Perinatal Outcomes in Women with Chronic Kidney Diseases

Resultados perinatais em mulheres com doenças renais crônicas

Marcus Vinicius Pinheiro Zilli1, Anderson Borovac-Pinheiro1, Maria Laura Costa1, Fernanda Garanhani Surita1

1 Department of Obstetrics and Gynecology, Universidade Estadual de Campinas, Campinas, SP, Brazil

Rev Bras Ginecol Obstet 2022;44(12):1094–1101.

Address for correspondence Fernanda Garanhani Surita, Full Professor, Department of Obstetrics and Gynecology, Universidade Estadual de Campinas, Rua Alexander Fleming, 101, Campinas, SP 13083-881, Brazil (e-mail: surita@unicamp.br).

Abstract

Objective To assess maternal and neonatal outcomes in women with chronic kidney disease (CKD) at a referral center for high-risk pregnancy.

Methods A retrospective cohort of pregnant women with CKD was followed at the Women’s Hospital of Universidade Estadual de Campinas, Brazil, between 2012 and 2020. Variables related to disease etiology, treatment duration, sociodemographic variables, lifestyle, other associated diseases, obstetric history, and perinatal outcomes were assessed. The causes of CKD were grouped into 10 subgroups. Subsequently, we divided the sample according to gestational age at childbirth, as preterm and term births, comparing maternal and neonatal outcomes, and baseline characteristics as well as outcomes among such groups.

Results A total of 84 pregnancies were included, in 67 women with CKD. Among them, six pregnancies evolved to fetal death, five to miscarriage, and one was a twin pregnancy. We further analyzed 72 single pregnancies with live births; the mean gestational age at birth was 35 weeks and 3 days, with a mean birth weight of 2,444 g. Around half of the sample (51.39%) presented previous hypertension, and 27.7% developed preeclampsia. Among the preterm births, we observed a higher frequency of hypertensive syndromes, longer maternal intensive care unit (ICU) stay in the postpartum period, higher incidence of admission to the neonatal ICU, higher neonatal death, lower 5-minute Apgar score, and lower birth weight.

Conclusion This study demonstrates increased adverse outcomes among pregnancies complicated by CKD and expands the knowledge on obstetric care among such women in an attempt to reduce maternal risks and identify factors related to prematurity in this population.

Keywords ► kidney disease ► high-risk pregnancy ► antenatal care ► perinatal outcomes
Introduction

Chronic kidney disease (CKD) is a global health problem that affects ~ 10% of the population. The prevalence has increased in recent decades and is higher among low- and middle-income countries. In Brazil, more than ten million people have CKD. Chronic kidney disease occurs in women and men equally, and reproductive function can be affected in women, in addition to influencing maternal and neonatal outcomes.

Approximately 3 to 4% of women of reproductive age and ~ 1 to 3% of pregnant women have CKD, regardless of the underlying cause. In these patients, there is a greater risk of maternal hypertensive complications, fetal growth restriction, and premature birth; therefore, there is a greater chance of hospitalization of the newborn in a neonatal intensive care unit (ICU), stillbirth and neonatal death, in addition to morbidities related to prematurity. Women with CKD are 10 times more likely to develop preeclampsia than women at usual risk, with a reported prevalence of preeclampsia of up to 40% among pregnant women with CKD.

The reported overall prevalence of preterm birth (before 37 weeks) in Brazil is around 10 to 12%. In other countries, rates vary according to other health indicators. One of the factors that can influence this is pregestational creatinine levels. A Canadian study of 56,000 pregnancies showed an increase in therapeutic preterm birth in women with pregestational creatinine above the 95th percentile (0.87 mg/dL), which was not observed in patients with spontaneous preterm births. In this same study, a graph was constructed that illustrates a J-curve supporting the association of serum creatinine and the probability of preterm delivery, with a 1.23-fold increase in the chance of preterm delivery in patients who had some renal dysfunction compared with pregnant women with normal renal function.

Despite the possible unfavorable perinatal outcomes in pregnant women with CKD reported in the literature, there is still a lack of Brazilian studies on the subject. The aim of this study is to evaluate the maternal and perinatal outcomes of women with CKD who underwent prenatal care and delivery at a single Brazilian reference center for high-risk pregnancies, and further compare cases with preterm and term childbirth.

Methods

We performed a retrospective cohort study at the Women’s Hospital of Universidade Estadual de Campinas, Brazil, a referral university hospital in southeast Brazil, accounting for a surrounding population of 3,100,000 inhabitants. This study was approved by the research ethics committee of the institution (CAAE report 15429419.5.0000.5404).

We included all pregnancies of women with a previous diagnosis of CKD who underwent prenatal follow-up at the specialized antenatal care (ANC) outpatient clinic and who
gave birth at the Women’s Hospital between 2012 and 2020. All patients with high risk of preeclampsia were given prophylaxis with low dose aspirin and calcium supplementation, as recommended by institutional protocol. We collected data from the medical records on an electronic system by completing a data collection form specifically created for the study.

We evaluated variables related to CKD etiology, duration of kidney disease treatment, sociodemographic variables, lifestyle variables, other associated diseases, and obstetric history and perinatal outcomes. In the case of patients with more than one pregnancy during the study period, each index pregnancy was considered, that is, the unit of study was the pregnancy. The data obtained were entered into a database created for this study, in Excel format, which was reviewed to identify inconsistencies. The underlying causes of kidney disease were later grouped into 10 subgroups according to similar characteristics and frequency of diagnoses.

To describe the profile of the sample according to the variables under study, frequency tables of categorical variables were made with absolute (n) and percentage (%) frequency values, and descriptive statistics of numerical variables, with mean values and standard deviation. Subsequently, women who had a viable pregnancy (excluding abortions and fetal deaths) were divided into 2 groups according to the occurrence or not of prematurity (gestational age [GA] < 37 weeks). To compare the categorical variables between pregnancies that ended in preterm birth, the chi-squared test or Fisher exact test (for expected values lower than 5) were used.

The significance level adopted for the statistical tests was 5%. The software used was the SAS System for Windows version 9.2. (SAS Institute Inc, 2002–2008, Cary, NC, USA).8–11

All Strengthening the Reporting of Observational Studies in Epidemiology (STROBE) requirements for an observational study were followed and verified in this article.

Results

A total of 84 pregnancies were included, in 67 women with CKD who underwent absolute neutrophil count (ANC) between 2012 and 2020. Of these, 6 pregnancies evolved with fetal deaths and 5 with abortion, totaling 11 gestational losses, which corresponds to 13.1% of this sample. A diamniotic dichorionic twin pregnancy occurred in a 32-year-old primigravid patient with a history of systemic lupus erythematosus (SLE), CKD on dialysis, kidney transplant in 2010 and viral infection with loss of the transplanted kidney. The patient progressed to preterm labor at 32 weeks and underwent a cesarean section; the newborns were born weighing 1,460 g and 1,197 g, with favorable neonatal outcomes. Considering the single pregnancies that progressed to childbirth (n = 72), the mean age of the pregnant women was 28.58 years (standard deviation [SD] = 6.34), with a mean time since diagnosis of CKD of 10.61 years (SD = 8.82). Most of the women were white, were in a stable relationship, + and had high school education; none of the participants reported using alcohol or illicit drugs. – Table 1 shows the sociodemographic data of the patients included in the study.

We grouped the main causes of CKD into 10 categories, presented from the most frequent to the least frequent: SLE (n = 21), glomerulopathy (n = 12), nephrotic syndrome (n = 11), transplant (n = 10), infection (n = 8), dialysis (n = 3), hypertension (n = 3), diabetes (n = 2), and other diseases with lower frequency that were included in the “others” group (n = 2: one patient with Wegener granulomatosis and the other with rheumatoid arthritis and nephrolithiasis). – Fig. 1 shows the cause of CKD of the patients included in the study.

Among the 72 pregnancies that resulted in live births, the mean gestational age at birth was 35 weeks and 3 days (SD = 7.21), with a mean birth weight of 2,443.7 g (SD = 722.48). The majority had a birth at term (52.78%) or late preterm (34.72%). Eight newborns were classified as small for gestational age (11.1%), and 5 of them were children.

| Table 1 | Characteristics of women with chronic kidney disease and singleton pregnancy that progressed to childbirth |
|---------|----------------------------------------------------------------------------------------------------------|
| Age (years) | N = 72 | % |
| < 20 | 3 | 4.17 |
| 20–29 | 35 | 48.61 |
| 30–39 | 31 | 43.05 |
| ≥ 40 | 3 | 4.17 |
| Age (mean) | 28.58 years |
| Marital status | Single | 26 | 36.11 |
| Stable relationship | 46 | 63.89 |
| Occupation* | Paid work | 11 | 15.28 |
| Unpaid work | 28 | 38.89 |
| Schooling** | Elementary | 13 | 13.06 |
| High school | 33 | 45.83 |
| University | 5 | 6.94 |
| Skin color*** | White | 46 | 63.89 |
| Non-White | 24 | 33.33 |
| Smoking | 2 | 4.17 |
| Number of Pregnancies | 1 | 31 | 43.05 |
| 2 | 16 | 22.22 |
| ≥ 3 | 25 | 34.73 |
| Previous miscarriage | 0 | 59 | 81.94 |
| ≥ 1 | 13 | 18.06 |

Frequency missing ‘33 ‘21 ‘332.
of mothers with SLE ($\frac{3}{4} = 62.5\%$). Five children were classified as low birth weight newborns (0.69%). Table 2 shows obstetric and neonatal data for patients with CKD included in the study. These women attended an average of 9.29 (SD = 3.88) prenatal consults. Approximately a quarter (20/72; 27.78%) of the study population developed preeclampsia during pregnancy.

Considering the high prevalence of preterm birth among the considered cases, and the burden of this condition for mothers and their children in the short and long term, we aimed to investigate conditions associated with this event. We compared the two groups according to the occurrence or not of preterm birth. There was a significant association between the occurrence of premature birth (gestational age $< 37$ weeks) and the need for a woman to be hospitalized in the ICU after childbirth, a higher occurrence of complications after childbirth, and a greater number of days of hospitalization (Table 3). The majority cause of preterm birth among our population was preeclampsia (12/34 = 35.3%) and premature rupture of membranes (4/34 = 11.7%). This population had more prevalence of ICU admission and more days staying in the hospital.

**Discussion**

The present study reports increased adverse maternal and neonatal outcomes among cases of CKD followed at a referral maternity hospital. Overall, around 13.1% of pregnancies progressed to abortion or stillbirth, and, among the cases of livebirths, almost half were preterm deliveries, with around one quarter complicated by preeclampsia. The cases of preterm delivery were associated with increased adverse outcomes, with 6 neonatal deaths.

According to international epidemiological data, the average rate of preterm births in the general population is 7 to 12% of births and, of these, $\sim 12\%$ occur due to preeclampsia. In our study, as expected, the incidence of premature births was much higher than in the general population (almost 50%), which is most likely due to CKD itself or secondary to the development of hypertensive syndromes and their consequences. The incidence of prematurity among pregnant women with CKD found in our study is similar to the data reported in the literature. A meta-analysis of 23 studies and 506,340 pregnant women concluded that CKD increased the risk of preeclampsia 10-fold, the risk of premature delivery and small-for-gestational-age newborns and led to a 3-fold increased risk of cesarean section.

Another study showed an association between CKD stage and its implications, with an increased incidence of prematurity as the CKD stage increased (CKD stage 1: 23.5% preterm; stage 2: 50.6%; stage 3: 78.4%; stage 4–5: 88.9%), using a serum creatinine threshold of 1.9 mg/dL, observed 93% newborn survival, 59% preterm birth, and fetal growth restriction of 37%. In our study, we did not distinguish the CKD stage of the patients; however, our data are similar to the data from this study, since the incidence is within this
Maternal ICU admission

None 53 76.61

Hospitalized in adult ICUs among those who had premature birth with more severe cases, with a greater number of women having been admitted to the ICU.

Maternal morbidities after birth

C-Section 49 68.06
Vaginal 23 31.94

Mode of birth

Without hypertension 35 48.61
Preeclampsia 8 11.11
Superimposed preeclampsia 12 16.67
Chronic hypertension 17 23.61
Hypertensive syndrome 2 2.78

Gestational age at birth

GA < 28 4 5.55
≥ 28 GA < 32 5 6.95
≥ 32 GA < 37 25 34.72
GA ≥ 37 38 52.78

Neonatal ICU admission

Yes 26 36.11
No 45 62.50

Neonatal death

Yes 6 8.33
No 62 86.11

Abbreviation: GA, gestational age; ICU, intensive care unit.

Missing ‘3’ ‘1’ ‘4’ (no information due to medical transfer of the newborn to another hospital) ‘*’ ‘**’ ‘***’ ‘****’ Others: one hypoglycemia, four with hypertensive spikes, one with hypervolemia and dialysis.

range. Other studies have already shown that there is an association between CKD and prematurity, and one showed a 1.23-fold increase in relative risk of preterm birth in patients with prepregnancy kidney dysfunction, compared with those with normal renal function.\(^2\,7\,13\,15\)

Our data show that prematurity was most likely associated with more severe cases, with a greater number of women hospitalized in adult ICUs among those who had premature births (around one-third of cases), while in patients with term birth, the number of ICU admissions was much lower (less than 10%). Maternal factors associated with CKD can increase the chance of patients being hospitalized in the ICU, due to the complexity of their cases, especially in those undergoing dialysis.

Epidemiologically, the most prevalent risk factors for CKD in the general population are arterial hypertension and diabetes mellitus; however, among pregnant women (mostly young women), other comorbidities are associated with the loss of renal function.\(^16\) In our study, the mean age of the patients was 28.5 years, with multiple other causes for CKD. Additionally, the study hospital is the referral hospital for some diseases, including SLE.

It was also possible to verify that CKD was a risk factor for the development of hypertensive syndromes during pregnancy, more frequently observed in the group of patients whose outcome was preterm birth.

It is known that CKD is a factor for hypertensive syndromes, as observed in several studies. One study of 778 women with CKD reported that 25.3% presented chronic arterial hypertension, and the incidence of preeclampsia was 9.3%.\(^17\) Another study reported an incidence of chronic arterial hypertension of 30.5% and of preeclampsia of 24.6%, with a higher rate of preeclampsia, explained in the study by almost a quarter of patients having CKD at more advanced stages (3–5).\(^18\) In addition, meta-analyses and cohorts show that women with CKD are 10 times more likely to develop preeclampsia, and up to 40% of patients with preeclampsia have had CKD previously.\(^2\,6\,19\,–\,22\)

Acetylsalicylic acid has been recommended as an effective intervention to reduce the incidence of preeclampsia, especially in women with known risk factors, including those with CKD, preferably introduced between 12 and 16 weeks.\(^19\,\,23\,\,24\) A study showed that its use may reduce the chance of developing preeclampsia and intrauterine growth restriction,\(^24\) while another showed a reduction in the incidence of severe preeclampsia among patients with CKD stages 3 to 5, with no evidence in this study among patients with CKD stages 1 to 2.\(^18\,\,24\) However, another controlled trial showed a reduction in preterm preeclampsia in patients that used aspirin.\(^25\) Our patients, guided by the institution protocol, used aspirin prophylaxis throughout the pregnancy, as well as calcium supplementation, as this is recommended in some groups of patients such as those with SLE.\(^26\)

The overall incidence of preeclampsia in the general population is ~4.6 to 8.1%, depending on the region.\(^5\,\,19\,\,27\)

In this study group, we saw a rate of 27.78%, high compared with the general population, but close to the values found in other studies that evaluated populations with CKD. We currently know that CKD, even in its early stages, is associated with the production of proinflammatory cytokines, which triggers endothelial inflammation and consequently increases the chance of developing hypertension. In a normal pregnancy, there is a balance between angiogenic factors (among them PlGF and VEGF) and anti-angiogenic factors (sFlt-1), favoring good placental implantation. However,
when there is an imbalance between these factors, poor implantation of the placenta or worsening of placental perfusion can occur, leading to a reduction in factors such as PlGF and VEGF and an increase in sFlt-1, causing endothelial inflammation and increasing the chances of developing preeclampsia.

Biomarker assessment may be an interesting way to adequately distinguish preeclampsia from other complications that can present with worsening proteinuria and hypertension.

Some of these studies report that the decline in kidney function is even worse among patients with CKD stages 3 to 5, while other studies did not see any worsening of renal function in stages 1 to 3. These data in the literature still lacks consensus and require further investigation.

Our study presented some limitations. Given that our data was from a single center, it was not possible to further investigate the association between the reported adverse outcomes and the diverse CKD reported. In addition, pregnant women were referred to our service after the diagnosis of kidney disease, implying that we had no data related to the kidney biopsy, details about the infections that caused kidney failure or previous treatments. Our patients had a very heterogeneous treatment during pregnancy based on the causes and evaluation of their kidney disease. It was possible only to the group of women who underwent dialysis. The others received basic care to treat the underlying

| Variables                              | Preterm birth (n = 34) | Term birth (n = 38) | p-value |
|----------------------------------------|------------------------|---------------------|---------|
| Maternal age (mean / SD)               | 29.03 (6.44)           | 28.18 (6.31)        | 0.560 * |
| Years since diagnosis (mean / SD)      | 10.00 (7.68)           | 11.13 (9.77)        | 0.894*  |
| Maternal hospitalization in days (mean / SD) | 4.85 (2.93)       | 3.13 (1.44)         | 0.001*  |
| Group of kidney disease                |                        |                     | 0.143** |
| SLE                                    | 8 (23.5%)              | 13 (34.2%)          |         |
| Glomerular disease                     | 4 (11.7%)              | 8 (21.1%)           |         |
| Nephrotic syndrome                     | 5 (14.7%)              | 6 (15.8%)           |         |
| Transplant                             | 4 (11.7%)              | 6 (15.8%)           |         |
| Infectious                             | 7 (20.6%)              | 1 (2.6%)            |         |
| Hemodialysis                           | 2 (5.9%)               | 1 (2.6%)            |         |
| Hypertensive nephropathy               | 2 (5.9%)               | 1 (2.6%)            |         |
| Diabetes nephropathy                   | 2 (5.9%)               | 0                   |         |
| Others                                 | 0                      | 2 (5.3%)            | 0.892** |
| Skin color                             |                        |                     |         |
| White                                  | 23 (67.3%)             | 23 (63.9%)          |         |
| Non-white                              | 11 (32.7%)             | 13 (36.1%)          |         |
| Hypertensive syndrome                  |                        |                     | 0.009** |
| yes                                    | 23 (67.7%)             | 14 (36.8%)          |         |
| no                                     | 11 (32.3%)             | 24 (63.2%)          |         |
| Preeclampsia                           |                        |                     | 0.178** |
| yes                                    | 12 (35.3%)             | 8 (21.0%)           |         |
| no                                     | 22 (64.7%)             | 30 (78.9%)          |         |
| Mode of birth                          |                        |                     | 0.663** |
| vaginal                                | 10 (29.4%)             | 13 (34.2%)          |         |
| cesarean                               | 24 (70.6%)             | 25 (65.8%)          |         |
| Postpartum maternal ICU                |                        |                     | 0.006** |
| yes                                    | 11 (34.4%)             | 3 (11.9%)           |         |
| no                                     | 21 (65.6%)             | 35 (92.1%)          | 0.029** |
| Adverse maternal outcome #             |                        |                     |         |
| yes                                    | 11 (35.5%)             | 5 (13.2%)           |         |
| no                                     | 20 (64.5%)             | 33 (86.8%)          |         |

Abbreviations: ICU, intensive care unit; SD, standard deviation; SLE.
*Kruskal-Wallis test, ** Qui-square test, *** Fisher test # adverse maternal outcome: bleeding, infection or other; #2 missing data.
disease, such as hypertension, lupus, and diabetes. However, our sample is relevant to referral centers in a middle-income setting.

In our study, 68.06% of the patients underwent cesarean section, a number higher than that recommended by the World Health Organization (WHO) and higher than the Brazilian average.\(^{30,31}\) However, in this case, these are patients with a greater possibility of complications, or acute or chronic fetal distress, which may be factors that increase the chances of opting for cesarean delivery. Even so, it is a high rate of cesarean sections, similar to that seen in other studies, which found cesarean section rates of 37 to 59%.\(^{13,14}\) This corroborates the results of the study by Zhang et al., which showed a 3-fold increase in the number of cesarean sections in patients with CKD.\(^6\)

**Conclusion**

These data reinforce that pregnancy complicated by CKD can present increased adverse maternal and perinatal outcomes, in addition to worsening the underlying disease or renal function. It is necessary to counsel these women on adequate family planning, to help plan their pregnancies when their kidney disease is stable and controlled. These are pregnant women who require multiprofessional evaluation at a referral center, with special attention and care during high-risk prenatal care. With a planned pregnancy, it is possible to better evaluate risk factors and prognosis, and evaluate the indication of prophylaxis for preeclampsia, in addition to undertaking maternal-fetal surveillance and monitoring.

**Contributors**

Marcus Vinicius Pinheiro Zilli: investigation, data curation and writing - original draft. Anderson Borac-Pinheiro: methodology, data curation and writing - original draft. Maria Laura Costa: investigation, review & editing. Fernanda Garanhani Surita: conceptualization, methodology, data curation, supervision, review & editing. All authors approved the final version to be published.

**Conflict of Interests**

The authors have no conflict of interests to declare.

**Acknowledgments**

The authors would like to thank Helymar Costa Machado, a statistician who made it possible to analyze the data of the study.

**References**

1. GBD Chronic Kidney Disease Collaboration. Global, regional, and national burden of chronic kidney disease, 1990-2017: a systematic analysis for the Global Burden of Disease Study 2017. Lancet. 2020;395(10225):709–733. DOI: 10.1016/S0140-6736(20)30045-3
2. Hui D, Hladunewich MA. Chronic kidney disease and pregnancy. Obstet Gynecol. 2019;133(06):1182–1194. DOI: 10.1097/AOG.0000000000003256
3. Williams D, Davison J. Chronic kidney disease in pregnancy. BMJ. 2008;336(7637):211–215. DOI: 10.1136/bmj.39406.652986.BE
4. Kumakura S, Okamoto K, Takeuchi S, Yoshida M, Nakamichi T, Nagasawa T, et al. Kidney function, blood pressure and proteinuria were associated with pregnancy outcomes of pregnant women with chronic kidney disease: a single-center, retrospective study in the Asian population. Clin Exp Nephrol. 2020;24(06):547–556. DOI: 10.1007/s10157-020-01865-0
5. Kattah A. Preeclampsia and kidney disease: deciphering cause and effect. Curr Hypertens Rep. 2020;22(11):91. DOI: 10.1007/s11906-020-01099-1
6. Zhang JJ, Ma XX, Hao L, Liu LJ, Lv JC, Zhang H. A systematic review and meta-analysis of outcomes of pregnancy in CKD and CKD outcomes in pregnancy. Clin J Am Soc Nephrol. 2015;10(11):1964–1978. DOI: 10.2215/CJN.09250914
7. Harel Z, Park AL, McArthur E, Hladunewich M, Dirk JS, Wald R, et al. Prepregnancy renal function and risk of preterm birth and related outcomes. CMAJ. 2020;192(30):E851–E857. DOI: 10.1503/cmaj.200089
8. Conover WJ. Practical nonparametric statistics. 3rd ed. New York: John Wiley & Sons; 1999
9. Siegel S, Castellan NJ Jr. Estatística não-paramétrica para ciências do comportamento. 2a ed. Porto Alegre: Art Med. 2006
10. Tabachnick BG, Fiddel LS. Using multivariate statistics. 4th ed. Boston: Allyn and Bacon; 2001
11. Fleiss JL, Levin B, Paik MC. Statistical methods for rates and proportions. 3rd ed. Hoboken: John Wiley & Sons; 2003
12. Goldenberg RL, Culhane JF, Iams JD, Romero R. Epidemiology and causes of preterm birth. Lancet. 2008;371(9606):75–84. DOI: 10.1016/S0140-6736(08)60074-4
13. Piccoli GB, Cabiddu G, Attini R, Vigotti FN, Maxia S, Lepori N, et al. Risk of adverse pregnancy outcomes in women with CKD. J Am Soc Nephrol. 2015;26(08):2011–2022. DOI: 10.1681/ASN.2014050459
14. Jones DC, Hayslett JP. Outcome of pregnancy in women with moderate or severe renal insufficiency. N Engl J Med. 1996;335(04):226–232. DOI: 10.1056/NEJM199607253350402
15. Cabiddu G, Castellino S, Gernone G, Santoro D, Moroni G, Giannattasio M, et al. A best practice position statement on pregnancy in chronic kidney disease: the Italian Study Group on Kidney and Pregnancy. J Nephrol. 2016;29(03):277–303. DOI: 10.1007/s40620-016-0285-6
16. Marinho AW, Penha AP, Silva MT, Galvão TF. Prevalência de doença renal crônica em adultos no Brasil: revisão sistemática da literatura. Cad Saúde Coletiva. 2017;25(03):379–88. DOI: 10.1590/1414-4627201700030134
17. Kendrick J, Sharma S, Holmen J, Palit S, Nuccio E, Chonchol M. Kidney disease and maternal and fetal outcomes in pregnancy. Am J Kidney Dis. 2015;66(01):55–59. DOI: 10.1053/j.ajkd.2014.11.019
18. Wang M, Chen S, He Y, Zhao M, Yang H, Chen Q. Low-dose aspirin for the prevention of severe preeclampsia in patients with chronic kidney disease: a retrospective study. This is the study for kidney and pregnancy. J Nephrol. 2021;34(05):1631–1639. DOI: 10.1007/s40620-021-01049-3
19. Wiles K, ChapPELL LC, Lightstone L, Bramham K. Updates in diagnosis and management of preeclampsia in women with CKD. Clin J Am Soc Nephrol. 2020;15(09):1371–1380. DOI: 10.2215/CJN.15121219
20. Brown MA, Magee LA, Kenny LC, Karumanchi SA, McCarthy FP, Saito S, et al; International Society for the Study of Hypertension in Pregnancy (ISSHP). The hypertensive disorders of pregnancy: ISSHP classification, diagnosis & management recommendations for international practice. Pregnancy Hypertens. 2018;13:291–310. DOI: 10.1016/j.preghy.2018.05.004
21. Bartsch E, Medcalf KE, Park AL, Ray J. High Risk of Pre-eclampsia Identification Group. Clinical risk factors for pre-eclampsia determined in early pregnancy: systematic review and meta-
analysis of large cohort studies. BMJ. 2016;353:i1753. Doi: 10.1136/bmj.i1753

22 Saudan P, Brown MA, Buddle ML, Jones M. Does gestational hypertension become pre-eclampsia? Br J Obstet Gynaecol. 1998;105(11):1177–1184

23 The National Institute for Health and Care Excellence. Hypertension in pregnancy: diagnosis and management [Internet]. 2019 [cited 2022 Feb 1]. (NICE guideline; 133). Available from: www.nice.org.uk/guidance/ng133

24 Ray JG, Bartsch E, Park AL, Shah PS, Dzakpasu S. Estimated reductions in provider-initiated preterm births and hospital length of stay under a universal acetylsalicylic acid prophylaxis strategy: a retrospective cohort study. CMAJ Open. 2017;5(02):E508–E516. Doi: 10.9778/cmao.20160092

25 Rolnik DL, Wright D, Poon LC, O’Gorman N, Syngelaki A, de Paco Matallana C, et al. Aspirin versus placebo in pregnancies at high risk for preterm preeclampsia. N Engl J Med. 2017;377(07):613–622. Doi: 10.1056/NEJMoa1704559

26 Pastore DEA, Costa ML, Parpinelli MA, Surita FG. A critical review on obstetric follow-up of women affected by systemic lupus erythematosus. Rev Bras Ginecol Obstet. 2018;40(04):209–224. Doi: 10.1055/s-0038-1625951

27 Giordano JC, Parpinelli MA, Cecatti JG, Haddad SM, Costa ML, Surita FG, et al. The burden of eclampsia: results from a multicenter study on surveillance of severe maternal morbidity in Brazil. PLoS One. 2014;9(05):e97401 Doi: 10.1371/journal.pone.0097401

28 de Jesús GR, Lacerda MI, Rodrigues BC, Santos FC, Nascimento AP, Cristóvão Porto L, et al. Soluble Flt-1, placental growth factor, and vascular endothelial growth factor serum levels to differentiate between active lupus nephritis during pregnancy and preeclampsia. Arthritis Care Res (Hoboken). 2021;73(05):717–721

29 Di Leo V, Capaccio F, Gesualdo L. Preeclampsia and glomerulonephritis: a bidirectional association. Curr Hypertens Rep. 2020;22(05):36. Doi: 10.1007/s11906-020-1033-9

30 Rudey EL, Leal MDC, Rego G. Cesarean section rates in Brazil: Trend analysis using the Robson classification system. Medicine (Baltimore). 2020;99(17):e19880. Doi: 10.1097/MD.0000000000019880

31 Boerma T, Ronsmans C, Melesse DY, Barros AJ, Barros FC, Juan L., et al. Global epidemiology of use of and disparities in caesarean sections. Lancet. 2018;392(10155):1341–1348. Doi: 10.1016/S0140-6736(18)31928-7