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Research Article

Pain relief in labour: tramadol versus pentazocine

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ABSTRACT

Background: The present study was undertaken to compare the effect of 100 mg intramuscular tramadol to 30 mg intramuscular Pentazocine for labour analgesia.

Methods: A total of 60 cases with 37-40 weeks pregnancy in labour, without any foetal or maternal complications were selected. Out of them Inj. Tramadol was given to 30 cases while rest of the 30 patients received injection Pentazocine.

Results: In Tramadol group pain relief was observed in 80% cases, effect started as early as 7-8 min and continued for 2.13 hrs. While in Pentazocine group pain relief was observed in only 60% cases with delayed onset (15-16 min), effect lasted for 2.67 hrs. Maternal and foetal complications were slightly more in Pentazocine group.

Conclusions: Tramadol is an effective and safe labour analgesic, producing moderate to satisfactory. Besides it also significantly shortens the duration of labour.

Keywords: Tramadol, Pentazocine, Labour analgesia, Pain

INTRODUCTION

Motherhood though rewarding seems to require sacrifice on part of the women, i.e. to bear the extremely painful process of child birth. This pain if not adequately controlled can lead to maternal and fetal sequelae because of widespread maternal sympathetic activation that causes increase in cardiac output, Blood Pressure and pulse rate of the mother. Effective analgesia prevents the pain included hyperventilation and hypocapnia which can be severe enough to produce tetany in painful labour. Painful labour also reduces uteroplacental blood flow by upto 25%. The requirements of a satisfactory analgesic in labour are safety and effective analgesia throughout the painful periods of labour with no unpleasant maternal side effects and no depressant effects on the baby or on the maternal cardiorespiratory system.⁸ To overcome this problem a number of pharmacological agents have been tried.

Tramadol, a certainly acting drug with low affinity for opioid receptors and pentazocine a weak antagonist and potent agonist for opioid receptors are currently being studied in our department for relief of labour pain. Progress of labour, foetal outcome and side effects were compared in both the groups.

METHODS

Present study was conducted in the department of Obstetrics and Gynecology, Dhiraj Hospital, S.B.K.S. MIRC, Piparia, Vadodara, Gujarat.

This series included 60 cases out of which 30 cases were treated with tramadol and in remaining 30 cases injection pentazocine was given.

Both primi and multigravida with 37 to 42 weeks pregnancy with vertex presentation in established labour,

i.e., effective uterine contractions, good cervical effacement and cervical dilatation not less than 3cm. were included in the study.

Patients with previous uterine scar, malpresentation, multiple presentations, absent membranes, antepartum haemorrhage, cephalopelvic disproportion, pre eclamptic toxemia and other medical and psychiatric diseases were excluded.

In well selected cases, a general and systemic examination was undertaken. Obstetric examinations, inclusive of abdominal and per vaginal examination were performed and informed consent was taken. All cases were done in consultation with anaesthetist and paediatrician.

In tramadol group, Inj. Tramadol 100 mg was given by deep intramuscular injection in upper and outer quadrant of gluteal region with 2 ml syringe. While in pentazocine group, Inj. Pentazocine 30 mg was given by deep intramuscular injection in the gluteal region.

Active labour was defined as dilatation of cervix > 3cm with one uterine contraction every 3 min. each lasting 30 seconds or more. Repeat dose of 50 mg tramadol or 15 mg pentazocine was injected intramuscular after 4 hours except in case where the women had reached the end of 1st stage or entered 2nd stage of labour. The following points were noted.

Time taken for onset of analgesic action was noted by verbal rating score. All resuscitative measures for mother and baby were kept ready. Vital parameter, foetal heart rate and progress of labour were checked repeatedly. Mode of delivery was noted, Apgar score of every infant was noted at 1 and 5 min. Any complication or emergency reactions in both groups were noted.

Observation

Both the groups were comparable in age, parity, socioeconomic status and locality of patients.

Among both groups 50% patients were primigravida while 50% were multigravida. Mean age of patients in tramadol group was 25 years, while in pentazocine group it was 24.63 years. Maximum patients were of low socioeconomic status and from rural locality.

Mean gestational age was 38.13 weeks in Tramadol and 38.00 weeks in pentazocine group.

By giving I/M tramadol 80% cases responded, response was excellent in 23.33% cases and average in 56.67% cases while in pentazocine group pain relief was observed in only 60% of cases with excellent pain relief in only 6.67% cases (Table 1).

Mean onset of analgesia was early in Tramadol group (7.40 min) in comparison to pentazocine group (15.60 min). However mean duration of analgesia was slightly more with pentazocine group (2.67 hrs.) in comparison to tramadol (2.13 hrs.) (Table 2).

On correlating degree of pain with different physical factors, it was seen that in both the groups pain relief was maximum with 20-25 yrs. Age group, multigravida, stable personality, low socioeconomic status and rural locality.

Tramadol analgesia causes no appreciable changes in maternal pulse rate and blood pressure while with pentazocine mean pulse rate was significantly increased 1 hr. after giving injection (9.80/min) both systolic and diastolic blood pressure were increased and changes in systolic blood pressure were significant (Table 3).

Mean duration of 1st stage of labour was more in pentazocine group (9.13 hrs) in comparison to tramadol group (8.90 hrs). Similarly mean duration of IInd stage was more in pentazocine group, while mean duration of IIIrd stage was less in pentazocine group (8.20 min) in comparison to tramadol group (9.03 min) (Table 4).

Table 1: Degree of pain relief in tramadol and pentazocine groups.

Degree of pain relief	Tramadol group		Pentazocine group	
	No.	%	No.	%
Poor (score I)	06	20	12	40
Average (score II)	17	56.67	16	53.33
Excellent (score III)	07	23.33	02	6.67
Mean		2.03		1.67
S.D.		0.67		0.60
T value	2.211		p<0.05	

Table 2: Onset and duration of analgesia.

Analgesia		Tramadol group	Pentazocine group	t	p
Onset (min)	Mean	7.40	15.60	7.918	<0.01
	SD	1.77	4.66		
Duration (hrs)	Mean	2.13	2.67	3.431	<0.05
	SD	0.58	0.38		

Table 3: Changes in maternal pulse rate, systolic and diastolic blood pressure.

Changes 1 hr after injection	Tramadol group Mean	Tramadol group SD	Pentazocine group Mean	Pentazocine group SD	t	p
Pulse rate /min	0.47	3.04	9.80	2.75	12.466	<0.01
Systolic B.P./min	1.30	4.52	9.53	4.49	5.019	<0.01
Diastolic B.P./min	0.53	4.56	5.37	5.03	3.905	<0.05

Table 4: Duration of stages of labour in tramadol and pentazocine groups.

Groups	Stage I (hrs)		Stage II (hrs)		Stage III (min)	
	Mean	SD	Mean	SD	Mean	SD
Tramadol group	8.90	3.71	0.68	0.52	9.03	4.86
Pentazocine group	9.13	3.41	0.73	0.69	7.20	3.28
t		0.250	0.317	0.317	1.710	
p	>0.05		>0.05		>0.05	

Table 5: Mode of delivery in different groups.

Mode of delivery	Tramadol group		Pentazocine group	
	No.	%	No.	%
Normal	26	86.66	24	80.00
Instrumental	02	6.67	02	6.67
Caesarean section	02	6.67	04	13.33

Table 6: Mean apgar score of neonates.

Time	Tramadol group		Pentazocine group	
	Mean	SD	Mean	SD
At 1 min.	7.63	1.67	7.73	1.26
At 5 min.	9.60	0.60	9.33	0.91

Table 7: Side effects in tramadol and pentazocine group.

Side effects	Tramadol group		Pentazocine group	
	No.	%	No.	%
Nausea	2	6.67	2	6.67
Vomiting	1	3.33	2	6.67
Drowsiness	1	3.33	4	13.67
Tachycardia	2	6.67	7	23.33
Increase in B.P.	2	6.67	10	33.33
Post partum Haemorrhage	2	6.67	2	6.67
Foetal distress	1	3.33	3	10

Normal delivery occurred in 86.66% cases of tramadol group in comparison to 80 % cases of pentazocine group. Operative interference was more in pentazocine group (20% cases) while in tramadol group it was needed in only 13.34% cases (Table 5).

Mean apgar score was more in tramadol group in comparison to pentazocine group. It was 7.93 and 7.73 at 1min. and 9.60 and 9.33 at 5 min. in tramadol group and pentazocine groups respectively (Table 6).

Side effect were more in pentazocine group in form of nausea, vomiting (13.34%), tachycardia (23.33%) increase in blood pressure (38.33%) and foetal distress (10% cases) (Table 7).

DISCUSSION

Prasertsawat et al (1986) observed that 100mg tramadol gives satisfactory to good effect in 78% patients.⁹ However, Suvonnakote et al (1986) reported that 59% patients did not obtain adequate analgesia with Tramadol.¹¹

Nawani et al (1996) observed excellent to average pain relief with tramadol in 80% cases while Sarkar and Mukhopadhyay (1997) observed satisfactory pain relief in 13% and average pain relief in 85% cases.^{8,10}

Mean onset of analgesia was 7.40 min. and effect lasted for 2.13 hrs. in present study. While Husslein et al (1987) concluded that distinct analgesic effect was observed after 10 min. and lasted for 2 hrs.⁴

Duration of different stages of labour was not increased after use of tramadol while in pentazocine group 1st stage of labour was significantly prolonged. Bitsch et al (1980) also observed that tramadol has no effect on duration and intensity of uterine contractions and on duration of different stages of labour.³

Maternal and foetal side effects were very few in tramadol group in comparison to pentazocine group.

In present study, we have found that tramadol is superior in all aspects, onset was earlier, degree of pain relief was

better and there was no danger of maternal and foetal neonatal respiratory depression. Operative interference and complications were more in pentazocine group. The only advantage of pentazocine was that mean duration of analgesia was more with pentazocine (2.67 hrs) in comparison to tramadol (2.13 hrs) although it was not significant.

All these findings confirm the superiority of tramadol over pentazocine for labour analgesia.

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Ethical approval: The study was approved by the Institutional Ethics Committee

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