



PROVIDE 2.0 Data: Definitions, Processes and Tools

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Partnering to Improve Health Care Quality
for Mothers and Babies



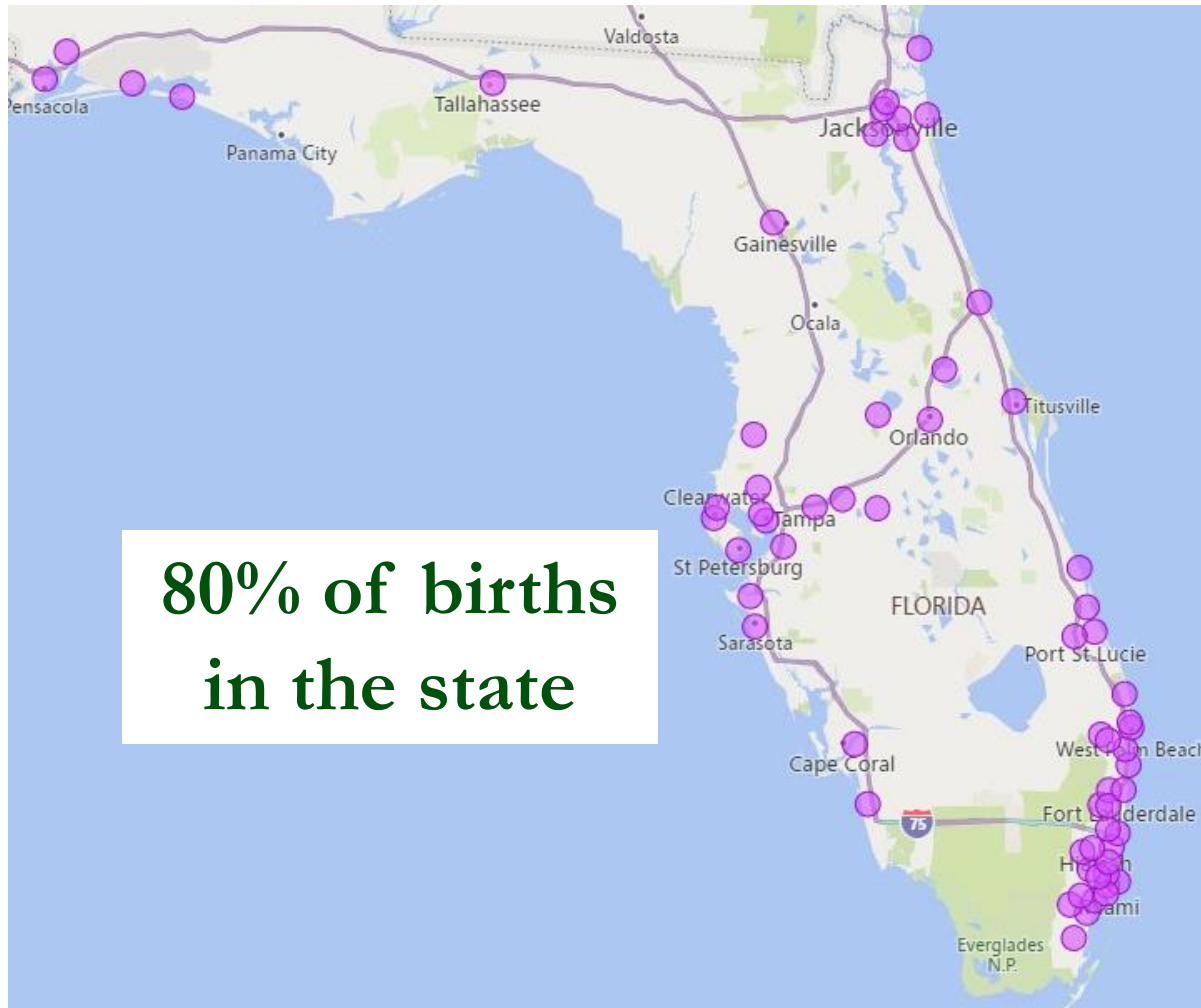


PROVIDE's Goal: To improve maternal and newborn outcomes by applying evidence-based interventions to promote primary vaginal deliveries at Florida delivery hospitals and ultimately reduce NTSV cesareans.

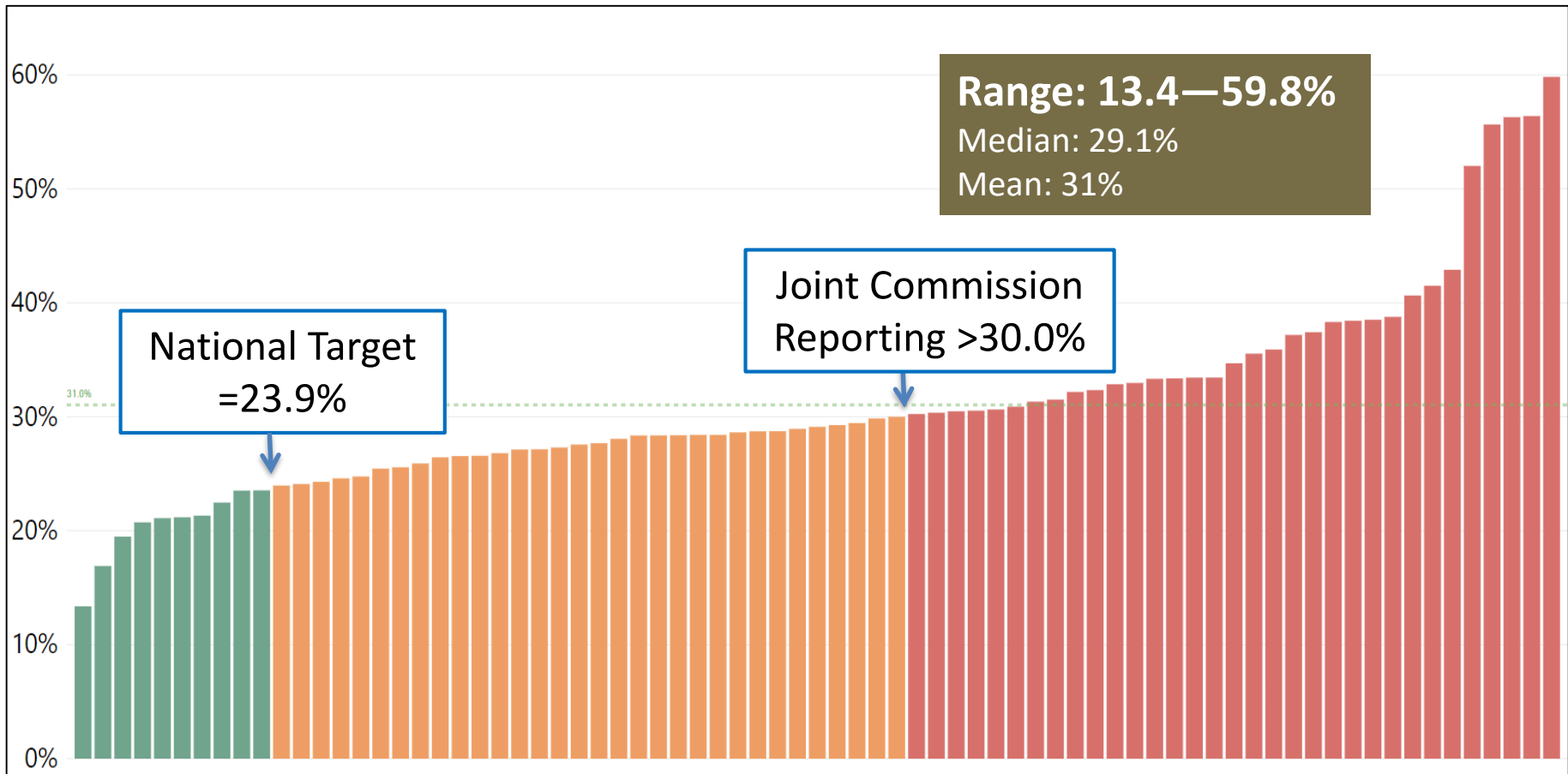
Partnering to Improve Health Care Quality
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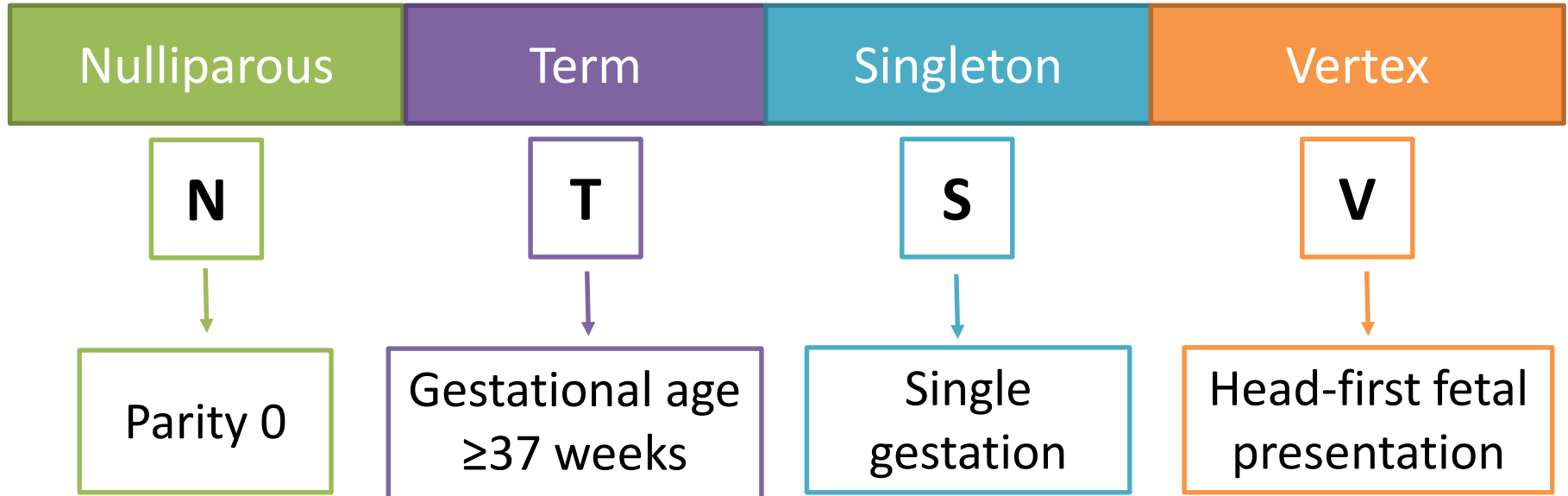
Low-Risk First-Birth (Nulliparous Term Singleton Vertex) Cesarean Rate, 76 PROVIDE Hospitals



Low-Risk First-Birth (Nulliparous Term Singleton Vertex) Cesarean Rate, 76 PROVIDE Hospitals



Our population



- The most favorable conditions for vaginal birth
- Most difficult labor management
- Largest contributor to the rise in cesarean rates



“Things get done only if the data we gather can inform and inspire those in a position to make a difference.” – [Mike Schmoker](#)

Processes for Data Collection

Data Collection Process

1

**Identify qualifying
NTSV cesarean**

Check Inclusion &
Exclusion Criteria

INCLUSION & EXCLUSION CRITERIA

INCLUDE

NTSV cesareans
per TJC

EXCLUDE

Birth weight \geq 4250g

Medical Indication for
Cesarean



1. Maternal or fetal hemorrhage
2. Hypertensive emergencies not responding to treatment
3. Abnormalities of placenta or umbilical cord
4. Fetal or maternal conditions that obstruct the pelvis
5. Active HSV lesions or HIV viral load $>$ 1000copies/ml
6. Other maternal medical indications (cardiac, neurological, orthopedic, pulmonary, malignancy, previous uterine surgery) that preclude vaginal delivery

How to identify NTSV cesareans?

INCLUDE

NTSV
cesareans per
TJC



Nulliparous

Term

**CHART
REVIEW
(EHR)**

Cesarean

Singleton

Vertex

**ICD-10
codes**

How to identify NTSV cesareans?

EXCLUDE

Birth weight \geq 4250g

Medical Indication for Cesarean

CHART REVIEW
(EHR)

AIM curated list
of ICD-10 codes

Data Collection Process

1

Identify qualifying NTSV cesarean

Check Inclusion & Exclusion Criteria

2

Chart Abstraction

FPQC PROVIDE 2.0 Initiative Chart Audit Sheet Study ID #: _____

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Baseline data collection: Hospital to audit up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)
 Prospective data collection: Hospital to audit up to 20 cases per month on 1 (or more if you choose) of the 3 primary indication areas

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other _____	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum	Gestation ____ weeks Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm
--	---	--	--

Only complete the section of the form that corresponds to the c/s category (e.g. if c/s category is "Induction" only complete the "Induction case audit" section; if c/s category is "Other" then only complete the section above.

INDUCTION CASE AUDIT
 Sample of cases that are NTSV per TIC and were induced labor and had a cesarean birth for labor arrest, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: *Fetal heart rate concern or *Medical indication for cesarean section

Dilation at start of induction: _____	Dilation at last exam before c/s: _____	Bishop Score as noted on chart: _____
<input type="checkbox"/> Unknown	<input type="checkbox"/> Unknown	<input type="checkbox"/> Unknown
Was Cervix 6 cm or greater at time of Cesarean? <input type="checkbox"/> If No, go to A. <input type="checkbox"/> If Yes, go to B.		A. If <6 cm, unable to generate regular contractions (every 3 minutes) and cervical change after oxytocin administered for at least 12-18 hours after membrane rupture? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A
Completed labor dystocia checklist by nurse and doctor <input type="checkbox"/> Yes <input type="checkbox"/> No		B. If ≥6cm, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin? <input type="checkbox"/> Yes <input type="checkbox"/> No
Completely dilated at time of Cesarean? <input type="checkbox"/> No <input type="checkbox"/> If Yes →		Were there 3 hours or more of pushing (4 hours with epidural)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown

LABOR DYSTOCIA/FAILURE TO PROGRESS CASE AUDIT
 Sample of cases that are NTSV per TIC and were spontaneous labor and had a cesarean for labor arrest, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: *Fetal heart rate concern or *Medical indication for cesarean section

Dilation at time of admission: _____	Dilation at time of cesarean: _____	Was cervix 6 cm or greater at time of cesarean? <input type="checkbox"/> Yes → <input type="checkbox"/> No
Completely dilated at time of cesarean? <input type="checkbox"/> No <input type="checkbox"/> If Yes →		Were there 3+hrs of pushing (4hrs with epidural)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown
Completed labor dystocia checklist by nurse and doctor <input type="checkbox"/> Yes <input type="checkbox"/> No		If Yes, please check <u>one</u> reason for cesarean that applies: <input type="checkbox"/> Membranes ruptured and no cervical change x4 hrs with adequate uterine activity (e.g., >200 MVU) <input type="checkbox"/> Membranes ruptured, Oxytocin administered, and no cervical change x6hrs with inadequate uterine activity (e.g., <200 MVU) <input type="checkbox"/> None of the above

FETAL HEART RATE CONCERN/INDICATIONS
 Sample of cases that are NTSV per TIC and had a cesarean for fetal heart rate (FHR) concern/indications, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: *Labor arrest / CPD

What was the FHR concern/indication? (Linked with specific corrective and evaluative measures)

<input type="checkbox"/> Antepartum testing results which precluded trial of labor <input type="checkbox"/> Category III FHR tracing <input type="checkbox"/> Category II FHR tracing (Were these specific types present?) <input type="checkbox"/> Clinically significant variable decelerations <input type="checkbox"/> Minimal/absent FHR variability without significant decelerations <input type="checkbox"/> Other concern: _____	Please check all corrective and evaluative measures used: <input type="checkbox"/> Basic resuscitation measures such as: Maternal position change, maternal fluid bolus, and/or administration of O2 <input type="checkbox"/> Reduced or stopped oxytocin or uterine stimulants <input type="checkbox"/> Used Amnioinfusion with significant variable decelerations after other measures failed <input type="checkbox"/> Elicited stimulation (scalp, vibroacoustic, or abdominal wall) with minimal or absent FHR variability
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Other labor issues:
 Did the mother have uterine tachysystole? Yes No
 Corrected uterine tachysystole: decrease or discontinue uterine stimulants, fluid bolus, terbutaline or nitroglycerin and/or other? Yes No



Chart Audit Sheet



www.fpqc.org

Complete audit form for each NTSV cesarean

What do I do with the Study ID#?

FPQC PROVIDE 2.0 Initiative Chart Audit Sheet

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Study ID #: _____

Baseline data collection: Hospital to audit up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)

Prospective data collection: Hospital to audit up to 20 cases per month on 1 (or more if you choose) of the 3 primary indication areas

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other (specify) _____	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum	Gestation ____ weeks	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm
		Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured	

- 👤 Study ID # : Start at 001 and add sequentially
- 👤 Every patient chart that you include for PROVIDE data submission should get a hospital assigned Study ID number

Keep an on-site log of the study ID number and the medical chart number and/or identifiable patient information for data verification

Which C/S Category to Choose?

FPQC PROVIDE 2.0 Initiative Chart Audit Sheet

Study ID #: ____

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Baseline data collection: Hospital to audit up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)

Prospective data collection: Hospital to audit up to 20 cases per month on 1 (or more if you choose) of the 3 primary indication areas

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other (specify) _____	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum	Gestation ____ weeks	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm
		Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured	

- 🕒 Select only one C/S category – the primary reason for the cesarean
- 🕒 If there is a cesarean for fetal heart rate concerns, then select “FHR concerns”
- 🕒 If not and mother induced, then select “Induction”
- 🕒 If neither and cesarean for labor dystocia or failure to progress, then select “Labor Dystocia” Otherwise, select “Other”

Which C/S Category to Choose?

FPQC PROVIDE 2.0 Initiative Chart Audit Sheet

Study ID #: _____

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Baseline data collection: Hospital to audit up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)

Prospective data collection: Hospital to audit up to 20 cases per month on 1 (or more if you choose) of the 3 primary indication areas

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other (specify) _____	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum	Gestation ____ weeks	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm
		Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured	

- 👉 “Other” reasons will only be an option during “Baseline”
- 👉 Specify if the cesarean was due to maternal request or the reason for the cesarean (birth defect, suspected macrosomia or trauma)

FPQC PROVIDE Initiative Chart Audit Sheet

Study ID #: _____

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Baseline data collection: Complete form for up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum		Gestation ____ weeks	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm		
			Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured			
INDUCTION CASE AUDIT <i>Sample of cases that are NTSV per TJC and were induced labor and had a cesarean birth for labor arrest, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: •Fetal heart rate concern or •Medical indication for cesarean section</i>						
Event	Dilation	Effacement	Station	Cervix Position	Cervix consistency	Bishop Score as noted on chart
At Start of Induction	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown
Last Exam before Delivery	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown
Was Cervix 6 cm or greater at time of Cesarean? <input type="checkbox"/> If No, go to A. <input type="checkbox"/> If Yes, go to B. <input type="checkbox"/> Unknown			A. If <6 cm, unable to generate regular contractions (every 3 minutes) and cervical change after oxytocin administered for at least 12-18 hours after membrane rupture?		<input type="checkbox"/> Yes <input type="checkbox"/> No	
If Bishop score ≤ 8 at start of induction, was cervical ripening used?		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	B. If ≥6cm, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin?		<input type="checkbox"/> Yes <input type="checkbox"/> No	
Completely dilated at time of Cesarean? No If Yes →		Were there 3 hours or more of pushing (4 hours with epidural)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown				
LABOR DYSTOCIA/FAILURE TO PROGRESS CASE AUDIT <i>Sample of cases that are NTSV per TJC and were spontaneous labor and had a cesarean for labor arrest, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: •Fetal heart rate concern or •Medical indication for cesarean section</i>						
Dilation at time of admission: _____ <input type="checkbox"/> Unknown		Was Cervix 6 cm or greater at time of Cesarean? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown	If Yes, please check the <u>one</u> reason for cesarean that applies:			
Dilation at time of cesarean: _____ <input type="checkbox"/> unknown			<input type="checkbox"/> Membranes ruptured and No cervical change x 4 hrs with Adequate Uterine activity (e.g., > 200 MVU) <input type="checkbox"/> Membranes ruptured, Oxytocin administered, and No cervical change x 6 hrs with Inadequate Uterine activity (e.g., < 200 MVU) <input type="checkbox"/> None of the above			
Completely dilated at time of Cesarean? No If Yes →		Were there 3 hours or more of pushing (4 hours with epidural)? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown				
FETAL HEART RATE CONCERN/INDICATIONS <i>Sample of cases that are NTSV per TJC and had a cesarean for fetal heart rate (FHR) concern/indications, excluding those with birth weight ≥ 4250g or with ICD-10 codes for: •Labor arrest / CPD</i>						
What was the FHR concern/indication? (Linked with specific corrective and evaluative measures) <input type="checkbox"/> Antepartum testing results which precluded trial of labor <input type="checkbox"/> Category III FHR tracing <input type="checkbox"/> Category II FHR tracing (Were these specific types present?) <input type="checkbox"/> Clinically significant variable decelerations <input type="checkbox"/> Minimal/absent FHR variability without significant decelerations <input type="checkbox"/> Other concern: _____			Please check all corrective and evaluative measures used: <input type="checkbox"/> Basic resuscitation measures such as: Maternal position change, maternal fluid bolus, and/or administration of O2 <input type="checkbox"/> Reduced or stopped oxytocin or uterine stimulants <input type="checkbox"/> Used Amnioinfusion with significant variable decelerations after other measures failed <input type="checkbox"/> Elicited stimulation (scalp, vibroacoustic, or abdominal wall) with minimal or absent FHR variability			
Other labor issues: Did the mother have uterine tachysystole? <input type="checkbox"/> Yes <input type="checkbox"/> No			Corrected uterine tachysystole: decrease or discontinue uterine stimulants, fluid bolus, terbutaline or nitroglycerin and/or other? <input type="checkbox"/> Yes <input type="checkbox"/> No			

Which section to complete?

- You only need to answer questions related to the C/S Category selected after the first (white) section.

Definitions and Clinical Criteria

NTSV = ≥37 weeks, parity 0, single gestation pregnancy, vertex fetal presentation

Study ID = Begins with 001 & numbers the patient charts consecutively. On site log, record patient's medical record number or identifying number next to the corresponding Study ID# to keep track and return for any needed case review.

CS Category = If the cesarean delivery has fetal heart rate concerns requiring delivery, then label "FHR Concerns." If not and had an induction, then "Induction." If neither of these and had labor dystocia, then "Labor Dystocia." Otherwise, mark the form as "Other."

Medical Indication for Cesarean (*chart review exclusion criteria, or "Other"*) include:

1. Maternal or fetal hemorrhage
2. Hypertensive emergencies not responding to treatment
3. Abnormalities of placenta or umbilical cord
4. Fetal or maternal conditions that obstruct the pelvis
5. Active HSV lesions or HIV viral load >1000copies/ml
6. Other maternal medical indications (cardiac, neurological, orthopedic, pulmonary, malignancy, previous uterine surgery) that preclude vaginal delivery

Primary Indication for NTSV Cesarean	Fall out if these not met:	Reference
Labor Dystocia/Failure to Progress	Chart Review: looking for Yes answers to the following (a no answer would indicate inconsistency with the ACOG guidelines): <ul style="list-style-type: none"> • If <6cm dilated, automatic fallout • If 6-10cm dilated, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin? • If completely dilated, was there 3h or more of active pushing (4h with epidural)? 	ACOG/SMFM criteria (Ob Gyn 2014;123:693-711) -CMQCC
Induction	Chart Review: looking for Yes answers to the following (a no answer would indicate inconsistency with the ACOG guidelines): <ul style="list-style-type: none"> • If <6cm dilated, were there at least 12 hours of oxytocin after rupture of membranes? • If 6-10cm dilated, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin? (identical to the question for Labor arrest/CPD above) • If completely dilated, was there 3h or more of active pushing (4h with epidural)? 	ACOG/SMFM criteria (Ob Gyn 2014; 123:693-711) CMQCC
Fetal Heart Rate Concern	Cesarean deliveries performed for "fetal heart rate concern" using listed resuscitation techniques listed below based on the FPQC FHR Concern algorithm: <ul style="list-style-type: none"> • Antepartum testing which preclude labor: no techniques required. • All Cat. II and III FHR concerns should use some techniques listed under "any intrauterine resuscitation efforts." • Category Cat. II FHR concerns should also use additional techniques if the following: <ul style="list-style-type: none"> ○ Receiving oxytocin—reduced or stopped oxytocin ○ Clinically significant variable decelerations—possibly Amnioinfusion (not required) ○ Minimal/absent variability—elicited stimulation if no significant decelerations ○ Uterine tachysystole—any combination listed to correct 	Spong et al (Ob Gyn 2012; 120:1181-93) Clark et al (AJOG 2013; 209:89-97) ACOG/SMFM criteria (Ob Gyn 2014; 123:693-711) CMQCC FPQC

How to Calculate a Bishop Score:

Cervical Exam	Points				SUBSCORE
	0	1	2	3	
Dilation	Closed	1-2 cm	3-4 cm	≥5 cm	
Effacement	0-30%	31-50%	51-80%	≥80%	
Station	-3	-2	-1, 0	+1, +2	
Consistency	Firm	Medium	Soft		
Position	Posterior	Mid	Anterior		
Bishop's Score =					

See back of the chart audit sheet for helpful definitions

Inductions?

- 👤 No ACOG definition for “failed induction”
- 👤 We have removed the term “failed.”
- 👤 Abstractors do not need to determine whether a failed induction or not.
- 👤 If there are no fetal heart rate concerns and there is an induction, choose “Induction.”
- 👤 Abstractors will only assess whether ACOG labor duration guidelines were followed or not.

Induction Case Audit

INDUCTION CASE AUDIT

Sample of cases that are NTSV per TJC and were induced labor and had a cesarean birth for labor arrest, excluding those with birth weight $\geq 4250g$ or with ICD-10 codes for: •Fetal heart rate concern or •Medical indication for cesarean section

Event	Dilation	Effacement	Station	Cervix Position	Cervix consistency	Bishop Score as noted on chart
At Start of Induction	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown
Last Exam before Delivery	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	_____ <input type="checkbox"/> unknown	
Was Cervix 6 cm or greater at time of Cesarean? <input type="checkbox"/> If No, go to A. <input type="checkbox"/> If Yes, go to B. <input type="checkbox"/> Unknown			A. If <6 cm, unable to generate regular contractions (every 3 minutes) and cervical change after oxytocin administered for at least 12-18 hours after membrane rupture?			<input type="checkbox"/> Yes <input type="checkbox"/> No
If Bishop score ≤ 8 at start of induction, was cervical ripening used?		<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A	B. If ≥ 6 cm, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin?			<input type="checkbox"/> Yes <input type="checkbox"/> No
Completely dilated at time of Cesarean? No If Yes →	Were there 3 hours or more of pushing (4 hours with epidural)?					<input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown

Labor Dystocia/Failure to Progress Case Audit

<p>Was cervix 6 cm or greater at time of cesarean? <input type="checkbox"/> Yes → <input type="checkbox"/> No</p>	<p>If Yes, please check <u>one</u> reason for cesarean that applies:</p> <p><input type="checkbox"/> Membranes ruptured and no cervical change x4 hrs with adequate uterine activity (e.g., >200 MVU)</p> <p><input type="checkbox"/> Membranes ruptured, Oxytocin administered, and no cervical change x6hrs with inadequate uterine activity (e.g., <200 MVU)</p> <p><input type="checkbox"/> None of the above</p>
<p>Were there 3+hrs of pushing (4hrs with epidural?) <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> Unknown</p>	
<p>urse and doctor <input type="checkbox"/> Yes <input type="checkbox"/> No</p>	

Fetal Heart Rate Concern Case Audit

FETAL HEART RATE CONCERN/INDICATIONS

Sample of cases that are NTSV per TJC and had a cesarean for fetal heart rate (FHR) concern/indications, excluding those with birth weight $\geq 4250g$ or with ICD-10 codes for: •Labor arrest / CPD

What was the FHR concern/indication? (Linked with specific corrective and evaluative measures)

- Antepartum testing results which precluded trial of labor
- Category III FHR tracing
- Category II FHR tracing (Were these specific types present?)
 - Clinically significant variable decelerations
 - Minimal/absent FHR variability without significant decelerations
- Other concern: _____

Please check all corrective and evaluative measures used:

- Basic resuscitation measures such as: Maternal position change, maternal fluid bolus, and/or administration of O2
- Reduced or stopped oxytocin or uterine stimulants
- Used Amnioinfusion with significant variable decelerations after other measures failed
- Elicited stimulation (scalp, vibroacoustic, or abdominal wall) with minimal or absent FHR variability

Other labor issues:

Did the mother have uterine tachysystole? Yes No

Corrected uterine tachysystole: decrease or discontinue uterine stimulants, fluid bolus, terbutaline or nitroglycerin and/or other? Yes No

Data Collection Process

1

Identify qualifying NTSV cesarean

Check Inclusion & Exclusion Criteria

2

Chart Abstraction

FPQC PROVIDE Initiative Chart Audit Sheet Study ID #: _____

Complete only for Nulliparous Term Singleton Vertex Cesarean Sections

Baseline data collection: Complete form for up to 20 NTSV C-sections per month for 3 months to determine hospital's main focus area(s)

C/S Category <input type="checkbox"/> Induction <input type="checkbox"/> Labor Dystocia <input type="checkbox"/> FHR Concerns <input type="checkbox"/> Other	Patient Status: <input type="checkbox"/> Admitted already in labor <input type="checkbox"/> Induced <input type="checkbox"/> Indicated augmented labor <input type="checkbox"/> Not in labor: spontaneous rupture of membranes <input type="checkbox"/> Previously admitted antepartum	Gestation ____ weeks Membranes on Admission <input type="checkbox"/> Intact <input type="checkbox"/> Ruptured	Oxytocin <input type="checkbox"/> None utilized <input type="checkbox"/> Induction <input type="checkbox"/> Augmentation at ____ cm
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INDUCTION CASE AUDIT
Sample of cases that are NTSV per TIC and were induced labor and had a cesarean birth for labor arrest, excluding those with birth weight 2-4250g or with ICD-10 codes for: *Fetal heart rate concern or *Medical indication for cesarean section

Event	Dilation	Effacement	Station	Cervix Position	Cervix consistency	Bishop Score as noted on chart
At Start of Induction	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown
Last Exam before Delivery	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown	<input type="checkbox"/> unknown

Was Cervix 6 cm or greater at time of Cesarean?
 If No, go to A.
 If Yes, go to B. Unknown

A. If <6 cm, unable to generate regular contractions (every 3 minutes) and cervical change after oxytocin administered for at least 12-18 hours after membrane rupture? Yes No

B. If ≥6cm, was there at least 4h with adequate uterine activity or at least 6h with inadequate uterine activity and with oxytocin? Yes No

If Bishop score ≤ 8 at start of induction, was cervical ripening used? Yes No N/A

Completely dilated at time of Cesarean? No If Yes → Were there 3 hours or more of pushing (4 hours with epidural)? Yes No unknown

LABOR DYSTOCIA/FAILURE TO PROGRESS CASE AUDIT
Sample of cases that are NTSV per TIC and were spontaneous labor and had a cesarean for labor arrest, excluding those with birth weight 2-4250g or with ICD-10 codes for: *Fetal heart rate concern or *Medical indication for cesarean section

Dilation at time of admission: <input type="checkbox"/> unknown	Dilation at time of cesarean: <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> unknown	Was Cervix 6 cm or greater at time of Cesarean? <input type="checkbox"/> Yes <input type="checkbox"/> No	If Yes, please check the one reason for cesarean that applies: <input type="checkbox"/> Membranes ruptured and No cervical change x 4 hrs with Adequate Uterine activity (e.g., > 200 MVU) <input type="checkbox"/> Membranes ruptured, Oxytocin administered, and No cervical change x 6 hrs with inadequate Uterine activity (e.g., < 200 MVU) <input type="checkbox"/> None of the above
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Completely dilated at time of Cesarean? No If Yes → Were there 3 hours or more of pushing (4 hours with epidural)? Yes No unknown

FETAL HEART RATE CONCERN/INDICATIONS
Sample of cases that are NTSV per TIC and had a cesarean for fetal heart rate (FHR) concern/indications, excluding those with birth weight 2-4250g or with ICD-10 codes for: *Labor arrest / CSO

What was the FHR concern/indication? (Linked with specific corrective and evaluative measures)
 Antepartum testing results which precluded trial of labor
 Category III FHR tracing
 Category II FHR tracing (Were these specific types present?)
 Clinically significant variable decelerations
 Minimal/absent FHR variability without significant decelerations
 Other concern: _____

Please check all corrective and evaluative measures used:
 Basic resuscitation measures such as: Maternal position change, maternal fluid bolus, and/or administration of O2
 Reduced or stopped oxytocin or uterine stimulants
 Used Amnioinfusion with significant variable decelerations after other measures failed
 Elicited stimulation (scalp, vibroacoustic, or abdominal wall) with minimal or absent FHR variability

Other labor issues:
 Corrected uterine tachysystole: decrease or discontinue uterine stimulants, fluid bolus, terbutaline or nitroglycerin and/or other? Yes No
 Did the mother have uterine tachysystole? Yes No

3

Enter data in the REDCap data portal

redcap.health.usf.edu

Florida Perinatal Quality Collaborative
Partnering to Improve Health Care Quality for Mothers and Babies

Promoting Primary Vaginal Delivery (PROVIDE) Data Entry Form

Baseline data collection: Please complete the PROVIDE Data Entry Form for up to 20 NTSV C-sections per month, for 3 months to determine your hospital's main focus area(s).

Prospective data collection: Please complete the PROVIDE Data Entry Form for up to 20 NTSV C-sections per month on 1 (or more if you choose) of the 3 primary indication areas.

Thank you for your commitment to and participation in the PROVIDE project.

You are submitting for the month of: Please select the month for which you are submitting data

You are submitting for the year of: Please select the year for which you are submitting data

Please select the hospital you are reporting for:

Study ID # Start with 001 and add consecutively for each mother

Demographics

Patient Status
 Admitted already in labor (not induced)
 Induced
 Indicated augmented labor
 Not in labor: Spontaneous rupture of membranes
 Previously admitted antepartum
 None of the above

Infant's Gestational Age
 Infant's gestational age can only be between 37 and 42 weeks

Data Submission

- 👤 Each hospital lead will receive the REDCap link for data submission
- 👤 Bookmark this link, you will use the same link throughout the initiative to submit your data



Baseline Data Collection

- 👤 Collect baseline data for **July, August & September 2019**
- 👤 Audit up to 20 NTSV cesareans per month for **all reasons** for a total of up to 60 charts
- 👤 Audit **the first 20 of each month** so your audit is not biased

Data Submission

- 👤 Each hospital lead will receive the REDCap hyperlink for data submission
- 👤 Enter forms into the online PROVIDE data portal (REDCap) **by Dec 1st**
- 👤 You are guaranteed to receive your hospital's report before the end of December if you submit by this deadline

What Next?

- You will receive a Baseline data report that includes:
 - NTSV cesarean rate report from January 2018 to September 2019
 - Baseline audit percentages for induction, labor dystocia, & fetal heart rate
- Attend the Webinar on **January 8th at 12:00 pm** - we will help you understand your baseline report
- Use your baseline report to choose your hospital focus area (Induction, Labor Dystocia or FHRC)



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QUESTIONS?



Prospective Data

- After choosing 1 (or more) focus area(s)
- Complete audit form for up to 20 NTSV C-sections per month for each focus area(s) you have chosen
- Follow the same REDCap hyperlink
- Complete the REDCap form in the same manner as Baseline
- “Other” will not be an option for prospective data

Baseline vs Prospective data

	BASELINE	PROSPECTIVE
Data collection	July-September 2019	Starts January 1st, 2020
Collect on	all NTSV cesareans up to 20/month	All NTSV cesareans in your hospital's focus area up to 20/month
Data submission due	December 1st, 2019	The 15th of the following month (e.g. January data is due February 15th)
Report due	December 30th, 2019	The 30th of the following month (e.g. receive January report by February 30th)

Structural Measures

Collected Every Month by Survey

- 🌀 A link will be sent to the project lead
- 🌀 Report on:
 - Labor guidelines, policy & procedures
 - EHR Integration
 - Multidisciplinary Case Review
 - Staff Education on ACOG/SMFM labor management guidelines and techniques to promote vaginal birth (Providers, Nurses)

Engage your Birth Certificate Clerks/Leads

Hospital Reporting of Delivery Attendant NTSV Cesarean Rates

“Sharing provider NTSV cesarean rates was one of the most important component in reducing our cesarean rates.” Elliott Main, CMQCC

Sample Report

NTSV CESAREAN DELIVERY BY ATTENDANT REPORT-2018 (Q3)

Hospital Code	X	Year 2018(Jan-Oct)			2018 Q3(Aug, Sep, Oct)				
<i>RandomID</i>	<i>Attendant Name</i>	<i>Running Yearly NTSV Ces</i>	<i>Running Yearly NTSV Births</i>	<i>Running Yearly Cesarean Rate</i>	<i>Quarterly NTSV Cesarean Total</i>	<i>Quarterly NTSV Births Total</i>	<i>Quarterly NTSV Cesarean Rate</i>	<i>Rank</i>	<i>Attendant Type</i>
4231	Van De Waal	11	37	28.5%	16	17	94.1%	1	
4213	Latif Abdul	36	43	82.7%	20	23	87.0%	2	
3241	Vasha Patel	31	39	79.6%	19	25	76.0%	3	
3124	Li Mark	19	27	71.1%	13	18	72.2%	4	
2341	Shanice Four	17	23	71.9%	10	15	66.7%	5	
2143	Jose Miguel	13	27	47.4%	15	27	55.6%	6	
1234	John Smith	10	22	45.8%	10	20	50.0%	7	
2314	Jose Martinez	18	37	47.4%	12	24	50.0%	8	
1423	Rose Jones	29	56	51.9%	23	47	48.9%	9	
1432	Rosa Three	19	45	42.1%	11	23	47.8%	10	
3421	James Junior	10	23	45.1%	11	23	47.8%	11	
4132	Jane McJonas	14	34	42.5%	12	28	42.9%	13	
2431	John Peery	15	43	35.5%	10	27	37.0%	14	
3412	Alex Peters	10	30	34.5%	12	34	35.3%	15	
1342	Tom Jones	12	38	32.5%	10	30	33.3%	16	
4123	Alex III Mark	2	9	23.7%	6	18	33.3%	17	
2134	Pierre Thomas	1	7	14.0%	2	11	18.2%	21	
4321	Okoye Oba	2	18	13.0%	3	23	13.0%	23	
3214	Vasha Peters	1	10	8.5%	2	23	8.7%	24	
1324	Sam Three	0	3	5.2%	0	13	0.0%	25	

Hospital Reporting of Delivery Attendant NTSV Cesarean Rates

Special Opportunity:

- 👤 DOH will train birth certificate leads and clean dataset
- 👤 Hospitals receive monthly named/coded attendant reports

Eligibility:

- 👤 Participate in required hospital report user training
- 👤 Provide feedback and suggestions to the FPQC

PROVIDE—Accuracy of Birth Certificate Data

- ➊ Number now living or dead
- ➋ Induction of labor
- ➌ Fetal presentation at birth
- ➍ Final route and delivery method
- ➎ Obstetric estimate of gestation
- ➏ Plurality
- ➐ Apgar Score

New Completion Guide



PROMOTING PRIMARY VAGINAL DELIVERIES (PROVIDE)

Completion Guide for Key Birth Certificate Data Reporting

The variables included in this manual are required to calculate several measures for PROVIDE. Please review this manual and collaborate with your teams and data abstractors to improve the completeness and accuracy of these birth certificate variables.

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Adapted from: - the NICHD, "Guide to Completing the Facility Worksheets for the Certificate of Live Birth and Report of Fetal Death" (2016 Revision)
- the Florida DDM "Electronic Birth Registration System Manual & 2014 Registration Handbook" (2014 Revision)

SECTION	ITEM	SUBITEM	SOURCES
Pregnancy History	Previous Live Births		1 st Prenatal care record 2 nd Labor and delivery nursing admission triage form 3 rd Admission history and physical (H&P)
		Number Now Dead	1 st Prenatal care record 2 nd Admission history and physical (H&P)
Medical and Health Information	Characteristics of Labor and Delivery	Induction of labor	1 st Delivery record 2 nd Physician progress note 3 rd Labor and delivery nursing admission triage form
		Augmentation of labor	1 st Delivery record under: 2 nd Physician progress note
	Method of Delivery	Fetal presentation at birth: - Cephalic - Breech - Other	1 st Delivery record
		Final Route and method of delivery - Vaginal/Spontaneous - Vaginal/Forceps - Vaginal/Vacuum - Cesarean	1 st Delivery record under 2 nd Newborn admission H&P 3 rd Recovery room record

DEFINITION	BC ITEM #	TIPS FOR ENTRY	KEYWORDS AND ABBREVIATIONS	NCHS RECOMMENDED SOURCE
1. Previous Live Birth				
<p>a. Number now living- total number of previous live-born infants who are still living.</p>	# 42a	<p><u>DO NOT include this child</u></p> <p>Do not include abortions (spontaneous miscarriages or therapeutic or elective abortions), fetal deaths/stillbirths.</p> <p>For multiple deliveries: Include all live-born infants before this infant in this pregnancy. If the first born, do not include this infant. If the second born, include the first born, etc. If no previous live-born infant now alive enter 00.</p>	<p>L–Now living G–Gravida–Total number of pregnancies P–Para–Previous live births and fetal deaths > 28 weeks of gestation T–Term–Delivered at 37 to 40 weeks gestation</p>	<p>1st Prenatal care record under:</p> <ul style="list-style-type: none"> - Intake information - Gravida section–L (living)–last number in series - Para section–L–last number in series - Pregnancy history information - Previous OB history - Past pregnancy history <p>2nd Labor and delivery nursing admission triage form under:</p> <ul style="list-style-type: none"> - Patient data <p>3rd Admission history and physical-H&P</p>

Thank you!

👉 Please don't hesitate to contact us if you have questions fpqc@health.usf.edu or

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