

Corelation between the Endoscopic and Histologic Diagnosis of Gastritis at the Ekiti State University Teaching Hospital, Ado Ekiti, Nigeria

Akande Oladimeji Ajayi^{1*}, Ebenezer Adekunle Ajayi¹, Olusoji Abidemi Solomon²,
Babatunde Duduyemi³, Emmanuel Abidemi Omonisi³, Olufemi Joshua Taiwo³

¹Department of Medicine, College of Medicine, Ekiti State University, Ado-Ekiti, Nigeria

²Department of Family Medicine, Ekiti State University Teaching Hospital, Ado-Ekiti, Nigeria

³Department of Anatomic Pathology, College of Medicine, Ekiti State University, Ado-Ekiti, Nigeria

Abstract **AIM AND OBJECTIVE:** Endoscopists make judgements on the presence or absence of gastritis on the basis of endoscopic appearances of the gastric mucosa. Most studies in literature have shown poor concordance between endoscopic and histologic diagnosis of gastritis with concordance rates of between 54-63%. The aim of this study was to evaluate the rate of concordance between endoscopic and histologic diagnosis of gastritis at our facility given the variable but generally poor concordance reported in literature. **MATERIALS AND METHODS:** One hundred and seventy three consecutive patients who underwent upper gastrointestinal endoscopy and biopsy for dyspepsia at Ekiti State University Teaching Hospital (EKSUTH), Ado Ekiti, Nigeria were recruited into this study. Ethical approval for the study was obtained from the hospital's Research and Ethics Committee and all the patients gave their individual written consent. **RESULTS:** The mean age of the studied population was 53.02±16.28 years (age range 17-87 years). Epigastric pain was the main (86.7%) indication for referral for upper gastrointestinal endoscopy, distantly followed by belching (5.8%), early satiety (5.2%) and bloating (2.3%) respectively. At endoscopy; diffuse erythema was found in 38.7% (67/173), followed in the descending order by linear erythema in 28.3% (49/173), atrophic gastritis in 13.9% (24/173), patchy erythema in 12.7% (22/173) respectively and normal findings in 6.4% (11/173). At histology; superficial chronic active gastritis was the most common findings 74.6% (129/173) of the patients, followed by chronic atrophic gastritis 15.6% (27/173) and normal findings in 9.8% (17/173). Of the 162 that were endoscopically diagnosed as gastritis, 153 were histologically confirmed (concordance rate of 94.4%). Also, of the 156 histologically diagnosed as gastritis, 153 were endoscopically diagnosed (concordance rate of 98.1%). In all, 153 of the 173 patients were both endoscopically and histologically diagnosed as gastritis (concordance rate of 88.4%). 76.3% (119/156) of those that were diagnosed as gastritis histologically were found to be *H.pylori* associated gastritis. **CONCLUSIONS:** In conclusion, endoscopy is a reliable predictor of histological gastritis in a population with a severe form of gastritis. We equally agreed with other authors that histology is the goal standard and mandatory for accurate diagnosis of gastritis in all cases.

Keywords Endoscopy, Histology, Gastritis, Concordance

1. Introduction

Endoscopists and Gastroenterologists make judgements on the presence or absence of gastritis on the basis of the endoscopic appearance of the gastric mucosa. The concept of "endoscopic gastritis" was given acknowledgement by the working party that formulated the Sydney System of classifying gastritis (endoscopic division) [1].

Most studies in literature have shown poor concordance between endoscopic and histologic diagnosis of gastritis with concordance rates of between 54-63% [2-4]. Very few studies however, have found good concordance with one in particular reporting a concordance rate of about 97% [5-6]. In the severe forms of gastritis, concordance is said to be good and that a normal endoscopy may actually excludes active gastritis [2, 7].

The aim of this study was to evaluate the rate of concordance between endoscopic and histologic diagnosis of gastritis at our facility given the variable but generally poor concordance reported in literature.

* Corresponding author:

dejiajayi2@yahoo.co.uk (Akande Oladimeji Ajayi)

Published online at <http://journal.sapub.org/ijim>

Copyright © 2015 Scientific & Academic Publishing. All Rights Reserved

2. Materials and Methods

Study location: This study was carried out at the Ekiti State University Teaching Hospital (EKSUTH), Ado-Ekiti, Nigeria from January 2012 to December 2013.

Study population: One hundred and seventy three consecutive patients who underwent upper gastrointestinal endoscopy for dyspepsia were recruited into this study.

Study type: This is a cross sectional prospective study.

Inclusion and exclusion criteria: All patients aged 16 years and above with signs and symptoms of dyspepsia were included in the study, while patients who refused endoscopy or their medical condition precluded a satisfactory endoscopic examination were excluded.

Data collection: The followings were extracted from the patients or the relations: age, gender, use of aspirin or NSAIDS, alcohol ingestion, use of native concoctions, haematemesis, melena, haematochezia, presenting signs and symptoms. Endoscopic findings and histologic reports were documented.

Methods and procedure: Upper gastrointestinal endoscopy was performed using Olympus EVIS GIF-140 under xylocaine anaesthetic spray of the oropharynx and where necessary with midazolam after an overnight fasting with the patient on the left lateral position. All the endoscopies were performed by a single trained Endoscopist who formed a global impression on the presence or otherwise of gastritis in the antrum of the stomach. Three gastric biopsies were taken from the antrum using a standard 7 FG sized biopsy forceps. All the biopsies were taken from the perceived “inflamed” areas and fixed immediately in 10% formalin. Endoscopic gastritis was classified according to the Sydney system [1] taking into consideration the following components oedema, punctate and confluent erythema, friability, punctate and confluent exudate, flat and raised erosions, rugal hyperplasia and atrophy, visibility of the vascular pattern, punctate and confluent intramural bleeding spots, and fine and coarse nodularity. The endoscopic appearances of the antrum were documented. The biopsies were cut into sections and stained with the routine Haematoxylin and Eosin (H & E) and also with Giemsa stain. The slides were assessed histologically by three Gastrointestinal Pathologists at different times and were blinded to the endoscopic features of the patients. The histological assessment was in accordance with the criteria described by the Sydney classification System (histologic division) [1, 8], which includes a subjective assessment of severity as mild, moderate or severe, and then classified into one of the following eight categories: erythematous/exudative gastritis, atrophic gastritis, raised erosive gastritis, flat erosive gastritis, haemorrhagic gastritis, rugal hyperplastic gastritis, enterogastric reflux gastritis, and congestive gastroenteropathy.

Ethical clearance: Ethical approval for the study was

obtained from the hospital’s Research and Ethics Committee and all the patients gave their individual written consent.

Statistical analyses: SPSS version 15.0 (SPSS, Inc., Chicago, Illinois, USA) was deployed for statistical analysis using the t-test for quantitative variables and χ^2 test for qualitative variables. Differences were considered to be statistically significant if P value was less than 0.05.

3. Results

The male: female ratio was 1:1. The mean age of the studied population was 53.02 ± 16.28 years (age range 17-87 years). Majority of the patients were in the age group 31-60 years (Table 1). Epigastric pain was the main (86.7%) indication for referral for upper gastrointestinal endoscopy, distantly followed by belching (5.8%), early satiety (5.2%) and bloating (2.3%) respectively (Figure 1).

Table 1. Baseline characteristics of the studied population

Physical measurement (mean \pm standard deviation):	
Age in years	53 \pm 16.28
Indication, n (%):	
Epigastric pain	150(86.7)
Belching	10(5.8)
Early satiety	9(5.2)
Bloating	4(2.3)
Age group in years, n (%):	
up to 20 years	1(.6)
21-30 years	18(10.4)
31-40 years	22(12.7)
41-50 years	36(20.8)
51-60 years	45(26.0)
61-70 years	25(14.5)
above 70 years	26(15.0)
Gender, n (%):	
Female	88(50.9)
Male	85(49.1)

Table 2. Showing the various endoscopic findings

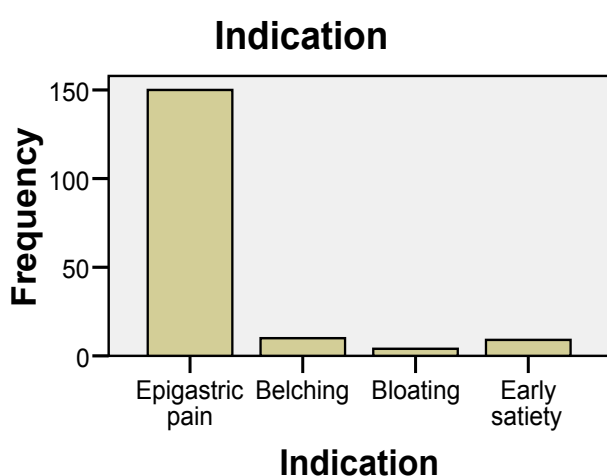
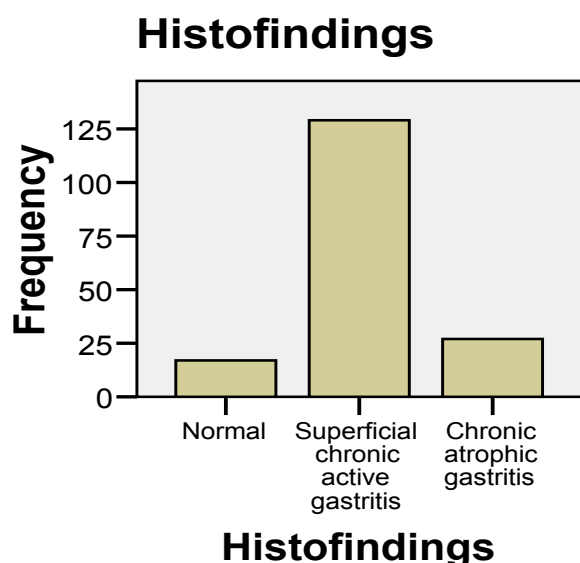
	Frequency	Percent	Valid Percent	Cumulative Percent
Normal	11	6.4	6.4	6.4
Patchy erythema	22	12.7	12.7	19.1
Linear erythema	49	28.3	28.3	47.4
Diffuse erythema	67	38.7	38.7	86.1
Atrophic changes	24	13.9	13.9	100.0
Total	173	100.0	100.0	

Table 3. Correlation between endoscopy and histology findings

		Histo		Total
		Normal	Gastritis	
Endofind	Normal	8	3	11
	Gastritis	9	153	162
	Total	17	156	173

Pearson's R = 0.551 P value = 0.001

At endoscopy; diffuse erythema was found in 38.7% (67/173), followed in the descending order by linear erythema in 28.3% (49/173), atrophic gastritis in 13.9% (24/173), patchy erythema in 12.7% (22/173) and normal findings in 6.4% (11/173) (Table 2).

**Figure 1.** Showing the indications for endoscopy**Figure 2.** Showing the various histological findings

At histology; superficial chronic active gastritis was the most common findings 74.6% (129/173) of the patients, followed by chronic atrophic gastritis 15.6% (27/173) and normal findings in 9.8% (17/173) [Figure II]. Of the 162 that were endoscopically diagnosed as gastritis, 153 were

histologically confirmed (concordance rate of 94.4%). Also, of the 156 histologically diagnosed as gastritis, 153 were endoscopically diagnosed (concordance rate of 98.1%). In all, 153 of the 173 patients were both endoscopically and histologically diagnosed as gastritis (concordance rate of 88.4%). Of the eleven that were found to be normal endoscopically, eight were confirmed to be histologically normal while the remaining three were diagnosed as superficial chronic active gastritis (concordance rate of 72.7%). On the other hand, however, eight of the seventeen found to be histologically normal were endoscopically found normal while the remaining nine were found to have patchy erythema endoscopically (concordance rate of 47.1%). The correlation coefficient was found to be 0.551 and p value was 0.001 (Table 3). 76.3% (119 /156) of those that were diagnosed as gastritis histologically were found to be *H.pylori* associated gastritis. Among the studied population, the prevalence of *H.pylori* infection was found to be 68.8%. Use of aspirin or NSAIDS, native concoctions and alcohol ingestion were not statistically significant in this study population.

4. Discussion

The Sydney classification of endoscopic gastritis [1] aims to standardize reporting by classifying endoscopic gastritis into seven categories, based on 15 endoscopic mucosal features: oedema, punctate and confluent erythema, friability, punctate and confluent exudate, flat and raised erosions, rugal hyperplasia and atrophy, visibility of the vascular pattern, punctate and confluent intramural bleeding spots, and fine and coarse nodularity. It also re-categorizes portal hypertensive gastropathy as congestive gastroenteropathy. This study was devised to correlate the endoscopic features described by the Sydney system with the histological findings.

Diffuse erythema was the commonest type of gastritis diagnosed endoscopically in this study 38.7% (67/173) while on the other hand; superficial chronic active gastritis was the most common histological diagnosis 74.6% (129/173). Of the 173 patients, 153 were diagnosed as gastritis both endoscopically and histologically giving a concordance of 88.4%. While a number of studies in literature have shown abysmal or poor concordance between endoscopic and histologic diagnosis of gastritis with concordance rates of between 54-63% [2-4, 9-13], a high concordance obtained in this study is similar to the 97% concordance obtained by Fung et al [5] and Zubair et al [6]. This might be due to the fact that most of our patients had severe forms of gastritis at the time of their presentation. Late presentation is the most common form of presentation of diseases in the third or under developed worlds. Studies have equally shown that concordance between endoscopy and histology may be good in the severe forms of gastritis [1] and that a normal endoscopy may in actual fact excludes active gastritis [7]. High resolution magnifying endoscopy has been shown to

have a better correlation with histologic findings and might indeed allow definitive diagnosis of gastritis to be made without the need for biopsy [14-15]. Similar conclusions have been reached for duodenitis [11, 16].

Out of the eleven that were found to be normal endoscopically, eight were confirmed to be histologically normal while the remaining three were diagnosed as superficial chronic active gastritis. This observation was similar to the findings in the study of Gad [17].

Of the 22 diagnosed endoscopically as patchy erythema, 13 were histologically found to have gastritis (concordance of 59%). This findings support the studies that found poor concordance in mild forms of gastritis showing the insensitivity of endoscopy in diagnosing gastritis in the mild forms of gastritis.

Endoscopically in this study, 80.9% (140/173) of the patients had severe forms of gastritis and all these patients had gastritis histologically. This finding is in keeping with the studies of Tytgat [1], and Toukan et al [7]. This might be due to the high prevalence of *H.pylori* infection in the third world countries. 93.6% of our study population had one form of gastritis or the other at endoscopy; this was in contrary to the 62.1% obtained by Jemilohun et al [18] in a similar population study. Again, this might be due to fact that most of our patients had severe forms of gastritis at the time of their presentation. 90.1% of our patients had histologically diagnosed gastritis similar to the 96.6% found by Jemilohun et al [18]. 88.4% of our patients were both endoscopically and histologically diagnosed as gastritis compared to the 60.9% in the Jemilohun et al study [18].

The prevalence of *H.pylori* associated gastritis in this study was 73.6% while the overall prevalence among the studied population was 68.8%. This is consistent with results of previous studies conducted in Nigeria and other parts of West Africa which have consistently shown a high prevalence of *H. pylori* with the use of biopsy based methods [19-24].

In conclusion, endoscopy is a reliable predictor of histological gastritis in a population with a severe form of gastritis. We equally agreed with other authors that histology is the goal standard and mandatory for accurate diagnosis of gastritis in all cases.

REFERENCES

- [1] Tytgat GNJ. The Sydney System: Endoscopic division. Endoscopic appearances in gastritis/ duodenitis. J Gastroenterol Hepatol 1991; 6:223-234.
- [2] Atkins L, Benedict E. Correlation of gross gastroscopic findings with gastroscopic biopsy in gastritis. N Engl J Med 1956; 254:641-644.
- [3] Kaur G, Raj M. A study of the concordance between endoscopic gastritis and histological gastritis in an area with a low background prevalence of *Helicobacter pylori* infection. Singapore Med J 2002; 43: 90-92.
- [4] Redeen S, Petersson F, Jonsson KA, et al. Relationship of gastroscopic features to histological findings in gastritis of *Helicobacter pylori* infection in a general population sample. Endoscopy 2003; 35:946-950.
- [5] Fung WP, Papadimitriou JM, Matz LR. Endoscopic, histological and ultrastructural correlations in chronic gastritis. Am J Gastroenterol 1979; 71(3):269-79.
- [6] Zubair M, Channa MA, Mirza MR, et al. Is biopsy needed in every gastritis found during endoscopy? Pak J Med Sci 2009; 25:849-51.
- [7] Toukan AU, Kamal MF, Amr SS, Arnaout MA et al. Gastroduodenal inflammation in patients with nonulcer dyspepsia. A controlled endoscopic and morphometric study. Dig Dis Sci 1985; 30(4):313-320.
- [8] Price B. The Sydney System: Histological division. J Gastroenterol Hepatol 1991; 6:209-222.
- [9] Kreuning J, Bosman FT, Kuiper G, Van der Wal AM, Lindman J. Gastric and duodenal mucosa in 'healthy' individuals. J Clin Pathol 1978; 31:69-77.
- [10] Jonsson KA, Gotthard R, Bodemar G, Brodin U. The clinical relevance of endoscopic and histologic inflammation of a gastroduodenal mucosa in dyspepsia of unknown origin. Scand J Gastroenterol 1989; 24:385-395.
- [11] Kaur G, Raj SM. A study of concordance between Endoscopic gastritis and Histologic gastritis in an area with a low background prevalence off *H.pylori* infecton. Singapore Med J 2002; 43(2): 090-092
- [12] Carr NJ, Leadbetter H, Marriott A. Correlation between the endoscopic and histologic diagnosis of gastritis. Annals of Diag Path 2012; 16: 13-15
- [13] Anagnostopoulos GK, Yao K, Kaye P, et al. High-resolution magnification endoscopy can reliably identify normal gastric mucosa, *Helicobacter pylori*-associated gastritis, and gastric atrophy. Endoscopy 2007; 39: 202-207.
- [14] Gonen C, Simsek I, Sarioglu S, et al. Comparison of high resolution magnifying endoscopy and standard videoendoscopy for the diagnosis of *Helicobacter pylori* gastritis in routine clinical practice: a prospective study. Helicobacter 2009;14: 12-21.
- [15] Lewis S, Stableforth W, Awasthi R, Awasthi A, Pitts N, et al. An examination of the relationship between the endoscopic appearance of duodenitis and the histological findings in patients with epigastric pain. Int J Clin Exp Pathol 2012;5(6):581-587.
- [16] Gad A. Erosion: a correlative endoscopic histopathologic muticenter study. Endoscopy 1986; 18: 76-79.
- [17] Jemilohun AC, Otegbayo JA, Ola SO, Oluwasola AO, Akere A. Correlation between Endoscopic and Histological Gastritis in South-Western Nigerians with Dyspepsia. Niger J Gastro Hepatol 2010. Vol. 2 (2): 73-76.
- [18] Poudel A, Regmi S, Poudel S, Joshi P. Correlation between endoscopic and histopathological findings in gastric lesions. Journal of Uni Coll of Med Sci. 2013. Vol.1 (3):37-41.
- [19] Ndububa DA, Agbakwuru AE, Adebayo RA, Olasode BJ,

- Olaomi OO, Adeosun OA. et al. Upper gastrointestinal findings and incidence of *Helicobacter pylori* infection among Nigerian patients with dyspepsia. *West Afr J Med*. 2001; 20(2):140–145.
- [20] Adesanya AA, Oluwatowaju IO, Oyediji KS, da Rocha-Afodu JT, Coker AO, Afonja OA. Evaluation of a locally-made urease test for detecting *Helicobacter pylori* infection. *Niger Postgrad Med J*. 2002;9(1):43–47.
- [21] Baako BN, Darko R. Incidence of *Helicobacter pylori* infection in Ghanaian patients with dyspeptic symptoms referred for upper gastrointestinal endoscopy. *West Afr J Med*. 1996;15(4):223–227.
- [22] Mustapha SK, Ajayi NA, Nggada HA, Pindiga UH, Bolori MT, Ndahi A. et al. Endoscopic findings and the frequency of *Helicobacter pylori* among dyspeptic patients in North-Eastern Nigeria. *Highland Medical Research Journal*. 2007;5(1):78–81.
- [23] Mbengue M, Diouf ML, Dangou JM, Ka MM, Ba-Seck A, Ndiaye MF. et al. [Frequency of *Helicobacter pylori* infection in symptomatic patients in Senegal]. *Med Trop (Mars)*. 1997;57(3):256–258.
- [24] Aduful HK, Naaeder SB, Darko R, Baako BN, Clegg-Lampsey JNA, Nkrumah KN. et al. Upper Gastrointestinal Endoscopy at the Korle Bu Teaching Hospital, Accra, Ghana. *Ghana Med J*. 2007; 41(1):12–16.