

Clinical Policy: Wireless Motility Capsule

Reference Number: LA.CP.MP.143

Date of Last Revision: 12/24

Coding Implications
Revision Log

See <u>Important Reminder</u> at the end of this policy for important regulatory and legal information.

Description

The wireless motility capsule (WMC) assesses gastroparesis or delayed gastric emptying.^{1,2} The WMC is an orally ingested, nondigestible, data-recording device that enables the simultaneous assessment of regional and whole gut transit.¹⁻³

Policy/Criteria

It is the policy of Louisiana Healthcare Connections that wireless motility capsule (WMC) is **not medically necessary** for the evaluation of suspected gastric and intestinal motility disorders, as well as all other indications. There is a paucity of peer-reviewed, evidence-based literature to determine that the diagnostic performance and clinical utility surpass conventional means of measuring gastric emptying.

Background

The U.S. Food and Drug Administration approved wireless motility capsule (WMC) for the evaluation of patients with suspected gastroparesis, even though there is no sign of a blockage. The WMC, which is a 26 x 13 mm size capsule with a battery life of five days, is also proposed to evaluate colonic transit time in patients with chronic idiopathic constipation. Additionally, the WMC is noted to continuously measure the temperature, pH, and pressure of its surrounding environment while traveling through the gastrointestinal tract, via gut peristalsis, until exiting the body through the anus. In the surrounding through the anus.

After eating a standard meal, the member/enrollee swallows the capsule and wears a small monitor that makes telemetry recordings. The established cutoff point for gastric emptying time is 300 minutes. Gastric emptying of the WMC seems to occur with the Phase III migrating motor complex, signifying completion of postprandial phase and return of the fasting state. It assesses small bowel transit time by a sharp increase in pH on entry into duodenum and by a fall in pH at the ileocecal junction. However, in 15% of patients, this pH drop is not observed, and this may be related to the ileocecal valve incompetence. An example of a wireless GI motility monitoring system is the SmartPill® GI monitoring system 2.0.

Advantages of the WMC include that it is wireless and painless and contains no radiation.³ Disadvantages of the capsule include failure to capture data that would require repeat testing, and delay or total failure to pass the capsule, requiring serial x-rays to document passage or endoscopic or surgical removal. Another disadvantage is that it should not be used in patients with a possible stricture, altered anatomy, or severe pyloric stenosis.⁷ Patients ideally should be able to tolerate not using proton pump inhibitors and histamine-2 blockers before testing.⁷

CLINICAL POLICY Wireless Motility Capsule



Agency for Healthcare Research and Quality (AHRQ)⁶

WMC is comparable in accuracy to current modalities in use for detection of slow-transit constipation and gastric emptying delay and is therefore another viable diagnostic modality. Little data are available to determine the optimal timing of WMC for diagnostic algorithms.

American College of Gastroenterology⁸

Scintigraphic gastric emptying of solids is the standard for the evaluation of gastric emptying and the diagnosis of gastroparesis. Alternative approaches for assessment of gastric emptying include WMC testing and 13C-spirulina breath testing. (Conditional recommendation, low quality of evidence).

American and European Neurogastroenterology and Motility Societies

Tests of gastrointestinal transit are available and useful in the evaluation of patients with symptoms suggestive of gastrointestinal dysmotility since they can provide objective diagnosis and a rational approach to patient management.⁹

Studies note that WMC is comparable in accuracy to current modalities in use for detection of slow-transit constipation and gastric emptying delay and is therefore another viable diagnostic modality. However, little data are available to determine the optimal timing of this device for diagnostic algorithms.¹⁰

Other studies have noted that the sensitivity and specificity of the WMC is comparable to radiopaque marker test and scintigraphic gastric emptying. WMC is well tolerated, has good compliance, and avoids the risk of radiation exposure, however, it is not clear if it provides added clinical value in most patients. 5,7,12

Coding Implications

This clinical policy references Current Procedural Terminology (CPT®). CPT® is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2023, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only and may not support medical necessity. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

NOTE: Coverage is subject to each requested code's inclusion on the corresponding LDH fee schedule. Non-covered codes are denoted (*) and are reviewed for Medical Necessity for members under 21 years of age on a per case basis.

CPT ®	Description
Codes	
91112*	Gastrointestinal transit and pressure measurement, stomach through colon, wireless
	capsule, with interpretation and report



HCPCS Codes	Description
N/A	

ICD-10-CM Code	Description
N/A	

Reviews, Revisions, and Approvals	Revision Date	Approval Date	Effective Date
Converted corporate to local policy.			
Annual review. Criteria section updated with wording for		1/14/23	
abbreviation. Background updated with no impact on criteria.			
References reviewed and updated. Specialist reviewed.			
Annual review. Background updated with no impact on criteria.	10/23	1/9/24	
References reviewed and updated. External specialist review.			
Added note for non-covered codes.			
Annual review. Removed ICD-10-CM verbiage and ICD-10	12/24	1/27/25	2/27/25
codes. References reviewed and updated.			

References

- 1. Rao SS, Camilleri M, Hasler WL, et al. Evaluation of gastrointestinal transit in clinical practice: position paper of the American and European Neurogastroenterology and Motility Societies. *Neurogastroenterol Motil*. 2011;23(1):8 to 23. doi:10.1111/j.1365-2982.2010.01612.x
- U.S. Food and Drug Administration 510(k) Premarket Notification Database. SmartPill GI Monitoring System. Version 2.0 Summary of Safety and Effectiveness No. K092342. Silver Spring, MD: FDA. July 29, 2009.
 - http://www.accessdata.fda.gov/cdrh_docs/pdf9/K092342.pdf. Accessed August 16, 2024.
- Local coverage determination: Wireless Gastrointestinal Motility Monitoring System (L33455). Centers for Medicare and Medicaid Services Web site. http://www.cms.hhs.gov/mcd/search.asp. Published October 01, 2015 (revised September 09, 2021. Accessed August 16, 2024.
- 4. Arora Z, Parungao JM, Lopez R, Heinlein C, Santisi J, Birgisson S. Clinical utility of wireless motility capsule in patients with suspected multiregional gastrointestinal dysmotility. *Dig Dis Sci.* 2015;60(5):1350 to 1357. doi:10.1007/s10620-014-3431-9
- 5. Lembo AJ. Overview of Gastrointestinal Motility Testing. UpToDate. www.uptodate.com Published July 26, 2023. Accessed August 20, 2024.
- 6. Stein E, Berger Z, Hutless S, et al. Wireless Motility Capsule Versus Other Diagnostic Technologies for Evaluating Gastroparesis and Constipation: A Comparative Effectiveness Review. Rockville (MD): Agency for Healthcare Research and Quality (US); May 2013.
- 7. Saad RJ. The Wireless Motility Capsule: a One-Stop Shop for the Evaluation of GI Motility Disorders. *Curr Gastroenterol Rep.* 2016 Mar;18(3):14. doi: 10.1007/s11894-016-0489-x
- 8. Camilleri M, Kuo B, Nguyen L, et al. ACG Clinical Guideline: Gastroparesis. *Am J Gastroenterol*. 2022;117(8):1197 to 1220. doi:10.14309/ajg.000000000001874f

CLINICAL POLICY Wireless Motility Capsule



- 9. 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(12):1269 to 1282. doi:10.1111/j.1365-2982.2008.01230.x
- 10. Farmer AD, Wegeberg AL, Brock B, et al. Regional gastrointestinal contractility parameters using the wireless motility capsule: inter-observer reproducibility and influence of age, gender and study country. *Aliment Pharmacol Ther*. 2018;47(3):391 to 400. doi: 10.1111/apt.14438
- 11. Lee AA, Rao S, Nguyen LA, et al. Validation of Diagnostic and Performance Characteristics of the Wireless Motility Capsule in Patients with Suspected Gastroparesis. *Clin Gastroenterol Hepatol.* 2019; 17(9):1770 to 1779.e2. doi:10.1016/j.cgh.2018.11.063
- 12. Tran K, Brun R, Kuo B. Evaluation of regional and whole gut motility using the wireless motility capsule: relevance in clinical practice. *Therap Adv Gastroenterol*. 2012;5(4):249 to 260. doi:10.1177/1756283X12437874

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. LHCC makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable LHCC administrative policies and procedures.

This clinical policy is effective as of the date determined by LHCC. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. LHCC retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible

CLINICAL POLICY Wireless Motility Capsule



for the medical advice and treatment of members/enrollees. This clinical policy is not intended to recommend treatment for members/enrollees. Members/enrollees should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom LHCC has no control or right of control. Providers are not agents or employees of LHCC.

This clinical policy is the property of LHCC. Unauthorized copying, use, and distribution of this clinical policy or any information contained herein are strictly prohibited. Providers, members/enrollees and their representatives are bound to the terms and conditions expressed herein through the terms of their contracts. Where no such contract exists, providers, members/enrollees and their representatives agree to be bound by such terms and conditions by providing services to members/enrollees and/or submitting claims for payment for such services.

©2023 Louisiana Healthcare Connections. All rights reserved. All materials are exclusively owned by Louisiana Healthcare Connections and are protected by United States copyright law and international copyright law. No part of this publication may be reproduced, copied, modified, distributed, displayed, stored in a retrieval system, transmitted in any form or by any means, or otherwise published without the prior written permission of Louisiana Healthcare Connections. You may not alter or remove any trademark, copyright or other notice contained herein. Louisiana Healthcare Connections is a registered trademark exclusively owned by Louisiana Healthcare Connections.