

Educational needs related to irritable bowel syndrome with constipation (IBS-C) management across disciplines: a comparison of nurse practitioners, physician assistants, and physicians

Emily Belcher¹, Gregory D. Salinas¹,
Brandon Coleman¹, David Rosenbaum²

1. CE Outcomes, LLC; Birmingham, AL, USA
2. Ardelyx, Inc; Waltham, MA, USA



Introduction and purpose

Despite availability of multiple treatment options for irritable bowel syndrome with constipation (IBS-C), optimal management of the condition remains challenging. This study was designed to investigate the educational needs and practice differences amongst primary care and gastroenterology nurse practitioners (NPs), physician assistants (PAs), and physicians, and highlight areas for continuing education.

Methodology

- A case-based survey was developed with input from a gastroenterology specialist and pilot tested with the target audience. Survey instruments and protocols were determined to be exempt from review by an independent IRB.
- Surveys were fielded to gastroenterology and primary care physicians, NPs and PAs currently practicing in the United States from January to February 2024 using national mailing lists and lists of clinicians who have previously opted-in for similar educational research.
- HCPs had to see patients with IBS-C to be included in the results. To determine differences in management and perceptions by clinical group, subanalyses were conducted.

HCP sample demographics

Responses from 410 HCPs, including 63 NP/PAs were analyzed.	Primary care provider physicians (PCPs) (n = 222)	Primary care provider NPPAs (PCP NPPAs) (n = 34)	Gastro physicians (GIs) (n = 125)	Gastro NPPAs (GI NPPAs) (n = 29)
Patients seen per week, mean (SD)	109 (56.8)	87 (48.8)	84 (45.5)	62 (36.8)
% in an academic practice setting	15%	3%	31%	24%
Years in practice, mean (SD)	23 (9.3)	18 (6.3)	22 (9.9)	12 (5.9)
Patients with IBS per month, mean	34 (47.0)	23 (18.1)	63 (49.8)	56 (36.0)
% with IBS-C predominant	42%	38%	48%	45%

Diagnostic testing

Patient Case 1 Summarized:

- Previously healthy 25-year-old with 6 months of abdominal pain 4-5 days per week and pain improvement after defecation
- Several days between bowel movements, hard and pellet-like stools. No family history of IBD, colorectal cancer, or celiac disease
- Healthy diet, unpredictable mealtimes, walks 3-4 times per week. Feels stressed but not increased.
- No weight loss, rectal bleeding, or nocturnal symptoms reported.

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**exclusive answer option*

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Anorectal manometry	0%	0%	0%	0%	0%	0%	0%
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Abdominal							

IBSRELA® (tenapanor) is indicated for treatment of irritable bowel syndrome with constipation (IBS-C) in adults

Important Safety Information

WARNING: RISK OF SERIOUS DEHYDRATION IN PEDIATRIC PATIENTS

- **IBSRELA is contraindicated in patients less than 6 years of age; in nonclinical studies in young juvenile rats, administration of tenapanor caused deaths presumed to be due to dehydration. *[see PI Contraindications (4), Use in Specific Populations (8.4)]*.**
- **Avoid use of IBSRELA in patients 6 years to less than 12 years of age. *[see PI Warnings and Precautions (5.1), Use in Specific Populations (8.4)]*.**
- **The safety and effectiveness of IBSRELA have not been established in pediatric patients less than 18 years of age. *[see PI Use in Specific Populations (8.4)]*.**

CONTRAINDICATIONS

IBSRELA is contraindicated in patients less than 6 years of age due to the risk of serious dehydration.

IBSRELA is contraindicated in patients with known or suspected mechanical gastrointestinal obstruction.

WARNINGS AND PRECAUTIONS

Risk of Serious Dehydration in Pediatric Patients

IBSRELA is contraindicated in patients below 6 years of age. The safety and effectiveness of IBSRELA in patients less than 18 years of age have not been established. In young juvenile rats (less than 1 week old; approximate human age equivalent of less than 2 years of age), decreased body weight and deaths occurred, presumed to be due to dehydration, following oral administration of tenapanor. There are no data available in older juvenile rats (human age equivalent 2 years to less than 12 years).

Avoid the use of IBSRELA in patients 6 years to less than 12 years of age. Although there are no data in older juvenile rats, given the deaths in younger rats and the lack of clinical safety and efficacy data in pediatric patients, avoid the use of IBSRELA in patients 6 years to less than 12 years of age.

Diarrhea

Diarrhea was the most common adverse reaction in two randomized, double-blind, placebo-controlled trials of IBS-C. Severe diarrhea was reported in 2.5% of IBSRELA-treated patients. If severe diarrhea occurs, suspend dosing and rehydrate patient.

MOST COMMON ADVERSE REACTIONS

The most common adverse reactions in IBSRELA-treated patients (incidence ≥2% and greater than placebo) were: diarrhea (16% vs 4% placebo), abdominal distension (3% vs <1%), flatulence (3% vs 1%) and dizziness (2% vs <1%).

For additional safety information, including the Boxed Warning, please see full Prescribing Information: [click here](#).