Semi-quantitative segmental perfusion scoring in myocardial perfusion SPECT: visual vs. automated analysis

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ABSTRACT

Introduction: It is recommended that the physician apply at least a semi-quantitative segmental scoring system in myocardial perfusion SPECT. We aimed to assess the agreement between automated semi-quantitative analysis using QPS (quantitative Perfusion SPECT) software and visual approach for calculation of summed stress score (SSS), summed rest score (SRS) and summed difference score (SDS).

Methods: We retrospectively studied 1782 consecutive patients who had undergone two-day stress-rest Tc99m-MIBI myocardial perfusion SPECT. Based on 17-segment 5-scale scoring system, SSS, SRS and SDS were calculated visually and using QPS software.

Results: There was good correlation between visual analysis and QPS software in calculation of SSS and SRS and a fair correlation for SDS. However, there was statistically significant difference between two methods. By Bland-Altman analyses mean value of the differences (estimated bias) differs significantly from 0 on the basis of 1-sample t-test. Based on bias, Precision and 95% limits of agreement, discrepancies between measurements indicate no agreement equally through the range of measurements, so there is a proportional bias. Based on SSS, SRS and SDS \leq 3 and SSS, SRS and SDS >3, there was fair concordance between the visual assessment and automated QPS calculation. Kappa statistics was 0.41 (P<0.001), 0.69 (P<0.001) and 0.25 (P<0.001) for SSS, SRS and SDS respectively.

Conclusion: Although semi-quantification sores by visual and automated analysis is correlated, the agreement assessed by Bland-Altman analysis is not high especially in more extensive perfusion defects. Semi-quantitative automated analysis should be used as a supplement to the visual assessment.

Key words: Myocardial perfusion SPECT; Semi-quantification; Visual analysis; Automated analysis; Scoring system

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INTRODUCTION

Perfusion assessment of myocardial perfusion SPECT (MPS) can be performed visual or by automated software [1]. Visual interpretation is subjective and is prone to intra-observer variability and to have lower repeatability than objective automated quantitative measurements [2, 3]. In addition to the qualitative evaluation of perfusion defects, it is recommended that the physician also apply at least a semi-quantitative segmental scoring system [4]. For this purpose, a semi-quantitative scoring system using a left ventricular-segmented model (17 or 20 segments) and scored with a five-point scale was introduced [3]. In order to facilitate consistency of nomenclature with other imaging modalities, the 17-segment model is preferred [4, 5].

Left ventricular myocardial walls in tomogram perfusion images are divided into a number of segments, e.g. 17 (Figure 1). Each segment is scored separately using a 5-point scoring system ranging from 0 (normal uptake) to 4 (uptake absent) [5-7]. The summed stress scores (SSS) equals the sum of the stress scores of all the segments and the summed rest score (SRS) equals the sum of the resting scores of all the segments. Difference between the SSS and SRS is summed difference score (SDS) and it is a measure of inducible ischemia [4]. This semiquantitative scoring system provides a standard and reproducible assessment of perfusion defect severity and extent. SSS, SRS and SDS can provide valuable diagnostic and prognostic parameters [3]. In addition to visual calculation of the SSS, SRS and SDS, computer software packages were developed for automatic generation of semi-quantitative segmental scores [3, 8-9]. Agreement for the visual or automated semi quantitative systems is essential and understanding of the differences is necessary. QPS (Quantitative Perfusion SPECT) is one of the widely used and commercially available software packages.

In this study, we aimed to assess the agreement and concordance between automated semi-quantitative analysis using QPS software and visual approach for calculation of SSS, SRS and SDS.

METHODS

Study population

We retrospectively studied 1782 consecutive patients (732 male, 1050 female) with mean age 59.71±12.12 (25-94) who had underwent MPS in our nuclear medicine department. They were referred for clinically stress/rest Tc-99m Sestamibi MPS.

Myocardial perfusion SPECT

All patients underwent two-day stress-rest Tc99m-MIBI (methoxyisobutyl-isonitrile) myocardial perfusion SPECT (MPS). In a two-day protocol, after intravenous injection of 740-925 MBq Tc99m-MIBI SPECT acquisition was performed with a Dual-head SPECT system with the detectors oriented at 90 Variable-Angle degrees ((Dual-Head E.CAM; Siemens) equipped with a low-energy-high resolution collimator. A 20% window with 140 keV energy peak was used. A total of 32 projections (stepand-shoot mode, 25 s per view) were obtained over a 180° arc commencing from the right anterior oblique to left posterior oblique view. We used a zoom factor of 1.45. The images were stored in a 64×64 matrix in the computer and reconstructed by filtered backprojection using a Butterworth filter (cut-off value was 0.55 cycle/cm for ungated data, order =5). No attenuation or scatter correction was applied. All reconstructed tomographic images were interpreted by consensus of 2 experienced physicians. Based on 17-segment model and 5-point scale system (0, normal perfusion; 1, mildly reduced uptake; 2, moderately reduced uptake; 3, severely reduced uptake; and 4, absent uptake) visual semi-quantitative scores: SSS, SRS and SDS were calculated (Figure 1) [6].



Fig 1. 17-segment 5-point scale for semi-quantitative assessment of myocardial perfusion.

Readers were blinded regarding clinical information, final diagnosis and results of automatic quantifications. Using QPS software, 17-segment automatic quantification of SSS, SRS and SDS was performed.

Statistical analysis

All continuous variables are reported as mean \pm SD. Univariate analyses of continuous variables were performed using the paired sample t test for the comparison of two studies and categorical variables were compared using the chi-squared test. Pearson correlation coefficients were calculated. Diagnostic concordance between the visual and quantitative methods was assessed with the kappa statistic. The Bland-Altman method was applied to determine the bias and agreement between quantitative and visual methods. P<0.05 was considered to represent a statistically significant difference.

Variable	Visual	QPS	t–test P value	Correlation: r (p value)
SSS	3.36±5.94	5.33±7.87	p<0.001	0.797 (p<0.001)
SRS	2.25±5.04	2.80±6.79	p<0.001	0.838 (p<0.001)
SDS	1.34±2.59	2.56±3.06	p<0.001	0.424 (p<0.001)

Table 1: Calculated summed stress score (SSS), summed rest score (SRS) and summed difference score (SDS) visually and using QPS (Quantitative Perfusion SPECT) software.

Table 2: Categorized summed stress score (SSS) calculated visually and using QPS (Quantitative Perfusion SPECT) software (P<0.001).

		QPS Total		QPS	
		SSS≤3	SSS>3		
	SSS≤3	944(68.9%)	427(31.1%)	1371 (100%)	
Visual	SSS>3	53(13.9%)	328(86.1%)	381 (100%)	
Т	otal	997	755	1752	

Table 3: Categorized summed rest score (SRS) calculated visually and using QPS (Quantitative Perfusion SPECT) software (P<0.001).

		QPS Total		Total
		SRS≤3	SRS>3	
	SRS≤3	1302(92.6%)	104(7.4%)	1406 (100%)
Visual	SRS>3	33(14.4%)	196(85.6%)	229 (100%)
Т	otal	1335	300	1635

Table 4: Categorized summed difference score (SDS) calculated visually and using QPS (Quantitative Perfusion SPECT) software (P<0.001).

		QPS		Total	
		SDS≤3	SDS>3		
	SDS≤3	1119 (78.0%)	315 (22.0%)	1434 (100%)	
Visual	SDS>3	80 (40%)	120 (60%)	200 (100%)	
Т	otal	1999	435	1634	

RESULTS

According to the perfusion patterns stress/rest tomograms, 1203(66.1%) had normal myocardial perfusion SPECT, 310(17.4%) patients had reversible defects, 72(4.0%) had fixed defects and 197(11.1%) patients had partially reversible perfusion defects. Table 1 showed SSS, SRS and SDS calculated visually as well as using QPS software in all patients. There was good correlation between visual analysis and QPS software in calculation of SSS and SRS and a fair correlation in calculation of SDS. However, there was statistically significant difference between two methods.

The differences of visual and automated QPS assessments by Bland-Altman analyses are shown in Figure 2. Mean value of the differences (estimated bias) differs significantly from 0 on the basis of 1sample t-test. Bias (mean difference), Precision (SD) and 95% limits of agreement (mean ±1.96SD) for SSS were -1.97, 4.77, and -11.31 to 7.37 respectively. Bias (mean difference), Precision (SD) and 95% limits of agreement (mean ± 1.96 SD) for SRS were -.54, 3.76, and -7.91 to 6.82 respectively. Bias (mean difference), Precision (SD) and 95% limits of agreement (mean ± 1.96 SD) for SDS were -1.22, 3.06 and -7.21 to 4.78 respectively. Discrepancies between measurements indicate no equal agreement through the range of measurements, so there is a proportional bias.

Based on SSS, SRS and SDS ≤ 3 and SSS, SRS and SDS >3, there was fair concordance between the visual assessment and automated QPS calculation (Table 2, 3 and 4). Kappa statistics was 0.41 (P<0.001), 0.69 (P<0.001) and 0.25 (P<0.001) for SSS, SRS and SDS respectively. These values suggest moderate to good agreement for SRS but weak agreement for SSS and SDS specially SDS.

DISCUSSION

Semi-quantitative analysis of myocardial perfusion adds useful information, including prognostic information, and improves reproducibility as well as reliability of interpretations [10]. A variety of automated semi-quantitative scores and polar maps have been developed. Abnormalities are defined by comparison with a gender-matched normal database [3, 11].

In scoring systems such as semi-quantitative 17segment system, it is possible to determine perfusion scores (SSS, SRS and SDS) based on visual assessment or using automated software programs. A number of programs for semi-quantitative analysis have been validated [8]. QPS is one of the most validated and common software used in this setting.



Fig 2. Bland-Altman analysis of calculated perfusion scores visually and using QPS (Quantitative Perfusion SPECT) software. A: Summed stress score (SSS), B: Summed rest score (SRS) and C: Summed difference score (SDS) visually and using QPS (Quantitative Perfusion SPECT) software.

We observed not high but weak to modest level of diagnostic agreement between the visual assessment of myocardial perfusion with 99mTc sestamibi SPECT and the computer derived automated SSS, SRS and SDS (weighted Kappa: 0.41, 0.69 and 0.25

respectively). Based on κ statistics, bias and precision in Bland-Altman analysis, SRS calculation had better values of agreement between visual and automated assessment.

Based on Bland-Altman analysis, we observed two remarkable points: first the agreement between two methods is not high and second the relationship is proportional: in low scores, agreement is higher than high scores indeed when there are multiple perfusion defects, agreement is lower.

Our findings are less compatible with the report from the group that developed the commercial software [9]. Leslie et al. [12] reported good concordance between the original visual interpretation and the automated SSS. In their study, they considered the automated derived $SSS \le 3$ as normal comparing with the visual findings.

Of the 388 patients having visually normal scans, 305 (79%) had SSS \leq 3 and of the 330 patients were visually abnormal, 268 (81%) with SSS>3. The kappa statistic was 0.60 (P<0.0001) [12].

Hsu et al. reported that diagnostic performance of automated semi-quantitative analysis with 4D-MSPECT was comparable with the visual approach [3]. However, they reported moderate agreement for stress and rest images (weighted κ =0.55, weighted κ =0.50 respectively) between visual and automated 4D-MSPECT segmental scores (based on scores 0, 1, 2, 3, 4).

In practice, nuclear medicine physicians integrate the patients' clinical data and review image quality and attenuation artifacts and put all findings into consideration for calculation of perfusion scores [3]. Semi-quantitative automated analysis should be used as a supplement to the visual assessment and should not be reported in isolation from the review of the images [10]. Despite the advantages of quantitative analysis, visual analysis is still an integral part of myocardial perfusion SPECT interpretation, since the quantitative analyses have not yet been refined to recognize a variety of artifactual patterns [11]. It is recommended that semi-quantitative visual analysis and automated analysis be assessed simultaneously [11]. Physicians should be aware of the variability that exists between visual analysis and software packages as well as different interpreting physicians and software factors [3]. Concordant quantitative findings with the clinical impression can result more confidence in the interpretation, however, in presence of discordant findings, more careful review of the scan is necessary [12].

The result of our study including discrepancy with previous reports and lack of expected agreement between visual and automated calculation of SSS, SRS and SDS raises questions which need explanations:

- In presence of multiple defects and high scores, variability between measurements would be high.
- Our incorporated normal data base in the computer software program may be not a good reference population.
- Software cannot recognize some of the artifactual patterns easily recognized by visual inspection.
- On the other hand, in automated programs the patient's data was compared with normal data base. So for calculation of perfusion scores in different segments (for example in the inferior and anteroapical segments), the software may consider zero score based on diaphragmatic or breast attenuations which are existing in normal data pool. On the other hand, in visual scoring we may consider no zero scores for these segments.

In our retrospective study, we didn't have the data related to segment-by-segment scores as a limitation of study. So we recommend performing a prospective study with a large population to compare the visual and automated analysis based on a segment-bysegment protocol to be able to define the regions with most variability.

CONCLUSION

The present study demonstrates that although semiquantification sores by visual and automated analysis more or less are in agreement but this correlation as assessed by Bland-Altman analysis is not high especially in more extensive perfusion defects. Semiquantitative automated analysis should be used as a supplement to the visual assessment and should not be reported in isolation from the review of the images.

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