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# Automatic Detection of Polyp Using Hessian Filter and HOG Features

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#### Abstract

An endoscope is a medical instrument that acquires images inside the human body. This paper proposes a new approach for the automatic detection of polyp regions in an endoscope image using a Hessian Filter and machine learning approaches. The approach improves performance of automatic detection of polyp detection with higher accuracy. The approach uses HOG feature as a local feature since the polyp and non-polyp region often have similar color information. The approach also uses Real Adaboost and Random Forests as classifiers which works efficiently even when the dimension of feature vector becomes large. It is suggested that Hessian filter can contribute to reducing the computational time in comparison with the case when only HOG features are used to detect the polyp region. K-means++ is introduced to integrate the detection results in the classification. It is shown that polyp detection with high accuracy is performed in the computer experiments with endoscope images.

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Keywords: Polyp Defection; Hessian Filter; HOG feature; Real Adaboost; Random Forests; k-means++

## 1. Introduction

Medicine is an important area of application for computer vision. Endoscopy allows medical practitioners to observe the interior of hollow organs and other body cavities in a minimally invasive way. Diagnosis involves both shape detection and the assessment of tissue state. Here, we consider a general purpose endoscope, of the sort

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still most widely used in medical practice. There are many different kinds of polyp shape and size in endoscope images. Polyps are usually found via endoscopy but polyps can be missed. The main factor to find polyp depends on the empirical skill of medical doctor. Automatic detection of polyps, with high accuracy, is an important aid to medical practice. Diagnosis typically requires polyp removal and biopsy.

Some previous approaches <sup>1</sup>, <sup>2</sup>, <sup>3</sup> introduce features of Color Wavlet Covariance(CWC), Local Binary Pattern(LBP), Gray-Level Co-Occurrence Matrix(GLCM), which is used for the texture analysis, respectively. Another approach <sup>4</sup> learns color and *xy* position coordinates partitioned in the local window, then classifies if polyp is included or not in the each region. These are patch-based approaches and extract features and perform classification. Patch-based approach may include the problem detection ratio depends on patch size, size and position of polyp.

A patch-based approach<sup>5</sup> uses capsule endoscope image as input and extracts the Rotational Invariant Uniform Local Binary Pattern (RIULBP), and statistical value obtained from intensity histogram as texture features. Classifier is designed to learn these features and polyp is detected. The approach has some difficulty that it is difficult to detect small polyp with non-textures.

On the other hand, the geometric feature is used in the approach of so that polyp appears with ellipse form in general. The approach detects polyp with only the ellipse fitting etc. without learning. Geometrical feature based approach is robust to the small polyp but some segmentation is necessary to detect polyp with edge information, that is the detection performance depends on the edge extraction level.

Another approach which authors have proposed is Hessian Filter based approach, where Hessian filter emphasizes the blob-like structure and extracts some polyp candidates with segmentation. Some texture-based features extracted from each polyp candidate region are classified with SVM and judged if polyp region or not. The approach is not patch-based approach and it makes possible to detect polyp region.

This paper improves the previous approach<sup>7</sup> to have the higher accuracy of polyp detection based on the edge-based features. It is shown that increasing the number of examples to learn the classifier using Real Adaboost and Random Forests have performed higher accuracy in classification.

## 2. Extraction of Polyp Candidate Regions Using Hessian Filter

The convexity and concavity of the image intensity surface itself is used to select candidate polyp regions. A Hessian filter is applied to the endoscope image to determine regions of convex and concave image intensity. The image is segmented based on the results of Hessian filtering. Low pass filtering, via Gaussian smoothing, is included to remove the effects of noise and texture.

As a pre-processing, Fourier transform is applied to the gray scale image converted from the original RGB image to remove the high frequency components. Low pass filter is applied in the frequency space and inverse Fourier transform is applied. The effects of noise and texture of the surface can be removed and original image and low pass filtered image after the removal of high frequency components are shown in Fig.1 and Fig.2, respectively.

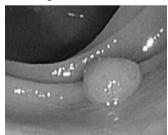


Fig. 1. Original Image

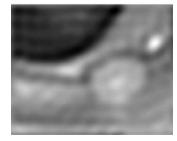


Fig. 2. Low Pass Filtered Image

In paper<sup>8</sup>, tube-like image features arising from blood vessels are enhanced according to the eigenvalues,  $\lambda_1$ ,  $\lambda_2$  of the associated Hessian matrix as shown in Table 1. Here, the procedure to enhance blob-like structure is as follows:

٠,	(H-Iligh, L-low, N-lloisy)					
	$\lambda_1$	$\lambda_2$	pattern and structure noisy, no preferred direction			
	N	N				
	L	$H_{-}$	tube structure (bright)			
	L	$H_{+}$	tube structure (dark)			
	$H_{-}$	$H_{-}$	blob-like structure (bright)			
	$H_{+}$	$H_{+}$	blob-like structure (dark)			

Table 1. Eigenvalues of the Hessian Matrix (H=high, L=low, N=noisy)

- Step 1. Generate a Gaussian filter, G(x, y), with (width) scale s.
- Step 2. Generate filters for the partial derivatives,  $G_x(x, y)$ ,  $G_y(x, y)$ , and the second partial derivatives,  $G_{xx}(x, y)$ ,  $G_{xy}(x, y)$  and  $G_{yy}(x, y)$ .
- Step 3. Compute  $L_{xx}(x,y)$ ,  $L_{xy}(x,y)$  and  $L_{yy}(x,y)$  as the convolution of input image, L(x,y), with  $G_{xx}(x,y)$ ,  $G_{xy}(x,y)$  and  $G_{yy}(x,y)$  respectively. That is

$$L_{xx}(x,y) = G_{xx}(x,y) * L(x,y)$$
(1)

$$L_{xy}(x,y) = G_{xy}(x,y) * L(x,y)$$
 (2)

$$L_{yy}(x,y) = G_{yy}(x,y) * L(x,y)$$
(3)

where \* denotes convolution.

Step 4. Compute the  $(2\times2)$  Hessian matrix image, **H**, from Eqs.(1-3).

$$\mathbf{H} = \begin{bmatrix} L_{xx} & L_{xy} \\ L_{xy} & L_{yy} \end{bmatrix} \tag{4}$$

Step 5. Calculate the eigenvalues,  $\lambda_1$ ,  $\lambda_2$ ,  $(\lambda_1 \le \lambda_2)$ , of **H** 

Step 6. Calculate the image, I(x, y), where

$$I(x,y) = \left(\lambda_1(x,y)^2 + \lambda_2(x,y)^2\right) \left(1 - \left(\frac{\lambda_1(x,y)}{\lambda_2(x,y)}\right)^2\right)$$
 (5)

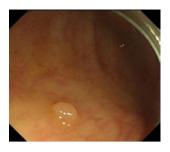
Again, the classification table for pattern and structure based on the eigenvalues of the Hessian matrix is shown in Table 1. Here, H corresponds to the high (largest in magnitude) eigenvalue and L corresponds to the low (smallest in magnitude) eigenvalue.  $H_+$  and  $H_-$  denote the sign of the high eigenvalue,  $H_+$  if H > 0 and  $H_-$  if H < 0, respectively. Recall that  $\lambda_1 \leq \lambda_2$ .

An example of an original endoscope image is shown in Fig.3. The corresponding image with blob-like structures enhanced is shown in Fig.4.

A Voronoi diagram based segmentation <sup>9</sup> is used to generate candidate polyp regions. Segmentation with conservative (i.e., lower) threshold values is shown in Fig.5. Red points correspond to points with highest values while points with blue points correspond to points with lowest value. Segmentation with higher threshold values is shown in Fig.6. The segmentation shown in Fig.6 determines the candidate polyp regions. Final classification for whether each candidate region is a polyp or not is determined subsequently.

# 3. Extracting HOG Features

Previous approach <sup>7</sup> tried the down sampling to have the patch with constant size so that the number of dimension of Histograms of Oriented Gradients (HOG) features <sup>10</sup> becomes constant. Proposed approach avoid down



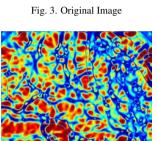


Fig. 5. Segmented Image

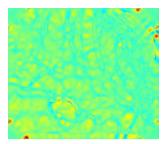


Fig. 4. Enhanced Image

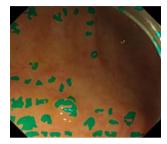


Fig. 6. Image of Candidate Polyp Regions

sampling and extracts local features with keeping the parameters of HOG features constant according to the size of candidate polyp region.

Endoscope image consists of polyp and non-polyp regions and change of color is small. Although it is not easy to treat the global edge features for the ellipse fitting, HOG features extract the oriented gradient of local intensity and it may be available to catch the difficult edge or shape conditions.

Gradient m(x, y) and its orientation  $\theta$  are calculated from the intensity I(x, u) at each pixel of input image.

$$m(x,y) = \sqrt{f_x(x,y)^2 + f_y(x,y)^2}$$
 (6)

$$\theta(x,y) = tan^{-1} \frac{f_y(x,y)}{f_x(x,y)} \tag{7}$$

$$\begin{cases}
f_x(x,y) = I(x+1,y) - I(x-1,y) \\
f_y(x,y) = I(x,y+1) - I(x,y-1)
\end{cases}$$
(8)

Weighted histogram of gradient of intensity is generated based on the gradient m and its direction  $\theta$  for each cell of  $p \times p$  pixels. Here  $\theta$  is quantized to N orientations, and the gradient m is added for each of orientation then histograms of oriented gradients are generated.

Finally, the histogram is normalized. Here each count of histograms are divided by the sum of HOG features in the cell  $q \times q$ .

$$v(n) = \frac{v(n)}{\sqrt{\sum_{k=1}^{q \times q \times N} v(k)^2 + \epsilon}}$$
(9)

where v(n) represents n-th HOG feature and  $\epsilon$  is a constant to avoid the division by 0. Moving this block with overlapping gives HOG features of input image.

For example, when the size of input image is  $h \times w$ , the number of dimension of HOG features become  $(\frac{w}{p} - q + 1)(\frac{h}{p} - q + 1) \times q^2 \times N$ . Visualization of HOG features is shown in Fig.7,8 as an example.



Fig. 7. Input Image



Fig. 8. HOG Features

#### 4. Selection of Classifiers

The number of dimensions becomes very large in HOG features and It was necessary for the approach, to introduce feature selection with SVM<sup>11</sup> to perform the classification. While in the proposed approach, RealAdaBoost<sup>12</sup> and Random Forests<sup>13</sup> are used as classifiers to perform efficient classification for the case of the larger number of dimension of features. This performance is evaluated for the comparison.

#### 4.1. RealAdaBoost

Multiple classifiers are used to the classification using boosting approach. Boosting is an approach to construct a strong classifier by combining multiple weak classifiers which compensate with other classifiers to reduce the misjudged data. Here RealAdaBoost is introduced as this purpose and probabilistic density function is used as a weak classifier.

#### 4.2. Construction of RealAdaBoost

Sample weight  $\vec{D_1}$  is initialized with  $\frac{1}{N}$  for the N learning samples of  $(\vec{x_i}, y_i)$ . Weighted histogram is generated for each feature of input data  $\vec{x_i}$ . Here features are quantized into N levels and sample weight  $D_t(i)$  is used as added weight.

$$W_{+}^{j} = \sum_{i,y_{i}=+1} D_{t}(i) \tag{10}$$

$$W_{-}^{j} = \sum_{i,v:=-1} D_{i}(i) \tag{11}$$

where  $W_+$  is weighted histogram of data for y = 1,  $W_-$  is weighted histogram of data for y = -1. When these two histograms are not similar, these are separable weak classifiers. Bhattacharyya distance is calculated for these two histogram and evaluation value for each weak classifier is calculated.

$$z = 1 - \sum_{j} \sqrt{W_{+}^{j} W_{-}^{j}} \tag{12}$$

Candidate weak classifier which gives the maximum of evaluation is taken the weak classifier at learning time t. Here the output of weak classifier is defined by Eq.(13) and  $\epsilon$  is a very small value to avoid division by 0.

$$h(x) = \frac{1}{2} \ln \frac{W_+^j + \epsilon}{W_-^j + \epsilon} \tag{13}$$

Weight of learning sample is updated and normalized using Eq.(14) according to the selected weak classifier.

$$D_{t+1}(i) = D_{t}(i)exp[-y_{i}h_{t}(x_{i})]$$

$$D_{t+1}(i) = \frac{D_{t+1}(i)}{\sum_{n=0}^{N} D_{t+1}(n)}$$
(14)

Sum of these weak classifiers becomes the output for unknown data.

#### 4.3. Random Forests

Random Forests is introduced to construct classifier with high accuracy by combining low correlated weak classifiers. Random forests use a decision tree as a weak classifier.

#### 4.3.1. Construction of Random Forests

The whole learning data is  $\vec{S}$  is learned with learning data  $\vec{S_0} \in \vec{S}$  which is randomly selected for each decision tree. Node of the learning data  $\vec{S_0}$  allows the overlapping of data. Split function at node j of learning data  $\vec{S_0}$  is given by Eq.(15).

$$h(\vec{\mathbf{v}}, \theta_i) \in \{0, 1\} \tag{15}$$

 $\vec{v}$  represents data which reached node j,  $\theta$  represents the parameter to decide the split function, where  $\theta = (\phi, \psi, \tau)$ .  $\phi$  means the filter which extracts several features from d-dimensional data  $\vec{v}$ ,  $\psi$  means the splitting criteria and  $\tau$  means the threshold for the split. Decision tree is constructed until data splitting cannot be done anymore.

Majority voting of these weak classifiers become the output for unknown data.

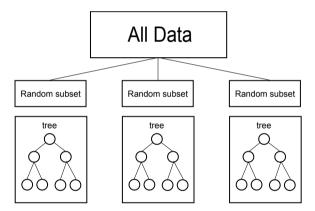


Fig. 9. Random Forests

#### 5. Integration of Detected Results

When the approach is applied for the detection of polyp in endoscope image, multiple rectangle for polyp candidate are detected. Here k-means++ is used to integrate the detection results for improving accuracy.

#### 5.1. k-means++

k-means++ <sup>14</sup> is an algorithm to find the cluster center. Cluster center corresponds to the point which minimizes the variance inside classes, in other words, the point which minimizes the squared sum of distance between the point and each data point inside a class. k-means++ is an improvement of k-means. Point is randomly selected

and cluster center is determined. After that, data which has not been selected to cluster center become cluster center with the probability which is proportional to the distance between its nearest neighbor cluster center and all data.

Example of result which k-means++ is applied to the random data in shown in Fig.10. Blue corresponds to input data, red corresponds to the cluster center and connected line corresponds to the result of clustering.

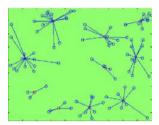


Fig. 10. k-means++

#### 6. Experiments

Endoscope image is used for evaluation of the proposed approach in the experiments.

#### 6.1. Data Set

156 Endoscope images are used in the experiments, where small size of polyp has around 1000 pixels while large size of polyp has around 13000 pixels. Endoscope image resolution consists of 1000×900 pixels. Mask image which was manually made is used, whose example is shown in Fig.12. Here one positive sample is generated from polyp region while four negative samples are generated from non-polyp region. HOG features are extracted from positive sample under the condition that the center of mask and the center of detection window becomes the same point. On the other hand, negative sample is randomly determined under the condition that mask is not included inside detection window, then HOG features are extracted.

Considering the variation for the appearance of polyp, important issue is data set was increased by rotating the endoscope image from 0 degree to 160 degrees with 20 degrees step. The number of positive samples was 1404 and that of negative samples was 5616 as data set.

1053 samples among 1404 positive samples were randomly used for learning data while the remained 351 positive samples were used for test data. 1053 samples among 5616 negative samples were randomly used for learning data while the remained 4563 negative samples were used for test data. The number of weak classifiers was set to be 6000 and quantization level was set to be 100 as learning parameters of Real Adaboost.

Number of decision tree was set to be 5000 and number of dimension of features used in the decision tree was 151.



Fig. 11. Endoscope Image

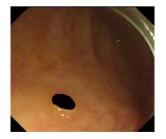


Fig. 12. Mask Image

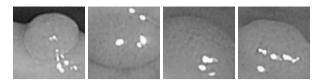


Fig. 13. Positive Samples

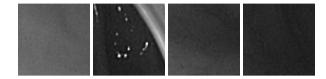


Fig. 14. Negative Samples

#### 6.2. Results

Experimental results are shown in Table 2 and Table 3. Sensitivity (Sen) represents the correct ratio for positive samples while Specificity (Spe) represents the correct ratio for negative samples. Learning data are randomly selected in each trial count.

$$Sensitivity = \frac{Number of Correct Positive Predictions}{Number of Positives}$$
$$S pecificity = \frac{Number of Correct Negative Predictions}{Number of Negatives}$$

Table 2. RealAdaBoost

a	daBoost							
	Times	Sen[%]	Spe[%]					
	1	100	97.7					
	2	99.1	96.5					
	3	99.1	97.2					
	4	100	97.4					
	5	98.5	97.1					
	6	100	96.8					
	7	98.8	97.7					
	8	98.2	99.1					
	9	99.1	99.1					
	10	100	97.4					

Table 3. RandomForests

l	irorests						
	Times	Sen[%]	Spe[%]				
	1	100	98.4				
	2	100	98.1				
	3	100	98.2				
	4	100	98.4				
	5	100	98.1				
	6	100	98.3				
	7	100	98.2				
	8	100	98.4				
	9	99.7	97.9				
	10	100	98.4				

It is shown that either of Real Adaboost and Random Forests performs high recognition ratio. Robust results are obtained based on the effect that the number of data set was increased with rotation of endoscope images which is robust to the appearance and various size of polyp.

## 6.3. Example of Polyp Detection

Polyp detection is shown as example. The same number of positive samples and negative samples were randomly selected as learning data set except the test image. Initial number of clusters was set to be 10 as

parameter of k-means++. Result of polyp detection before applying k-means++ is shown in Fig.15 and Fig.17, while that after applying k-means++ is shown is Fig.16 and Fig.18.

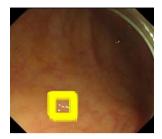


Fig. 15. Result before Integration



Fig. 16. Result after Integration

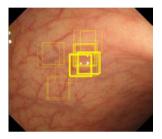


Fig. 17. Result before Integration

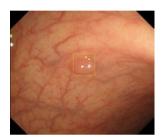


Fig. 18. Result after Integration

It is shown that polyp was correctly detected from endoscope image. It is also shown that integration of detection region by k-means++ works well.

#### 6.4. Incorrect Detection of Polyp

Examples of incorrect detection of polyp are shown in Fig.19 and Fig.20.



Fig. 19. Failed Example 1

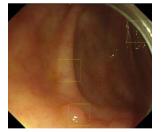


Fig. 20. Failed Example 2

The limitation of this proposed approach includes that negative samples are randomly selected from out of mask image from endoscope image. This sometimes affects the negative samples with low quality data. Region with specular reflectance or region with edge except polyp are some candidate where detection fails. It is considered that the situation will be improved if these incorrect detection results are used for learning and recognition again

#### 7. Conclusion

This paper proposed automatic detection of polyp using Hessian filter and HOG features. The approach introduced Real Adaboost and Random Forests as classifiers and it was shown that the approach performs the effective classification with the large dimensional number of features. k-means++ is also introduced to integrate the detection results with clustering. Increasing data with rotation of endoscope images are effective to give robust results for polyp detection.

As the approach tends to detect the polyp with over detection, further subjects remains for preparing the data set for negative samples including the points with specular reflectance points or candidate points of edge except polyp regions to save the over detection of polyp.

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