

The Galaa Study
Normal Labour Facility Practices
VS
Evidence-based Obstetrics

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Justification

- Why study normal labour ?
- Why study hospital practices for normal labour?
- Why study normal labour at El Galaa hospital ?

Background

Maternal mortality in Egypt

- 49 % of births take place in facilities
- 69 % are medically assisted (by trained providers)
- Egypt's maternal mortality rate of 74 deaths per 100,00 live births is relatively high
- Postpartum hemorrhage is the leading direct cause of maternal deaths in Egypt.

(MOHP, 2001)

Justification

Why study normal labour in Egypt?

- The **majority** (85%) of all deliveries worldwide are normal.
- Substandard care **by the obstetric team** was the leading avoidable factor contributing to the maternal deaths identified by the last mortality survey. (43 % of avoidable deaths)
- 49 % of deaths occurred during delivery/within 24 hrs

(MOHP, 2001)

Risk of Maternal Death by Day of Pregnancy, Egypt 2000 (MOHP, 2001)



- Neonatal morbidity and mortality- illness and death- can occur after normal labour and not just after complicated deliveries.
- While normal labour is the most common event, most studies focus on high risk or complicated labour and delivery.

Why study hospital practices for normal labour?

Deliveries in hospitals are increasing.

59% of all deliveries in Egypt occur in facilities
(EIDHS, 2003)

There is **no documentation** of facility practices for normal labour in our region.

Why study normal labour in El-Galaa hospital?

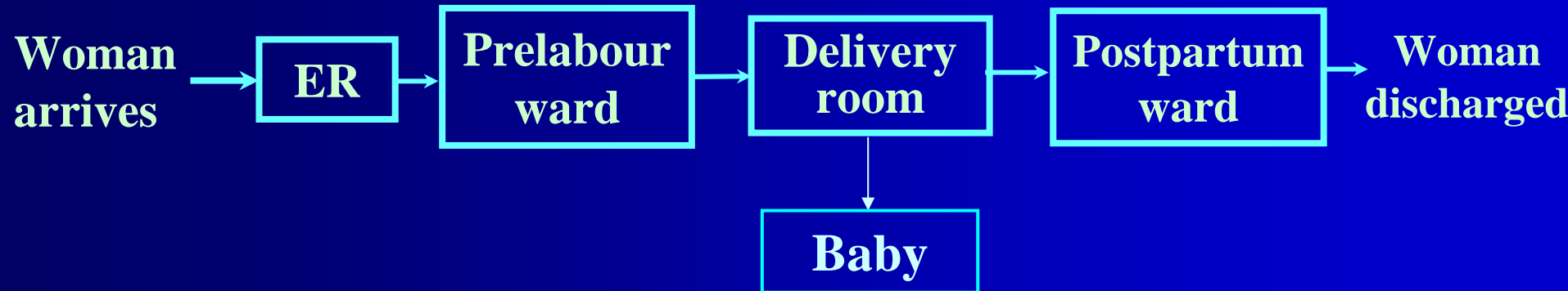
- It delivers 25,000 women per year.
- . It is an important obstetrical training center. (at least 225 young doctors are trained at El Galaa every year).

The Galaa Study

- The research team comprehensively documented practices for normal labour at El Galaa hospital, for the first time, in 2001.
- The labours and delivery of 176 women from admission to discharge were individually observed using a multifaceted observational methodology.
- Data was collected quantitatively and qualitatively by directly observing women for their entire labour and delivery; by documenting ward activities, interviewing the delivered women and sharing the findings with the providers.

Our study documented:

Every practice for which there is evidence for a role or an effect during normal birth.



This involved:

- a) Observation of one woman and neonate at a time using a 165-item observational checklist. 672 continuous hours of observation
- b) Observation of ward activities
- c) A postpartum interview exploring women's perceptions of their experience.
- d) Feedback from providers.
- e) Lessons from the pilot study.

We were not able to identify any other studies that quantified facility procedures comprehensively for normal labour.

Main Findings

18 Main issues were identified:

1. Demographic variables
2. Caseload
3. Facility preparedness
4. Initial assessment
5. Recording
6. Management of the 1st stage
7. Management of the 2nd stage
8. Management of the 3rd stage
9. Immediate postpartum care
10. Neonatal care
11. Postpartum care
12. Discharge procedures
13. Supervision of doctors in training
14. Communication
15. Continuous monitoring
16. Infection control
17. Pain relief
18. Hospital policies from the woman's perspective

Main General Findings

- Many practices were acceptable and followed evidence-based obstetrics.
- Facility preparedness was good.
- Some practices were observed for which insufficient evidence exists.
- However many observed practices were **not** in line with evidence-based obstetrics.

Were the women assessed
properly on arrival?

The general examination in the ER & the prelabour ward

N=188

Blood pressure

Pulse

Temperature

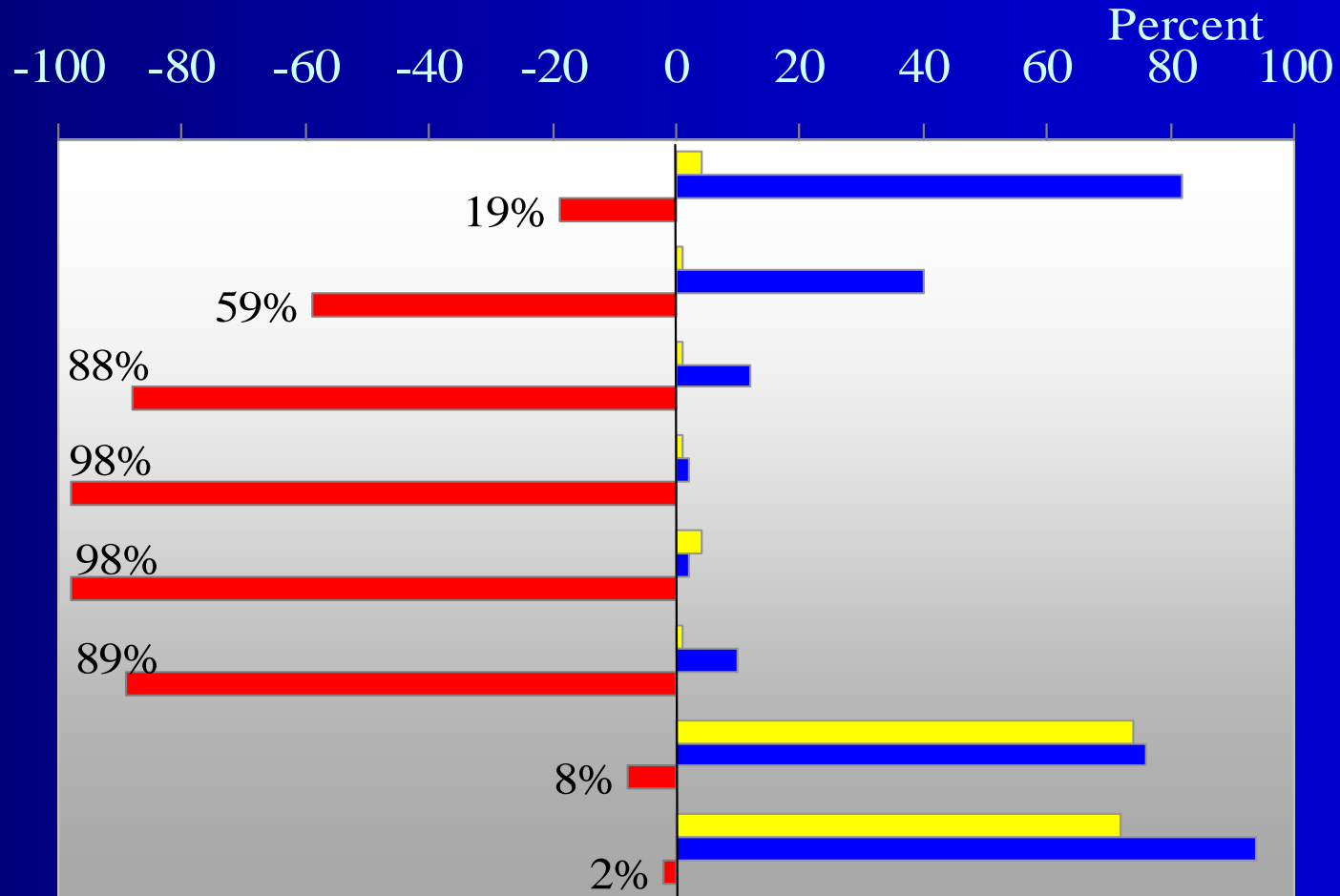
Chest

Heart

Legs

Abdomen

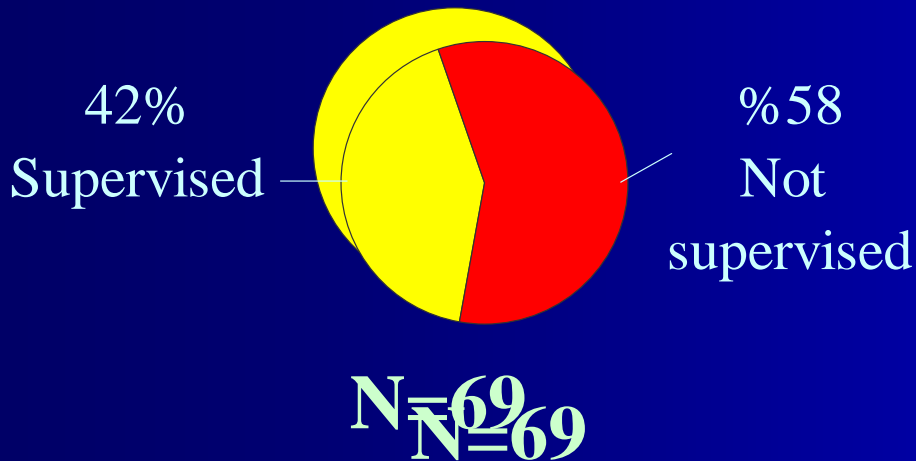
FHS



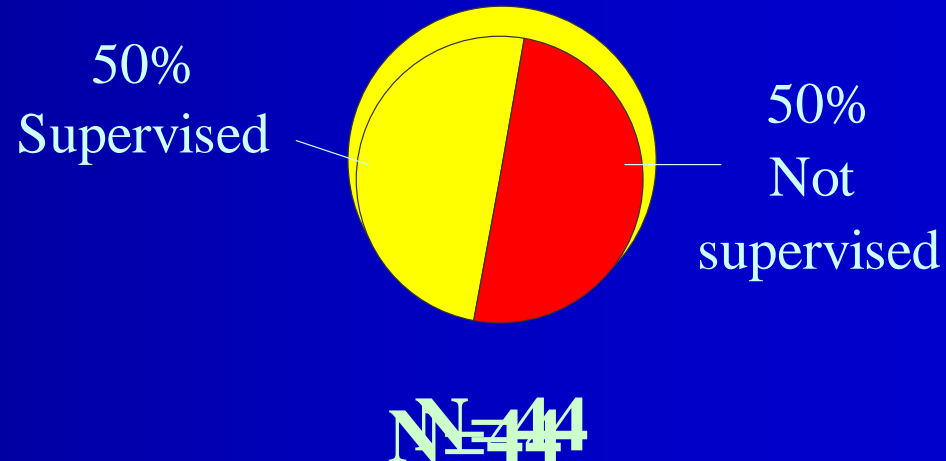
■ Examined in the ER ■ Examined in the prelabour ward ■ Not examined in either

Supervision of house officers

A house officer conducted the delivery in 69 deliveries

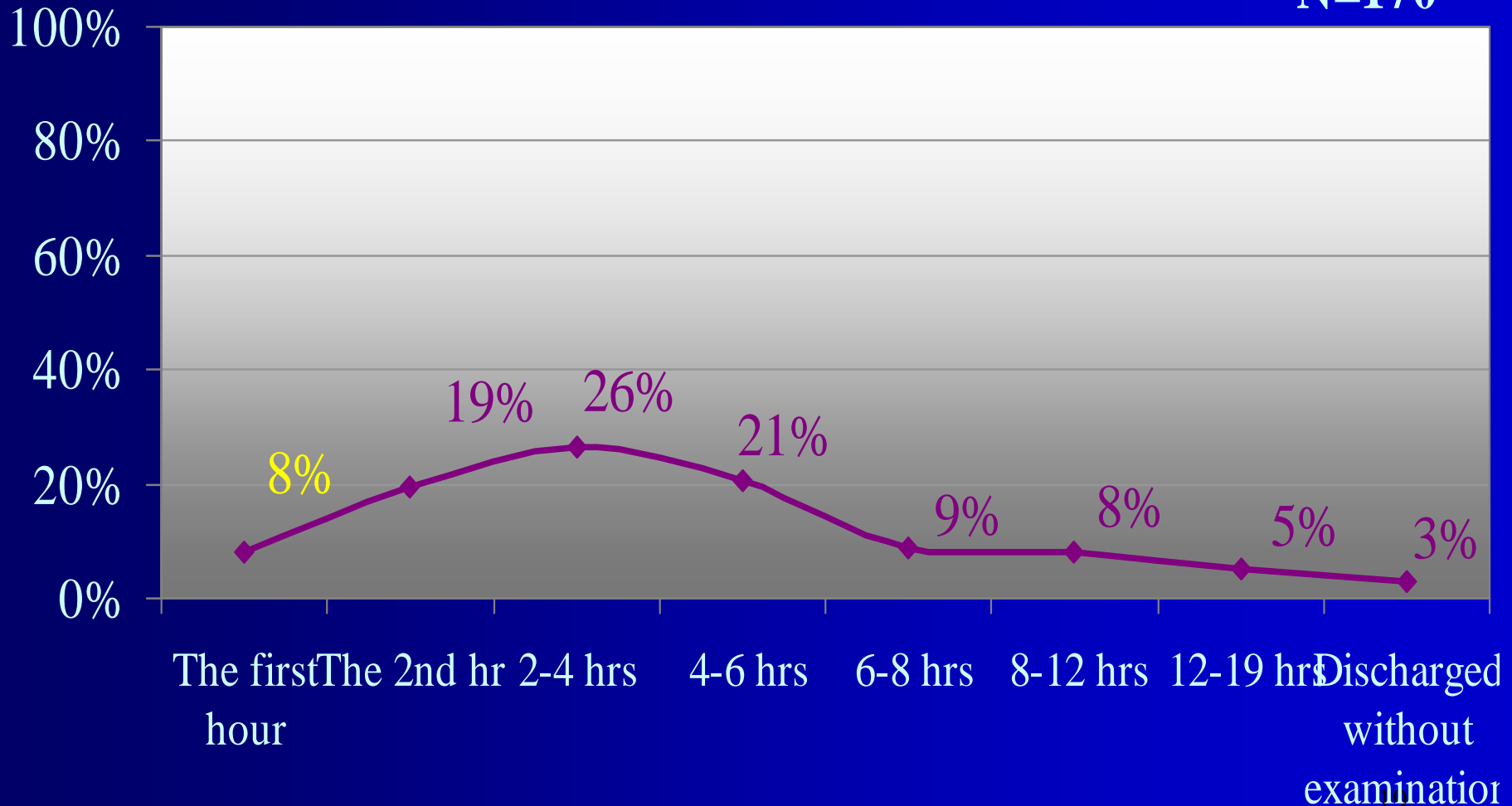


A house officer repaired the episiotomy or tear in 44 deliveries



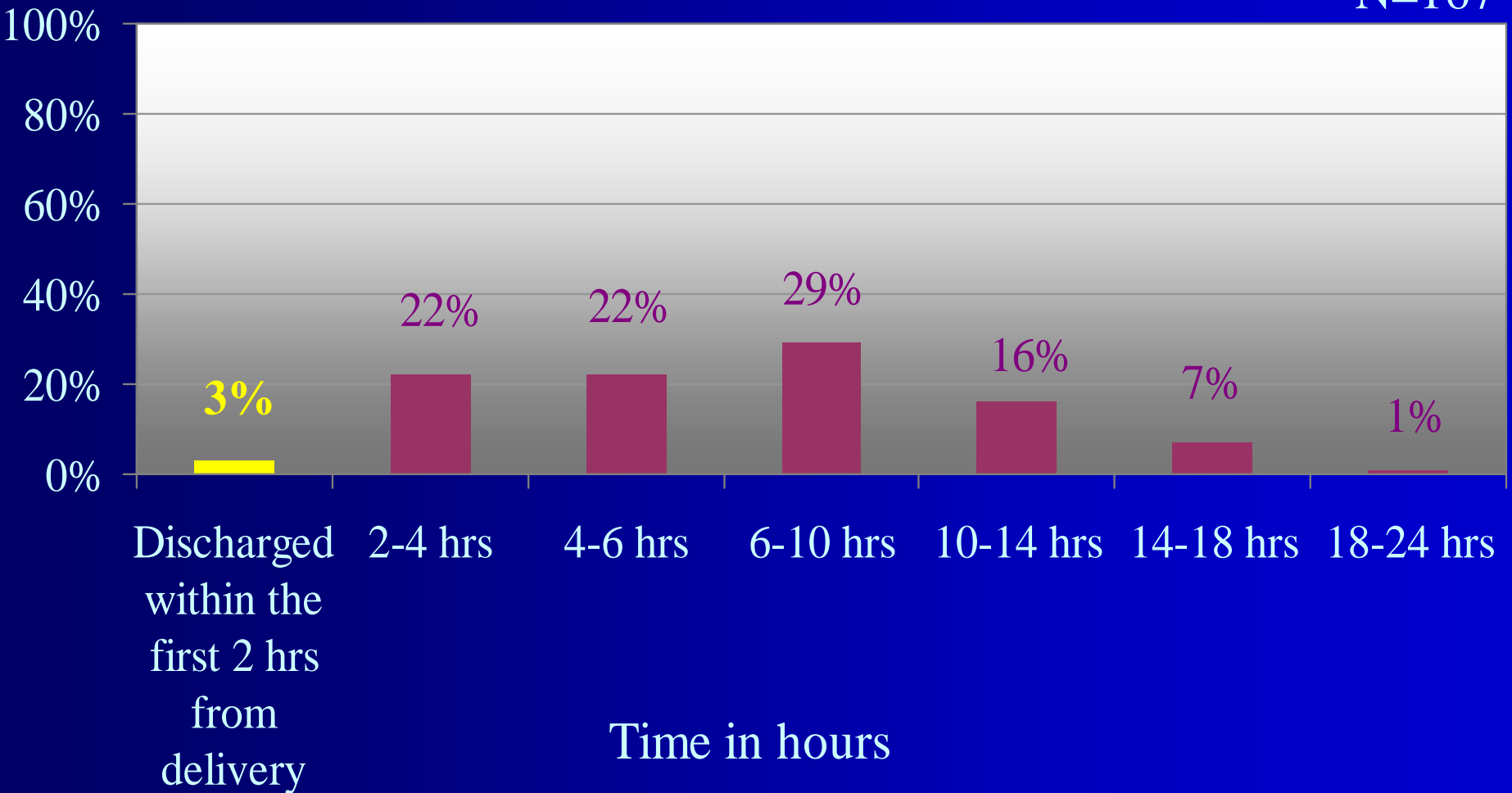
Duration between time of arrival to the postpartum ward until the time of the first exam

N=170



Hours between delivery and leaving the hospital

N=167



Some findings in detail

Detailed examination of oxytocin usage

Oxytocin is a drug that stimulates the uterine contractions, and can be given when the progress of labour is slow.

Side-effects of oxytocin can include uterine hyperstimulation, fetal distress, fetal asphyxia and uterine rupture. (ACOG, 1995)

Oxytocin is also associated with neonatal hyperbilirubinemia. (Kruse, 1986)

“When used carefully, oxytocin is a very safe pharmacologic agent.

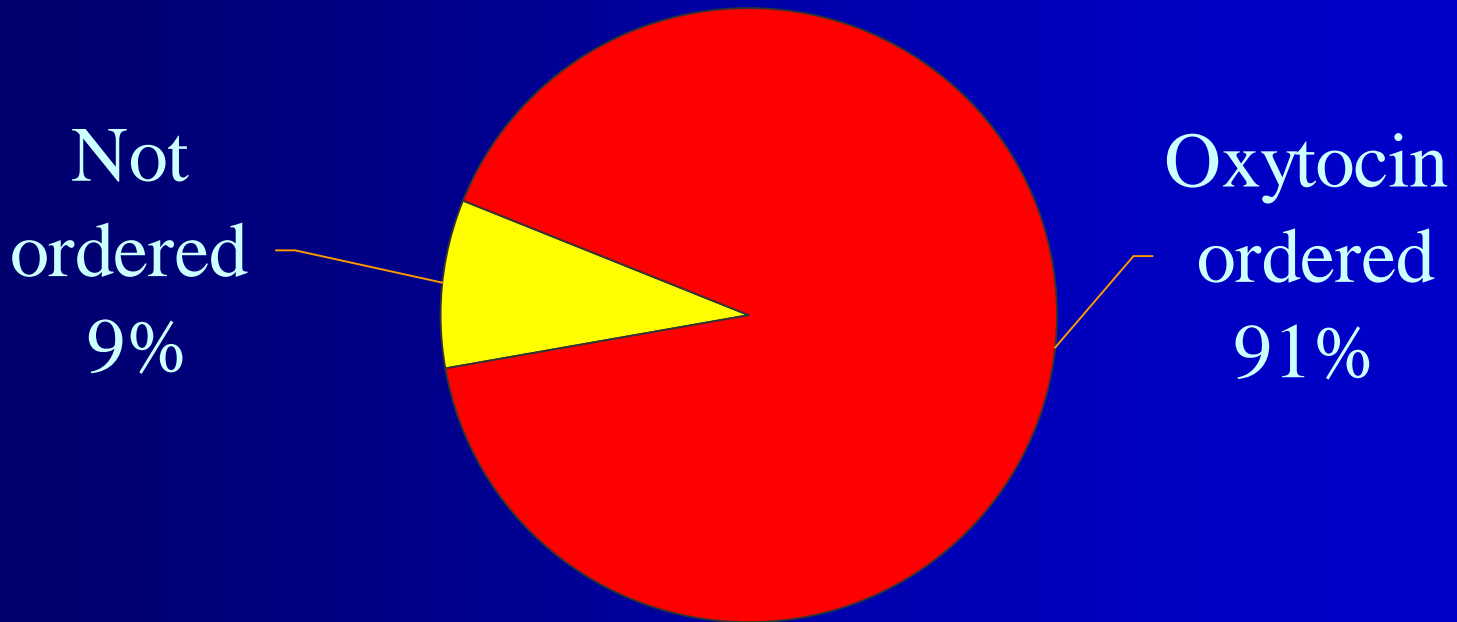
Complications caused by oxytocin are almost always due to **improper doses or inadequate supervision.**”

(Kruse, 1986)

It is estimated that approximately **15 %** of all women in labour in the United States receive oxytocin for augmentation. (Merrill et al., 1999)

There is **no documentation** of either the magnitude or the pattern of oxytocin usage for normal labour in Egyptian obstetric facilities.

What proportion of our sample was given oxytocin ?



N=181

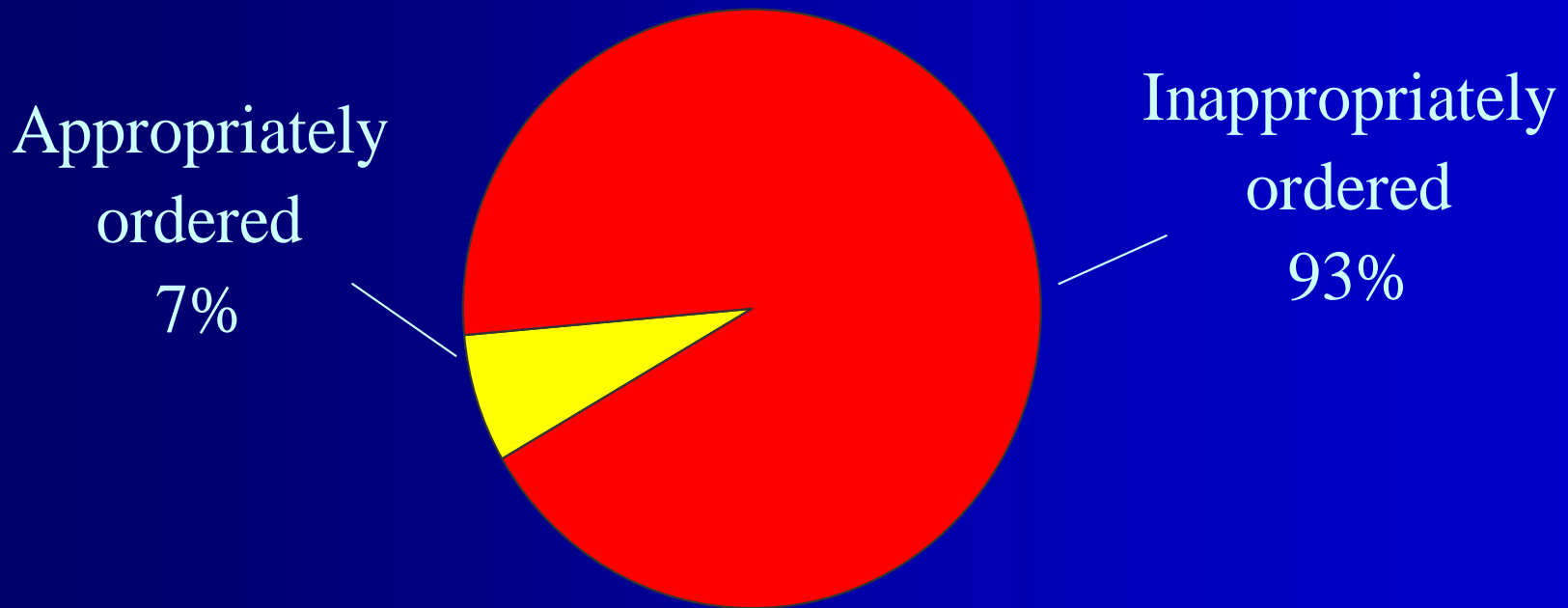
“When oxytocin is given continuous monitoring of the fetal heart rate is important to such an extent that if it cannot be performed, serious consideration should be given to abandoning the effort”

(Stubbs, et al., 2000)

Fetal heart sounds were heard appropriately when oxytocin was given

20%

Was the oxytocin order appropriate?



N=165

Reason for categorizing the oxytocin order as inappropriate

N=154

0% 20% 40% 60% 80% 100%

Ordered at time of 1st vaginal exam

41%

Ordered at the same time as
ARM/SROM (or < 1 hr)

42%

Ordered with the membranes intact

36%

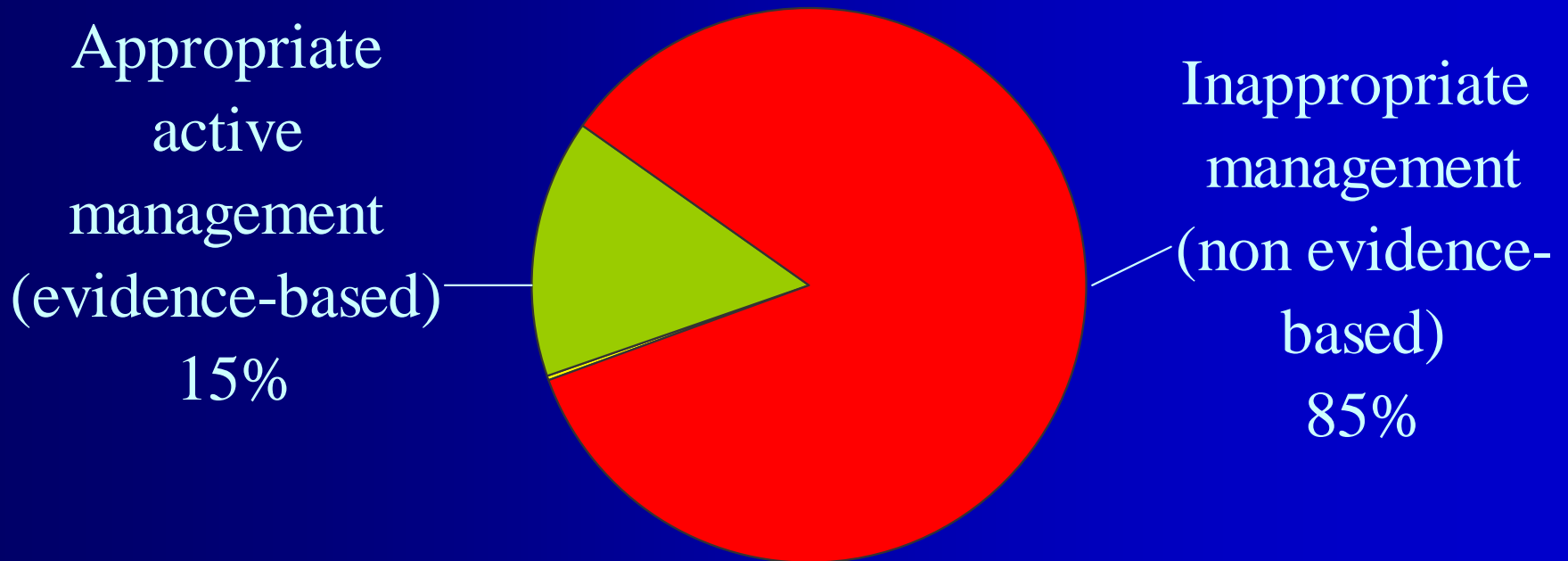
Ordered in spite of good progress

24%

These do not add up to a 100 as there can be more than one reason (45% had 2 reasons)

Oxytocin was ordered for the **majority** of the women observed, and in most cases the order was both **inappropriate** and its use was **not monitored**.

How was the 3rd stage managed ?



N=176

Appropriate (evidence-based) passive management = 0%

Reason for categorizing the active management as inappropriate

N=150

0% 20% 40% 60% 80% 100%

No ecbolic was given



No cord traction



No early clamping of the cord



Ecbolics more than 1 min. after delivery of baby



Ecbolics after the placenta



These do not add up to a 100 as there can be more than one reason (36% had 2 reasons)

The **protective effect** that active management can potentially provide against postpartum hemorrhage **was lost to 85 %** of the women we observed.

What factors could be contributing to the practices we documented?

- Heavy caseload
average deliveries per day = 55
- Providers focus on high-risk cases
- Unavailability of written protocol for normal labour
- Junior providers unaware of factors contributing to maternal mortality in Egypt

The Galaa Intervention Study

2004

In collaboration with the facility administration, the team designed and carried out a training program to address a selection of problematic practices

This was done as a case-control study.

Objectives

The intervention aimed to improve:

- **The third-stage management** (delivery of the placenta) as this reduces post partum hemorrhage or bleeding after delivery.
- **Postpartum** assessment and care
- **Information-sharing**
- **Patient-provider communication**

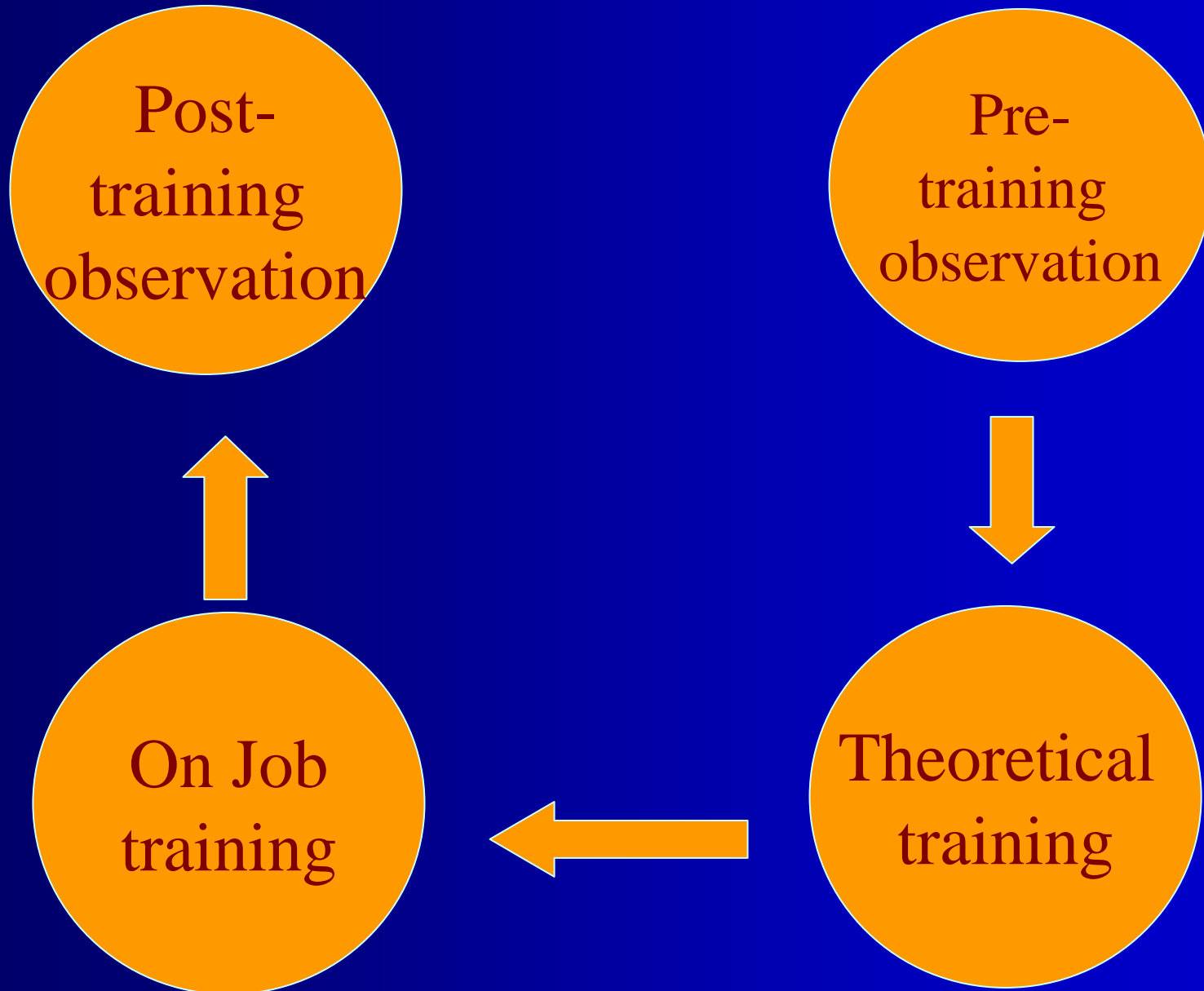
Why did we choose these practices?

- It is **impossible to intervene in all areas** and all practices at the same time.
- We chose practices that might **contribute to the leading cause of maternal mortality in Egypt** (postpartum hemorrhage).
- We found that the **communication** and the **information-sharing** were very deficient.

- We set targets we hoped were **achievable**:

We decided to try to reduce the inappropriate clinical practices (third stage management & postpartum care) by **60%** and the attitude (communication & information-sharing) by **40%**.

Process of the intervention



There are 6 departments in the hospital.

The intervention was carried out in 3 departments (**target** departments).

The 3 other departments were observed as **controls**.

Pre-training observation

A second pre-training observation was conducted in 2004.

Despite great improvement in the physical structure and facilities, the practices were **very similar** to the 2001 findings.

This similarity between 2001 and 2004 shows the need for an intervention.

Theoretical training

- **Three days** of presentations and discussions were carried out.
- A package of session objectives, reading materials and hard copies of all presentations were given as **handouts**.
- **32 obstetricians** were involved: residents, assistant specialists, specialists and heads of departments.
- **Staff specialists** were assigned as trainers for the **on-job training**.
- One social worker and all the **nurses** in charge of the delivery room and the postpartum ward were trained.

On-job training

- Took place in the three target departments over **four weeks.**
- Providers, including doctors & nurses, were trained during the course of their scheduled shifts in the labour ward.
- Providers were trained on appropriate third-stage management, improving the postpartum care and assessment, and enhancing their interpersonal communication skills and information-sharing with women in the postpartum ward.

Intervention outcomes:

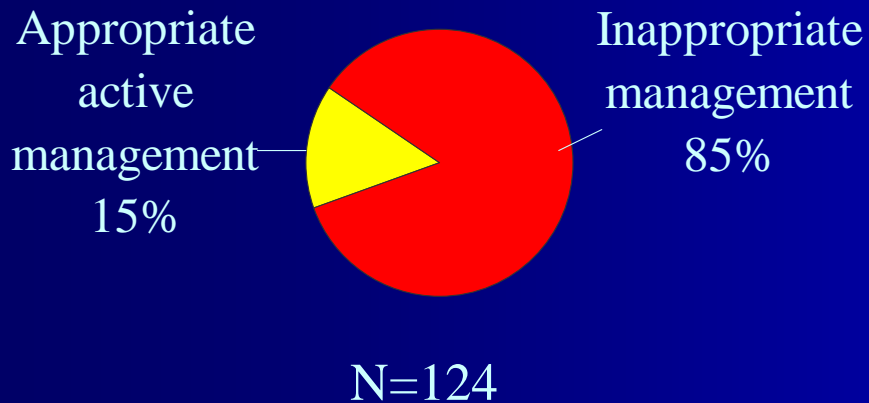
Post-training practices as compared
to pre-training practices

All practices improved, although not all the intervention targets were met

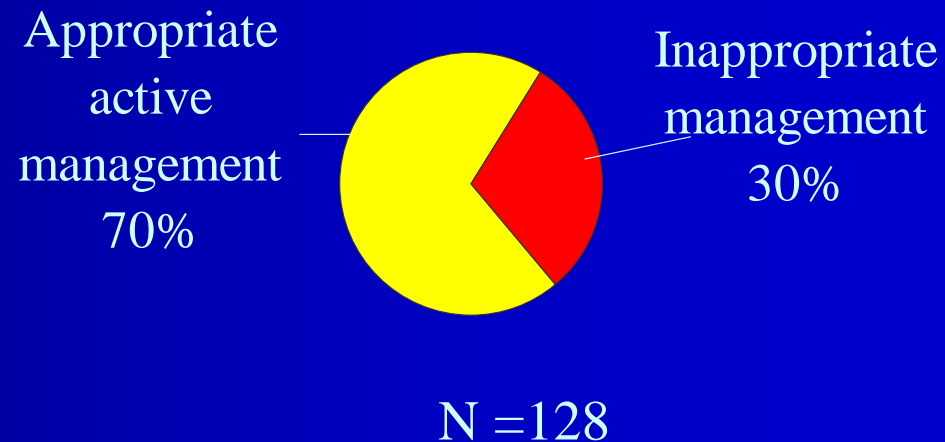
Management of the third stage of labour

Management of the 3rd stage showed improvement, exceeding our target.

Pre-intervention



Post-intervention



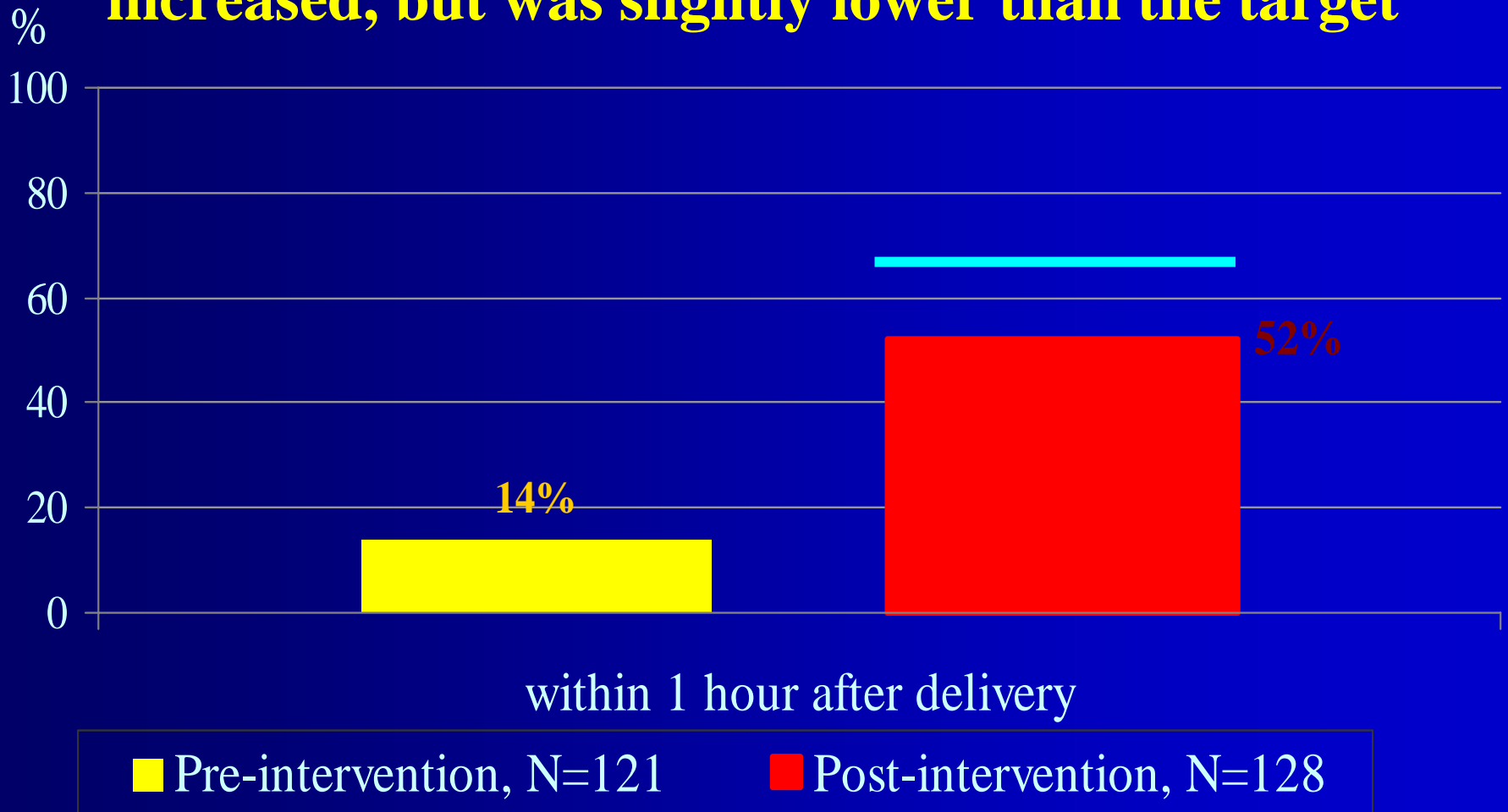
Target = 66%

Achieved=70%

p<0.001

Postpartum Care

Percent of women examined in 1st hour after delivery increased, but was slightly lower than the target

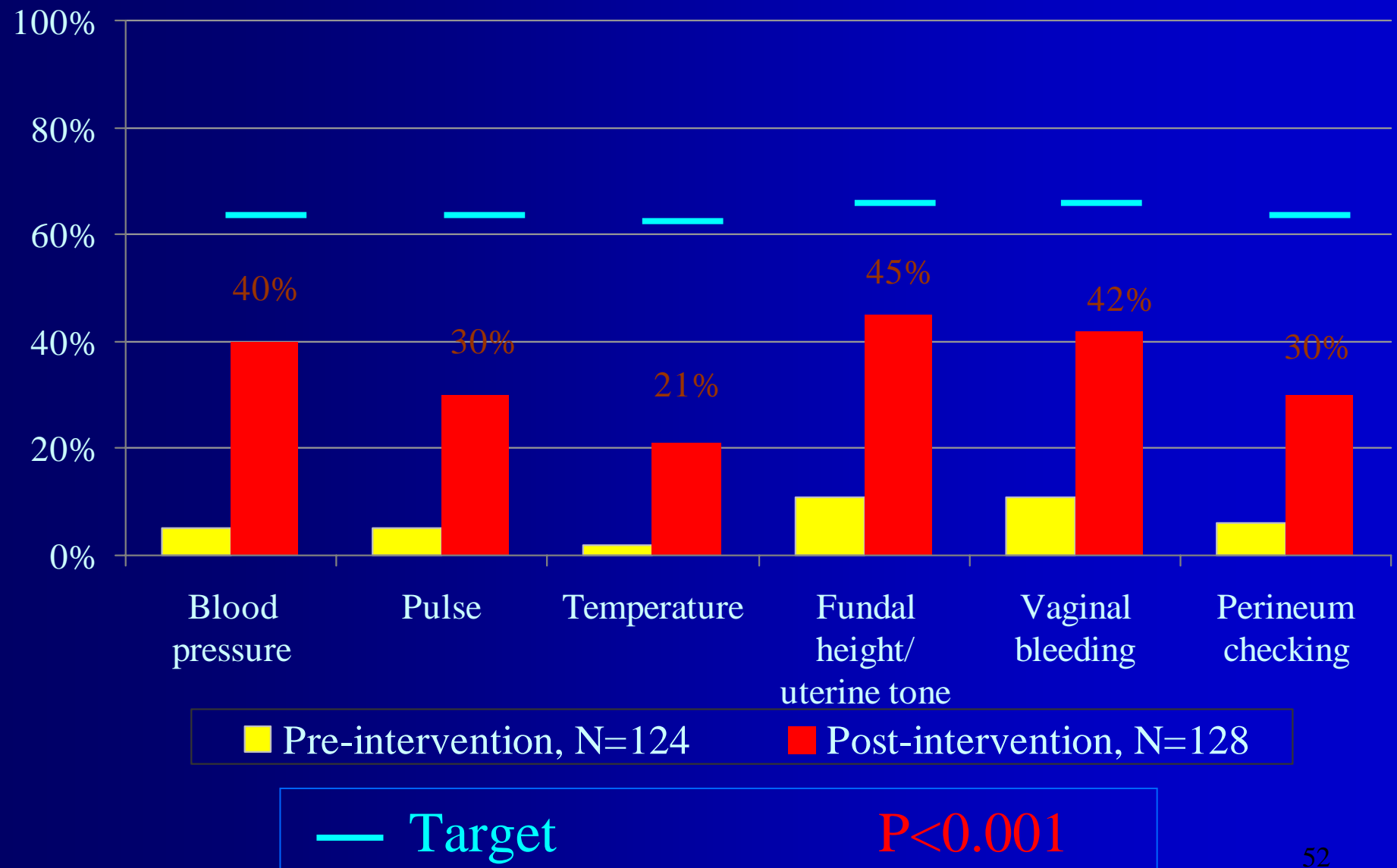


Target = 66%

Achieved=52%

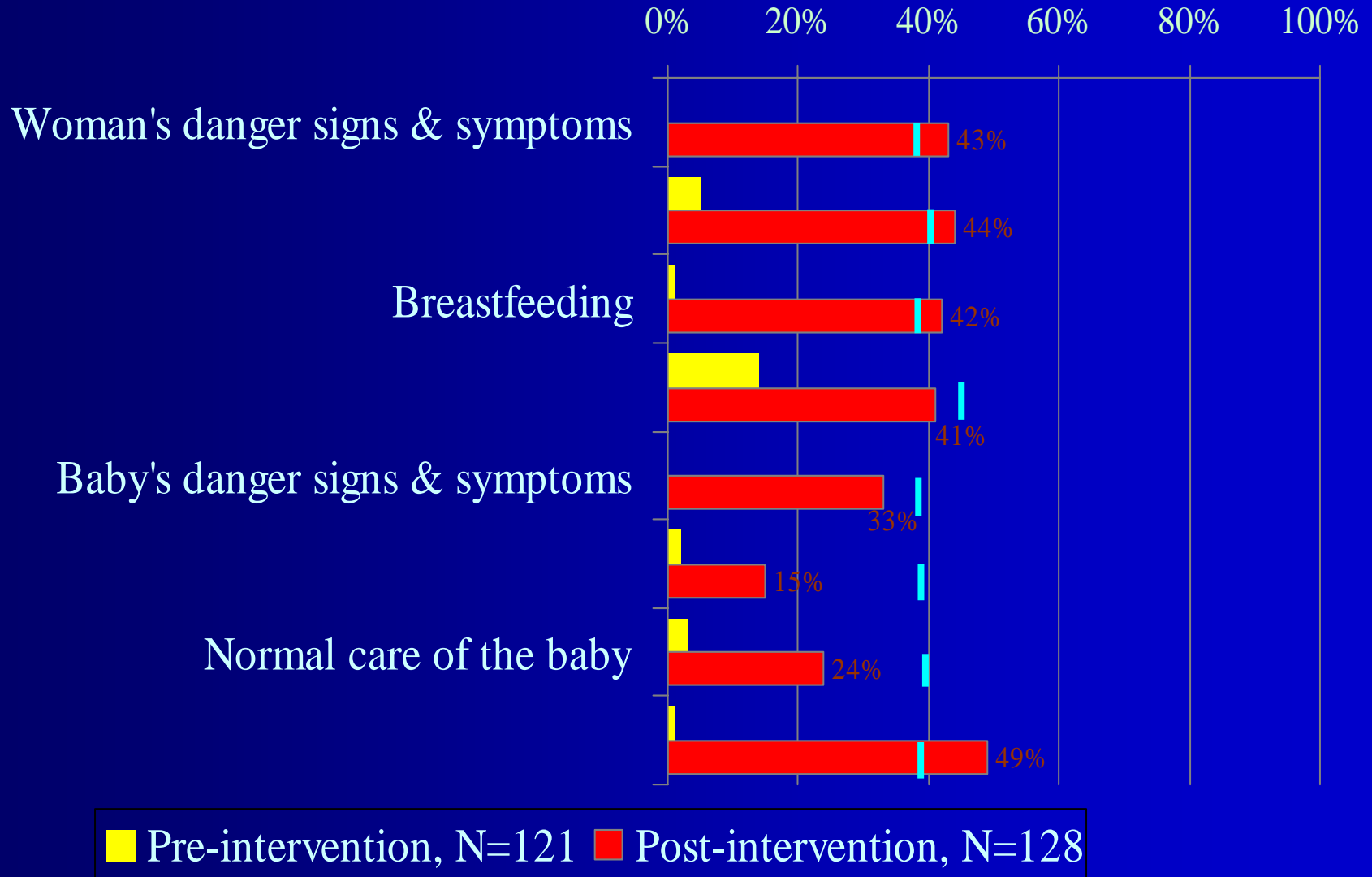
$p < 0.001$

Items checked in the first hour after delivery increased



Information-sharing with the women

Information given to the lady about her health and the baby's health and care increased



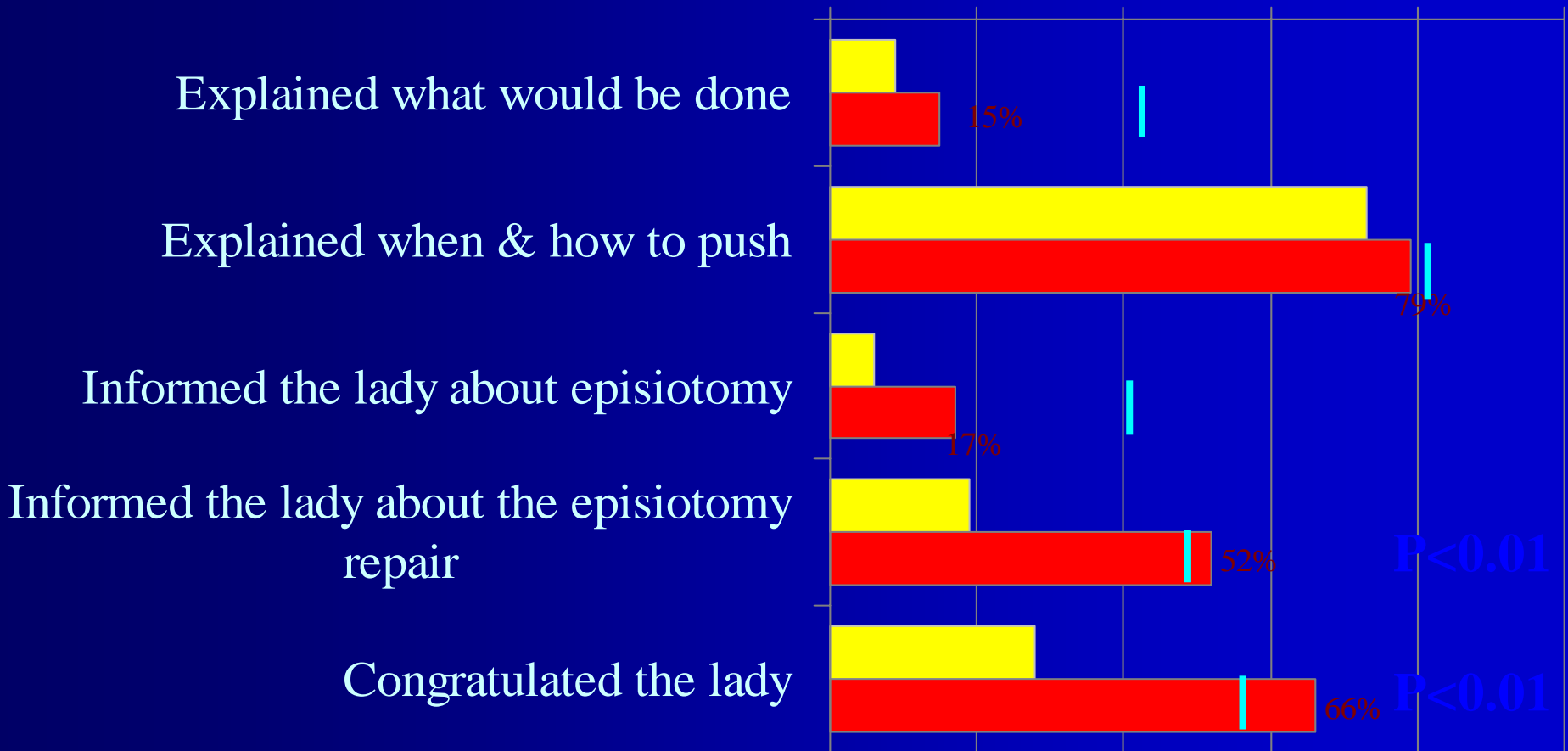
— Target **P<0.01**

Communication

Communication in the delivery room improved

— Target

0% 20% 40% 60% 80% 100%



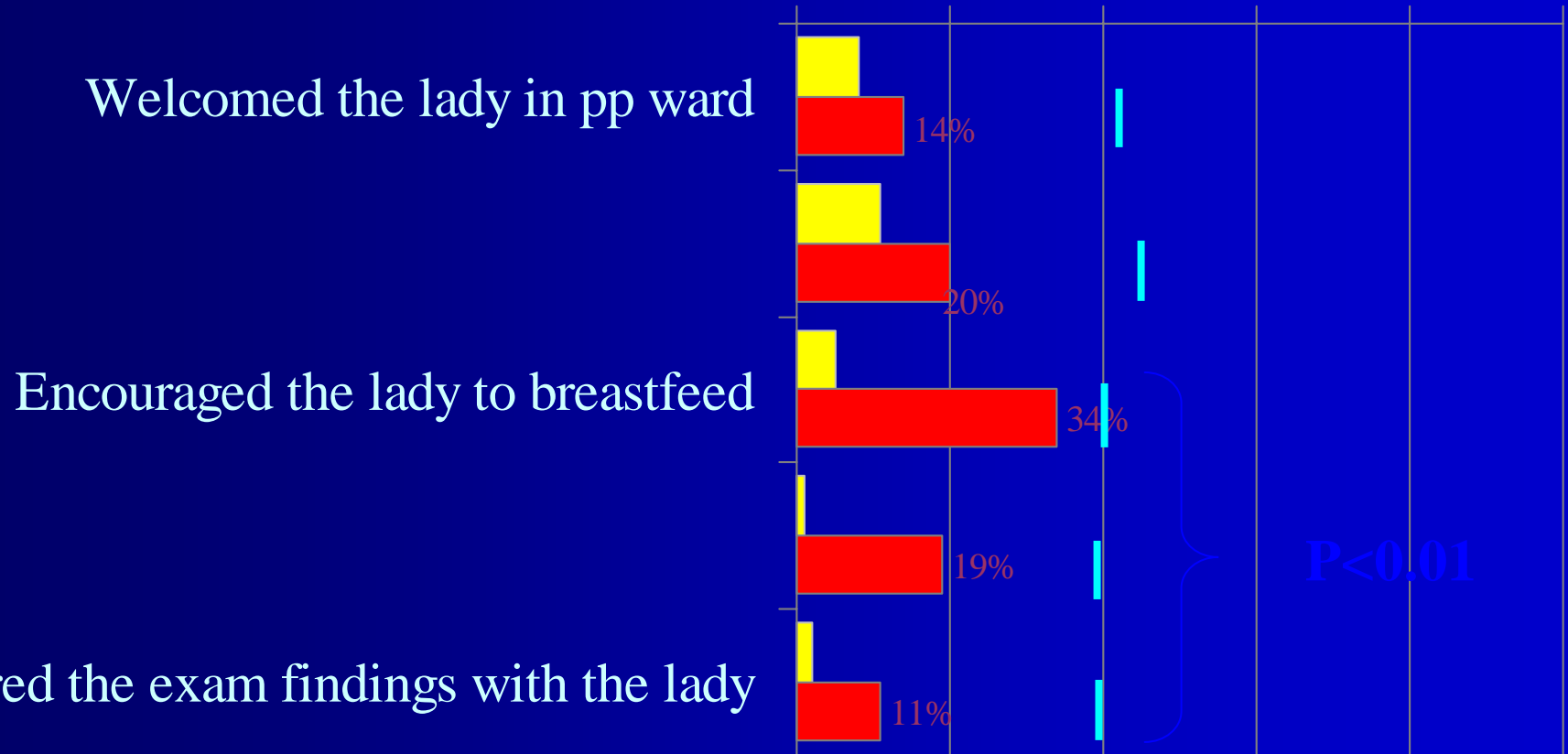
■ Pre-intervention, N=124

■ Post-intervention, N=128

Communication in the postpartum ward improved

— Target

0% 20% 40% 60% 80% 100%



$P < 0.01$

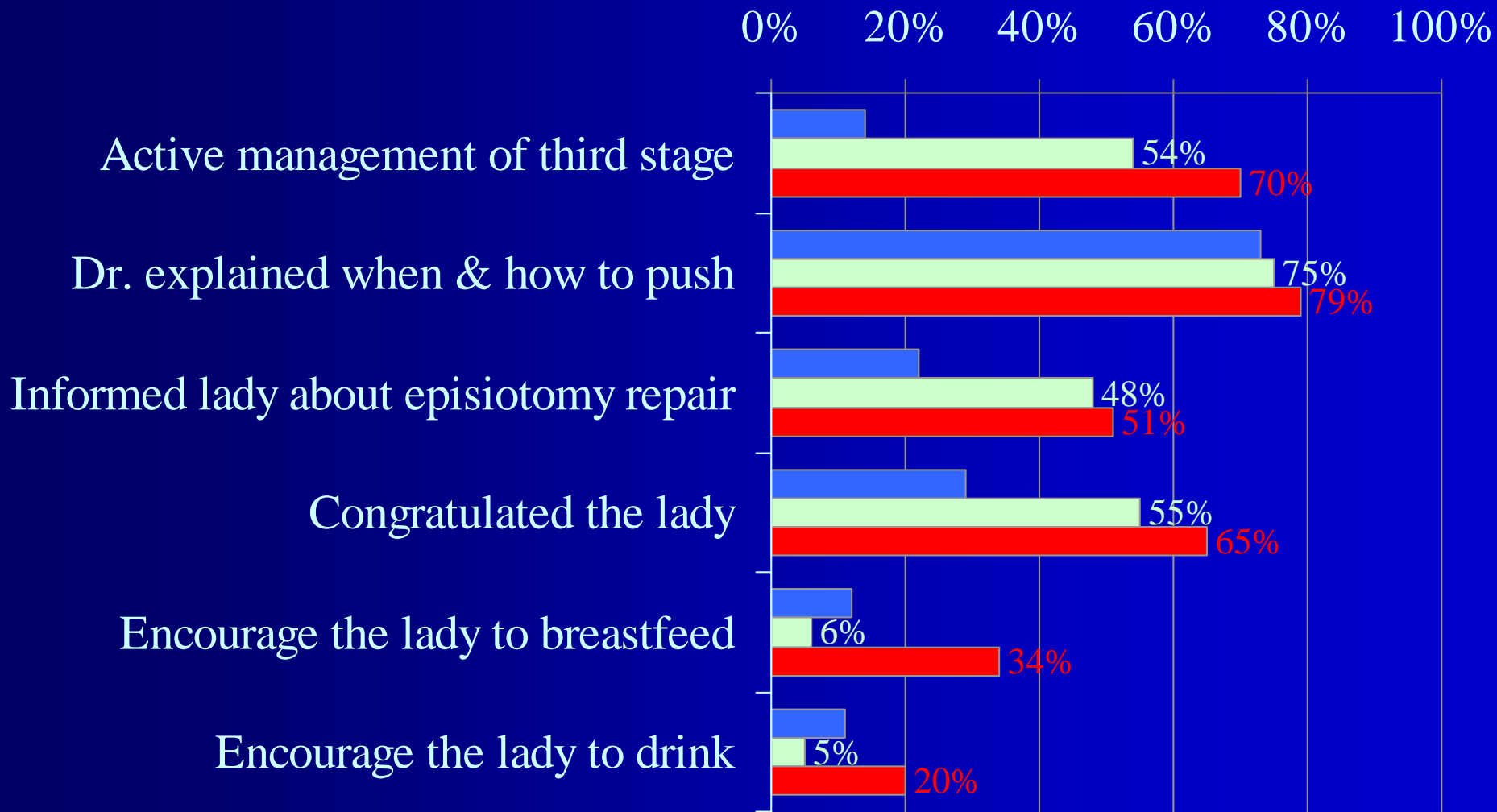
■ Pre-intervention, N=121

■ Post-intervention, N=128

You have seen improvements in the target groups.

What about the departments that were controls and did not receive training?

Practices observed in the control and target departments

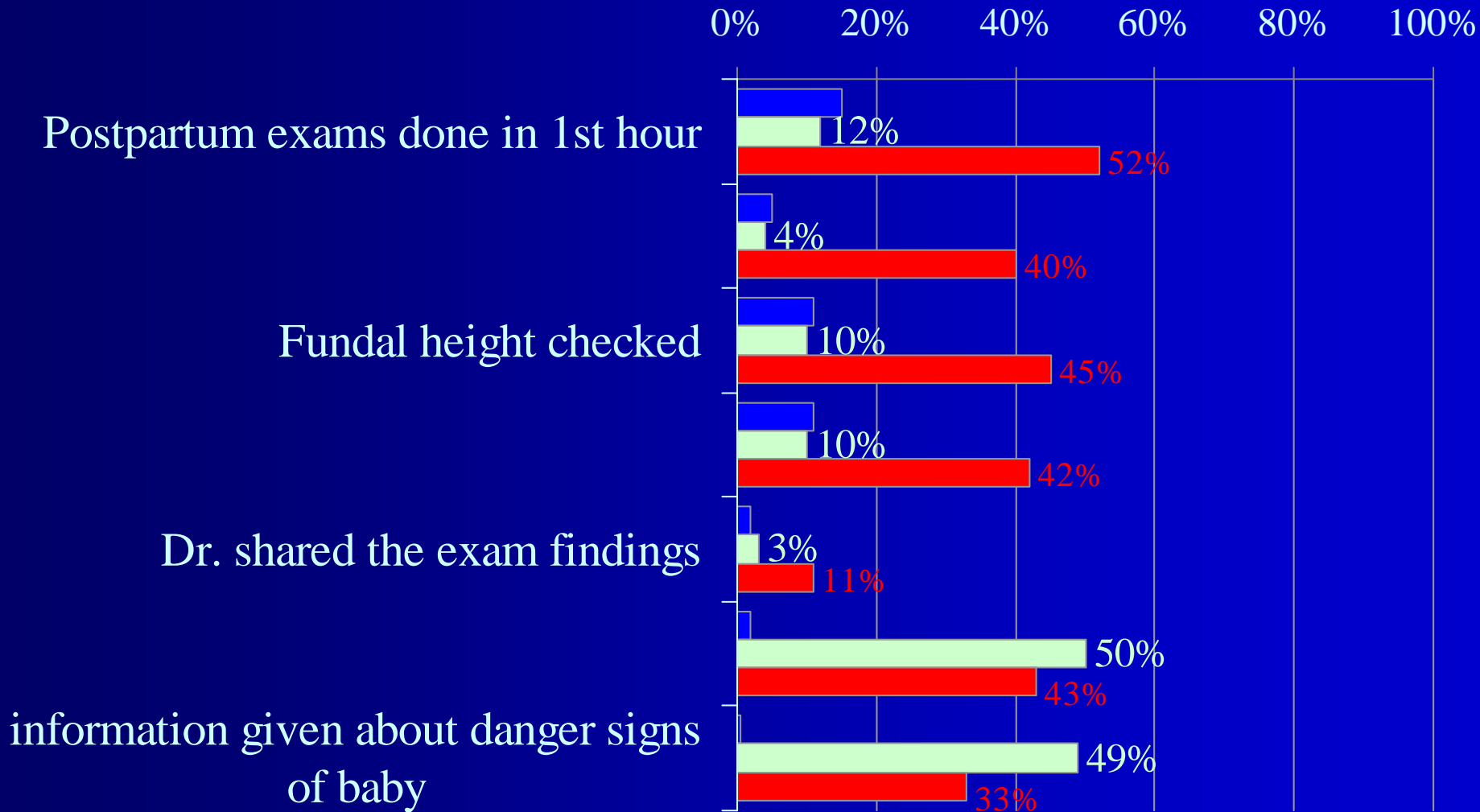


■ pre-intervention, N=124

■ Control, N=97

■ Target, N=128

Practices observed in the control and target departments



Pre-intervention, N=124

Control, N=97

Target, N=128

Why did some of the practices improve in the control group?

Reasons for improvement of the control groups

- **The most remarkable improvement happened in information-sharing which depended mainly on nurses. They were part of both the control and target groups -this could not be avoided for staffing reasons.**
- **There were some doctors from the target departments who also served in the control departments due to shortage of staff.**
- **There was a poster in the delivery room demonstrating the steps of appropriate management of the 3rd stage which the control group shared.**

Overall conclusions

- **Directly observing** practices using a specially developed checklist is the only way to accurately record actual practices.
- Direct observations revealed a **divergence between actual practices and evidence-based medicine**
- Short, targeted training **can improve** both intrapartum and postpartum facility care.

- Practices did not change between 2001 and 2004 despite great improvements to the facility's physical infrastructure.
- This indicates that interventions in the form of regular provider training and assessment are much needed as well.
- Trained departments showed improvement after the intervention

- Some practices did not meet the targets we set but significantly improved (e.g. postpartum assessment and care)
- Communication items were the most difficult to change (e.g. explaining when and how to push)
- Control departments also improved and showed improvement due to contamination

Points for discussion

- Need for another observation at a later time to check the sustainability of the intervention.
- Need for more or repeated trainings to sustain the improvement over time
- Regular, in-house training should be included in all facilities, with periodic assessments of provider performance.

The study was supported by:

- The Reproductive Health Working Group
- The Ford Foundation
- The Population Council
- The Social Research Center, American University in Cairo

For further information please refer to these Galaa study publications:

- Sholkamy H, Khalil K, Cherine M, Elnoury A, Breebaart M, Hassanein N. An observation checklist for facility-based normal labor and delivery practices: The Galaa study. *Monographs in Reproductive Health* No. 5. Cairo: Population Council, 2003.
- Khalil K, Cherine M, Elnoury A, Sholkamy H, Breebaart M, Hassanein N. Labor augmentation in an Egyptian teaching hospital. *Int. J Gynecol Obstet* 2004, 85:74-80.

- Cherine et al. Management of the third stage of labor in an Egyptian teaching hospital. *Int. J Gynecol Obstet* 2004; 87:54-58.
- Hospital practices vs. Evidence-Based medicine: Categorization of normal birth practices in an Egyptian Teaching Hospital. K Khalil, A Elnoury, M Cherine, N Hassanein, H Sholkamy, M Breebaart and A Shoubary. *Birth: Issues in Perinatal Care* 2005; 32: 284-291.
- Choices and Challenges in Changing Childbirth Research Network.. Routines in facility-based maternity care: Evidence from the Arab World. *British Journal of Obstetrics and Gynecology* 2005; 112:1-7.