Rituximab, MS, and pregnancy

Jessica B. Smith, MPH, Kerstin Hellwig, MD, Katharina Fink, MD, PhD, Deirdre J. Lyell, MD, Fredrik Piehl, MD, PhD, and Annette Langer-Gould, MD, PhD

Neurol Neuroimmunol Neuroinflamm 2020;7:e734. doi:10.1212/NXI.0000000000000734

Correspondence
Dr. Langer-Gould
Annette.M.Langer-Gould@kp.org

Abstract

Objective
To describe the safety and efficacy of rituximab (RTX) in MS and pregnancy, we conducted a retrospective cohort study of 74 pregnancies among 55 women treated with RTX for MS and their offspring.

Methods
We used prospectively collected information from the electronic health record at Kaiser Permanente Southern California between 2012 and 2019 of mother and baby to identify treatment history, pregnancy outcomes, and relapses.

Results
Last RTX exposure before conception occurred between 1.8 and 5.2 months in 32 (49%) of 65 pregnancies and accidentally during the first trimester in 9 (12%). Among 38 live births, adverse pregnancy outcomes were as follows: 3 preterm deliveries (including 1 set of twins), 1 neonatal death (preterm twin), and 1 perinatal stroke (full-term). No stillbirths, chorioamnionitis, or major malformations were found. Fifteen (27%) women had at least one first-trimester miscarriage, of whom 8 (53%) had a history of infertility. Cumulative dose or timing of last RTX infusion was not associated with an increased risk of miscarriage. Only 2 (5.4%) women experienced relapses, one during pregnancy and the other postpartum.

Conclusion
We observed no increase in adverse pregnancy outcomes compared with expected national incidence rates and remarkably little disease activity in RTX-treated women with MS, particularly when compared with periconceptional natalizumab-treated cohorts. However, larger studies are needed to fully assess the safety of RTX use before pregnancy, especially risks associated with prolonged B-cell depletion and hypogammaglobulinemia. Until these data are available, we recommend restricting RTX use before pregnancy to women who require highly effective MS treatments.

Classification of evidence
This study provides Class IV evidence that for pregnant women with MS, RTX controls disease activity and does not increase adverse pregnancy outcomes.
Neurology: Neuroimmunology & Neuroinflammation | Volume 7, Number 4 | July 2020 Neurology.org/NN

We recently found that 67% of women with MS entered pregnancy with suboptimally controlled disease in our large, population-based cohort. These findings led to the development of a pregnancy-specific MS treatment algorithm in KPSC that prioritizes the use of rituximab (RTX) in these women over fingolimod, natalizumab, or other highly effective disease-modifying treatments (DMTs).

RTX is an attractive treatment option in women with MS who desire pregnancy and require a highly effective DMT because its immunomodulatory effects last long after the drug has been eliminated, and RTX is not associated with drug-cessation relapses. We recommend starting RTX before pregnancy, typically 500 mg every 6 months, holding infusions during pregnancy and resuming infusions typically 6–12 months postpartum or sooner if disease activity returns or is desired by the mother.

We based these recommendations mainly on the high risks of using other highly effective DMTs and the low biological plausibility of harm with relatively low doses of RTX before pregnancy. However, data regarding safety and efficacy of RTX in human pregnancies are scant, particularly in MS. No studies were adequately designed to assess the effect of RTX on miscarriages or stillbirths. This is important because a plausible risk of prolonged B-cell depletion from high-dose RTX or other B-cell depleting agents could increase the risk of systemic maternal infections or chorioamnionitis, leading to miscarriages, preterm birth, or stillbirths. It is concerning that the drug company’s global safety database reported a high number of preterm births; however, these data are difficult to interpret because they combined outcomes from women with a wide variety of underlying diseases, some of which were predisposed to poor pregnancy outcomes, received RTX during late pregnancy, and/or were coadministered teratogenic medications. Other significant methodological limitations precluding clear interpretation of the manufacturer’s study and a subsequent systematic review include reporting bias, selection bias, missing data, and reliance on case reports.

The objective of this study was to assess the risk of miscarriages, adverse pregnancy and neonatal outcomes, and maternal disease control in a well-defined cohort of women with MS treated with RTX.

Methods

Study population

Pregnant women with MS were identified through the membership of Kaiser Permanente Southern California (KPSC). We searched electronic databases to identify members who received RTX infusions for MS or its potential precursors, clinically isolated syndrome (CIS) or radiologically isolated syndrome (RIS), between January 2012 and November 2019. We used a combination of International Classification of Diseases, 9th and 10th Revisions codes for MS, optic neuritis, and transverse myelitis and pregnancy to identify potentially eligible subjects. The complete electronic health records (EHRs) were then reviewed to determine eligibility by an MS expert (A.L.-G.). All KPSC members who met the 2017 McDonald criteria for MS, CIS, or RIS at the onset of pregnancy and had received at least 1 RTX infusion before conception or within 2 months of last menstrual period (LMP) were included. Women who had no history of RTX infusions before or around the time of conception or had conditions other than MS, CIS, or RIS were excluded.

KPSC is a large prepaid healthcare organization that provides comprehensive health care, including inpatient, outpatient, pharmacy, laboratory, and radiology services to over 4 million members in Southern California. The membership of KPSC is representative of the general Southern California population. KPSC uses an integrated EHR system that includes all inpatient and outpatient encounters, diagnostic tests, diagnoses and medications, and some demographic and behavioral characteristics.

Standard protocol approvals, registrations, and patient consents

The study protocol was approved by the KPSC institutional review board.

Data collection

Symptom onset, relapses, MRI disease activity, and disability level were abstracted from the mother’s EHR by an MS expert (A.L.-G.). Relapses were defined as the occurrence, reappearance, or worsening of symptoms of neurologic dysfunction lasting for 48 hours or more and needed to be documented by a treating physician during a physical examination. Symptoms that occurred within 1 month of each other were part of the same relapse. MRIs were obtained as part of routine clinical care. MRI disease activity was defined as an unequivocally new or enlarging T2 lesion, new diffusion-restricting or gadolinium-positive lesion(s). Comparisons were made with previous

Glossary

BMI = body mass index; CIS = clinically isolated syndrome; DMT = disease-modifying treatment; EDD = expected delivery date; EHR = electronic health record; gw = gestational week; IQR = interquartile range; KPSC = Kaiser Permanente Southern California; LMP = last menstrual period; RA = rheumatoid arthritis; RCT = randomized controlled trial; RIS = radiologically isolated syndrome; RTX = rituximab.
for comparison with other populations.1,20 For graphical de-
excluded from the maternal disease course outcomes to allow
fi
Pregnancies ending in the
nancy during the study period to account for women with
comes, we describe the maternal characteristics at
statistician (A.L.-G.). Twenty of 34 (58.8%) postpartum MRIs were
fied before or during each pregnancy.

Risk factors for adverse pregnancy outcomes (infertility, comorbidities, body mass index (BMI) at conception ≥25,15
smoking, black race, substance use, thrombophilia, thyroid disease, and infection), LMP, expected delivery date (EDD),
gravity, parity and pregnancy, and labor and delivery outcomes were abstracted from maternal EHRs by a re-
arch professional (J.B.S.). Neonatal outcomes, breast-
feeding, formula feeding, and introduction of solid food were
abstracted from the infant’s medical record as previously
described.1

Statistical analyses
The primary purpose of analyses was to describe adverse pregnancy outcomes and maternal disease control in women
with MS being treated with RTX. To allow for comparison
with the national and California (whenever available) rates,
we used the Centers for Disease Control definitions for low
birthweight (less than 2,500 g),16 very low birthweight (less
than 1,500 g),16 small for gestational age (SGA, birthweight at
or below the 10th percentile for a given gestational age),17
stillbirth (death or loss of baby after 20 weeks completed
pregnancy weeks),18 preterm delivery (less than 37 weeks
gestational age),19 and major malformations. We were also
interested in chorioamnionitis or systemic maternal infections
that could potentially lead to adverse pregnancy outcomes
given the mechanism of action of RTX. Therefore, we de-
scribe RTX dosing before or during each pregnancy.

When examining the risk factors for adverse pregnancy out-
comes, we describe the maternal characteristics at first preg-
nancy during the study period to account for women with
multiple pregnancies.

Pregnancies ending in the first-trimester miscarriages were
excluded from the maternal disease course outcomes to allow
for comparison with other populations.1,20 For graphical de-
piction purposes, we divided the prepartum and postpartum
years into 3-month intervals.

To examine the potential impact of resuming RTX on breast-
feeding choices, we used previously published definitions.1,21
Breastfeeding was defined as exclusive (no regular formula
feeding for at least the first 2 months postpartum), nonexclusive
(breastfeeding and regular formula feeding within 2 months),
or none.

The baby of one woman who delivered outside of KPSC, but
whose postpartum neurology notes and date of delivery were
available, was excluded from descriptions of birthweight,
major malformations, and breastfeeding because we were not
able to access the infant records.

Women or pregnancies that were not eligible for the outcome
described were excluded from the denominator when calcu-
ling percentages. For instance, first-trimester miscarriages,
elective abortions, and ongoing pregnancies were excluded
when describing infant outcomes.

The means and SDs of normally distributed variables were
compared using 2 sample t tests; for variables with non-
normal distributions, the Wilcoxon rank-sum test was used;
and for binary or categorical variables, χ² with the Fisher exact
test was used to compare periconceptional RTX exposure
characteristics in pregnancies ending or not ending in mis-
carriages and clinical characteristics of the mothers with or
without at least one post-RTX exposure miscarriage. Statis-
tical significance was set a priori at p = 0.20 to detect potential
confounders, given the small sample and purpose of analyses
to detect any signal of adverse pregnancy outcomes.22 No
adjustment for multiple comparisons was made. All statistical
analyses were performed using SAS version 9.4 (SAS Institute
Inc, Cary, NC).

Data availability
Owing to KPSC’s institutional review board, data would be
available upon reasonable request.

Results
Disease characteristics, risk factors for adverse pregnancy outcomes, and periconceptional RTX exposure
We identified 55 women with MS who conceived 74 pregnan-
cies after exposure to at least one dose of RTX during the study
period. Seventeen women had 2 pregnancies, and one had 3
pregnancies. General characteristics of the women with MS at
the onset of their first RTX-exposed pregnancy are presented in
table 1. Many women with MS (n = 40) had previously been
treated with other DMTs, of whom 16 (29.1%) were clinically
and radiographically stable on their previous DMT and switched
to RTX solely for pregnancy planning. Thirty-four (62%) women were nulliparous. Sixteen women (29%) had been
diagnosed with infertility, 5 from polycystic ovarian syndrome, 2
from structural causes, and 1 each from premature ovarian fail-
ure, age-related infertility, and factor V Leiden deficiency. In 6
women, the cause of infertility was unknown.

Forty-six (84%) women had at least one risk factor for
adverse pregnancy outcomes, including miscarriages be-
fore starting RTX (n = 11, 20%), overweight (BMI ≥25,
n = 33, 60%), age ≥35 at EDD (n = 15, 27%), being black
(n = 10), gestational diabetes (n = 5), thyroid dysfunction
(n = 2), and 1 each of the following: diabetes before
pregnancy, smoking during pregnancy, twin pregnancy, or
retained intrauterine device.
Most pregnancies (n = 59, 80%) were conceived within 6 months after the last RTX infusion, including 9 pregnancies with accidental 1st trimester exposure (table 2). Only 6 pregnancies occurred more than 12 months after the last dose, including a woman with RIS who received one infusion with plans for further treatment only on return of disease activity. RTX exposure during pregnancy was uncommon, occurring accidentally in 9 (12%) pregnancies during the first trimester with a median time from LMP of 2 weeks, ranging from 0.5 to 5 weeks after LMP. No pregnancies were exposed to RTX infusions during the second or third trimester. Half of the pregnancies were conceived after 3 or more RTX infusions, with a median cumulative dose of 2,000 mg (table 2).

Pregnancy, labor, delivery, and neonatal outcomes

Pregnancy outcomes after at least 1 dose of RTX are shown in figure 1. Fifteen (27%) women had 23 first-trimester miscarriages; 6 women had 2 and 1 had 3. No miscarriages after 12 weeks or stillbirths occurred. A diagnosis of infertility was the only adverse pregnancy risk factor measured that differed significantly between women who had pregnancies ending in miscarriages (53%, n = 8) compared with those who did not (20%, n = 8; p = 0.0223). Cumulative RTX dose, duration of RTX, the number of infusions, accidental 1st trimester exposure, and timing of the last infusion before pregnancy were all similar (p > 0.20) among pregnancies ending in miscarriages compared with those ending in full-term or preterm births or elective abortions (appendix e-1, links.lww.com/NXI/A246). Five (33%) of the 15 women with miscarriages had successful pregnancies on RTX, and another 2 (13%) women are currently in the third trimester. None of the pregnancies were discontinued RTX because of miscarriages or other adverse pregnancy outcomes.

Of the 45 pregnancies that advanced past the first trimester, 37 resulted in 38 live births and 8 are ongoing (3 in the 2nd trimester and 5 in the 3rd trimester; figure 1). Most of the live births were normal, full-term babies (n = 33, 89%). The average birthweight among full-term neonates was 3,217.2 g (SD 346.3). None of the pregnancies were complicated by chorioamnionitis.

| Table 1 Characteristics of women with MS at the onset of first RTX-exposed pregnancy |
|---------------------------------|------------------|
| **n = 55**                      |                  |
| **Age, median (IQR), y**        | 31.1 (28.0–34.5) |
| **Age ≥35, n (%)**              | 15 (27.3)        |
| **Race/ethnicity, n (%)**       |                  |
| White                           | 21 (38.2)        |
| Hispanic                        | 22 (40.0)        |
| Black                           | 10 (18.2)        |
| Asian                           | 2 (3.6)          |
| **Obesity, n (%)**              |                  |
| BMI at conception, median (IQR) | 26.6 (22.9–30.5) |
| Normal weight (<25)             | 22 (40.0)        |
| Overweight (≥25 to <30)         | 18 (32.7)        |
| Obese Class I (≥30 to <35)      | 9 (16.4)         |
| Obese Class II (≥35 to <40)     | 2 (3.6)          |
| Obese Class III (≥40)           | 4 (7.3)          |
| **Infertility, n (%)**          | 16 (29.1)        |
| **Previous miscarriages, n (%)**| 11 (20.0)        |
| **Gravidity, n (%)**            |                  |
| 1                               | 26 (47.3)        |
| 2                               | 16 (29.1)        |
| 3+                              | 13 (23.6)        |
| **Parity, n (%)**               |                  |
| 0                               | 34 (61.8)        |
| 1                               | 14 (25.5)        |
| 2+                              | 7 (12.7)         |
| **MS subtype, n (%)**           |                  |
| RRMS                            | 54 (98.2)        |
| RIS                             | 1 (1.8)          |
| **Disease duration, median (IQR), y** | 5.3 (2.6–8.7)  |
| **No. of relapses 2 year prepregnancy, n (%)** | 0 (50.9) |
| 1                               | 16 (29.1)        |
| ≥2                              | 11 (20.0)        |
| **EDSS, median (IQR)**          | 2.0 (1.0–3.0)    |
| **DMT use before RTX, n (%)**   |                  |
| None                            | 15 (27.3)        |
| Fingolimod                      | 4 (7.3)          |
| Natalizumab                     | 14 (25.5)        |

Abbreviations: BMI = body mass index; DMT = disease-modifying treatment; EDSS = expanded disability status scale; IQR = interquartile range; IFN = interferon; RIS = radiologically isolated syndrome; RRMS = relapsing-remitting MS; RTX = rituximab.

*n = 2 dimethyl fumarate, n = 1 cyclophosphamide and n = 1 azathioprine.
Three pregnancies (8.1%) ended in preterm deliveries of 4 babies, including one set of 27-gestational-week (gw) twins. The other preterm infants were noted as a normal spontaneous vaginal delivery (36 gw) or premature rupture of membranes from incompetent cervix (34 gw).

The 27-week twins were both <1,500 g, and 1 twin died 6 days after delivery from an intraventricular hemorrhage. Another preterm baby weighed <2,500 g, but no babies were SGA. In addition to these 3 preterm, low birthweight babies, another baby was admitted to the neonatal intensive care unit for perinatal ischemic stroke and seizures after vacuum-assisted delivery for prolonged labor and fetal distress after a 40-week pregnancy complicated only by mild pre-eclampsia. Extensive workup to identify a cerebrovascular, cardiac, or genetic etiology of perinatal stroke was negative.

The last RTX dose before pregnancy was 2.3 months for the mother of the twins, 9.5 months for the mother of the neonate with stroke and seizures, and 2 days after LMP for the mother of the 34 gw preterm baby. The mother with twins had a history of infertility from premature ovarian failure and had failed multiple rounds of in vitro fertilization before starting RTX.

Caesarean sections were performed in 13 (35%) pregnancies, of which 6 were emergencies. The remaining 24 (65%) pregnancies ended with vaginal deliveries.

**Maternal disease course, pregnancy, breastfeeding, and postpartum RTX use**

Very few relapses or MRI disease activity occurred after the first dose of rituximab prepartum, intrapartum, or during the postpartum period in women with live births (figure 2). Only 1 (2.7%) relapse occurred during pregnancy with the most recent RTX dose 3 months before conception and 1 (2.7%) during the 12-month postpartum where the most recent RTX dose was 4 months before conception. These 2 women with relapses resumed RTX postpartum after their relapses. No postpartum MRI disease activity was detected among the 22 (59%) women who had an MRI in the postpartum year.

RTX was resumed after 24 (65%) pregnancies ending in live births within 12 months postpartum (figure 2). RTX was not resumed after 9 pregnancies (24%) within 12 months postpartum, none of whom had a relapse or active MRI, and 4 have not resumed RTX but are not yet 1 full year postpartum. Among those resuming RTX in the postpartum year, the median time to restarting was 2.2 months (interquartile range [IQR] 1.0–5.7) postpartum and the median interval from last RTX dose to first postpartum dose was 14.6 months (IQR 11.3–18.4).

Most babies were breastfed (89%, n = 34), of which 13 (35%) were breastfed exclusively for at least 2 months or more. Eight (22%) babies were still being breastfed when their mothers resumed RTX infusions.

No postpartum relapses occurred after the 23 first-trimester miscarriages during a median follow-up time of 12 months (IQR 4–12). RTX was resumed during this timeframe after 13 (56.5%) miscarriages.

**Discussion**

Our findings show that using RTX 500 mg every 6–12 months before pregnancy and simply holding infusions during pregnancy results in remarkably well-controlled MS disease activity without a signal for increased risk of harm to the baby. RTX was used primarily in women with suboptimally controlled MS or in those controlled on fingolimod or natalizumab who desired pregnancy, consistent with the KPSC algorithm. The findings are reassuring for these women who are at high risk of pregnancy-related relapses and/or long-term MS-related disability but are insufficient to recommend expanding routine RTX use to women at low risk of long-term disability or pregnancy-related relapses. Specifically, the women in this study received relatively few and low doses of RTX; thus, potential risks of longer term, higher dose use of RTX on adverse pregnancy outcomes, particularly maternal or neonatal infections, remain unknown. Our findings should also not be extrapolated to other B-cell depleting treatments, where the risk of hypogammaglobulinemia appears to be significantly higher and occurs earlier than with the doses of RTX used in this study.

The KPSC MS and Pregnancy Treatment algorithm was developed because, in 2017, we found that 67% of women in KPSC and KP Northern California had suboptimally controlled disease entering pregnancy, yet very few had been treated with
highly effective DMTs. Optimizing disease control before pregnancy is important because MS-related disability is a predictor of poor long-term prognosis and an independent predictor of postpartum relapses, along with the prepregnancy relapse rate and relapses during pregnancy. We prioritize RTX use in these women because the other commonly used, highly effective DMTs, natalizumab and fingolimod, are associated with an increased risk of rebound disease activity. These rebound relapses can be severe and, if they occur during pregnancy, necessitate treatments that carry some risk of adverse pregnancy and neonatal outcomes, including repeated courses of glucocorticoids, resuming natalizumab, or starting RTX during late pregnancy. In addition, fingolimod is a weak teratogen with a long half-life; thus, discontinuing before conception is imperative. Although we and others have continued natalizumab throughout pregnancy when the risk of a severe MS-rebound relapse was judged to be high, this is not an optimal approach because natalizumab use during the third trimester can result in neonatal cytopenia with uncertain risks, including intraventricular hemorrhages. Therefore, we reasoned that it is more prudent to switch women who desire pregnancy from fingolimod or natalizumab to RTX before conception.

In this study, we show that periconceptional use of RTX resulted in significantly fewer relapses during pregnancy and the postpartum period compared with our previously published contemporary cohort (2.7% vs 8.6% during pregnancy and 2.7% vs 26.4% postpartum, respectively). This remarkably well-controlled disease activity starkly contrasts with the relatively high risk of pregnancy-related relapses with periconceptional natalizumab use, where 36.5% relapsed during pregnancy and 22% relapsed in the first 3 months postpartum alone. We strongly encourage women to breastfeed exclusively, given the plethora of maternal and infant health benefits. Although exclusive breastfeeding appears to reduce the risk of postpartum MS relapses, the rates of exclusive breastfeeding were similar in this cohort compared with our previous study; thus, an unlikely explanation for the disease protection afforded by RTX.

We consider RTX to be compatible with breastfeeding and recommend restarting infusions 6–12 months postpartum or sooner if disease activity returns. However, in this cohort, half the women restarted RTX within 6 months postpartum.

Figure 1 Pregnancy outcomes after RTX exposure in women with MS

Depicted are the pregnancy outcomes of women with MS who were exposed to at least 1 dose of RTX before known pregnancy. Fifteen women had 23 first-trimester miscarriages. No miscarriages occurred during the second trimester. One preterm birth at 36w, followed a normal pregnancy with NSVD. Two pregnancies ended with PROMs, resulting in delivery of 3 preterm infants, all of whom were low birthweight but were not small for gestational age. The twins were of very low birthweight. One twin died from an intraventricular hemorrhage 6 days after birth. Among the full-term infants, 1 born by vacuum-assisted vaginal delivery had a stroke and seizures. There were no cases of chorioamnionitis and no adverse pregnancy outcomes due to systemic maternal infections.

In this study, we show that periconceptional use of RTX resulted in significantly fewer relapses during pregnancy and the postpartum period compared with our previously published contemporary cohort (2.7% vs 8.6% during pregnancy and 2.7% vs 26.4% postpartum, respectively). This remarkably well-controlled disease activity starkly contrasts with the relatively high risk of pregnancy-related relapses with periconceptional natalizumab use, where 36.5% relapsed during pregnancy and 22% relapsed in the first 3 months postpartum alone. We strongly encourage women to breastfeed exclusively, given the plethora of maternal and infant health benefits. Although exclusive breastfeeding appears to reduce the risk of postpartum MS relapses, the rates of exclusive breastfeeding were similar in this cohort compared with our previous study; thus, an unlikely explanation for the disease protection afforded by RTX.

We consider RTX to be compatible with breastfeeding and recommend restarting infusions 6–12 months postpartum or sooner if disease activity returns. However, in this cohort, half the women restarted RTX within 6 months postpartum.
although only one had signs of the return of disease activity. Similarly, this was because of a cautionary approach among clinicians and/or patients regarding the risks of being untreated. To what extent resuming RTX influenced the quiescence of disease activity we observed 6–12 months postpartum is, therefore, unclear.

Similar to other studies, we did not observe any rebound relapses in RTX-treated women, even when treatment intervals exceeded 2 years. That extending RTX intervals beyond 1 year is effective was first described in the RTX relapsing-remitting MS randomized controlled trial (RCT) where efficacy was sustained in most patients after 2,000 mg induction. A recent RCT found that 77% of patients were still relapse-free 3 years after a single dose of 1,000 mg, followed by glatiramer acetate. What role an intervening pregnancy plays in the efficacy of extended dosing intervals we observed is unclear.

Findings from previous studies strongly suggested that every 6-month infusion of RTX at doses higher than those used in this study increases the risk of harm without added benefit in most patients. Safety advantages of using lower doses and longer dosing interval for RTX and other B-cell depleting agents are fewer infections and less hypogammaglobulinemia. Maternal hypogammaglobulinemia would be expected to increase the risk of neonatal and maternal infections because the placental transfer of maternal antibodies is a critical source of humoral immunity in neonates. B cells play a broader, yet unclear, role in maintaining or disrupting normal pregnancies. Recent advances in studies of regulatory B-cell populations indicate shifting dynamics during pregnancy. That this could affect repopulation of specific B-cell subsets after pregancy B-cell depletion seems plausible, but whether this would result in improved or worsened pregnancy outcomes remains murky. Future studies examining immunoglobulin levels, B-cell subset repopulation, and pregnancy and neonatal outcomes are needed.

We did not find any concerning safety signals. None of the women developed serious infections or chorioamnionitis. The rates of other pregnancy complications potentially signaling occult infections, including late miscarriages, stillbirths, or low birthweight, were comparable with, or lower than, the...
expected rates. Specifically, we observed 8.1% of pregnancies ending in preterm births and 7.9% low birthweight neonates with only 1 (2.8%) low birthweight singleton. The expected national and California-specific rates are 10% and 8.8% for preterm births and 8.3% and 7.0% for low birthweight babies, respectively, and 6.6% for low birthweight singletons nationally. No major malformations (3% expected nationally), terminations because of congenital anomalies, or stillbirths (1% expected nationally) were found. Two cases of stillbirths after B-cell depleting treatments in women with MS have been reported, 1 following RTX and another with ocrelizumab, but whether these women had other risk factors for adverse pregnancy outcomes and whether this exceeds the expected rates are unknown. The 2 significant adverse pregnancy outcomes, neonatal death and perinatal stroke, do not appear to be RTX related. Neonatal death is a known complication of extremely preterm twin pregnancies. Perinatal ischemic stroke occurs in 1 per 3,500 live births, and an idiopathic etiology is common. Future studies should continue to examine these potential risks.

RTX is our treatment of choice in women with active MS who are struggling with infertility because it is unclear if or when they will become pregnant. This is why the proportion of RTX-treated women with infertility in our study is relatively high. This is also the most likely explanation for why we observed a relatively high rate of miscarriages because infertility and miscarriages are comanifestations of subfertility. Thus, it is not surprising that we found that a history of infertility was more common in women with miscarriages than those without. We believe it is probably not related to RTX exposure because the characteristics of RTX use did not differ between pregnancies ending in miscarriages and those advancing to the third trimester and beyond. It is also reassuring that many women with miscarriages went on to have successful pregnancies despite continued RTX treatment. Nevertheless, given the possible role of regulatory B-cell subpopulations in implantation and maintenance of successful pregnancies, more studies are needed to accurately assess this potential risk.

The main limitation of this study is the relatively small sample size. We believe this is the most likely explanation for the lower than expected numbers of major malformations, stillbirths, preterm births, and low birthweight babies. The rates of pregnancy complications are particularly low when considering that a high proportion of women had at least 1 risk factor for adverse pregnancy outcomes. Alternatively, high-quality prenatal care could explain these findings. Strengths of this study are a population-based sample with comprehensive medical records of the mother and infant, allowing unbiased and thorough assessment of maternal and infant outcomes and risk factors.

Although our findings are reassuring for women with suboptimally controlled MS who desire pregnancy, we caution against extending these findings to all women with MS or to treatment with ocrelizumab. Women diagnosed with MS today appear to have milder disease before, during, and after pregnancy compared with historical cohorts, and some are adequately controlled on glatiramer acetate, interferon-beta, or no treatment. Exposing these women to even a small increased risk of adverse pregnancy outcomes from routine RTX infusions likely outweighs any benefit they would experience. The risk of maternal and neonatal infections and ensuing adverse pregnancy outcomes could be higher with ocrelizumab compared with our findings. This is quite plausible because RTX is FDA-approved for rheumatoid arthritis (RA) without significant complications, but the ocrelizumab RA development program was ended because of the increased risk of serious infections. The FDA-approved MS ocrelizumab dose of 600 mg every 6 months is substantially higher than those used in the RA development program.

Taken together, our findings indicate that periconceptional RTX is a highly effective and safe treatment of women with suboptimally controlled MS or those on fingolimod or natalizumab. These women should start or switch to RTX before pregnancy, should be encouraged to breastfeed exclusively to maximize the general maternal and infant health benefits, and may not need to resume RTX until much later in the postpartum period. Future studies to determine the minimum RTX dosing interval are needed to optimize safety, particularly in young women who will want to get pregnant in the more distant future.

Acknowledgment
We thank Dr. Dean P. Sarco (Los Angeles Medical Center, Department of Neurology, Southern California Permanente Medical Group) for assistance reviewing the record of the infant with perinatal stroke.

Study funding
This study was supported in part by Patient-Centered Outcomes Research Institute (PCORI/MS-1511-33,196 PIs: Langer-Gould and Piehl).

Disclosure
J. Smith reports no disclosures relevant to the manuscript. K. Hellwig has received speaker and research honoraria from Biogen Idec, Germany, Teva, Sanofi Genzyme, Novartis, Bayer Healthcare, Merck Serono, and Roche. K. Fink has received research grants from Biogen, served on advisory boards for Roche and Alexion, and has held a lecture for Biogen. D.J. Lyell serves as a consultant for Bloomlife Technology and an investor for Zenflow. F. Piehl has received research grants from Genzyme, Merck KGaA, and Novartis and fees for serving as Chair of DMC in clinical trials with Parexel. A. Langer-Gould has received grant support and awards from the National Institutes of Health, Patient-Centered Outcomes Research Institute, and the National MS Society. She currently serves as a voting member on the California Technology Assessment Forum, a core program of the Institute for Clinical and Economic Review (ICER). She
has received sponsored and reimbursed travel from ICER. Go to Neurology.org/NN for full disclosures.

Publication history
Received by Neurology: Neuroimmunology & Neuroinflammation December 26, 2019. Accepted in final form March 24, 2020.

Appendix Authors

| Name                | Location                                                                 | Contribution                          |
|---------------------|--------------------------------------------------------------------------|---------------------------------------|
| Jessica B. Smith, MPH | Department of Research & Evaluation, Kaiser Permanente Southern California, Pasadena, CA | Collected, analyzed, and interpreted the data; revised the manuscript for content |
| Kerstin Hellwig, MD | Department of Neurology, St. Joseph Hospital, Bochum, Germany            | Interpreted the data; revised the manuscript for content |
| Katharina Fink, MD, PhD | Department of Clinical Neuroscience, Karolinska Institutet and Center for Neurology, Academic Specialist Center, Stockholm, Sweden | Interpreted the data; revised the manuscript for content |
| Deirdre J. Lyell, MD | Department of Obstetrics & Gynecology – Maternal Fetal Medicine, Stanford University School of Medicine, Palo Alto, CA, USA | Interpreted the data; revised the manuscript for content |
| Frederik Pielh, MD, PhD | Department of Clinical Neuroscience, Karolinska Institutet and Department of Neurology, Karolinska University Hospital, Stockholm, Sweden | Interpreted the data; revised the manuscript for content |
| Annette Langer-Gould, MD, PhD | Los Angeles Medical Center, Department of Neurology, Southern California Permanente Medical Group, Los Angeles, CA, USA | Designed and conceptualized the study; collected, analyzed, and interpreted the data; drafted and revised the manuscript for intellectual content; had full access to all the data and takes responsibility for the integrity of the data and accuracy of the data analysis |

References

1. Langer-Gould A, Smith JB, Albers KB, et al. Pregnancy-related relapses and breastfeeding in a contemporary multiple sclerosis cohort. Neurology Epub 2020 April 13.
2. Langer-Gould AM. Pregnancy and family planning in multiple sclerosis. Continuum (Minneap Minn) 2019;25:773–792.
3. Carton G, Blasco H, Painaud G, Waubant E, Tzou M, Le Guellec C. Pharmacokinetics of rituximab and its clinical use: thought for the best use? Crit Rev Oncol Hematol 2007;62:43–52.
4. Juto A, Fink K, Al Nimer F, Piehl F. Interrupting rituximab treatment in relapsing-remitting multiple sclerosis; no evidence of rebound disease activity. Mult Scler 2019;25:1360.
5. Honce JM, Nair KV, Sillu S, et al. Rituximab vs placebo induction prior to glatiramer acetate monotherapy in multiple sclerosis. Neurology 2019;92:e232–e32.
6. Hauser SL, Waubant E, Arnold DL, et al. B-cell depletion with rituximab in relapsing-remitting multiple sclerosis. N Engl J Med 2008;358:676–688.
7. Meint I, Havla J, Hohlfeld R, Kumpfel T. Recurrence of disease activity during pregnancy after cessation of fingolimod in multiple sclerosis. Mult Scler 2018;24:991–994.
8. Chakravarty EF, Murray ER, Kelman A, Farmer P. Pregnancy outcomes after maternal exposure to rituximab. Blood 2011;117:1499–1506.
9. Das G, Damotte V, Gelfand JM, et al. Rituximab before and during pregnancy: a systematic review, and a case series in MS and NMO/MS. Neurol Neuroimmunol Neuroinflamm 2018;5:e453. 10.1212/NXI0000000000453.
10. Beaber BE, Chi MD, Braza SM, Zhang JH, Langer-Gould AM. Immunomodulatory agents and risk of postpartum multiple sclerosis relapses. Perin J 2014;18:9–13.
11. Thompson AJ, Banwell Bi, Barkhof F, et al. Diagnosis of multiple sclerosis: 2017 revisions of the McDonald criteria. Lancet Neurol 2018;17:162–173.
12. Langer-Gould A, Braza SM, Beaber BE, Zhang JL. The incidence of clinically isolated syndrome in a multi-ethnic cohort. J Neurol 2014;261:1349–1355.
13. Okuda DT. Radiologically isolated syndrome: MR imaging features suggestive of multiple sclerosis prior to first symptom onset. Neuroimaging Clin N Am 2017;27:267–275.
14. Koebnick C, Langer-Gould A, Gould MK, et al. Do the sociodemographic characteristics of members of a large, integrated health care system represent the population of interest? Perin J 2012;16:37–41.
15. Metwally M, Ong KJ, Ledger WL, Li TC. Does high body mass index increase the risk of miscarriage after spontaneous and assisted conception? A meta-analysis of the evidence. Fertil Steril 2008;90:714–726.
16. Martin JA, Hamilton BE, Osterman MJ, Driscoll AK, Drake P. Births: Final Data for 2018. Natl Vital Stat Rep 2019;68(8).
17. Physical status: the use and interpretation of anthropometry. Report of a WHO Expert Committee. World Health Organ Tech Rep Ser 1995;85:41–452.
18. Centers for Disease Control and Prevention. What is Stillbirth? [online]. Available at: www.cdc.gov/nchcddl/stillbirth/facts.html. Accessed November 26, 2019.
19. Centers for Disease Control and Prevention. Preterm Birth [online]. Available at: www.cdc.gov/reproductivehealth/maternalinfanthealth/pretermbirth.htm. Accessed November 26, 2019.
20. Confavreux C, Hutchinson M, Hours MM, Cortinovis-Tourniaire P, Moreau T. Rate of pregnancy-related relapse in multiple sclerosis. Pregnancy in Multiple Sclerosis Group. N Engl J Med 1998;339:285–291.
21. Hellwig K, Rockhoff M, Herbstrit S, et al. Exclusive breastfeeding and the effect on postpartum multiple sclerosis relapses. JAMA Neurol 2015;72:1132–1138.
22. Greenland S, Pearce N. Statistical foundations for model-based adjustments. Annu Rev Public Health 2015;36:89–108.
23. Derfus T, Weber MS, Hughes R, et al. Serum immunoglobulin levels and risk of serious infections in the pivotal Phase III trials of ocrelizumab in multiple sclerosis and their open-label extensions. Abstract presented at the ECTRIMS 2019 Congress; September 11–13, Stockholm, Sweden. ECTRIMS Online Library 2019.
24. Hellborg S, Borelmalm M, Evertsson B, et al. Risk of hypogammaglobulinemia in long-term treatment with rituximab in multiple sclerosis. Abstract presented at the ECTRIMS 2019 Congress; September 11–13, Stockholm, Sweden. ECTRIMS Online Library 2019.
25. Langer-Gould A, Popat RA, Huang SM, et al. Clinical and demographic predictors of long-term disability in patients with relapsing-remitting multiple sclerosis: a systematic review. Arch Neurol 2006;63:1686–1691.
26. Portaccio E, Ghezzi A, Hakibi K, et al. Postpartum relapses increase the risk of disability progression in multiple sclerosis: the role of disease modifying drugs. J Neurol Neurosurg Psychiatry 2014;85:845–850.
27. Vukusic S, Hutchinson M, Hours M, et al. Pregnancy and multiple sclerosis (the PRMS study): clinical predictors of post-partum relapse. Brain 2004;127:1353–1360.
28. Hughes SE, Spelman T, Gray OM, et al. Predictors and dynamics of postpartum relapses in women with multiple sclerosis. Mult Scler 2014;20:739–746.
29. Haghiakia A, Langer-Gould A, Rellensmann G, et al. Natalizumab use during the third trimester of pregnancy. JAMA Neurol 2014;71:791–795.
30. Novi G, Ghezzi A, Pizzorno M, et al. Dramatic rebounds of MS during pregnancy following fingolimod withdrawal. Neurol Neuroimmunol Neuroinflamm 2017;4:e377. 10.1211/NXI0000000000377.
31. Portaccio E, Mosela L, Martellini V, et al. Pregnancy decision-making in women with multiple sclerosis treated with natalizumab: II. maternal risks. Neurology 2018;90:832–839.
32. Victoria CG, Bah R, Barros AJ, et al. Breastfeeding in the 21st century: epidemiology, mechanisms, and lifelong effect. Lancet 2016;387:475–490.
33. Dunn N, Juto A, Ryner M, et al. Rituximab in multiple sclerosis: frequency and clinical relevance of anti-drug antibodies. Mult Scler 2018;24:1224–1233.
34. Emery P, Fleischmann R, Filipowicz-Sonnowska A, et al. The efficacy and safety of rituximab in patients with active rheumatoid arthritis despite methotrexate treatment: results of a phase IIIb randomized, double-blind, placebo-controlled, dose-ranging trial. Arthritis Rheum 2006;54:1390–1400.
35. Salter J, Eversen-Bornbak J, Alping P, et al. Rituximab in multiple sclerosis: a retrospective observational study on safety and efficacy. Neurology 2016;87:2074–2081.
36. Esteve-Sole A, Luo Y, Vlaga A, et al. Regulatory cells: players in pregnancy and early life. Int J Mol Sci 2018;19:E2099.
37. Martin JA, Hamilton BE, Osterman MJ, Driscoll AK, Drake P. Births: final data for 2018. Table I-19. Preterm births by race and Hispanic origin of mother: United States, each state and territory, 2018. Natl Vital Stat Rep 2019;68. Available at: www.cdc.gov/nchs/data/nvss/nvss8/nvss8_13_tables-508.pdf. Hyattsville, MD: National Center for Health Statistics.
38. Centers for Disease Control and Prevention. What is Stillbirth? [online]. Available at: www.cdc.gov/reproductivehealth/maternalinfanthealth/pretermbirth.htm. Accessed November 26, 2019.
39. Centers for Disease Control and Prevention. Data & Statistics on Birth Defects [online]. Available at: www.cdc.gov/ncbddd/birthdefects/data.html. Accessed December 12, 2019.
40. Fink K, Gorczyca A, Smith JB, Puehl F, Langer-Gould A. Rituximab, multiple sclerosis and pregnancy. Poster presented at the ECTRIMS 2019 Congress; September 11–13; Stockholm, Sweden. ECTRIMS Online Library 2019.
41. Wray S, Kappos L, Vukusic S, et al. An Update on Pregnancy Outcomes Following Ocrelizumab Treatment in Patients with Multiple Sclerosis and Other Autoimmune Diseases. Poster presented at the 2018 Annual Meeting of the Consortium of Multiple Sclerosis Centers (CMSC); May 30–June 2; Nashville, TN. 2018.
42. Agrawal N, Johnston SC, Wu YW, Sidney S, Fullerton HJ. Imaging data reveal a higher pediatric stroke incidence than prior US estimates. Stroke 2009;40:3415–3421.
43. Agnord A, Bhattacharya S. Infertility and miscarriage: common pathways in manifestation and management. Womens Health (Lond) 2015;11:527–541.
44. Emery P, Rigby W, Tak PP, et al. Safety with ocrelizumab in rheumatoid arthritis: results from the ocrelizumab phase III program. PLoS One 2014;9:e87379.
Rituximab, MS, and pregnancy
Jessica B. Smith, Kerstin Hellwig, Katharina Fink, et al.
Neurol Neuroimmunol Neuroinflamm 2020;7;
DOI 10.1212/NXI.0000000000000734

This information is current as of May 1, 2020

Updated Information & Services
including high resolution figures, can be found at:
http://nn.neurology.org/content/7/4/e734.full.html

References
This article cites 34 articles, 5 of which you can access for free at:
http://nn.neurology.org/content/7/4/e734.full.html##ref-list-1

Citations
This article has been cited by 2 HighWire-hosted articles:
http://nn.neurology.org/content/7/4/e734.full.html##otherarticles

Subspecialty Collections
This article, along with others on similar topics, appears in the following collection(s):
Multiple sclerosis
http://nn.neurology.org/cgi/collection/multiple_sclerosis

Permissions & Licensing
Information about reproducing this article in parts (figures, tables) or in its entirety can be found online at:
http://nn.neurology.org/misc/about.xhtml##permissions

Reprints
Information about ordering reprints can be found online:
http://nn.neurology.org/misc/addir.xhtml##reprintsus

Neurol Neuroimmunol Neuroinflamm is an official journal of the American Academy of Neurology. Published since April 2014, it is an open-access, online-only, continuous publication journal. Copyright Copyright © 2020 The Author(s). Published by Wolters Kluwer Health, Inc. on behalf of the American Academy of Neurology. All rights reserved. Online ISSN: 2332-7812.