

1 **Pain Intensity predicts Pain Catastrophizing during the Postpartum Period: A**

2 **Longitudinal Random Intercept Cross-Lagged Panel study**

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20 Running title: Pain Intensity and Pain Catastrophizing

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## 24 **Abstract**

### 25 **Objective**

26 Pain catastrophizing is an important psychological predictor of pain. Recent evidence suggests  
27 the relationship between catastrophizing and pain intensity could be bidirectional, but most  
28 studies have been conducted on chronic pain patients and using criticized statistical methods. The  
29 present study aimed to examine if the relationship between pain intensity and catastrophizing was  
30 bidirectional in the context of childbirth.

### 31 **Methods**

32 A total of 504 women without chronic pain were recruited on their 32-37 gestational week. They  
33 completed measures of catastrophizing and pain intensity on the first encounter and then again at  
34 1, 3, and 6 months postpartum. The temporal relationship between the variables was assessed  
35 using a random intercept cross-lagged panel model.

### 36 **Results**

37 The hypothesis of reciprocal association did not receive support, as pain intensity predicted  
38 catastrophizing during the postpartum period, but catastrophizing did not show an effect over  
39 pain intensity at any moment.

### 40 **Conclusions**

41 Pain intensity predicting catastrophizing is consistent with previous literature, while the lack of  
42 effect of catastrophizing over pain intensity is an unexpected result, which may suggest that  
43 catastrophizing plays a different role in the postpartum period. These results highlight the  
44 importance of timely efforts for pain management during the postpartum period and contribute to  
45 the theoretical conceptualization of catastrophizing.

### 46 **Key words**

47 Catastrophizing; pain intensity; pregnancy; postpartum; Random intercept cross-lagged panel  
48 model

## 49 INTRODUCTION

50 Pain is a subjective experience shaped by biological, psychological, and social factors(1). Among  
51 the psychological factors, pain catastrophizing (PC) has received substantial attention in the past  
52 two decades, being considered as a prominent predictor of diverse pain outcomes(2), such as pain  
53 severity and disability on chronic pain patients(3–5) and persistence of pain and recovery in  
54 postoperative settings(6–11).

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57 PC comprises magnification of the painful stimulus threat value, feelings of helplessness, and  
58 constant worry about pain(2). According to the Fear and Avoidance Model of pain, people who  
59 respond to pain with catastrophic thoughts may develop a series of maladaptive emotions and  
60 cognitions towards pain (i.e. fear, anxiety, and hypervigilance), promoting an avoidance behavior  
61 which can lead to disuse, disability, depression and, as a consequence, more pain(12). In this  
62 model, PC is considered a response to pain; however, research has typically conceptualized it as  
63 an enduring personality factor preceding or having a causal influence on the pain  
64 experience(13,14).

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67 Only recently, the direction of the relationship between PC and pain intensity (PI) has begun to  
68 be tested longitudinally. For instance, clinically significant changes in PC over time predicted  
69 subsequent changes in PI and pain interference in patients with chronic musculoskeletal pain(15).  
70 Conversely, changes in PI have also shown to predict prospectively subsequent changes in  
71 PC(13,16). Consequently, the relationship between PI and PC could be bidirectional.

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74 In fact, five studies have tested this hypothesis using a cross-lagged panel model analysis  
75 (CLPM)(17–21). From these five studies, only the most recent of them (and the one with the  
76 biggest sample size, N= 538) supported a reciprocal relationship(17). The other four studies did  
77 find a prospective influence of PC over PI, but the converse relationship was non-significant.  
78 However, the samples used in these studies were relatively small ( $n \leq 90$ ), which renders the test  
79 of the hypothesis underpowered. Furthermore, except for one study (conducted with healthy  
80 subjects exposed to a painful stimulus)(20), all of them considered patients with chronic pain  
81 conditions undergoing multidisciplinary treatments. These patients had likely experienced pain  
82 and PC since long before the studies took place, making it difficult to elucidate their temporal  
83 relationship. To the best of our knowledge, no studies with acute pain samples have yet been  
84 conducted in this regard. Finally, the studies conducted until now have used traditional CLPM,  
85 which has been recently criticized(22). Studies examining the relationship between PC and PI  
86 need to be conducted using better statistical procedures (i.e., random intercept cross-lagged  
87 model) and samples without chronic pain. A better understanding of how these two variables  
88 influence each other longitudinally on non-chronic pain patients, not only will contribute to the  
89 theoretical framework of PC but will provide insight on the best time to intervene on catastrophic  
90 cognitions that may influence the experience of pain.

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93 The context of childbirth is very suitable for studying the temporal relationship between pain and  
94 PC. During the prenatal and the postnatal period, women commonly experience different kinds of

95 pains, and for a considerable percentage of these women (some estimates as high as 22%) the  
96 pain may become chronic(23). In this context, PC has shown to predict (i) low back PI during  
97 pregnancy(24), (ii) pain anticipation and PI – as well as the request for pain relief – during  
98 labor(25,26), (iii) postoperative pain one and two days after cesarean section(27), (iv) perineal  
99 pain in the first 24 hours after vaginal delivery(28), and (v) increased risk of persistent  
100 postpartum perineal(28) and lumbopelvic pain(29). The influences of PI on PC on the context of  
101 childbirth remain to be studied. Therefore, the main goal of the present study was to examine  
102 whether the relationship between PI and PC is bidirectional during the perinatal and postpartum  
103 periods on women without a history of chronic pain. We expected to find that PI and PC are  
104 reciprocally related.

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## 107 **METHODS**

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### 110 **Participants and procedure**

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113 Between 2017 and 2019, we invited pregnant women attending four medical centers of a  
114 university health network in Santiago, Chile to participate in the study. Inclusion criteria were  
115 being at 32–37 weeks of gestation of a viable pregnancy, being older than 18, and understanding  
116 the Spanish language. Women with a history of chronic pain condition before pregnancy were  
117 excluded from the study. Recruitment strategies included research assistants approaching

118 potential participants in the waiting rooms and dissemination of study information by health  
119 professionals and administrative staff. We did not keep a record of the number of pregnant  
120 women visiting the four centers during the study period or how many women were invited to the  
121 study. Nonetheless, we registered how many women gave their contact to recruiters and staff and  
122 accepted to be contacted to receive more information about the study. Of the 882 women who  
123 gave their contact information, 57% finally accepted to participate in the study.

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126 After obtaining informed consent, we applied a battery of questionnaires either in a face-to-face  
127 interview or via telephone, according to the participant's preferences. We contacted them again  
128 by phone at one, three, and six months after childbirth and asked them to complete a shorter set  
129 of questionnaires. As an exception, we offer the possibility to complete the questionnaires via  
130 email to those participants who were not able to answer by phone; less than 0.7% (n=10) of all  
131 the surveys across the waves were completed by this method. Participants received a gift card for  
132 10.000 Chilean pesos (approx. 14 US dollars) at the first assessment and 5.000 Chilean pesos  
133 (approx. 7 US dollars) at each follow-up. The study protocol was approved by the ethical  
134 committees of the host university and the funding agency.

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137 Figure 1 shows a flow diagram of enrollment, attrition, and drop out at each assessment. At the  
138 beginning of the study, 504 participants were recruited and 349 participants (69% of the original  
139 sample) provided data at all waves. We had missing data of 1, 87, 68, and 87 participants for the  
140 PCS and 2, 87, 69, and 87 participants for the Composed Pain Intensity Index at the first, second,

141 third, and fourth assessment respectively. The main reason for missing follow-up evaluations was  
142 the inability to contact the participants (by phone or email) and only 34 participants (7% of the  
143 total sample) refused to continue in the study because of lack of time or interest. Chi-squares test  
144 showed that those who completed all the follow-up assessments had a slightly higher educational  
145 level than the dropouts ( $\chi^2(4) = 14.09, p = .007$ ), but groups were no different in levels of PC ( $M_{diff}$   
146  $= -0.19, t(283) = -0.18, p = .855$ ) or PI ( $M_{diff} = -0.02, t(309) = -0.14, p = .887$ ).

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## 149 **Measures**

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152 We collected information about demographics, including age, educational level, family income,  
153 health care system, number of previous children, and marital status. Type of health care insurance  
154 (i.e. public versus private) was used as a proxy for socioeconomic status due to the strong  
155 association of these variables in Chile(30). Besides, after childbirth, we asked about the type of  
156 delivery and if there was any complication during the pregnancy/delivery or if the newborn  
157 presented any health problem at birth.

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161 Composed Pain Intensity Index (31).

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164 We asked participants about the perception of any pain during the last week, regardless of its  
165 etiology. Using four items, participants were asked to rate their current, worst, minor, and  
166 average pain on scales from 0 (no pain) to 10 (worst pain imaginable). The items were averaged  
167 to obtain a composite pain intensity index. This composed index has shown to be a more reliable  
168 way of assessing PI than individual numerical rating scales(31).

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171 Pain Catastrophizing Scale (PCS)(32)

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174 The PCS is a 13-item questionnaire that measures catastrophic thoughts and feelings about pain,  
175 on a scale from 0 (not at all) to 4 (all the time) (e.g. “I worry all the time about whether the pain  
176 will end”, “I become afraid that the pain may get worse”). The original structure(32) comprises  
177 three subscales assessing rumination, magnification, and helplessness. The sum of all the items  
178 provides a global score on catastrophizing. In the present study, we used only the global score.  
179 Higher scores represent greater PC. The Spanish version of the PCS has an adequate internal  
180 consistency (Cronbach alpha = 0.79) and test-retest reliability (intraclass correlation coefficient =  
181 0.84)(33). The internal consistency of the scale in our sample was excellent, with Cronbach  $\alpha$   
182 ranging from 0.95 to 0.96 on the four assessments.

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**185 Data Analytic Plan**

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188 To test the bidirectional relation between PC and PI, we estimated a Random Intercept Cross-  
189 Lagged Panel Model (RI-CLPM), a type of analysis that overcomes the limitations of the  
190 traditional cross-lagged models(22). As demonstrated by Hamaker et al. (22), when estimating  
191 the individuals' rank-order stability of a construct over time, part of this stability is due to stable,  
192 trait-like individual characteristics. When estimating regular CLPM, trait-like stability is not  
193 accounted for. In contrast, RI-CLPM allows parsing the part of the variability of a measure that is  
194 invariant, trait-like, from genuine autoregressive stability, which results in a more precise  
195 estimation of the cross-lagged relationships. Additionally, using multigroup analyses we tested  
196 the invariance of parameters across the type of health insurance, type of delivery (vaginal versus  
197 cesarean), primiparous versus multiparous participants, and pregnancy/delivery complication or  
198 newborn health problem. Analyses were run using full information maximum likelihood to make  
199 use of all available data. Besides the chi-square, traditional fit indices were used as criteria to  
200 determine the fit of the model; including the Comparative Fit Index (CFI), Tucker-Lewis Index  
201 (TLI), and Root Mean Square Error of Approximation (RMSEA). Because PC and PI were  
202 positively skewed, variables were log-transformed before the analyses.

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**205 RESULTS**

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208 Descriptive statistics are summarized in Table 1. Women's mean age was 31.2 years ( $SD=5.3$ ;  
209 range = 18-44), most of them were in a relationship (94.6%) and had tertiary education (87.7).  
210 The participants of our study presented similar characteristics to the women from the  
211 metropolitan area of Santiago, with the exception that our sample presented a higher educational  
212 level and was married in a greater proportion(34).  
213 Means and correlations between PC and PI scores are presented in Table 2. Except for PC at one  
214 month postpartum and PI at six months postpartum, all pairs of variables showed low to moderate  
215 significant positive correlations.

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## 218 **RI-CLPM**

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221 Figure 2 presents the parameters for the fitted RI-CLPM of PC and PI. The model had a good fit  
222 to the data;  $\chi^2(9)=14.59$ ,  $p=0.10$ , CFI=0.99, TLI=0.98, RMSEA= 0.04. The correlation between  
223 the trait-like PC and PI latent variables was .48 ( $p < .001$ ), suggesting a moderate association  
224 between these two stable characteristics. Autoregressive parameters of both variables were  
225 significant except between T1 and T2, showing low to moderate intra-individual stability.  
226 Our results did not support our bidirectional hypothesis, as none of the cross-lagged effects from  
227 PC to PI were significant. In contrast, we found a positive and significant effect from PI to PC  
228 between T2 and T3 (unstandardized  $\beta=0.24$ ,  $SE=0.10$ ,  $p=.015$ ), and between T3 and T4  
229 (unstandardized  $\beta=0.35$ ,  $SE=0.10$ ,  $p<.001$ ). No differences were found in autoregressive and

230 cross-lagged parameters on the separate multi-group analyses by type of health insurance  
231 ( $\Delta\chi^2(\Delta 12) = 13.131, p = 0.360$ ), type of delivery ( $\Delta\chi^2(\Delta 12) = 3.818, p = 0.987$ ), primiparous versus  
232 multiparous participants ( $\Delta\chi^2(\Delta 12) = 14.943, p = 0.245$ ), or pregnancy or delivery complication or  
233 newborn health problem ( $\Delta\chi^2(\Delta 12) = 9.553, p = 0.655$ ).

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## 236 **DISCUSSION**

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239 The present study aimed to test whether the relationship between PC and PI was bidirectional  
240 throughout the perinatal and postpartum periods. Our hypotheses were partially supported as PI  
241 predicted within-individual changes in PC during the postpartum period, but PC did not predict  
242 within-individual changes in PI between any of the adjacent assessment periods.

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245 Our results are in contrast with those reported by Racine et al.(17), who recently found a  
246 bidirectional relationship between PC and PI in patients with neuropathic pain. Our findings are  
247 also in contrast with the results of several other studies(18,19,21) in which only PC showed to  
248 predict PI, but the reverse association was not found. The discrepancies in the results may be  
249 related to methodological differences between studies. First, the bidirectionality of the  
250 relationship on previous studies was tested using statistical methods that have been widely  
251 criticized (i.e., traditional cross-lagged analyses), and may lead to overestimation of the mutual  
252 influences between the variables(22). By contrast, our use of RI-CLPM allows us to control for

253 variability due to stable individual characteristics. Specifically, the RI-CLPM model separates the  
254 between-person variance through the inclusion of a random intercept, allowing that the lagged  
255 relationship actually pertains to within-person dynamics(22). This distinction is especially needed  
256 when the stability of the constructs is to some extent of a trait-like, time-invariant nature, like has  
257 been suggested for PC(14). Our results show that, although the stable, trait-like PC and PI are  
258 moderately correlated, the within-individual PC variability (state-like) does not predict within-  
259 individual variability in PI over time.

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261

262 Second, previous studies were conducted with chronic pain patients(17–19,21) or healthy  
263 subjects exposed to a painful stimulus(20). By contrast, we recruited a sample of pregnant  
264 women, which did not present chronic pain before their pregnancy, and assessed how PC and PI  
265 influenced each other before and after giving birth. To our knowledge, this is the first study  
266 testing the mutual influence between PC and PI in the context of childbirth. On one hand, it is  
267 possible that the subjective experience of pain during the peri- and the post-natal period, when  
268 pain may be expected and normalized and is often associated with a positive outgrowth (the birth  
269 of a child), differs from other contexts where pain may be interpreted as a health threat(25). On  
270 the other hand, our results may indicate that PC plays a different role in acute pain conditions  
271 than in long-term persistent pains. This idea has been previously raised by Buenaver et al.(35)  
272 who conceptualizes PC as an inactive factor, which would need to be triggered by a pain  
273 condition of a certain duration or severity to exert its adverse influence over PI and other  
274 outcomes. This conceptualization is congruent with the results of a systematic review and meta-  
275 analysis, which found that the proportion of studies showing a significant association between

276 preoperative PC (and anxiety) and postoperative chronic pain was larger on the subgroup of  
277 musculoskeletal surgery patients, compared with other types of surgeries(6). Also, among  
278 musculoskeletal surgery patients, the maximum effect ORs were higher, compared with other  
279 types of surgeries(6), which led the authors to argue that, for musculoskeletal patients, pain after  
280 surgery may be a perpetuation of preexisting pain rather than the onset of new pain, which could  
281 suggest that PC (and anxiety) contribute less to eliciting (emerging) pain than to prolong  
282 consolidated pain(6). Similarly, and consistent with our results, Richez et al.(36) examined the  
283 role of different psychological predictors on persistent pain at three and six months after cesarean  
284 delivery; finding no association between presurgical PC and postsurgical persistent pain on any  
285 assessment. The authors compare their results with those reported by Pinto et al. (37) who found  
286 that PC predicted postsurgical persistent pain in women undergoing hysterectomy due to benign  
287 causes. According to Richez et al. (36), the similarities between both procedures allow suggesting  
288 that discrepancies between study findings may be due to differences in the surgical context,  
289 including that hysterectomy is often performed on women reporting preoperative pain. Likewise,  
290 Olsson et al. (29) found that PC at pregnancy was predictive of lumbopelvic pain at six-month  
291 postpartum, but only in participants with lumbopelvic pain present at the pregnancy assessment.  
292 Taken together, these studies (including the present work) suggest that the duration of the pain  
293 could play an important role in the relationship between PC and pain. This theory will need to be  
294 tested with longer longitudinal studies, that could examine if and when the role of catastrophizing  
295 changes over the onset of chronic pain.

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298 Regarding autoregressive relationships, the fact that PC in the postpartum period was predicted  
299 by PC in the same period, but not by PC before delivery, may suggest that catastrophic thoughts  
300 during both periods could be related to different psychological processes. In fact, a post hoc  
301 repeated-measure ANOVA analysis showed that PC before delivery was significantly higher than  
302 PC on the postpartum period ( $F(3, 1041) = 73.35, p < 0.001$ ). One possible explanation could be  
303 that PC on T1, being evaluated so close to childbirth, may be influenced by apprehensions  
304 regarding the delivery process. In order to explore this hypothesis, we ran an additional post hoc  
305 analysis to see if PC on T1 correlated with a measure of delivery expectancies in the same period.  
306 Specifically, we used four questions rated from 0 to 10, that have shown to predict labor pain in  
307 previous works(38, 39): 1)“How anxious are you feeling about your delivery?”, 2)“How much  
308 pain do you think you will feel during the procedure?”, 3) “How much fear do you have of  
309 experiencing pain during the delivery or due to your C-section?” and 4) “How much analgesia  
310 do you think you will require during and after your delivery?”. Previous exploratory factor  
311 analysis supported a single factor structure, so we average the items to obtain a global score. This  
312 scale showed a positive low correlation with PC ( $r = 0.344, p < .001$ ), supporting the idea that PC  
313 in this period is related to some extent to anxiety and fears associated with the forthcoming  
314 childbirth. This hypothesis should be further examined in future confirmatory studies.

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316  
317 In accordance with our expectations, we found that PI predicted PC during the postpartum period.  
318 This result is in line with the findings of numerous longitudinal studies that have supported this  
319 relationship throughout the recovery of surgery or injuries(13,17,40,41). However, our prediction  
320 that PI assessed during pregnancy will predict within-individual changes in PC at one-month

321 postpartum was not supported. Interestingly (and similar to what happened with PC), unlike  
322 when PI was assessed during the postpartum period, PI at pregnancy was not related to PI  
323 assessed one-month after delivery either, suggesting that this variable changed considerably from  
324 the prenatal to the postnatal period. This could be explained by the relatively high prevalence of  
325 pain experienced during pregnancy compared to the postpartum period(42). A post hoc repeated-  
326 measure ANOVA analysis showed that PI before delivery was significantly higher than PI on the  
327 postpartum period ( $F(3, 1035)= 67.17, p<0.001$ ). Likewise, it is possible that given that pain  
328 during pregnancy is more expected than in the postpartum, it could be less linked to catastrophic  
329 thoughts during this period.

330  
331  
332 The results of our study have both, practical and theoretical implications. First, they may suggest  
333 that it is important to timely manage pain during the postnatal period. An appropriate pain control  
334 could prevent catastrophic thoughts from growing and avoid the entrance on the vicious circle  
335 described in the Fear and Avoidance Model(12). This may be especially relevant in the context of  
336 childbirth, where the pain is sometimes naturalized(43) and the risk perception of analgesic  
337 consumption can be overestimated by both pregnant women and health professionals(44,45).  
338 On the other hand, our results also contribute to a better understanding of the construct of PC and  
339 its relationship with PI. Traditionally, PC has been considered as a relatively stable precursor of  
340 pain(14). The present research supports the idea that part of the PC variance corresponds to a trait  
341 like nature and another part seems to depend on other circumstances, such as PI. Moreover, our  
342 findings highlight that the relationship between PC and PI varies throughout different pain

343 conditions and contexts. The specific factors that may activate PC to play a predictive role over  
344 PI remain to be studied.

345

346

347 This study has some limitations. First, we assessed PI regarding any pain reported by the  
348 participant. That means that pain referred by some participants could not have been related to  
349 pregnancy or postpartum factors and could vary from one assessment to another. Even though  
350 “any pain” after childbirth is a common postpartum health problem reported, a study found that  
351 perineal pain had a different effect over other pain-related outcomes than “any pain”(46).

352 Therefore, whether the relationship between PC and PI varies among different pains, particularly  
353 those related to pregnancy, remains to be studied. Second, our study assessed variables until the  
354 first six months postpartum. Future research may examine if the relationship between PC and PI  
355 changes beyond that period, being possible that longer pains may activate the adverse effect of  
356 PC. Thirdly, we did not keep a record of the number of pregnant women visiting the recruitments  
357 centers during the study period or how many women were invited to the study. This limitation  
358 must be considered when generalizing the results of the study. Despite this, apart from  
359 educational level and marriage, participants of this study presented similar characteristics to the  
360 women from the metropolitan area of Santiago. Finally, our sample consists mostly of women  
361 from a middle socioeconomic class. Furthermore, we found that participants who missed at least  
362 one follow-up assessment had a lower educational level than completers. This limitation is  
363 particularly relevant considering that education has been identified as a significant moderator on  
364 the relationship between PC and PI(47). Given that the main reason for missing follow-up  
365 evaluation was the inability to contact the participant, it is possible that women with a lower

366 educational level had less access to a personal phone. This limitation could be addressed in future  
367 longitudinal research by adding additional contact methods as could be network contacts. It  
368 remains to be examined if the relationship between PC and PI is replicated in samples of women  
369 with a different educational background (lower socioeconomic status).

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## 372 **Conclusions**

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375 Despite these limitations, the results of the present study support the conclusion that during the  
376 postpartum period, changes in PI predict subsequent changes on PC, but this relationship is not  
377 reciprocal. The incongruence between these findings and those found in other cross-lagged  
378 studies may suggest that the relationship between PC and PI varies depending on the pain  
379 characteristics. Finally, these results highlight the importance of timely pain management  
380 strategies during the postnatal period, to prevent the growth of catastrophic thoughts and  
381 postpartum recovery difficulties.

382

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385

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393 **Conflict of Interest/disclosure Summary.**

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533 **Figure 1.** Study flow diagram. Abbreviations: FU= follow up.

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536 **Figure 2.** Random intercept Cross-Lagged Model of Pain Catastrophizing (PC) and Pain

537 Intensity (PI). Dashed lines indicate non-significant coefficients. Standardized coefficients in

538 parenthesis. All coefficients shown are significant  $p < .05$ . Abbreviations: PC= Pain

539 catastrophizing, PI = Pain Intensity.