

# Updates in Management of Irritable Bowel Syndrome

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## Functional GI Disorders (FGIDs)

- Chronic recurrent digestive symptoms with no structural abnormalities
- Common FGIDs
  - Functional heartburn
  - Functional dysphagia
  - Functional chest pain
  - Functional dyspepsia (FD)
  - Belching disorder
  - Functional abdominal pain
  - Functional constipation
  - Irritable bowel syndrome (IBS)
  - Functional defaecation disorder

#### **Rome IV Criteria for IBS**

Recurrent abdominal *pain*, on average, at least <u>1 day per</u> <u>week</u> in the last 3 months, associated with 2 or more of the following criteria:

- 1. Related to defecation
- 2. Associated with a change in frequency of stool
- 3. Associated with a change in form (appearance) of stool

Criteria fulfilled for the last 3 months with symptom onset at least 6 months before diagnosis.

Lacy et al. Gastroenterol 2016

#### **Rome IV Criteria for IBS in Children**

Abdominal pain at least 4 days per month associated with one or more of the following:

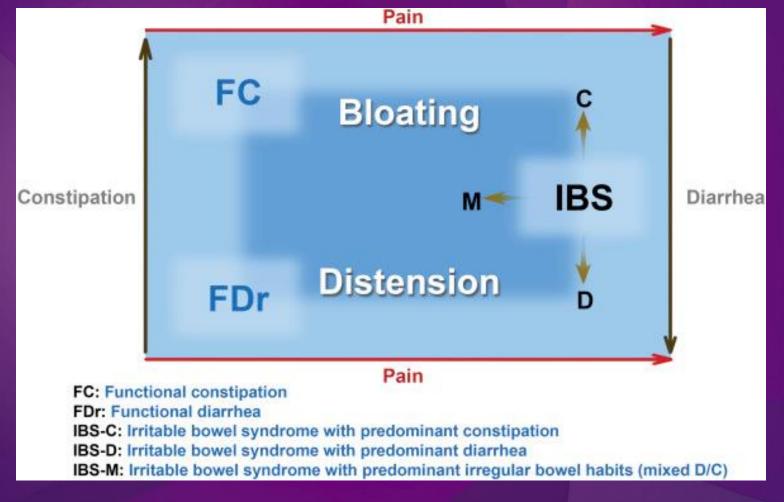
- 1. Related to defecation
- 2. A change in frequency of stool
- 3. A change in form (appearance) of stool

In children with constipation, the pain does not resolve with resolution of the constipation

Above criteria needs to be fulfilled for at least 2 months before diagnosis

Hyams et al. Gastroenterol 2016

# IBS and Functional Bowel Disorders are spectrum disorders differentiated by pain

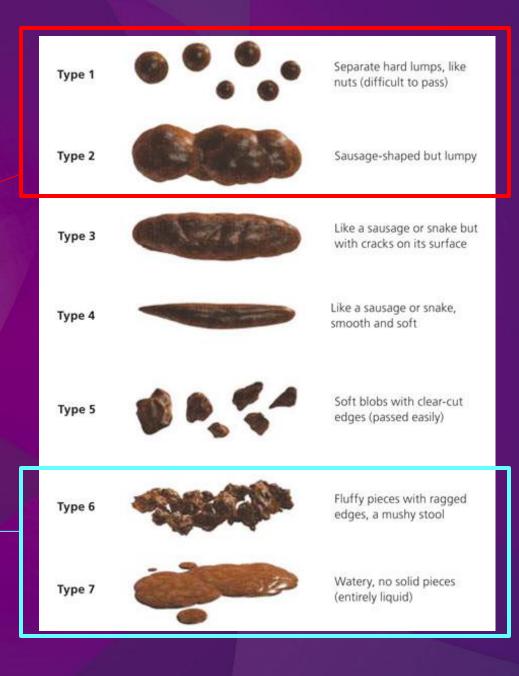


#### Lacy et al. Gastroenterol 2016

# IBS Subtyping by Stool Consistency

#### Constipation

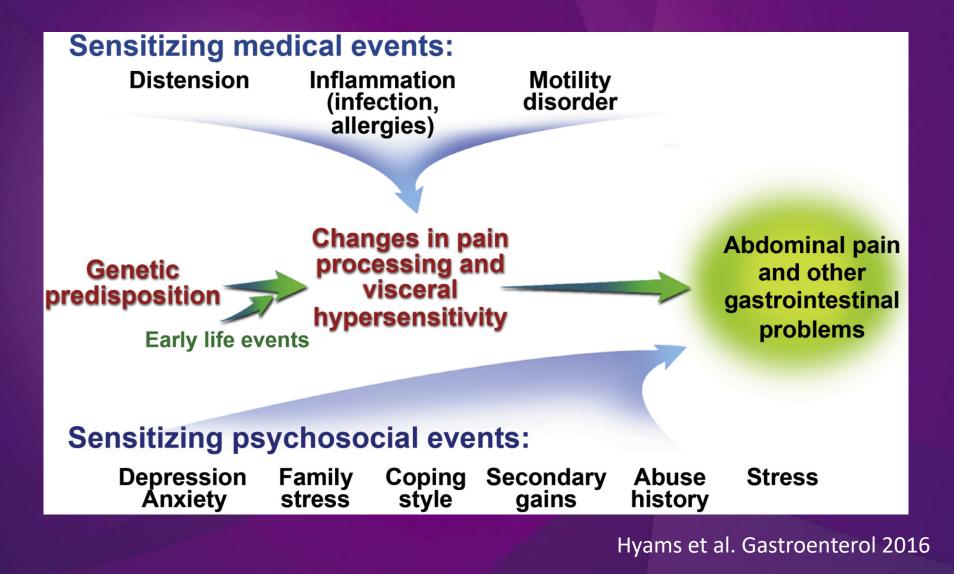
>25% Type 1 or 2: IBS-C
>25% Type 6 or 7: IBS-D
>25% in both: IBS-mixed
<25% in both: IBS-unclassified</li>



**Bristol Stool Form Scale** 

Diarrhea

#### Pathophysiology of IBS in Children



#### Important differential diagnoses in Asia

- Inflammatory bowel disease
- Cancers: colorectal, ovarian, endometrial
- Enteric infection and small intestinal bacterial overgrowth
- Metabolic: Thyrotoxicosis, hypercalcemia, etc.
- Gynecological condition: endometriosis, fibroid, pelvic inflammatory disease
- Medications: NSAID, metformin, etc.
- Ischemic bowel disease
- Celiac disease
- Heavy metal poisoning

## "Alarm features" in children

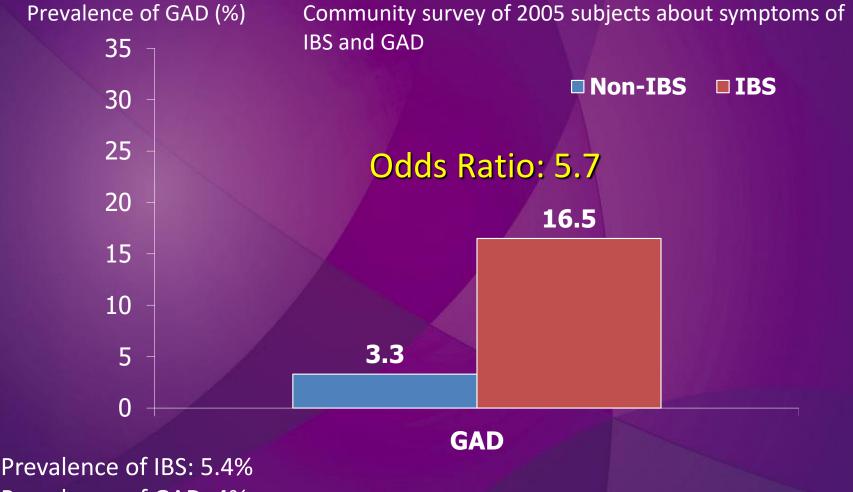
- Family history of inflammatory bowel disease, celiac disease or peptic ulcer disease
- Persistent right upper or lower abdominal pain
- Dysphagia and odynophagia
- Persistent vomiting
- Gastrointestinal blood loss
- Nocturnal diarrhea
- Peri-rectal disease

- Arthritis
- Unexplained fever
- Involuntary weight loss
- Deceleration of linear growth
- Delayed puberty

#### **Irritable Bowel or Irritable Brain?**



# IBS is associated with Generalized Anxiety Disorder (GAD) in the community

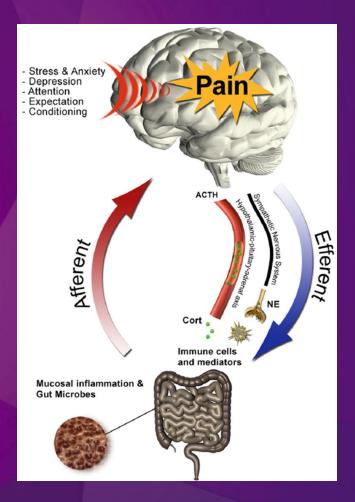


Prevalence of GAD: 4%

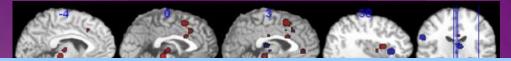
Lee et al. Aliment Pharmacol Ther 2009

#### **IBS: A Disorder of Gut-Brain Interaction (DGBI)**

- Motility dysfunction
- Visceral hypersensitivity
- Psychiatric comorbidity
- Neuroendocrine dysfunction
- Genetics and epigenetics
- Immune activation
- Dysbiosis
- Diet



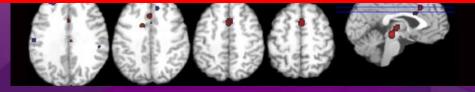
# Aberrant emotional arousal and central pain modulation in IBS



IBS: pregenual anterior cingulate cortex, amygdala (Emotional arousal) and midbrain cluster (Endogenous pain modulation)



No difference in visceral afferent processing (ie, thalamus, insula, anterior mid-cingulate)

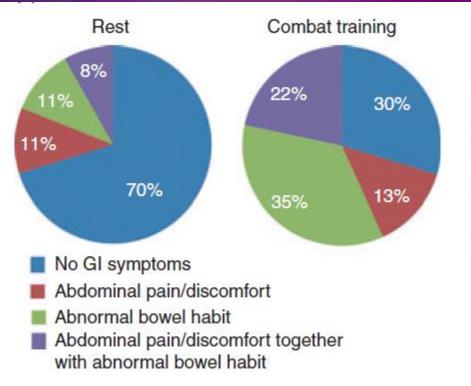


Quantitative meta-analysis of neuroimaging studies

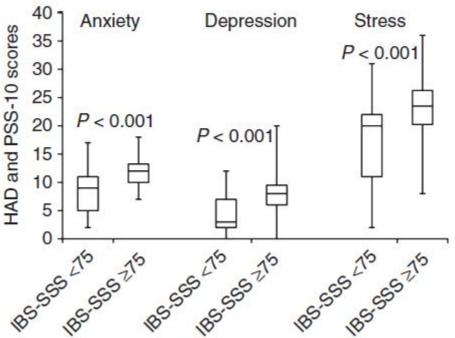
Tillisch et al. Gastroenterol 2011

# Psychological stress is the primary event of both immune activation and GI symptoms

#### 37 male soldiers underwent high-intensity combat training



HAD and PSS-10 scores in soldiers with IBS-SSS  $\geq$  75 or <75 during combat training



#### Li et al. Aliment Pharmacol Ther 2013

# "Hints" for psychiatric comorbidity

- Sleep disturbance
- Fleeting somatic pain syndromes
- Chronological association between stress / major life event and symptoms
- Significant functional impairment
- Significant avoidance behavior
- Health anxiety
  - Repeated investigations
  - Relentless search for health information

## Investigations

- Routine investigations are not recommended in patients aged<50 without alarm symptoms</li>
- Blood tests: Complete blood counts, ESR, C-reactive protein, thyroid function, IgA tissue Transglutaminase (tTG)
- Stool tests: Culture, microscopy, C. difficile enterotoxin, faecal calprotectin
- Imaging: Colonoscopy, CT abdomen and pelvis

#### **Management of IBS**

- Therapeutic relationship
- Dietary advice
- Medical therapy
- Psychological intervention

#### **IBS-associated Diet**

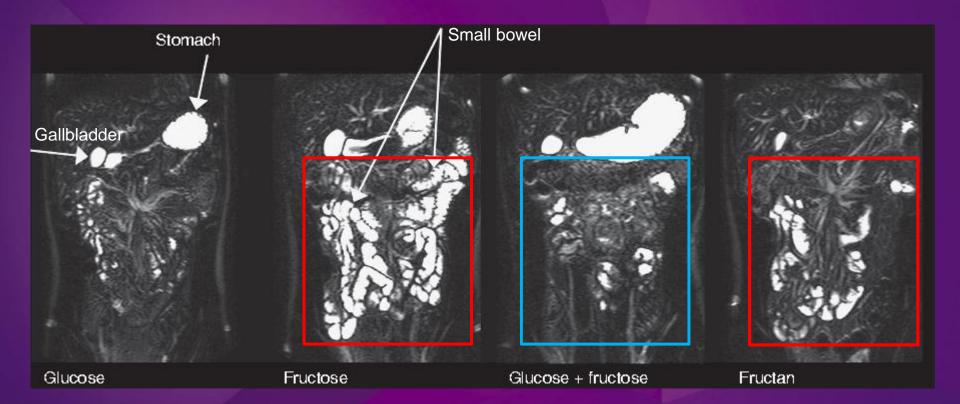


- FODMAP
- Sugar intolerance
- Fat
- Chili
- Gluten

#### Low FODMAP diet: Emerging 1<sup>st</sup> line treatment

- Fermentable Oligo-Di-Monosaccharides And Polyols
  - Fructose: Honey, watermelon
  - Lactose: Milk, ice-cream
  - Fructans: Wheat, garlic, onion
  - Galactans: Legumes, e.g. Beans, lentils, soybeans, cauliflower
  - Polyols: Sweeteners containing isomalt, mannitol, sorbitol, stone fruits (e.g. apples, apricots, cherries, peaches, plums)
- Low FODMAP diet improves IBS symptoms and quality of life
- ↓ Osmotic diarrhea, fermentation, altered gut microbiota, ↓ immune activation and visceral sensitivity
- $\downarrow$ Compliance to 50% in 8 weeks

#### "Sugar Intolerance" in IBS



MRI done 75 min after test drinks

Major et al. Gastroenterol 2017

## **Bulking agents and Antispasmodics**

	No. of studies	No. of patients	Relative risk (95% CI) of global improvement	P value
Bulking agents	11	565	1.10 (0.91-1.33)	0.32
Antispasmodics	22	1983	1.49 (1.25-1.77)	<0.0001

- Bulking agents or dietary fibre
  - Helps improve constipation
  - > No benefit or may worsen abdominal pain
- Antispasmodics
  - > NNT: 7 (pain), 5 (global improvement), 3 (symptom score)

Ruepert et al. Cochrane Review 2012; Bijkerk et al. Aliment Pharmacol Ther 2004

## Antispasmodic remains the 1<sup>st</sup> line Medical Treatment

- Anticholinergic antispasmodics (e.g. Hyoscine)
- Non-anticholinergic antispasmodics (e.g. Otilonium, mebeverine)
- Peppermint oil

## **Efficacy Vs Side effects of Antispasmodics**

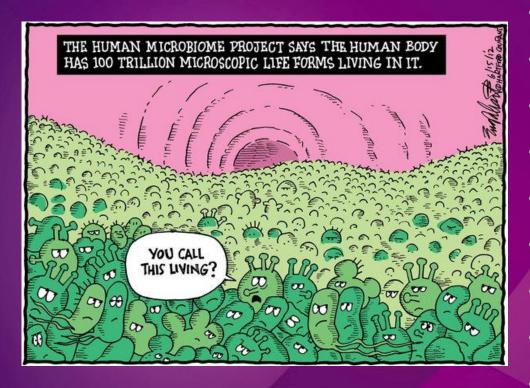
	No with syn abdominal pair				
Subcategory and study	Treatment group	Control group	Relative risk (random) (95% CI)	Weight (%)	Relative risk (random) (95% C
Cimetropium	10000 C				
Centonze 1988 <sup>w11</sup>	4/24	19/24 -		2.72	0.21 (0.08 to 0.5)
Passaretti 1989 <sup>w13</sup>	7/20	12/20		3.89	0.58 (0.29 to 1.1)
Dobrilla 1990 <sup>w12</sup>	4/35	11/35 —		2.25	0.36 (0.13 to 1.0
ubtotal (95% CI)	79	79		8.87	0.38 (0.20 to 0.7
yoscine					
Ritchie 1979 <sup>w33</sup>	8/12	12/12		6.38	0.67 (0.45 to 0.9
Nigam 1984 <sup>w35</sup>	11/21	21/21		oscine	0.52 (0.35 to 0.7
Schafer 1990 <sup>w10</sup>	44/182	64/178		JSLIFE	0.67 (0.49 to 0.9
ubtotal (95% CI)	215	211	→ 1	19.86	0.63 (0.51 to 0.7
naverium					
_evy 1977 <sup>w22</sup>	6/25	18/25 -		3.62	0.33 (0.16 to 0.7
Delmont 1981 <sup>w27</sup>	6/30	13/30		3.15	0.46 (0.20 to 1.0
Viral 1987 <sup>w24</sup>	14/39	26/39		5.66	0.54 (0.34 to 0.8)
ubtotal (95% CI)	94	94		12.44	0.47 (0.33 to 0.6
imebutine					
Moshal 1979 <sup>w15</sup>	3/10	4/10		1.78	0.75 (0.22 to 2.5
ielding 1980 <sup>w20</sup>	17/30	13/30		5.29	1.31 (0.78 to 2.1
bhidini 1986 <sup>w14</sup>	8/30	10/30		3.39	0.80 (0.37 to 1.7
ubtotal (95% CI)	70	70		10.46	1.08 (0.72 to 1.6
ebeverine					
(ruis 1986 <sup>w34</sup>	35/40	28/40		8.06	1.25 (0.99 to 1.5
ibtotal (95% CI)	40	40		8.06	1.25 (0.99 to 1.5
tilonium					
Arienzo 1980 <sup>w23</sup>	1/14	4/14 -		0.70	0.25 (0.03 to 1.9
Baldi 1983 <sup>w25</sup>	3/15	7/15 -		1.95	0.43 (0.14 to 1.3
astiglione 1991 <sup>w26</sup>	8/30	20/30	AL!	1.24	0.40 (0.21 to 0.7
ilende 2002 <sup>w16</sup>	99/157	124/160			0.81 (0.70 to 0.9
ubtotal (95% CI)	216	219			0.43 (0.14 to 1.3 0.40 (0.21 to 0.7 0.81 (0.70 to 0.9 0.55 (0.31 to 0.9
lverine					
Mitchell 2002 <sup>w18</sup>	26/53	31/54		6.82	0.85 (0.60 to 1.2
ubtotal (95% CI)	53	54		6.82	0.85 (0.60 to 1.2
cycloverine (dicyclomine)					
cyclovernie (arcyclonnie)	e				
mber of pa	tiont no	odod to		mber	of patie
					•
		4 0)	ne	ed to	harm: 1
treat: 5 (9	5% CI:	4-91			indirini .
	0 /0 011	• • • •			. 7 21)
ubtotal (95% CI)	39			5%0 CI	: 7-21)
ociverine					-
ahidini 1986 <sup>w14</sup> ubtotal (95% Cl)	11/30 30	10/30 30		3.93 3.93	1.10 (0.55 to 2.1 1.10 (0.55 to 2.1
otal (95% CI) otal events: 350 (treatment	905 ), 495 (control)	903		100.00	0.68 (0.57 to 0.8
est for heterogeneity: $\chi^2=52$ est for overall effect: $z=4.22$	8.88, df=22, P<0.00	01, I <sup>2</sup> =62.6% -0.1 -	0.2 -0.5 0 2 5	10	
est for overall effect: z=4.22	2, PR0.001	Favours	Favo	ours	

Favours treatment

#### Ford et al. BMJ 2008

control

## How Dysbiosis leads to IBS?



- P Fermentation
- Immune activation
- ↑ Intestinal permeability
- Visceral hypersensitivity
- Motility dysfunction
- Malabsorption
- Altered metabolome
- Mood regulation
- Epigenetic changes

- IBS-Diarrhea: ↓ Lactobacillus

Kassinen et al. Gastroenterol 2007; Chatterjee et al. Am J Gastroenterol 2007

#### **Probiotics: Positive results are consistent**

Probiotic	No. of studies	No. of subjects	Relative risk (95% C.I.) of persistent IBS symptoms
Combination (e.g. VSL#3)	12	609 (control: 588)	0.81 (0.67-0.98)
Lactobacillus	6	213 (control: 209)	0.75 (0.54-1.04)
Bifidobacterium	2	330 (control: 154)	0.71 (0.44-1.16)
Escherichia	2	208 (control: 210)	0.86 (0.79-0.93)
Streptococcus	1	32 (control: 22)	0.72 (0.53-0.99)
ALL	23	1392 (control: 1183)	0.79 (0.70-0.89)

Unresolved: mechanism, best regimen, optimum delivery mode, durability of efficacy

Ford et al. Am J Gastroenterol 2014

## **Rifaximin reduces bloating in IBS**

#### Poorly-absorbed, luminal-active antibiotic 5 Studies (N= 1,803)

Study ID	OR (95% CI)	% Weight	Study ID	OR (95% CI)	% Weight
Pimentel <i>et al,</i> Lembo <i>et al,</i>		5.16 25.83	Sharara <i>et al,</i> Pimentel <i>et al,</i> Lembo <i>et al,</i>	+ 3.70 (0.92, 14.89) + 4.83 (1.44, 16.18) + 1.39 (0.93, 2.07)	3.06 3.99 26.01
Target 1	1.62 (1.16, 2.27)	33.84	Target 1	1.52 (1.09, 2.11)	33.20
Target 2 Overall	1.48 (1.07, 2.05)           1.55 (1.23, 1.96)	35.17 100.00	Target 2 Overall	1.44 (1.04, 2.00)           1.57 (1.22, 2.01)	33.73 100.00
	1 2 3			১ ১ <sub>২</sub> ৬ ৫ ৬	

Bloating: OR = 1.55 (95 % CI = 1.23 – 1.96); therapeutic gain = 9.9% NNT = 10.1 Global improvement OR = 1.57 (95 % CI = 1.22 – 2.01); therapeutic gain = 9.8% NNT = 10.2

Age strongly predictive of response (correlation coefficient of 0.97)

Menees et al. Am J Gastroenterol. 2012

## **Antibiotics for IBS: Unresolved issues**

#### **Unclear mechanism**

 Animal studies: 

 mucosal inflammation and visceral hyperalgesia but microbiota data is lacking

#### **Unclear long-term safety**

- Good short-term safety profile: serious AEs (1.5% vs. 2.2%), discontinuation due to drug-related AEs (0.8% vs. 0.8%)
- Antibiotic resistance
- Clostridium difficile infection
- **Unclear long-term efficacy**
- Lack of long-term data or change in disease course

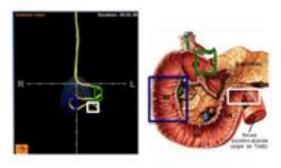
Xu et al. Gastroenterol 2014; Schoenfeld et al. Aliment Pharmacol Ther 2014

## FMT for non-constipation IBS with bloating

62 patients with refractory IBS symptoms and predominant abdominal bloating (ROME III criteria), aged 18-75 years and without constipation were randomly assigned (2:1) to transplantation with fresh donor stool or with placebo

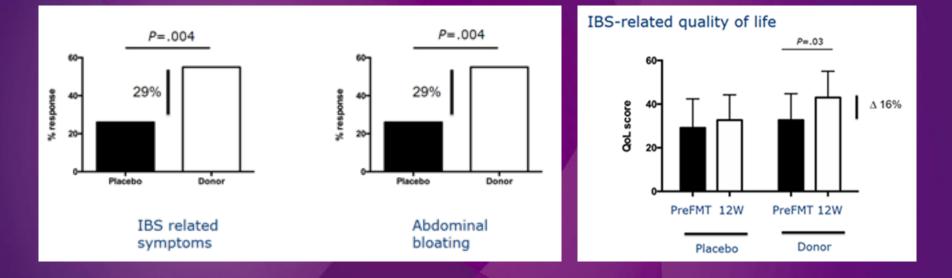
- FMT
  - Nasojejunal/duodenal administration
  - Cortrak<sup>™</sup> probe
- 2 healthy donors
  - Rich microbial diversity
  - Proven efficacy in pilot trial
  - Screened repeatedly for infectious diseases
  - Fresh stools
- Placebo
  - Patients' own frozen stool





#### Holvoet et al. Digestive Disease Week 2018

#### FMT reduces IBS symptoms and bloating



#### 27% responders in remission at 1 year

Holvoet et al. Digestive Disease Week 2018

## Serotonin (5-HT) receptor modulators

Class	Drug	IBS subtypes with efficacy reported	Side effects
5HT <sub>3</sub> antagonist	Alosetron	IBS-D	Complicated constipation, ischemic colitis
	Cilansetron	IBS-D	
	Ramosetron	IBS-D	
5HT <sub>4</sub> agonist	Tegaserod	IBS-C, IBS-mixed	Myocardial infarction, stroke
	Prucalopride	Constipation	
5HT <sub>4</sub> agonist/ 5HT <sub>3</sub> antagonist	Renzapride	IBS-C	Ischemic colitis

Camilleri et al. Lancet 2000; Kellow et al. Gut 2003; Camilleri et al. N Engl J Med 2008 Lembo et al. Aliment Pharmacol Ther 2010; Chey et al. Am J Gastroenterol 2008

# Selective Serotonin Reuptake Inhibitor (SSRI) is the antidepressant of choice for IBS

Meta-analysis: 230 non-anxiety/depression IBS patients randomized to SSRI Vs placebo in 5 RCTs (Mostly from referral centers)

Study or sub-category	Treatment n/N	Control n/N	RR (random) 95% Cl
02 selective serotonin re-upt	ake inhibitors		
Kuiken 2003 <sup>54</sup>	9/19	12/21	
Tabas 2004 <sup>55</sup>	25/44	36/46	-8-
Vahedi 2005 <sup>52</sup>	6/22	19/22	
Tack 2006 <sup>57</sup>	5/11	11/12	
Talley 2008 <sup>51</sup>	5/17	5/16	
Subtotal (95% CI)	113	117	◆
			0,1 0,2 0,5 1 2 5 10
			Favours treatment Favours control

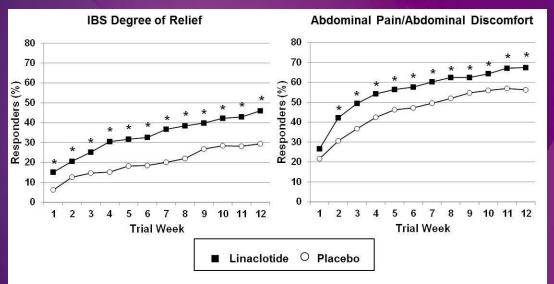
Relative risk of persistent IBS symptoms: 0.62 (95% CI, 0.45-0.87)

Number of patients needed to treat: 3.5 (95% CI: 2-14)

Ford et al. Gut 2009

# Secretagogue: Linaclotide (Guanylate cyclase C agonist) for IBS-C

# 839 IBS-C (Rome III) patients randomized to 12-week treatment with once-daily oral LIN 290 μg or placebo (PBO) (Phase 3 Trial)



Analysis uses the weekly responder criteria for each of the co-primary endpoints.

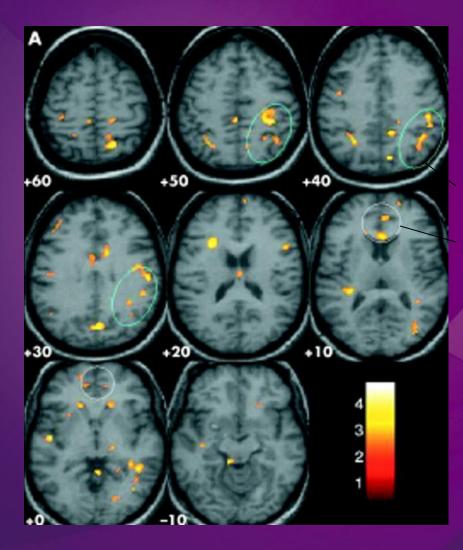
\* P<0.05; nominal P values based on a comparison of LIN vs PBO using a Cochran-Mantel-Haenszel test controlling for geographic region.

Table. Fillinary and Secondary Endpoint Results (111 Fopulation)					
	PBO (N=422)	LIN (N=417)	P value <sup>d</sup>		
Co-primary Endpoints (% Responders) <sup>a</sup>					
≥30% decrease in abdominal pain or discomfort, with neither score worsening from baseline, for ≥6/12 weeks	48.8	60.0	0.0010		
IBS degree of relief score ≤2 for ≥6/12 weeks <sup>b</sup>	15.4	31.7	<0.0001		
Secondary Endpoints (LS Mean Change from Baseline) <sup>c</sup>					
CSBMs/week	1.0	1.9	< 0.0001		
SBMs/week	1.5	3.0	< 0.0001		
Stool consistency (7-point BSFS)	0.8	1.5	<0.0001		
Straining (5-point ordinal scale)	-0.7	-1.0	<0.0001		
Abdominal bloating (11-point NRS)	-0.9	-1.5	<0.0001		
Abdominal pain (11-point NRS)	-1.1	-1.6	<0.0001		
Abdominal discomfort (11-point NRS)	-1.0	-1.5	<0.0001		

- Diarrhea reported in 9.4% of LIN patients Vs 1.2% of PBO patients.
- Low discontinuation rates due to diarrhea were low (0.7% LIN, 0.2% PBO).

Yang et al. J Gastroenterol Hepatol. 2018

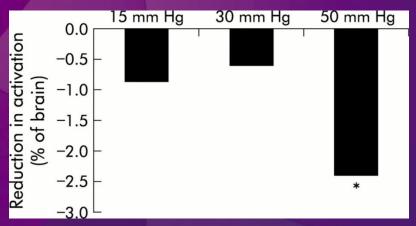
# Tricyclic antidepressant \$\frac{1}{2}\$ stressinduced visceral hypersensitivity



Net difference in brain activation between placebo and TCA during rectal distension under stress

Left posterior parietal cortex

#### **Perigenual anterior cingulate cortex**



#### Morgan et al. Gut 2005

## **Therapist-based psychotherapies are effective**

Psychotherapy	No. of studies	No. of subjects	Relative risk (95% C.I.) of persistent IBS symptoms
Cognitive behavioural therapy	9	349 (control: 261)	0.60 (0.44-0.83)
Dynamic psychotherapy	2	138 (control: 135)	0.60 (0.39-0.93)
Hypnotherapy	5	141 (control: 137)	0.74 (0.63-0.87)
Multi-component psychotherapy	5	168 (control: 167)	0.72 (0.62-0.83)
Mindfulness mediation training	1	36 (control: 39)	0.57 (0.32-1.01)
Self-administered CBT	3	73 (control: 71)	0.53 (0.17-1.66)
Internet-based CBT	2	71 (control: 69)	0.75 (0.48-1.17)
Relaxation training	6	133 (control: 122)	0.77 (0.57-1.04)
Stress management	2	59 (control: 39)	0.63 (0.19-2.08)
ALL		1232 (control: 1102)	0.68 (0.61-0.76)

Ford et al. Am J Gastroenterol 2014

## **Conflicting results for Chinese Medicine**

• Only individualized treatment group has maintained improvement at 14 weeks after completion of treatment

Bensoussan et al. JAMA 1998

 No difference in global or individual IBS symptoms, or quality of life for standardized herbal formula

Leung et al. Am J Gastroenterol 2006

• Acupuncture is not more effective than sham treatment

Manheimer et al. Am J Gastroenterol 2012

## Summary

- Clinical diagnosis based on symptom criteria
- Complex pathophysiology implies the need of multitarget treatment
- Antispasmodic remain the 1<sup>st</sup> line treatment
- Trial of probiotics is safe and may be effective
- Short-term efficacy for low-FODMAP diet and Rifaximin in selected patients
- Good efficacy for SSRIs as off-label use
- Role of new drugs yet to be defined