the median newborn weight was 2.520±832 gram. There was one abortion, two pregnancy terminations, and 17 deliveries (3 vaginal and 14 cesareans). Cesarean sections (11/14; 78.6%) were mostly performed under spinal block and general anesthesia was performed in three (21.4%) women. Epidural analgesia for vaginal delivery was recorded in one parturient.

Conclusion: Despite the presence of preterm delivery and comorbidities, antenatal/peripartum follow-up and analgesia/anesthesia management of renal transplant recipients revealed good perinatal outcomes.

Keywords: Anesthesia, spinal, general, surgery, cesarean/labor and delivery, renal transplantation

Öz

Amaç: Son zamanlarda böbrek nakli sonrası başarılı gebeliklerin artması nedeniyle, yüksek maternal, fetal ve/veya neonatal risk ile ilişkili olarak ekip yaklaşımı gerektiren vajinal veya sezaryen doğum olan böbrek transplant alıcıları artmıştır. Renal transplantasyonlu gebelerin antenatal takibi, perinatal sonuçları ve anestezi yönetimi güncel literatürle karşılaştırarak bunun değerlendirilmesi amaçlandı.

Gereç ve Yöntemler: Etiq kurul onayından sonra, Ocak 2010 - Aralık 2019 tarihleri arasında hastanemizde doğum yapan böbrek nakli alıcıları bu retrospektif çalışmada belgelendi. Demografik özellikler, komorbiditeler, antenatal takip, anestezi yönetimi, maternal, fetal ve neonatal sonuçlar sunuldu.

Bulgular: Böbrek nakli yapılan toplam 20 gebe belirlendi. Doğum yapanların ortalaması yaş 31±5 yıldı. Transplantasyondan gebe kalmaya kadar geçen medyan aralık 8.15±4.8 yıldı. Antenatal ortalamal serum kreatinin düzeyi ve proteinürü sırasıyla 1.48±1.39 mg/dL ve 1.397±1.316 mg/dL idi. Allograft reddi izlenmedi. Hipertansiyon (n=12), preeklampsia (n=6) ve erken doğum (n=10) gibi komorbiditeler izlenildi. Ortanca gebelik yaş 35±3 hafta ve ortanca yeni doğan ağırlığı 2.520±832 gramdı. Bir dilatasyon kuretaj, 2 gebelik terminasyonu ve 17 doğum (3 vajinal ve 14 sezaryen) vardı. Sezaryenlerin çoğunlukla 11/14 (%78,6%) spinal anestezi altında yapıldı ve üç (%21,4) gebeye genel anestezi uygulandı. Doğum eylemini ve vajinal doğum için epidural analjezi 1

Conclusion: Despite the presence of preterm delivery and comorbidities, antenatal/peripartum follow-up and analgesia/anesthesia management of renal transplant recipients revealed good perinatal outcomes.

Keywords: Anesthesia, spinal, general, surgery, cesarean/labor and delivery, renal transplantation

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Introduction

After the first successful pregnancy in a renal transplant recipient in 1958(1), a recent meta-analysis including 4,706 pregnancies among 3,570 renal transplant recipients reported a live birth rate of 73.5%(2). Pregnancy after renal transplantation carries maternal (e.g. allograft rejection, infection, hypertension or preeclampsia), fetal (spontaneous abortion, premature delivery, fetal growth restriction, and intrauterine fetal death) and neonatal (recurrence risk of the underlying maternal delivery, fetal growth restriction, and intrauterine fetal death) outcomes because this particular group of patients is under immunosuppressive therapy, which may increase infection risk(3-6). As the number of renal transplantations and parturients with renal transplant have increased, there is an anticipated high risk in perinatal outcomes because this particular group of patients is under immunosuppression treatment and has several comorbidities. Therefore, we aimed to revisit the risks and characteristics of these recipients by documenting our experience to provide updated national information for comparison with international data.

Materials and Methods

After obtaining approval from the ethics committee (decision number: 503, dated: 20.07.2020), the database of our institution was searched between January 1st, 2010, and December 31st, 2019, for renal transplants and pregnancy. We specifically identified post-renal transplant, pregnant patients. Data including maternal age, the time elapsed from renal transplant to delivery, reason for transplantation, maternal comorbidities, gestational age, gravity and parity, and immunosuppressant medication were documented. Antenatal follow-up of renal functions in terms of serum creatinine (SCr) levels and estimated glomerular filtration rate (eGFR) were recorded from the records to elucidate the incidence of allograft rejection. Antenatal or peripartum anesthesia data were the mode of delivery, analgesia or anesthesia provided, anesthesia monitoring used, and complications related to obstetrics or anesthesia. Obstetric outcomes in terms of maternal, fetal and neonatal were noted from the records. Maternal outcomes included the rate of maternal mortality, cesarean section (CS), preeclampsia and/or proteinuria (total urine protein >0.3 g/day during pregnancy), hypertension, gestational diabetes mellitus, and anemia (hemoglobin <10.5 g/dL and <11 g/dL in the 2nd and 3rd trimester, respectively). All patients’ blood pressures were under control with an antihypertensive agent. Fetal outcomes consisted of spontaneous abortion, stillbirth, preterm birth, intrauterine growth retardation, perinatal death, newborn weight, and Apgar scores (at 1 and 5 minutes).

Statistical Analysis

Statistical analysis was conducted using the SPSS version 21.0 package. After performing descriptive statistics, data were presented as number (n), rate (%), mean and standard deviation or median where appropriate. Neonatal data according to maternal eGFR groups were compared using the t-test. The number of patients with three different eGFR intervals was compared using the chi-square test. A p-value <0.05 was considered statistically significant.

Results

The demographic characteristics of parturients (as renal transplant recipients), those who underwent normal spontaneous vaginal deliveries (NSVD) or CS, are presented in Table 1. The mean maternal age was 31±5 years, the rate of nulliparity was 45%, and two patients became pregnant via in vitro fertilization. Renal transplant recipients (two of whom had two deliveries) included three NSVD, 14 cesarean deliveries, two pregnancy terminations, and one abortion. The mean interval from transplantation to conception was 8.15±4.83 years. The causes of end-stage renal failure requiring transplant and immunosuppressive regimen of the patients are presented (Table 1).

Antenatal Follow-up

Antenatal follow-up of renal outcomes is presented in Table 2. The mean SCr level in the 1st trimester was 1.48±1.39 mg/dL and the mean 1st-trimester eGFR was 78.1±8.3 mL/min/1.73 m². After delivery, the mean SCr level and eGFR were 1.42±0.22 mg/dL and 64.7±8 mL/min/1.73 m², respectively. These parameters were not different between the 1st trimester and after delivery.

Then, 1st-trimester eGFR was assigned as <60, 60-89 and ≥90 mL/min/1.73 m². The number of patients with each eGFR is presented in Table 2. The mean 24-h urine protein was 1.397±1.316 g/day; 60% of them had >0.3 g/day proteinuria. No allograft rejection records were observed in any of the recipients, either before or after pregnancy.

Perinatal Outcome

The obstetric outcomes, maternal comorbidities before pregnancy and peripartum morbidity and mode of delivery,
indications for CS, and type of anesthesia are shown in Table 3. There were 17 singleton live births, 14 of which were CS and three were NSVD. Eleven parturients received single-shot spinal anesthesia using 10 mg of hyperbaric bupivacaine with opioids (fentanyl 10 µg and morphine 100 µg). Three patients received general anesthesia. General anesthesia induction was provided using intravenous (IV) 5 mg/kg of thiopental and 1 mg/kg succinylcholine followed by 1 MAC of sevoflurane in 50% oxygen-air mixture until delivery of the newborn, and then by adding 0.2 µg/kg/h of IV remifentanil infusion. One of the NSVDs received epidural analgesia using 0.125% bupivacaine with 2 µg/mL fentanyl.

Peripartum fluid administration was 78% saline and 21% Ringer's lactate. The mean perioperative arterial pressure (MAP) ranged between 77±12 to 105±14 mm Hg. Records revealed that spinal anesthesia-induced hypotension was treated with ephedrine or noradrenaline. Post-dural puncture headache was reported in four out of 14 patients who underwent CS.

### Neonatal Outcome

The median duration of gestation was 35±3 weeks (abortion and terminations were not included). There were 10 (58%) preterm deliveries. Preterm birth indications were ablatio placenta (n=2), spontaneous preterm birth (n=3), premature rupture of the membranes (n=2), and severe preeclampsia (n=3). The median birth weight was 2,520±832 grams (10 females and 7 males), with five newborns (29%) having low birth weight and two newborns (11%) having very low birth weight. Two newborns (11%) had Apgar scores <7 at 1 minute and 1 newborn (5%) had an Apgar score <7 at 5 minutes. The mean umbilical artery pH was 7.37±0.05.

We assigned our patients into three groups according to their eGFR as <60, 60-90, >90 mL/min/1.73 m² (Table 4). Then, we compared their perinatal results including gestational week at delivery, birth weight, umbilical artery pH, and Apgar scores. In group eGFR >90 mL/min/1.73 m², we observed higher birth weight, gestational week at delivery, and 1st minute Apgar score, but there were no significant differences between the groups (p=0.580, p=0.788, and p=0.715 respectively). The neonatal intensive care unit admission rate was 41.2%; the indications were tachypnea (57.1%) and low Apgar score (28.6%).

### Discussion

We have reported antenatal follow-up, and anesthesia and/or analgesia management and perinatal outcomes in parturients with renal transplantation who underwent CS or NSVD.
Transplantation Association, pregnancy should occur at least 112 years after renal transplantation. The rate of 32.7% of preterm birth among 453 pregnant women who had undergone renal transplantation. Bramham et al. found a rate of 38.5% of preterm birth among 50 patients with renal transplantation and pregnancy, which is very much acceptable and reliable. Pregnancies after transplantation are considered as high risk because of the higher incidence of pregnancy complications such as miscarriage, preeclampsia, gestational diabetes, need for CS, and premature birth compared with otherwise healthy pregnant women. Gill et al. found a rate of 32.7% of preterm birth among 453 pregnant women who had undergone renal transplantation. Bramham et al. observed preeclampsia at a rate of 30% but no increased risk of gestational diabetes. The CS rate was 82% among 105 pregnant women with renal transplantation. Similarly, in our study, we documented gestational diabetes, preeclampsia and miscarriage, in addition to rates of CS and preterm delivery of 82.4% and 58%, respectively.

National authors have shown that maternal and fetal outcomes of renal transplant recipients were mostly dependent on transplanted kidney function determined by eGFR in the 1st trimester of pregnancy. In our study, eGFR >90 mL/min/1.73 m² were recorded in 9 out of 20 patients in the 1st trimester, whereas it was in 5 out of 17 parturients after delivery. Immunosuppressive medication including tacrolimus, azathioprine, and low-dose corticosteroids are well tolerated in pregnancy without significant risk in renal transplant recipients, but mycophenolate mofetil, sirolimus, and mammalian target of rapamycin inhibitors are contraindicated in pregnancy. Sixty percent of our patients were using a combination of tacrolimus, azathioprine, and prednisolone, and none of the chemotherapeutics that are contraindicated was documented and used in our audit. Immunosuppressive medication can manifest serious adverse effects or complications, steroids and tacrolimus may cause commonly leukopenia and rarely epidural abscess. Currently, no such serious adverse effects or complications have been documented. Regarding the use of neuraxial analgesia and/or anesthesia techniques in renal transplant recipients, obeying a strict aseptic technique is a must as stated. Although there is no particular recommendation for prophylactic antibiotic use, it is routinely administered before the skin incision in our obstetric anesthesia practice. In a retrospective analysis of a multicenter cohort study, 83 renal transplant recipients had vaginal and cesarean delivery with a mean gestation of 36±0.5 (range, 25-40) weeks. Then, 44 out of 83 (53%) parturients underwent CS; 33 out of 44 (75%) of these parturients received neuraxial anesthesia and 11 out of 44 (25%) underwent general anesthesia. The choice of neuraxial techniques includes single-shot spinal (56.8%), epidural (13.6%), and CSE anesthesia (4.6%).

Table 3. Renal recipient data, type of delivery and anesthesia (mean ± SD, range, n or %)

| Comorbidity before pregnancy (n=20) | | |
|-----------------------------------|---|---|
| Hypertension                      | 4 (20%) | |
| Type 2 DM                        | 1 (5%) | |

| Maternal data (n=17) | | |
|----------------------|---|---|
| Preeclampsia         | 6 (35%) | |
| Gestational DM       | 2 (11%) | |
| Anemia               | 12 (70%) | |
| •No treatment (n)    | 3 | |
| •IV Iron (FCM) replacement (n) | 2 | |
| •Blood product (ES) Use (n) | 7 | |

| Mode of delivery (n) | | |
|----------------------|---|---|
| CS                   | 14 (82.4%) | |
| NSVD                 | 3 (17.6%) | |

| Indication for CS | | |
|-------------------|---|---|
| Previous CS       | 7 (50%) | |
| Maternal medical indication | 3 (21.4%) | |
| Presentation abnormality | 2 (14.3%) | |
| Placental abruption | 2 (14.3%) | |

| Anesthesia type for CS (n=14) | | |
| Spinal (n=11)                | 11 (78.6%) | |
| General (n=3)                | 3 (21.4%) | |

| Epidural Analgesia for NSVD (n) | | |
|---------------------------------|---|---|
| DM: Diabetes mellitus, IV: Intravenous, FMC: Ferric carboxymaltose, ES: Erythrocyte suspension, CS: Cesarean section, NSVD: Normal spontaneous vaginal delivery, SD: Standard deviation | | |

Table 4. Neonatal data according to maternal eGFR (mL/min/1.73 m²) (mean ± SD)

| Gestation | eGFR <60 | eGFR 60-89 | eGFR≥90 | p |
|-----------|----------|------------|---------|---|
| 34.2±3.7  | 34.68±4.1| 36.67±1.8  | 0.788   | |

| Delivery (week) | | |
|-----------------|---|---|
| UA pH           | 7.3±0.1  | 7.4±0   | 7.3±0   | 0.702 | |
| Birth weight    | 2.296±0.1| 1.650±1.4| 2.506±4.76| 0.380 | |
| Apgar at 1 min  | 7.9±3    | 6.5±3.5  | 8.2±0.8  | 0.715 | |
| Apgar at 5 min  | 9±2      | 8±1.4    | 9±1      | 0.723 | |

| UA: Unbilical artery, SD: Standard deviation, Abortion and termination were not included in the average delivery week of gestation | | |

Perinatologists and anesthesiologists need to deal with those patients more frequently because the number of pregnant women who previously underwent renal transplantation has increased. According to the report of the American Transplantation Association, pregnancy should occur at least a year after renal transplantation, though highest maternal complications and the least favorable birth outcomes were observed in the <2-year interval between renal transplantation and pregnancy. In the present study, we audited an average interval of 8 years (from 2 to 19 years) between renal transplantation and pregnancy, which is very much acceptable and reliable.
center retrospective analysis including 17 deliveries (17.6% and 82.4% were NSVD and CS, respectively) at 35±3 weeks of gestation, neuraxial anesthesia was the most commonly chosen technique for CS (11 out of 14; 78.6%); the remainder (21.4%) received general anesthesia. In contrast to the multicenter study including three major tertiary hospitals in Israel[7], our CS rate was very much higher (82.4% vs 53%) due to the comorbidities and previous CS indication that required CS. Additionally, single-shot spinal anesthesia was the only technique we chose, accounting for 78.6%, and higher than the 56.8% rate reported by Ioscovich et al.[7].

According to national data related to pregnancy and renal transplantation, eight pregnancies in eight renal transplant recipients were reported between 1975 to 2003 (=28-year transplantation, eight pregnancies in eight renal transplant recipients). Herein, 17 pregnant renal transplant recipients delivered success fully with both predelivery and postdelivery good graft function during a 10-year period. Similarly, the present mean gestation time for all these parturients was 35±3 (range 28.6-40) weeks, which was comparable to previous studies[7,14]. We documented that single-shot spinal anesthesia was provided with hyperbaric bupivacaine plus opioids. The average intraoperative fluid administration in our study was 1.2-1.3 L of saline and/or lactated Ringer’s solution. In this population, adequate volume replacement has been recommended as a prerequisite to maintain placental perfusion and to avoid critical renal hypoperfusion[2]. Spinal anesthesia-induced hypotension is a known common complication, and we needed to treat only four patients (the rest were mostly hypertensive) with vasopressors to maintain maternal renal and placental perfusion. We provided a strict aseptic technique during neuraxial anesthesia practice to avoid spinal/epidural abscess or any other infectious complications because these patients are also at high risk because of their ongoing immunosuppressive therapy. In the current retrospective analysis, despite the presence of preterm delivery and comorbidities, antenatal and peripartum follow-up and analgesia/anesthesia management of parturients as renal transplant recipients were uneventful and revealed good perinatal outcomes, consistent with both international and national analyses.

**Study Limitations**

Our study has some limitations. First, it has a retrospective design and the second is that it is conducted in a single institution. Studies involving many centers and evaluating more patients will improve our knowledge about kidney transplant patients.

**Conclusion**

When we evaluated all these obstetric and neonatal data in our study, we observed that renal transplant patients have been found to have higher obstetric complications like preterm delivery than the normal population. However antenatal/peripartum follow-up and analgesia/anesthesia management of renal transplant recipients revealed good perinatal outcomes.

**Ethics**

**Ethics Committee Approval:** The local ethics committee approved the study (decision number: 503, dated: 20.07.2020).

**Informed Consent:** Retrospective study.

**Peer-review:** Externally and internally peer-reviewed.

**Authorship Contributions**

Concept: D.B.G., M.B., E.T., Design: D.B.G., M.B., E.T., Data Collection or Processing: S.K., B.B., Analysis or Interpretation: D.K., G.I., Literature Search: D.B.G., G.I., Writing: E.T., D.B.G., G.I., B.B.

**Conflict of Interest:** The authors report no conflict of interest.

**Financial Disclosure:** Authors have no financial interests about the research.

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