Administration of Tocolytic Agents is not Only unuseful but also Culprit to Prevent Preterm Labor

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Abstract

In Japan, long term tocolytic therapy for threatened preterm labor is still performed at many perinatal centers despite of less evidence for prevention against preterm delivery. There are some reports which tocolytic agents can prolong preterm delivery during antenatal corticosteroid therapy for maturation of fetal lung, or for maternal transport to a tertiary perinatal care center and which the administration of tocolytic agents to patients with threatened premature delivery was restricted due to side effects such as maternal lung edema or rhabdomyolysis. In addition to these side effects, long term administration for mother with intrauterine exposure for fetus in beta-2 receptor agonist including ritodrine hydrochloride is risk factor of childhood asthma. It was reviewed two reports about tocolytic agents including beta-2 receptor agonist to reveal that there is no evidence performing long term tocolytic therapy.

Keywords: Tocolytic therapy; Ritodrine hydrochloride; Asthma; Side effect

Needless of Long Term Tocolytic Therapy

Short-term tocolysis using ritodrine and/or magnesium sulfate for threatened preterm labor is not common in Japan. Long term tocolytic management strategy to prevent preterm birth may contribute to the epidemiological data that the Japanese preterm delivery rate was less than 6% in 2007, with the lowest perinatal mortality in the world [1]. However, there is few reports which long-term tocolysis was useful to prolong the pregnancy period [2].

Figure 1A: Previous protocol for using tocolytic agents (long term protocol)
Natural preterm delivery due to onset of labor is thought to be caused by some inflammations in uterine cave as a maternal protective reaction. Such a onset of labor which would be occurred term or preterm cannot be controlled using tocolytic agents including ritodrine hydrochloride and magnesium sulfate. Nakamura et al. changed the management protocol for threatened preterm birth in January 2014 from long-term tocolysis to short-term tocolysis (Figure 1). There were 1548 and 1444 deliveries before after 2014 during the study period. There was no significant change Gestational age (week) at delivery, neonatal body weight, and Apgar score (1/5 minutes) between the two groups, respectively. The outcomes of the subjects in both groups are demonstrated in Table 1. No significant differences were observed between the two groups for any of the parameters in the study. Japan currently has the lowest perinatal mortality worldwide [4]. Japanese obstetricians believe that long-term tocolysis has a definite contribution to the low preterm delivery and perinatal mortality rates. Although the rates of singleton preterm delivery in Estonia, Finland, Ireland, Lithuania and Sweden were less than 5% among European countries [5,6].

Table 1: Outcomes of the subjects before and after the change of the management protocol for preterm labor

| Before (1548) | After (1444) | After (1444) | p-value |
|-------------|-------------|-------------|---------|
| Delivery weeks’ gestation (w) | 38±3 | 38±3 | 0.975 |
| Gestational age at diagnosis of preterm labor(W) | 26±5 | 28±4 | 0.221 |
| Admission due to preterm labor | 65 (4.2 %) | 42 (2.9 %) | 0.062 |
| Duration of hospitalization due to preterm labor (days) | 44±37 | 34±30 | 0.117 |
| Preterm birth (<37 weeks) | 182 (11.8 %) | 153 (10.6 %) | 0.324 |
| Preterm birth (<28 weeks) | 20 (1.3 %) | 17 (1.2 %) | 0.869 |
| Preterm premature rupture of membrane | 57 (3.7 %) | 41 (2.8 %) | 0.217 |
| Preterm delivery due to preterm labor | 29 (1.9 %) | 20 (1.4 %) | 0.316 |
| Prolonged gestational days from admission | 54±37 | 48±34 | 0.432 |
| Fetal fibronectin positive | 27.7% (18/65) | 19.4% (8/42) | 0.361 |
| Neutrophil elastase | 2.6±2.8 | 1.9±2.6 | 0.194 |
| Nugent score | 2±1 | 2±1 | 0.887 |
Cervical length of the preterm labor | 28.5±14.1 | 25.4±10.8 | 0.248
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Male (neonates) | 815 (52.6 %) | 726 (50.3 %) | 0.2
Neonatal body weight (g) | 2855±75 | 2868±585 | 0.542
Apgar score | 8 (0-10) | 8 (0-10) | 0.634
1 min | 8 (0-10) | 9 (0-10) | 0.543
5 min | 137 (8.4%) | 129 (8.5%) | 0.488
Admission to NICU | 59±42 | 56±55 | 0.632

Data indicates frequencies, median (range) or mean±standard deviation.

Because there was no significant change of perinatal outcomes before and after 2014, it was no need at least long term tocolytic therapy to extend a delivery period in cases with threatened preterm labor.

**Maternal and Childhood Risks of Long Term Tocolytic Therapy**

It has been reported that the frequencies of side effects of tocolytic agents were lower than those of the FDA recommendation6. Although it was reported that there are some 18.7% of side effects with ritodrine hydrochloride such as elevated liver enzyme, rhabdomyolysis, edema of lung, granulocytopenia, palpitation tachycardia, parotid bubo and hyperamylasemia in Japan7, the frequencies of side effects might be lower than in Western countries due to a lower prescribed dosage. Because long term tocolytic therapy could not improve the outcomes in cases with threatened preterm labor as mentioned previously, tocolytic agents might give a risk of side effect to such cases.

Ogawa K. et al. [8] have reported that beta-2 receptor agonist (B2RA) exposure to fetus in utero performed intravenous administration is a risk factor for childhood asthma until 5 years old. 1158 children were analyzed in this study. Children exposed to B2RA in utero with current asthma at 5 years old were increased to adjusted odds ratio 2.04 (95% confidence interval 1.02-4.05) compared to them non-exposed group. The risk for current asthma at 5 years old was total dose dependent during pregnancy according to the point which B2RA administration with low-dose was not significant compared to non-exposed group. In addition, Ogawa K et al. [9] reported that as dosage of oral administration was much smaller than that of intravenous, oral B2RA administration during pregnancy would not be a risk for current asthma at 5 years old. European Medicine Agency had decided to restrict the period of usage in Ritodrine hydrochloride and to revoke insurance approbation for oral administration of Ritodrine.

**Discussion**

It would be needed to be reconsidered the management using tocolytic agents for threatened preterm labor to the aspects of the side effects, cost and the bed for pregnant women who would need the perinatal management without threatened preterm labor.

**Conclusion**

In conclusion, merit of tocolysis for threatened preterm labor not only was small but also have some problems for maternal, fetal and childhood periods.

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