

POSTER PRESENTATION

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Simultaneous flow dynamics in small and great thoracic vessels during physiological stress tests and normal breathing using real-time cardiac magnetic-resonance

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Background

Aim of this study was to measure blood flow dynamics in the descending aorta (AD) and azygos vein (AzV) using Muller (MM) and Valsalva (VM) maneuver as physiological stress tests and normal breathing in real-time cardiac magnetic resonance imaging (RT-CMR).

Methods

Blood flow was measured in $n = 20$ healthy volunteers in the AD and AzV with RT-CMR including an image acquisition of under 40 ms and undersampled further development of established FLASH sequences. The volunteers were instructed to perform MM as well as VM in a 3T MR system, TR/TE were 3.44/2.76 ms. Pressure changes were digitally monitored to ensure sufficient pressure gradients.

Results

Muller maneuver: At the beginning of MM, flow per heart beat (hb) in the AD (56 ± 9.4 SD ml/hb $\sim 100\%$) decreased significantly ($p < .001$, to 46 ± 10.4 ml/hb $\sim 82\% \pm 13$ ml/hb at early strain phase). In the AzV flow per hb increased significantly ($p < .001$) during MM (normal breathing 1.8 ± 1.5 ml/hb $\sim 100\%$ vs. early strain 5.5 ± 3.6 ml/hb $\sim 309\% \pm 555$ ml/hb). Heart rate increased significantly ($p < .001$, normal breathing 74 ± 9.9 bpm $\sim 100\%$ vs. early strain 79 ± 12.6 bpm $\sim 108\% \pm 8$ bpm). Average peak

velocity decreased significantly ($p < .001$) in AD (normal breathing 62 ± 11.8 cm/s $\sim 100\%$ vs. early strain 52 ± 11.6 cm/s $\sim 84\% \pm 10$ cm/s) and increased significantly ($p < .01$) in the AzV (normal breathing 16 ± 4.4 cm/s $\sim 100\%$ vs. early strain 25 ± 8.4 cm/s $\sim 162\% \pm 56$ cm/s). Average area in AD ($p < .001$) and AzV ($p < .05$) increased during MM. Valsalva maneuver: During VM flow per hb in the AD (regular breathing 56 ± 9.8 ml/hb $\sim 100\%$) decreased significantly ($p < .001$) until the end of VM (late strain 23 ± 10.9 ml/hb $\sim 36\% \pm 18$). Similar significant changes could be found during VM for average peak velocity (regular breathing 62 ± 12.9 cm/s $\sim 100\%$ vs. 42 ± 11.8 cm/s late strain $\sim 61\% \pm 25$). Heart rate increased significantly compensatory ($p < .001$) from 75 ± 9.1 bpm during regular breathing ($\sim 100\%$) to 99 ± 17 bpm at late strain phase ($\sim 119\% \pm 43$). Flow per hb changed significantly ($p < .001$) in the AzV from 2.2 ± 1.4 ml/hb ($\sim 100\%$) to 2.0 ± 2.1 ml/hb ($\sim 111\% \pm 223$). Average peak flow velocity changed less significant ($p < .05$) from 17 ± 3.4 cm/s ($\sim 100\%$) to 14 ± 5.7 cm/s ($\sim 75\% \pm 43$).

Table 1 Patient's characteristics

Type	Total \pm SD
Age	33.86 \pm 12.75
Weight	71.45 \pm 13.69
Height	1.76 \pm 0.11
Body mass index	22.82 \pm 2.62
Cardiovascular risk factors	$n = 1^*$
Gender	male (12/8)

* One proband with nicotine abuse (many pack years)

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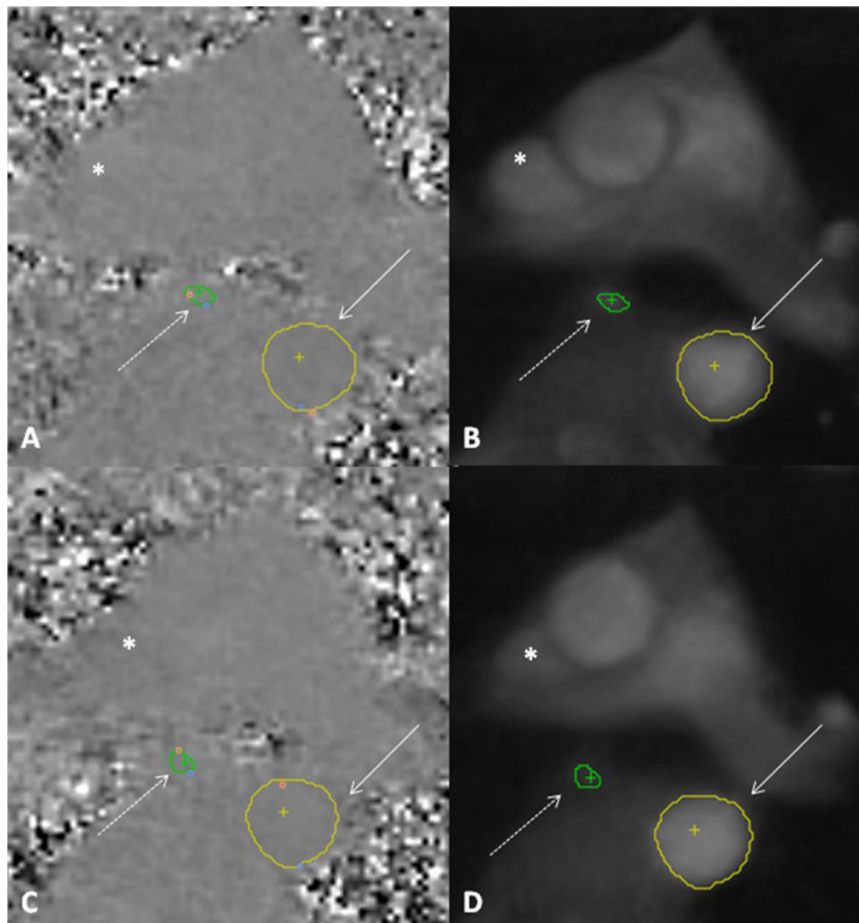


Figure 1 Flow in the aorta descendens (arrow, yellow contour) and vena azygos (dashed arrow, green contour), images in systole (28-years old healthy male volunteers). Normal breathing (A-B) and Valsalva maneuver (C-D). Lumen of the aorta und vena azygos is compressed during Valsava (C, D), especially reduced in the superior vena cava (white stars). The luminal area and diameter of thoracic vessels is reduced under elevated intra-thoracic pressure. Scan area (localizers) were adapted to Valsalva and normal breathing, because mediastinal and heart structures are moved during variation of the intra-thoracic pressure.

The luminal area of the AzV changed non-significantly, in AD significantly ($p < .001$). All values are with standard deviation, \pm SD.

Conclusions

RT-CMR reliably and simultaneously determines flow dynamics during physiological stress. Compared to routinely used Cine-CMR it is able to show beat to beat variability in great and small thoracic vessels in detail. In comparison to echocardiography, it enables the investigator to measure flow simultaneously in thoracic descending aorta and azygos vein. CMR might provide new diagnostics insights into aortic diseases or portocaval hypertension in future.

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