**Name of journal:** *World Journal of Gastroenterology*

**ESPS Manuscript NO: 13417**

**Columns: ORIGINAL ARTICLE**

***Retrospective Study***

**Psychosocial mechanisms for the transmission of somatic symptoms from parents to children in the setting of irritable bowel syndrome**

van Tilburg MAL *et al*. Intergenerational transmission of illness behaviors

Miranda AL van Tilburg, Rona L Levy, Lynn S Walker, Michael Von Korff, Lauren D Feld, Michelle Garner, Andrew D Feld, William E Whitehead

**Miranda AL van Tilburg, William E Whitehead,** University of North Carolina, Center for Functional GI and Motility Disorders, Chapel Hill, NC 27599, United States

**Rona L Levy,** University of Washington, School of Social Work, Seattle, WA 98101, United States

**Lynn S Walker,** Vanderbilt University School of Medicine, Department of Pediatrics, Nashville, TN 37201, United States

**Michael Von Korff,** GroupHealth Research Institute, Seattle, WA 98101, United States

**Lauren D Feld,** Mount Sinai School of Medicine, New York, NY 10001, United States

**Michelle Garner,** University of Washington Tacoma, Department of Social Work, Tacoma, WA 98417, United States

**Andrew D Feld,** Group Health Cooperative, Spokane, WA 99255, United States

**Author contributions:** van Tilburg MAL was involved in writing of the manuscript and analysis of the data; Levy RL and Whitehead WE designed the study and were involved in writing of the manuscript and analysis of the data; Walker LS, Von Korff M were involved in study design and reviewed the manuscript; Feld LD and Garner M were involved in data collection and reviewed the manuscript.

**Supported by** grants from NIH, RO1 HD36069, RO1 DK31369, and R24 67674

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**Correspondence to: Miranda AL van Tilburg, PhD,** Division of Gastroenterology and Hepatology and Center for Functional Gastrointestinal and Motility Disorders, University of North Carolina at Chapel Hill, Campus Box 7080, Chapel Hill, NC 27599, United States. tilburg@med.unc.edu

**Telephone:** +1-919-8430688

**Fax:** +1-919-8432793

**Received:** August 22, 2014

**Peer-review started:** August 23, 2014

**First decision:** September 27, 2014

**Revised:** November 19, 2014

**Accepted:** December 20, 2014

**Article in press:**

**Published online:**

**Abstract**

**AIM:** To test multiple mechanisms to explain familial aggregation irritable bowel syndrome (IBS) including parental reinforcement/modeling of symptoms as well as family aggregation of coping, psychological distress, and exposure to stress.

**METHODS**: Mothers of children between the ages of 8 and 15 and with and without IBS were identified through the Group Health Cooperative of Puget Sound. Mothers completed questionnaires including child behavior checklist (child psychological distress), the family inventory of life events (family exposure to stress), SCL-90R (mother psychological distress) and the pain response inventory (beliefs about pain). Children were interviewed separately from their parents and completed the pain beliefs questionnaire (beliefs about pain), pain response inventory (coping) and child symptom checklist (gastrointestinal symptoms). In addition, health care utilization data was obtained from the automated database of Group Health Cooperative. Mothers with IBS (*n* = 207) and their 296 children were compared to 240 control mothers and their 335 children while controlling for age and education.

**RESULTS:** (1) Hypothesis 1, reinforcement of expression of gastrointestinal (GI) problems is related to GI but no other symptom (cold) reporting in children**:** There was no significant correlation between parental reinforcement of symptoms and child expression of GI or other (cold) symptoms; (2) Hypothesis 2, modeling of GI symptoms is related to GI but not non-GI symptom reporting in children**:** Children of parents with IBS reported more non-gastrointestinal (8.97 *vs* 6.70, *P <* 0.01) as well as more gastrointestinal (3.24 *vs* 2.27, *P <* 0.01) symptoms. Total health care visits made by the mother was correlated with visits made by the child (rho = 0.35, *P <* 0.001 for cases, rho = 0.26, *P <* 0.001 for controls); (3) Hypothesis 3, children learn to share the methods of coping with illness that their mothers exhibit:Methods used by children to cope with stomachaches differed from methods used by their mothers. Only 2 of 16 scales showed statistically significant but very weak correlations (Stoicism *r* = 0.13, *P <* 0.05 and Acceptance (*r =* 0.13, *P <* 0.05); (4) Hypothesis 4, mothers and children share psychological traits such as anxiety, depression, and somatization: Child psychological distress was correlated with mother’s psychological distress (rho = 0.41, *P <* 0.001 for cases, rho = 0.38, *P <* 0.001 for controls); (5) Hypothesis 5, stress that affects the whole family might explain the similarities between mothers and their children: Family exposure to stress was not a significant predictor of children’s symptom reports; and (6) Hypothesis 6, the intergenerational transmission of GI illness behavior may be due to multiple mechanisms: Regression analysis identified multiple independent predictors of the child’s gastrointestinal complaints, which were similar to the predictors of the child’s non-gastrointestinal symptoms (mother’s IBS status, child psychological symptoms, child catastrophizing and child age).

**CONCLUSION:** Multiple factors influence the reporting of children's gastrointestinal and non-gastrointestinal symptoms. The clustering of illness within families is best understood within a model that incorporates all these factors.

**Key words:** Abdominal pain; Coping; Illness behaviors; Psychological distress; Social learning; Stress

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**Core tip:** Irritable bowel syndrome (IBS) tends to run in families. IBS clusters within families. In previous studies, we found that this phenomenon could be explained by reinforcement and modeling of gastrointestinal illness behavior by parents. In the current study we want to extend our previous reports by examining various psychosocial influences on intergenerational transmission. We found that multiple psychosocial similarities between mother and child may explain familial aggregation, including: Mother’s modeling of IBS symptoms, shared psychological distress and shared family stress.

van Tilburg MAL, Levy RL, Walker LS, Von Korff M, Feld LD, Garner M, Feld AD, MD, Whitehead WE. Psychosocial mechanisms for the transmission of somatic symptoms from parents to children in the setting of irritable bowel syndrome. *World J Gastroenterol* 2015; In press

**Introduction**

Irritable bowel syndrome (IBS) tends to cluster in families[1-5]. The children of mothers with IBS are more likely than the children in control families to report bothersome gastrointestinal (GI) symptoms, to miss school for GI symptoms, and to be brought to the doctor for GI symptoms[1,2] We have referred to this phenomenon as the intergenerational transmission of GI illness behavior.

In a series of studies extending over more than 30 years, we[1-3,6-11] and others[12-15] have attempted to identify psychosocial factors that could explain how GI illness behavior is passed from one generation to another. In previous studies, we found that this phenomenon could be explained by social learning in the form of reinforcement and modeling of GI illness behavior by parents[1,2,6-8]. In the current study we want to extend our previous reports by examining various psychosocial influences on intergenerational transmission. These include: (1) Reinforcement of illness behaviors: Multiple studies have shown that parental protective behaviors can increase pain and disability in children with GI symptoms[16-20]. This suggests that selective attention to expression of GI problems or limiting the child’s normal activities and responsibilities may unintentionally reinforce illness behaviors; (2) Modeling of Illness behaviors: The observation that children of mothers who have IBS report more GI symptoms than the children of control mothers suggests that social learning through modeling processes may also contribute to the intergenerational transmission of GI illness behavior[2]; (3) Shared methods of coping with illness: Children may learn to interpret somatic symptoms in the same way as their parents (*i.e.*, as threatening or non-threatening) and to cope with abdominal pain in a similar way; (4) Shared psychological phenomena: Psychological distress is known to be associated with somatic symptoms. Mothers and their children may have similar psychological traits that predispose them to report somatic symptoms; and (5) Psychological stress: Studies in adults and children have shown that exposure to stress is associated with an increased tendency to report GI and other somatic symptoms[21-27]; thus, stress affecting the whole family could explain similarities in symptom reports between mothers and children.

A previous report of this study has been published[1] which supported the social learning hypothesis (a above) by showing that when parents respond solicitously to their children’s GI complaints their children are more likely to describe themselves as having serious GI symptoms, and they are more likely to miss school because of reported GI symptoms.

The aim of the analyses presented in this report is to look more broadly at several psychosocial influences on symptom reporting by examining several variables together. First, we will determine if transmission of mother to child is specific to GI symptoms or generalizes to other symptoms as well. Our first hypothesis is that reinforcement of expression of GI problems is related to GI but no other symptom reporting in children. Secondly, we hypothesize that modeling of GI symptoms is related to GI but not non-GI symptom reporting in children. For this we will examine if IBS mothers have children who not only report increased GI symptoms, but also increased cold symptoms compared to controls. The focus on cold symptoms is in line with previous research in adults[6] which found that IBS patients recalled parents to responds with empathy and concern to both GI symptoms and cold symptoms. Research in children has focused exclusively on GI symptoms so far. Our next three hypotheses examine similarities between mother and child which may explain shared GI symptoms: Our third hypothesis is that children learn to share the methods of coping with illness that their mothers exhibit. Our fourth hypothesis states mothers and children share psychological traits such as anxiety, depression, and somatization and our fifth hypothesis examines if family stress might explain the GI similarities between mothers and their children. Lastly we will examine if the intergenerational transmission of GI illness behavior may be due to multiple mechanisms (hypothesis 6).

**MATERIALS AND Methods**

This study is a secondary data analysis of an existing dataset. The methods described below are based on the original data collection and recruitment methods.

***Subjects***

The electronic medical records of Group Health Cooperative of Puget Sound were used to identify all women who had received a diagnosis of IBS or abdominal pain at a clinic visit during the preceding two years and who had one or more children between the ages of 8 and 15. Women with a diagnosis of IBS were sent a letter explaining the study and notifying them that they would be contacted by telephone unless they declined by returning a postcard. Women with a diagnosis of abdominal pain received a letter explaining the study, but were not contacted unless they returned a postcard confirming that they had symptoms consistent with IBS (we assumed that less than half would satisfy diagnostic criteria for IBS). Potential subjects were screened by phone to insure that the mother met Rome I criteria for IBS diagnosis[28]. These criteria are as follows: At least three months or longer of continuous or recurrent abdominal pain that is relieved with defecation and/or, changes in stooling frequency or consistency. In addition, two or more of the following at least on one-fourth of occasions or days: Altered stool frequency, altered stool form, altered stool passage, passing mucus, and/or bloating[29].

Control families were selected from the electronic medical records by identifying women who did not have a clinic visit during the preceding two years for IBS, abdominal pain, constipation, or diarrhea; and who had children within the 8-15 years old age range. Controls were also screened by telephone interview to insure that neither the mother nor any adult family member met Rome I criteria for IBS.

Additional inclusion criteria that applied to both cases and controls were ascertained in the telephone interview: (1) Mother was child’s legal guardian and child lived with her at least half of the time during the preceding two years; (2) Child did not have developmental disability requiring full-time special education; and (3) No family member with a diagnosis of ulcerative colitis or Crohn’s disease. At the conclusion of the telephone interview, subjects were scheduled for in-person interview. All eligible children willing to participate were enrolled in the study, allowing inclusion of multiple children per family.

This study was reviewed and approved by the Institutional Review Boards of Group Health Cooperative, the University of Washington, and the University of North Carolina.

***Procedure and measures***

Children and their mothers completed a battery of tests[1]. Children were interviewed separately from their parents by a trained interviewer.

**Child measures**: Pain beliefs questionnaire (CPBQ). This questionnaire measures children’s beliefs about their own abdominal pain[30]. This measure contains 32 statements about stomach aches. A trained interviewer asked the child to respond to each statement by rating how true it was for their stomach aches. There were 5 possible answers ranging from “not at all true” to “very true.” Out of 7 different subscales we included only the subscales for condition seriousness and condition frequency. The CPBQ has been found to be valid and reliable[31]. A sample question is: “My stomachaches mean I have a serious illness”.

Pain response inventory (PRI) – child version. This questionnaire measures how children cope with abdominal pain[32]. A sample question is: “If I have a stomachache or other stomach problems, I try hard to do something about it”. The PRI assesses three broad pain coping factors – active, passive, and accommodative, each with subscales representing specific strategies for coping with pain, including a 5-item catastrophizing subscale. The PRI has good internal reliability and validity[32].

**Child symptom checklist:** The child symptom checklist (CSCL) lists the 17 most common symptoms, including 5 GI symptoms, from the Children’s Somatization Inventory[34]. The child is asked to rate on a scale from 0-4 how much he or she was bothered by each symptom in the past two weeks. For example: “In the last 2 wk how much were you bothered by pain in your stomach or abdomen?” The Children Somatization Inventory has excellent validity and reliability[34,35]. For this study, we added 7 upper-respiratory symptoms: having a cold, runny nose, stuffy nose, sneezing, sore throat, coughing, and chest congestion. The questions coincide with respiratory questions on the Common Cold Questionnaire (general symptoms on this questionnaire such as ‘fevers’ were not included)[36] which has been validated in children with asthma. We added the item ‘having a cold’. The use of cold questions was modeled after research in adults, in which childhood reinforcement of cold symptoms reinforced adult cold symptoms but were independent of adult bowel symptoms[6,7]. Given that reinforcement of cold symptoms has previously been shown to be unrelated to reinforcement of bowel symptoms, the use of cold symptoms can measure the specificity of learned illness behaviors in children.

The CSCL inventory was scored for 3 subscales: GI symptoms, cold Symptoms, and non-GI, non-cold symptoms. Scale scores were computed by summing ratings for all items on the scale. Psychometric properties for the original questionnaire are good[34,37].

**Parental report of child distress:** Child behavior checklist (CBCL). The mother was asked to complete the CBCL for each child who participated in the survey[38]. This inventory contains 112 symptoms or behavior problems, and mothers were asked to respond by indicating whether these occurred “never, sometimes, or often”. A sample question is: “Acts too young for his/her age”. The CBCL can be scored for two global scales – internalizing and externalizing – as well as 9 subscales. In this study, the internalizing scale was used because it is conceptually most similar to the global symptom index of the SCL-90R used to test the mothers, and the T-scores transformations[38] were used so that boys and girls could be pooled together in the same analyses. The CBCL was also scored for the somatization, anxiety and depression subscales for comparison to adult psychological scales. The CBCL was been widely validated[38].

**Parent questionnaires:** The family inventory of life events (FILE). This is a parent-report form that assesses potentially stressful negative life events experienced by family members in the previous year[39]. For example: “Increased conflict with in-laws or relatives”. McCubbin *et al*[39] reported that internal consistency is 0.81 and four-week test-retest reliability is 0.80. Agreement between family members on the occurrence of events supports the validity of the FILE.

SCL-90R. The anxiety, depression, and somatization subscales of the SCL-90R was used to assess psychological distress in the mother[40]. Derogatis[40] reports high test-retest reliability and convergent validity. Derogatis and Cleary[41] also demonstrated a theoretically consistent factor structure. A sample item is “How much were you bothered by feeling fearful”.

 The pain response inventory (see above) – adult version. The PRI was reworded to assess the mother’s characteristic responses to her own abdominal pain[32].

Adult response to child symptoms (ARCS). Solicitous responding to gastrointestinal and cold complaints were measured by the Protective subscale of the ARCS[42], which was completed by the mother. Sample questions from this scale were “When your child has a stomachache or abdominal pain/cold, how often do you tell your child he/she doesn’t have to finish all of his/her homework?” The ARCS has high internal consistency and validity has been established for the Protective subscale[43,44].

***Health care utilization data***

The automated database of Group Health Cooperative was used to calculate the total number of ambulatory visits made by the child and by the mother for any illness and also the total number of visits for a lower gastrointestinal indication during a three-year period. This period extended from the date of the interview back two years in time and forward one year in time.

***Data analysis***

Many analyses below compare children of case and control mothers. Since these groups were significantly different for age and education, these two variables were entered as covariates in statistical tests comparing the two groups to control for these differences.

**Hypothesis 1:** Which states reinforcement of expression of GI problems is related to GI but not other symptom reporting in children, was evaluated by comparing cases and controls with respect to the scales of the ARCS measuring solicitous responses to expression of GI symptoms and solicitous responses to cold symptoms. Nonparametric (Spearman) correlations were used to compare mother’s scores on the solicitousness scales of the ARCS to symptom reports of children.

**Hypothesis 2:** States that modeling of GI symptoms is related to GI but not non-GI symptom reporting in children was evaluated by F-tests comparing case and control mothers and their children with respect to the frequency of medical clinic visits for different categories of illness. Nonparametric (Spearman) correlations were used to compare health care visits in children to health care visits by their mothers.

**Hypothesis 3***:* Which states that children learn to share the methods of coping with illness that their mothers exhibit, was tested by computing nonparametric correlations between the coping scale scores of children on the pain response inventory to the coping scale scores of their mothers on the adult version of the same inventory.

**Hypothesis 4:** States mothers and children share psychological traits such as anxiety, depression, and somatization, was tested by computing nonparametric correlations between children and their mothers on these psychological scales.

**Hypothesis 5:** Suggests that stress that affects the whole family might explain the similarities between mothers and their children; this was assessed by computing the correlation between the FILE35 and the GI and non-GI subscales of the child symptom checklist. All children were included in these analyses.

**Hypothesis 6:**The intergenerational transmission of GI illness behavior may be due to multiple mechanisms and some of the factors discussed above (*e.g.*, stress and anxiety) may be correlated with each other. Therefore, separate regression analyses were used to identify independent child and mother characteristics that predict GI and non-GI symptom reporting in children. To reduce the number of independent variables in these analyses, we first computed the nonparametric correlations of the dependent measures with all psychosocial variables measured and selected the variables with significant (*P <* 0.05) correlations with either dependent variable for inclusion in the regression analyses. The same set of predictor variables were used in both regression analyses.

**Results**

***Demographics***

Table 1 gives the demographic characteristics of the children and shows that the two groups were comparable. The average age of the 240 control mothers was 45.5 ± 0.4 years, and average age of the 207 IBS mothers was 41.8 ± 0.5 years (*P <* 0.01). Sixty percent of control mothers had completed college compared to 36% of IBS mothers (*P <* 0.001). Age and education of mothers was controlled for in between-group comparisons. The average number of children in the family was 2.1 ± 0.07 for controls and 2.0 ± 0.07 for case families.

Table 2 shows that symptoms and health care visits are highly correlated among children and parents, supporting the intergenerational transmission of expression of GI problems symptoms. The next set of analyses will aim to determine the psychosocial mechanisms by which this transmission may happen.

***Do mothers with IBS exclusively reinforce expression of GI problems?***

We investigated whether case mothers only behave in a solicitous fashion in response to illness behaviors related to GI symptoms. Contrary to hypothesis, there were no differences between case mothers and control mothers for solicitous responses to expression of GI symptoms (29.10 ± 0.64 *vs* 28.40 ± 0.63, F (1/608) = 0.23, *P =* 0.631), or cold symptoms (29.39 ± 0.64 *vs* 29.29 ± 0.59, F (1/611) = 1.542, *P =* 0.215). There was no significant correlation between the ARCS Protective scale for GI illness behavior (rho = 0.01; *P =* 0.76) and scores on this GI symptom scale, nor was there a significant correlation between the ARCS Protective scale for cold illness behavior and scores on the cold symptom scale (rho = 0.06; *P =* 0.11).

***Do mothers with IBS exclusively model GI symptoms?***

The question addressed was whether case mothers modeled only GI illness behavior (*i.e.*, modeled a unique class of illness behaviors) or whether they also displayed other types of illness behavior for possible imitation by their children. Table 2 shows that, in addition to excess clinic visits for GI symptoms, case mothers made more clinic visits than control mothers for all types of disorders evaluated – upper respiratory symptoms, asthma, headaches, and back pain. In a three-year period, the case mothers made three times as many health care visits overall compared to the control mothers, and visits with a GI diagnosis accounted for only 3 of 19 excess health care visits (Table 2).

Table 2 also provides data on whether case children showed selective increases in medical clinic visits for GI symptoms *vs* whether they showed increases in all types of health care visits. Case children had significantly more health care visits overall (12.90 visits per 3 years in case children *vs* 8.45 visits in controls), and they also had significantly more clinic visits for upper respiratory infections. In addition, Table 2 shows (last column) that the frequency of clinic visits made by children was significantly correlated with the number of clinic visits made by their parents across all disease categories – a finding that is consistent with social learning but does not show it to be specific to GI complaints.

***Do children model the coping mechanisms and other illness-related cognitions of their mothers?***

Correlations between children and their mothers on the scales of the Pain Response Inventory were generally very low. Among case children and their parents, only 2 of 16 scales showed statistically significant correlations; these were stoicism (*r =* 0.13; *P <* 0.05) and acceptance (*r =* 0.10; *P <* 0.05).

***Do children share psychological traits with their mothers?***

Table 3 shows the average scores on the psychological dimensions of overall psychological distress, anxiety, depression, and somatization in both mothers and children. Case mothers had significantly higher scores on all scales compared to control mothers, and case children had significantly higher scores on all scales compared to control children. For each of these psychological scales, the scores of children were significantly correlated with the scores of their mothers.

**Effects of shared, family stress on the child’s symptom reports:** The FILE score for family stress correlated significantly with the GI Symptom subscale of the Child Symptom Checklist (rho = 0.10, *P <* 0.05). However, a correlation of rho=0.10 is unlikely to be clinically meaningful.

**Regression analysis to identify contributors to child symptom reports:** In order to determine which child and parental factors are most important contributors to child expression of GI and non-GI problems, two regression analysis was performed: One with the GI Symptom subscale and the other with the other symptom subscale of the child symptom checklist as the dependent variable. Variables were included independent of whether mother-child overlap was found. This was in order to determine which of the variables were the most likely to influence reported symptoms. The variables which were found to make significant independent contributions to the symptom reports of the child are shown in Figure 1. The total variance that could be explained by this combination of variables was 12% for GI symptom reports and 16% for non-GI symptom reports. In general, the same variables predicted both GI symptom reports and non-GI symptom reports of the child, and the standardized betas show that these variables contributed to both types of symptom reports in a similar way. An exception was child’s age, which was positively correlated only with non-GI symptoms. In addition, mother’s IBS severity contributed only to child GI symptoms.

**Discussion**

There is a striking tendency for IBS to cluster within families. We evaluated five psychosocial hypotheses that may contribute to the intergenerational transmission of illness behaviors.

***Modeling***

 We hypothesized that a child may learn expression of GI symptoms through the observation of a model’s behavior (*i.e.*, the mother). This implies that the role model displays a specific behavior pattern and that the child emulates selectively this behavior pattern. When we tested the specificity of modeling by mothers, we found little evidence for it: mothers with IBS consulted physicians more often than control mothers for a wide range of somatic complaints, and only about 16% of their excess health care utilization was for GI symptoms. It was therefore not surprising to find that the children of IBS mothers reported excess somatic complaints that were not specific to GI as well: only about 10% of their excess visits were for GI diagnoses. The correlations between the number of health care visits by mothers and their children for all types of illness were more robust than the correlations between mother and child for GI visits. These data are consistent with the hypothesis that the intergenerational transmission of illness behavior may occur through modeling, but a diffuse pattern of comorbidity dominates the association. Many studies have previously found that pain and IBS in particular clusters in families[2,45] corroborating the current findings.

***Reinforcement of illness behaviors***

Another way a child may learn from a parent is when the parent reinforces expression of GI problems in their children by giving special attention, gifts *etc*. when the child complains of a stomachache. Multiple studies have shown that parental protective behaviors can increase pain and disability in children with GI symptoms[16-20,45]. What is new in the current study is that we predicted that the correlation between mother’s solicitous responses and child reported symptom would be stronger for GI symptoms than for cold symptoms. However, neither correlation exceeded 0.10, and there was no difference between them. We also previously found that parental solicitous response to child GI symptoms is not associated with reported abdominal pain frequency, but rather with the child’s perception of the seriousness of the symptoms[1]. The current study only focused on the reported occurrence of symptoms and did not examine which factors influence interpretation of the seriousness of symptoms. These differences in factors measured in this *vs* our earlier work may explain the difference in findings on the relationship between solicitousness and reported GI symptoms reported here. Furthermore, behavioral theory predicts that solicitousness is associated with changes in pain behaviors, but this study did not directly measure pain behavior per se, but reported symptoms.

The finding that both modeling and reinforcement of somatic symptoms was not different for cold and GI symptoms suggests that the intergenerational transmission of illness behavior from IBS mothers to their children is the transmission of a general pattern of somatic complaints and not a specific tendency to report GI symptoms.This is in contrast to studies in which adults were asked about their childhood experiences. In this case, specific patterns of symptoms were reported: Adults who remember their parents reinforcing cold symptoms, reported more cold symptoms but not GI symptoms, while adults who remember parents reinforcing GI symptoms reported more GI but not cold symptoms[7]. It needs to be investigated if this discrepancy with our current findings is due to recall issues in adulthood or due to developmental differences in the effects of parental reinforcement.

***Cognitions and coping styles***

We found no support for the hypothesis that intergenerational transmission of GI illness behavior is explained by the child adopting the methods used by the mother to cope with the threats posed by illness. There were no meaningful correlations between mothers and their children on the pain response inventory. The mother’s use of passive coping strategies was found to be negatively correlated with symptom reporting by the child. This implies its obverse, namely that mothers who are not passive but rather actively seek to solve their own stomach aches, are more likely to have children who make multiple somatic complaints.

***Psychological traits***

Both adults[46] and children[47] with functional GI disorders are known to have higher levels of psychological distress, and it has been suggested that psychological symptoms such as depression, anxiety, and somatization may predispose patients to report more somatic symptoms of many kinds. We found that mothers and their children are remarkably similar to each other in the psychological symptoms they display, even though there are marked methodological differences between the SCL-90R used to assess psychological symptoms in adults and the CBCL used to assess behavior problems in children. These findings are in line with studies that have shown intergenerational transmission of psychological disorders such as anxiety and depression[48,49].

It is unknown how children come to exhibit the same psychological symptoms as their mothers, although heredity and social learning through such mechanisms as modeling are possibilities. It is also unclear how these similarities in psychological traits can explain the intergenerational transmission of gastrointestinal complaints. It is possible that children learn to emulate their mothers’ hyper-vigilance for noticing somatic sensations and their tendency to attribute disease significance to these sensations. Consistent with this speculation, there was a positive correlation between mothers and child for somatization. However, the correlations for somatization were no greater than the correlations for anxiety and depression or for psychological symptom severity in general. Purposefully designed studies are needed to address the ways in which psychological traits are transmitted between generations and the ways in which they influence GI symptom reporting.

***Stress***

Family stress was significantly correlated to GI symptom reports by the child; however, the correlations were relatively weak (rho = 0.10) and likely of little clinical significance.

***Multiple factors predicting child GI symptoms***

Regression analyses show that there are multiple psychosocial factors that make independent contributions to the prediction of children’s symptom reports. The same psychosocial factors predict both GI symptom reports and non-GI symptom reports by the child. These include both characteristics of the mother - her IBS status and severity - and characteristics of the child – his or her overall level of psychological distress and tendency catastrophize. We would like to emphasize that catastrophizing about stomachaches by the child was associated with an increased tendency to report both GI and non-GI symptoms. This has been reported for adults[50] but has not previously been confirmed in children.

***Study limitations***

A limitation of this study was a modest participation rate. The necessity of obtaining consent from two (or more) family members (*i.e.*, the mother and each eligible child) was an important factor influencing the level of participation. A second limitation was that the control group of mothers was older and more educated than the cases. We controlled statistically for this limitation in our analyses, but we cannot rule out that it may still have affected our outcomes. A third limitation was that only mothers were studied; other research has demonstrated that the health status of the father may have as great an influence on the illness behavior of the child as the mother 1. Moreover, this study was designed with the intent of explaining the intergenerational transmission of illness behavior. However, our study was retrospective in nature and, therefore, a cause and effect cannot be confirmed. The findings we report in this manuscript should be regarded as exploratory, and should be confirmed in further prospective studies.

Other limitations relate to the challenges of assessing symptoms in children and the fact that methods used to assess psychological symptoms and coping in children were only approximately equivalent to methods used in their mothers. These methodological limitations may have contributed to the relatively low correlations and regression coefficients we observed.

**CONCLUSION**

In our previous publications, we emphasized the significance of modeling and reinforcement for the clustering of functional GI symptoms within families. The important contribution of the present study is that we expanded our understanding of the many ways multiple factors, including psychological phenomena, can influence a child’s response to somatic feelings. Mothers with IBS and their children are alike in their tendency to report most types of somatic symptoms, including both GI and non-GI symptoms. Children are also similar to their mothers in the severity and types of psychological traits they have, and psychological traits in the child are associated with increased symptom reports. Mothers and children in this study were not found to share cognitions and coping styles with pain. However, even if parents and children did not share similar coping, we found that less passivity on the part of the mother and a tendency to catastrophize about illness on the part of the child was associated with more somatic symptom reports. Thus, it appears there are multiple independent factors or behavior patterns that may be passed from, or shared by parent and child that contribute to the clustering of illness within families. Future research, and the potential for this research to contribute to our understanding, would benefit from including this more comprehensive, multidimensional perspective on this phenomenon.

**COMMENTS**

***Background***

Irritable bowel syndrome (IBS) is a chronic condition characterized by abdominal pain and changes in stool. IBS often runs in families. One reason for this is that children may inadvertently learn symptoms from their parents. Research has shown that when parents are sick themselves or respond with a lot of empathy and sympathy to the child’s symptoms, they may communicate that gastrointestinal symptoms are important and should be attended to.

***Research frontiers***

Parents play an important role in child’s symptoms. Parents drive a large part of the disability because they take the decision to take the child out of school and/or consult a doctor. In addition, the way parents react to their own symptoms and their child’s symptoms has been found to predict child’s abdominal pain level. Current research is exploring the many ways in which parents contribute to child’s symptoms and how to help parents in managing their child’s symptoms.

***Innovations and breakthroughs***

Previous studies have mainly focused on children learning through observing parental reactions to their own pain and to the child’s pain. However, children may learn from or share other characteristics with their parents that may put them at risk for developing IBS. For example, stress, psychological symptoms and how one copes with the gastrointestinal symptoms have been found to be important predictors of symptom severity. It is not known if these characteristics are shared by parent and child.

***Applications***

The current study explored if multiple psychosocial mechanisms may explain how gastrointestinal symptoms are transmitted from parent to child. We found that learning of symptoms was not specific to gastrointestinal symptoms but to other symptoms as well, indicating that children of parents with IBS learn to attend to all symptoms not just IBS symptoms. In addition, we found that parents and children shared similar levels psychological symptoms, which may indirectly increase gastrointestinal symptoms. Parent and children did not share the same methods of coping with symptoms, nor did family stress predict child symptoms. Thus, children of parents with IBS may be at increased risk of gastrointestinal symptoms because they learn to attend to all symptoms from their parents and because they share similar psychological symptoms with their parents which may indirectly increase gastrointestinal symptoms. The transmission of parent to child is best understood within a model that incorporates both these factors.

***Terminology***

Modelling is learning that occurs when a child observes and copies parental reactions to parent symptoms. Reinforcement is learning that occurs when the child receives positive or negative reactions of the parent to the child’s symptoms.

***Peer review***

This is an interesting article looking at the relationship between maternal and child somatic symptoms in the setting of IBS.

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**P-Reviewer:** McCloskey K **S-Editor:** Yu J **L-Editor: E-Editor:**

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**Figure 1** **Comparison of regression analyses used to identify independent predictors of children’s expression of gastrointestinal symptoms (right side of figure) and children’s non-gastrointestinal symptoms (left side of figure)**. Numbers beside arrows represent the beta weights for these predictors.

**Table 1 Characteristics of the sample**

|  |  |  |
| --- | --- | --- |
|  | **Case** | **Control** |
| Children, *n* | 296 | 335 |
| Age, Mean ± SD (yr) | 11.89 ± 2.62 | 11.84 ± 2.49 |
| Gender (male%) | 48.6 | 49.0 |
| Ethnicity (%) |  |  |
|  Caucasian | 77.5 | 82.6 |
|  African American | 5.8 | 5.7 |
|  Asian | 6.1 | 8.7 |
|  Hispanic | 4.8 | 2.4 |
|  Native American | 0.7 | 0.3 |

**Table 2 Number of medical clinic visit for each symptom category by mothers and children in case and control families, and correlation between number of mother and child visits**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Case families** | **Control families** | **Combined samples** |
|  | Mother | Child |  | Mother | Child |  | All Mothers | All Children |  |
| Symptom Category | Mean ± SE | Mean ± SE | *r* | Mean ± SE | Mean ± SE | *r* | Mean ± SE | Mean ± SE | *r* |
| Gastrointestinal | 3.27 ± 0.18c | 0.61 ± 0.16a | 0.064 | 0.20 ± 0.03 | 0.20 ± 0.05 | 0.042 | 1.62 ± 0.11 | 0.39 ± 0.08 | 0.219e |
| Upper Respiratory Infection | 2.07 ± 0.14c | 1.97 ± 0.15c | 0.290e | 0.74 ± 0.06 | 1.08 ± 0.09 | 0.326e | 1.37 ± 0.08 | 1.50 ± 0.09 | 0.355e |
| Asthma | 0.60 ± 0.13c | 0.48 ± 0.10 | 0.203e | 0.09 ± 0.02 | 0.30 ± 0.08 | 0.180e | 0.32 ± 0.06 | 0.37 ± 0.06 | 0.204e |
| Headache | 1.47 ± 0.42b | 0.25 ± 0.06 | 0.056 | 0.22 ± 0.06 | 0.14 ± 0.04 | 0.097 | 0.76 ± 0.20 | 0.19 ± 0.03 | 0.089d |
| Back Pain | 1.43 ± 0.23c | 0.17 ± 0.04 | 0.168d | 0.31 ± 0.05 | 0.12 ± 0.03 | -0.038 | 0.77 ± 0.09 | 0.14 ± 0.02 | 0.080d |
| All Outpatient Visits | 30.25 ± 1.53c | 12.90 ± 0.80c | 0.348e | 10.74 ± 0.43 | 8.45 ± 0.49 | 0.260e | 19.80 ± 0.083 | 10.50 ± .0.46 | 0.375e |

a*P* < 0.05 case *vs* control; b*P <* 0.01 case *vs* control; c*P* *<* 0.001 Case *vs* Control; d*P* < 0.05 mother *vs* child; e*P* < 0.001 mother *vs* child.

**Table 3 Psychological traits in children and their mothers**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Case families** | **Control families** | **Combined samples** |
|  | Mother | Child |  | Mother | Child |  | All Mothers | All Children |  |
| Psychological Trait | Mean ± SE | Mean ± SE | Rho | Mean ± SE | Mean ± SE | rho | Mean ± SE | Mean ± SE | rho |
| Global psychological distress  | 2.81 ± 0.10c | 55.16 ± 0.62c | 0.414e | 1.22 ± 0.07 | 48.46 ± 0.53 | 0.376e | 1.99 ± 0.07 | 51.65 ± 0.42 | 0.484e |
| Anxiety | 0.66 ± 0.04c | 4.64 ± 0.25c | 0.314e | 0.30 ± 0.02 | 2.65 ± 0.17 | 0.354e | 0.46 ± 0.02 | 3.60 ± 0.15 | 0.393e |
| Depression | 1.01 ± 0.04c | 2.20 ± 0.15c | 0.282e | 0.48 ± 0.03 | 1.29 ± 0.09 | 0.301e | 0.77 ± 0.03 | 1.72 ± 0.09 | 0.345e |
| Somatization | 1.15 ± 0.04c | 3.24 ± 0.16c | 0.366e | 0.45 ± 0.02 | 1.73 ± 0.10 | 0.157e | 0.78 ± 0.03 | 2.43 ± 0.10 | 0.374e |

c*P <* 0.001 Case *vs* Control; e*P <* 0.001 mother *vs* child.