Preterm birth is strongly affected by the glucocorticoid dose during pregnancy in women complicated by systemic lupus erythematosus

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Abstract

**Background:** This study aimed to investigate the effect of glucocorticoid doses on adverse pregnancy outcomes (APOs) in women complicated by systemic lupus erythematosus (SLE).

**Methods:** We investigated 74 pregnancies complicated by SLE or SLE-dominant mixed connective tissue disease. The pregnancies were managed from conception to delivery in our institution. We retrospectively evaluated whether the mean glucocorticoid dose during pregnancy is associated with APOs, including preterm birth (PB), low birth weight (LBW), and light-for-date (LFD). We also calculated the cut-off dose of glucocorticoid that affected APOs.

**Results:** All APOs occurred in 35 (50.7%) patients, with 14 cases of PB, 23 cases of LBW, and 10 cases of LFD. Patients with all APOs or PB had a higher dose of glucocorticoid during pregnancy than patients without all APOs or with full-term birth ($P = 0.03$, $P < 0.01$, respectively). Logistic regression analysis for all APOs and PB showed that the cut-off values of the mean glucocorticoid dose were 6.5 and 10.0 mg/day, respectively. Patients who delivered LBW or LFD newborns showed no significant difference in the glucocorticoid dose used during pregnancy than patients without LBW or LFD newborns. Patients who delivered LBW newborns were more likely to have used glucocorticoids during pregnancy ($P < 0.01$).

**Conclusions:** In pregnancies complicated by SLE, a relatively lower dose of glucocorticoid than previously reported is significantly related to APOs, especially PB. Therefore, the disease activity of patients with SLE should be managed with the appropriate lower dose of glucocorticoid during pregnancy.

**Keywords:** Glucocorticoid, Systemic lupus erythematosus, Adverse pregnancy outcome, Preterm birth

Background

Systemic lupus erythematosus (SLE) occurs in women of child-bearing age. Recently, patients with SLE have been able to be diagnosed earlier according to the diagnosis or classification criteria of each disease and achieve long-term remission by treatment agents, including glucocorticoids and several immunosuppressants. Therefore, women with SLE who used to avoid...
pregnancy because of their disease, can currently conceive and become mothers. However, women with SLE have more difficulty in achieving a successful pregnancy than healthy women. Women with SLE have a higher risk of adverse pregnancy outcomes (APOs), including preterm birth (PB), light-for-date (LFD), premature rupture of the membranes (PROM), and preeclampsia [1–6]. These APOs are related to uncontrolled high disease activity [7–10]. Therefore, SLE disease activity needs to be strictly controlled with glucocorticoids and immunosuppressants, which are tolerable in pregnancy.

Glucocorticoids are the most commonly used agents for maintaining and managing high aggressive disease activity during pregnancy. The updated British Society for Rheumatology guideline, the European League Against Rheumatism recommendation, and the 2020 American College of Rheumatology (ACR) guideline showed that prednisolone was compatible with each trimester of pregnancy [11–13]. After completion of placental development, maternal glucocorticoids are metabolized by 11-beta-hydroxylase in the placenta, and <10% of the dose of prednisone or prednisolone crosses the placenta. Several studies have shown that glucocorticoids are not associated with major abnormalities, especially in oral cleft [14, 15]. However, a prolonged high dose of glucocorticoid use during pregnancy increases the risk of APOs. Reinisch et al. showed that intrauterine growth restriction was associated with glucocorticoids, and it was independent of maternal disease activity [16]. Many studies have also shown that steroid use during pregnancy is a risk factor for PB and low birth weight (LBW) [15, 17–19]. Chakravarty et al. found that preterm delivery was also associated with glucocorticoid use in pregnancies complicated by SLE [20]. In addition, high doses of glucocorticoids increase the risk of serious infection during pregnancy [21].

Several reports have also described that using <20 mg/day of prednisolone during pregnancy and sometimes higher doses of prednisolone are appropriate for aggressive disease activity [22–24]. Palmsten et al. showed that high doses of prednisolone (>20 mg/day) early and late in pregnancy were associated with PB [25]. Additionally, the ACR guideline showed that glucocorticoids should be tapered to <20 mg/day by adding pregnancy-compatible immunosuppressants [11]. However, whether a glucocorticoid dose of <20 mg/day is appropriate for mothers with SLE and infants has not been clarified.

Therefore, this study aimed to investigate the effect of glucocorticoid doses on adverse pregnancy outcomes (APOs), including PB, LBW and LFD, in women complicated by SLE. We also evaluated which glucocorticoid dose can cause these APOs.

Methods

Patients and data collection

Our study used data from patients with SLE and those with SLE-dominant mixed connective tissue disease (MCTD) who were treated in Kagawa University Hospital from March 2006 to April 2021. These patients were diagnosed with the 1997 ACR revised criteria for SLE [26] or the 2019 Japanese Ministry of Health, Labour and Welfare diagnostic criteria of MCTD [27]. MCTD was diagnosed with any two characteristics of SLE, myositis, and systemic sclerosis. SLE-dominant MCTD was defined as patients with MCTD who mainly had symptoms of SLE and immunological abnormalities, such as erythema, low complement levels, or a high titer of anti-double stranded antibody. The patients were treated at Kagawa University Hospital from pre-conception counseling to pregnancy and delivery. We investigated these patients’ clinical background (age at conception disease duration, which was defined as the period from disease onset to conception, and parity), autoantibody profiles (including anti-SS-A/B antibodies and antiphospholipid antibodies), SLE disease activity scores and parameters, and the status of treatment agents (immunosuppressants and glucocorticoid used). The disease activity score was calculated using the SLE Disease Activity Index (SLEDAI) [28, 29] and the lupus low disease activity state (LLDAS) [30]. The parameters of the SLE disease activity score included complement levels of C3, C4, and CH50, and the titer of anti-dsDNA antibody. All laboratory tests were performed using standard methods.

We examined glucocorticoid use at conception or during pregnancy, and the mean dose of glucocorticoid, which was defined as the average glucocorticoid dose from conception to delivery. Additionally, we investigated whether the dose of glucocorticoid needed to be increased during pregnancy.

We evaluated the association between glucocorticoid use and APOs, which included PB, LBW, and LFD. PB was defined as delivery before 37 gestational weeks. LBW was defined as newborns who were born weighing <2500 g. LFD was defined as newborns whose birth weight was lower than the 10th percentile. PROM was defined as rupture of the membranes (amniotic sac) before labor began, and preterm PROM occurred before 37 gestational weeks.

All data were retrospectively collected from medical records at Kagawa University Hospital. This was a retrospective, observational study. Therefore, we did not require ethical approval. We did not obtain informed consent from each patient, however, we disclosed the information of our research.
Statistical analysis
Values are presented as mean ± standard deviation or number (%). We performed univariate analysis to determine the associations between APOs and glucocorticoid use, mean glucocorticoid dose, and increasing the dose of glucocorticoid during pregnancy. A logistic regression analysis of the associations between the glucocorticoid dose and APOs, PB, LBW, and LFD was performed. The receiver operating characteristic (ROC) curve was used to determine the cut-off value for the mean glucocorticoid dose that affected APOs. Descriptive statistics were compared using the Wilcoxon rank sum test for continuous variables and Fisher’s exact test for categorical variables. A two-sided P value of < 0.05 was considered significant. We evaluated parameters with significant differences (P < 0.05) as risk factors. All analyses were conducted using JMP for Mac, Version 13.0.0 (SAS Institute, Japan).

Results
Patients’ characteristics and treatments
Our study used data from 74 pregnancies in 52 patients with SLE and those with SLE-dominant MCTD. The patients’ characteristics, disease activities, treatment agents, and pregnancy outcomes are shown in Table 1. The mean age at delivery was 31.8 ± 4.4 years and the mean disease duration was 10.0 ± 5.1 years. Forty-two (56.8%) patients were positive for anti-SS-A antibody and 29 (39.2%) were positive for antiphospholipid antibodies. With regard to disease activity parameters, the mean SLEDAI score in the first and third trimesters was 1.8 ± 2.1 and 1.2 ± 1.9, respectively. Forty (59.7%) achieved an LLDAS and 50 (74.6%) achieved an LLDAS without a glucocorticoid dose at conception. Complement levels were normal (C3: 89.8 ± 21.4, C4: 18.2 ± 6.7, CH50: 41.0 ± 9.6), and the titer of anti-dsDNA antibody was almost normal (10.8 ± 32.6). Regarding treatment agents used at conception and during pregnancy, immunosuppressants were administered in 16 (21.6%) patients at conception. Glucocorticoids were administered in 57 (77.0%) patients at conception and in 61 (82.4%) patients during pregnancy, and all of them took prednisone. The mean dose of glucocorticoid at conception was 6.5 ± 3.2 mg/day, and that during pregnancy was 8.4 ± 4.8 mg/day. Fifteen (20.3%) patients needed to increase their dose of glucocorticoid during pregnancy because the SLE disease activity was elevated.

Pregnancy outcomes
Table 1 also shows the pregnancy outcomes in all patients. Fifteen (20.3%) patients experienced loss of the newborn because of eight (10.8%) cases of spontaneous abortion, two (2.7%) cases of stillbirth, and five (6.8%) cases of induced abortion. In 59 (79.7%) patients with a live birth, the mean gestational age at delivery was 37.5 ± 3.0 weeks. The mean birth weight of newborns was 2643.3 ± 665.2 g. APOs occurred in 35 (50.7%) women, and these included spontaneous abortion, stillbirth, 14 (23.7%) cases of PB, 23 (39.0%) cases of LWB, 10 (16.9%) cases of LFD, 7 (11.9%) cases of preterm PROM, and 5 (8.6%) cases of preeclampsia.

Analysis of risk factors for APOs
We investigated the associations between glucocorticoid use and all APOs, PB, LBW, and LFD in all patients (Table 2). All APOs were evaluated in 69 patients after excluding 5 cases of induced abortion. Additionally, PB, LBW, and LFD were examined in 59 patients after excluding 8 cases of spontaneous abortion and 2 cases of stillbirth. The mean glucocorticoid dose during pregnancy was significantly higher in patients who had all APOs or PB than in patients without all APOs or with full-term birth (P = 0.03 and P < 0.01, respectively). The rate of increasing the glucocorticoid dose was also significantly higher in patients with all APOs or PB than in patients without all APOs or with full-term birth (P = 0.01 and P < 0.01, respectively). We performed logistic regression analysis of the mean glucocorticoid dose for APOs and PB (Fig. 1). The ROC curve showed that the area under the curve (AUC) was 0.674 and 0.808 for all APOs and PB, respectively. The cut-off value of the mean glucocorticoid dose was 6.5 and 10.0 mg/day, respectively (P = 0.01 and P < 0.01). In patients who had a newborn with a LBW, the rates of glucocorticoid use and increasing doses of glucocorticoids during pregnancy were significantly higher than those in patients with newborns without LBW (both P < 0.01). However, there was no significant difference in the mean glucocorticoid dose between these two groups (P = 0.05). There were no significant differences in the rate of glucocorticoid use, mean glucocorticoid dose, or the rate of increasing dose of glucocorticoid during pregnancy between patients with newborns who were LFD and those with newborns who were not LFD (P = 0.19, P = 0.26, and P = 1.00, respectively). In logistic regression analysis for LBW and LFD, the ROC curve showed that the AUC was 0.662 and 0.615, respectively, but they were not significant (P = 0.06 and P = 0.18, respectively, Fig. 1). Additionally, the cut-off value of the mean glucocorticoid dose was 6.5 and 6.7 mg/day, respectively. In patients who had preterm PROM or preeclampsia, there was no significant difference in the rate of glucocorticoid use or the mean glucocorticoid dose during pregnancy compared with those without preterm PROM or preeclampsia (Additional file 1).
Table 1  Patients’ characteristics, disease activities, treatment agents, and pregnancy outcomes

| Patients’ characteristics |  |  |
|---------------------------|---|---|
| Mean age at delivery, years | 31.8 ± 4.4 |  |
| Mean disease duration, years | 10.0 ± 5.1 |  |
| Parity, n (%) | 40 (54.1) |  |

| Autoantibody positivity |  |  |
|-------------------------|---|---|
| Anti-SS-A antibody, n (%) | 42 (56.8) |  |
| Antiphospholipid antibodies, n (%) | 29 (39.2) |  |
| LAC/anti-cardiolipin/anti-CLβ2GP1 | 23 (31.1)/10 (13.5)/3 (4.1) |  |
| Number of positive antibodies (single/double/triple) | 22 (29.7)/6 (8.1)/1 (1.4) |  |

| Disease activity parameters | First trimester (at conception) | Third trimester |
|-----------------------------|-------------------------------|----------------|
| Mean SLEDAI score | 1.8 ± 2.1 | 1.2 ± 1.9 |
| Achievement of LLDAS, n (%) | 40 (59.7) | 35 (58.3) |
| Achievement of LLDAS without a glucocorticoid dose, n (%) | 50 (74.6) | 50 (83.3) |
| C3, mg/dl | 89.8 ± 21.4 | 101.9 ± 23.1 |
| C4, mg/dl | 18.2 ± 6.7 | 17.9 ± 7.2 |
| CH50, IU/ml | 41.0 ± 9.6 | 42.7 ± 10.1 |
| Titer of anti-dsDNA antibody, IU/ml | 10.8 ± 32.6 | 5.8 ± 12.3 |

| Status of treatment agents |  |  |
|----------------------------|---|---|
| Immunosuppressant use at conception, n (%) | 16 (21.6) |  |
| Hydroxychloroquine use during pregnancy, n (%) | 10 (13.5) |  |

| Glucocorticoid use |  |  |
|--------------------|---|---|
| Use at conception, n (%) | 57 (77.0) |  |
| Mean dose at conception, mg/day | 6.5 ± 3.2 |  |
| Use during pregnancy, n (%) | 61 (82.4) |  |
| Mean dose during pregnancy, mg/day | 8.4 ± 4.8 |  |
| Increasing doses used, n (%) | 15 (20.3) |  |

| Pregnancy outcomes |  |  |
|-------------------|---|---|
| Neonatal loss, n (%) | 15 (20.3) |  |
| Spontaneous abortion, n (%) | 8 (10.8) |  |
| Stillbirth, n (%) | 2 (2.7) |  |
| Induced abortion, n (%) | 5 (6.8) |  |
| Live birth, n (%) | 59 (79.7) |  |
| Cesarean section, n (%) | 16 (27.1) |  |
| Gestational age at delivery, weeks | 37.5 ± 3.0 |  |
| Birth weight of newborns, g | 2643.3 ± 665.2 |  |
| APOs, n (%) | 35 (50.7) |  |
| Spontaneous abortion, n (%) | 8 (10.8) |  |
| Stillbirth, n (%) | 2 (2.7) |  |
| PB, n (%) | 14 (23.7) |  |
| LBW newborns, n (%) | 23 (39.0) |  |
| LFD newborns, n (%) | 10 (16.9) |  |
| Preterm PROM, n (%) | 7 (11.9) |  |
| Preeclampsia, n (%) | 5 (8.6) |  |
| NICU administration, n (%) | 17 (28.8) |  |

Values are shown as mean ± standard deviation or n (%). LAC lupus anti-coagulant, CLβ2GP1 cardiolipin-beta 2 glycoprotein 1, SLEDAI Systemic Lupus Erythematosus Disease Activity Index, LLDAS lupus low disease activity state, APOs adverse pregnancy outcomes, PB preterm birth, LBW low birth weight, LFD light-for-date, PROM premature rupture of the membranes, NICU neonatal intensive care unit.
Table 2  Associations of glucocorticoid use with APOs, PB, LBW, and LFD

| APOs | PB | LBW | LFD |
|------|----|-----|-----|
|      | APOs (+) | APOs (−) | P value | PB (+) | PB (−) | P value | LBW (+) | LBW (−) | P value | LFD (+) | LFD (−) | P value |
|      | (n = 35) | (n = 34) | (n = 14) | (n = 45) |         |         | (n = 23) | (n = 36) |         |         |         |         |
| Glucocorticoid use, n (%)## | 31 (88.6) | 25 (73.5) | 0.13 | 13 (92.9) | 36 (80.0) | 0.42 | 23 (100.0) | 26 (72.2) | < 0.01* | 10 (100.0) | 39 (79.6) | 0.19 |
| Mean glucocorticoid dose, mg/day# | 9.6 ± 5.0 | 6.6 ± 3.7 | 0.03* | 12.3 ± 5.5 | 6.7 ± 3.3 | < 0.01* | 9.5 ± 4.9 | 7.1 ± 4.2 | 0.05 | 100 ± 5.7 | 7.8 ± 4.3 | 0.26 |
| Increase in the glucocorticoid dose, n (%)## | 11 (31.4) | 2 (5.9) | 0.01* | 9 (64.3) | 3 (6.7) | < 0.01* | 9 (39.1) | 3 (8.3) | < 0.01* | 2 (200) | 10 (20.4) | 1.00 |

*Values are shown as mean ± standard deviation or n (%). ##Wilcoxon rank sum test; #Fisher's exact test; *P < 0.05. APOs adverse pregnancy outcomes, PB preterm birth, LBW low birth weight, LFD light-for-date.
We also investigated the associations between disease activity parameters and all APOs, PB, LBW, and LFD in all patients (Table 3). All APOs were significantly associated with achievement of an LLDAS without a glucocorticoid dose at the third trimester ($P = 0.01$), the SLEDAI score at the first and third trimesters ($P = 0.01$ and $P = 0.02$, respectively), and C3 levels at the first trimester ($P = 0.01$). PB was significantly associated with achievement of an LLDAS at the third trimester ($P = 0.01$), achievement of an LLDAS without a glucocorticoid dose at the first and third trimesters (both of $P < 0.01$), the SLEDAI score at the first trimester ($P < 0.01$), C3 levels at the first and third trimesters ($P = 0.02$ and $P < 0.01$, respectively), CH50 levels at the third trimester ($P = 0.02$), and the titer of anti-dsDNA antibody at the first trimester ($P = 0.01$). LBW was significantly associated with achievement of an LLDAS without glucocorticoid at the third trimester ($P < 0.01$), the SLEDAI score at the first and third trimesters ($P < 0.01$), C3 and CH50 concentrations at the first trimester (both $P = 0.01$). However, there was no significant association between LFD and any disease activity parameters.

**Fig. 1** ROC curves based on logistic regression analysis of cut-off values for the mean glucocorticoid dose. A. ROC curve for APOs ($P = 0.01$). The AUC was 0.674 and the cut-off value for the mean prednisone dose was 6.5 mg/day. B. ROC curve for PB ($P < 0.01$). The AUC was 0.808 and the cut-off value for the mean prednisone dose was 10.0 mg/day. C. ROC curve for LBW ($P = 0.06$). The AUC was 0.662 and the cut-off value for the mean prednisone dose was 6.5 mg/day. D. ROC curve for LFD ($P = 0.18$). The AUC was 0.615 and the cut-off value for the mean prednisone dose was 6.7 mg/day. ROC: receiver operating characteristic; AUC: area under the curve; TPF: true positive fraction; FPF: false positive fraction; APOs: adverse pregnancy outcomes; PB: preterm birth; LBW: low birth weight; LFD: light-for-date.
Table 3  Association between disease activity parameters and APOs, PB, LBW, and LFD

|                      | APOs                  | P value | PB                   | P value | LBW                   | P value | LFD                   | P value |
|----------------------|-----------------------|---------|----------------------|---------|-----------------------|---------|-----------------------|---------|
|                      | APOs (+) (n = 35)     | APOs (−) (n = 34) | PB (+) (n = 14) | PB (−) (n = 45) | LBW (+) (n = 23) | LBW (−) (n = 36) | LFD (+) (n = 10) | LFD (−) (n = 49) |
| Achievement of LLDAS, n (%)## | 1st trimester 17 (56.7) | 21 (65.6) | 0.60 | 5 (41.7) | 29 (70.7) | 0.09 | 12 (63.2) | 22 (64.7) | 1.00 |
|                      | 3rd trimester 12 (46.2) | 23 (67.7) | 0.12 | 4 (28.6) | 31 (68.9) | 0.01* | 11 (47.8) | 24 (66.7) | 0.18 |
| Achievement of LLDAS without a glucocorticoid dose, n (%)## | 1st trimester 20 (66.7) | 27 (84.4) | 0.14 | 5 (41.7) | 36 (87.8) | < 0.01* | 13 (68.4) | 28 (82.4) | 0.31 |
|                      | 3rd trimester 18 (69.2) | 32 (94.1) | 0.01* | 7 (50.0) | 42 (93.3) | < 0.01* | 15 (65.2) | 34 (94.4) | < 0.01* |
| SLEDAI score#       | 1st trimester 25 ± 2.1 | 1.0 ± 1.4 | 0.01* | 3.5 ± 2.9 | 1.1 ± 1.3 | < 0.01* | 2.3 ± 2.1 | 1.3 ± 2.0 | 0.03* |
|                      | 3rd trimester 1.9 ± 2.4 | 0.7 ± 1.2 | 0.02* | 2.4 ± 3.0 | 0.8 ± 1.2 | 0.05 | 2.0 ± 2.5 | 0.7 ± 1.2 | 0.02* |
|                      | 3rd trimester 21.2 ± 2.5 | 10.7 ± 20.0 | 0.09 | 85.5 ± 23.4 | 108.2 ± 20.1 | < 0.01* | 93.7 ± 24.9 | 108.3 ± 20.0 | 0.07 |
|                      | 1st trimester 82.4 ± 18.9 | 99.8 ± 21.6 | 0.01* | 77.3 ± 19.0 | 95.5 ± 20.9 | 0.02* | 80.5 ± 16.3 | 97.8 ± 22.3 | 0.01* |
|                      | 3rd trimester 94.2 ± 25.2 | 107.7 ± 20.0 | 0.09 | 85.5 ± 23.4 | 108.2 ± 20.1 | < 0.01* | 93.7 ± 24.9 | 108.3 ± 20.0 | 0.07 |
|                      | 1st trimester 171 ± 6.7 | 196 ± 6.8 | 0.24 | 161 ± 9.0 | 195 ± 6.2 | 0.14 | 17.6 ± 6.3 | 194 ± 7.4 | 0.47 |
|                      | 3rd trimester 167 ± 8.4 | 189 ± 6.0 | 0.19 | 151 ± 6.8 | 191 ± 7.0 | 0.11 | 169 ± 8.7 | 189 ± 5.9 | 0.21 |
|                      | 3rd trimester 139 ± 10.6 | 430 ± 8.4 | 0.11 | 373 ± 10.6 | 415 ± 8.4 | 0.13 | 365 ± 6.6 | 427 ± 9.5 | 0.01* |
|                      | 3rd trimester 40.2 ± 11.5 | 446 ± 8.7 | 0.18 | 365 ± 10.6 | 450 ± 9.0 | 0.02* | 40.2 ± 11.6 | 448 ± 8.6 | 0.14 |
|                      | 1st trimester 15.6 ± 4.4 | 70 ± 126 | 0.39 | 32.5 ± 68.5 | 5.7 ± 11.2 | 0.01* | 20.7 ± 55.8 | 70 ± 123 | 0.40 |
|                      | 3rd trimester 8.7 ± 17.0 | 34 ± 6.1 | 0.38 | 14.4 ± 21.4 | 31 ± 5.7 | 0.05 | 9.1 ± 17.7 | 36 ± 6.1 | 0.52 |
|                      | 1st trimester 17.0 ± 2.4 | 1.0 ± 1.4 | 0.01* | 3.5 ± 2.9 | 1.1 ± 1.3 | < 0.01* | 2.3 ± 2.1 | 1.3 ± 2.0 | 0.03* |
|                      | 3rd trimester 1.9 ± 2.4 | 0.7 ± 1.2 | 0.02* | 2.4 ± 3.0 | 0.8 ± 1.2 | 0.05 | 2.0 ± 2.5 | 0.7 ± 1.2 | 0.02* |
|                      | 3rd trimester 21.2 ± 2.5 | 107.7 ± 20.0 | 0.09 | 85.5 ± 23.4 | 108.2 ± 20.1 | < 0.01* | 93.7 ± 24.9 | 108.3 ± 20.0 | 0.07 |
|                      | 1st trimester 82.4 ± 18.9 | 99.8 ± 21.6 | 0.01* | 77.3 ± 19.0 | 95.5 ± 20.9 | 0.02* | 80.5 ± 16.3 | 97.8 ± 22.3 | 0.01* |
|                      | 3rd trimester 94.2 ± 25.2 | 107.7 ± 20.0 | 0.09 | 85.5 ± 23.4 | 108.2 ± 20.1 | < 0.01* | 93.7 ± 24.9 | 108.3 ± 20.0 | 0.07 |

Values are presented as mean ± standard deviation or number (%). Wilcoxon rank sum test; #Fisher’s exact test; *P < 0.05. LLDAS low lupus disease activity status. SLEDAI Systemic Lupus Erythematosus Disease Activity Index, APOs adverse pregnancy outcomes, PB preterm birth, LBW low birth weight, LFD light-for-date.
Discussion
In this retrospective study, we showed that the mean glucocorticoid dose used during pregnancy was significantly related to APOs, especially in patients who had PB, in pregnancies complicated by SLE and in those with SLE-dominant MCTD. Additionally, women who delivered LBW newborns had a higher rate of glucocorticoid use during pregnancy than those without LBW. We found that the cut-off doses of glucocorticoid that affected all APOs and PB were 6.5 and 10.0 mg/day, respectively. Additionally, a relatively lower dose of glucocorticoid used during pregnancy than that previously reported [22–25] could cause APOs, especially PB. Furthermore, some of the disease activity parameters were associated with APOs, PB, and LBW, but not with LFD.

Previous studies have shown a significant association between connective tissue disease (CTD) and APOs, including PB, LBW, PROM, and preeclampsia [1–4]. In particular, patients with SLE have an increased risk of APOs, which are associated with a high disease activity and disease flare-ups during pregnancy [5–10]. Therefore, the disease activity of SLE needs to be maintained during pregnancy with therapeutic agents that do not affect fetal development. Glucocorticoids are most frequently used for maintenance and flare-up of CTD during pregnancy. In our study, glucocorticoids were administered in approximately 80% of women at conception and during pregnancy. Some patients needed to increase the dose of glucocorticoid and continue a high dose during pregnancy because of their elevated disease activity.

However, prolonged glucocorticoid use during pregnancy is also a risk factor for APOs. Reinisch et al. showed that oral glucocorticoids (especially prednisone) were associated with intrauterine growth restriction in humans and mice, and this was independent of maternal disease [16]. Furthermore, many studies have also shown that steroid use during pregnancy is a risk factor for PB and low birth weight (LBW) [15, 17–19]. Similar to these previous studies, our study showed that women who delivered LBW newborns were more likely to have used glucocorticoids during pregnancy. Additionally, we showed that the mean glucocorticoid dose used during pregnancy was significantly associated with all APOs and PB. However, these APOs were strongly associated with women who had increased doses of glucocorticoids because of an elevated disease activity. We found that some disease activity parameters were significantly related to all APOs, PB, and LBW, but not to LFD. Therefore, APOs are associated not only with glucocorticoid use and the mean glucocorticoid dose, but also with SLE disease activity. Determining whether glucocorticoids or disease activity is the most strongly associated with these APOs is difficult because of the small number of patients and outcomes in our study.

The glucocorticoid dose that can be tolerated in mothers with CTD and their fetus during pregnancy is unclear. Some reports have suggested that the prednisone dose should be < 20 mg/day [22–25]. The 2020 ACR guideline for the management of reproductive health also recommends tapering the prednisone dose to < 20 mg/day by adding pregnancy-compatible immunosuppressants [11]. However, whether < 20 mg/day of prednisone affects APOs is unclear. We found that the cut-off doses of glucocorticoid that affected all APOs and PB were 6.5 and 10.0 mg/day, respectively. The dose that could cause all APOs and PB in our study was relatively lower than that previously reported [22–25]. Therefore, an even lower dose of glucocorticoid < 20 mg/day could be a risk factor for APOs and PB.

Recently, some immunosuppressants have been recognized as safe and compatible during pregnancy. The ACR, European League Against Rheumatism, and British Society for Rheumatology guidelines show the availability of each drug in the period of preconception, during pregnancy, and at breastfeeding [11–13]. These drugs might be helpful in reducing the risk of APOs.

Our study has several limitations. First, our sample comprised a small number of patients, and there was also a small number of outcome events, which might have resulted in a low statistical power. Second, we could not fully exclude the heterogeneity of MCTD. In our study, patients with SLE-dominant MCTD who mainly had SLE symptoms and immunological abnormalities were enrolled, but these patients had some features of myositis or sclerosis. Third, this study was conducted in only one expert institution. Therefore, we might have collected data on patients with relatively high levels of disease activity and risk.

Conclusions
Continuing glucocorticoid use and the glucocorticoid dose used during pregnancy are significantly associated with APOs, including PB and LBW, in patients with SLE. Additionally, the cut-off doses of glucocorticoid that can cause all APOs and PB are 6.5 and 10.0 mg/day, respectively. Additionally, a relatively lower dose of glucocorticoids used during pregnancy than previously reported is strongly associated with all APOs and PB. Rheumatologists should pay attention to the risk of glucocorticoids in patients with SLE and manage their disease activity with an appropriate dose of glucocorticoid during pregnancy.

Abbreviations
APOs: Adverse pregnancy outcomes; ACR: American College of Rheumatology; AUC: Area under the curve; CLβ2GP1: Cardiolipin-beta 2 glycoprotein
1. Ohyama A, Tsuboi H, Noma H, Terasaki M, Shimizu M, Toko H, et al. Associations between maternal clinical features and fetal outcomes in pregnancies of mothers with connective tissue diseases. Mod Rheumatol. 2019;29:344–50.

2. Sugawara E, Kato M, Fujieda Y, Oku K, Bohgaki T, Yasuda S, et al. Pregnancy outcomes in women with rheumatic diseases: real-world observational study in Japan. Lupus. 2019;28:1407–16.

3. Kither H, Heazzel A, Bruce IN, Tower C, Crocker I. Adverse pregnancy outcomes and subsequent development of connective tissue disease in the UK: an epidemiological study. BJOG. 2020;127:941–9.

4. Tsuda S, Sameshima A, Sekine M, Kawaiuchi H, Fujita D, Makino, et al. Pre-conception status, obstetric outcome and use of medications during pregnancy of systemic lupus erythematosus (SLE), rheumatoid arthritis (RA) and inflammatory bowel disease (IBD) in Japan: multi-center retrospective descriptive study. Mod Rheumatol. 2019;24:1–10.

5. Clowse ME, Jamison M, Myers E, James AH. A national study of the complications of lupus in pregnancy. Am J Obstet Gynecol. 2008;199:127 e1–6.

6. Bundhun PK, Soogund MZ, Huang F. Impact of systemic lupus erythematosus on maternal and fetal outcomes following pregnancy: a meta-analysis of studies published between years 2001–2016. J Autoimmun. 2017;79:17–27.

7. Buyon JP, Kim MY, Guerra MM, Laskin CA, Petri M, Lockshin MD, et al. Predictors of pregnancy outcomes in patients with lupus: a cohort study. Ann Intern Med. 2015;163:153–63.

8. Deguchi M, Maesawa Y, Kubota S, Morizane M, Taniruma K, Ebina Y, et al. Factors associated with adverse pregnancy outcomes in women with systemic lupus erythematosus. J Reprod Immunol. 2018;125:39–44.

9. Robertson SE, Schumacher HR, Battle RM, et al. Biologic agents and pregnancy: a prospective study. J Rheumatol. 2014;41:1053–61.

10. Shimada H, Kaemda T, Kanenishi K, Miyatake N, Nakashima S, Wakiya R, et al. Factors affecting the Apgar score of offsprings born to mothers suffering from systemic lupus erythematosus. Medicine (Baltimore). 2020;99:e22843.

11. Sammaritano LR, Bermas BL, Chakravarty EE, Chambers C, Clowse ME, Lockshin MD, et al. 2020 American College of Rheumatology guideline for the management of reproductive health in rheumatic and musculoskeletal diseases. Arthritis Care Res. 2020;72:461–88.

12. Fung JT, Panchal S, Hurrell A, van de Venne M, Gayed M, Schreiber K, et al. BSR and BHPR guideline on prescribing drugs in pregnancy and breastfeeding-part I: standard and biologic disease modifying anti-rheumatic drugs and corticosteroids. Rheumatology (Oxford). 2016;55:1693–7.

13. Skorpen CG, Høienbein M, Tincani A, Fischer Betz R, Elalfet E, Chambers C, et al. The EULAR points to consider for use of antirheumatic drugs before pregnancy, and during pregnancy and lactation. Ann Rheum Dis. 2016;75:795–810.

14. Gur C, Diov-Citrin O, Shechtman S, Arnon J, Ormoy A. Pregnancy outcome after first trimester exposure to corticosteroids: a prospective controlled study. Reprod Toxicol. 2004;18:93–101.

15. Bandoli G, Palmsten K, Forbess Smith CJ, Chambers CD. A review of systemic corticosteroid use in pregnancy and the risk of select pregnancy and birth outcomes. Rheum Dis Clin N Am. 2017;43:489–502.

16. Reinisch JM, Simon NG, Karow RG, Gandelman R. Prenatal exposure to prednisone in humans and animals retards intrauterine growth. Science. 1978;202:436–8.

17. Shiozaki A, Yoneda S, Nakabayashi M, Takeda Y, Takeda S, Sugimura M, et al. Multiple pregnancy, short cervix, past gestation, and birth outcomes. Rheum Dis Clin N Am. 2017;44:53–61.

18. Palmsten K, Rolland M, Hebertt MF, Clowse ME, Schatz M, Xu R, et al. Patterns of prednisone use during pregnancy in women with rheumatoid arthritis: daily and cumulative dose. Pharmacoeconomics Drug Saf. 2018;27:430–8.

19. Shimada H, Kaemda T, Kanenishi K, Miyatake N, Nakashima S, Wakiya R, et al. Effect of biologic disease-modifying anti-rheumatic drugs for patients with rheumatoid arthritis who hope to become mothers. Clin Rheumatol. 2019;38:1453–8.

20. Charkvayt DF, Colon I, Langen ES, Nix DA, El-Sayed YY, Genovese MC, et al. Factors that predict prematurity and preeclampsia in pregnancies that are complicated by systemic lupus erythematosus. Am J Obstet Gynecol. 2005;192:1897–904.

21. Desai R, Bateman BT, Huybrechts KF, Patorno E, Hernandez-Diaz S, Park Y, et al. Risk of serious infections associated with use of immunosuppressive agents in pregnant women with autoimmune inflammatory conditions: cohort study. BMJ. 2017;356:j895.

Supplementary Information

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Authors’ contributions

HS and HD designed the study; HS and NM analyzed the data; HS wrote the manuscript; HD and NK revised the manuscript; KK, TK, SN, MMFM, RW, MK, TM, KS, YU, RM, and MM assisted in collecting the data. All authors read and approved the final manuscript.

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Availability of data and materials

The data that support the findings in this study will be shared on reasonable request to the corresponding author.

Declarations

Ethics approval and consent to participate

We did not obtain patients’ informed consent or study approval because this was a retrospective, observational study.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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References

1. Ohyama A, Tsuboi H, Noma H, Terasaki M, Shimizu M, Toko H, et al. Associations between maternal clinical features and fetal outcomes in pregnancies of mothers with connective tissue diseases. Mod Rheumatol. 2019;29:344–50.

2. Sugawara E, Kato M, Fujieda Y, Oku K, Bohgaki T, Yasuda S, et al. Pregnancy outcomes in women with rheumatic diseases: real-world observational study in Japan. Lupus. 2019;28:1407–16.
22. Østensen M, Forger F. How safe are anti-rheumatic drugs during pregnancy? Curr Opin Pharmacol. 2013;13:470–5.
23. Ponticelli C, Moroni G. Immunosuppression in pregnant women with systemic lupus erythematosus. Expert Rev Clin Immunol. 2015;11:549–52.
24. Mitchell K, Kaul M, Clovse ME. The management of rheumatic diseases in pregnancy. Scand J Rheumatol. 2010;39:99–108.
25. Palmsten K, Bandoli G, Watkins J, Vazquez-Benitez G, Gilmer TP, Chamber CD. Oral corticosteroids and risk of preterm birth in the California Medicaid. J Allergy Clin Immunol Pract. 2021;9:375–84 e5.
26. Hochberg MC. Updating the American College of Rheumatology revised criteria for the classification of systemic lupus erythematosus. Arthritis Rheum. 1997;40:1725.
27. Tanaka Y, Kuwana M, Fujiwara T, Kameda H, Muro Y, Fujio K, et al. Diagnostic criteria for mixed connective tissue disease (MCTD) from the Japan research committee of the ministry of health, labor, and welfare for systemic autoimmune diseases. Mod Rheumatol. 2019;7:1–5.
28. Bombardier C, Gladman DD, Urowitz MB, Caron D, Chang CH. Derivation of the SLEDAI, a disease activity index for lupus patients. The committee on prognosis studies in SLE. Arthritis Rheum. 1992;35:630–40.
29. Griffiths B, Mosca M, Gordon C. Assessment of patients with systemic lupus erythematosus and the use of lupus disease activity indices. Best Pract Res Clin Rheumatol. 2005;19:658–80.
30. Franklyn K, Lau CS, Navarra SV, Louthrenoo W, Latee A, Hamijoya L, et al. Definition and initial validation of a lupus low disease activity state (LLDAS). Ann Rheum Dis. 2016;75(9):1615–21.

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