

Abdominal massage as an adjunctive therapy for pediatric functional constipation: a randomized controlled trial

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ABSTRACT

Background. Chronic functional constipation affects approximately 3% of children globally, leading to painful defecation, fecal incontinence, and abdominal discomfort. Abdominal massage may improve gastrointestinal motility by stimulating vagal activity and reducing abdominal muscle tension. This study aimed to evaluate the effectiveness of abdominal massage therapy as an adjunct treatment for chronic functional constipation in children.

Methods. This randomized controlled trial included 61 children aged 4–10 years (mean age 6.36 ± 1.77) diagnosed with functional constipation. Participants were randomly assigned to two groups: the control group, receiving standard drug therapy, and the intervention group, receiving 12 sessions of Swedish abdominal massage involving effleurage and gentle pressing, vibration of the small and large intestines, kneading of the abdomen, and clockwise circular movements in addition to drug treatment. Outcomes assessed included stool consistency (using the Bristol Stool Scale), constipation severity (measured by the Constipation Assessment Scale), and associated symptoms.

Results. Both groups showed improvements in stool consistency; however, no statistically significant difference was found between them. The intervention group demonstrated a significantly greater reduction in constipation symptoms (Constipation Assessment Scale scores decreased from 14.70 ± 1.29 to 10.21 ± 1.45 , $P < 0.001$.) and fewer episodes of fecal incontinence (from 3.82 ± 1.33 to 2.70 ± 1.33 days/week, $P < 0.001$) compared to the control group.

Conclusion. A 12-session abdominal massage therapy program appears to be an effective adjunct treatment to standard pharmacological therapy for alleviating constipation-related symptoms in children. Larger, multicenter trials are needed to confirm these findings.

Key words: Swedish abdominal massage, children, chronic constipation, functional constipation, Bristol Stool Scale, Constipation Assessment Scale.

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Received 29th May 2025, revised 3rd July 2025, accepted 8th August 2025.

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Chronic functional constipation is one of the most common gastrointestinal disorders in children, with a global prevalence ranging from 7% to 30%. It often presents with symptoms such as painful defecation, abdominal discomfort, fecal incontinence, bloating, and reduced appetite. These symptoms can significantly impair a child's physical, emotional, and social well-being, interfering with daily activities, school performance, and peer relationships. Constipation accounts for approximately 3–5% of general pediatric consultations and 10–25% of referrals to pediatric gastroenterologists.¹⁻⁵

The management of chronic constipation typically involves pharmacological and non-pharmacological approaches. Pharmacological interventions include the use of laxatives such as stimulant laxatives, stool softeners, and fiber supplements, though long-term use may result in side effects and dependency.^{6,7} Non-pharmacological approaches aim to modify diet and behavior and include increased fiber and water intake, behavioral therapy, and complementary methods such as abdominal massage.⁸⁻¹² Abdominal massage has been proposed as a supportive therapy that enhances colonic motility and facilitates bowel movements. Its physiological mechanism is thought to involve stimulation of the parasympathetic nervous system and vagal activity, leading to improved colonic transit and reduced fecal retention.¹³⁻¹⁵ Furthermore, it may help reduce dependence on laxatives and improve comfort during defecation.¹⁶⁻¹⁹ Despite emerging evidence, abdominal massage therapy remains underexplored as a treatment for chronic functional constipation in children. High-quality randomized controlled trials are needed to determine its efficacy as a supportive therapy in children with functional constipation.

This study aims to assess the effectiveness of abdominal massage therapy as an adjunct pharmacological treatment for chronic functional constipation in children, based on a randomized controlled trial conducted at Bahrami Children's Hospital.

Materials and Methods

Study design and participants

This randomized controlled trial was conducted to assess the effects of abdominal massage therapy on children aged 4 to 10 years diagnosed with chronic functional constipation. Participants were recruited from the pediatric gastroenterology clinic at Bahrami Children's Hospital between 2022 and 2023. Diagnosis was confirmed according to the Rome IV criteria for functional constipation.^{5,19} Written informed consent was obtained from the parent or legal guardian of each participant.

Inclusion and exclusion criteria

Children were eligible if they experienced fewer than three spontaneous bowel movements per week, exhibited symptoms such as painful defecation, large-diameter stools, fecal incontinence, and difficulty during bowel movements. Exclusion criteria included metabolic, endocrine, or anatomical causes of constipation, neurological disorders, Hirschsprung's disease, severe systemic diseases (liver, kidney, heart), and current use of medications that may interfere with bowel function.

Sample size calculation

Sample size was calculated based on a power analysis from a previous study by Van et al.^{20,21} considering a power of 80%, a confidence interval of 95%, and an improvement rate of 63% in the control group and 93% in the intervention group. The calculated sample size was 26 per group, resulting in a total of 28 participants in the control group and 33 participants in the intervention group.

Intervention protocol

Eligible participants were randomly assigned to either the control or intervention group using a Google Random Generator software to ensure equal allocation. The randomization process was not blinded, and both the participants and

researchers were aware of group assignments. To minimize potential bias resulting from the lack of blinding, outcome assessors and data analysts were blinded to group allocation. All statistical analyses were conducted using de-identified and coded data. The control group received standard medical care, which included laxatives (polyethylene glycol) at a dose of 0.5 g/kg, adjusted based on individual needs. The intervention group received 12 sessions of Swedish abdominal massage therapy, in addition to the same drug treatment. The massage sessions were conducted twice a week for six weeks by a trained therapist with over four years of pediatric massage experience. This therapy involved Swedish abdominal massage, following a protocol described by Sinclair.²² Daily dietary intake of all participants was documented using food diaries throughout the study. Although no formal statistical comparison of dietary fiber or fluid intake was conducted between the groups, all participants and their families received standardized dietary advice. This included recommendations for adequate intake of fruits, vegetables, and fluids, and the avoidance of low-fiber foods and unhealthy snacks. Families were instructed not to make major dietary changes during the study or use dietary supplements unless approved by the study clinicians. We monitored the diet by reviewing the food diaries weekly. Laxative was administered based on each child's clinical requirement, with the frequency monitored through follow-up consultations. Adverse events were monitored throughout the study using standardized forms completed by caregivers and verified by study clinicians. Events were classified as mild, moderate or severe based on predefined criteria.

Outcome measures

Primary outcomes included stool consistency, assessed using the Bristol Stool Scale (BSS)²³, which categorizes stool types from 1 (hard lumps, severe constipation) to 7 (watery, severe diarrhea)²⁴; and the severity of constipation symptoms, assessed using the Constipation

Assessment Scale (CAS).²⁵ CAS is a validated tool consisting of 8 items, each scored from 0 to 2. The total score ranges from 0 (no symptoms) to 16 (most severe symptoms). Higher scores indicate greater severity of constipation-related complaints, such as painful or infrequent defecation, bloating, and straining. The scale was completed at baseline and post-intervention to evaluate changes in symptom severity.²⁵ Secondary outcomes included the frequency of fecal incontinence and the time until initiation of bowel movements. Symptoms were also monitored by evaluating straining, painful defecation, and withholding behavior.

Statistical analysis

All data were analyzed using SPSS version 22. Descriptive statistics, including means ($\pm$ standard deviation) for continuous variables and frequencies (percentages) for categorical variables, were computed. The chi-square test was used for categorical data, and repeated measures ANOVA was performed to analyze differences in primary outcomes between groups over time. Non-parametric tests were applied for non-normal distributions. A p-value of less than 0.05 was considered statistically significant. All statistical analyses were conducted by analysts blinded to group allocation.

Ethical considerations

The study was approved by the Research Council of the Faculty of Medicine at Tehran University of Medical Sciences, and the ethical code (IR.TUMS.MEDICINE.REC.1401.268) was obtained. The study was also registered in the Iranian Registry of Clinical Trials (IRCT20230929059549N1). Informed consent was obtained from the parents or legal guardians, and confidentiality of patient information was maintained throughout the study. The research adhered to the ethical guidelines outlined in the Declaration of Helsinki.

Results

A total of 66 participants were enrolled in the study with parental consent, with 5 children discontinuing participation (Fig. 1). The baseline demographic and clinical characteristics, including age, gender, height, and weight, showed no significant differences between the control and intervention groups ($P > 0.05$), indicating balanced allocation (Table I). Among the 61 participants, 41% were female and 59%

were male, with an average age of 6.36 ± 1.77 years.

At baseline, no significant differences were observed between the groups in terms of stool consistency, constipation severity, frequency of straining, or frequency of retentive fecal incontinence ($P > 0.05$) (Table II). The frequency of laxative use was also similar between the groups ($P > 0.05$).

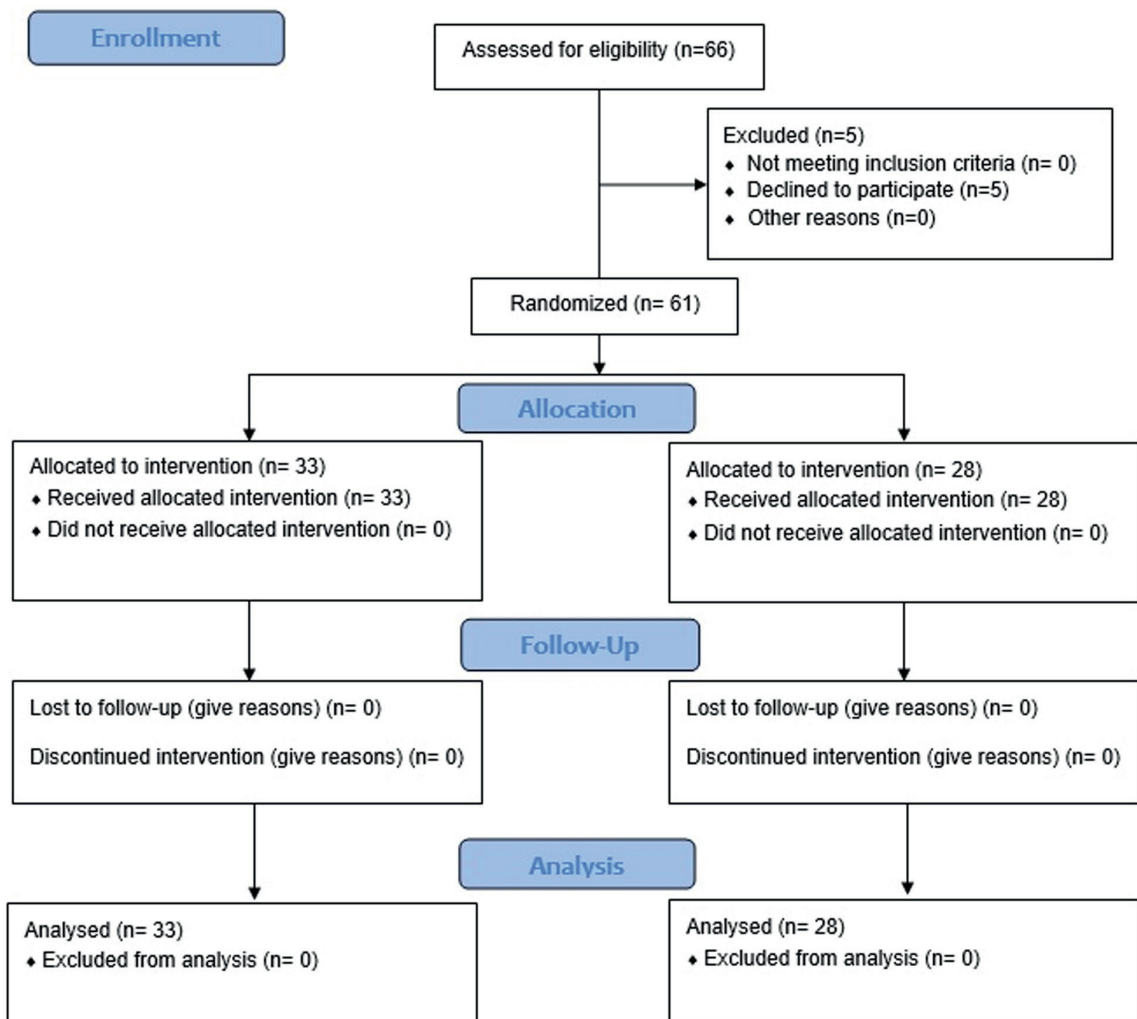


Fig. 1. Study population (CONSORT Flow Diagram).

Table I. Demographic characteristics of participants

Characteristic	All (n=61)	Control group (n=28)	Intervention (massage) group (n=33)	P-value
Age (years)	6.36 ± 1.77	6.36 ± 1.75	6.43 ± 1.81	0.897
Female gender	25 (41%)	11 (39.3%)	14 (42.4%)	0.506
Height (cm)	118.49 ± 19.482	117.93 ± 20.80	118.97 ± 18.60	0.839
Weight (kg)	20.29 ± 5.44	20.34 ± 5.28	20.24 ± 5.65	0.94

Table II. Analysis of stool variables

Variable	All (n=61)	Control group (n=28)	Intervention (massage) group (n=33)	P-value between groups
Bristol Stool Scale				
Before treatment				0.589
Type 1	26 (42.6%)	12 (42.9%)	14 (42.4%)	
Type 2	35 (57.4%)	16 (57.1%)	19 (57.6%)	
After treatment				0.138
Type 3	38 (62.3%)	20 (71.4%)	18 (54.5%)	
Type 4	23 (37.7%)	8 (28.6%)	15 (45.5%)	
P-value within group		<0.001	<0.001	
Constipation Assessment Scale				
Before treatment	14.62±1.42	14.54±1.57	14.70±1.29	0.661
After treatment	12.03±2.49	14.18±1.56	10.21±1.453	<0.001
P-value within group		0.398	<0.001	
Straining				
Before treatment	50 (82)	23 (82.1%)	27 (81.8%)	0.620
After treatment	24 (39.3)	11 (39.3%)	13 (39.4%)	0.601
P-value within group		0.001	<0.001	
Painful defecation				
Before treatment	44 (72.1)	20 (71.4%)	24 (72.7%)	0.567
After treatment	16 (26.2)	7 (25%)	9 (27.3%)	0.538
P-value within group		0.001	<0.001	
Retentive fecal incontinence				
Before treatment	3.80±1.44	3.79±1.59	3.82±1.33	0.932
After treatment	3.10±1.33	3.57±1.20	2.70±1.33	0.009
P-value within group		0.573	0.001	
Withholding behavior				
Before Treatment	40 (65.6%)	18 (64.3%)	22 (66.7%)	0.529
After Treatment	19 (31.1%)	9 (32.1%)	10 (30.3%)	0.548
P-value within group		0.016	0.003	

Control group only received drug treatment. Intervention group additionally received 12 sessions of Swedish abdominal massage.

Stool consistency

Stool consistency was assessed using BSS. At baseline, stool consistency was predominantly classified as type 1 or type 2, indicative of constipation. After treatment, both groups showed improvement in stool consistency, with a shift towards types 3 (sausage-shaped, with cracks) and 4 (smooth and soft, resembling a sausage). However, there was no significant difference in stool consistency between the two groups ($P > 0.05$) (Table II).

Constipation severity

Constipation severity was assessed using CAS, where higher scores indicate more severe symptoms. In the control group, there was no significant reduction in constipation severity, with average CAS scores decreasing slightly from 14.54 ± 1.57 to 14.18 ± 1.56 ($P = 0.398$). Conversely, the intervention group showed a significant decrease in constipation symptoms, with the average CAS score reducing from 14.70 ± 1.29 to 10.21 ± 1.45 ($P < 0.001$) (Table II).

Straining and painful defecation

The frequency of straining decreased in both groups, from 82.1% to 39.3% in the control group and from 72.7% to 39.3% in the intervention group; however, the difference between groups was not statistically significant ($P = 0.601$). The frequency of painful defecation decreased significantly in both groups. In the control group, the percentage decreased from 71.4% to 25% ($P < 0.001$), while in the intervention group, it decreased from 72.7% to 27.3% ($P < 0.001$). No significant difference in the frequency of painful defecation was found between the two groups ($P = 0.538$) (Table II).

Retentive fecal incontinence and withholding behavior

The frequency of retentive fecal incontinence decreased significantly in the intervention group, from 3.82 ± 1.33 days per week at baseline to 2.70 ± 1.33 days per week post-treatment ($P < 0.001$). In the control group, this change was not

significant, decreasing from 3.79 ± 1.59 to 3.57 ± 1.20 days per week ($P = 0.372$). Withholding behavior also decreased in both groups: from 64.3% to 32.1% in the control group ($P = 0.016$) and from 66.7% to 30.3% in the intervention group ($P = 0.003$), but there was no significant difference between the groups ($P = 0.548$) (Table II).

Adverse events

In the intervention group, 5 out of 33 participants (15.2%) reported mild abdominal discomfort during massage sessions, which resolved without intervention. No serious adverse events were reported in either group. In the control group, 2 out of 28 participants (7.1%) experienced mild diarrhea, likely related to laxative use.

Discussion

The findings demonstrated that abdominal massage significantly improved CAS and reduced fecal incontinence compared to pharmacological treatment alone. However, no significant difference was observed in stool consistency between the groups ($P > 0.05$), indicating that the effect of abdominal massage on stool consistency remains inconclusive based on our results. Therefore, claims regarding improvements in stool consistency should be made cautiously and require further investigation.

The therapeutic efficacy of abdominal massage may be attributed to its potential to enhance gastrointestinal motility and stimulate bowel movements, although the exact physiological mechanisms in children with chronic functional constipation require further research. Previous studies have reported that abdominal massage can improve colonic transit time and reduce constipation symptoms, supporting our findings.^{15,26}

The abdominal massage intervention was well-tolerated, with no serious adverse effects reported. Mild abdominal discomfort

observed in a few children was transient and did not necessitate discontinuation of therapy. These results align with prior studies suggesting that abdominal massage is a safe complementary therapy for children with functional constipation. A 2019 systematic review confirmed the effectiveness and safety of several complementary interventions, including massage, without any reported adverse effects.²⁷

Our results are consistent with a randomized controlled trial comparing drug therapy with manual physiotherapy—including abdominal massage—in children with functional constipation, which demonstrated symptom relief and improved stool consistency, although no statistically significant difference between treatment groups was found.²⁸ Moreover, systematic reviews have highlighted abdominal massage as a useful intervention to alleviate constipation severity and associated symptoms such as bloating and pain.²⁸ Participants in our study showed improvement in stool form, transitioning from severe constipation types (1 and 2) to more normal stool types (3 and 4) after treatment, indicating clinical benefit. This is aligned with findings from other pediatric studies, including research on infants with early-onset constipation who experienced symptom relief following abdominal massage.²⁹ However, our study focused on older children with more established behavioral and dietary patterns, suggesting that the benefits of massage extend across a wider pediatric age range.

A randomized controlled trial comparing drug therapy to manual physiotherapy in children with functional constipation found both methods effective in symptom relief and stool consistency improvement, although no statistically significant superiority was observed.³⁰ Although quality of life (QoL) and psychosocial factors were not formally assessed in our study, these are important dimensions to consider in future research. Future research should incorporate validated QoL assessments to examine this potential benefit more rigorously.

Functional constipation is influenced by complex interactions among physiological, psychological, social, and cultural factors. A systematic review and meta-analysis revealed that children with functional constipation report a lower quality of life compared to their healthy peers.³¹ Previous literature highlights that parental education, psychological health, and caregiving strategies significantly affect constipation management outcomes.^{32,33} In our study, parents were provided with information about the pathophysiology of constipation and the mechanisms through which massage could help, which likely enhanced their engagement. Although no formal satisfaction assessment was conducted, the observed clinical improvements and regular follow-up sessions may have contributed to maternal satisfaction. Furthermore, a recent systematic review underscored the association between pediatric constipation and exposure to stress, suggesting that life stressors—both at home and in school—can act as contributing factors.³⁴ In our context, it is reasonable to assume that symptom relief positively affected maternal emotional well-being, as improvements in the child's condition may reduce caregiver stress.

Limitations

Several limitations should be noted. First, the absence of a massage-only group limits the ability to isolate the specific effects of abdominal massage apart from pharmacological treatment. Second, the lack of a sham massage (placebo control) group reduces the ability to distinguish true therapeutic effects from placebo responses. Third, QoL and caregiver satisfaction were not evaluated. Lastly, the relatively small sample size and single-center design may limit generalizability. Larger, multicenter studies with longer follow-up and more comprehensive outcome measures are needed to validate and expand upon these findings.

Conclusion

In conclusion, this study provides evidence that abdominal massage combined with

conventional drug therapy is an effective safe complementary intervention for children with chronic functional constipation. The underlying physiological mechanisms—such as parasympathetic stimulation and enhanced gastrointestinal motility—support its therapeutic potential. While further research is needed to address study limitations, our findings support abdominal massage as a safe, cost-effective, and non-invasive complementary strategy. Future investigations should evaluate long-term efficacy, impacts on quality of life, and caregiver-administered protocols to improve adherence and accessibility.

Ethical approval

The study was approved by the Research Council of the Faculty of Medicine at Tehran University of Medical Sciences, and the ethical code (IR.TUMS.MEDICINE.REC.1401.268) was obtained. The study was also registered in the Iranian Registry of Clinical Trials (IRCT20230929059549N1). Informed consent was obtained from the parents or legal guardians, and confidentiality of patient information was maintained throughout the study. The research adhered to the ethical guidelines outlined in the Declaration of Helsinki.

Author contribution

The authors confirm contribution to the paper as follows: Study conception and design: AM, MM, HSM, MTOY, KE; data collection: AM, MA, MM, HSM, KE; analysis and interpretation of results: AM, MA, MM, MTOY, KE; draft manuscript preparation: AM, MA, MM, HSM, MTOY, KE. All authors reviewed the results and approved the final version of the article.

Source of funding

The authors declare the study received no funding.

Conflict of interest

The authors declare that there is no conflict of interest.

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