

Research Article

Mean Platelet Volume and Platelet Distribution Width as Markers in the Diagnosis of Acute Gangrenous Appendicitis

Zhe Fan,¹ Jiyong Pan,¹ Yingyi Zhang,¹ Ziyi Wang,² Ming Zhu,² Baoshun Yang,² Lei Shi,³ and Huirong Jing²

¹Department of General Surgery, The Third People's Hospital of Dalian, Non-Directly Affiliated Hospital of Dalian Medical University, Dalian 116033, China

²Department of General Surgery, The Second Hospital of Dalian Medical University, Dalian 116023, China ³Department of General Surgery, The Second Hospital of Dalian Medical University, Northern Yard, Dalian 116023, China

Correspondence should be addressed to Huirong Jing; jinghuirongdalian@hotmail.com

Received 10 July 2015; Revised 16 October 2015; Accepted 28 October 2015

Academic Editor: Irene Rebelo

Copyright © 2015 Zhe Fan et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

Introduction. Acute gangrenous appendicitis (AGA) is a common medical condition; however, the grade of appendicitis usually cannot be established preoperatively. We have attempted to identify some indicators, such as the mean platelet volume (MPV) and the platelet distribution width (PDW), to diagnose AGA. *Aims.* To evaluate whether or not the MPV and PDW are suitable markers to diagnose AGA. *Methods.* A retrospective study of 160 patients with AGA and 160 healthy patients was undertaken. Disease diagnosis was confirmed based on the pathologic examination of surgical specimens. Patient white blood cell (WBC) count, neutrophil ratio (NR), platelet (PLT) count, MPV, PDW, and hematocrit (HCT) were analyzed. Receiver operating characteristic (ROC) curves were used to evaluate the sensitivity and specificity of these indices in AGA. *Results.* There were no significant differences between the AGA and control groups in age and gender. Compared to the control group, the WBC count, NR, and PDW were significantly higher (P < 0.001, resp.) and the MPV and HCT were significantly lower (P < 0.001, resp.) in the AGA group. The diagnostic specificities of the WBC count, NR, PLT count, MPV, PDW, and HCT were 86.3%, 92.5%, 58.1%, 81.7%, 83.9%, and 66.3%, respectively. Therefore, the NR had the highest diagnostic specificity for the diagnosis of AGA. *Conclusions.* This is the first study to assess the MPV and PDW in patients with AGA. Our present study showed that the MPV is reduced and the PDW is increased in patients with AGA; the sensitivity of PDW was superior to the MPV. A decreased MPV value and an increased PDW could serve as two markers to diagnose AGA. The NR had the highest specificity for the diagnosis of AGA.

1. Introduction

Acute appendicitis is a common medical condition encountered in general surgery. There are three grades to illustrate acute appendicitis (focal appendicitis, suppurative appendicitis, and gangrenous appendicitis) [1, 2]. The clinical diagnosis of acute appendicitis is often based on medical history, physical examination, blood tests, and iconographic examination [3]. Timely emergency surgery should be carried out to avoid peritonitis, especially in patients with suppurative or gangrenous appendicitis, before perforation occurs [1]; however, the grade of appendicitis cannot usually be evaluated based on the above examinations. Platelets (PLTs) have effects on haemostasis and regulate inflammatory events. PLTs are more highly activated when inflammatory mediators are released [4]. The mean platelet volume (MPV) and platelet distribution width (PDW) are two PLT parameters of the complete blood count [5]. After PLT production increases, MPV changes accordingly [4–6]. It has been reported that the MPV is affected in patients with acute pancreatitis, ulcerative colitis, rheumatoid arthritis, and ankylosing spondylitis [4, 6, 7].

In the present study, we focused on the relationship between MPV and PDW and acute gangrenous appendicitis (AGA). Dinc et al. [8] suggested the PDW as a new index in the diagnosis of acute appendicitis, and we verified changes in the PDW in patients with AGA.

2. Materials and Methods

One hundred and sixty patients with AGA and 160 healthy people were enrolled in our study. AGA was diagnosed based on postoperative pathologic examination. The patients in the AGA group were treated at The Third People's Hospital of Dalian between June 2011 and June 2014 and healthy patients (control group) with normal physical examinations were enrolled from our physical examination center during the same period. This retrospective study was approved by the hospital ethics committee.

Patients with the following conditions were excluded from the study: <15 years of age; alcohol consumption; cigarette smoking; diabetes mellitus; hypertension; morbid obesity; and severe comorbidities (heart failure, peripheral vascular disease, hematologic disorders, acute or chronic infections, cancer, and hepatic disease) [9, 10].

All blood samples were collected into tubes containing EDTA (potassium ethylenediaminetetraacetate) through the cephalic vein and assayed using internationally certified devices. All results were available in <10 min [8].

The white blood cell (WBC) count, NR, PLT count, MPV, PDW, and hematocrit (HCT) were collected from AGA and control groups.

The reference values were $4-10 \times 10^9$ /L for WBC, 40% - 70% for NR, $100-300 \times 10^9$ /L for PLT, 7.6–13.2 fL for the MPV, 12%–16.5% for the PDW, and 40%–50% for the HCT.

2.1. Statistical Analysis. Research data were analyzed using SPSS 20.0 software (SPSS for Windows; SPSS, Inc., Chicago, IL, USA). The continuous data are presented as the mean \pm standard deviation (SD), Student's *t*-test was used for the comparison between the two groups, and a χ^2 test was used for comparing count data between the two groups. A normal distribution was analyzed using binary logistic analysis. Logistic analysis and receiver-operating curve (ROC) analysis were used to describe the parameters in the AGA and control groups. The results were examined within the 95% CI, and a P < 0.05 was considered statistically significant.

3. Results

The mean ages of the patients were 45.6 ± 19.6 years (range, 14–89 years) in the AGA group and 43.0 ± 12.5 years (range, 15–87 years) in the control group. There were no significant differences between the AGA and control groups with respect to age and gender (Table 1).

The mean WBC counts in the AGA and control groups were $13.06 \pm 4.64 \times 10^9$ /L (range, $1.81-25.66 \times 10^9$ /L) and $6.12 \pm 1.62 \times 10^9$ /L (range, $3.26-11.49 \times 10^9$ /L; *P* < 0.001).

NR in the AGA and control groups was $84.21 \pm 9.34\%$ (range, 43.3-97.24%) and $54.57\pm8.11\%$ (range, 32.62-75.71%; P < 0.001).

The PLT counts in the AGA and control groups were $210.29 \pm 60.41 \times 10^9$ /L (range, $90-428 \times 10^9$ /L) and $220.09 \pm 45.28 \times 10^9$ /L (range, $105-348 \times 10^9$ /L; P = 0.102 > 0.05).

TABLE 1: Demographic data for patients with AGA and control groups.

Characteristic	Control group	AGA group	P value
Age	43.0 ± 12.5 years	45.6 ± 19.6 years	0.155
Gender (male/female)	99/61	102/58	0.644

TABLE 2: Laboratory results for patients with AGA and control groups.

Indexes	Control group	AGA group	P value
WBC	$6.12 \pm 1.62 \times 10^9/L$	$13.06 \pm 4.64 \times 10^9/L$	0.000
NR	$54.57 \pm 8.11\%$	$84.21 \pm 9.34\%$	0.000
PLT	$220.09 \pm 45.28 \times 10^9/L$	$210.29 \pm 60.41 \times 10^9 / L$	0.102
MPV	$10.91 \pm 2.72 \times 10^9/L$	$9.21 \pm 1.38 \times 10^9 / L$	0.000
PDW	$12.50 \pm 1.93 \times 10^9/L$	$15.25 \pm 1.90 \times 10^9/L$	0.000
HCT	$43.92 \pm 3.77\%$	$40.93\pm5.48\%$	0.000

The MPV in the AGA and control groups were 9.21 \pm 1.38 × 10⁹/L (range, 6.6–12.9 × 10⁹/L) and 10.91 \pm 2.72 × 10⁹/L (range, 8.82–43.6 × 10⁹/L; *P* < 0.001).

The PDW in the AGA and control groups was $15.25 \pm 1.90 \times 10^9$ /L (range, 9.9–18.1 × 10⁹/L) and $12.50 \pm 1.93 \times 10^9$ /L (range, 1.3–19.1 × 10⁹/L; *P* < 0.001).

The HCT in the AGA and control groups was 40.93 \pm 5.48% (range, 24.82–52.5%) and 43.92 \pm 3.77% (range, 33.4–52.4%; *P* < 0.001) (Table 2).

Based on the results in Table 2, the WBC counts, NR, and PDW were significantly increased (P < 0.001) and the MPV and HCT were significantly decreased (P < 0.001) in the AGA group compared to the control group. There were no significant changes in PLT count (P > 0.05).

Binary logistic analysis was carried out and the results were as follows: MPV (P = 0.000); PDW (P = 0.000); PLT (P = 0.012); HCT (P = 0.001); neutrophil ratio (NR) (P = 0.026); and WBC (P = 0.024) (P < 0.05, resp.).

The ROC curves were analyzed for the following indices, as shown in Figure 1:

Area under curve (AUC): MPV (0.817), PDW (0.839), PLT count (0.581), HCT (0.663), NR (0.975), and WBC count (0.923);

Cut-off value: MPV ($9.6 \times 10^{9}/L$), PDW ($15.1 \times 10^{9}/L$), PLT count ($179 \times 10^{9}/L$), HCT (40.3 %), NR (69.5 %), and WBC count ($8.45 \times 10^{9}/L$);

Sensitivity and specificity: [MPV (66.25% and 91.19%), PDW (76.3% and 93.1%), PLT count (33.13% and 83.02%), HCT (41.88% and 82.39%), NR (92.5% and 96.9%), and WBC count (86.3% and 92.5%)] (Table 3).

4. Discussion

Acute appendicitis often presents as an acute abdomen. In recent studies [9, 11, 12], researchers considered the MPV to be helpful in diagnosing acute appendicitis; however, opinions have not been consistent. Previous studies [8, 9, 11] have concluded that acute appendicitis can induce changes

TABLE 3: ROC curves/cut-off values/sensitivity/specificity withAGA and control groups.

Indexes	AUC	Cut-off value	Sensitivity	Specificity
WBC	0.923	$8.45 \times 10^9/L$	86.3%	92.5%
NG	0.975	69.5%	92.5%	96.9%
PLT	0.581	$179 \times 10^9/L$	33.13%	83.02%
MPV	0.817	$9.6 \times 10^9 / L$	66.25%	91.19%
PDW	0.839	$15.1 \times 10^{9}/L$	76.3%	93.1%
HCT	0.663	40.3%	41.88%	82.39%

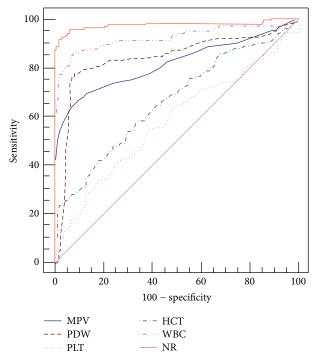


FIGURE 1: ROC curves for MPV, PDW, PLT count, HCT, NR, and WBC count.

in the MPV or PDW; however, the conclusions drawn in the two studies about MPV were not consistent. Perhaps there is another reason to account for the discrepant results, such as ethnicity and geographic influence. Erdem et al. [9] reported that the MPV was markedly lower in acute appendicitis groups compared to control groups. In contrast, Narci et al. [11] concluded that the MPV was significantly higher in acute appendicitis groups compared with control groups. Uyanik et al. [12] concluded that the MPV has no diagnostic value with respect to acute appendicitis. Based on our analysis of the recent literature, we suggest that there is no relationship between the MPV and acute appendicitis. Classic symptoms and physical examination findings could be used to diagnose acute appendicitis; however, it is difficult to distinguish gangrenous cases from several types of appendicitis. Therefore, we focused on the MPV in the diagnosis of AGA. A MEDLINE search for articles in the English language from 1981 to 2014 with the terms "MPV"/"PDW" and "acute gangrenous appendicitis" revealed no entries.

MPV has been studied as an inflammatory marker in several diseases. MPV represents an index of PLT function. An increase in "young" platelets and an aggregation of large platelets could lead to higher MPV values. PLT size and activity are influenced by cytokines, such as IL-3 or IL-6 [11]. In many chronic diseases, the MPV increases, while in many acute diseases the MPV decreases [9]. Specifically, the MPV decreases in patients with ulcerative colitis, rheumatoid arthritis, and ankylosing spondylitis [8, 13, 14], and the MPV increases in patients with ankylosing spondylitis, familial Mediterranean fever, Behcet's disease, and psoriasis [15, 16]. In the current study, the MPV was markedly decreased in patients with AGA, in agreement with Erdem et al. [9]. In our research, the sensitivity of MPV in the current study was slightly lower than that of PDW, which indicates that the MPV could be as a diagnostic marker for acute appendicitis or AGA. The PDW can represent the heterogeneity of thrombocyte volume [10]. There are two studies [10, 17] which illustrate the correlation between the PDW and acute appendicitis; both studies demonstrated an increase in the PDW. Our study was the third study to show an increase in the PDW in patients with acute appendicitis and to show an increase in the PDW in patients with AGA.

In the present study, a decrease in the HCT was shown in patients with AGA. This is the first study to show a relationship between HCT and AGA. A MEDLINE search for "hematocrit" and "inflammation" revealed no entries. Indeed, alternate mechanisms may exist and additional research should be conducted.

The increases in the WBC counts and neutrophil-tolymphocyte ratio have been used for the diagnosis of acute appendicitis [1, 9, 18]. Similarly, our research showed that the NR has the best sensitivity (92.5%); the sensitivities of the WBC count and PDW were 86.3% and 76.3%, respectively.

Based on the work of Albayrak et al. [10] and our results herein, the PDW may be a new marker to diagnose acute appendicitis. In the present study, the PDW increased in patients with AGA. A new diagnostic algorithm for the diagnosis of AGA may include the WBC count, NR, and PDW.

Although some factors have been excluded from the study, there were some limitations, including the small sample size, and the onset of acute appendicitis was not recorded.

5. Conclusion

Based on the current study, the MPV is a new index for diagnosing AGA. While the MPV was clearly lower in patients with AGA, the MPV did not have a higher sensitivity compared with the PDW. We confirmed that the PDW is a new, highly sensitive parameter with which we diagnose AGA. The WBC count and NR have high sensitivity in diagnosing AGA.

Abbreviations

- MPV: Mean platelet volume
- PDW: Platelet distribution width
- PLT: Platelet count
- NR: Neutrophil ratio.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

Authors' Contribution

Zhe Fan and Jiyong Pan are co-first authors with equal contribution to the present study.

References

- M. Ishizuka, T. Shimizu, and K. Kubota, "Neutrophil-tolymphocyte ratio has a close association with gangrenous appendicitis in patients undergoing appendectomy," *International Surgery*, vol. 97, no. 4, pp. 299–304, 2012.
- [2] S. Emil, F. Gaied, A. Lo et al., "Gangrenous appendicitis in children: a prospective evaluation of definition, bacteriology, histopathology, and outcomes," *Journal of Surgical Research*, vol. 177, no. 1, pp. 123–126, 2012.
- [3] P. Chaudhary, A. Kumar, N. Saxena, and U. C. Biswal, "Hyperbilirubinemia as a predictor of gangrenous/perforated appendicitis: a prospective study," *Annals of Gastroenterology*, vol. 26, no. 4, pp. 325–331, 2013.
- [4] A. Incebiyik, A. Seker, M. Vural, N. Gul Hilali, A. Camuzcuoglu, and H. Camuzcuoglu, "May mean platelet volume levels be a predictor in the diagnosis of pelvic inflammatory disease?" *Wiener Klinische Wochenschrift*, vol. 126, no. 13-14, pp. 422–426, 2014.
- [5] Y. Beyazit, A. Sayilir, S. Torun et al., "Mean platelet volume as an indicator of disease severity in patients with acute pancreatitis," *Clinics and Research in Hepatology and Gastroenterology*, vol. 36, no. 2, pp. 162–168, 2012.
- [6] F. Catal, C. Tayman, A. Tonbul et al., "Mean Platelet Volume (MPV) may simply predict the severity of sepsis in preterm infants," *Clinical Laboratory*, vol. 60, no. 7, pp. 1193–1200, 2014.
- [7] J. J. Yang, S. Y. Cho, H. J. Ahn, H. J. Lee, W. I. Lee, and T. S. Park, "Mean platelet volume in acute appendicitis: a gender difference," *Platelets*, vol. 25, no. 3, pp. 226–267, 2014.
- [8] B. Dinc, A. Oskay, S. E. Dinc, B. Bas, and S. Tekin, "New parameter in diagnosis of acute appendicitis: platelet distribution width," *World Journal of Gastroenterology*, vol. 21, no. 6, pp. 1821–1826, 2015.
- [9] H. Erdem, R. Aktimur, S. Cetinkunar et al., "Evaluation of mean platelet volume as a diagnostic biomarker in acute appendicitis," *International Journal of Clinical and Experimental Medicine*, vol. 8, no. 1, pp. 1291–1295, 2015.
- [10] Y. Albayrak, A. Albayrak, F. Albayrak et al., "Mean platelet volume: a new predictor in confirming acute appendicitis diagnosis," *Clinical and Applied Thrombosis/Hemostasis*, vol. 17, no. 4, pp. 362–366, 2011.
- [11] H. Narci, E. Turk, E. Karagulle, T. Togan, and K. Karabulut, "The role of mean platelet volume in the diagnosis of acute appendicitis: a retrospective case-controlled study," *Iranian Red Crescent Medical Journal*, vol. 15, no. 12, Article ID e11934, 2013.
- [12] B. Uyanik, C. Kavalci, E. D. Arslan et al., "Role of mean platelet volume in diagnosis of childhood acute appendicitis," *Emergency Medicine International*, vol. 2012, Article ID 823095, 4 pages, 2012.
- [13] O. Yüksel, K. Helvac, Ö. Başar et al., "An overlooked indicator of disease activity in ulcerative colitis: mean platelet volume," *Platelets*, vol. 20, no. 4, pp. 277–281, 2009.

- [14] B. Kisacik, A. Tufan, U. Kalyoncu et al., "Mean platelet volume (MPV) as an inflammatory marker in ankylosing spondylitis and rheumatoid arthritis," *Joint Bone Spine*, vol. 75, no. 3, pp. 291–294, 2008.
- [15] A. Y. Gasparyan, L. Ayvazyan, D. P. Mikhailidis, and G. D. Kitas, "Mean platelet volume: a link between thrombosis and inflammation?" *Current Pharmaceutical Design*, vol. 17, no. 1, pp. 47–58, 2011.
- [16] D. S. Kim, J. Lee, S. H. Kim, S. M. Kim, and M. G. Lee, "Mean platelet volume is elevated in patients with psoriasis vulgaris," *Yonsei Medical Journal*, vol. 56, no. 3, pp. 712–718, 2015.
- [17] A. Aydogan, S. Akkucuk, S. Arica et al., "The analysis of mean platelet volume and platelet distribution width levels in appendicitis," *Indian Journal of Surgery*, 2013.
- [18] M. Keskek, M. Tez, O. Yoldas et al., "Receiver operating characteristic analysis of leukocyte counts in operations for suspected appendicitis," *The American Journal of Emergency Medicine*, vol. 26, no. 7, pp. 769–772, 2008.



The Scientific World Journal



Gastroenterology Research and Practice





Journal of Diabetes Research



Disease Markers



Immunology Research









BioMed **Research International**





Computational and Mathematical Methods in Medicine





Behavioural Neurology



Complementary and Alternative Medicine











Oxidative Medicine and Cellular Longevity