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Efficiency comparison of selected endoscopic video analysis algorithms

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ABSTRACT: In the paper, selected image analysis algorithms were examined and compared in the task of identifying informative frames, blurry frames, colorectal cancer and healthy tissue on endoscopic videos. In order to standardize the tests, the algorithms were modified by removing from them parts responsible for the classification, and replacing them with Support Vector Machines and Artificial Neural Networks. The tests were performed in an unified manner on a common, large movie database of real endoscopy videos. The test results often do not seem to confirm the high efficiency declared by their authors. A maximum of 80% sensitivity and specificity was achieved, while the authors often declared as much as 90%.

KEYWORDS: endoscopy, video analysis, algorithms, comparison, efficiency

I. INTRODUCTION

Since last several years, endoscopic movie analysis algorithms (for gastroscopy, colonoscopy and wireless capsule endoscopy, WCE) gained much popularity. These algorithms were designed for recognizing informative and non-informative frames, and various diseases or healthy tissues. Algorithms found in the literature are claimed by their authors to give high performance (in terms of accuracy, sensitivity, specificity etc.) results [1]. However, the publications' flaw is often the lack of comparative tests of different algorithms (or lack of any comparison at all). One of the reasons of such situation is the lack of a good public database of medical gastrointestinal endoscopy images prepared for algorithm testing purposes.

This article focuses on a comparison of selected endoscopic image analysis algorithms. To allow the comparative analysis, it was necessary to establish common conditions for algorithms' operation. Algorithms were modified so to unify their operation, and then, the comparative tests were carried out on identical sets of data, measuring algorithms' performance in detecting informative and non-informative frames, colorectal cancer and normal tissue.

II. ALGORITHMS

In the article, selected image analysis algorithms were tested and compared, as in table I.

III. TEST PROCEDURE

Algorithms were compared in two main tasks: efficiency in distinguishing (a) cancer from normal tissue of the large intestine, and (b) informative / non-informative (e.g. distorted by the movement of the endoscope, poor lighting, liquid covering the camera of the endoscope, etc.).

For this purpose, all tests were performed on a common database of real endoscopic videos of the colon [2]. To unify algorithms' operation, their parts responsible for classification were removed, leaving only the core – feature vector extraction. All feature vectors were also normalized so that every feature had the mean of 0 and standard deviation 1 over the whole database. For classification, Artificial Neural Networks (ANN) and Support Vector Machines (SVM) were used to test algorithms' efficiency (all the classifiers were trained and tested the same way on the same data).

Classifier training was carried out on the database of [1] endoscopic endoscopic videos, fully labeled by an expert for the content of each frame. The expert gave every frame of every video one of three labels: [*blurry*], [*sharp, cancer*] or [*sharp, healthy*]. Due to the different length of the videos and different proportions of labels, from each film maximum

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Table I: Algorithms tested

Ref.	Algorithm	ID	Objects detected	Feature vector
[3]	Autocorrelation Homogeneous Texture	AHT	tumor in chromoendoscopy and narrow-band imaging	AHT
[4]	Discrete Fourier Transform – Homogeneous Texture	DFT-HT	textures	HT
[5]–[7]	Gastropathy	Gastrop.	portal hypertensive gastropathy in gastroscopy	% of edge pixels, histogram P+2 bins (only P=16 tested) or $LBP_{P,R}^{riu2}$, % of blocks with local brightness maximas
[8]	Baopu Li	BaopuLi	adenoma, adenocarcinoma in Wireless Capsule Endoscopy (WCE)	10-bin histograms for 9 channels and DWT = 630 values
[9]	Poh Chee Khun	PCK-C	informative frames, bleeding in WCE	histogram for each block for each quantized value
		PCK-T		CWC – 72 features + 8 additional
[10]	Local Color Vector Pattern	LCVP	textures in magnification endoscopy	histogram of LCVP – 256 – features
[11]	Multi-scale Block LBP	MB-LBP-G	patterns for face recognition	histogram of MB-LBP – 256 features
[12]	LBP	MB-LBP-C	polyps in endoscopy	2D-histogram of color MB-LBP – up to 256^2 features (tested up to 256)
[13], [21]		Test	Test 1	tumor, polyps, informative frames in endoscopy
	Test 2		36 statistical features	
	Test 3		90 statistical features	
[16]	Kodogiannis	Kodog.	normal/abnormal tissue in WCE	54 statistical features from N_{TU}
[14]	Magoulas	Mago. 1	normal/abnormal tissue in colonoscopy	16 features from GLCM
[15]		Mago. 2		48 features from DWT and GLCM

of 30 frames (possibly far from each other) were selected for further processing for each label. Total number of selected frames was ≈ 4750 for blur recognition and ≈ 2750 for cancer recognition.

Two main types of tests were performed: (a) identify clear (informative) / blurry (non-informative) frames, and (b) identify healthy / cancerous tissue. For each test type, the input data was divided into eight sets, preserving the ratio of classes, and so that images of one patient were placed always in the same set (set assignment was performed with an algorithm described in [19]). Such set balancing is recommended in medical research [1].

Prepared sets were used to 8-fold cross-validation. For each classifier, a set of its parameters was selected, and then their optimization was performed by algorithm *CRS* [17] from *NLopt* library [18], with a time limitation of 8 hours (usually it resulted in 5000–50000 iterations of the algorithm). During the tests, following efficiency parameters were measured:

- 1) Sensitivity — performance at recognizing positive samples
- 2) Specificity — performance at recognizing negative samples
- 3) Accuracy — performance at giving correct answer
- 4) Smoothness — smoothness of the classifier's output [20]
- 5) Overall score — weighted harmonic mean of sensitivity, specificity and smoothness values

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IV. RESULTS

This section contains the test results of all tested algorithms. The tests were performed in the same manner, on the same hardware, in the same conditions, and with the same data (as described in the previous section).

Table II: Efficiency in **blur** recognition with ANN

Algorithm	Sens.	Spec.	Acc.	Video	Overall
BaopuLi	78.4%	79.4%	79.0%	92.1%	78.7%
LCVP	77.8%	80.5%	79.2%	88.7%	78.1%
MB-LBP-G	77.4%	79.5%	78.5%	88.9%	77.7%
MB-LBP-C	78.1%	77.2%	77.6%	95.3%	77.5%
Kodo	74.4%	77.6%	76.1%	89.9%	74.8%
DFT-HT	74.5%	75.6%	75.1%	89.0%	74.7%
Gastropathy	73.7%	80.8%	77.4%	88.2%	74.5%
PCK-T	72.9%	76.9%	75.1%	85.5%	73.4%
Mag2	66.5%	69.3%	68.0%	72.2%	66.8%
AHT	66.6%	66.5%	66.6%	94.1%	66.7%
Mag1	65.8%	65.9%	65.8%	90.1%	66.0%
PCK-C	61.4%	61.5%	61.4%	93.7%	61.6%
T3	60.4%	60.7%	60.6%	93.3%	60.6%
T2	59.6%	60.8%	60.3%	92.4%	60.0%
T1	57.9%	54.7%	56.2%	92.0%	55.2%

Tables II – III and figures 1 – 2 present the results of the recognition of blurry/clear (informative/non-informative) frames with the Artificial Neural Networks and Support Vector Machines. In this task, the neural network performed significantly better than SVM. The results are relatively consistent with expectations and with the descriptions of the authors of the original publications (if present). *Test* algorithms performed far worse than the others. In the task of blurry frames recognition, the best algorithms were: BaopuLi, MB-LBP-C, Kodo, LCVP.

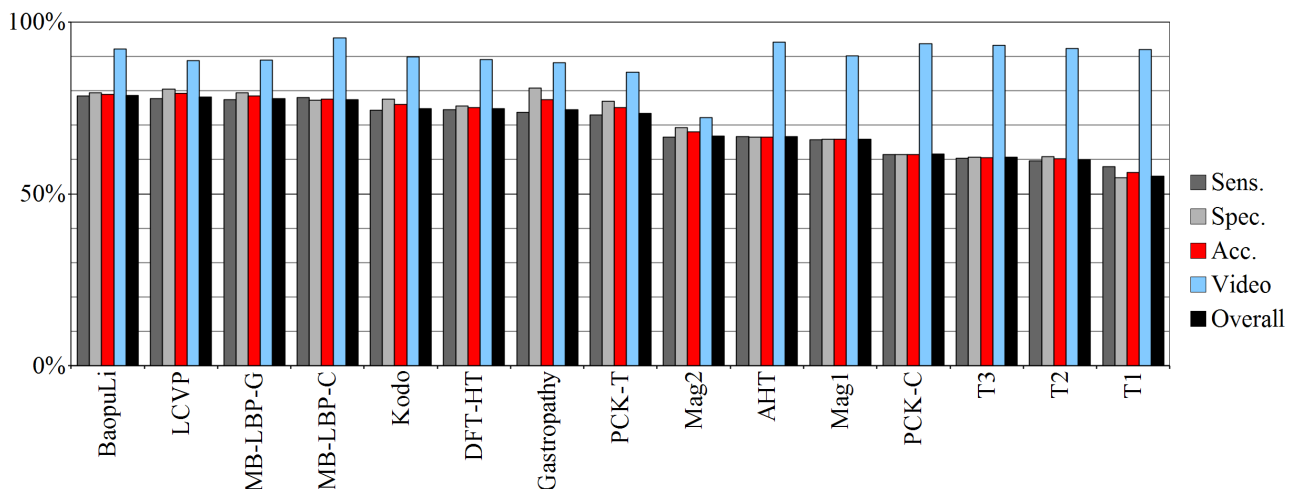


Figure 1: Algorithms' efficiency in **blur** recognition with ANN

Tables IV – V and figures 3 – 4 present the results of recognition of colorectal cancer / normal tissue, with the ANN and SVM. In this task, the neural network performed also better than SVM, though not as clearly as in blur recognition.

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Table III: Efficiency in **blur** recognition with SVM

Algorithm	Sens.	Spec.	Acc.	Video	Overall
MB-LBP-C	70.5%	71.4%	71.0%	78.7%	70.7%
Kodo	67.1%	67.8%	67.5%	83.4%	67.3%
LCVP	64.1%	65.4%	64.8%	78.3%	64.4%
DFT-HT	62.7%	64.7%	63.7%	79.5%	63.0%
PCK-C	61.5%	62.1%	61.8%	88.7%	61.8%
Gastropathy	61.0%	63.3%	62.2%	79.8%	61.4%
PCK-T	61.3%	61.0%	61.1%	44.4%	60.8%
Mag2	61.6%	60.5%	61.0%	59.6%	60.6%
BaopuLi	59.9%	62.8%	61.4%	70.7%	60.3%
T2	59.6%	61.0%	60.4%	89.0%	59.9%
AHT	60.4%	59.4%	59.8%	93.3%	59.7%
Mag1	57.4%	58.3%	57.9%	66.4%	57.6%
T1	56.3%	56.2%	56.2%	89.7%	56.4%
MB-LBP-G	52.9%	67.1%	60.4%	87.0%	54.5%
T3	51.0%	51.1%	51.0%	97.1%	51.2%

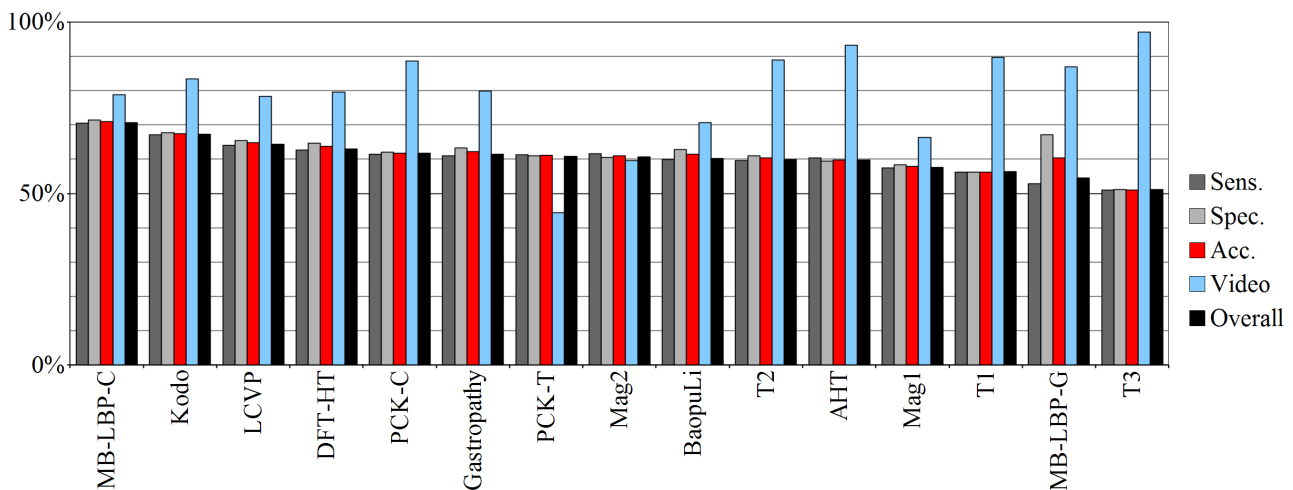


Figure 2: Algorithms' efficiency in **blur** recognition with SVM

Test algorithms also performed usually worse than most other algorithms. In the task of identifying cancerous tissue, the best algorithms were: BaopuLi, MB-LBP-C, LCVP, DFT-HT, AHT. However, these results are far below declared by the authors of the original publications of efficacy (often over 90 %) [1].

V. CONCLUSION

In the article, selected endoscopic image algorithms were tested and compared in the tasks of detection of blurry and clear (informative/non-informative) frames, colorectal cancer and healthy colon. Tests were performed on a large endoscopic video database, under the same conditions for all algorithms. The efficiency of recognizing diseases clearly differed from those declared by the authors. In the task of blur recognition, the algorithms performed similarly (or slightly better).

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Table IV: Efficiency in cancer recognition with ANN

Algorithm	Sens.	Spec.	Acc.	Video	Overall
BaopuLi	79.5%	79.3%	79.4%	76.3%	79.3%
MB-LBP-C	79.5%	78.5%	79.0%	78.0%	78.6%
DFT-HT	77.4%	77.8%	77.6%	91.1%	77.6%
LCVP	75.2%	75.4%	75.3%	83.2%	75.3%
AHT	74.6%	74.6%	74.6%	92.7%	74.7%
Gastropathy	74.8%	74.1%	74.5%	92.2%	74.3%
MB-LBP-G	73.1%	73.4%	73.2%	87.2%	73.2%
T3	72.8%	73.1%	72.9%	92.6%	72.9%
Kodo	70.5%	70.2%	70.3%	87.6%	70.4%
T2	68.1%	67.5%	67.8%	91.1%	67.7%
Mag1	67.3%	67.9%	67.6%	90.0%	67.5%
T1	65.1%	69.2%	67.0%	94.9%	65.7%
Mag2	64.1%	65.0%	64.5%	78.0%	64.3%
PCK-T	63.7%	62.1%	63.0%	50.9%	62.1%
PCK-C	62.2%	60.8%	61.5%	94.4%	61.1%

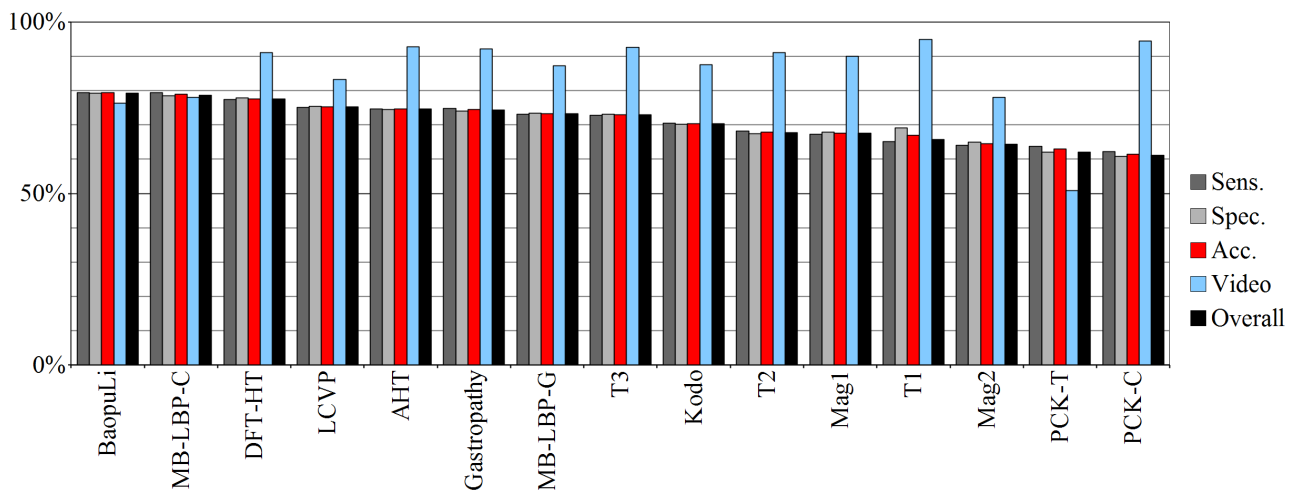


Figure 3: Algorithms' efficiency in cancer recognition with ANN

These results indicate the need for greater comparative tests across the field of the endoscopic image analysis. Such tests should be performed on a single shared database, in the same way. The previous approach of the authors in the field, consisting of only testing on their own (often small) data sets seems to be insufficient.

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Table V: Efficiency in **cancer** recognition with SVM

Algorithm	Sens.	Spec.	Acc.	Video	Overall
BaopuLi	76.5%	77.6%	77.0%	70.4%	76.6%
LCVP	72.8%	70.2%	71.6%	65.4%	70.4%
MB-LBP-C	70.2%	70.3%	70.2%	76.1%	70.3%
DFT-HT	70.0%	71.1%	70.5%	78.5%	70.2%
AHT	69.6%	69.4%	69.5%	90.9%	69.6%
MB-LBP-G	66.9%	69.6%	68.2%	72.9%	67.2%
Gastropathy	66.4%	67.6%	66.9%	64.2%	66.5%
Kodo	65.7%	66.5%	66.1%	78.8%	65.9%
T3	65.6%	65.5%	65.6%	91.2%	65.7%
Mag1	66.1%	65.2%	65.7%	84.2%	65.5%
PCK-C	64.5%	65.4%	64.9%	87.6%	64.8%
T2	64.2%	64.1%	64.2%	86.0%	64.2%
Mag2	58.8%	58.7%	58.8%	52.2%	58.7%
T1	58.0%	57.2%	57.6%	93.2%	57.5%
PCK-T	52.8%	52.2%	52.5%	94.3%	52.5%

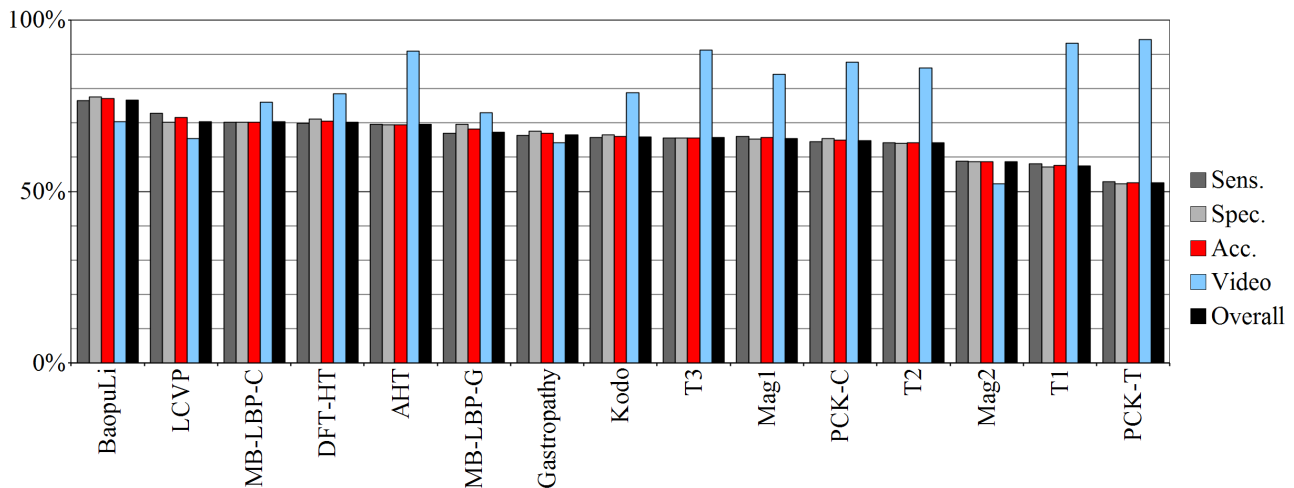


Figure 4: Algorithms' efficiency in **cancer** recognition with SVM

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