

PRULAREN

Prucalopride 1 mg & 2 mg Tablets

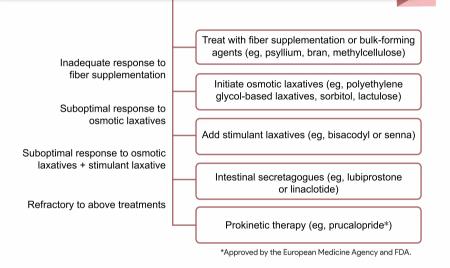
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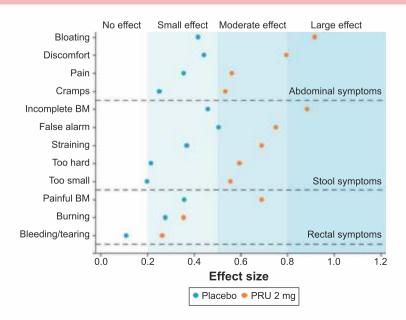
BACKGROUND:

- Chronic idiopathic constipation (CIC) is the common GI disorder. In general population, the prevalence of constipation worldwide ranges from 0.7 to 79% (median 16%).
- Prucalopride, a selective serotonin receptor agonist, treats chronic constipation in adults unresponsive to laxatives. It's high selectivity for the 5-HT4 receptor and potent enterokinetic properties make it more effective than traditional laxatives.²

THERAPEUTIC APPROACH FOR CHRONIC CONSTIPATION³



IMPACT OF PRUCALOPRIDE ON CHRONIC CONSTIPATION SYMPTOMS³





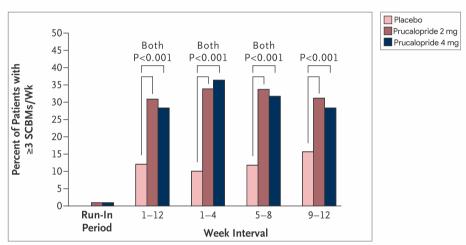
CLINICAL EVIDENCE:

1. A PLACEBO-CONTROLLED TRIAL OF PRUCALOPRIDE FOR SEVERE CHRONIC CONSTIPATION. 4

Patients: 620 patients (Prucalopride 2 mg: 207, Prucalopride 4 mg: 204, Placebo: 209)

Dose: Prucalopride 2 mg or 4 mg & Placebo Duration: 12 weeks

Result: Increase in bowel movements over 12 weeks (P<0.001)



Duration: 4 weeks

Fig. The Primary Efficacy End Point in the Intention-to-Treat Population.

CONCLUSION: Over 12 weeks, prucalopride significantly improved bowel function and reduced the severity of symptoms in patients with severe chronic constipation.

2. EFFICACY AND SAFETY OF PRUCALOPRIDE IN PATIENTS WITH CHRONIC NONCANCER PAIN SUFFERING FROM OPIOID-INDUCED CONSTIPATION.⁵

Patients: 196 patients

Result:

Dose: Prucalopride 2 mg or 4 mg & Placebo

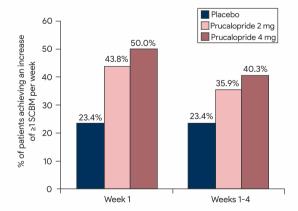


Fig. Effect of Prucalopride on the Primary Efficacy End Point

CONCLUSION: Prucalopride showed effectiveness in improving bowel movement frequency and reducing symptoms, with a favourable safety profile in patients with opioid-induced constipation.

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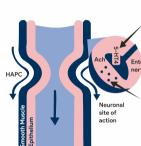
DESCRIPTION:

Prularen-1/2 contains prucalopride and is available as a tablet, in strengths of 1 mg and 2 mg. Prucalopride is a highly selective, high affinity 5-HT4 receptor agonist that activates both cholinergic and non-adrenergic, non-cholinergic neurotransmission.

INDICATION:

Indicated in the treatment of chronic idiopathic constipation (CIC).

MECHANISM OF ACTION:



Activate: Prucalopride selectively binds and activates 5-HT4 receptors

Release: Activation of colonic 5-HT4 receptors triggers the release of acetylcholine.

Move: Resulting in increased colonic peristalsis and enhanced bowel motility.

KEY BENEFITS OF PRUCALOPRIDE:

- → Superior Efficacy: Prucalopride is more effective than traditional laxatives in increasing stool frequency and improving consistency.2,6
- → Targeted Action: It selectively stimulates the 5-HT4 receptor in the gut, leading to efficient bowel movement stimulation.²
- Low Interaction Risk: With minimal protein binding, it poses a lower risk of drug interactions.
- Rapid Relief: Prucalopride achieves peak levels (2-3 hours) quickly after administration, providing fast relief from constipation.7

DOSAGE:

The recommended dose of prucalopride is 2 mg-4 mg once a day or as directed by the physician.

PRESENTATION:

Prularen 1 mg & 2 mg is available as a strip of 10 tablets.

References:

- 1. Approved by the European Medicine Agency and FDA
- Alfiment Pharmacol Ther 2010; 32: 1113–1123
 3. © 2013 The Authors. Neurogastroenterology & Motility published by John wiley & Sons Ltd. 60 Abbreviations
 4. n engl j med 358;22, May 29, 2008
- 5. Dig Dis Sci (2010) 55:2912–2921 6. https://doi.org/10.1080/17512433.2019.1620104
- 7. Motegrity (prucalopride) Prescribing Information. Lexington, MA: Takeda Pharmaceuticals America, Inc.

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