A 26-Year Follow-up of a Woman With 5 Consecutive Children Exposed to Risperidone During Pregnancy and Breastfeeding

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There is a large body of literature on gestational and neurodevelopmental outcomes after antipsychotic exposure during pregnancy1–4 but, to our knowledge, no report on outcomes following several successive pregnancies and breastfeeding with exposure to the same antipsychotic and anticholinergic drug. We therefore present the unusual case of a woman with continuous use of risperidone and trihexyphenidyl (THP) from 1995 to the present day, during which period she gave birth to 5 healthy children who have now reached adolescence or adulthood.

Case Report

Ms A, a 23-year-old woman, presented to the psychiatry clinic in 1995 with a 1-year history of hearing voices, smiling and laughing to herself, poor personal care, and other behavioral disturbances. There was no history of alcohol or substance use. She was diagnosed with undifferentiated schizophrenia (DSM IV criteria) and was prescribed risperidone uptitrated to 5 mg/d and THP 2 mg/d. She improved with treatment and conceived 6 months later. The pregnancy was unplanned. Her attitude toward the pregnancy was positive. She and her husband chose to continue treatment during pregnancy because they did not wish to risk illness relapse. She complied with all antenatal guidance.

At full term, Ms A delivered a healthy girl at home; the delivery was uneventful and was conducted by a midwife. Six months later, the dose of risperidone was reduced to 4 mg/d; THP was continued at 2 mg/d. This was the treatment regimen on which she conceived for the second time in 1999. The medications were again continued all through pregnancy. The pregnancy and delivery of a girl (again at home) were uneventful.

Six months later, there was a transient relapse into psychosis; this was successfully managed with the addition of olanzapine 5 mg/d. Olanzapine was withdrawn uneventfully 6 months later. While on risperidone 4 mg/d and THP 2 mg/d, she again conceived in 2001, 2004, and 2008 and delivered male children. All deliveries were conducted at home and, again, there were no clinically notable antenatal, intranatal, or postnatal complications.

All 5 children were breastfed for approximately 20–24 months. As far as could be ascertained during follow-up visits, there were no untoward events during pregnancy, delivery, or afterward in any of the pregnancies. All babies were healthy, fed well, and developed well. There were no evident congenital malformations, delays in development, or emotional, behavioral, or neurologic disorders.

At present, Ms A’s children are 26, 23, 22, 19, and 15 years old. The first 3 children have completed their education with good grades and are gainfully employed. The remaining 2 children continue their studies and are faring well academically.

Ms A continues to take risperidone 4 mg/d and THP 2 mg/d. She has maintained excellent adherence to these 2 drugs over the past 28 years.

Discussion

It is generally considered that, in women with schizophrenia, antipsychotic drugs are best continued during pregnancy because the risks associated with antipsychotic exposure are less than the risks associated with untreated psychosis.5 Our patient is unique in the literature for gestational and breastfeeding exposure to risperidone 4 to 5 mg/d and THP 2 mg/d across 5 consecutive pregnancies, in which all deliveries were conducted at home, assisted only by a midwife.

Although antipsychotic drug exposure during pregnancy has been associated (without certainty of causality) with a range of adverse gestational and neurodevelopmental
outcomes,1–4 there were no adverse outcomes that drew clinical attention in this woman and her offspring. We acknowledge that there is no information on the exact duration of gestation, Apgar score at birth, birth weight, developmental milestones, and other specific details. However, in terms of real-world outcomes, such as need for clinical attention, educational attainment, and occupational functioning, all has apparently been well.

There is no systematic study of THP during pregnancy and lactation, but the anecdotal literature is reassuring.6–8 Of note, risperidone raises serum prolactin and predisposes to menstrual irregularities, including amenorrhea and resultant infertility.9 This was clearly not the case with the patient reported here. Finally, one of us (D.M.) previously reported the uneventful use of risperidone in a woman across 2 consecutive pregnancies.10

References