



Effect of Ursodeoxycholic Acid (USDA) on Indirect Hyperbilirubinemia in Neonates Treated with Phototherapy

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Abstract

Objective: To investigate the effect of USDA in the treatment of indirect hyperbilirubinemia of neonates. Study design: This randomized clinical trial was conducted on neonates with indirect hyperbilirubinemia whom were hospitalized at Ardabil city hospital. 100 newborns were enrolled in the study and randomly divided into two groups. The treatment group (n=50) received 10 mg/kg/day divided q12h USDA in addition to phototherapy, while the control group (n=50) only received phototherapy. Total bilirubin levels were measured every 12 hours until reaching bilirubin below 10mg/dL. Also, duration of phototherapy and hospitalization in both groups were recorded. Data were analyzed in SPSS.21 using statistical methods.

Results: There was no difference in the mean of total bilirubin between treatment and control groups at baseline, 12, 24, 36, and 48 hours after admission.

Conclusions: Our study revealed that adding USDA to phototherapy in treating indirect hyperbilirubinemia of newborns had no preference to just phototherapy treatment.

Keywords: Phototherapy; Indirect Hyperbilirubinemia; Ursodeoxycholic Acid; Neonate

Introduction

Hyperbilirubinemia is the most common cause of hospitalization among healthy and premature infants. Hyperbilirubinemia is prevalent and was seen in 60% of term neonates and 80% of preterm neonates [1]. Increase of indirect bilirubin is harmful and toxic to the brain and may lead to kernicterus which is a lasting and severe damage to the infant's nervous system [2]. Therefore, early treatment of hyperbilirubinemia is important. Typically, phototherapy is widely used as an effective method for the treatment of indirect hyperbilirubinemia in neonates, but has side effects such as fever, dehydration, diarrhea, maculopapular rash, sunburn baby syndrome, higher costs, time consuming and disruptions in the relationship between mother and child because of limiting the baby to incubators and covering his eyes. Studies have shown that in cases with bilirubin more than 20 mg/dL, the power of phototherapy is decreased and reached to 10%-17% [3,4]. So far, drugs such as oral activated charcoal, Metalloporphyrin, D-penicillamine, Phenobarbital and Clofibrate, have been used for the treatment of indirect hyperbilirubinemia [5,6]. Some studies

showed that Phenobarbital has been effective in reducing indirect hyperbilirubinemia and length of phototherapy but has many side effects such as lethargy, reduction of breast-feeding, dehydration and neurological disorders [6,7]. Therefore, doing further studies in relation to medicines with fewer side effects seems necessary. Ursodeoxycholic acid (UDCA) is a bile acid that is widely used in the treatment of Cholestatic liver disorders. UDCA protect the liver against oxidative stress, prevents cell apoptosis and has main role in the stimulation of bile flow, and suppression of treating factors in immunological mechanisms [8]. Several studies have shown that the use of UDCA at a dose of 20-10 mg / kg / day is well tolerated by children and without any special side effects. 9-12 Recently two studies have shown that UDCA, without any special side-effects, considerably reduces the time of reaching bilirubin to normal levels and the infants' hospitalization length [9,10]. The causes of the elevation of indirect hyperbilirubinemia may be physiological jaundice, Crigler-Najjar syndrome, Gilbert's syndrome, breast milk jaundice, blood group incompatibility of Rh and ABO blood groups

and Glucose-6-Phosphate Dehydrogenase (G6PD) deficiency. This study aimed to investigate the effect of UDCA in reducing indirect hyperbilirubinemia of neonates treated with phototherapy.

Materials and Methods

Type of Study and Patients

This randomized clinical trial was conducted on 100 neonates with jaundice, who had received phototherapy in the Ardabil city hospital.

Experiments

All infants divided randomly in two intervention (n=50) and control group (n=50). The intervention group, in addition to phototherapy received UDCA at a dose of 10 mg / kg / day divided every 12 hours (UDCA, capsule 300 mg; provided by DrAdibi Company, Tehran, Iran), with water and was used according to the weight of the infant and the recommended dose and the control group received only phototherapy. On the first day of hospitalization, Total and direct bilirubin, reticulocyte count, Coombs test, G6PD level, complete blood count (CBC), RH, and blood group experiments were performed for both groups. Also, the total bilirubin level of each infant was measured every 12 hours until their bilirubin levels reach below 10 mg / dL and phototherapy stopped. The duration using of phototherapy for neonatal was calculated by the timer.

Statistical Analysis

Collected data entered in a checklist and analyzed using statistical methods and all statistical analysis were performed using SPSS.21. The P<0.05 was considered statistically significant.

Inclusion And Exclusion Criteria

The inclusion criteria for the study were birth weights of 2500 to 4000 gr, breast-feed, gestational age of 38 to 41 weeks,

Table 1: The mean of total bilirubin (mg/dL) in two groups by the time periods.

Groups	Test 1	Test 2	Test 3	Test 4	p-value
Intervention	66/1±75/17	15/2±34/15	85/1±58/12	58/1±65/10	001/0
Control	06/2±34/17	15/2±87/14	80/1±61/12	84/1±50/10	001/0

Table 2: Comparison of the mean of total bilirubin (mg/dL) at different times in two groups.

Groups	Intervention	Control	p-value
At the time of hospitalization	2.25±16.7	2.25±16.7	0.057
12 h after hospitalization	14.19±2.44	14.19±2.44	0.93
24 h after hospitalization	16.71±2.25	16.71±2.25	0.621
36 h after hospitalization	10.5±1.84	10.5±1.84	0.706
48 h after hospitalization	9.78±1.29	9.78±1.29	0.255

Table 3: Comparison of duration of phototherapy and hospitalization time in two groups.

Groups	Intervention	Control	p-value
Phototherapy time(min)	17.37±52.9	14.75±59.56	0.065
Time of hospitalization (day)	0.77±2.82	0.51±3.06	0.07

Discussion

Honar et al. in 2015 in a study, which investigated the effect of USDA plus phototherapy compared to single phototherapy in the treatment of indirect hyperbilirubinemia of neonates, showed that

age over 3 days old, total bilirubin level of 14 to 20 mg / dL, and direct bilirubin level < 2 mg / dL. Infants with ABO blood groups and RH incompatibility, G6PD deficiency, direct hyperbilirubinemia, septicemia and diseases leading to hyperbilirubinemia (Crigler-Najjarsyndrome, Gilbert's syndrome, hypothyroidism/hyperthyroidism, liverdiseases, etc.), premature infants and the infants of diabetic mothers were excluded from the study.

Ethical Approve

Written informed consent was taken from infant's parents and this study was approved in ethics committee of university.

Results

Of all infants, 57% were female and 43% were male. The mean birth weight of infants in treatment and control groups were 3079.8±371.33 and 3231±367.3 kg, respectively. The mean age of infants at the time of hospitalization in the treatment and control groups were 5.64 ± 2.4 and 6.3 ± 2.6 days, respectively. The mean duration of hospitalization in the treatment and control groups was 2/82 and 3/06 days, respectively. The mean duration of phototherapy was 53 and 59 hours in the treatment and control group. There was no significant difference in the average of total bilirubin of infants between treatment and control groups in different time periods (Table 1). There was no significant difference between the two study groups in terms of the mean total bilirubin of infants at the baseline, 12, 24, 36 and 48 hours after admission (Table 2). There was no significant difference in the mean duration of phototherapy and hospitalization between two groups (Table 3). There was no significant difference between boys and girls in term of the mean total bilirubin at baseline, 12, 24, 36 and 48 hours after admission. The average length of hospitalization and the duration of phototherapy in both groups was no significant in boys and girls.

the speed of the reduction of bilirubin concentration in the group taking USDA was considerably high and the mean of required time for bilirubin concentration reduction below 10mg/dl and, as a result, the mean of hospitalization in the experimental group was

considerably less than the control group [10]. In this study, no complications were observed in infants caused by the use of USDA. In this regard, Hassan et al [9]. In a similar study in 2015 examined the effect of USDA in reducing indirect hyperbilirubinemia in infants. The findings of their study revealed that using USDA along with phototherapy is more effective in bilirubin level reduction and quick treatment of infants compared to only phototherapy. In this study, also, no complications were observed in infants using USDA. Our study showed that the mean of bilirubin total in any of 12, 24, 36, and 48 hours after hospitalization between two groups did not differ. It seems that use of USDA with phototherapy does not lead to a more rapid decline in the average of total bilirubin compared to control group. So far, in the world, few studies have been conducted on the effect of USDA on indirect hyperbilirubinemia. Unlike the results of our study, the study Honar et al. and the study Hassan et al. showed that USDA caused a significant reduction in total bilirubin in neonates with indirect hyperbilirubinemia in the first 24 and 48 hours of hospitalization, respectively [9,10]. The findings of this study showed that the mean duration of phototherapy and length of hospitalization in the experimental group was lower than control group but not significant. Honar et al. and Hassan et al., in two studies showed that there was no significant difference in the reduction of total bilirubin, phototherapy and hospitalization time between male and female infants which in line with our study results. 9-10 The findings of this study did not reveal any complication resulting from the use of USDA among infants. This finding was somewhat expected because most previous studies have shown that the consumption of USDA at a dose of 10-20 mg / kg / day are well tolerated by children without any special side effects [9-12]. Although the study of Kotb stated that using USDA (at a dose of 20 mg / kg / day) caused unwanted side effects such as diarrhea outbreaks among consumers. However, since USDA use in our study and other studies was short-term, approving and establishing a definite safety of this drug in infants require long-term follow-up [13].

Conclusion

Our study revealed that adding USDA to phototherapy in treating indirect hyperbilirubinemia of newborns had no preference to single phototherapy treatment and it doesn't have a significant impact in reducing the duration of infants' phototherapy and hospitalization time. For future, doing multicenter studies, with a larger sample size and long-term follow-up are essential about possible side effects of this drug.

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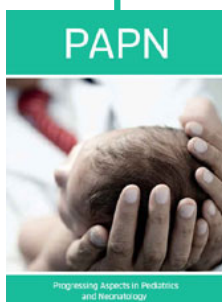
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