

Randomized Control Trial of Postpartum Visits at 2 and 6 Weeks

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Background

Background

- Postpartum care is an integral transition from pregnancy to well-women care
- More than half of maternal deaths occur postpartum¹, with suicide and self harm as some leading causes²
- Average postpartum (PP) visit attendance nationwide 60-90%³
 - Lower among minorities, younger women, higher parity, limited prenatal care

Background

Interventions studied to increase PP attendance

- Incentives
- Home visits
- Group prenatal care
- Patient education
- Patient reminders
- Scheduling visit upon hospital discharge
- Timing of visits?

Table 2 Antenatal interventions to increase attendance at postpartum follow-up visit (PPV)

Author, year	Target population	Population size	Intervention type	Results	Study design
Stevens-Simon et al. 1994 [41]	Low-income adolescents in Colorado	240	Incentives	Women given incentive of infant carrier at PPV more likely to attend PPV compared to control group (82.4 and 65.2 %, respectively; $p = .003$)	Randomized Controlled Evaluation
Laken and Ager, 1995 [30]	Medicaid-eligible women in Michigan	205	Incentives	No significant difference in PPV attendance (55 % overall) between control groups, women who received \$5.00 gift card or entrance in \$100 raffle for attending visit	Randomized Controlled Evaluation
<i>Patient incentives: inconclusive findings, possible increase in attendance for adolescents</i>					
Belizan et al. 1995 [5]	Socially high-risk women in Latin America	2235	Home visits, support services	No significant difference in PPV attendance between control group and women who received intervention (32.8 and 36.8 %, respectively)	Randomized Controlled Evaluation
Bensusen-Walls and Saewyc, 2001 [6]	Adolescents in Washington State	106	Teen-centered prenatal care	Adolescents in intervention group more likely to attend PPV than adolescents who received traditional prenatal care (70 and 77.8 % for adolescent clinics and <33.3 and 44 % for adult, $p < .05$)	Retrospective Matched Evaluation
Grady and Bloom, 2004 [19]	Adolescents in St. Louis, Missouri	124	Group prenatal care	87 % of adolescents in group prenatal care (CenteringPregnancy Model) returned for PPV	Descriptive Evaluation
Tandon et al. 2013 [43]	Hispanic women in Palm Beach County, Florida	176	Group prenatal care	Women in group prenatal care more likely to attend PPV than comparison group (99 and 94 %, respectively; $p = .04$)	Quasi-Experimental Evaluation
Trudnak et al. 2013 [44]	Spanish-speaking and Hispanic women	487	Group prenatal care	Women in group prenatal care had increased odds of attending PPV compared to women in traditional prenatal care (86.7 and 74.6 %, respectively; $aOR^* = 2.20 [1.20-4.05]$)	Retrospective Cohort Evaluation
Meghea et al. 2013 [34]	Medicaid-eligible women in Michigan	32,088	Home visits, support services	Women who received home visits through the Maternal and Infant Health Program were significantly more likely to attend their postpartum visit than their matched counter parts (OR = 1.50 [1.43, 1.57], $p < .05$)	Quasi-Experimental Evaluation
<i>Enhanced prenatal care interventions: inconclusive findings, possible increase in attendance for group prenatal care</i>					
Jones and Mondy, 1990 [26]	Low-income adolescents in Texas	399	Incentives, patient education	Women in high-treatment group (≥ 8 lessons) more likely to attend PPV compared to low-treatment group (< 8 lessons) (87 and 73 %, respectively; $p < .011$) and comparison group (87 and 71 %, respectively; $p < .002$)	Quasi-Experimental Evaluation

Timing of Postpartum Visits

- Worldwide guidelines
- Recommendations for timing and frequency of PP visits varies from 48 hours to 8 weeks postpartum
 - Based on “available literature, expert opinion, clinical practice”

Table 1 Guidelines for timing and frequency of postpartum visit (PPV)

Organization	Year	Guidelines development process	Guidelines for timing and frequency of PPV	Evidence for timing and frequency of PPV guideline	Quality of evidence as provided by guideline authors
Department of Health, Reproductive Care Program of Nova Scotia, Canada [34]	2002	A postpartum services review working group gathered information from literature review, clinical practice, current statistics and used information to develop guidelines then approved by an action group at the Reproductive Care Program of Nova Scotia and a program delivery group at the Department of Health	Follow up visit: 6 weeks postpartum <i>Follow-up care can be added to based on family care plan</i>	Available literature, expert opinion, and clinical practice	Specific quality level not stated
National Institute for Health and Care Excellence, UK [15]	2006	A Technical Team gathered information from economic and clinical databases for review by a Guideline Development Group. Final recommendations agreed upon by Group following comments from stakeholders	First visit: within 1st week postpartum Remaining visit(s): 2-8 weeks postpartum <i>Care should be individualized for each woman and determined in the antenatal period</i>	Available literature and expert opinion/panel	Recommendation based only on experiences of Guideline Development Group
American College of Obstetric and Gynecology [7]	2012	Developed by AAP Committee on Fetus and Newborn and the ACOG Committee on Obstetric Practice based on most up-to-date scientific information, clinical practice, and expert opinion	Follow up visit: 4-6 weeks postpartum <i>Interval can be modified by needs of patient</i>	Available literature, expert opinion, and clinical practice	Specific quality level not stated
Michigan Quality Improvement Consortium, USA [35]	2012	Evidence gathered through literature review and evaluated by type of study used to establish guidelines to be reviewed by committee until group consensus reached	Follow up visit: 3-8 weeks postpartum	Available literature and expert opinion/panel	Specific quality level not stated
Institute for Clinical Systems Improvement, USA [1]	2012	Evidence gathered through literature review evaluated using GRADE methodology used to establish set of guidelines reviewed and approved by a series of medical committees and relevant stakeholders	Follow up visit: 4-6 weeks postpartum <i>Discussion of postpartum care initiated in antenatal period</i>	Available literature and expert opinion/panel	Specific quality level not stated
Association of Reproductive Health Professionals, USA [7]	2013	Based on expert opinion and available literature	Follow up visit: 4-6 weeks postpartum	Not provided	Specific quality level not stated
World Health Organization [47]	2013	Guideline Development Group reviewed and evaluated evidence gathered through systematic reviews using GRADE profile and analysis of benefits, risks, and costs of implementation. Findings were used to draft recommendations by a WHO steering group and finalized through group consensus/vote	First Visit: 48-72 h postpartum Second Visit: 7-14 days postpartum <i>A minimum of three visits</i>	Available literature and expert opinion/panel	Low quality evidence
French College of Gynecologists and Obstetricians [51]	2016	A steering committee established research questions and assigned experts to conduct literature reviews related to these questions. This information was then used by the steering committee to develop guidelines	Follow up visit: 6-8 weeks postpartum	Not provided	Specific quality level not stated

Background

- ACOG highlights importance of “fourth trimester”
 - Timing of 6-week visit is arbitrary and should be individualized and woman centric
 - “all women should ideally have contact with a maternal care provider within the first three weeks postpartum...ongoing care as needed...comprehensive postpartum visit no later than 12 weeks after birth”
- Derived on expert opinion



The American College of
Obstetricians and Gynecologists
WOMEN'S HEALTH CARE PHYSICIANS

Postpartum Emergency Department Usage

Are patients utilizing the ED instead of attending clinic?

- One in twenty women will use the ED within 6 weeks postpartum¹
- Predictors of ED usage
 - Public insurance, young age, cesarean delivery, severe maternal morbidity, antepartum complications² and mood disorders³

Objective

- To determine if shortening the time to initial postpartum visit from six weeks to two weeks can increase clinic visit attendance and decrease usage of the emergency department

Methods

Methods – Trial Design

PUnCTuAL: Postpartum Care Timing: A Randomized Trial

- Followed CONSORT Guidelines
- Simple parallel randomization
- Non-blinded
- 1:1 allocation
- Clinical Trials NCT03733405
- Funding: N/A

Methods - Participants

- Study Population

- Publicly insured population at tertiary academic medical center

- Eligibility Criteria

- 18+ years old
- English or Spanish speaking
- 35'0 weeks of gestation
- Planning to continue their intrapartum and postpartum care at UCLA

- Exclusion Criteria

- Cognitive impairment or language barrier that limits ability to provide informed consent

Methods - Interventions



Control: Arm 1

6- week postpartum visit (29-56 days PP)



Intervention: Arm 2

2- week postpartum visit (8-28 days PP)

AND

6- week postpartum visit (29-56 days PP)

Methods - Outcomes

- Primary outcome:
 - Attendance at one or more routine postpartum clinic visits
- Secondary outcome:
 - Emergency room visit within 30 days of delivery
 - Chart review
 - Patient reported on postpartum survey
 - Attendance at non-routine postpartum clinic visit
 - Provider recommended
 - Patient initiated

Methods - Power

- Power calculation:
 - Baseline clinic attendance 70%
 - Alpha 0.05, beta 0.80
 - To increase clinic attendance from 70% to 85%, require 240 patients
 - To account for post-randomization drop-out, goal 250 patients

Methods – Recruitment

- Recruitment and consent documents in English and Spanish
- Simple computer randomization
- Allocation concealment
- No blinding

UCLA RESEARCH STUDY



PUnCTuAL

Postpartum Care Timing: A Randomized Trial

We are looking for pregnant women to participate in a study to determine the best time for a follow up visit after their baby is born.


Speak to any staff member for details.
UCLA OB/GYN Clinic, 1010 Veteran Ave, LA, CA 310-825-7955

Methods - Implementation

- Scheduling
 - Patient responsible for scheduling appointments (routine practice)
- Additional clinic visits continued to be scheduled as indicated by her intrapartum course, designated urgent or “non-routine visit”
 - Provider recommended blood pressure or wound check
 - Patient initiated visit

Methods – Postpartum visits

- Routine history and physical
- Patient survey



Postpartum Patient Survey

1. Did you go to the **emergency department** since having your baby?

- a. No
- b. Yes. UCLA.
- c. Yes. Other hospital. Please briefly explain why and what date.

2. Are you satisfied with your **prenatal** care at this UCLA OBGYN clinic?
(circle one)

Very Much	Somewhat	Undecided	Not Really	Not at All
5	4	3	2	1

3. Are you satisfied with your **postpartum** care (appointments, phone calls, etc) at this UCLA OBGYN clinic? (circle one)

Very Much	Somewhat	Undecided	Not Really	Not at All
5	4	3	2	1

4. When do you think is the **best time** for a woman to come back for her postpartum visit?

- a. Before 2 weeks
- b. Just 2 weeks
- c. Just 6 weeks
- d. Both 2 and 6 weeks
- e. Other: _____

Methods – CoVID-19 adjustments

- Due to CoVID-19 pandemic, after March 16, 2020, postpartum visits were adjusted to telephone visits when available
- Attendance was defined as answering the phone and completing appointment at prescheduled time and date

Methods – Data Analysis

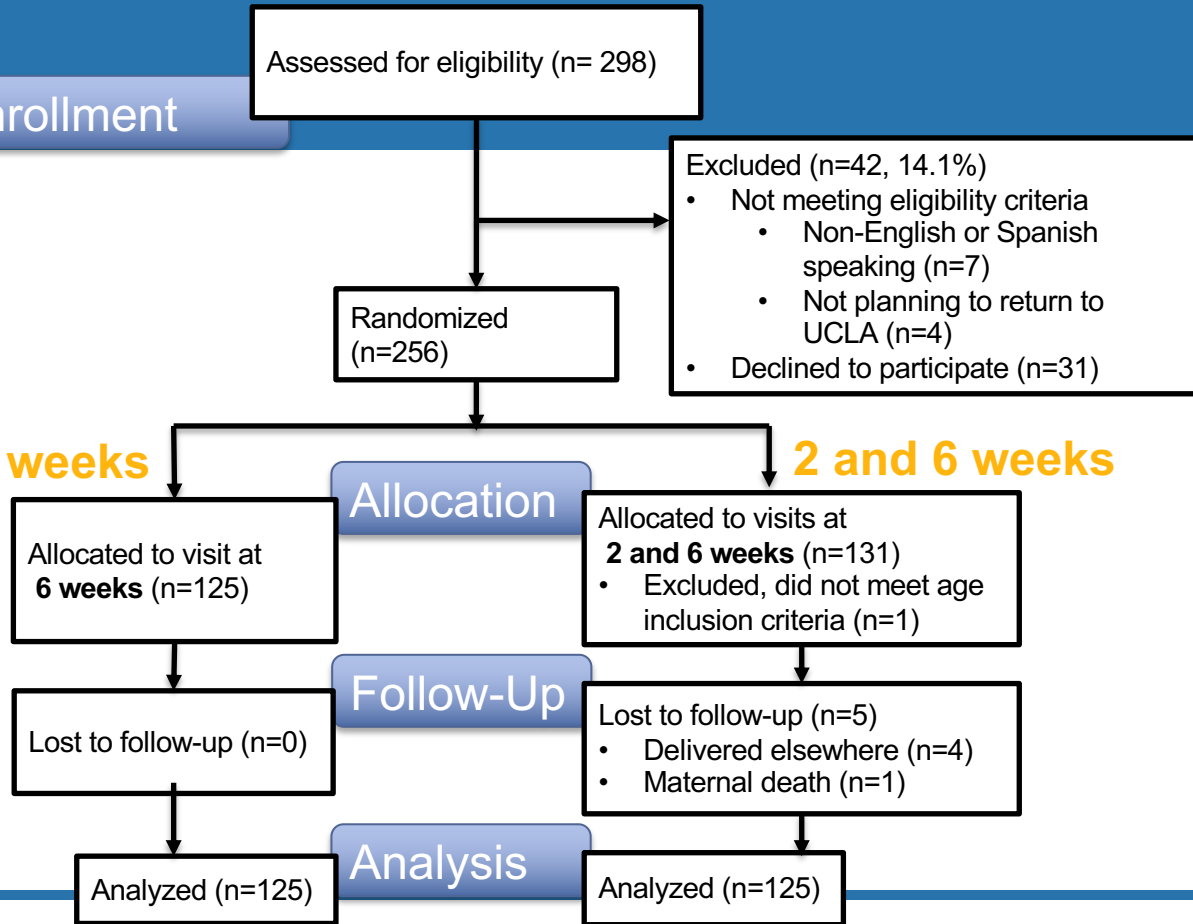
- Chi square or Fisher's exact for categorical variables
- T-test or Wilcoxon rank-sum for continuous variables
- Multivariable logistic regression using backwards elimination method to adjust for confounders of attendance at clinic and ED
- Univariable ROC curves
- Spearman correlation
- Analysis on Stata 15.1 and SAS 9.4

**Participant Flow
Chart Recruitment
Nov 7, 2018 –
Mar 4, 2020**

Enrollment

6 weeks

2 and 6 weeks



Results

Demographics

Variable	Arm 1 (n=125)	Arm 2 (n=125)	P-value
Age	31.0 +/- 6.1	29.8 +/- 5.8	0.117*
Race			
Hispanic	69 (55.2)	71 (56.8)	0.687†
White	17 (13.6)	11 (8.8)	
Asian	11 (8.8)	13 (10.4)	
Black	10 (8.0)	14 (11.2)	
Other	18 (14.4)	16 (12.8)	
Marital status	N=124	N=124	
Single	68 (54.8)	70 (56.4)	0.798†
Married	56 (45.2)	54 (43.6)	
Education	N=89	N=90	
<9 years	3 (3.4)	1 (1.1)	0.434‡
9-11 years	6 (6.7)	10 (11.1)	
12-16 years	58 (65.2)	62 (68.9)	
>16 years	22 (24.7)	17 (18.9)	
Tobacco use	2 (1.6)	5 (4.0)	0.446‡
Domestic violence	N=92	N=93	
	4 (4.4)	8 (8.6)	0.372‡
Obese at consent	69 (55.2)	64 (51.2)	0.526†

No differences in
baseline characteristics
between study groups

Data are mean+/- SD; *t-test; † Chi square test ‡ Fisher's exact

Demographics

Variable	Arm 1 (n=125)	Arm 2 (n=125)	P-value
Nulliparous	49 (39.2)	54 (43.2)	0.521†
Singleton	121 (96.8)	123 (98.4)	0.684‡
Twin	4 (3.2)	2 (1.6)	
Distance >20 miles	15 (12.0)	16 (12.8)	0.848†
Gestational age at intake (weeks)	20.1 (10.1-32.4)	16.7 (10.3-29.0)	0.297 §
Gestational age at randomization (weeks)	36.6 (35.7-37.4)	36.1 (35.6-36.7)	0.020 §
High Risk OB Clinic	50 (40.0)	50 (40.0)	0.999†
Low Risk OB Clinic	75 (60.0)	75 (60.0)	
Maternal Comorbidities			
Diabetes	38 (30.4)	36 (28.8)	0.782†
Mental health disorders	28 (22.4)	28 (22.4)	0.999†
Hypertensive disorders	21 (16.8)	20 (16.0)	0.864†
History of preterm birth	5 (4.0)	5 (4.0)	0.999†
Autoimmune disease	5 (4.0)	4 (3.2)	0.999‡
Cardiac disease	2 (1.6)	4 (3.2)	0.684‡
Renal disease	1 (0.8)	2 (1.6)	0.999‡
Major fetal anomaly	6 (4.8)	7 (5.6)	0.776†
	<i>N=80</i>	<i>N=92</i>	
Antepartum EPDS ≥ 10	18 (22.5)	17 (18.5)	0.513†
Antepartum EPDS ≥ 13	11 (13.8)	12 (13.0)	0.892†

No clinically important differences in baseline characteristics between study groups

Data are n(%), median (25-75% IQR); *t-test; † Chi square test ‡ Fisher's exact; § Wilcoxon rank sum

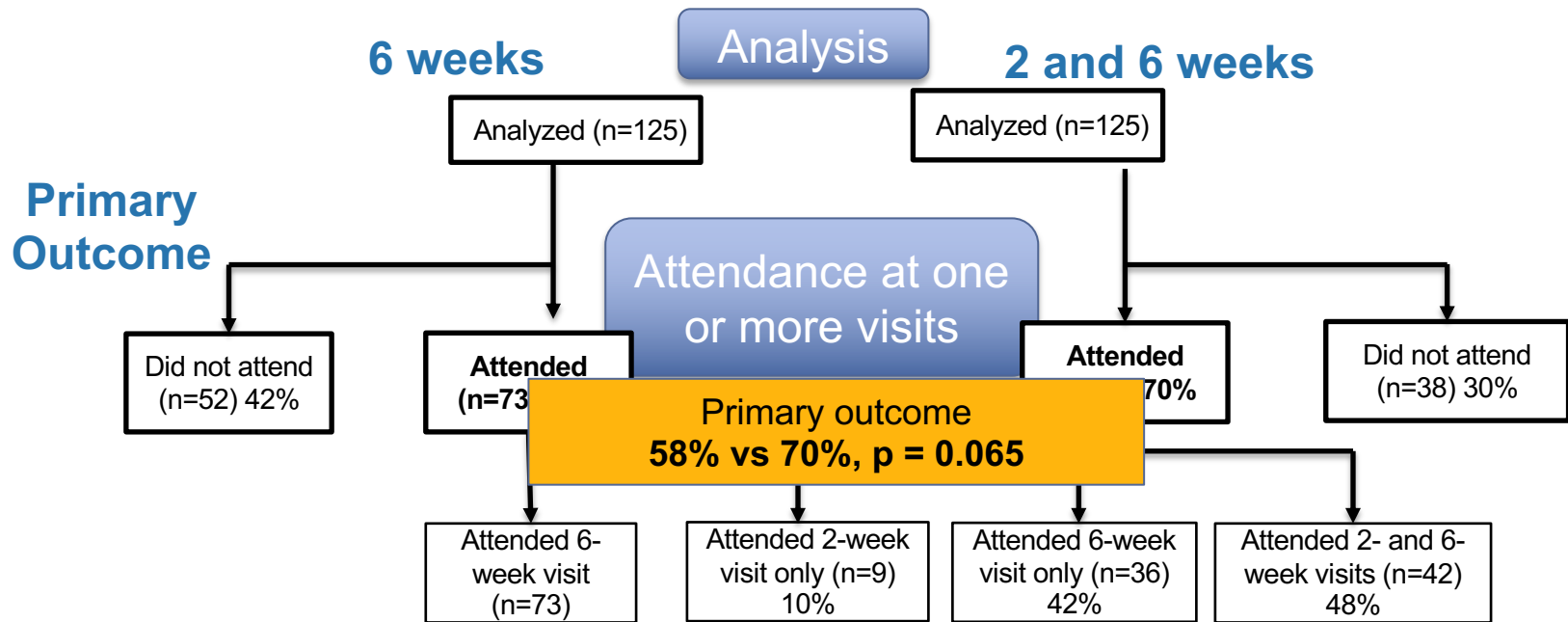
Delivery Characteristics

Variable	Arm 1 (n=125)	Arm 2 (n=125)	P-value
Method of delivery			
SVD	85 (68.0)	85 (68.0)	0.999‡
Cesarean	37 (29.6)	37 (29.6)	
Operative VD	3 (2.4)	3 (2.3)	
Gestational age at delivery (d)	39.1 (38.4-39.9)	39.0 (38.0-39.4)	0.063 §
Prolonged maternal length of stay	14 (11.2)	9 (7.2)	0.274‡
Hypertension without SF	31 (24.8)	37 (29.6)	0.394‡
Postpartum hemorrhage	14 (11.2)	10 (8.0)	0.390‡
Chorioamnionitis	12 (9.6)	11 (8.8)	0.827‡
Shoulder dystocia	3 (2.4)	0 (0)	0.162‡
Composite severe intrapartum complication**	6 (4.8)	9 (7.0)	0.451‡

No differences in delivery characteristics between study groups

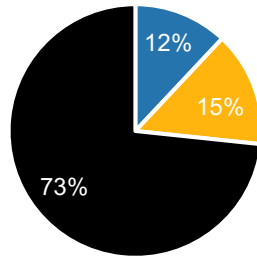
**severe hypertension, wound infection, hemorrhage >2L, Hg <7, blood transfusion, Bakri, foley at discharge, ICU, 3rd or 4th degree laceration, bowel injury, IR, hysterectomy

Primary Outcome



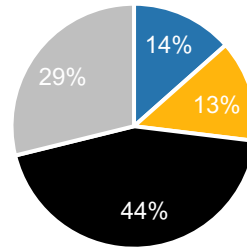
Reasons for Non-Attendance at Clinic Visit

2- week
n=74
Arm 2



- Cancelled
- No show
- Never scheduled
- Outside of time

Arm 1

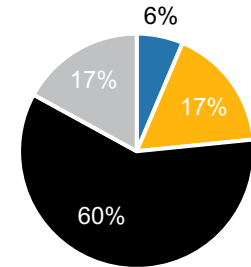


- Cancelled
- No show
- Never scheduled
- Outside of time

6- week
n=99

p=0.35†

Arm 2



- Cancelled
- No show
- Never scheduled
- Outside of time

CoVID-19 Effects on Postpartum Clinic Attendance

- After March 26, 2020, 25 (10%) of patients were eligible and had not yet attended a postpartum visit, defined as “post CoVID”
- There was no difference in primary outcome among those pre and post CoVID
 - 16/25 (64%) vs 144/225 (64%), $p= 0.374$

Clinic Visit Attendance

Variable	Attendance n=160	No Attendance n=90	P value
Age	30.9+/- 5.8	29.5 +/- 6.3	0.077*
Non-white race	143 (89.4)	79 (87.8)	0.701†
Single	83 (52.2)	55 (61.8)	0.145†
Less than high school	N=124 9 (7.3)	N=55 11 (20.0)	0.013 †
Distance >20 miles	15 (9.4)	16 (17.8)	0.053†
Nulliparous	71 (44.4)	32 (35.6)	0.174†
Gestational age at intake	15.3 (9.3-28.8)	23.8 (12.9-34.1)	0.001 §
Obese at consent	80 (50.0)	53 (58.9)	0.176†
High risk clinic	53 (33.1)	47 (52.2)	0.003 †
Any hypertension	67 (41.9)	31 (34.4)	0.248†
Diabetes	44 (27.5)	30 (33.3)	0.332†
Mental health disorder	29 (18.1)	20 (22.2)	0.433†
	N=113	N=59	
Antepartum EPDS ≥10	22 (19.5)	13 (22.0)	0.692†
Antepartum EPDS ≥13	16 (14.2)	7 (11.9)	0.675†
Cesarean	46 (28.8)	28 (31.1)	0.695†
Severe intrapartum complications	7 (4.4)	10 (11.1)	0.042 †
NICU admission	15 (9.4)	11 (12.2)	0.479†
	N=160	N=88	
Inpatient EPDS ≥10	16 (10.0)	10 (11.4)	0.737†
Inpatient EPDS ≥13	9 (5.6)	6 (6.8)	0.706†
Social work inpatient	47 (29.4)	33 (36.7)	0.235†

Predictors of Clinic Non-Attendance: Multivariable

Variable	Attendance n=160	No Attendance n=90	P value	OR 95% CI	aOR 95% CI
Age	30.9+/- 5.8	29.5 +/- 6.3	0.077	1.04 (0.99-1.09)	1.08 (1.00-1.15)
Non-white race	143 (89.4)	79 (87.8)	0.701	1.17 (0.52-2.62)	1.22 (0.36-4.13)
Single	83 (52.2)	55 (61.8)	0.145	0.68 (0.39-1.15)	0.63 (0.30-1.32)
Less than high school	N=124 9 (7.3)	N=55 11 (20.0)	0.013	0.31 (0.12-0.81)	0.42 (0.15-1.18)
Distance >20 miles	15 (9.4)	16 (17.8)	0.053	0.48 (0.22-1.02)	1.54 (0.44-5.39)
Nulliparous	71 (44.4)	32 (35.6)	0.174	1.45 (0.84-2.46)	3.09 (1.34-7.15)
Gestational age at intake	15.3 (9.3-28.8)	23.8 (12.9-34.1)	0.001	0.96 (0.94-0.99)	0.99 (0.95-1.03)
Obese at consent	80 (50.0)	53 (58.9)	0.176	0.70 (0.41-1.18)	0.92 (0.43-1.99)
High risk clinic	53 (33.1)	47 (52.2)	0.003	0.45 (0.27-0.77)	0.34 (0.16-0.72)
Severe intrapartum complications					0.42 (0.12-1.52)

Younger age aOR **1.08 (1.00-1.15)**
 Multiparity aOR **3.09 (1.34-7.15)**
 High-risk aOR **2.94 (1.39-6.25)**
 remained predictive for clinic visit NON-attendance after
 adjusting for confounders

Secondary Outcomes

Secondary Outcome – ED Visits

Variable	Arm 1 (n=125)	Arm 2 (n=125)	P-value
ED visit within 30 days of delivery	10 (8.0)	8 (6.4)	0.635†
ED visit postpartum days	<i>N</i> =10 14.5 (8-21)	<i>N</i> =8 7 (7-13.5)	0.284 §

Secondary Outcome ED Visits
8.0% vs 6.4%, p = 0.635

Data are n(%), median (25-75% IQR); *t-test; † Chi square test ‡ Fisher's exact; § Wilcoxon rank sum

Predictors of ED Visit

Variable	ED Visit (n=18)	No ED Visit (n=232)	P-value
Age	30.4 +/- 7.9	30.4 +/- 5.8	0.958*
Non-white race	16 (88.9)	206 (88.8)	0.999†
Single	10 (55.6)	128 (55.6)	0.994†
Less than high school	N=12 1 (8.3)	N=167 19 (11.4)	0.999†
Distance >20 miles	2 (11.1)	29 (12.5)	0.999†
Nulliparous	7 (38.9)	96 (41.4)	0.836†
Gestational age at intake	15.5 (10.1-28.1)	19.1 (10.1-31)	0.512†
Obese at consent	11 (61.1)	122 (52.6)	0.485†
Any hypertension	9 (50.0)	89 (38.4)	0.330†
Diabetes	4 (22.2)	70 (30.2)	0.477†
Mental health disorder	7 (38.9)	49 (21.1)	0.082†
	N=17	N=155	
Antepartum EPDS ≥ 10	8 (47.1)	27 (17.4)	0.004†
Antepartum EPDS ≥ 13	5 (29.4)	18 (11.6)	0.041†
Cesarean	6 (33.3)	68 (29.3)	0.719†
Severe intrapartum complications	3 (16.7)	14 (6.0)	0.112‡
NICU admission	3 (16.7)	23 (9.9)	0.412†
	N=18	N=230	
Inpatient EPDS ≥ 10	4 (22.2)	22 (9.6)	0.104‡
Inpatient EPDS ≥ 13	3 (16.7)	12 (5.2)	0.084‡
Social work inpatient	13 (72.2)	67 (28.9)	<0.001†

Not significant:

- Demographics, obesity, hypertension, diabetes, cesarean or NICU

Trending significant

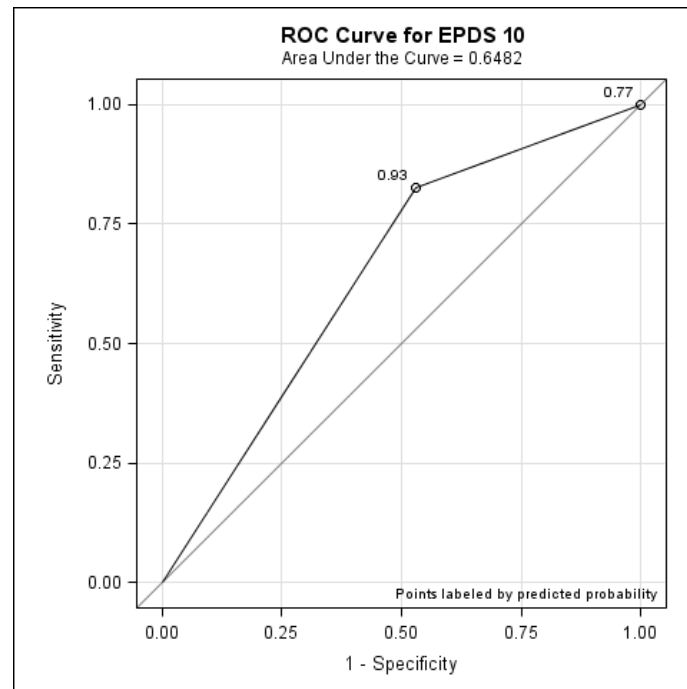
- Mental health disorder
- Severe intrapartum complications
- Inpatient EPDS ≥ 13

Significant

- Antepartum EPDS score ≥ 10
- Social work consult inpatient

Univariable Analysis

- EPDS ≥ 10 in antepartum period predicting ED usage within 30 days postpartum
 - AUC 0.648



Predictors of ED Visit: Multivariable Analysis

Variable	ED Visit (n=18)	No ED Visit (n=232)	P-value	OR 95% CI	aOR 95% CI
Age	30.4 +/- 7.9	30.4 +/- 5.8	0.958	1.00 (0.93-1.09)	1.01 (0.93-1.10)
Mental health disorder					0.47 (0.11-2.04)
Social work consult remained predictive for ED visit after adjusting for confounders aOR 5.43 (1.38-21.4)					
Antepartum EPDS ≥10	8 (47.1)	27 (17.4)	0.004†	4.21 (1.49-11.91)	2.87 (0.83-9.97)
Cesarean	6 (33.3)	68 (29.3)	0.719	1.21 (0.43-3.34)	1.06 (0.33-3.36)
Severe intrapartum complications	3 (16.7)	14 (6.0)	0.112‡	3.11 (0.80-12.04)	2.75 (0.58-13.04)
	<i>N</i> =18	<i>N</i> =230			
Inpatient EPDS ≥13	3 (16.7)	12 (5.2)	0.084‡	3.63 (0.92-14.28)	1.46 (0.29-7.47)
Social work inpatient	13 (72.2)	67 (28.9)	<0.001†	6.14 (2.10-17.89)	5.43 (1.38-21.4)

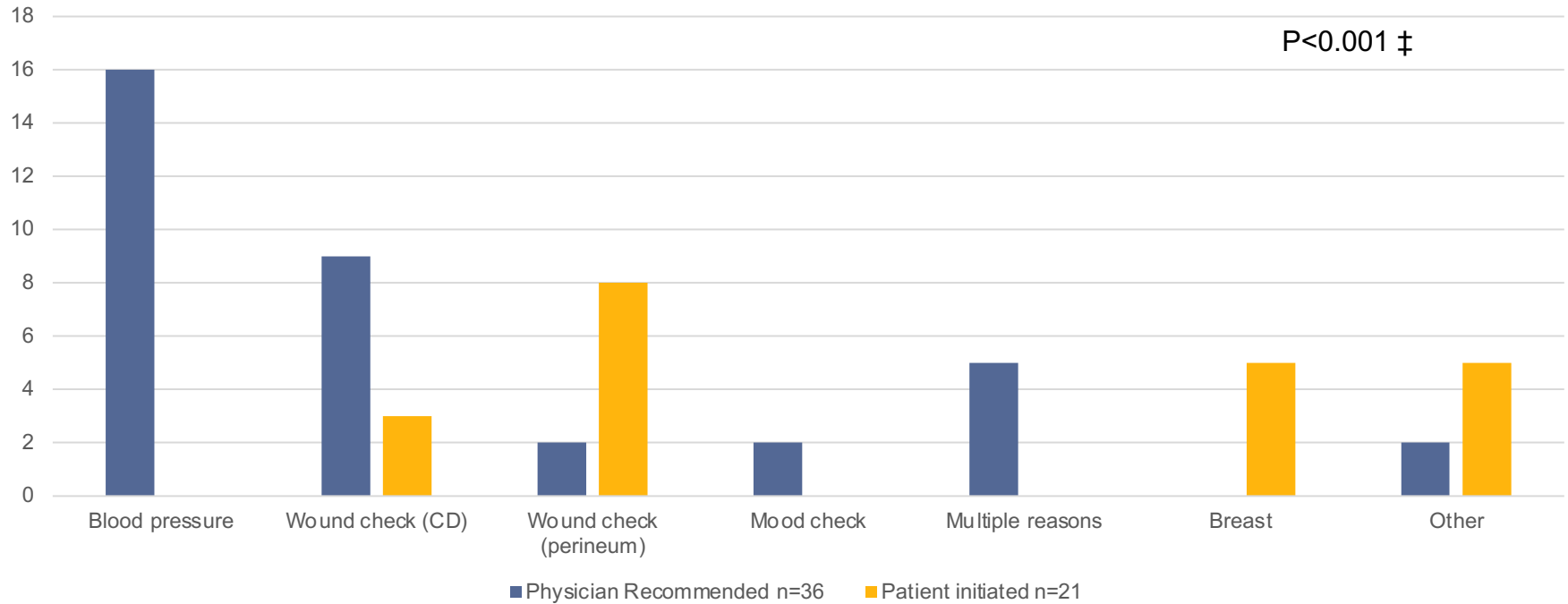
Secondary Outcome – Attendance at Non-Routine Visits

Variable	Arm 1 (n=125)	Arm 2 (n=125)	P-value
Attendance at non-routine visit	37 (29.6)	20 (16.0)	0.010†
Physician recommended	27 (73.0)	9 (45.0)	0.037†
Patient initiated	10 (27.0)	11 (65.0)	
Non-routine visit postpartum days	<i>N</i> =37 12 (9-17)	<i>N</i> =20 10.5 (8-22)	0.834 §

Data are n(%), median (25-75% IQR); *t-test; † Chi square test ‡ Fisher's exact; § Wilcoxon rank sum

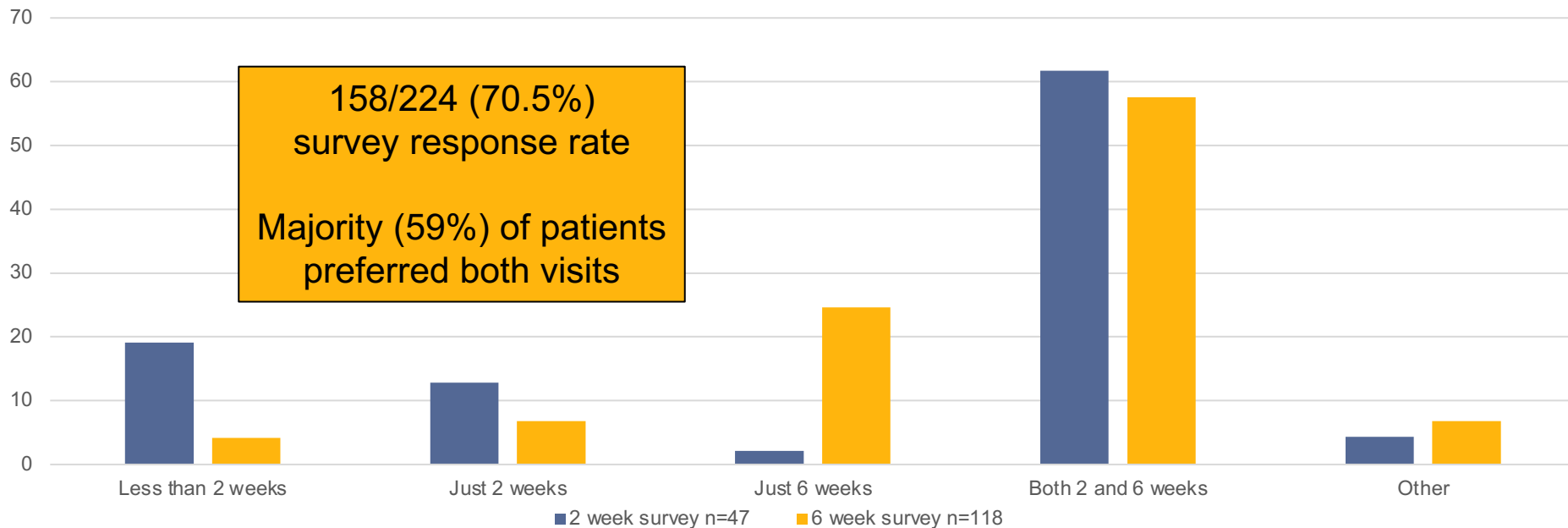
Secondary Outcome Non-Routine Visits
30% vs 16%, p = 0.010
Driven by more Physician recommended visits in Arm 1

Indications for Non-Routine Clinic Visit



Survey Results – Patient Preference

Patient Preferences of Timing of Postpartum Visit



Discussion

Conclusions

- In an obstetric clinic caring for a medically complex population with public insurance, the addition of a 2-week postpartum visit to the routine 6-week postpartum visit:
 - Did not significantly increase the likelihood of attendance at one or more routine postpartum visits (58% vs 70%, $p=0.065$)
 - Did not reduce the percent of women who presented to the ED within 30 days of delivery (8% vs 6%, $p=0.635$)
 - Did reduce the amount of non-routine clinic visits (30% vs 16%, $p=0.010$)

Strengths

- Randomized nature
- Minimal lost to follow up
- Comprehensive delivery data
- Sensitivity analysis demonstrated low risk of bias due to missing data

Limitations

- No blinding
- High rate of physician recommended non-routine visits potentially mitigating the effect of the earlier routine visit
- Population with high rate of medical comorbidities limiting generalizability
- Many non-routine clinic visits and ED visits were prior to two weeks

Future Directions

- Many women needed an early visit regardless of study arm suggesting that a universal versus targeted approach to an early postpartum visit may be optimal in a high-risk population, especially among those with psychosocial stressors

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