

## Comparison between Induction of Labor and Expectant Management in Post-Date Pregnancy

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### Abstract

**Background:** Post-date pregnancy is associated with increased perinatal morbidity and mortality. Therefore post-date pregnancy is considered as a high-risk condition which requires specialist surveillance and induction of labor at some stage. The aim of this work is to evaluate if a policy of induction of labor at 41 GW is superior, in terms of neonatal and maternal outcomes, as compared to expectant management in healthy women with a low risk singleton pregnancy.

**Aim of Study:** The aim with this study is to evaluate if a policy of induction of labor at 41 GW is superior, in terms of neonatal and maternal outcomes, as compared to expectant management in healthy women with a low risk singleton pregnancy.

**Patients and Methods:** A prospective case control study A total number of 100 pregnant women will be included in the study divided into 2 groups: Group (1): Consists of 50 pregnant women who undergo induction of labor at 41 + 0 or 41 + 1 weeks. Group (2): Consists of 50 women who undergo expectant management await spontaneous onset of labor until 42 weeks.

**Results:** This study shows that Meconium Aspiration Syndrome (MAS) was significantly associated with expectant group as 14% of this group had it in their children while only 2% in the induction group, also shows that there are no significant difference between the two groups regarding other perinatal outcomes also shows that rate of CS was significantly associated with induction group 34% while the rate of CS in expectant group only 16%, also the rate of using analgesia was significantly associated with induction group 44% while only 22% in expectant group.

**Conclusion:** Labor induction at 41 completed weeks should be offered to low risk women. The message from this review is that such a policy is associated with fewer deaths although the absolute risk is small. However, this policy may increase the rate of CS or need of analgesia.

**Key Words:** *Post-date pregnancy – Induction of labor – Post-term pregnancy.*

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### Introduction

**ACCORDING** to World Health Organization (WHO), post-date or post-term pregnancy is defined as pregnancy duration of 294 days or longer i.e. Gestational Week (GW) 42 and 0 days (42 + 0) or more measured from the first day of the last menstrual period.

Post-date pregnancy is associated with increased perinatal morbidity and mortality [1]. Therefore post-date pregnancy is considered as a high-risk condition which requires specialist surveillance and induction of labor at some stage.

The etiology of post-date birth is largely unknown. Some rare, known causes of post-date birth are fetal anencephaly, fetal adrenal hypoplasia or insufficiency and placental sulphatase deficiency. Risk factors for post term birth include: Primiparity, advanced maternal age, maternal obesity, heredity, previous post term pregnancy, and a male fetus [2].

Perinatal Mortality (PNM) is defined as the prevalence of stillbirth (after GW 28 + 0) and neonatal mortality within 7 days after birth [3]. PNM increased in women with post-date pregnancies as compared to women with term pregnancies [4].

The risk of perinatal complications such as Meconium Aspiration Syndrome (MAS), umbilical cord complications, asphyxia, pneumonia, sepsis, convulsions, shoulder dystocia, traumatic injuries and peripheral nerve damage is higher in post-date deliveries than in deliveries at term [4]. Also a higher risk of neonatal encephalopathy in children born post-date [5].

Maternal complications increase from GW 40. The risk of puerperal infections, postpartum bleeding, disproportion, labor dystocia, emergency caesarean sections, and cervical lacerations was higher for post-date than for term pregnancies [4].

### Patients and Methods

The study is performed at Ahmed Maher Teaching Hospital during the period between September 2016 and October 2017. A total number of 100 pregnant women will be included in the study divided into 2 groups:

- *Group (1):* Consists of 50 pregnant women who undergo induction of labor at 41 + 0 or 41 + 1 weeks women with a cervix that is judged to be 'ripe' at vaginal examination (Bishop score of 6 or more), will have labor induced with amniotomy followed by intravenous oxytocin. In case of unripe cervix, cervical ripening will be accomplished by vaginal dinoprostone.
- *Group (2):* Consists of 50 women who undergo expectant management await spontaneous onset of labor until 42 weeks. Monitoring can consist of consultations, electronic fetal heart rate monitoring and ultrasound assessment of amniotic fluid.

#### Outcome measures:

Primary outcome will be a composite of perinatal mortality and neonatal morbidity (meconium aspiration syndrome, birth trauma, and perinatal asphyxia and/or NICU admission).

Secondary outcomes will be maternal outcomes such as operative delivery (operative vaginal delivery, caesarean section), need for analgesia (epidural, remifentanyl, pethidin), postpartum hemorrhage and severe perineal injury (third-or fourth-degree perineal tear).

#### Inclusion criteria:

Obstetrical low risk women  $\geq 18$  years with singleton pregnancy in stable cephalic position. Gestational age of 40 + 5-41 + 0 without contraindications for expectant management until 42 weeks.

#### Exclusion criteria:

Age  $< 18$  years, uncertain gestational age, high risk pregnancy (e.g. hypertension, proteinuria ( $\geq 3$  g/L), pre-existent maternal heart or kidney diseases, gestational diabetes, previous caesarean section, multiple pregnancy, intra-uterine growth retardation and non-reassuring fetal status (no fetal movements, abnormal fetal heart rate, known fetal abnormalities

which could influence perinatal outcome, including abnormal karyotype, ruptured membranes at time of randomization and a non-reassuring fetal status at time of randomization).

### Results

Table (1): Demographic data of the studied cases.

		No.=100
<i>Age:</i>		
Mean $\pm$ SD		26.34 $\pm$ 6.82
Range		18-42
<35 years		82 (82%)
>35 years		18 (18%)
<i>BMI:</i>		
Mean $\pm$ SD		23.70 $\pm$ 3.32
Range		19-32
<25		61 (61%)
>25		39 (39%)
<i>Parity:</i>		
Primigravida		49 (49%)
Multigravida		51 (51%)
<i>History of past date pregnancy among multigravida:</i>		
No		21 (41.2%)
Yes		30 (58.8%)

Table (2): Maternal demographics comparing spontaneous and induction of labor for women delivering at 41 to 41 + 6 weeks.

	Expectant No.=50	Induction No.=50	Test value	p- value	Sig.
<i>Age:</i>					
Mean $\pm$ SD	27.74 $\pm$ 7.14	24.94 $\pm$ 6.24	2.087*	0.039	S
Range	18-42	18-39			
<35 years	38 (76%)	44 (88%)	2.439*	0.118	NS
>35 years	12 (24%)	6 (12%)			
<i>BMI:</i>					
Mean $\pm$ SD	24.26 $\pm$ 3.31	23.14 $\pm$ 3.27	1.703*	0.092	NS
Range	19-30	19-32			
<25	26 (52%)	35 (70%)	3.405*	0.065	NS
>25	24 (48%)	15 (30%)			
<i>Parity:</i>					
Primi gravida	18 (36%)	31 (62%)	6.763 *	0.009	HS
Multiparous	32 (64%)	19 (38%)			
<i>History of past date pregnancy among multigravida:</i>					
No	14 (43.8%)	7 (36.8%)	0.235*	0.628	NS
Yes	18 (56.2%)	12 (63.2%)			

This table shows that there was no significant difference between groups regard BMI and the history of post-date pregnancy among multiparous women, expectant group was significantly higher than induction group regard age as they were 27.74  $\pm$  7.14 and 24.94 $\pm$ 6.24 respectively and regard parity as multiparous significantly high in expectant group.

Table (3): Neonatal outcomes comparing expectant management and induction of labor for women delivering at 41 to 41+6 weeks.

	Expectant		Induction		Test value*	p-value	Sig.
	No.	%	No.	%			
<b>Perinatal mortality:</b>							
No	49	98	50	100	1.010	0.315	NS
Yes	1	2	0	0			
<b>MAS:</b>							
No	43	86	49	98	4.891	0.027	S
Yes	7	14	1	2			
<b>Birth trauma:</b>							
No	49	98	50	100	1.010	0.315	NS
Yes	1	2	0	0			
<b>Perinatal asphyxia:</b>							
No	49	98	48	96	0.344	0.558	NS
Yes	1	2	2	4			
<b>NICU admission:</b>							
No	43	86	46	92	0.919	0.338	NS
Yes	7	14	4	8			

This table shows that MAS was significantly associated with expectant group as 14% of this group had it in their children while only 2% in the induction group, Also shows that there are no significant difference between the two groups regarding other perinatal outcomes.

Table (4): Maternal outcomes comparing expectant management and induction of labor for women delivering at 41 to 41+6 weeks.

	Expectant		Induction		Test value*	p-value	Sig.
	No.	%	No.	%			
<b>Caesarean section:</b>							
No	42	84	33	66	4.320	0.038	S
Yes	8	16	17	34			
<b>Operative vaginal delivery:</b>							
No	39	7	38	76	0.056	0.812	NS
Yes	11	22	12	24			
<b>Need for analgesia:</b>							
No	39	78	28	56	5.473	0.019	S
Yes	11	22	22	44			
<b>PPH:</b>							
No	46	92	46	92	0.000	1.000	NS
Yes	4	8	4	8			
<b>Severe perineal injury:</b>							
No	49	98	49	98	0.000	1.000	NS
Yes	1	2	1	2			

This table shows that rate of CS was significantly associated with induction group 34% while the rate of CS in expectant group only 16%, also the rate of using analgesia was significantly associated with induction group 44% while only 22% in expectant group.

## Discussion

The main finding in this study is that there is no significant difference in perinatal mortality between induction of labor at 41 weeks' gestation or later as compared to expectant management (test value 1.010, *p*-value 0.315), there was only one perinatal death in this study 2ry to asphyxia in the expectant group. This results is in keeping with results obtained from Mahomed et al., [6] who reported that there is no significant difference between the two groups regarding stillbirths as the percent of stillbirths in the 2 groups were 0.02. However in 2016 a paper published in ELSEVIER Sexual & Reproductive Healthcare journal titled (has perinatal outcome improved after introduction of a guideline in favor of routine induction and increased surveillance prior to 42 weeks of gestation?) Which show that the perinatal mortality rate remained steady in 2009, 2010 and 2011 (0.10%), but was reduced from 60% from 10 cases in 2010 to three cases in 2012. However, this reduction was not statistically significant (*p*=0.10) [7].

In this study, we also found that induction of labor compared with expectant management was associated with a significantly lower risk of meconium aspiration syndrome (test value 4.891, *p*-value 0.027). This results is in keeping with results from Wennerholm et al., who reported that induction of labor was associated with fewer infants with meconium aspiration syndrome compared with expectant management [8].

However, meconium aspiration syndrome is a poor indicator of neonatal stress, and most newborns with meconium aspiration syndrome recover and remain healthy. So There were no significant differences in intensive care unit admissions between induction of labor or expectant management groups (test value 0.919 *p*-value 0.338). These results are in keeping with results from Burgos et al., [9] and Abraham et al., [10] who reported that no significant difference between the two groups regarding admission of the newborn to NICU.

This study shows no significant difference between the two groups regarding perinatal asphyxia (test value 0.344, *p*-value 0.558), APGAR score less than 7 at 5th minute after delivery (test value -1.158#, *p*-value 0.247) or the rate of birth trauma (test value 1.010, *p*-value 0.315) these results are in keeping with results obtained from Gülmezoglu et al., [11] which found that there is no significant difference between the two groups regarding perinatal asphyxia, APGAR score at 5 minute and the

rate of birth trauma in women who complete 41 weeks and 42 weeks.

The rate of cesarean section in this study is significantly higher in the induction group than the expectant group (test value 4.320,  $p$ -value 0.038) these results are similar to results obtained from Thangarajah et al., [12] which found that the rate of the cesarean deliveries was significantly higher in the induction group (33.8% Vs. 21.1%,  $p$ -value 0.001). These results also are in keeping with results from Mahomed et al., [6] which found that the incidence of CS was significantly higher in the induction group, 22.2% versus 12.1% (OR 2.06; 95% CI 1.93-2.2). Results from Abraham et al., [10] are also similar to this study results regarding the higher cesarean delivery rate ( $p < 0.0001$ ) when compared to expectant management. However, results obtained from Burgos et al., [13] which compares expectant management and induction at 42 week with induction of labor at 41 week show that the rates of caesarean sections in the two groups were 14.1% and 11.4%, respectively ( $p = 0.01$ ).

This study shows significant difference between the two groups regarding the need for analgesia (epidural, remifentanyl, pethidin) there were high need for analgesia in the IOL group 44% compared with 22% for the expectant group (test value 5.473,  $p$ -value 0.019). These results are in keeping with results from Mahomed et al., [6] which show significant difference in the epidural use between the IOL and expectant groups (33.5% versus 21.9%), but differ from results from Abraham et al., [10] which show no significant difference between the two groups regarding epidural use ( $p$ -value 0.55).

The other maternal outcomes in this study show no significant difference between the two groups: Operative vaginal delivery (test value 0.056,  $p$ -value 0.812), PPH (test value 0.000,  $p$ -value 1.000) and perineal injury (test value 0.000,  $p$ -value 1.000). These results apart from perineal lacerations are in keeping with results from Thangarajah et al., [12] which shows no significant difference between the two groups regarding PPH and operative vaginal delivery but show significantly higher perineal injury in the IOL group 38.1% compared with 26.4% in the expectant group ( $p$ -value 0.002). Results from Mahomed et al., [6] show no significant difference between the two groups regarding PPH and 3<sup>rd</sup> or 4<sup>th</sup> degree perineal tear which are similar to this study results. Also, results obtained from Burgos et al., [9] show no significant difference between the two groups regarding instrumental deliveries ( $p$ -value 0.69). Sanne et al., [7] found

that there was no significant difference between the two groups regarding vacuum extraction ( $p$ -value 0.15). The results of this study regarding PPH and operative vaginal delivery also similar to results from Gülmezoglu et al., [11] which show no significant difference between the two groups (assisted vaginal delivery ( $p = 0.65$ ), PPH ( $p = 0.99$ )).

#### Conclusion:

Labor induction at 41 completed weeks should be offered to low risk women. The message from this review is that such a policy is associated with fewer deaths although the absolute risk is small. However, this policy may increase the rate of CS or need of analgesia.

There does not seem to be any increased risk of assisted vaginal delivery, perinatal asphyxia, NICU admission, perineal injury or birth trauma.

If the woman chooses to wait for spontaneous labor onset it would be prudent to have regular fetal monitoring as longitudinal epidemiological studies suggest increased risk of perinatal death by increasing gestational age.

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## مقارنة بين تحريض المخاض وانتظار المخاض الطبيعي في حالات الحمل الممتد لما بعد التاريخ المتوقع للولادة

هذه الدراسة الوصفية المستقبلية على ١٠٠ سيدة حامل تقوم بالمتابعة في العيادات الخارجية وإستقبال الطوارئ بمستشفى أحمد ماهر التعليمي، أجريت الدراسة لتقييم ما إذا كان تحريض المخاض عند إتمام ٤١ إسبوع حمل ينتج عنه أن تكون حالة المولود في الفترة المحيطة بالولادة وحالة الأم أفضل مما ستكون عليه حال إنتظار المخاض حتى إتمام ٤٢ إسبوع.

شملت الدراسة مجموعتين: المجموعة الأولى وعددهن (٥٠) تضم السيدات اللاتي إنتظرن الولادة التلقائية حتى إتمام ٤٢ إسبوع، والمجموعة الثانية وعددهن (٥٠) تضم السيدات اللاتي أجرى لهن تحريض مخاض عند إتمام ٤١ إسبوع إلى ٤١ إسبوع و٦ أيام.

بعد الحصول على الموافقة المستنيرة من الحالات تم أخذ التاريخ المرضي للحالات وإجراء الفحوصات اللازمة لكل حالة.

تم تحديد العمر الحمل من اليوم الأول لآخر دورة حيض (للساء الحوامل نوات الدورة المنتظمة ولا يستعملن موانع الحمل لأكثر من ثلاثة أشهر قبل الولادة) وتم تأكيده بالفحص بالموجات فوق الصوتية إذا كان موجوداً مع الحالة.

تم تقييم عنق الرحم عن طريق معادلة بيشوب المعدلة (على أساس إتساع عنق الرحم، وإمحاء عنق الرحم، والإتساق، والوضع والدموج).

تم عمل موجات فوق صوتية حالية (أثناء هذه الدراسة) للتحقق من وجود كمية سائل أمينوسى مناسب. تم عمل رسم نبض جنين للتقييم والإطمئنان على حالة الجنين.

في المجموعة التي خضعت للتدخل، تم إختيار ٥٠ سيدة عمر حملهن ٤١ إسبوع أو أكثر عشوائياً وتمت إحالتهم إلى المستشفى لإجراء تحريض مخاض.

تم إجراء تحريض المخاض للمجموعة التي خضعت للتدخل بإستخدام قرص دينوبروستون مهبلى ٣مجم (يكرر بعد ٦-٨ ساعات إذا لم تستجب الحالة لأول جرعة والتي لا تتجاوز ٦مجم) تبعه أوكسيتوسين بالتنقيط الوريدي.

في المجموعة المرجعية، ٥٠ سيدة اللاتي تم تخصيصهن للولادة في الموعد المتوقع إنتظرن بدء الولادة التلقائية حتى تمام ٤٢ إسبوع وتمت متابعتهم بإجراء فحوصات الموجات فوق الصوتية ورسم نبض جنين مرتين إسبوعياً. في المجموعة المرجعية، إذا لم تلد المرأة بنهاية ٤٢ إسبوع حمل تخضع لولادة قيصرية. تم جمع بيانات مراحل الحمل الأولى والثانية والثالثة من خلال مخطط المخاض، وتم تسجيل معدل وفيات الأمهات والولدان في الفترة المحيطة بالولادة ومعدل الأمراض.

أجريت الدراسة لتقييم ما إذا كان تحريض المخاض عند إتمام ٤١ إسبوع حمل ينتج عنه أن تكون الفترة المحيطة بالولادة أفضل مما ستكون عليه حال إنتظار المخاض حتى إتمام ٤٢ إسبوع.

أظهرت هذه الدراسة أنه لا يوجد فارق ذو أهمية بين المجموعتين فيما يتعلق بالوزن وتاريخ إمتداد الحمل لما بعد التاريخ المتوقع للولادة بالنسبة للسيدات اللاتي سبق لهن الولادة، والمجموعة التي إنتظرت المخاض سجلت معدل أعلى بشكل ملحوظ من المجموعة التي خضعت لتحريض مخاض فيما يتعلق بالعمر وعدد مرات الولادة حيث أن السيدات متعديدات الولادة أكثر بكثير في المجموعة التي إنتظرت المخاض.

كما أظهرت هذه الدراسة أنه هناك فرق كبير لصالح المجموعة التي إنتظرت المخاض فيما يتعلق بعمر الحمل، ولم يوجد فرق جوهري بين المجموعتين فيما يتعلق بمعامل أبغار. أظهرت هذه الدراسة أيضاً أن متلازمة شفت العقى مرتبطة بشكل كبير بالمجموعة التي إنتظرت المخاض حيث أن ١٤٪ من هذه المجموعة أصيب أطفالهن بهذه المتلازمة مقارنة ب ٢٪ في المجموعة التي خضعت لتحريض مخاض، وأظهرت الدراسة أنه لا يوجد فرق ذو أهمية بين المجموعتين فيما يتعلق بالنتائج الأخرى للفترة المحيطة بالولادة فيما يخص حالة المواليد.

وأظهرت الدراسة أن معدل العمليات القيصرية أعلى بكثير في المجموعة التي خضعت لتحريض مخاض بنسبة ٣٤٪ مقارنة ب ١٦٪ في المجموعة التي إنتظرت المخاض، وأن معدل إستعمال التسكين كان أعلى بكثير في المجموعة التي خضعت لتحريض مخاض.

وكشفت الدراسة أنه ينبغي أن يجرى تحريض المخاض عند إتمام ٤١ إسبوع أو أكثر للحالات نوات معدل خطورة منخفض. والغرض من هذا الإستعراض هو أن هذه السياسة يعزى إليها حالات وفاة أقل مع أن معدل الإختطار المطلق منخفض، ومع ذلك فإن هذه السياسة قد تزيد معدل العمليات القيصرية أو الحاجة إلى التسكين.