

**Clinical trial results:
Ropivacaine Perineal Infiltration for Postpartum Pain Management in
Episiotomy Repair: a Double-blind, Randomised, Placebo-controlled
Trial****Summary**

EudraCT number	2016-002786-62
Trial protocol	FR
Global end of trial date	04 November 2020

Results information

Result version number	v1 (current)
This version publication date	03 September 2022
First version publication date	03 September 2022

Trial information**Trial identification**

Sponsor protocol code	CHD065-16
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Additional study identifiers

ISRCTN number	-
ClinicalTrials.gov id (NCT number)	NCT03084549
WHO universal trial number (UTN)	-

Notes:

Sponsors

Sponsor organisation name	Centre Hospitalier Départemental Vendée
Sponsor organisation address	Bd Stephane MOREAU, La Roche sur Yon, France, 85925
Public contact	Dorion Agnès, Centre Hospitalier Départemental Vendée, +33 251446380, agnes.dorion@ght85.fr
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Notes:

Paediatric regulatory details

Is trial part of an agreed paediatric investigation plan (PIP)	No
Does article 45 of REGULATION (EC) No 1901/2006 apply to this trial?	No
Does article 46 of REGULATION (EC) No 1901/2006 apply to this trial?	No

Notes:

Results analysis stage

Analysis stage	Final
Date of interim/final analysis	24 June 2021
Is this the analysis of the primary completion data?	Yes
Primary completion date	06 May 2020
Global end of trial reached?	Yes
Global end of trial date	04 November 2020
Was the trial ended prematurely?	No

Notes:

General information about the trial

Main objective of the trial:

The primary outcome was the rate of perineal pain, defined by a Numerical Pain Rating Scale (NPRS) exceeding 3/10, in the mid-term (day 7) postpartum period.

Protection of trial subjects:

All adverse events or reactions (except those specified in the protocol), whether expected or unexpected, serious or not, were collected in the eCRF.

The follow-up of events or adverse reactions, serious or not, was ensured until resolution or consolidation

Background therapy: -

Evidence for comparator: -

Actual start date of recruitment	24 October 2017
Long term follow-up planned	Yes
Long term follow-up rationale	Efficacy
Long term follow-up duration	6 Months
Independent data monitoring committee (IDMC) involvement?	No

Notes:

Population of trial subjects

Subjects enrolled per country

Country: Number of subjects enrolled	France: 272
Worldwide total number of subjects	272
EEA total number of subjects	272

Notes:

Subjects enrolled per age group

In utero	0
Preterm newborn - gestational age < 37 wk	0
Newborns (0-27 days)	0
Infants and toddlers (28 days-23 months)	0
Children (2-11 years)	0
Adolescents (12-17 years)	0
Adults (18-64 years)	272

From 65 to 84 years	0
85 years and over	0

Subject disposition

Recruitment

Recruitment details:

Inclusion period: 36 months

First inclusion: 2017/20/24

Last inclusion: 2020/04/29

Pre-assignment

Screening details:

Women who had given birth to a singleton foetus in cephalic presentation at 37 weeks of gestation or more, with spontaneous or operative vaginal delivery under epidural anaesthesia and who had received mediolateral episiotomy were eligible for enrolment in the study.

Period 1

Period 1 title	Overall trial (overall period)
Is this the baseline period?	Yes
Allocation method	Randomised - controlled
Blinding used	Double blind
Roles blinded	Subject, Investigator

Arms

Are arms mutually exclusive?	Yes
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Arm title	Ropivacaine
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Arm description:

Eligible women were receive 75 mg of ropivacaine (Fresenius Kabi) in a 20-mL syringe (10-mL of ropivacaine 7.5mg/mL and 10-mL of normal saline)

Arm type	Experimental
Investigational medicinal product name	Ropivacaine
Investigational medicinal product code	N01BB09
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Injection

Dosage and administration details:

syringe with 10-mL of ropivacaine 7.5mg/mL and 10-mL of normal saline

One injection

Arm title	Placebo
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Arm description:

20 mL of normal saline

Arm type	Placebo
Investigational medicinal product name	Chlorure de sodium
Investigational medicinal product code	B05XA03
Other name	
Pharmaceutical forms	Solution for injection
Routes of administration	Injection

Dosage and administration details:

20 mL

one injection

Number of subjects in period 1	Ropivacaine	Placebo
Started	135	137
immediate postpartum period H12,H24,H48	135	137
day 7 postpartum	135	137
long-term postpartum M3	135	137
long-term postpartum M6	135	137
Completed	135	137

Baseline characteristics

Reporting groups

Reporting group title	Ropivacaine
Reporting group description: Eligible women were receive 75 mg of ropivacaine (Fresenius Kabi) in a 20-mL syringe (10-mL of ropivacaine 7.5mg/mL and 10-mL of normal saline)	
Reporting group title	Placebo
Reporting group description: 20 mL of normal saline	

Reporting group values	Ropivacaine	Placebo	Total
Number of subjects	135	137	272
Age categorical			
Units: Subjects			
In utero	0	0	0
Preterm newborn infants (gestational age < 37 wks)	0	0	0
Newborns (0-27 days)	0	0	0
Infants and toddlers (28 days-23 months)	0	0	0
Children (2-11 years)	0	0	0
Adolescents (12-17 years)	0	0	0
Adults (18-64 years)	135	137	272
From 65-84 years	0	0	0
85 years and over	0	0	0
Age continuous			
Units: years			
arithmetic mean	30.0	30.3	-
standard deviation	± 4.5	± 4.6	-
Gender categorical			
Units: Subjects			
Female	135	137	272
Male	0	0	0

End points

End points reporting groups

Reporting group title	Ropivacaine
Reporting group description:	Eligible women were receive 75 mg of ropivacaine (Fresenius Kabi) in a 20-mL syringe (10-mL of ropivacaine 7.5mg/mL and 10-mL of normal saline)
Reporting group title	Placebo
Reporting group description:	20 mL of normal saline

Primary: Perineal pain

End point title	Perineal pain
End point description:	The primary outcome was the analgesic efficacy of ropivacaine at day 7 postpartum (medium term) measured with the numerical pain rating scale (NPRS), exceeding 3/10 in the perineal repair area
End point type	Primary
End point timeframe:	day 7 postpartum

End point values	Ropivacaine	Placebo		
Subject group type	Reporting group	Reporting group		
Number of subjects analysed	114	115		
Units: number	39	35		

Statistical analyses

Statistical analysis title	Perineal Pain
Comparison groups	Ropivacaine v Placebo
Number of subjects included in analysis	229
Analysis specification	Pre-specified
Analysis type	superiority
P-value	≤ 0.05
Method	Regression, Linear

Adverse events

Adverse events information

Timeframe for reporting adverse events:

data were collected from randomisation to the immediate postpartum period at H48

Assessment type	Systematic
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Dictionary used

Dictionary name	MedDRA
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Dictionary version	24.0
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Reporting groups

Reporting group title	Ropivacaine
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Reporting group description: -

Reporting group title	Placebo
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Reporting group description: -

Serious adverse events	Ropivacaine	Placebo	
Total subjects affected by serious adverse events			
subjects affected / exposed	3 / 135 (2.22%)	4 / 137 (2.92%)	
number of deaths (all causes)	0	0	
number of deaths resulting from adverse events	0	0	
Injury, poisoning and procedural complications			
Incorrect route of product administration			
subjects affected / exposed	0 / 135 (0.00%)	1 / 137 (0.73%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Incision site haemorrhage			
subjects affected / exposed	0 / 135 (0.00%)	1 / 137 (0.73%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Pregnancy, puerperium and perinatal conditions			
Perineal haematoma			
subjects affected / exposed	1 / 135 (0.74%)	0 / 137 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Postpartum haemorrhage			

subjects affected / exposed	0 / 135 (0.00%)	2 / 137 (1.46%)	
occurrences causally related to treatment / all	0 / 0	0 / 2	
deaths causally related to treatment / all	0 / 0	0 / 0	
Reproductive system and breast disorders			
Lactation disorder			
subjects affected / exposed	1 / 135 (0.74%)	0 / 137 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	
Renal and urinary disorders			
Urinary retention			
subjects affected / exposed	0 / 135 (0.00%)	1 / 137 (0.73%)	
occurrences causally related to treatment / all	0 / 0	0 / 1	
deaths causally related to treatment / all	0 / 0	0 / 0	
Infections and infestations			
Genital abscess			
subjects affected / exposed	1 / 135 (0.74%)	0 / 137 (0.00%)	
occurrences causally related to treatment / all	0 / 1	0 / 0	
deaths causally related to treatment / all	0 / 0	0 / 0	

Frequency threshold for reporting non-serious adverse events: 5 %

Non-serious adverse events	Ropivacaine	Placebo	
Total subjects affected by non-serious adverse events			
subjects affected / exposed	116 / 135 (85.93%)	129 / 137 (94.16%)	
Injury, poisoning and procedural complications			
Incision site haematoma			
subjects affected / exposed	4 / 135 (2.96%)	1 / 137 (0.73%)	
occurrences (all)	4	1	
Incision site haemorrhage			
subjects affected / exposed	1 / 135 (0.74%)	1 / 137 (0.73%)	
occurrences (all)	1	1	
Incision site pain			
subjects affected / exposed	97 / 135 (71.85%)	107 / 137 (78.10%)	
occurrences (all)	129	135	
Postoperative wound complication			

subjects affected / exposed occurrences (all)	3 / 135 (2.22%) 3	1 / 137 (0.73%) 1	
Suture related complication subjects affected / exposed occurrences (all)	2 / 135 (1.48%) 2	1 / 137 (0.73%) 1	
Vascular disorders Haematoma subjects affected / exposed occurrences (all)	21 / 135 (15.56%) 21	15 / 137 (10.95%) 15	
Nervous system disorders Headache subjects affected / exposed occurrences (all)	1 / 135 (0.74%) 1	3 / 137 (2.19%) 3	
Pregnancy, puerperium and perinatal conditions Perineal haematoma subjects affected / exposed occurrences (all)	8 / 135 (5.93%) 8	14 / 137 (10.22%) 14	
Postpartum haemorrhage subjects affected / exposed occurrences (all)	15 / 135 (11.11%) 15	17 / 137 (12.41%) 17	
Blood and lymphatic system disorders Anaemia subjects affected / exposed occurrences (all)	6 / 135 (4.44%) 6	6 / 137 (4.38%) 6	
General disorders and administration site conditions Pain subjects affected / exposed occurrences (all)	8 / 135 (5.93%) 9	11 / 137 (8.03%) 12	
Pyrexia subjects affected / exposed occurrences (all)	2 / 135 (1.48%) 2	1 / 137 (0.73%) 1	
Oedema subjects affected / exposed occurrences (all)	7 / 135 (5.19%) 7	7 / 137 (5.11%) 7	
Reproductive system and breast disorders			

Oedema genital subjects affected / exposed occurrences (all)	3 / 135 (2.22%) 3	8 / 137 (5.84%) 8	
Gastrointestinal disorders Haemorrhoids subjects affected / exposed occurrences (all)	1 / 135 (0.74%) 1	4 / 137 (2.92%) 4	
Musculoskeletal and connective tissue disorders Coccydynia subjects affected / exposed occurrences (all)	0 / 135 (0.00%) 0	2 / 137 (1.46%) 2	
Infections and infestations Amniotic cavity infection subjects affected / exposed occurrences (all) Urinary tract infection subjects affected / exposed occurrences (all)	1 / 135 (0.74%) 1 2 / 135 (1.48%) 2	1 / 137 (0.73%) 1 1 / 137 (0.73%) 1	

More information

Substantial protocol amendments (globally)

Were there any global substantial amendments to the protocol? Yes

Date	Amendment
22 May 2017	Modification of the dose and administered concentration of Ropivacaine
21 March 2018	Modification of the labelling of the treatment boxes
12 September 2018	Modification of AE notification procedure updated consent form (implementation of REGULATION (EU) 2016/679 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL)
26 March 2019	extension of study period (+18 months)
02 March 2020	update Ropivacaine's Summary of Product Characteristics

Notes:

Interruptions (globally)

Were there any global interruptions to the trial? No

Limitations and caveats

None reported

Online references

<http://www.ncbi.nlm.nih.gov/pubmed/35876236>