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A study investigating variability of left ventricular ejection fraction using manual and automatic processing modes in a single setting

P. Bresser ^{a, *}, J. de Beer ^{a, b}, Y. de Wet ^{a, c}

^a Department of Radiography, Faculty of Health Sciences, School of Healthcare Sciences, University of Pretoria, South Africa

^b Dr G Lange Inc, Little Company of Mary Hospital, Pretoria, South Africa

^c Department of Nuclear Medicine, Steve Biko Academic Hospital, Pretoria, South Africa

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ABSTRACT

Purpose: A planar multi-gated cardiac blood pool acquisition is a non-invasive technique commonly used to measure left ventricular ejection fraction (LVEF). It is essential that the calculation of LVEF be accurate, repeatable and reproducible for serial monitoring of patients. Different processing modes may be used in calculating the LVEF which require various degrees of manipulation. In addition, different operators with varying levels of experience may process the same data set. It is not known whether the inter-operator variability of LVEF values within a single nuclear medicine department has the potential to affect the calculated LVEF and in turn affect patient management. The aim of the study was to determine variability of LVEF values among operators with different levels of experience using two processing modes.

Methods: A descriptive cross-sectional study was carried out in a single setting. Four operators with varying levels of experience analysed 120 left anterior oblique projections using manual and automatic processing modes to calculate the LVEF. Inter- and intra-operator correlation was determined.

Results: Manual processing showed moderate to strong agreement (r1 = 0.653) between operators. Automatic processing indicated almost perfect (r1 = 0.812) inter-operator correlation. Intra-operator correlation demonstrated a trend of decreasing variability between processing modes with increasing levels of experience.

Conclusion: Despite the overall inter-operator agreement, significant intra-operator variability was evident in results from operators with less experience. However, the discrepancies were such that the differences in LVEF would not play a role in patient management. It is recommended that automatic processing be used for determining LVEF to limit inter-operator variability. Additionally operator experience should be considered in the absence of standardised processing protocols when different processing modes are available in a single setting.

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Introduction

A planar multi-gated cardiac blood pool acquisition (MUGA) is considered as an accurate, safe, non-invasive method commonly used to evaluate left ventricular ejection fraction (LVEF).^{1–3} Any significant change in LVEF (decrease of 10%) from baseline is an early indicator of cardiac failure and may precede any symptoms of cardiac disease.^{3–6} Early detection of changes allows for intensified monitoring to prevent further complications, initiation of preventative measures or may implicate changes in cancer treatments or

E-mail address: pippa.bresser@up.ac.za (P. Bresser).

patient management.^{7.8} After gated image acquisition, regions of interest (ROIs) are drawn over the left ventricle at end systole and end diastole using processing software and algorithms are applied to calculate the LVEF.^{9,10} Previously, only manual processing modes were used to determine LVEF.^{1,11} Technological improvements led to the introduction of automatic and semi-automatic processing modes and the evolution of various processing software. However, these developments also increased the probability of variation in determining the LVEF. Sources of variability in manual and automatic processing modes include the following:

- (i) Differences between software packages and applied algorithms to calculate LVEF;^{1,2,4,12}
- (ii) Human detection inaccuracies of the true end systole and end diastole images;^{1,13}

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^{*} Corresponding author. Department of Radiography, University of Pretoria, Steve Biko Academic Hospital, Level F5, Gezina, Pretoria 0002, South Africa. Tel.: +27 (0) 12 354 4348.

- (iii) Differences in software edge-detection algorithms;¹
- (iv) Arrhythmias (e.g. ectopic beats or atrial fibrillation) which lead to artificially reduced counts in frames later in the cardiac cycle (end-diastole);¹
- (v) Inclusion of other anatomic structures within the left ventricle (LV) ROI due to poor positioning or anatomic variations;^{1,8}
- (vi) Poor labelling of the red blood cells leading to reduced count rates;¹
- (vii) Increased background counts leading to improper detection of the LV edge and reduced accuracy of the LVEF;¹³
- (viii) Different operators with various levels of experience processing the same data set which formed the basis of the current study.^{1–3,14}

The extent of variability in LVEF results in a particular setting should be considered where multiple operators and processing modes are available.^{13,15} The calculation of LVEF must be accurate, repeatable and reproducible, as serial monitoring of LVEF variations has the potential to affect patient treatment and management.^{4,11} 'Repeatability' of LVEF is an expression of the variability of repeated measurements of the same acquisition as opposed to 'reproducibility' which includes both repeated acquisition and measurement preferably at different times and places.¹⁴ Inter- and intra-operator variability in MUGA processing and data manipulation is a principal concern in nuclear medicine departments where the possibility of errors or discrepancies should be kept to a minimum.^{2–4} Hains et al. compared the LVEF values from three operators using three different processing modes (manual, semiautomated and regional) and established that there was no significant inter-operator variability.² This was supported by Bailey and Bailey in a study across multiple settings.³ However, these studies did not consider the different levels of experience of the operators in determining inter-operator variability. The intraoperator variability determined by Bailey and Bailey did include the different levels of experience of the operators but the results were based on the use of a single processing mode. It is thus contended that inter- and intra-operator differences in LVEF results may occur where operators in the nuclear medicine department have varying levels of clinical experience in processing MUGA examinations using different processing modes. The purpose of this study was to investigate the extent of inter- and intra-operator variability in calculating LVEF through automatic and manual processing modes with multiple operators within a single setting.

Methods

A descriptive cross-sectional study was carried out in a nuclear medicine department in Pretoria, South Africa where retrospective analysis of MUGA examinations was undertaken. Ethical approval to conduct this study was granted by the Ethics Committee of the Faculty of Health Sciences, at the University of Pretoria. The data collected from the picture-archiving system of the hospital consisted of the LAO projection of all patients that underwent a MUGA examination from October to April 2013 until a consecutive sample of 120 data sets was obtained. The images had been acquired using the General Electric (GE), Millennium, single head gamma camera (GE Healthcare) fitted with a low-energy, high-resolution collimator. All studies were acquired with a 64 × 64 matrix in the LAO projection using 24 frames.

Four operators with varying levels of experience and clinical expertise in the processing of nuclear medicine examinations volunteered to process the data. The 4 operators included one nuclear medicine registrar (doctor) with four years' experience in reporting nuclear medicine examinations, one nuclear medicine radiographer with 12 years' experience post-graduation (senior), one junior nuclear medicine radiographer with 3 years' experience post-graduation and one nuclear medicine student in the first year of training. The 4 operators were blinded to the patient information and were blinded to the results of the other operators. The data was processed using Xeleris 2.0 software (GE Healthcare). Each operator processed 120 data sets both manually and automatically to determine the LVEF. Manual processing required the operator to visually identify and manually draw the region of interest around the edge of the left ventricle. Automatic processing required the operator to select the centre of the left ventricle and the processing software identified the edge of the ventricle through the use of an edge-detection algorithm. Manual and automatic processing was performed at different time points in order to minimise the risk of operator bias in trying to replicate results using different processing modes. The results of the manual and automatic LVEF values were recorded on separate data collection sheets. Upon completion of processing with one mode, the data collection sheet was placed in a sealed box where after processing using the other mode was performed. The authors were not involved in the processing or collecting of data. Procedural bias was avoided in that operators could select the two time points that was most appropriate for them to complete the processing depending on their work load and personal preference. Operators were volunteers and were not given any incentives to participate.

Inter-operator variability was considered as the amount of variation between LVEF results obtained by the different operators processing the same data set. Intra-operator variation was considered as the variation of LVEF value calculated by one operator processing the same data set using two different processing modes.

Inter-operator correlation coefficient (r1) was calculated for manual and automatic processing modes to determine the agreement of LVEF values among the 4 operators. The variability between manual and automatic LVEF among the operators was described using summary statistics. Additionally, intra-operator correlation was determined using the Wilcoxon signed rank test to determine any discrepancies when the same data was processed multiple times by an individual, using different processing modes.

Results

Table 1

Inter-operator intra-class correlation coefficient (*r*1) was calculated with a 95% confidence interval (CI) as seen in Table 1. Manual processing showed a significant moderate to strong agreement between operators. Automatic processing indicated an almost perfect and significant inter-operator correlation.

LVEF values ranged from 20% to 84%. Table 2 illustrates the variability of LVEF values among the operators. Intra-operator correlation among the four operators was determined using the Wilcoxon signed rank test. The LVEF values from the doctor and senior radiographer did not show a significant difference (p > 0.05) between the manual and automatic results. A significant intra-operator difference existed, however, between the manual and automatic LVEF values obtained by the student and junior radiographer. There was a 6.97% deviation of LVEF between the manual

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Inter-operator	correlation	of LVEF to	or manual an	id automatic	processing.

	ICC	95% CI ranges		p-Value
Manual	r1 = 0.65	0.49	0.77	0.00*
Automatic	r1 = 0.81	0.76	0.86	0.00*

 $p^* \leq 0.05$ considered significant.

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