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Partograph versus no partograph: effect on labour progress and delivery

outcome: a comparative study

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Abstract:

Background:Worldwide, more than a million women between the ages of 15 and 49 years die each year from complications of pregnancy and childbirth. The use of partograph(or labour chart) to monitor the progress of labour is one of the globally recognized tools for reducing maternal mortality.

Aim:To evaluate the effect of use of Partograph on progress of labour and on delivery outcome.

Methodology: Prospective randomized comparative study.Thestudywas conducted in the Department of Obstetrics and Gynecology, Benghazi Medical Center,From1stofAprilto30thofJune2020.Pregnantwomenwererandomlyassigned to two groups, of 200 each,after satisfying the inclusion and exclusion criteria.

Results: Use of partograph significantly reduced the total duration of labour (active phase duration and second stage duration) p-value =0.0001.Augmentation of labour by Oxytocin and ARM were slightly reduced in group one (p value = 0.059). Inthe presentstudynormal vaginal deliverywas recorded in96% offirst group and 93% of second group, C/S done to 4% of 1st group and 7% of 2nd group , this difference was not statistically significant p value was 0.273.Neonatal out comes were better in partograph group . Mean fetal heart record was more in 1st group (pvalue = 0.0001).

RecommendationThe use of partograph is actually required to get a healthy mother and a healthy baby ,and in early identification of slow progress in labour ,so use of partograph should be included as an essential prerequisite while conducting deliveries in all labour wards.

Key words :partograph, progress of labour, Apgar score.

Introduction:

Worldwide, more than a million women between the ages of 15 and 49 years die each year from complications of pregnancy and child birth. About 500,000 women die annually with a huge number left with injury as a result of pregnancy related causes (World Health Organization; 2007).' For each maternal death more women suffer serious complications.

Unfortunately, developing countries disproportionately bear this burden despite global attention and efforts. Poor outcome during labour accounts for about 19% of maternal deaths in these countries. Maternal mortality remains between 500-1000 deaths per 100,000 live births in developing countries (Opiah MM, 2012).

Although the maternal mortality ratio (MMR) has dropped by approximately 45% in the last two decades, around 300,000 women continue to die each year globally due toavoidable pregnancy related complications (World Health Organization; 2014). Obstructed labour is a leading cause of maternal and neonatal mortality, especially in developing countries(Harrison MS, Asibong U, Mathai M.)

Globally, it is estimated that obstructed labour occurs in 5% of pregnancies and accounts for an estimated 8% of maternal deaths (Mathai M., Kayiga H,Kabakyenga JK^{).}

Obstructed labour may result in serious complications such as obstetric fistula, uterine rupture, puerperal sepsis and postpartum haemorrhage(Mukasa PK et all ,Kushwah B et all)

Evidencepointstoa44% declineintheglobalmaternalmortalityrate, from 385 to 216 deathsper 100000 live births betwe en 1990 and 2015 (WHO, 2015).¹² This significant achievement is attributed inpart to the use of the Partogram to improve labourmanagement. The Safe Motherhood Initiative concluded that use of the Partogram reduced maternal and foetal morbidity and mortality, especially in under-resourced countries (Mathibe-Neke JM, 2009, Health Tech USAID, Yisma E 2014).

The partogram is a graphical representation of the events in labour. As part of the Safe Motherhood Initiative launched in 1987, the World Health Organization (WHO) hasproduced, promoted, modified thepartogram and recommends itsuse in labour in order to improve labour management and reduce maternal and fetal morbidity and mortality. Partogram use in the active management of labour has been reported to reduce the occurrence of prolonged labour and need forcaesarean section (Okusanya BO et al, 2018).

Thepartographisapre-printedpaperwithavisual/graphicalrepresentationobservationsmadeonawomanandfoetusduringthecourseoflabour.Theobservationsarecomprisedoftheprogressoflabour,maternalvitalsignsandfoetalheartcondition.Theseobservationsandquickreviewofon-goinglabourandtimingofmanagementdecisions.The

partographisusedasatoolforriskassessmentandiseffectiveindetecting abnormal labour during the first stage of labour. When used correctly, the partographhelpstoidentifyproblemsandinterventionscanbetimelyinitiated thereby preventing morbidity and mortality(WHO,jhpiego,

A World Health Organization (WHO) study in South East Asia involving 35,484women foundthatusingapartograph contributed toreduced (a)prolonged labourfrom6.4% to3.4%,(b)needforaugmentationoflabourwithoxytocinfrom20.7% to9.1%,(c)occurrenceofcaesa rean sectionsfrom9.9% to8.3%, and(d) intrapartum stillbirths from 0.5% to 0.3%. Based on these findings, in 1994, the WHO declared universal use of the partograph in all settings in monitoring labour to help identify abnormal progress and provide timely intervention when required(Mandiwa et al 2017).

History of the partograph The partograph use dates back to the 1950s. It was developed by Friedman, an obstetrician, who had used it to monitor cervical dilation and called it the cervicograph (Friedman EA,1955).

In 1972; Philpott further developed the cervicograph into the partograph which became a practical tool for recording all intrapartum observations in addition to cervical dilation. In Philpott's partograph, he designed alert and action lines which helped to capture prolonged labour. (Philpott RH; Castle WM, 1972)In 1988, Safe Motherhood Initiative launched the use of partograph as an international standard practical tool to monitor labour and prevent prolonged labour. In 1994, WHO extensively tested its efficacy and established its scientific basisandrationale foritsuseinprevention ofprolonged labor(Lancet;1994). Itsusereduces the incidence of prolonged/ obstructed labour and can also detect foetal heart abnormalities which can result in intrapartum foetal hypoxia. In 1994 WHO declared universal application of the partographin all settings.

 ${\bf Aim:} To evaluate the effect of use of Partograph on progress of labour and on delivery outcome.$

Methodology:

Typeofthestudy: Prospectiverandomized comparative study.

Place: Departmentof Obstetrics and and Gynaecology, Benghazi Medical Center.

Durationofthestudy:From1stofAprilto30thbofJune 2020.

Patients: Pregnant women were randomly assigned totwogroups, of200each, aftersatisfying theinclusion and exclusion criteria. Women assigned toGroup1 hadtheiractivelabourusingpartograph whereasthoseassignedtoGroup2were not monitored using the partograph.

Inclusioncriteria:Age:18yearsandabove,Gestationalage:37-42weeks,Singleviablepregnancy,Cephalicpresentation, Cervicaldilatation4cmorbeyond.Figure 100 (200)

Exclusioncriteria: Non-cephalicpresentation, uterinescar and other contraindications for vaginal delivery.

Patients who satisfied the inclusion criteria and had given their consent tobe included into the studywere randomly allotted eitherinto, Group 1: Patients who were tobe monitored in the active phase of labour using Partograph. Group 2: Patients whose active labour was not monitored using Partograph, All the patients were routinely examined and tailed history was taken aspert prepared proforma. Patients fulfilling the inclusion criteria, after being admitted were questioned and thoroughly examined with pre-designed pre-tested proforma. General, systemic and obstetric examinations were done.

Per-abdominalexamination:heightofuterus,presentation,engagementandfetal heart rate were noted.

Per-vaginalexamination:presentation,position,engagement,cervicaldilatation, Effacement, station, status of membranes, color of liquor (if membrane were absent), adequacy of pelvis was done.Wheninactivelabour,thedetailsoflabourandotherrelevantdetailswere recorded on the Partograph in group1.Durationofactivephase,Durationsecondstage,Totaldurationoflabour,Needforaugmentation

(ARMandoxytocin),Modeoftermination and intervention required, Apgar score at birth, NICU admissions.Activephaseofgroup2patientswasmonitoredarbitrarily without recordingtheir findings on Partograph.

Statisticalmethods:

Statistical analysis onstudyresultswasper-formed bytheapplication of the SPSS version 22. For comparison of mean of two groups independent T-test was applied. For categorical variables, chi-square test wasapplied. P<0.05 was determined to be statistically significant. The variables in the two groups were tabulated and compared. Figure weredone by Microsoft office Excel 2010.

Results

Statistical analysis onstudyresultswasper-formed bytheapplication of the SPSS version 22. For comparison of mean of two groups independent T-test was applied. For categorical variables, chi-square test wasapplied. P < 0.05 was determined to be statistically significant. The variables in the two groups were tabulated and compared. Figure weredone by Microsoft office Excel 2010.

	Partogramgroup		NoPartogramgroup	
Durationofactive phase/hours	No.	0 p	No.	00
0.3-1.3	20	10.4	3	1.6
1.31—2.3	45	23.4	19	10.2
2.31-3.3	38	19.8	20	10.8
3.31—4.3	42	21.9	46	24.7
4.31—5.3	32	16.7	34	18.3
5.31—6.3	10	5.2	18	9.7
6.31—7.3	3	1.6	24	12.8
7.31—8.3	2	1	16	8.6
8.31—9.3	0	0	6	3.3
Total	192	100	186	100

Table 1: Duration of active phase in Partogram group and no Partogram group.

Chi-square = 64.385 with 8 degrees of freedom; P = 0.0001(Highly

significant.

Partogram:Mean=3.4hours.Std. Deviation=1.5hours.Median=3.2hours. Minimum=1hour.Maximum=8.30hours.

No Partogram: Mean =5.1 hours.Std.Deviation =2 hours.Median=5 hours. Minimum =1.30 hours. Maximum =9.30 hours.

t= -9.518with376degreesoffreedom;P=0.0001(Highlysignificant).

Table2: Total duration of labour (active phase duration and second stage duration)inPartogramgroupandnoPartogramgroup.

	Partogramgroup		Nopartogramgroup	
Timeinterval/ hours	No.	00	No.	00
1-2.3	43	22.4	10	5.4
2.31–4	59	30.7	22	11.8

4.01—5.30	64	33.3	56	30.1
5.31–7	22	11.5	41	22
7.01—8.3	2	1	36	19.4
8.31-10	2	1	21	11.3
Total	192	100	186	100

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Chi-square =89.756 with 5 degrees of freedom; P = 0.0001 (Highly significant).

Partogram: Mean =3.7.Std.Deviation =1.5.Median=3.5. Minimum =One hour. Maximum =9.10.

Nopartogram:Mean=5.7.Std. Deviation=2. Median=5.5.Minimum=1.50. Maximum =9.50.

t=-11.198with376degreesoffreedom;P=0.000(Highlysignificant)

Table3: Comparisonbetween Partogram group and no Partogram group based on mode of delivery.

	Partogramgroup		Nopartogramgroup	
Modeofdelivery	No.		No.	%
	1.00	%	1.0.	,,,
Normalvaginal	192	96	186	93
delivery				
C/S	8	4	14	7
Total	200	100	200	100

Chi-square=1.203with1degreeoffreedom;P=0.273(Notsignificant)



Chi-square=2.373 with 1 degree of freedom; P=0.123 (Not significant).

Fig.1:Comparison of APGAR score at birth of new-born between Partogram group and no Partogram group.



Partogram:Mean=6.7times.Deviation=2.9timesMedian=6timesMinimum =2times.Maximum=16times.

Nopartogram:Mean=3.1times.Std.Deviation=1.8times Median=3timesMinimum =Once. Maximum = 10times.

t=14.678with376degreesoffreedom;P=0.0001(Highlysignificant).

Fig.2: Comparisonbetween partogram group and no partogram group based on number of records of fetal heart.

D. A	Partogramgroup		Nopartogramgroup	
Presence of meconium	No.	%	No.	%
Yes	12	6	15	7.5
No	188	94	185	92.5
Total	200	100	200	100

 Table 4: Comparisonbetween Partogram group and no Partogram group based on presence of meconium.

Chi-square=0.159with1degreeoffreedom;P=0.690(Notsignificant).

Table 5: Comparisonbetween Partogram group and no Partogram group based on no significant difference

Variable	Group 1 (partograph use)	Group 2 (No partograph use)	P-value
Maternal Age	18-46	18-47	0.081
Gestational age (mean)	39.5	39.6	0.475
Presence of meconium	6%	7%	0.690

Table 6: Comparisonbetween Partogram group and no Partogram group based on significant difference

Variable	Group 1 (partograph use)	Group 2 (No partograph use)	P-value
Duration of active phase	3.4±1.5hours	5.1hours ±2hours	0.0001
Total duration of labour(mean)	3.7	5.7	0.0001
Need for augmentation of labour	30% of patients	39.5% of patients	0.059
Oxytocin use	16.7%	14%	
ARM	33.3%	35.4%	
ARM + Oxytocin	50%	50.6%	0.059

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Normal vaginal delivery	96%	93%	0.273
8 5			
Cesarean section rate	4%	7%	0.273
	170	770	0.275
Apgar<7			0.123
npgai </td <td></td> <td></td> <td>0.125</td>			0.125
Number of recorded fetal heart	67	3.1	0.0001
rumber of recorded retaineart	0.7	5.1	0.0001.
(mean)			
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Discussion:

The study included 200 patients done partogram (1st group) and 200 patientsnot done (2nd group), the maternal age of 1st group was ranged betweenl8years and 46 years, while the second group their ageranged between 18 and 47 years, there was no significant difference between the mean age of both group p=0.081. Gestational age <40 weeks constitute to 49% of 1st group and 45% in 2nd group, gestational age ñ40 weeks was 51% in 1st and 55% in 2nd group, these differences were not statistically significant p value =0.475.

group,thisdifferencewasnotstatisticallysignificantpvalue=0.690.

Duration ofactive phaseinpartogram group was ranged from 1hour to8.30hours, with mean equal to 3.4+1.5 hours, while range in no partogram group was 5.1hours to 9.30hours, with mean equal to 5.1hours +2hours, this difference between the two group was highly significant pvalue was 0.0001. This result was comparable to studies done by Sharma AK et al,2016²⁵ and study by AmedBenazir etal,2016

Duration of second stage in partogram group was ranged from 2.3minute to 60 minutes, with mean equal to 24.5 minute +12.4 minute, and in no partogram the duration of second stage was ranged from 10 minutes to 70minuts, with range equalto42.3minute+18minutes,thisdifferencewashighlysignificantpvalue=0.0001. AmedBenazir etal(2017)²⁴foundthemeanduration of second stageingroup1 was34.78+20.59minutes and ingroup2was 56.46+23.94 minutes, whileSharmaAK et al,² reported the mean duration of 2nd stage a labour in his study using Partograph to be 33.64+23.85minutes.(Sharma AK, 2016)

Total duration of labour (active phase duration and second stage duration) in partogram group was ranged from one hour to 9.10 hours, the mean was 3.7 hours+1.5 hours, while the total duration of labour in no partogram was ranged from 1.50 hours to 9.50 hours , the mean was 5.7 hours +2 hours, this difference was highly statistically significantp value0.0001. This result was comparable to study conducted by Pinky Rachhoya Peta 1²⁶ and study by A med Be nazir etal²⁴

Augmentation was needed by30% of patients in 1^{st} group and 39.5% of 2^{nd} group, ARM was done to 33.3% of 1^{st} group and 35.4% of 2^{nd} group, Oxytocin was given to 16.7% of 1^{st} group and 14% of 2^{nd} group, while ARM + Oxytocin recorded in 50% of 1^{st} group and 50.6% of second group, this difference was slightly significant pvalue=0.059, which was parable tostudybyAmed Benazir et al²⁴were the augmentation was required to 33.5% in group with partogram compared to 44% of no partogram.

In the present study normal vaginal delivery was recorded in 96% of first group and 93% ofsecond

group, C/S done to4% of1st group and 7% of2nd group, this difference wasnotstatistically significant pvaluewas0.273.Inotherstudy there were 91% normal vaginal deliveries in group 1 and 81.5% in group 2.

StudybyDivyaSetalreported83.8% spontaneous vaginaldeliveries inpatients followed by Partograph (cases) and 69.4% normal vaginal deliveries in patients not monitored by Partograph (controls). ²⁷Apgar score was>7in88% in 1stgroup and 82% of 2ndgroup, this difference was not statistically significant p value = 0.123.

Inother study, 6.5% Sharma AK et al,in his study demonstrated 6% of new borns with Apgar score <7 at birth which was comparable to the present study.²⁵

Pinky R et al. demonstrated an Apgar score of <7 at birth in 2.4% of the newborns born to mother monitored using partogram. ²⁸

Admission to NICU was recorded in 11.5% of 1^{st} group and 13% of 2^{nd} group, 34.8% of 1^{st} group had Apgar score ñ7 and 65.2% had Apgar score <7, while in 2^{nd} group 3.8% had Apgar score ñ7 and 96.2% Apgar score <7, this difference between the two group was statistically significant pvalue =0.015.

The present study was comparable to the study conducted by Surekha Tetal who found asignificant reduction in NICU admissions of control group not monitored by Partograph (17%) as compared to the cases $(6\%)^{29}$

Mean fetal heart records in 1^{st} group was 6.7times + 2.9times, wand ranged from 2times to l6times, while in 2^{nd} group themean was 3.1times +1.8times with range one to 10 times, these differences were highly significant p value = 0.0001Inothersstudyamong100ofthemodifiedWHOpartographsreviewed,foetal heartratewasnotrecordedin9(9%)andwassub-standardin13(13%)while monitored up to the

recommendedstandard in 78(78%) of the partographs. ³⁰Presence of meconium was present in 6% of 1^{st} group and 7.5% in 2^{nd} group, this difference was not statistically significant pvalue =0.6

Conclusion:

This study shows that the Partograph helps in reducing the active phase of labour, second stage of labour and hence the total duration of labour. It also effective in reducing the need for augmentation and allows the labour to progress spontaneously without the need of unnecessary interventions.

Proper and correct interpretation of Partograph increases the number of normal vaginal deliveriesbyreducingtheunnecessaryinterventions thatwouldhavebeen taken when Partograph is not used.

The neonatal condition, as assessed by Apgar score after the baby is born,

isalsobetterwhenthelabourismonitored usingPartograph. Admission of babiesto NICU also less in partograph group than no partograph group, so partograph improving the outcome of mothers and babie

Recommendation:

Theuseofpartograph isactually required togetahealthymother and a healthy baby, and in early identification of slow progress in labour, so use of partograph should be included as an essential pre-requisite while conducting deliveries in all labour wards.

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