

Minireview Article

EXPLORING THE FACTS REGARDING COLON TRANSIT STUDY IN CHILDREN

ABSTRACT

Aim: Constipation is a problem faced by many children around the world. There are different ways of diagnosis and management of this condition. The key to the right diagnosis is based on the series of investigation that differs with children. The subject of interest is often the amount of time it takes for a substance to move through the colon which is known as Colon Transit Time[CTT]. Colon transit study or radio-opaque marker [ROM] study otherwise known as pellet study is one of the first choices of treatment to identify the CTT. Describing the colon transit study based on the established findings is the aim of this study.

Method: The MeSH terms used for literature retrieval were 'marker studies in children', 'pellet study in children', 'colon transit study in children'. The retrieval was performed based on pubmed, EMBASE, Web of Science as well as official websites. The search found 31 studies published from 2002 to 2022, as being met the inclusion criteria.

Results: The existing studies have identified a systematic way of performing the colon transit studies in children in many different ways. The colon transit differs with the segment of the colon and the age group of children.

Conclusion: Although the colon transit study requires series of intervention and investigation, the exact way of performing the same depends on the policies and protocols adopted by the individual specialist health care setting.

Key-Words: Colon Transit Time, Colon Transit study, pellet study

ABBREVIATIONS

CTT- Colon Transit Time

ROM- Radio-opaque Marker

VCE- Video Capsular Endoscopy

ARM-Ano-rectal manometry

1.INTRODUCTION

Constipation is a common problem in children and adults with a prevalence of 3% worldwide, but it is surprising that even about 29% of population has been reported to have the same globally¹. Most of the children with chronic constipation are managed in primary and secondary health care centres. However, a major proportion of children require referrals to specialist centres, which further brings an inflation to the cost of healthcare². Majority of the children have functional constipation which satisfies two or more of the criteria defined by Rome IV consensus³. These children do not need specialised investigations³. But, thirty-three percentage of children with constipation do not respond positively, so that they require diagnostic evaluation⁴. These kids are said to have intractable constipation, which is defined as poor response to optimum therapy for at least 3 months⁵. The consequence of this on the children and their families is detrimental. Therefore, this needs careful assessment and evaluation to bring out positive results. However, prior to adopting diagnostic strategies, it is essential to determine the effectiveness and compliancy with the use of laxatives.

The pathophysiology of constipation throws light on the coordination and integrity of the nerves and muscles in the colon, rectum and anus; how quickly the bolus passes through the colon and the way the faeces are expelled from the rectum⁶.

There are different methods to assess each of the physiological processes, at the same time, there are limitations for each of them. Manometry studies of the colon permits the neuromuscular contraction studies, but it does not evaluate the problems with faecal expulsion or transit times⁷. There are radiopaque marker (ROM) studies which is advantageous in this regard, but has the risk of exposure to radiation and also the markers advance through the colon differently in comparison to the faeces⁸. Video Capsule Endoscopy [VCE] and scintigraphy, which are the other set of investigations,

can only be performed in highly specialised centres. Ano- Rectal Manometries [ARM] are also performed but this evaluates only the anus and the rectum.

As there are no tests which can be described as complete, nine reviews identify the tests to determine the function of the colon as - first line screening and in-depth second line investigations. This is divided so, based on the complexity, accessibility and successful nature of the investigations. This article is about the colon transit marker studies which is categorised as one of the first line investigations as per ten different researches. Any searches with unavailability of the full texts or complete information have been excluded from the study.

There are a number of indications for performing the colon transit studies according to eight studies, which were finally included in the review. Identification of children with faecal retention and to exclude them from the non-retentive faecal incontinence is an aim of transit marker study^{5,6,9,10}. It also helps in determining the type of constipation as to whether it is slow transit, segmental or recto-sigmoid delay¹¹⁻¹⁴. Marker studies are also used as a screen before any major second-line tests of motility are done¹⁵. Also, when there are uncertainties in the diagnosis after the symptoms are revealed or to detect the effectiveness of medications, the marker studies are carried out^{5,16-19}.

2.METHOD OF PERFORMING COLON TRANSIT MARKER STUDIES

The method of colon transit marker studies have been established by eight different authors. It can involve ingestion of single or sequential capsules which contain similar shaped or different shaped markers. This would follow X-rays which can either be single or in series.

With the single capsule technique, the patient consumes a single capsule containing 20 markers on day 1. The X-rays are done on Day 3 and Day 5 of the capsule swallow²⁰. Otherwise, X-rays can be done every 24 hours. The latter method helps evaluate the total and segmental transit times. But, to schedule an X-ray same time every day is practically inconvenient and has high risk of radiation exposure in smaller children. Sometimes, X-rays are done only on Day 5. However, this does not assess fast transit and has no quantitative evaluation of Colon Transit Time[CTT]. This method is also differently done with the ingestion of a single capsule containing 24 markers followed by X-rays only

on Day⁵. The most common method is ingestion of capsules containing 20 markers each on days 1, 2 and 3 followed by single X-ray on Day 4 or sequential X-rays on Day 4 and 7 and if more than 20% of the markers are remaining, further X-ray investigations are done²¹. There are uncommon methods of ROM as well. In one of them, a capsule containing 10 ring shaped markers is to be taken for 6 consecutive days followed by an additional capsule containing 20 rod shaped markers on days 4,5 and 6²². One single X-ray shot is taken on day 7, 24 hours after ingestion of the last capsule. However, this method is highly impractical in children.

As individual capsules contain pellets of markers, this test is also called pellet study. It is important that the children consume capsules the same time each day²³. Also, if they cannot swallow capsules as a whole, the pellets can be mixed with food items and then ingested. Impacted **colon** is found to affect the study^{24,25}, but disimpaction can be a challenge in practice. Any medication that affects the gastric motility [For example, laxatives] may need to be withheld depending on the individual need of the child, unless the test is done to detect the effect of medication.

3.INTERPRETATION OF THE RESULTS

The evidence for interpretation of the results for colon transit studies have been identified in nine different studies. The results of the marker studies can be interpreted both qualitatively and quantitatively^{18,19}.

To evaluate the marker studies **qualitatively**, the colon is divided into projection zones as right, left and rectosigmoid colon. This is done by dividing the abdominal radiograph such that a straight line overlays the spinal processes. Two imaginary lines from the fifth lumbar vertebra extending respectively to the left iliac crest and right pelvic outlet intersects the straight line such that the three quadrants are created²⁶. Visual inspection of the X-ray for the presence of markers in each of the zones differentiates between the various types. If more than 80 percent of the markers are evacuated, the study results are said to be normal. Slow transit constipation is qualitatively diagnosed if the markers are found scattered in the different zones²⁷. If more than 50% of the markers are confined to one of the zones, it becomes segmental delay²⁸.

Qualitative assessment is necessary to determine the need for further X-rays in the study.

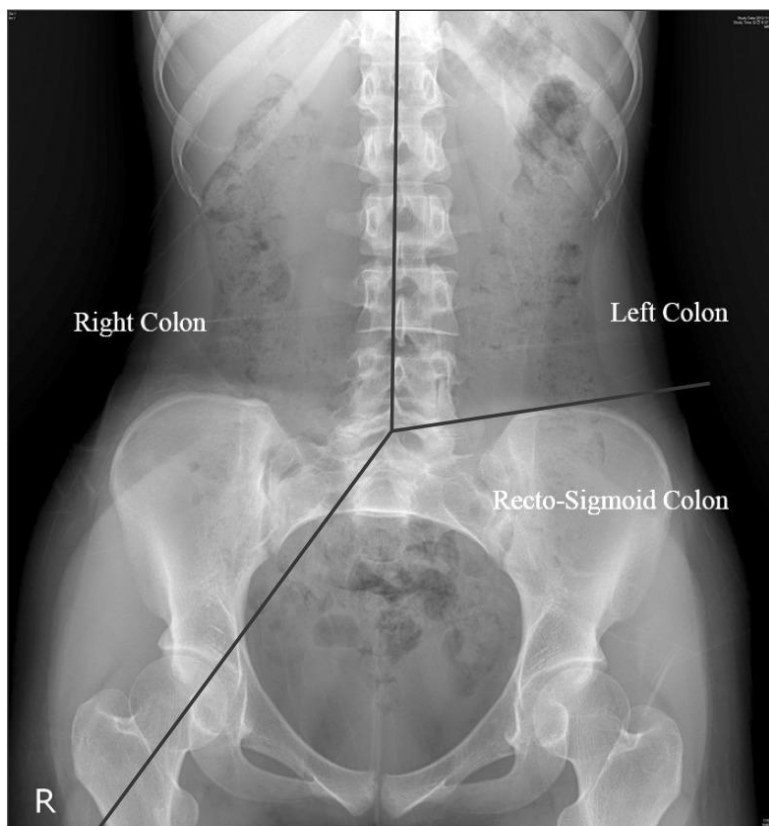


Fig1: Segmental division of Colon for CTT

Quantitative assessment measures the total and segmental colon transit in hours. Transit time is calculated by a specialised formula. The total number of markers in each colon segment or the entire colon is divided by a constant. The constant is obtained by dividing the time interval between the ingestion of capsules [in hours] by the total number of markers in each capsule²⁹. For example, in the common method whereby the child consumes one capsule containing 20 markers every day, the constant is 1.2 [24/20]. However, this formula cannot be used if the child did not evacuate all the markers in 3 days. Therefore, there is the need for another X-ray²⁹.

Various studies have established the normative values of colon transit time in children. Park (2004) has described a CTT with 3.1 ± 4.2 [mean \pm SD] hours for right colon, 5.1 ± 4.9 [mean \pm SD] hours for left colon and 7.4 ± 4.9 [mean \pm SD] hours for rectosigmoid region for children in the age of 2-10 years²⁶. Wagener defines a CTT of 5.5 ± 4.4 [mean \pm SD] hours for the ascending, 10.9 ± 9.6 [mean \pm SD] for the

transverse and 6.1 ± 5.4 [mean \pm SD] hours for the descending for children between 4 and 15 years³⁰. Another study published by Gutierrez et al explains a CTT of 29.08 ± 8.3 hours with 7.25 ± 5.75 [mean \pm SD], 6.6 ± 6.2 [mean \pm SD] and 14.96 ± 8.7 [mean \pm SD] hours respectively for right, left and rectosigmoid colon³¹. The same study has also proved that age and gender has an effect on the CTT among children between the age of 2 and 14 years³¹.

4.LIMITATIONS

Even though the marker studies are cheap and easily accessible, there is a higher risk of exposure to radiation with children than adults when the body surface area is considered. Also, the motion dynamics of markers is different from that of the faeces making the diagnosis different²³. The diagnosis can vary between experts as there is interobserver variability. Moreover, the formula used to calculate the CTT may not always be the same when the circumstances are taken into consideration²³. The compliance of patients can affect the study results as well. The study protocols differ from centre to centre.

When error count of the radio-opaque markers was considered, there was preference on barium suspension over X-rays as identified by 2 studies. However, as the searches done for this study comprise of selected databases and time periods, meta-analyses on the same might bring possible alternatives.

Although, this is the current scenario, marker studies still prove to be one of the best available first line investigation especially with regard to the assessment of constipation.

REFERENCES

1. Koppen IJN, Vriesman MH, Saps M, et al.. Prevalence of functional defecation disorders in children: a systematic review and meta-analysis. *J Pediatr* 2018;198:121–30. 10.1016/j.jpeds.2018.02.029

2. The Lancet Gastroenterology Hepatology . The cost of constipation. *Lancet Gastroenterol Hepatol* 2019;4:811. 10.1016/S2468-1253(19)30297-3
3. Hyams JS, Di Lorenzo C, Saps M. Functional disorders: children and adolescents. *Gastroenterology* 2016;S0016-5085. 10.1053/j.gastro.2016.02.015
4. Bongers MEJ, van Wijk MP, Reitsma JB, et al.. Long-Term prognosis for childhood constipation: clinical outcomes in adulthood. *Pediatrics* 2010;126:e156–62. 10.1542/peds.2009-1009
5. Tabbers MM, DiLorenzo C, Berger MY, et al.. Evaluation and treatment of functional constipation in infants and children: evidence-based recommendations from ESPGHAN and NASPGHAN. *J Pediatr Gastroenterol Nutr* 2014;58:258–74. 10.1097/MPG.0000000000000266
6. Vriesman MH, Koppen IJN, Camilleri M, et al.. Management of functional constipation in children and adults. *Nat Rev Gastroenterol Hepatol* 2020;17:21–39. 10.1038/s41575-019-0222-y
7. Li Y-W, Yu Y-J, Fei F, et al.. High-Resolution colonic manometry and its clinical application in patients with colonic dysmotility: a review. *World J Clin Cases* 2019;7:2675–86. 10.12998/wjcc.v7.i18.2675
8. Rao SSC, Kuo B, McCallum RW, et al.. Investigation of colonic and whole-gut transit with wireless motility capsule and radiopaque markers in constipation. *Clin Gastroenterol Hepatol* 2009;7:537–44. 10.1016/j.cgh.2009.01.017
9. de Lorijn F, van Rijn RR, Heijmans J, et al.. The leech method for diagnosing constipation: intra- and interobserver variability and accuracy. *Pediatr Radiol* 2006;36:43–9. 10.1007/s00247-005-0031-z
10. Southwell BR, Clarke MCC, Sutcliffe J, et al.. Colonic transit studies: normal values for adults and children with comparison of radiological and scintigraphic methods. *Pediatr Surg Int* 2009;25:559–72. 10.1007/s00383-009-2387-x
11. Frattini JC, Noguerras JJ. Slow transit constipation: a review of a colonic functional disorder. *Clin Colon Rectal Surg.* 2008;21:146–52.
12. Bharucha AE, Pemberton JH, Locke GR 3rd. American Gastroenterological Association technical review on constipation. *Gastroenterology.* 2013;144:218–38.

13. Lundin E, Graf W, Garske U, et al.. Segmental colonic transit studies: comparison of a radiological and a scintigraphic method. *Colorectal Dis* 2007;9:344–51. 10.1111/j.1463-1318.2006.01153.x
14. Maurer AH, Camilleri M, Donohoe K, et al.. The SNMMI and EANM practice guideline for small-bowel and colon transit 1.0. *J Nucl Med* 2013;54:2004–13. 10.2967/jnumed.113.129973
15. Tipnis NA, El-Chammas KI, Rudolph CD, et al.. Do oro-anal transit markers predict which children would benefit from colonic manometry studies? *J Pediatr Gastroenterol Nutr* 2012;54:258–62. 10.1097/MPG.0b013e31822bbcd8
16. Gould M, Renji E, Parmar R.P53 The effectiveness of colonic transit studies in the optimisation of the management of chronic constipation. *Frontline Gastroenterology* 2021;12:A44.
17. Bonapace ES, Maurer AH, Davidoff S, et al.. Whole gut transit scintigraphy in the clinical evaluation of patients with upper and lower gastrointestinal symptoms. *Am J Gastroenterol* 2000;95:2838–47. 10.1111/j.1572-0241.2000.03195.x
18. Kim ER, Rhee P-L. How to interpret a functional or motility test - colon transit study. *J Neurogastroenterol Motil* 2012;18:94–9. 10.5056/jnm.2012.18.1.94
19. Camilleri M, Bharucha AE, di Lorenzo C, et al.. American neurogastroenterology and motility Society consensus statement on intraluminal measurement of gastrointestinal and colonic motility in clinical practice. *Neurogastroenterol Motil* 2008;20:1269–82. 10.1111/j.1365-2982.2008.01230.x
20. Zaslavsky C, De Barros SG, Gruber AC, Maclel AC, Da Silveira TR (2004) Chronic functional constipation in adolescents: clinical findings and motility studies. *J Adolesc Health* 34(6): 517–522
21. Benninga MA, Voskuijl WP, Akkerhuis GW, Taminiu JA, Buller HA (2004) Colonic transit times and behaviour profiles in children with defecation disorders. *Arch Dis Child* 89(1):13– 16
22. Freedman PN, Goldberg PA, Fataar AB, Mann MM (2006) A comparison of methods of assessment of scintigraphic colon transit. *J Nucl Med Technol* 34(2):76–81
23. Popescu M, Mutalib M. Bowel transit studies in children: evidence base, role and practicalities. *Frontline Gastroenterol*. 2021 May 10;13(2):152-159. doi: 10.1136/flgastro-2020-101719. PMID: 35300467; PMCID: PMC8862445.

24. Sloots CEJ, Felt-Bersma RJF. Effect of bowel cleansing on colonic transit in constipation due to slow transit or evacuation disorder. *Neurogastroenterol Motil* 2002;14:55–61. 10.1046/j.1365-2982.2002.00304.x
25. Quitadamo P, Thapar N, Staiano A, et al.. Effect of bowel cleansing on colonic transit time measurement in children with chronic constipation. *J Pediatr* 2015;167:1440–2. 10.1016/j.jpeds.2015.09.035
26. Park ES, Park CI, Cho SR, Na SI, Cho YS (2004) Colonic transit time and constipation in children with spastic cerebral palsy. *Arch Phys Med Rehabil* 85(3):453–456. doi:10.1016/S0003-9993(03)00479-9 50.
27. Okawa, Y. Development of colonic transit time and ultrasound imaging tools as objective indicators for assessing abnormal defecation associated with food intake: a narrative review based on previous scientific knowledge. *BioPsychoSocial Med* **15**, 20 (2021). <https://doi.org/10.1186/s13030-021-00222-1>
28. Pomerri F, Frigo AC, Grigoletto F, et al.. Error count of radiopaque markers in colonic segmental transit time study. *AJR Am J Roentgenol* 2007;189:W56–9. 10.2214/AJR.07.2278
29. Bharucha AE, Anderson B, Bouchoucha M. More movement with evaluating colonic transit in humans. *Neurogastroenterol Motil* 2019;31:e13541. 10.1111/nmo.13541
30. Wagener S, Shankar KR, Turnock RR, Lamont GL, Baillie CT (2004) Colonic transit time—what is normal? *J Pediatr Surg* 39(2):166–169. doi:10.1016/j.jpedsurg.2003.10.002 (discussion 166–9)
31. Gutiérrez C, Marco A, Nogales A, et al.. Total and segmental colonic transit time and anorectal manometry in children with chronic idiopathic constipation. *J Pediatr Gastroenterol Nutr* 2002;35:31–8. 10.1097/00005176-200207000-00008