Tissue Doppler Imaging in Cardiology Nowadays: Clinical Applications

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ABSTRACT

Echocardiography is a routine test in cardiology nowadays and Tissue Doppler imaging (TDI) is particularly useful for evaluating myocardial tissue motion; compared to strain imaging, TDI has a higher temporal resolution, is less time consuming, also it has specific clinical applications and benefits: assessment of diastolic function, assessment of left and right ventricular function, evaluation of ventricular dyssynchrony, and evaluation of constrictive/restrictive syndromes.

Keywords: Tissue Doppler, myocardial tissue motion, ventricular dyssynchrony, diastolic function, constrictive, restrictive

INTRODUCTION

Tissue Doppler Imaging (TDI) has made a long way in echocardiography since the first investigations on tissue characterization and cardiac tissue motion were documented; indices on deformation imaging (scattering characteristics, tissue motion and velocity) were evoked in investigational studies more than 20 years ago [1,2].

TDI measures myocardial velocity and velocity gradient, which are indirect indices of myocardial deformation whereas strain imaging is a more sensitive technique for quantifying regional myocardial deformation. Besides being a more sensitive technique and angle independent, strain imaging allows early detection of many subclinical myocardial diseases such as idiopathic dilated cardiopathy. However, TDI has a higher temporal resolution than strain imaging, it is a less time-consuming technique and it has specific and relevant clinical applications: assessment of diastolic dysfunction, assessment of left ventricular (LV) and right ventricular (RV) systolic function, evaluation of ventricular dyssynchrony, assessment of constriction versus restriction. In addition, TDI has other uses in today echocardiography, like study of left atrial mechanics and prediction of myocardial viability, though these advanced techniques are not commonly used in every day practice.

Assessment of diastolic function

the latest guidelines on diastolic function study focus on TDI both for diagnosis and grading of diastolic dysfunction [3]. According to these guidelines, evaluation of diastolic function could not be made without TDI; among the 5 parameters used for diagnosis of diastolic dysfunction, TDI parameters comprise 3 of them (early diastolic septal velocity, medial E’; early diastolic lateral wall velocity, lateral E’; E/E’). Of note, acquisition of E’ is made via spectral TDI, and it is feasible with relatively high relevance even when sonographic signal is poor. However, a good alignment is a pre-requisite in order to avoid under-estimation of E’ value, given that TDI is an angle-dependent technique. TDI for assessment of diastolic function is a routine test in every day echocardiography, however it is particularly important for patients with preserved ejection fraction heart failure and for evaluation of LV filling pressure.

Assessment of systolic function

The most commonly used parameter is longitudinal septal systolic velocity (Sm) for LV systolic function and it is acquired as for E’ with the same spectral TDI mode. As for the E’ acquisition, Sm acquisition is relatively feasible even when sonographic signal is relatively poor, however a good alignment is necessary given the angle-dependency.
For the RV, TDI is used usually in conjunction with TAPSE (tricuspid annulus plane systolic excursion) at the level of basal RV free wall; contrary to the Sm value, TDI of the lateral RV free wall is indicative of the global RV function and not only local wall motion [4]. TDI assessment of LV systolic function is particularly important in subclinical LV systolic dysfunction given that Sm showed to be a sensitive marker of early LV systolic dysfunction.

**Assessment of synchrony**

Whether at the level of intraventricular or interventricular motion, TDI with its high temporal resolution allows a sensitive estimation of all synchronization parameters. Of note, colour TDI is a qualitative technique that allows visual estimation of synchrony, using different coded colours that may be translated into curves (semi-quantitative technique) showing the earliest and latest segment contraction. Moreover, spectral TDI is useful for tissue synchronization studies, and commonly used parameters are “time to onset” (from beginning of QRS to onset of systolic motion on spectral curve), “time to peak” (from onset to peak of motion on spectral curve). Ventricular synchrony study is particularly useful for patients with systolic dysfunction and wide QRS for whom resynchronization therapy is evoked, also it has a prognostic value during the follow-up of patients with a cardiac resynchronization device [5].

**Assessment of constriction and restriction**

TDI is a valuable method to differentiate restrictive cardiomyopathy from constrictive pericarditis. Values of septal E’ above 8 cm/s strongly favour the diagnosis of constriction while values less than 6 cm/s favour constriction. Values of medial E’ ≥6 and ≤8 may represent mixed constriction and restriction, and therefore additional parameters may help to clear the situation: annulus reversus (medial E’ > lateral E’) and hepatic veins expiratory end-diastolic reversal velocity strongly suggest constriction [3].

**Other uses of TDI**

Many other uses of TDI have been suggested, namely to assess left atrial mechanics and risk of supraventricular arrhythmia [6,7]. Moreover, TDI coupled with strain rate imaging helps to assess myocardial function and viability in ischemic cardiomyopathy; however, strain imaging is more sensitive in these indications given its angle independency and its higher spatial resolution [8].

**CONCLUSION**

TDI is a relatively simple echographic modality, relevant even when sonographic signal is moderate, and has special applications in many cardiovascular conditions like assessment of diastolic function, systolic longitudinal LV and global systolic RV function, inter- and intraventricular synchrony, constriction, and restriction.

**CONFLICT OF INTEREST**

The author(s) declared no potential conflicts of interest with respect to the authorship, and/or publication of this article.

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