

## Relationship between esophageal clinical symptoms and manometry findings in patients with esophageal motility disorders: a cross-sectional study

Hashem FakhreYaseri<sup>\*1</sup>, Ali Mohammad FakhreYaseri<sup>2</sup>, Ali Baradaran Moghaddam<sup>3</sup>  
Seyed Kamran Soltani Arabshhi<sup>4</sup>

Received: 19 November 2014

Accepted: 6 June 2015

Published: 3 October 2015

### Abstract

**Background:** Manometry is the gold-standard diagnostic test for motility disorders in the esophagus. The development of high-resolution manometry catheters and software displays of manometry recordings in color-coded pressure plots have changed the diagnostic assessment of esophageal disease. The diagnostic value of particular esophageal clinical symptoms among patients suspected of esophageal motor disorders (EMDs) is still unknown. The aim of this study was to explore the sensitivity, specificity, and predictive accuracy of presenting esophageal symptoms between abnormal and normal esophageal manometry findings.

**Methods:** We conducted a cross-sectional study of 623 patients aged 11-80 years. Data were collected from clinical examinations as well as patient questionnaires. The sensitivity, specificity, and accuracy were calculated after high-resolution manometry plots were reviewed according to the most recent Chicago Criteria.

**Results:** The clinical symptoms were not sensitive enough to discriminate between EMDs. Nevertheless, dysphagia, noncardiac chest pain, hoarseness, vomiting, and weight loss had high specificity and high accuracy to distinguish EMDs from normal findings. Regurgitation and heartburn did not have good accuracy for the diagnosis of EMDs.

**Conclusion:** Clinical symptoms are not reliable enough to discriminate between EMDs. Clinical symptoms can, however, discriminate between normal findings and EMDs, especially achalasia.

**Keywords:** Esophageal motility disorders, Gastroesophageal reflux, Achalasia.

*Cite this article as:* FakhreYaseri H, FakhreYaseri AM, Baradaran Moghaddam A, Soltani Arabshhi SK. Relationship between esophageal clinical symptoms and manometry findings in patients with esophageal motility disorders: a cross-sectional study. *Med J Islam Repub Iran* 2015 (3 October). Vol. 29:271.

### Introduction

The esophageal motor function is evaluated using a variety of techniques, including barium radiography, radionuclide transit studies, manometry with or without impedance testing, and more recently impedance planimetry (1). High-resolution esophageal pressure topography is a new technology based on a combination of high-resolution manometry and esophageal pres-

sure topography (EPT) for the examination of esophageal motor dysfunctions (EMDs). EPT plots are color-coded pressure representations on a spatiotemporal field generated by sophisticated software-based algorithms for visualizing and analyzing manometric data, the most recent classification scheme of which is intended to diagnosis of primary EMDs (2). The first step of the Chicago Classification described abnormal

<sup>1</sup>. (Corresponding author) Assistant Professor, Internist, Gastroenterologist, Research Center for Gastroenterology and Liver Disease, Department of Internal Medicine and Gastroenterology, Firoozgar Hospital, Iran University of Medical Sciences, Tehran, Iran. [hfyaseri@yahoo.com](mailto:hfyaseri@yahoo.com)

<sup>2</sup>. General physician, School of Medicine, Tehran University of Medical Sciences, Tehran, Iran. [fyasseri@mihanmail.ir](mailto:fyasseri@mihanmail.ir)

<sup>3</sup>. MSc in Clinical Microbiology, Department of Microbiology, School of Medicine, Iran University of Medical Sciences, Tehran, Iran. [ak.baradaran@gmail.com](mailto:ak.baradaran@gmail.com)

<sup>4</sup>. Professor, Internist, Department of Internal Medicine, Firoozgar Hospital, Iran University of Medical Sciences, Tehran, Iran. [soltarab34@gmail.com](mailto:soltarab34@gmail.com)

esophagogastric junction deglutitive relaxation in terms of an eSleeve 4-second nadir pressure. The latest classification system, proposed by Pandolfino et al, includes contraction patterns and peristalsis integrity based on integrated relaxation pressure 4 (IRP4) (3). The esophagus is a hollow muscular tube coursing through the posterior mediastinum joining the hypopharynx to the stomach with a sphincter at each end. The lower esophageal sphincter (LES) and the distal one half to two-thirds of the esophageal body are composed of smooth muscle. The distal esophagus and the LES are controlled by excitatory (cholinergic) and inhibitory (nitric oxide) myenteric plexus neurons (4). The major esophageal function is the transition of food and fluid between these two ends; it, otherwise, remains empty. Esophageal peristalsis results from the sequential contraction of circular muscles, which serves to push the ingested food bolus toward the stomach with minimal stasis in the esophageal body. Therefore, esophageal motility testing aims to investigate the esophageal function and to reveal any disorders to explain individual symptoms and provide a rationale for treatment. The diagnostic value of particular esophageal symptoms among patients with suspected esophageal motility abnormality is still unknown (5), although Mikieli reported there is a good correlation between the intensity of the motor abnormality and symptom severity (6). It has been suggested that patients with achalasia are at risk for the development of dysplasia and neoplasia, especially in patients with long-standing disease (7). The major esophageal symptoms are dysphagia, non-cardiac chest pain, regurgitation, heartburn, hoarseness, vomiting, and weight loss (5,8).

The aim of this study was to determine the sensitivity, specificity, and predictive accuracy of esophageal clinical symptoms in the diagnosis of variant types of EMDs in Iranian patients.

## Methods

We conducted a cross-sectional study of

patients with upper gastrointestinal symptoms (persist for > 8 weeks) from September 2012 to September 2014. The study population consisted of 623 patients (range 11–80 years old). Patients with dysphagia, noncardiac chest pain, regurgitation, heartburn, hoarseness, vomiting, and weight loss (>5kg) were included. The exclusion criteria were comprised of history of malignant disease, previous foregut surgery, cardiovascular diseases, large hiatal hernia, esophagitis of grade C or D according to the Los Angeles Classification, eosinophilic esophagitis, and Barrett's esophagus on pathology. All the patients provided informed consent and accepted to complete a standard questionnaire form. Esophagogas-

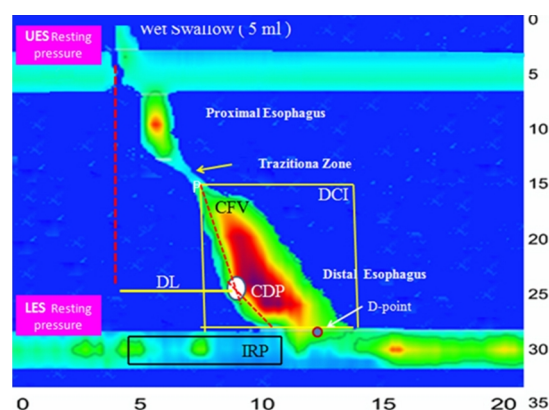


Fig. 1. A sample of the esophageal pressure topography of a swallow spanning the entire esophagus, from the pharynx to the stomach, of a normal subject with normal peristalsis and abnormal esophagogastric junction relaxation. Time is on the X-axis and distance from the nares is on the Y-axis. A normal Clouse plot indicates the locations of the upper esophageal sphincter, lower esophageal sphincter, and the two muscle contraction segments, proximal and distal. Integrated relaxation pressure 4 (IRP4) is the lowest mean abnormal esophagogastric junction pressure for four contiguous or non-contiguous seconds of relaxation. The contractile deceleration point (CDP) represents the inflexion point in the contractile front propagation. It is localized by fitting two tangential lines to the initial and terminal portions of the 30-mmHg isobaric contours and noting the intersection of the lines (red dot). The contractile front velocity (CFV) is calculated by taking the best-fit tangent of the 30-mmHg isobaric contour between P and CDP. The distal latency (DL) is measured from the upper esophageal sphincter relaxation to the CDP. The distal contractile integral is calculated by amplitude  $\times$  duration  $\times$  length (mmHg-s-cm) of the distal esophageal contraction greater than 20 mmHg from the proximal (P) to distal (D) pressure troughs.

trooduodenoscopy was done for all the patients in the same center by expert endoscopists. The procedures were performed by a trained esophageal laboratory nurse in collaboration with an expert gastroenterologist. Before each procedure, transducers were calibrated to 0 and 100 mmHg using externally applied pressure. The studies were conducted with the patient in the supine position after at least a 6-hour fast, and medications that could affect the esophageal motor function (e.g. Metoclopramide, anticholinergics, and smooth muscle relaxants) were discontinued for 5–7 days prior to the study. The catheter used was a 23-channel silicone-customized water-perfused catheter, with an outside diameter of 3.8 mm (manufactured by Mui Scientific, Ontario, CA). The catheters had 1 distal channel for gastric recording, 5 channels 1 cm apart for the LES pressure, and 16 proximal channels each 2 cm apart. Microlumina was perfused with a pneumohydraulic perfusion system (MMS software) at a water perfusion rate of 0.15 ml /min. Pressure data were acquired and shown using software specially designed for high-resolution manometry (MMS v 8.23), which displays isobaric contour plots. After topical anesthetic was applied into the nostril, the high-resolution manometry assembly was passed trans-nasally and the sensors were positioned to record from the hypopharynx to the stomach. After the LES was detected via the stationary pull-through method, the catheter was fixed in place by taping it to the nose. Then, 10 swallows of 5 mL ambient-temperature water spaced more than 20s apart were recorded. The pressure topography metrics utilized in the Chicago Classification is depicted in Figures 1 and 2. The definition based on this classification is shown in Table 1.

The data were entered into SPSS v.18 after encoding for each subject. Age is reported with mean  $\pm$  standard deviation. Hypercontractile esophagus and absent peristalsis were excluded in the statistical analysis due to rare findings in this study. The clinical symptoms and the results of

Table 1. Definitions of the contractile pattern based on the Chicago Classification

Contractile Pattern	Code	Contractile Pattern Definition
Absent peristalsis	AP	100% failed peristalsis with minimal (<3 cm) integrity of the 20 mmHg IBC* distal to the proximal pressure trough (P)
Frequent failed peristalsis	FFP	> 3 but <10 swallows with failed peristalsis
Panesophageal pressurization	PP	$\geq$ 20% of swallows with uniform pressurization of 30 mmHg IBC from the UES to the EGJ
Premature contraction	PC	$\geq$ 20% of swallows with DL < 4.5 s
Jackhammer	JH	Swallow with DL > 4.5 s and DCI > 8000 mmHg.s.cm
Rapid contraction	RC	$\geq$ 20% of swallows with contractile front velocity (CFV) > 9 cm/s and DL > 4.5 s
Hypertensive	HT	Mean DCI > 5000 but no swallow with value > 8000 mmHg.s.cm
Weak peristalsis	WP	> 20% swallows with large breaks in the 20 mmHg IBC (> 5 cm in length) or > 30% swallows with small breaks in the 20 mmHg IBC (2–5 cm in length)
Normal peristalsis	NP	$\geq$ 60% of swallows with an intact 20 mmHg IBC (or no break > 2 cm) not meeting any other code

\*IBC, Isobaric contour

the esophageal manometry findings (normal and abnormal) were compared, and the sensitivity [true positives/(true positive + false negative)], specificity [true negatives/(true negative + false positives)], and accuracy [(true positive + true negative)/(true positive + true negative + false positive + false negative)] of the clinical symptoms were calculated for a correct diagnosis of the manometry findings. The results are presented with a confidence interval (CI) of 95%. Comparison of statistical significance was made between the symptom categories and manometry findings using either the Chi-square test or the Fisher's exact probability test. A *P* value less than 0.05 was considered statistically significant.



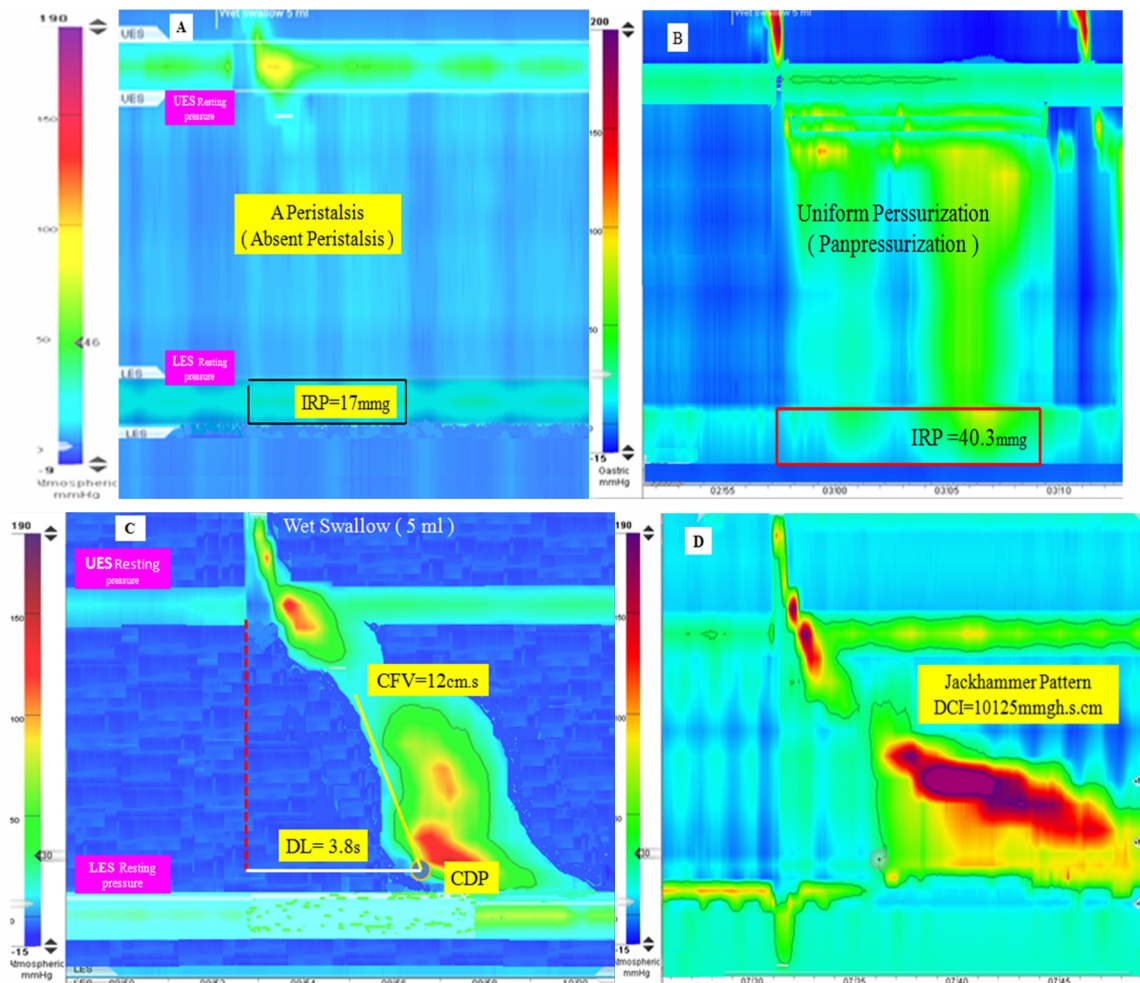


Fig. 2. EPT plots of the four types of manometric findings: (A) type 1 achalasia: all swallows with failed peristalsis, IRP4 (showed IRP)= 17 mmHg; (B) type 2 achalasia with uniform pressurization seen in the 40 mmHg isobaric contour, IRP4 =40.3 mmHg; (C) type 3 achalasia: premature contraction as demonstrate by DL=3.8s, CFV=12 cm.s, and IRP4=31 mmHg; and (D) hypercontractile esophagus: defined by DCI  $\geq$ 8000 mmHg.s.cm. IRP4, Integrated relaxation pressure 4; CFV, Contractile front velocity; DCI, Distal contractile integration

## Results

This study was performed on 623 patients, who met our inclusion criteria. The mean $\pm$ SD age of the patients was 40.2  $\pm$  12.9 (10-80) years, and 60.5% (n=377) of the patients were female. The dominant presenting symptoms were heartburn (66.7%) and regurgitation (63.7%). Almost all the patients had one to three of the symptoms as part of their initial presentation. The mean IRP4 was 17.6 $\pm$ 5.4 (range = -1 to 42) (Table 2). The prevalence rates of achalasia, normal peristalsis, and borderline motor function were 13.6% (85/623), 41.6% (212/510), and 40.8% (208/510), respectively (Fig. 3).

Tables 3 and 4 compare the sensitivity, specificity, and accuracy of the clinical

symptoms with the manometry findings. The prevalence rates of dysphasia (range = 39.4%-97.6%), noncardiac chest pain (range = 51.5%-87.8%), and heartburn (range = 61%-82.9%) were high but sensitivity was low for all the symptoms (range = 6.6%-66.6%) in all the types of abnormal manometry findings.

All the symptoms were highly specific (range = 43.6%-99.3%) for the diagnosis of patients with EMDs. Dysphasia (range= 55.2%-75.1%), noncardiac chest pain (range= 66%-72%), hoarseness (range= 49.7%-82.1%), asthma (range= 53.8%-76.7%), and weight loss (range= 52.1%-89.7%) had good accuracy, whereas regurgitation (range= 35.6%-50.7%) and heartburn (range= 39.5%-56.4%) had low accu-

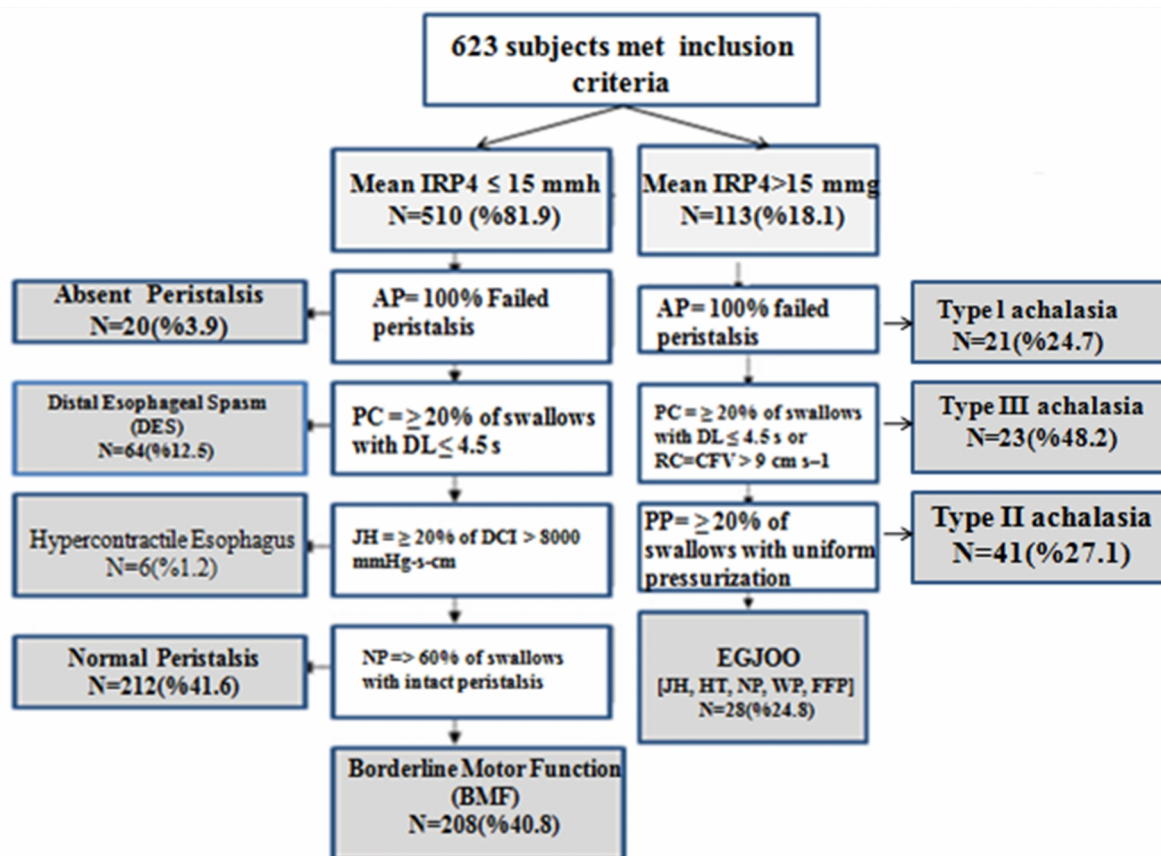


Fig. 3. Algorithm of the study analysis based on the Chicago Classification  
 IRP4, Integrated relaxation pressure 4; AP, Absent peristalsis; PC, Premature contractions; DL, Distal latency; RC, Rapid contraction; CFV, Contractile front velocity; JH, Jackhammer; DCI, Distal contractile integral; PP, Panesophageal pressurization; NP, Normal peristalsis; HT, Hypertensive peristalsis; WP, Weak peristalsis; FFP, Frequent failed peristalsis

racy for the diagnosis of patients with EMDs.

### Discussion

Normal peristalsis and borderline motor function (BMF) constituted the most preva-

lent pattern in this study (Fig. 3), while hypercontractile esophagus and absent peristalsis were rare patterns. The females accounted for the highest frequency of clinical presentations in all types of EMDs. With the exception of BMF, absent

Table 2. Demographic information of patients with variant types of esophageal motility disorder in our center for two years

Findings	N(%)	Age (years)		Sex		IRP4 (mmg)	
		Mean±SD	Range	Female (%)	p	(Mean± SD)	Range
IRP>15	113(18.1)	39.1±13.1	16-80	71(63)	-	29.6±6.7	17-42
Achalasia	85(75.2)	39.3±15.5	21-80	54(63.2)	0.000	29.8±6.8	17-42
Type I	21 (24.7)	38.6±8.2	25-59	12(57)	0.005	29.5±7.9	17-41
Type II	41(48.2)	40.5±15	20-80	26(63)	0.000	29.6±7	17-42
Type III	23(27.1)	39.3±15.5	21-75	16(69.5)	0.002	30.3±5.4	22-40
EGJOO	28(24.8)	38.2±13.7	16-80	17(61)	0.001	28.9±6.5	19-41
IRP<15	510(81.9)	41.2±12.8	10-73	306(60)	-	5.6±4.1	-1-14
NP	212(41.6)	41±21.5	11-72	136(64.1)	0.206	4.4±4	-1-13
DES	64(12.5)	41.2±15.7	10-73	39(61)	0.000	7.7±3.1	0-14
AP	20(3.9)	42.7±7.3	30-55	12(60)	0.999	5.6±4	0-12
HE	6(1.2)	35±3.3	32-51	2(33.3)	0.999	6.5±6	1-12
BMF	208(40.8)	46±16.4	23-68	117(56.2)	0.204	4±3.2	0-13
Total	623	40.2±12.9	10-80	377(60.5)	-	17.6±5.4	-1-42

IRP4, Integrated relaxation pressure 4; EGJOO, Esophagogastric junction outflow obstruction; NP, Normal peristalsis; DES, Distal esophageal spasm; AP, Absent peristalsis; HE, Hypercontractile esophagus; BMF, Borderline motor function

Table 3. Relationship between the clinical symptoms and the esophageal manometric findings in the patients with IRP4 >15 mmg

Symptoms	NP N=212		Achalasia(N=85)											EGJOO (N=28)			
			Type 1 (N=21, 24.7%)			Type 2 (N=41, 48.2%)			Type 3 (N=23, 27.1%)								
	N (%)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95 %CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)
Dysphagia	62 (29.2)	20 (9.52)	24.4 (22.8-25.8)	99.3 (97.5-99.5)	73 (68.7-77.5)	40 (97.6)	39.2 (37.4-41.7)	99.3 (99.2-100)	75.1 (70.8-79.8)	22 (9.56)	26.2 (24.5-27.6)	99.3 (98.3-100)	73.2 (68.7-77.5)	23 (82.1)	27 (25.5-28.8)	96.8 (94.6-98.5)	72.1 (69.4-75.1)
NCCP	66 (31.1)	18 (8.57)	21.4 (20.5-22.2)	98 (91.8-99.5)	70.4 (66-74.4)	36 (87.8)	35.3 (33.1-37.3)	96.7 (94.6-98.6)	72 (67.3-76)	18 (8.2)	21.4 (20.1-22.6)	86.4 (82.9-89.8)	69.8 (65.3-73.7)	16 (57.1)	19.5 (18.7-20.3)	92.4 (86.5-95.6)	67.5 (63.4-71.5)
Regurgitation	138 (65.1)	9 (4.28)	6.1 (5.7-6.4)	86 (82.2-89.1)	35.6 (33.4-38.7)	22 (53.6)	13.7 (13.2-14.2)	79.6 (76.7-83)	38 (35.9-40.4)	18 (8.2)	11.5 (11-12)	93.7 (90-97.5)	39.1 (36.6-41.2)	17 (60.7)	11 (10.5-11.3)	87 (83.1-90)	38 (35.8-40.4)
Heartburn	131 (62)	11 (5.24)	7.7 (7.2-8.2)	89 (85.6-92.7)	39.5 (37.2-41.9)	34 (82.9)	20.6 (19.7-21.3)	92 (88.2-95.6)	45.4 (42.5-47.9)	15 (6.52)	10.3 (9.9-10.7)	91 (87.3-94.6)	40.8 (38.5-43.4)	23 (82.1)	15 (14.3-15.5)	94 (91.8-95.6)	43.3 (40.7-45.9)
Hoarseness*	20 (9.4)	6 (2.86)	23.1 (21.7-24.5)	92.7 (89.1-96.5)	85 (81.4-88.2)	7 (17)	25.9 (24.3-27.4)	84.9 (81.4-88.2)	78.6 (75.2-81.4)	0 (0)	-	89.3 (85.6-92.7)	81.7 (78.2-84.8)	5 (17.8)	20 (19.1-20.7)	89.3 (85.6-92.7)	82.1 (78.2-84.7)
Asthma	14 (6.6)	4 (1.9)	22.2 (21.3-23.1)	92.1 (90-93.7)	86.7 (83.1-90)	3 (7.3)	17.6 (16.8-18.2)	83.9 (80.5-87.3)	79.4 (75.9-82.2)	0 (0)	-	89.6 (85.6-92.7)	84.2 (80.6-87.3)	5 (17.8)	26.3 (24.5-27.7)	89.6 (85.6-92.7)	84.5 (80.6-87.3)
Vomiting	72 (33.9)	9 (4.28)	11.1 (10.7-11.6)	92.1 (88.2-95.6)	64 (60-68)	29 (70.7)	28.7 (26.8-30.3)	92.1 (88.2-95.6)	66.8 (62.8-70.8)	17 (73.9)	19.1 (18.3-19.9)	95.9 (93.7-97.5)	66.8 (62.8-70.8)	10 (35.7)	12.2 (11.7-12.7)	88.6 (84.8-91.8)	62.5 (58.8-66.3)
Weight loss	14 (6.6)	11 (5.24)	44 (41.2-46.5)	95.2 (92.7-96.5)	89.7 (85.6-92.7)	26 (63.4)	65 (60.9-68.7)	92.9 (89.1-96.5)	88.5 (84.7-91.8)	1 (4.3)	6.6 (6.2-7)	90 (86.4-93.6)	84.7 (81.4-88.2)	3 (10.7)	17.6 (16.8-18.2)	88.8 (84.8-91.8)	83.7 (79.8-86.4)

NP, Normal peristalsis; EGJOO, Esophago gastric junction outflow obstruction; NCCP, Noncardiac chest pain-- \*Hoarseness, permanent throat clearing, post-nasal drip.

Table 4. Relation between the clinical symptoms and the esophageal manometric findings in the patients with IRP4 <15 mmg

Symptoms	NP N=212		DES (N=64)			AP (N=20)			HE (N=6)			BMF (N=208)					
	N (%)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)	N (%)	Sensitivity (95% CI)	Specificity (95%CI)	Accuracy (95% CI)
Dysphagia	62 (29.2)	43 (67.2)	41 (38.4-43.4)	87.7 (83.9-91)	69.9 (67.3-72.9)	15 (75)	19.5 (18.9-21.3)	96.8 (94.6-98.5)	71.1 (66.7-75.2)	3 (50)	4.6 (4.4-4.7)	2 (1.9-2.1)	70.2 (66-74.4)	82 (39.4)	57 (54.6-59.1)	54.3 (51.9-56.2)	55.2 (52.9-57.4)
NCCP	66 (31.1)	33 (51.5)	33 (31.2-35.2)	82.5 (78.2-84.8)	64.8 (62.2-67.3)	9 (45)	12 (11.5-12.4)	93 (81.5-99.5)	66.8 (62.8-70.8)	6 (100)	8.3 (8.1-8.4)	100 (95.6-103.5)	69.7 (65.3-73.7)	131 (63)	66.5 (64.1-69.4)	65.5 (64.7-68.7)	66 (63.4-68.7)
Regurgitation	138 (65.1)	48 (75)	25.8 (24.7-26.8)	82.2 (78.2-84.7)	44.2 (41.7-47)	8 (40)	5.5 (5.1-5.8)	86 (82.3-89.1)	35.5 (33.1-37.3)	3 (50)	2.1 (1.9-2.2)	96.1 (88.2-97.5)	35.3 (33.1-37.3)	139 (66.8)	50.2 (47.5-75.7)	51.7 (47.5-51.4)	50.7 (48.6-52.7)
Heartburn	131 (62)	39 (61)	23 (22-23.8)	76.4 (73-79)	43.5 (40.8-46)	6 (30)	4.4 (4.1-4.7)	85.2 (81.4-88.2)	37.5 (35.1-39.5)	6 (100)	4.4 (4.1-4.7)	0 (0)	40 (37.3-42)	156 (75)	54.3 (52-56.3)	61 (58.5-63.4)	56.4 (54-58.5)
Hoarseness	20 (9.4)	5 (7.8)	20 (19.1-20.7)	76.5 (73-79)	71.4 (68.5-74.3)	0 (0)	-	90.6 (86.5-93.7)	82.7 (79.4-86)	3 (50)	13 (12.4-13.5)	98.5 (97.5-99.5)	89.4 (85.6-92.7)	17 (8.2)	46 (44.2-48)	50.1 (48-60)	49.7 (47.4-51.4)
Asthma	14 (6.6)	4 (6.2)	22.2 (21.3-23.1)	76.7 (72.9-79)	73.2 (70.1-75.9)	0 (0)	-	91 (89.1-92.7)	85.3 (81.4-88.2)	0 (0)	-	97 (95.6-97.5)	90.8 (87.3-94.6)	28 (13.5)	66.6 (64.1-69.4)	52.4 (49.9-54)	53.8 (51.4-55.7)
Vomiting	72 (33.9)	20 (31.2)	21.7 (20.9-22.6)	76.1 (73-79)	57.9 (54-60.9)	0 (0)	-	87.5 (84-91)	60.3 (56.8-64.1)	0 (0)	-	95.9 (91.8-99.5)	64.2 (60.3-68)	41 (19.7)	36.3 (34.8-37.7)	45.6 (43.4-47)	43.1 (41.3-44.8)
Weight loss	14 (6.6)	5 (7.8)	26.3 (25.3-27.4)	77 (73.7-79.8)	73.5 (70.1-75.9)	0 (0)	-	91 (87.3-94.6)	85.3 (82.2-89.1)	0 (0)	-	97 (94.5-98.5)	90.8 (87.3-93.6)	21 (10.1)	60 (57.4-62.2)	43.6 (41.7-45.1)	52.1 (49.8-54)

DES, Distal esophageal spasm; AP, Absent peristalsis; HE, Hypercontractile esophagus; BMF, Borderline motor Function

\* Due to the low number of subjects

peristalsis, and hypercontractile esophagus, all types of manometric patterns were more prevalent in the male patients.

Dysphagia and non-cardiac chest pain (NCCP) were more prevalent in the patients with EMDs. Regurgitation and heartburn known as the diagnostic symptoms of gastroesophageal reflux disease (GERD) Hoarseness and asthma are known as the complications of GERD (10). We found that regurgitation was not a sensitive symptom for EMDs diagnosis. Although vomiting was prevalent in achalasia (types 2 and 3) and weight loss was prevalent in type 2 achalasia (6); these two symptoms lacked enough sensitivity for the differentiation between motility disorders among our study population.

The manometry is the gold-standard investigation of motility disorders in the esophagus. The transport of bolus is "successful" when minimal bolus material is retained within the esophageal body (11). The relationship between esophageal motility and transit is complex because factors such as bolus shape, surface, and consistency cannot be measured with manometry (12).

The LES relaxation does not seem to be only a major factor in determining bolus stasis. This may be justified by the observation that stasis most often occurs in the proximal and mid portions of the esophagus and that the bolus often does not reach the distal esophagus due to failed or incomplete peristalsis (12). However, the IRP4 value is also influenced by distal esophageal contractility (3). It has been suggested that IRP4 is the optimal measure of abnormal esophagogastric junction relaxation (6,13).

Dysphagia is usually mild in patients with ineffective esophageal motility (now called weak peristalsis) and abnormal esophageal propagation velocity (14). Although our study showed that only dysphagia and NCCP were more prevalent and highly specific in EMDs, their accuracy could not discriminate between the various types of EMDs. Regurgitation can be a problematic symptom inasmuch as it may lead to aspira-

tion (6). It has been suggested that regurgitation has a significant correlation with the LES relaxation pressure when compared to other individual symptoms (6). Heartburn is a symptom complex that has traditionally been accepted as an acid-mediated event and a reliable indicator of GERD. It may occur in other conditions such as stress or smoking (6). GERD is prevalent in asthma (10). In our study, regurgitation, heartburn, asthma, and weight loss lacked enough accuracy to distinguish EMDs.

Achalasia is to date the best described manometric abnormality with the most well-defined treatment options from all EMDs. In our study, the prevalence of achalasia was similar to chimed in with that previously reported in the literature (6). It has been posited that esophagogastric junction outflow obstruction (EGJOO) is due to the contraction of the crural diaphragm and might also represent an achalasia variant in some cases (4). In this study, the patients with EGJOO presented with dysphagia and chest pain (15). Distal esophageal spasm (DES) is an uncommon EMD. The distal latency seems to be a more reliable measure of premature contractions presenting with dysphagia and chest pain; however, if the reduced distal latency is associated with a high IRP, it is termed "spastic achalasia" (3). A review of 1070 consecutive interpretable EPT studies revealed that all 24 patients with reduced distal latency had a dominant symptom of dysphagia or chest pain and were diagnosed and managed as distal esophageal spasm or spastic achalasia (9). Rapid contraction is defined as an increased contractile front velocity (CFV) ( $>9\text{cm.s}$ ). However, although the CFV is a regional variability in contractile velocity, the correlation of symptoms with this "spastic" pattern is unclear (5). The previous Chicago Classification defined hypercontractile disorders in terms of mean distal contractile integration (DCI), a DCI greater than  $8000\text{ mmgh.cm.s}$  was called "hypercontractile (nicknamed "Jackhammer") esophagus". This pattern of EMDs is rare and reported only in between 3% and



4.1% of cases, and it is universally associated with dysphagia and/or chest pain (3,6,16).

In this study, the prevalence of hypercontractile esophagus was low and it presented with dysphagia and hoarseness. A wide range of motility disorders with a normal IRP do not have the criteria for major motility disorders and are referred to as BMF. In our study, BMF was more prevalent than the other types of EMDs.

In Iran, this is the first cross-sectional study on the relationship between the sensitivity, specificity, and accuracy of symptoms in patients with EMDs and normal esophageal manometry findings, based on the most recent Chicago Classification. Nonetheless, this study had some limitations, first and foremost among which is that the motility patterns may differ between liquid and solid boluses. It is worthy of note, however, that the technical limitations are the consequence of the patients' condition (e.g. achalasia) or issues related to anatomy (e.g. hernia).

### Conclusion

Our findings suggest that clinical esophageal symptoms may not be reliable enough for the differentiation of EMDs from one another. However, clinical symptoms can discriminate between abnormal and normal esophageal motility findings, especially achalasia. Furthermore, abnormal esophagogastric junction relaxation during swallowing is not an accurate diagnostic criterion for predicting clinical symptoms, which makes defining the peristalsis pattern mandatory. Other modalities such as barium esophagography and esophageal endosonography are required to correct diagnosis and management. There is no doubt that findings of the present study should be further analyzed by future studies on larger sample volumes.

### Acknowledgments

We thank the Department of Gastroenterology, Endoscopy, and Pathology as well as the Motility Disorders Laboratory of

Firoozgar Teaching Hospital.

### References

1. Gyawali CP, Bredenoord AJ, Conklin JL, Fox M, Pandolfino JE, Peters JH, et al. Evaluation of esophageal motor function in clinical practice. *Neurogastroenterology & Motility* 2013; 25: 99–133.
2. Kahrilas PJ. Esophageal motor disorders in terms of high-resolution esophageal pressure topography: what has changed? *Am J Gastroenterol* 2010; 105(5):981–7.
3. Bredenoord AJ, Fox M, Kahrilas PJ, Pandolfino JE, Schwizer W, Smout AJ. Chicago classification criteria of esophageal motility disorders defined in high resolution esophageal pressure topography. *Neurogastroenterology & Motility* 2012; 24(Suppl 1):57–65.
4. Pandolfino JE, Leslie E, Luger D, Mitchell B, Kwiatek MA, Kahrilas PJ. The contractile deceleration point: an important physiologic landmark on oesophageal pressure topography. *Neurogastroenterol Motil* 2010; 22(4): 395–400, e90.
5. Pandolfino JE, Kahrilas PJ. AGA technical review on the clinical use of esophageal manometry. *Gastroenterology* 2005; 128(1):209–24
6. Mikieli J, Islami F, Malekzadeh R. Achalasia: A review of Western and Iranian experiences. *World J Gastroenterol* 2009; 15(40):5000–5009
7. Rohof WO, Bergman JJ, Bartelsman JF. Screening for dysplasia in idiopathic achalasia using Lugol staining. *Gastroenterology* 2011; 140 (Suppl): S–227.
8. Mikieli J, Farrokhi F, Bishehsari F, Mahdavinia M, Malekzadeh R. Gender effect on clinical features of achalasia: a prospective study. *BMC Gastroenterol* 2006; 6:12.
9. Pandolfino JE, Roman S, Carlson D, Luger D, Bidari K, Boris L, et al. Distal Esophageal Spasm in High-Resolution Esophageal Pressure Topography: Defining Clinical Phenotypes. *Gastroenterology* 2011; 141(2):469–475.
10. Liang B, Yi O, Feng Y. Association of gastroesophageal reflux disease with asthma control. *Diseases of the Esophagus* 2013; 26:794–8
11. Ghosh SK, Janiak P, Schwizer W, Hebbard GS and Brasseur JG. Physiology of the esophageal pressure transition zone: separate contraction waves above and below. *Am J Physiol Gastrointest Liver Physiol* 2006; 290:G568–G76.
12. Bogte A, Bredenoord AJ, Oors J, Siersema PD, Smout AJPM. Relationship between esophageal contraction patterns and clearance of swallowed liquid and solid boluses in healthy controls and patients with dysphagia. *Neurogastroenterology & Motility* 2012; 24:e364–e72.
13. Ghosh SK, Pandolfino JE, Rice J. Impaired deglutitive EGJ relaxation in clinical esophageal manometry: a quantitative analysis of 400 patients



and 75 controls. *Am J Physiol Gastrointest Liver Physiol* 2007;293(4):G878–85.

14. Sankineni A, Salieb L, Harrison M, Fisher RS, Parkman HP. Slow esophageal propagation velocity: association with dysphagia for solids. *Neurogastroenterology & Motility* 2013;25(1):e44-51.

15. Scherer JR, Kwiatek MA, Soper NJ, Pandolfino JE, Kahrilas PJ. Functional esophagogastric junction obstruction with intact peristalsis: a heter-

ogeneous syndrome sometimes akin to achalasia. *J Gastrointest Surg* 2009;13(12):2219-25.

16. Roman S, Pandolfino JE, Chen J, Boris L, Luger D, Kahrilas PJ. Phenotypes and clinical context of hypercontractility in high resolution esophageal pressure topography (EPT). *Am J Gastroenterol* 2012;107(1):37–45.