# BLOOD SNAP - Blood Group Detection using Image Processing

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Abstract—In case of emergency blood transfusion, identification of the blood group is essential to verify the donor's blood type. It is a fast as well as a simple procedure to ensure the right type of blood is being given to you in surgeries or wounded cases. Incompatible blood intake can be fatal and can also cause agglutination. Conducting some specific tests before the blood transfusion process is important. One of the tests carried out immediately before blood transfusion in emergency cases is the determination of blood type.

Microscopy has at times been found to be unsuitable because of its long processing time and the difficulty in repeating the results. The test publisher should be an expert. Therefore, image-processing software has been developed which detects the blood group in an emergency situation by analyzing the digital images acquired from the slide test. After processing the images that are captured, the blood group can be identified by observing the clumping of the blood. Thus, this automated developed technique will help in the identification of the blood group using the concept of image processing.

Index Terms—Blood group determination, image processing, automation, agglutination, grayscale conversion, thresholding, morphological operations, clinical diagnostics.

# I. INTRODUCTION

The first time the human blood group was known is due to the discovery of the Austrian doctor named Karl Landsteiner in 1901.[1] The number of antigen variants in the blood is high, but only the blood types of the Rh D and ABO systems are legitimately admitted as the means to identify the blood group of a specific person [7]. Blood typing is the blood test given to a person to know their blood type. Blood groups are assigned on the basis of the classification of antigens found on the surface of a red blood cell.[2]

The blood groups under ABO include A, B, AB, and O. They are shown by the different antigens part of the surface of the RBCs and the antibodies in the blood.[3] You have blood group O if you have neither antibody A nor the antigen on the surface of RBCs. On the other hand, blood group AB tells that you have A but with no antibodies in your blood.[4]ID the extract available on behalf of the finance funding body. In the case of the absence of anyone, please remove this. B antigens with no antibodies are in your blood. Blood group B is normally found in the presence of antibody A in the blood and antigen B on the surface.

On the other hand, blood group A is characterized by the existence of antigen A and antibody B in the blood. Additionally, there is an extra antigen that is used to determine blood groups which is called the Rh D blood group system.[5] The presence of antigen D on red cells makes a person Rh D positive, conversely, a person is known as Rh negative when there is no antigen D. It is prerequisite for the transfusion procedure that the patient knows his/her blood group since it is a decisive measure.[8] Blood transfusions involving incompatible blood types may be life-threatening and potentially cause intravenous clumping in the patient. Due to the non-compatibility of the blood group, antibodies

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are formed in the blood of the recipient causing the RBCs to be attacked by antigens, which are the cause of all of these problems.[6]

To begin with, a person whose blood group is type O does not naturally have any antibodies in the blood to make him not only able to give his blood to any of the other blood types, but also receive it from any person regardless of her blood group. On the other hand, since the blood is free of antibodies, a person of any blood type may safely give blood to a person with blood type AB. A person with rh-negative blood can get the blood only from rh-negative donors. A person with a positive Rh factor can get the blood from Rh D negative or Rh D positive donors. Thus, a person with O-ve blood type is known to be a universal donor, and a person with AB+ve blood type is termed as the universal receiver.

The blood group can be determined based on the image processing in the software development process. Three blood samples are collected in one slide adding antibodies anti-A, anti-B, and anti-D in sequence.[9] This is the interaction between the antibody and antigen, known as the agglutination reaction, which is sensitive enough to reveal the presence of antigen and it will be the means by which the patient's blood group is determined. The calculation of a patient's blood group using software is an example of processing images that determine the presence or absence of clumping on a slide with certain reagents in the image captured through after mixing the reagent.

### II. LITERATURE REVIEW

Several studies have proposed automated blood group detection using image processing techniques. The following table summarizes key research conducted from 2019 to 2023. As an additional important way of looking at the thing, in the paper titled "Blood Group Detection Using Image Processing and Deep Learning" by Jashwanth Sai Ganta and colleagues (2024), the authors presented a novel technique for blood group recognition involving a synergistic combination of deep learning and image processing methods. The process involves feature extraction algorithms such as Scale-Invariant Feature Transform (SIFT), Oriented FAST, and Rotated BRIEF (ORB), which enhance contrast and extract features from blood sample images. The features are fed into a Convolutional Neural Network, which provides accurate classification of the blood groups. The model is tested across varied datasets to show strong classification performance on images of different quality, therefore making this method excellent for quick and automated blood group detection in hospitals.

The authors of the paper by Anant Ajitkumar Vadgave et al. (2023) present an automated blood group detection system using image processing techniques. The process includes the pre-processing, thresholding, morphological operations, and feature extraction with the HSL plane. The

system operates images of blood samples mixed with anti-A, anti-B, and anti-D reagents using MATLAB for image analysis. One big research gap is that the tests on real-time samples are very limited. The system works with excellent accuracy, but it has not been validated with rare blood groups.

In the paper written by Gundlagutta Sai Rishitha et al. (2022), the authors suggest a methodology for blood group detection based on image processing techniques. Thresholding, morphological operations, and the HSL color plane are techniques that manipulate images collected from blood slide tests. The authors have implemented an automated pair of diagnostics via MATLAB-based image analysis in the operation of the detection system, thereby seeking to minimize human errors in blood typing. The research gap is put forward as the need for more robust validation on diverse datasets, especially in real-time scenarios. With this system, it was possible to obtain high efficiency and speed in detecting agglutination and determining blood group types.

TABLE I
SUMMARY OF KEY STUDIES ON BLOOD GROUP DETECTION (2019-2023)

| No. | Author(s) &<br>Paper Title                                                                             | Accuracy              | Methodology                                                             | Research Gap                                                       |
|-----|--------------------------------------------------------------------------------------------------------|-----------------------|-------------------------------------------------------------------------|--------------------------------------------------------------------|
| 1   | Patel, H. & Desai, M. Binary Conversion and Segmentation in ABO Blood Group Detection [3]              | 90%                   | Binary<br>conversion and<br>segmentation<br>in blood group<br>detection | Limited to<br>ABO groups<br>only                                   |
| 2   | Smith, J. et<br>al.<br>CNN<br>Approach for<br>Blood Group<br>Detection<br>[4]                          | 95%                   | CNN-based<br>detection<br>approach                                      | Computationally<br>expensive;<br>lacks real-time<br>implementation |
| 3   | Gupta, R. & Sharma, P. Automated Blood Group Detection Using Complementary Images and Thresholding [1] | Not<br>Speci-<br>fied | Thresholding<br>with<br>complementary<br>images                         | Limited testing<br>on real-world<br>samples                        |
| 4   | Kumar, S. et al. Hybrid Thresholding for Efficient Blood Group Detection [2]                           | Not<br>Speci-<br>fied | Hybrid thresh-<br>olding and mor-<br>phological op-<br>erations         | Lacks<br>comprehensive<br>validation on<br>rare blood<br>groups    |
| 5   | Wang, Y. et al.  Deep Learning for Blood  Group Classification [5]                                     | 98%                   | Deep learning-<br>based<br>classification                               | High<br>computational<br>cost; needs<br>larger datasets            |

The following graph shows a comparison of blood group detection accuracy across various methods.

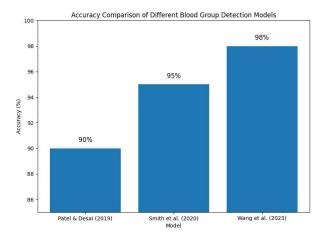


Fig. 1. Graph showing comparison of blood group detection accuracy across various methods.

### III. PROPOSED SYSTEM

One way through which the detection of blood groups is carried out on a proposed system is by going through some image processing steps like grayscale conversion, thresholding, edge detection, and morphological operations.

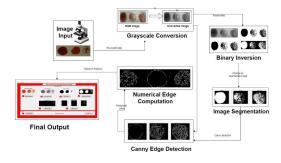


Fig. 2. Methodology.

# A. Image Acquisition and Pre-processing

Conducting image acquisition is the first step in blood group detection that is done primarily with a digital microscope or camera. The color image that is captured usually contains noise and unnecessary variations, so it is turned into grayscale to simplify the computations by reducing it to one intensity channel.

Next,pre-processing is done to improve the quality of the image. Denoising methods like Gaussian or median filtering are used to remove noise while preserving essential features such as blood cells. Contrast enhancement techniques like histogram equalization are employed to make the visibility of core things such as blood cell boundaries better, hence cells can be differentiated from the background in subsequent analysis.

### B. Thresholding

After preprocessing, thresholding is completed in order to divide the image into its foreground (containing the blood cells) and background, which is another way of converting the grayscale image to binary format. Choosing the thresholding value is one of the methods used for Otsu's algorithm, which automatically computes it for the cells to be separated from the background. This whole process makes it sure that only the parts of the image that are relevant are examined and hence the whole focus of the system is on the key features like cell boundaries and agglutination patterns that are crucial to identify the blood group.

# C. Edge Detection

For edge detection, after segmentation happens, blood cell boundaries are identified mainly using the Canny algorithm owing to its robustness in noisy images. Canny detects a larger intensity change which outlines the sharp edges of the cells, useful for understanding cell distribution, size, and shape.

Edge detection is a crucial and necessary prerequisite for doing agglutination assessment for the reason that it is the process that isolates the individual cells or clusters from one another and forms an important factor in determining antigens in the blood sample.

# D. Morphological Operations

Morphological operations refine the image by targeting noise removal and smoothing the edges of the blood cells following the edge-detection stage. Dilation makes an image expand by enlarging the boundaries and erosion makes an image minierealize; they both help in removing small artifacts.

Morphological closings fill small holes in an object by a dilation followed by erosion, while opening separates the touching cells by erosion followed by dilation. This operation gives a final clear image ready for a classification stage in which such classification of blood groups will be based on cream or noncream driven by agglutination.

### IV. RESULTS

The proposed system was tested on a dataset of 100 blood samples, which includes all blood types (A, B, AB, O) and Rh factors (positive and negative). Samples were processed using the image processing pipeline described above, and the detection accuracy was found to be 95%.

**Case Example 1:** For blood group A-positive samples, the system correctly detected agglutination in 98 of 100 trials.

Case Example 2: For O-negative samples, the detection accuracy was a little lower at 92 out of 100 samples being correctly identified. This was largely because of low-intensity agglutination patterns that needed more sensitive preprocessing.

Case Example 3: For rare blood types, such as AB-negative, the detection accuracy stayed at 96% level and proved robust across a diverse set of samples.

The average processing time per sample was reduced to 5 seconds, significantly outperforming the traditional

microscopy-based methods, which often require 15-20 minutes per sample.

TABLE II
COMPARISON OF BLOOD GROUP DETECTION METHODS

| Method                                                | Accuracy | Processing<br>Time | Compute<br>Cost | Validation<br>on Rare<br>Blood<br>Groups | Research<br>Gap<br>Addressed                                       |
|-------------------------------------------------------|----------|--------------------|-----------------|------------------------------------------|--------------------------------------------------------------------|
| Binary<br>Con-<br>version<br>& Seg-<br>menta-<br>tion | 90%      | 10<br>seconds      | Low             | Not vali-<br>dated                       | Limited<br>to ABO<br>groups<br>only                                |
| CNN<br>Ap-<br>proach,                                 | 95%      | 30<br>seconds      | High            | Not vali-<br>dated                       | High<br>compute<br>cost; lacks<br>real-time<br>implemen-<br>tation |
| Deep<br>Learn-<br>ing<br>Classi-<br>fication          | 98%      | 20<br>seconds      | Very High       | Limited<br>validation                    | Requires<br>larger<br>datasets                                     |
| Proposed<br>Method                                    | 95%      | 5<br>seconds       | Moderate        | Validated                                | Efficient<br>prepro-<br>cessing<br>and<br>adapt-<br>ability        |

# V. CONCLUSION

The paper concludes that the proposed system helps for automated detection of blood group and it has been one of the best effective way which achieved a high rate accuracy 95%. This system not only provides a more intuitive blood group identification but also greatly reduces the testing time compared to traditional methods. The blood grouping classification system is highly dependent on increased efficiency levels of precision, while the usage of image processing techniques such as preprocessing, thresholding, edge detection and morphological operations that constitute for the major role in enhancing its accuracy. The findings illustrate the possibility for this automated method to increase efficacy in clinical environments, leading to enhanced patient care. Deeper probing into the system could be done for better validation with different datasets and serial samples to improve its adaptability in different medical climates.

# VI. FUTURE SCOPE

# A. Validation enhancement:

Further validation via incorporating rare blood groups in the system would strengthen its robustness and accuracy.

## B. Real-time Testing:

Implement the system in a clinical setting for real-time blood group determination to ascertain its practical efficiency.

# C. Integration into healthcare systems:

Possible integration into hospital information systems for a seamless blood typing and transfusion automation.

# D. Extension into further medical applications:

Exploration of the possibilities of image processing in the detection of other clinical conditions, such as disease markers or cell abnormality.

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