

An Adjunctive Technique in the Detection of Pancreaticobiliary Malignancy: Computer-Assisted Wide Area Transepithelial Sampling

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Abstract

Background/Aims: Computer assisted full thickness brush biopsy has been used to identify intestinal metaplasia and dysplasia in the esophagus. Its efficacy in diagnosing pancreatic and biliary neoplasia has not previously been evaluated. The objective of this study was to compare the sensitivity of the Cdx Wide Area Transepithelial Sampling (WATS3D®) Brush Biopsy in the diagnosis of pancreatic and biliary malignancies to standard brush cytology and forceps biopsy during Endoscopic Retrograde Cholangiopancreatography (ERCP).

Materials and Methods: Forty-six patients underwent ERCP for evaluation of biliary dilatation, stricture, or mass in the pancreas or biliary tree found on radiological imaging (CT scan, Ultrasound, or MRI). All patients underwent standard brush cytology, forceps biopsy, and computer assisted full thickness brush biopsy. The primary outcome measured was the detection of neoplasia with each modality. All pathology was reviewed by pathologists at New York Presbyterian Queens, NY, with final readings categorized as malignant, nonmalignant (metaplasia, hyperplasia), or atypical (dysplasia).

Results: The CDx WATS3D® brush biopsy was positive for malignant cells in 39.13% patients. This was a statistically significant finding when compared to the detection rate of malignant cells by traditional brushing of 21.74% ($p = 0.035$). Forceps biopsy was positive for malignant cells in 26.1% ($p = 0.09$).

Conclusions: WATS3D demonstrates enough potential for use in the diagnosis of pancreatobiliary malignancy to warrant additional studies with larger cohorts to clarify the role of this tissue acquisition technique.

Keywords: ERCP; Brush Biopsy; Pancreaticobiliary Neoplasms; Forceps Biopsy; WATS3D

Abbreviations

ERCP: Endoscopic Retrograde Cholangiopancreatography; WATS3D®: Wide Area Transepithelial Sampling

Introduction

Endoscopic Retrograde Cholangiopancreatography (ERCP) has long been used in the evaluation and treatment of biliary strictures and masses. In addition to imaging the biliary tree and pancreatic duct, ERCP allows for tissue acquisition. Both cytology and forceps biopsies can be used. Studies have shown the sensitivity of standard brush cytology to be 35 - 70% [1-3] and forceps biopsy to be 40 - 80% [4,5]. Wide-Area Transepithelial Sampling 3Dimensional Analysis (WATS3D®) Brush Biopsy, CDx, Suffrin, NY) computer assisted full thickness

brush biopsy is a method of tissue collection that samples the total thickness of the epithelium. Improved diagnostic accuracy is thought to be due to the abrasive, rigid structure of the brush which is passed through the working port of the endoscope. Tissue acquisition is enhanced utilizing a continuous, up-and-down motion with gentle pressure that captures deeper layers that may be missed with standard forceps or brushings. Abnormal cells are then identified through a computer assisted analysis which includes 3-dimensional scanning to evaluate each portion of the full thickness specimen. When utilized to obtain biopsies of esophageal tissue, WATS3D, in conjunction with forceps biopsies, has resulted in a statistically significant increase in the number of patients diagnosed with esophageal intestinal metaplasia and dysplasia [6,7]. There is a paucity of literature that describes the use of WATS3D in the diagnosis of biliary malignancies. In this study, we compared the sensitivity and specificity of the WATS3D to standard brush cytology and forceps biopsy in diagnosing pancreaticobiliary malignancies at ERCP.

Materials and Methods

This was a retrospective case series design that included 40 patients (Males 18, Females 22) with a mean age of 80 (range 55 - 96). All patients underwent ERCP to evaluate pancreaticobiliary abnormalities suspicious for malignancy found on radiographic imaging. All procedures were performed at New York Presbyterian Queens between April 2012 and December 2013. Patients underwent standard cholangiography followed by sequential traditional brushing, WATS3D cytology and forceps biopsy, or traditional brushing and WATS3D cytology only without forceps biopsy. The primary outcome measured was the detection of neoplasia with each modality. All patients underwent standard brushing during ERCP with a 2.1 mm x 8 French cytology brush (RX Cytology Brush, Boston Scientific; Natick, Massachusetts) over a 0.035 inch guidewire (Hyrda Jagwire™, Boston Scientific; Natick, Massachusetts). Each stricture, mass, or area of dilatation was then brushed 20 times. After brushing was completed with the standard cytology brush, WATS3D was then used. Finally, in some patients, a forceps biopsy with a 2.0 mm biopsy forceps (Radial Jaw (™), Boston Scientific, Natick Massachusetts) was used. All pathology was reviewed by pathologists at New York Presbyterian-Queens, NY, with final readings categorized as malignant, nonmalignant (metaplasia, hyperplasia), or atypical (dysplasia). Definitive diagnoses were obtained by forceps biopsy. Cytologic findings obtained by WATS3D and brushings were compared with the final diagnosis. A positive finding from any of the three modalities (forceps, brushing or WATS3D) was considered a positive finding and was treated as such. Atypical findings were considered positive.

Informed consent was not obtained, as the study is a retrospective case series design; all patient information was de-identified prior to data analysis. Prior to data collection, internal review board approval was obtained for chart review. All listed authors verify their full and honest academic involvement with regard to initiation of study, review of current literature regarding pancreaticobiliary malignancy detection, and collection and analysis of the data presented in this manuscript.

Statistical Analysis

Statistical analysis was performed using SPSS statistics version 21.0 (IBM corporation, Armonk, NY, USA). The simple kappa coefficient was derived using the SPSS software to obtain measure of agreement. Sensitivities and specificities were obtained using SPSS software.

Results

In the period between April 2012 and December 2013, 40 patients with average age of 80 (range 55 - 96) underwent ERCP for biliary stricture as seen on radiographic imaging, with sequential biopsies procured using brushing, WATS3D and/or forceps biopsy. Of the 40 patients, 18 underwent brushing and WATS3D CDx followed by forceps biopsy and 22 patients underwent brushing and CDx without forceps biopsy. Of the 18 patients who underwent brushing, CDx and forceps biopsy, Cholangiocarcinoma was detected in 8 patients (with one result indeterminate) as diagnosed with forceps biopsy, the gold standard. The sensitivity and specificity of CDx as compared to forceps biopsy was 75% and 67%, respectively. The sensitivity and specificity of cytologic brushings was 25% and 89%, respectively (Table 1).

Summary	Brush Biopsy	WATS CDx
Sensitivity	25%	75%
Specificity	89%	67%

Table 1: Sensitivity and Specificity of WATS CDx and brush biopsy.

There was moderate concordance between forceps biopsy and CDx ($\kappa = 0.414$). Of the negative forceps biopsy results, WATS3D identified one malignancy, and 2 atypical specimen results. CDx and Brushing cytology showed moderate concordance ($\kappa = 0.4738$), and when used in conjunction, specificity was 100%. Although brushing cytology was specific, it showed only slight concordance when compared with forceps biopsy ($\kappa = 0.1439$) In the 22 patients who did not undergo the gold standard of forceps biopsy, WATS3D identified 3 additional malignancies, and 2 additional atypical results. WATS3D failed to identify one malignancy that was positive using forceps biopsy.

Discussion

ERCP is a technique used in the evaluation and treatment of biliary strictures and masses, in which tissue is obtained via brush and forceps biopsy. Although these modalities have a high specificity, sensitivity remains poor. Brush cytology, which is the most commonly used method for tissue sampling, has an estimated sensitivity of 35-70% when used alone and a specificity of over 90% [4,5,8]. In a recent study including 75 specimens, sensitivity of brush cytology was 66% [9]. The sensitivity of forceps biopsy ranges from 43 - 80% [8,10-12]. Though the combination of brushing and biopsy has been previously evaluated, the data has been mixed. One study suggested that the combination could increase the sensitivity by 15 - 25% compared to either option alone [13-15]. However, other studies suggest that the combination of brushing and biopsies were more sensitive, and the risks of doing both was not warranted [11,13]. This demonstrates the need for a safe, reliable, and sensitive technique for diagnosing pancreaticobiliary neoplasia.

The WATS3D system has been shown to improve diagnostic accuracy when used as an adjunct to forceps biopsy in detecting Barrett's Metaplasia [7]. In a recent randomized clinical trial, 160 patients underwent esophageal biopsy followed by WATS3D, or vice-versa. The absolute diagnostic yield increased 14.4% when WATS3D was added to standard biopsy [17]. The use of Wide-Area Transepithelial Sampling outside of Barrett's Metaplasia has not been studied. Due to the importance of prompt surgical resection in the management of biliary malignancies, an accurate tissue diagnosis is important.

WATS3D displayed more sensitivity when compared to standard brushing cytology. Additionally, our data show that WATS3D, while not as specific a test as compared to brush cytology, did show moderate concordance. This is encouraging, as it is expected that the studies would be, at best, moderately concordant considering the apparent increased sensitivity of WATS3D.

Conclusion

WATS3D is a specialized form of tissue acquisition that has been previously utilized for analysis of Barrett's metaplasia. Our study is the first to report the use of this technology in the diagnosis of pancreatobiliary malignancy.

The advantages of WATS3D over traditional cytology include stiffer brushes allowing for deeper tissue acquisition and greater sampled surface area, as well as computer aided analysis of cells. The technical limitation to the use of WATS3D is the lack of guidewire utilization, which may limit the reach of the brush beyond certain strictures.

Our study was limited by the small sample size which most likely contributed to the poor sensitivity of brush cytology. Additionally, the sensitivity of each method might have been affected by the order of tissue acquisition, which was not analyzed in this study. Despite this, the promising results demonstrated by this study warrants further investigation. Additional studies with larger cohorts have the potential to clarify the role of Wide-Area Transepithelial Sampling brush biopsy in the diagnosis of pancreaticobiliary neoplasms.

Competing Interests

The authors declare that they do not have any conflicts of interest.

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Bibliography

1. Bain VG, *et al.* "Prospective study of biliary strictures to determine the predictors of malignancy". *Canadian Journal of Gastroenterology* 14.5 (2000): 397-402.
2. Adamsen S, *et al.* "Endobiliary brush biopsy: Intraand interobserver variation in cytological evaluation of brushings from bile duct strictures". *Scandinavian Journal of Gastroenterology* 41.5 (2006): 597-603.
3. Glasbrenner B, *et al.* "Prospective evaluation of brush cytology of biliary strictures during endoscopic retrograde cholangiopancreatography". *Endoscopy* 31.9 (1999): 712-717.
4. Kurzawinski TR, *et al.* "A prospective study of biliary cytology in 100 patients with bile duct strictures". *Hepatology* 18.6 (1993): 1399-1403.
5. Schöfl R. "Diagnostic endoscopic retrograde cholangiopancreatography". *Endoscopy* 3.2 (2001): 147-157.
6. Anandasabapathy S, *et al.* "Computer Assisted Brush Biopsy Analysis for the Detection of Dysplasia in a High Risk Barrett's Esophagus Surveillance Population". *Digestive Diseases and Sciences* 56.3 (2011): 761-766.
7. Johanson JF, *et al.* "Computer Assisted Analysis of Abrasive Transepithelial Brush Biopsies Increases the Effectiveness of Esophageal Screening: A Multicenter Prospective Clinical Trial by the EndoCDx Collaborative Group". *Digestive Diseases and Sciences* 56.3 (2011): 767-772.
8. Trent V, *et al.* "Diagnostic accuracy and clinical utility of endoscopic bile duct brushing in the evaluation of biliary strictures". *Archives of Pathology and Laboratory Medicine* 123.8 (1999): 712-715.
9. Eiholm S, *et al.* "Endoscopic brush cytology from the biliary duct system is still valuable". *Danish Medical Journal* 60.7 (2013): A4656.
10. Lee JG, *et al.* "Benign, dysplastic, or malignant making sense of endoscopic bile duct brush cytology: results in 149 consecutive patients". *American Journal of Gastroenterology* 90.5 (1995): 722-726.
11. Logrono R, *et al.* "Analysis of false negative diagnoses on endoscopic brush cytology of biliary and pancreatic duct strictures: the experience at 2 university hospitals". *Archives of Pathology and Laboratory Medicine* 124.3 (2000): 387-392.
12. Rumalla A, *et al.* "Improved diagnostic yield of endoscopic biliary brush cytology by digital image analysis". *Mayo Clinic Proceedings* 76.1 (2001): 29-33.
13. Foutch PG. "Diagnosis of cancer by cytologic methods performed during ERCP". *Gastrointestinal Endoscopy* 40 (1994): 249-252.
14. Parasher VK and Huibregtse K. "Endoscopic retrograde wire guided cytology of malignant biliary strictures using a novel scraping brush". *Gastrointestinal Endoscopy* 48.3 (1998): 288-290.
15. Fogel EL and Sherman S. "How to improve the accuracy of diagnosis of malignant biliary strictures". *Endoscopy* 31.9 (1999): 758-760.
16. Vennalaganti PR, *et al.* "Increased Detection of Barrett's Esophagus-associated Neoplasia Using Wide-Area Transepithelial Sampling: A Multicenter, Prospective, Randomized Trial". *Gastrointestinal Endoscopy* 87.2 (2017): 348-355.

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