

Cardiac Contractility Assessment in Rotary Blood Pump Recipients Derived from Pump Flow

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Abstract—A new cardiac contractility index derived from pump flow (I_Q) has been developed for rotary blood pumps (RBPs) recipients, to determine preservation and eventual recovery of the remaining cardiac function. Pulse flow indices were used for comparison with I_Q during pump speed changes.

Pump flow was recorded in animal experiments and clinically in RBP recipients (MicroMed DeBakey LVAD[®]) at different pump speeds. I_Q was derived from the maximal derivative of pump flow versus QP2P relationship (dQ/dt_{\max} vs. QP2P) during speed variations. I_Q was compared to classical indices. Further, simple currently used parameters such as peak to peak of pump flow (QP2P) and pulsation index (PI = QP2P divided by mean pump flow) were calculated.

I_Q was speed-independent for both pumps, and correlated well in animal experiments with classical invasively measured contractility index. Simple parameters (QP2P and PI) depended on speed and could be used only for estimation of percentage of support, but not for characterization of cardiac contractility.

In conclusion the cardiac contractility index I_Q can be derived from pump flow (I_Q) only. It allows easy and non-invasive continuous access to cardiac function.

Keywords— Cardiac contractility index, Rotary blood pump, Cardiac recovery, Pulsatility index and LVAD

I. INTRODUCTION

Nowadays, rotary blood pumps (RBPs) are not only used as a left ventricular assist devices (LVADs) for bridge to transplantation in end stage heart failure patients but also for a bridge to recovery (BTR) and destination therapy (DT) [1-9]. A reliable monitoring tool for cardiac contractility in RBP recipients is extremely important for diagnostics and therapeutic decisions on heart protection and eventual weaning strategies.

Several cardiac contractility indices have been established: the maximal derivative of left ventricular pressure (dP/dt_{\max}), the dP/dt_{\max} versus instantaneous developed isovolumetric pressure (dP/dt_{\max} vs. IP), the dP/dt_{\max} versus end diastolic volume (dP/dt_{\max} vs. EDV), the preload recruitable stroke work (PRSW), the end systolic elastance (E_{es}), and the maximal elastance (E_{\max}) [10-16]. They all require

the determination of intraventricular pressure and/or volume, which needs complex and mostly invasive procedures.

However, with current RBPs technology, the pump flow signal is available either measured or estimated in most of the pumps. We could develop a new cardiac contractility index (I_Q) derived from this signal only. In a previous paper, we demonstrated the correspondence of I_Q to classical indices both in computer simulation and animal experiments [17]. In the following, these results are summarized and first clinical data are presented.

II. MATERIALS AND METHODS

A. Definition of I_Q

The new pump flow derived contractility index (I_Q) is the slope of the linear relationship between the maximal derivative of pump flow and the peak-to-peak valued of pump flow (dQ/dt_{\max} vs. QP2P). The I_Q is calculated as:

$$I_Q = \frac{dQ/dt_{\max}}{QP2P - Q_0} \quad [s^{-1}] \quad (1)$$

In the formula, dQ/dt_{\max} is the maximal derivative of pump flow, QP2P is the peak-to-peak value of the pump flow in each heart cycle, and Q_0 is the extrapolated value-axis intercept of dQ/dt_{\max} vs. QP2P regression line. Q_0 is calculated via speed variation.

B. In-vivo study

In animal experiments, a MicroMed-DeBakey LVAD[®] was implanted in seven healthy sheep (102 ± 20 kg). The study protocol was approved by the animal ethics committee of the Medical University of Vienna. The sheep were instrumented with a pressure-tip catheter (Millar Instruments Inc, Houston, USA) to measure the LVP, an ultrasonic flow probe (Transonic Systems Inc, Ithaca, USA) to measure pump flow, and a conductance catheter (CD Leycom, Zoetermeer, The Netherlands). Cardiac contractility was changed by pharmacological interventions with verapamil, first to reduce the contractility from the control condition and then, with noradrenalin to increase contractility

again. The detail of experiment setup was presented in our previous work [17-18].

The I_Q was compared then with a ventricular-pressure derived index that has been reported to be relatively load independent [14]:

$$dP/dt_{\max} \text{ vs. } IP = \frac{dP/dt_{\max}}{IP - b_0} \text{ [s}^{-1}\text{]} \quad (2)$$

In the formula, dP/dt_{\max} is the maximal derivative of invasively measured LVP, IP is the instantaneous developed isovolumetric pressure (or pressure at time of dP/dt_{\max}) in every heart cycle, and b_0 is the extrapolated value-axis intercept of dP/dt_{\max} vs. IP regression line.

C. Application to clinical data

Subsequently, I_Q was derived from clinical data, coming from a previous study with the Micromed-DeBakey LVAD[®] [19]. Patient data included reactions to speed adjustment within 7.7 to 9.46 krpm. The flow signal was recorded from the built-in flowmeter using data acquisition hardware (DS1103, dSPACE, Paderborn, Germany) at a sampling frequency of 100 Hz and resolution of 16 bits. The flow signal was filtered with a FIR filter (with a Hamming window of 32 samples) at a cutoff frequency of 15Hz.

I_Q was also compared to flow pulsatility and flow pulsatility index, which in daily clinical practice are used as indicators of remaining heart action:

$$QP2P = Q_{\max} - Q_{\min} \quad [\text{l/min}] \quad (3)$$

$$PI = \frac{Q_{\max} - Q_{\min}}{Q_{\text{mean}}} \quad [-] \quad (4)$$

Here, $QP2P$ is the peak-to-peak value of pump flow, Q_{\max} is the maximal pump flow, Q_{\min} is the minimal pump flow, PI the pulsatility index, and Q_{mean} the average of pump flow in each heart cycle.

III. RESULTS

Both, in the animal experiment and in human data, the independency of the index to speed variation could be proven by the linear relationship between dQ/dt_{\max} vs. $QP2P$ at different speed levels (Example fig. 1). In animal experiments, I_Q was proven to be sensitive to pharmacologically induced changes of cardiac contractility in a manner similar to dP/dt_{\max} vs. IP (Fig. 2).

The plots of I_Q , $QP2P$ and PI versus pump speed (calculated beat by beat) are shown in Fig. 3, Fig.4 and Fig.5, respectively. The independent of mean I_Q on speed variation

and the strong dependence of $QP2P$ and PI on such variation can be seen.

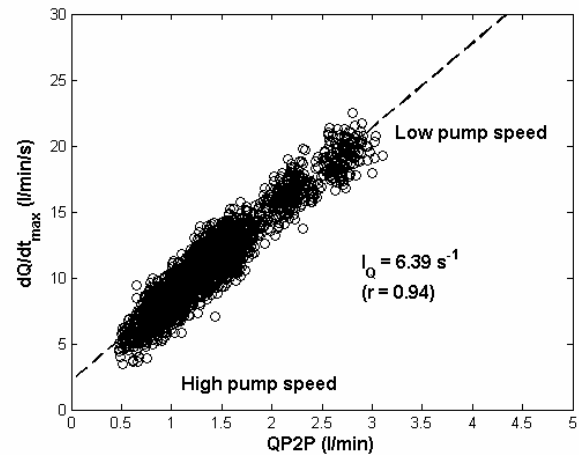


Fig. 1 dQ/dt_{\max} vs. $QP2P$ relationship in a patient at varying speed (7.7-9.5 krpm),

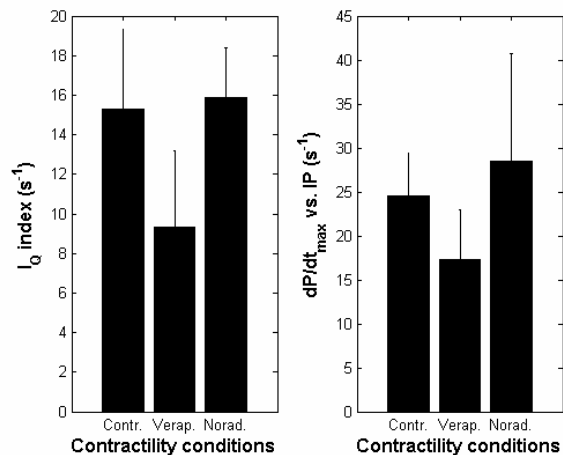


Fig. 2 I_Q compared to the classic index dP/dt_{\max} vs. IP in the animal experiment with pharmacologically induced varying contractility (Contr.: control condition; Verap.: verapamil administration; Norad.: noradrenalin administration).

IV. DISCUSSION AND CONCLUSION

This study demonstrates that the I_Q index is sensitive to cardiac contractility changes, but independent on pump speed and independent on preload and afterload changes (for details see [17]), which are inevitable prerequisites for a contractility index. Certainly the clinical data presented here, are preliminary and further clinical studies in various states and types of underlying disease should be performed. Certainly, the method can be applied also to pumps with

indirectly determined flow pattern, provided that the frequency contents of the calculated flow signal allow proper estimation of dQ/dt_{max} .

Advantageously, the index can be derived purely from pump-data, only requiring minor speed variations from time to time. Due to this simple procedure, it can be applied even quasi-continuously. It can certainly not substitute occasional Ultrasound inspections, which deliver far more details on the local distribution of contractile muscular tissue, but should allow new and immediate insights into the trends and variability of cardiac contractility in different therapeutically regimens.

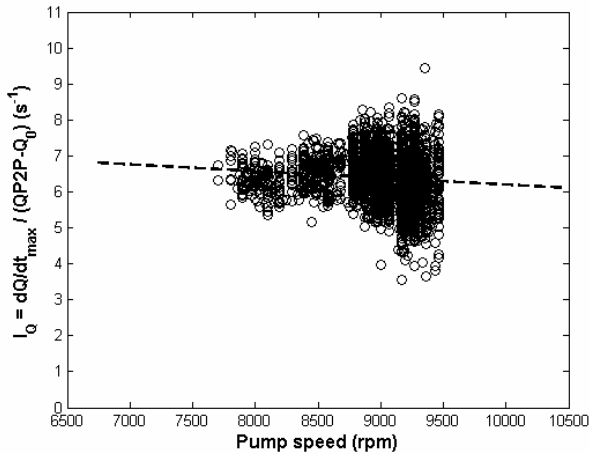


Fig. 3 Derived from the same speed data as shown in fig 2, the independence of the index to pump speed is demonstrated. Due to beat-to-beat variation the mean over several beats should be taken.

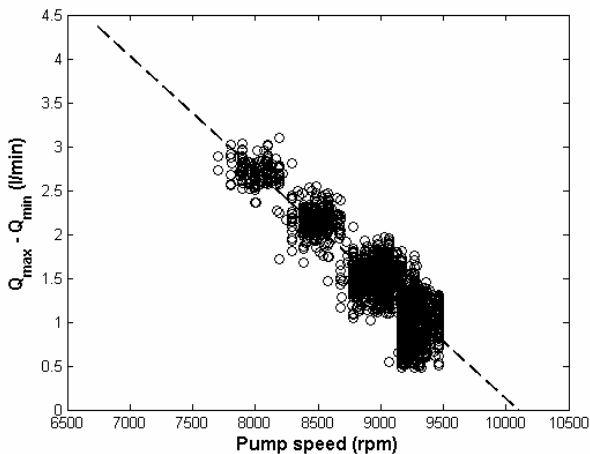


Fig. 4 Peak-to-peak of pump flow ($Q_{max}-Q_{min}$) at stable cardiac condition depends strongly on pump speed. ($r = -0.88, p < 0.001$).

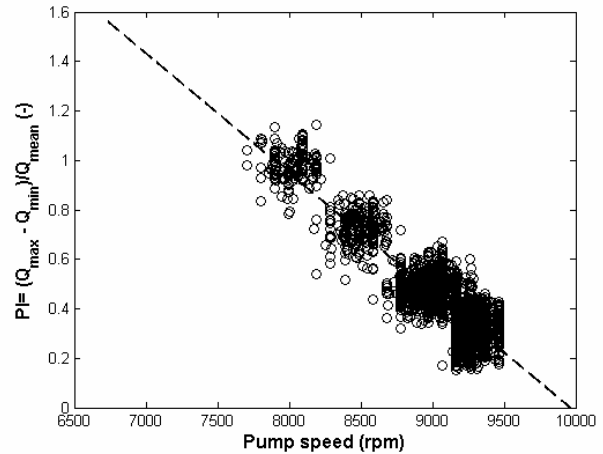


Fig. 5 Also pulsatility index PI depends at stable cardiac condition strongly on pump speed ($r = -0.89, p < 0.001$).

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