



ELSEVIER

# Revista Brasileira de Hematologia e Hemoterapia

## Brazilian Journal of Hematology and Hemotherapy

[www.rbhh.org](http://www.rbhh.org)

### Update article

# Are the review criteria for automated complete blood counts of the International Society of Laboratory Hematology suitable for all hematology laboratories?



Samuel Ricardo Comar\*, Mariester Malvezzi, Ricardo Pasquini

Universidade Federal do Paraná (UFPR), Curitiba, PR, Brazil

#### ARTICLE INFO

##### Article history:

Received 13 October 2013

Accepted 2 January 2014

Available online 3 April 2014

##### Keywords:

Automation

Blood cell count

Validation studies

Hematology

#### ABSTRACT

**Objective:** to verify whether the review criteria for automated blood counts suggested by the International Consensus Group for Hematology Review of the International Society for Laboratory Hematology are suitable for the Hematology Laboratory of Hospital de Clínicas, Universidade Federal do Paraná.

**Methods:** initially, the review criteria of the International Society for Laboratory Hematology were adapted due to limitations in the Institution's electronic hospital records and interfacing systems. The adapted review criteria were tested using 1977 samples. After this first assessment, an additional 180 inpatient samples were analyzed to evaluate the screening criteria of the review criteria in conjunction with positive smear findings established by the institution. The performance of the review criteria was verified by determining false positive, false negative, true positive and true negative rates, sensitivity, specificity, positive predictive value, negative predictive value, microscopic review rate and efficiency.

**Results:** initial analysis showed false negatives = 6.73%, false positives = 23.27%, microscopic review rate = 46.03% and efficiency = 70.0%. An evaluation of the screening criteria adapted from the review criteria together with the positive smear findings of the institution showed false negatives = 15.5%, false positives = 10.5%, microscopic review rate = 37.3% and efficiency = 73.8%. In both situations the safety limit (false negative <5%) recommended by the review criteria was exceeded.

**Conclusions:** the review criteria adapted from the International Society for Laboratory Hematology are neither suitable nor safe for use in the hematology laboratory of the Hospital de Clínicas. This implies a need to develop and validate institution-specific review criteria in order to decrease false negative results to an acceptable and safe rate for patients.

© 2014 Associação Brasileira de Hematologia, Hemoterapia e Terapia Celular. Published by Elsevier Editora Ltda. All rights reserved.

\* Corresponding author at: Rua Padre Camargo, 280 – Alto da Glória, Laboratório de Hematologia, Unidade de Apoio Diagnóstico, Hospital de Clínicas, Universidade Federal do Paraná (UFPR), 80.060-240 Curitiba, PR, Brazil.

E-mail address: [srcomar@ufpr.br](mailto:srcomar@ufpr.br) (S.R. Comar).

<http://dx.doi.org/10.1016/j.bjhh.2014.03.011>

1516-8484/© 2014 Associação Brasileira de Hematologia, Hemoterapia e Terapia Celular. Published by Elsevier Editora Ltda. All rights reserved.

## Introduction

In 2005, the International Society for Laboratory Hematology (ISLH) through the International Consensus Group for Hematology Reviews, founded by hematologist Berend Houwen, published a set of 41 rules applicable as criteria for the review of automated complete blood counts (CBCs) and leukocyte differential results of automated hematology analyzers, i.e., review criteria for automated complete blood counts (RC).<sup>1</sup> These guidelines were formulated with the aims of reducing costs and the turnaround time of the results without sacrificing their quality, and justifying the performance and skills of the multiparametric hematology analyzers.<sup>2-4</sup> Since then, the rules suggested by the ISLH<sup>2</sup> have been considered the international standard to indicate situations requiring a blood smear review (BSR). They take into account the age and gender of patients, whether the request for CBC is the initial or a subsequent one to monitor CBCs, or whether there are significant differences between the current results, and previously validated and released results.<sup>2,4</sup> In practice, they are based on the set of screening thresholds for the results given by the analyzers and in the presence or absence of suspect flags. The aim is to distinguish samples with a high probability of containing relevant morphological alterations for the diagnosis and treatment of patients. When the CBC results do not meet the screening criteria, there are recommended procedures to follow, specifically to prepare an adequate peripheral blood smear for microscopic analysis.<sup>2</sup>

Hospital de Clínicas of the Universidade Federal do Paraná (HC-UFPR) is a general Class IV hospital according to the hospital classification system of Brazil's publically funded healthcare system (SUS); it is the largest provider of government healthcare services in the State of Paraná with 510 beds. Moderately to highly complex procedures are carried out in 59 departments. Approximately 61,000 consultations are made per month. The clinical hematology laboratory is located in the Diagnosis Support Service and contains two types of hematology analyzers: the Sysmex XE-2100D and XT-2000i (Sysmex Corporation, Kobe, Japan). Approximately 500 samples are sent for CBCs daily. Prior to the development of the RC, 100% of CBCs were analyzed microscopically, which led to delays in the release of the results even when performed by experienced professionals.

According to Bain<sup>5</sup> because BSRs and manual differential leukocyte counts (MDLCs) are laborious and expensive, they should be based on the RC. Thus, all hematology laboratories must be encouraged to establish locally valid protocols indicating when a BSR and MDLC should be performed. The guidelines suggested by the ISLH can be the starting point as long as they are interpreted in consideration of the experience of the laboratory staff, sophistication of the hematology analyzers and the laboratory's electronic records system, and incidences of abnormalities and variations in reference values of the population being tested.<sup>6,7</sup> Thus, this study evaluated the implementation of the RC suggested by the ISLH in the HC-UFPR Hematology Laboratory in order to determine automated thresholds such that microscopic analyses are performed only under special circumstances. In addition, the study aimed to define whether such guidelines could be tailored to the

population served or whether there is a need to establish and evaluate specific RC for this Institution.

## Methods

### Study site and sample preparation

The investigation was conducted in the Hematology Laboratory of HC-UFPR after approval by the local Ethics Committee. The samples were obtained in two stages. First, for five consecutive days, all laboratory samples were collected after the release of the results into the electronic hospital records system. A total of 1977 whole-blood samples in ethylenediaminetetraacetic acid (EDTA)-K<sub>2</sub> (1.8 mg/mL) were analyzed within 3 h of collection. Of these, 1573 and 404 were analyzed using the XE-2100D and XT-2000i hematology analyzers, respectively. Furthermore, to evaluate the screening criteria adapted from the ISLH together with positive smear findings (PSFs) of the HC-UFPR, an additional 180 inpatient samples were collected randomly and analyzed using the XT-2000i device; these samples were more likely to have PSFs because they also had abnormal CBC results. The PSFs elaborated by the HC-UFPR were intended to ensure clinically significant abnormalities were not omitted from the results, thereby establishing a minimum threshold of information that should be reported in the CBC results according to local consensus. All numerical data and information from suspect flags and blood smear findings were recorded. Approximately 70% of the samples tested were from outpatients, many of whom were having their first blood count. The other 30% were from inpatients from various hospital units (e.g., hematology, chemotherapy, infectious diseases, intensive care units, emergency care), many of whom had their blood counts monitored daily.

### Adaptation of the review criteria of the International Society for Laboratory Hematology according to local requirements

In order to determine whether the performance of these RC met local requirements or indicated the need to develop specific RC, the screening criteria and PSFs suggested by the ISLH were initially evaluated. However, changes were made to tailor the ISLH screening criteria to the hematology analyzers used in this study and particularly to adapt them to the electronic hospital records system. The main adaptations were associated with Delta Check rules, which were not possible to implement because of limitations of the institution's electronic hospital records and interfacing systems. The adapted ISLH screening criteria concerning the possibility of local implementation are shown in [Table 1](#). [Table 2](#) shows the PSFs recommended by the ISLH.<sup>2</sup> PSFs that differ from those recommended by the ISLH were also created for the HC-UFPR ([Table 3](#)) in an attempt to meet local requirements.

### Sample classification criteria

The criteria followed to select samples for review were compared with the findings of the peripheral BSR. A sample was classified as true positive (TP) if it was positive for a particular

**Table 1 – Adapted International Society of Laboratory Hematology review criteria for automated complete blood counts.**

Criterion or rule	Parameter	Adapted ISLH criteria	Action proposed
1	WBC	$<4.0 \times 10^3/\mu\text{L}$ or $>30.0 \times 10^3/\mu\text{L}$	BSR – confirm leukopenia or leukocytosis by blood smear estimate
2	Neutrophils <sup>#</sup>	$<1.0 \times 10^3/\mu\text{L}$ or $>20.0 \times 10^3/\mu\text{L}$	BSR and if necessary MDLC
3	Lymphocytes <sup>#</sup>	$>5.0 \times 10^3/\mu\text{L}$ (adults) and $>7.0 \times 10^3/\mu\text{L}$ (children aged $\leq 12$ years)	
4	Monocytes <sup>#</sup>	$>1.5 \times 10^3/\mu\text{L}$ (adults) and $>3.0 \times 10^3/\mu\text{L}$ (children aged $\leq 12$ years)	
5	Eosinophils <sup>#</sup>	$>2.0 \times 10^3/\mu\text{L}$	
6	Basophils <sup>#</sup>	$>0.5 \times 10^3/\mu\text{L}$	
7	Hemoglobin	$<7.0$ g/dL or $>18.5$ g/dL	BSR – check RBC morphology
8	MCV	$<75$ fL or $>105$ fL	BSR – check microcytosis and macrocytosis
9	MCHC	$<30$ g/dL	BSR – check hypochromia, verify time passed after sample collection
10	MCHC	$>36.5$ g/dL	BSR – check lipemia, hemolysis, hyperleukocytosis, RBC agglutination, and blood: anticoagulant relation and presence of spherocytes
11	RDW-CV	$>22.0\%$	BSR – check anisocytosis
12	Reticulocytes	$>100 \times 10^3/\mu\text{L}$	BSR – check polychromatophilia
13	Platelets	$<100 \times 10^3/\mu\text{L}$ or $>1000 \times 10^3/\mu\text{L}$	BSR – estimate platelet count on blood smear to confirm thrombocytopenia or thrombocytosis
14	MPV	$<5.0$ fL or $\geq 12.5$ fL	BSR – check platelet size and morphology
15	Suspect flags	<b>Immature Granulocytes?; Immature Granulocytes present</b> (Promyelocytes, myelocytes, and metamyelocytes)	BSR and if necessary MDLC
		<b>Left shift?</b>	
		<b>Atypical lymphocytes?</b>	
		<b>Abnormal Lymphocytes/blasts?; Blasts?</b> (Myeloblasts, lymphoblasts, or atypical lymphocytes)	
		<b>Nucleated RBC?</b>	BSR and if positive, count erythroblasts/100 leukocytes. If $>10\%$ , correct total leukocyte count
		<b>Fragments?</b> (Schistocytes)	BSR – check presence of schistocytes
		<b>Dimorphic RBC population</b> (Erythrocyte population heterogeneous in size)	Check RBC histogram, observe RDW value and verify anisocytosis on blood smear
		<b>RBC lyse resistance</b>	Rerun sample. If positive, BSR and check for nucleated RBC and abnormal RBC morphology
		<b>Platelet clumps</b> (generated by scattergram)? and <b>Platelet clumps</b> (generated by the impedance channel)?	Search for clots in the sample and platelet aggregates on blood smear
		<b>Platelet clumps</b> (generated by scattergram)?: verify at the bottom of the scatter plot	
	<b>Platelet clumps</b> (generated by the impedance channel)?: check platelet histogram. There is a possibility of error in counting by impedance		
	Turbidity/Hemoglobin interference?; Hemoglobin defect? (Spurious hemoglobin measurement)	Check lipemia in sample. If the sample is lipemic, remove the plasma and replace it with the same volume of analyzer diluents. Rerun only considering hemoglobin and recalculate red blood cell indices	
16	ALL	If newborn	Perform BSR and MDLC
17	(*) and (----)	These symbols beside the counts on the readout indicate that automated counts are not reliable or not available for the sample in question, respectively	

MCV: mean corpuscular volume; MCHC, mean corpuscular hemoglobin concentration; RDW-CV: red blood cell distribution width coefficient of variation; RBC: red blood cells; WBC: white blood cells; BSR: blood smear review; MDLC: manual differential leukocyte count.

<sup>#</sup> refers to absolute leukocyte counts

**Table 2 – Criteria for a positive smear recommended by the International Society of Laboratory Hematology.**

<b>Morphology</b>
<b>RBC</b>
2 <sup>+</sup> /moderate or greater. The only exception is malaria, where any finding will be considered a positive finding
<b>Platelets</b>
Giant platelets at either 2 <sup>+</sup> /moderate or greater
Platelet clumps at > rare or occasional
<b>WBC</b>
Döhle bodies at either 2 <sup>+</sup> /moderate or greater
Toxic granulation at either 2 <sup>+</sup> /moderate or greater
Vacuoles at either 2 <sup>+</sup> /moderate or greater

**Abnormal cell types**

Blasts ≥ 1%
Myelocytes/promyelocytes ≥ 1%
Metamyelocytes > 2%
Atypical lymphocytes > 5%
Nucleated RBC ≥ 1%
Plasma cells ≥ 1%

The International Society of Laboratory Hematology recommends that the use of band cell counts and left shift suspect flag (Left Shift?) should be in accordance to laboratory standard operating procedures. Thus, the Left Shift? Suspect flag was used as a screening criterion in this study and the band count was considered a positive smear finding when it was > 8%.  
RBC: red blood cells; WBC: white blood cells; SOP: standard operating procedure.

screening criterion (Table 1) and the microscopic analysis produced some PSF (Tables 2 and 3). Meanwhile, a sample was classified as false positive (FP) if it was positive for a particular screening criterion with no PSF in the microscopic analysis. A sample was classified as false negative (FN) if it was negative for all screening criteria and contained some PSF in microscopic analysis. Finally, a sample was classified as true negative (TN) if it was negative for all screening criteria and the BSR did not show any PSFs.<sup>2,8</sup>

**Considerations**

Good laboratory practices and procedures for quality assurance and quality control in hematology were followed to ensure good performance. All settings and configurations of the hematology analyzers followed the manufacturers' recommendations. For all samples, the BSR and MDLC were performed according to an adapted version of the H20-A2 Clinical and Laboratory Standards Institute (CLSI),<sup>9</sup> because each BSR and each MDLC was carried out by a single observer. In most cases, 100 leukocytes were counted. In some cases with leukocyte counts  $< 0.03 \times 10^3/\mu\text{L}$ , leukocytes were not observed and only a BSR could be performed. In other cases with leukocyte counts above  $50.0 \times 10^3/\mu\text{L}$ , the MDLC reached 200 leukocytes. Furthermore, in all samples, the BSR was performed to identify any qualitative or quantitative changes in red and white blood cells as well as platelets. Guidelines were formulated to standardize the quantification of morphologic alterations and which terms are used to report changes in CBC results. These guidelines were codified as standard operating procedures (SOP).

**Table 3 – Criteria for a positive smear recommended by the Hospital de Clínicas da Universidade Federal do Paraná.**

Red blood cells		
Anisocytosis ≥ 2 <sup>+</sup>	Microcytes ≥ 2 <sup>+</sup>	Macrocytes ≥ 2 <sup>+</sup>
Poikilocytosis ≥ 2 <sup>+</sup>	Elliptocytes ≥ 3 <sup>+</sup>	Stomatocytes ≥ 3 <sup>+</sup>
Codocytes ≥ 2 <sup>+</sup>	Dacryocytes ≥ 2 <sup>+</sup>	Schistocytes ≥ 1 <sup>+</sup>
Acanthocytes ≥ 2 <sup>+</sup>	Drepanocytes: present	Spherocytes ≥ 1 <sup>+</sup>
Howell-Jolly: present	Cabot ring: present	Basophilic stippling ≥ 1 <sup>+</sup>
Rouleaux ≥ 1 <sup>+</sup>	Hypochromia ≥ 2 <sup>+</sup>	Polychromatophilia ≥ 2 <sup>+</sup>
RBC agglutination: present	Hemoglobin C crystals: present	Hematozoa: present
White blood cells		
Döhle bodies ≥ 1 <sup>+</sup>	Toxic granulation ≥ 1 <sup>+</sup>	Cytoplasmic vacuoles ≥ 1 <sup>+</sup>
Polylobocytes ≥ 1 <sup>+</sup>	Hyposegmented neutrophils: ≥ 2 <sup>+</sup>	Neutrophil hypo/degranulation: present
Auer rod: present	Pseudo-Pelger-Huët: present	Dysplastic cells: present
Platelets		
Giant platelets ≥ 1 <sup>+</sup>	Microplatelets ≥ 2 <sup>+</sup>	Platelet aggregates: present
Platelet anisocytosis ≥ 3 <sup>+</sup>	Degranulated platelets: present	Gray platelets: present
Abnormal cell types		
Blasts ≥ 1%	Promyelocytes ≥ 1%	Myelocytes ≥ 1%
Metamyelocytes ≥ 2%	Bands > 8%	NRBC ≥ 1/100 leukocytes
Plasma cells ≥ 1%	Prolymphocytes ≥ 1%	Atypical lymphocytes ≥ 5%

Source: adapted from Barnes et al.,<sup>2</sup> Gulati et al.,<sup>3</sup> and Hyun et al.<sup>10</sup>

RBC: red blood cells; NRBC: nucleated red blood cells.

**Statistical analysis**

The sensitivity, specificity, positive predictive value, negative predictive value, microscopic review rate, and efficiency of the RC adapted from the ISLH were calculated as follows: sensitivity (%) =  $TP/(TP + FN) \times 100$ ; specificity (%) =  $TN/(TN + FP) \times 100$ ; positive predictive value (%) =  $TP/(TP + FP) \times 100$ ; negative predictive value (%) =  $TN/(TN + FN) \times 100$ ; microscopic review rate (%) =  $(TP + FP)/(TP + FP + FN + TN) \times 100$ ; and efficiency (%) =  $(TP + TN)/(TP + FP + FN + TN) \times 100$ .<sup>8</sup>

**Results**

Out of the 1977 samples used in the initial investigation of the adapted RC, 583 (29.49%) were positive and 1394 (70.51%) were negative for relevant microscopic results, i.e., for PSFs. Most samples failed to meet some screening criteria; usually

**Table 4 – Truth table of the adapted International Society of Laboratory Hematology criteria.**

Parameters (%)	XE-2100D (n = 1573)	XT-2000i (n = 404)	Total (n = 1977)
True positive	21.93	25.99	22.76
False positive	21.93	28.46	23.27
True negative	48.77	41.34	47.24
False negative	7.37	4.21	6.73
Specificity	68.97	59.22	67.00
Sensitivity	74.84	86.06	77.19
Positive predictive value	50.00	47.73	49.45
Negative predictive value	86.86	90.76	87.54
Efficiency	70.69	67.32	70.00
Microscopic review rate	43.86	54.45	46.03

Note: the adapted ISLH screening criteria were tested considering the ISLH criteria for a positive smear as shown in Table 2.

**Table 5 – Incidence of false negatives: the adapted International Society of Laboratory Hematology criteria considering the International Society of Laboratory Hematology criteria for a positive smear.**

False negatives	n	%
Left shift >8% bands	114	5.76
RBC morphology $\geq 2^+$	13	0.65
Myelocytes/promyelocytic $\geq 1\%$	13	0.65
Giant platelets $\geq 2^+$	1	0.05
Total	141 <sup>a</sup>	6.73% = 133 samples of 1977

RBC: Red blood cell.

<sup>a</sup> Some samples showed more than one positive smear finding.

from one to four criteria but up to 15. The results obtained for the validation of the screening criteria tailored from the ISLH criteria, considering the positive smear criteria in Table 2, are shown in Table 4.

The microscopic review rate during the initial investigation (Table 4) was quite high (46.03%) compared to the 30% recommended by the American College of Pathologists.<sup>11</sup> The main rules producing FP results in the initial assessment were as follows: leukocytes  $<4.0 \times 10^3/\mu\text{L}$  (16.7% of total FPs), platelets  $<100 \times 10^3/\mu\text{L}$  (13.3% of total FPs), and suspect flags (30% of total FPs). In the initial assessment of the RC adapted from the ISLH criteria, there were 133 FN samples (Table 5) including 114 with nuclear shift to the left in >8% of bands, 13 myelocytes/promyelocytes  $\geq 1\%$ , 13 alterations in red blood cells, and one alteration in platelets.

The evaluation results for the screening criteria adapted from the ISLH in conjunction with the PSFs of the HC-UFPR are shown in Tables 6 and 7. From the 180 samples analyzed, 76 contained PSFs. Using PSFs specific for the HC-UFPR, which were tailored to meet the requirements of the authors' Institution, caused the screening criteria adapted from the ISLH to be incapable of filtering samples with relevant morphological alterations (FN = 15.5%), including one case with blasts present from a patient with acute leukemia.

**Table 6 – Truth table of the adapted International Society of Laboratory Hematology criteria considering the Hospital de Clínicas da Universidade Federal do Paraná criteria for a positive smear.**

Parameters (%)	XT-2000i (n = 180)
True positive	26.7
False positive	10.5
True negative	47.3
False negative	15.5
Specificity	81.7
Sensitivity	63.2
Positive predictive value	71.6
Negative predictive value	75.2
Efficiency	73.9
Microscopic review rate	37.2

**Table 7 – Incidence of false negatives: the adapted International Society of Laboratory Hematology screening criteria considering the Hospital de Clínicas da Universidade Federal do Paraná criteria for a positive smear.**

False negatives	n	%
Left shift >8% bands	8	4.44
Atypical lymphocytes $\geq 5\%$	1	0.55
RBC morphology	10	5.55
Toxic granulation $\geq 1^+$	6	3.33
Cytoplasmic vacuoles $\geq 1^+$	1	0.55
Giant platelets $\geq 1^+$	6	3.33
Platelet aggregates: present	6	3.33
Blasts $\geq 1\%$	1	0.55
Total	39 <sup>a</sup>	15.5% = 28 samples in 180

RBC: Red blood cell.

<sup>a</sup> Some samples showed more than one positive smear finding.

In both analyses, the percentage of FN results exceeded the reliability threshold of 5% established by the ISLH, causing cases with serious illnesses to be overlooked.

## Discussion

Despite the extensive capabilities of the latest-generation multiparametric hematology analyzers, microscopic review of blood smears still plays an important role in hematology laboratories. The use of RC, which would allow the release of automated counts without a BSR, is neither a widespread nor standardized procedure. In practice, many laboratories simply adopt published criteria or alter some criteria without empirical evidence.<sup>12</sup>

Establishing and assessing a specific RC for a particular laboratory means that the ISLH screening and positive smear criteria must first be validated, especially because these criteria are international standards. Considering the actual demands and capabilities of each laboratory, this initial validation clarifies whether such criteria require optimization or not.

In this study, after the initial evaluation of the RC adapted from the ISLH, 6.73% and 23.27% of FN and FP results were

verified, respectively. Moreover, the microscopic review rate was 46.03% and efficiency was 70.0%. The microscopic review rate with the XT-2000i analyzer (54.45%) was higher than that with the XE-2100D (43.86%). This is because the former analyzer was used by staff members on duty, who only attend inpatients and emergency patients, who evidently have CBC results that differ substantially from those of outpatients.

The negative predictive value determined in the initial evaluation using the screening criteria and positive smear criteria showed that in 87.54% of the times in which the screening criteria adapted from ISLH did not indicate the need for BSR, the sample analyzed really did not contain any PSFs. The observed sensitivity was 77.19%, indicating that out of 583 samples, 450 with PSFs were correctly screened by applying the RC adapted from the ISLH. As mentioned above, the percentage of microscopic reviews (46.03%) greatly exceeded the 30% recommended by the American College of Pathologists. The fact that 30% of the FP results were of samples with suspect flags with a nonexistent microscopic counterpart indicated that the hematology analyzers used were guilty of over-flagging, i.e., they gave more warnings than necessary. As the sensitivity of the suspect flags had been adjusted by technicians of the manufacturer, we suggest that each laboratory should evaluate the efficiency of each suspect flag from the analyzers, thereby making proper adjustments to the sensitivity of the hematology analyzer or defining whether a suspect flag is actually useful as a screening criterion.

A total of 13.3% of samples had a platelet count  $<100 \times 10^3/\mu\text{L}$  in the initial evaluation of the RC adapted from the ISLH; this is relatively high considering the sampling was representative of the local reality. Therefore, the microscope estimate of the platelet count should always be performed on samples with this profile in order to verify how well it complies with automated counting and to search for platelet aggregates and giant platelets, which are factors that produce underestimates.<sup>13</sup>

Moreover, the screening criteria adapted from the ISLH were tested according to the PSFs of the HC-UFPR; 15.5% and 10.5% of FNs and FPs were verified, respectively. In addition, the microscopic review rate was 37.3%, and the efficiency was 73.8%. One FN sample contained blasts. It is unacceptable to fail to detect cases of undiagnosed hematological malignancies; therefore, each institution should evaluate the need to perform BSRs in all patients in the hematology unit even at the expense of an increased microscopic review rate.

Despite some studies suggesting poor clinical practice when counting bands,<sup>14-16</sup> in this study, a band count  $>8\%$  was chosen as the positive smear criterion, particularly because a substantial proportion of doctors in Brazil posit there are associations of a nuclear shift to the left of neutrophils with infectious and inflammatory conditions. The FN analysis in the present study revealed a high occurrence of a nuclear shift to the left  $>8\%$  bands. Nevertheless, the technical limitations of MDLC must be taken into account when interpreting these results, especially results regarding band counts and variations occurring as a result of age, gender, and conditions such as pregnancy and physical exercise.<sup>15,17</sup>

The greatest amendment made to the RC of the ISLH was regarding the Delta check rules. These rules recognize morphological abnormalities detected and validated in previous

analyses as well as discrepancies between the results of the current analysis and previous results from up to five days earlier. The Delta check limit threshold for a particular parameter is the amount by which a result can differ from a previous one; this difference can be expressed as the percentage difference or the absolute value in the unit of the hematological parameter in question. Delta limits should be established by each laboratory taking into account the physiopathological aspects and technical characteristics of the hematology analyzers used.

Although the Delta check rules play important roles in the efficiency and reliability of the CBC results directly released without a BSR, many clinical laboratories are incapable of implementing them in their electronic records or interfacing systems because of high software development costs. Even in developed countries, some laboratories are still incapable of modifying their electronic record systems to incorporate such regulations.<sup>18</sup>

Another equally important factor is how unfamiliar laboratory professionals are with these rules, which make their dissemination and implementation difficult. The ISLH does not suggest specific Delta check limits, leaving them to the discretion of the laboratory. This could lead to small differences in the efficiency of the ISLH criteria when applied by different laboratories using different Delta limits since the International Consensus Group for Hematology Reviews only suggests specific actions for situations in which the Delta limits individually established by the laboratories are exceeded.

---

## Conclusions

Neither assessment carried out in this study proved to be reliable, because the FN rates exceeded 5%. Moreover, the microscopic review rates were high, which may pose problems when attempting to decrease the turnaround time of results and when laboratories have an insufficient number of experienced professionals to perform the BSR. Therefore, in conclusion, new criteria should be developed and evaluated taking into account local peculiarities, requirements, and opportunities while aiming not to overlook samples with PSFs. To this end, additional studies are being carried out by the authors with the aim of developing a method to help establish and evaluate the efficiency of criteria for reviewing automated CBC results.

---

## Conflict of interest

The authors declare no conflicts of interest.

---

## REFERENCES

1. International Society for Laboratory Hematology [Internet]. Available from: [www.islh.org](http://www.islh.org) [accessed on 22.01.14].
2. Barnes PW, McFadden SL, Machin SJ, Simson E. The international consensus group for hematology review: suggested criteria for action following automated CBC and WBC differential analysis. *Lab Hematol.* 2005;11:83-90.

3. Gulati GL, Alomari M, Kocher W, Schwarting R. Criteria for blood smear review. *Lab Med.* 2002;33:374-7.
4. Pierre RV. Peripheral blood film review: the demise of the eyecount leukocyte differential. *Clin Lab Med.* 2002;22:279-97.
5. Bain BJ. Diagnosis from the blood smear. *N Engl J Med.* 2005;353:498-507.
6. Wang H, Sun F, Yu G, Wang M, Liu N, Jin Y. Applications of international slide review criteria for actions among Chinese populations. *Int J Lab Hematol.* 2008;30 Suppl. 1:65-6.
7. Peterson P, McNeill S, Gulati G. Cellular morphologic analysis of peripheral blood. In: Kottke-Marchant K, Davis BH, editors. *Laboratory Hematology Practice.* Singapore: Wiley-Blackwell; 2012. p. 10-25.
8. Ruzicka K, Veitl M, Thalhammer-Scherrer R, Schwarzwinger I. The new hematology analyzer Sysmex XE-2100: performance evaluation of a novel white blood cell differential technology. *Arch Pathol Lab Med.* 2001;125:391-6.
9. Clinical and Laboratory Standards Institute (CLSI). Reference Leukocyte (WBC). In: *Differential count (proportional) and evaluation of instrument methods; approved standard.* 2nd ed. Wayne, PA: CLSI Document H20-A2; 2007.
10. Hyun BH, Gulati GL, Ashton JK. Differential leukocyte count: manual or automated, what should it be? *Yonsei Med J.* 1991;32:283-91.
11. Novis DA, Walsh M, Wilkinson D, Louis MS, Ben-Ezra J. Laboratory productivity and the rate of manual peripheral blood smear review: a College of American Pathologists Q-Probes study of 95,141 complete blood count determinations performed in 263 institutions. *Arch Pathol Lab Med.* 2006;130:596-601.
12. Cui W, Wu W, Wang X, Wang G, Hao Y, Chen Y, et al. Development of the personalized criteria for microscopic review following four different series of hematology analyzer in a Chinese large scale hospital. *Chin Med J (Engl).* 2010;123:3231-7.
13. Comar SR, Danchura HSM, Silva PH. Contagem de plaquetas: avaliação de metodologias manuais e aplicação na rotina laboratorial. *Rev Bras Hematol Hemoter.* 2009;31: 431-6.
14. Ward MJ, Fertel BS, Bonomo JB, Smith CL, Hart KW, Lindsell CJ, et al. The degree of bandemia in septic ED patients does not predict inpatient mortality. *Am J Emerg Med.* 2012;30: 181-3.
15. Cornbleet PJ. Clinical utility of the band count. *Clin Lab Med.* 2002;22:101-36.
16. Van der Meer W, Van Gelder W, De Keijzer R, Willems H. Does the band cell survive the 21st century? *Eur J Haematol.* 2006;76:251-4.
17. Rüemke CL. Laboratory aids. Variability of results in differential cell counts on blood smears. *Triangle.* 1960;4:154-8.
18. Galloway MJ, Charlton A, Holland D, Trigg G, Gibson R. An audit of the implementation of the international consensus group's guidelines on reporting of blood films. *J Clin Pathol.* 2010;63:351-4.