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This is the author's manuscript
Original Citation:
Availability:
This version is available http://hdl.handle.net/2318/1654949 since 2023-06-02T12:56:48Z
Published version:
DOI:10.1002/jum.14235
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# Vena cava responsiveness to controlled isovolumetric respiratory efforts

## Manuscript category: Original research

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Running title: IVC response to respiratory efforts

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# 1 ABSTRACT

Objective. Respirophasic variation of inferior vena cava (IVC) size is affected by large variability
 with spontaneous breathing. This study aims at characterizing the dependence of IVC size on
 controlled changes in intra-thoracic pressure.

5 Methods. Ten healthy subjects, in supine position, performed controlled isovolumetric respiratory 6 efforts at functional residual capacity, attaining positive (5, 10, 15 mmHg) and negative (-5, -10, -15 7 mmHg) alveolar pressure levels. The isovolumetric constraint implies that equivalent changes are 8 exhibited by alveolar and intrathoracic pressures during respiratory tasks.

**Results.** The IVC cross sectional area (CSA) equal to  $2.88\pm0.43$  cm<sup>2</sup> at baseline (alveolar pressure 9 10 = 0 mmHg) was progressively decreased by both expiratory and inspiratory efforts of increasing 11 strength, with diaphragmatic efforts producing larger effects than thoracic ones: -  $55\pm15\%$  decrease, 12 at +15 mmHg of alveolar pressure (p<0.01), -80±33±12% at -15 mmHg diaphragmatic (p<0.01), -13 33±12% at -15mmHg thoracic. Significant IVC changes in size (p<0.01) and pulsatility (p<0.05), 14 along with non significant reduction in the response to respiratory efforts, were also observed 15 during the first 30 min of supine rest, detecting an increase in vascular filling, taking place after 16 switching from the standing to the supine position.

17 Conclusion. This study quantified the dependence of IVC CSA on controlled intra-thoracic 18 pressure changes and evidence the stronger influence of diaphragmatic over thoracic activity. 19 Individual variability in thoracic/diaphragmatic respiratory pattern should be considered in the 20 interpretation of the respirophasic modulations of IVC size.

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KEYWORDS Inferior vena cava; caval index; breathing pattern; alveolar pressure; IVC
 collapsibility; Valsalva Maneuver.

#### 24 INTRODUCTION

25 Ultrasonographic monitoring of the inferior vena cava (IVC) is a noninvasive, widely adopted procedure to derive information about the volume status of patients as well as their possible 26 27 responsiveness to fluid therapy (1-4). IVC diameter varies during the respiratory cycle due to 28 pressure changes in the thorax and abdomen (5, 6) and depending on vessel compliance, which in 29 turn depends on filling pressures and volume status (5). These variations are quantified by the caval 30 index (CI), defined as the difference between expiratory and inspiratory IVC diameters divided by 31 the expiratory diameter. However, proposed diagnostic cut-offs for spontaneously breathing patients 32 vary considerably in the literature and caution in relying on CI for fluid therapy management is 33 generally recommended (3, 5).

34 Several factors, mostly related to the variability of spontaneous breathing, may affect assessment of 35 the CI and potentially undermine its reliability. 1) Spontaneous breathing is intrinsically irregular in 36 both amplitude and frequency (7) and is affected by emotional status, pain, and pathology (8). 2) 37 The respiratory pattern changes depending on gender and age (9, 10), also in terms of its 38 thoracic/diaphragmatic components, which affect in different ways abdominal pressure and IVC 39 diameter (6). 3) The IVC moves considerably during respiration (11), introducing errors in the 40 assessment of CI, unless some advanced image analysis is implemented (12). 4) Often, the cross-41 section of the IVC is not circular, and may exhibit anisotropic changes in size due to modifications 42 of filling pressure or breathing (11, 13). Consequently, the arbitrary choice of a given section for the measurement of IVC diameter may not be adequate to assess changes in its size and pulsatility and 43 44 the assessment of the whole cross-sectional area has instead been proposed (14).

This study aims at characterizing the actual dependence of IVC size on changes in intrathoracicpressure, in the absence of the above described confounding factors.

47 To this aim IVC deformations were assessed in static conditions (short apnea), during controlled

48 isovolumetric respiratory efforts and quantified in terms of changes in cross-sectional area (CSA).

In particular, attention was focused on 1) the IVC CSA response to controlled positive and negative changes in alveolar pressure, as produced by expiratory and inspiratory efforts, respectively, 2) the specific effect of thoracic/diaphragmatic involvement in inspiratory efforts, 3) testing whether such respiratory maneuvers may potentially be employed to detect changes in vascular filling, which spontaneously occur when switching from the standing to the supine position.

54

#### 55 MATERIALS AND METHODS

#### 56 Subject selection

Subjects were recruited among PhD students and partly among amateur swimmers. Out of 16 subjects screened, 6 were excluded because of low quality of the ultrasonographic images (n=4), inability to correctly perform respiratory maneuvers (n=1), and for complete collapse of the IVC already at the low-pressure maneuvers (n=1). The study was then conducted on 10 healthy volunteers (4 males and 6 females, age 27  $\pm$ 7, BMI 20.7 $\pm$ 1.6). The study was approved by the Ethics Committee of Turin University. All participants gave their informed consent according to the principles of Helsinki Declaration.

#### 64 **Experimental setup**

During the whole experimental procedure, participants maintained a supine position with head slightly raised with respect to the body. They were asked to either relax and breathe normally or perform respiratory efforts. The upper part of the IVC was visualized by subxyphoid or right lateral intercostal approach by a single echographer (PP), taking into account anatomical landmarks as the left portal branches and the ligamentum venosum.

Video clips of the IVC were recorded in the transversal plane (see below) with an ultrasonographic
unit (Mylab25 Gold ESAOTE; Genoa, Italy) equipped with a 2-5 MHz convex probe.

72 Air pressure during respiratory maneuvers was measured at the mouth level by a pressure monitor

- 73 (BP-1 pressure monitor, World Precision Instruments, Florida, USA) equipped with a mouthpiece,
- 74 providing no air leakage. In the absence of airflow, the pressure measured at the mouth during

respiratory efforts coincides with the alveolar pressure, provided the glottis remains open. The analog output of the device was digitally acquired (sampling frequency 200 Hz, CED 1401micro, and Spike2 acquisition software, Cambridge, UK) and displayed on a monitor to provide a visual feedback to the subject. In order to synchronize the recording of alveolar pressure signal with the IVC video clip, a digital trigger signal generated by the program (Spike2, CED, UK) was acquired with alveolar pressure and fed to the ECG input of the echograph, thus being displayed and recorded in the video clip.

### 82 **Respiratory maneuvers**

83 In a preliminary session, the subjects were invited to practice diaphragmatic and thoracic breathing 84 and learned to perform the controlled isovolumetric respiratory efforts while maintaining the glottis 85 open. Respiratory maneuvers consisted of isovolumetric respiratory efforts conducted at functional 86 residual capacity (FRC) as follows. At the end of a spontaneous expiration a trigger signal was 87 manually generated by the experimenter and, after 4 s of apnea (basal condition), the subjects 88 performed the controlled expiratory/inspiratory effort through the mouthpiece according to pre-89 defined positive/negative target levels, and maintained them for 10 seconds (Fig. 1). At the end of 90 each maneuver, the mouthpiece was removed and the subject could relax and breathe normally. 91 While during expiratory efforts both abdominal and thoracic muscles were simultaneously 92 recruited, inspiratory efforts were performed by selectively activating the diaphragm or thoracic 93 muscles. The accuracy of the inspiratory maneuver was checked by the experimenter based on visual inspection of thoracic and abdominal movements. 94

## 95 **Protocol**

A resting period of 30 minutes in supine position was allowed to stabilize transcapillary fluid exchange. During this time, respiratory efforts at -5 and +5 mmHg were performed (at 0, 15 and 30 min) with the aim of testing the effect of possible changes in blood volume with time. To the same purpose, 30-s video clips of IVC cross section were recorded during spontaneous breathing.

After this time, a sequence of both thoracic and diaphragmatic inspiratory efforts at -5, -10 and -15 mmHg was performed along with isovolumetric expiratory efforts at 5, 10, and 15 mmHg according to the diagram in Fig. 2. Twenty-second video clips in the transversal plane of the IVC were taken at each respiratory maneuver, allowing to monitor its cross-section before and during the maintained change in alveolar pressure. Resting intervals of at least 30 s were allowed between consecutive maneuvers.

## 106 Image processing

107 All videos were processed by a custom-made software (implemented in Matlab 2015a, The 108 Mathworks) based on automated detection of the IVC wall, providing continuous assessment of the 109 IVC cross sectional area (CSA, Fig. 3) (manuscript in preparation). The CSA was estimated for 110 each frame and a time series obtained with sampling frequency equal to the video frame rate, 111 between 11 Hz and 19 Hz (depending on current echographic settings).

The trigger signal recorded in the video clip was automatically detected and used to re-align in time the CSA signal with the alveolar pressure recording, separately acquired. The CSA signal was then low-pass filtered with cut-off frequency of 2 Hz (Butterworth anti-causal IIR filter of order 4), which preserved both cardiac and respiratory oscillatory components and re-sampled at 200 Hz, as the alveolar pressure signal.

The responses to respiratory maneuvers were analyzed in terms of changes in IVC CSA.. Average CSA was computed over 4-s intervals, before (baseline) and during the respiratory effort (effect), after alveolar pressure reached the target level (see Fig. 1). In baseline intervals, for each heartbeat, the cardiac caval index was calculated as: CCI = (max(A)-min(A))/max(A), where A is the IVC-CSA computed by the algorithm. An average CCI was then obtained for each interval.

Responses to the respiratory maneuvers were calculated as relative changes referred to baseline[(effect-baseline)/baseline].

Assessment of the "classical" caval index, CI= [max(D)-min(D)]/max(D), D being the IVC
diameter, was computed off-line by the same echographer (PP), from the 30s video clips recorded

during spontaneous breathing using the "frame-by-frame" method, as the average of 3
measurements collected on 3 different respiratory excursions (MyLabDesk, Esaote).

#### 128 Statistical analysis

129 Data are expressed as mean±SD in the text and displayed as mean±SEM in bar diagrams. Statistical 130 significance of respiratory effects was assessed by repeated-measures one-way ANOVA (factor: pressure level), for expiratory efforts, by two-way ANOVA (factors: pressure level and 131 132 diaphragmatic/thoracic pattern) for inspiratory efforts, with Bonferroni correction for multiple 133 comparisons, and by the Dunnett's test for comparison with the basal value. Statistical significance of changes during the stabilization phase with respect to initial condition (t=0) was assessed with 134 135 the Dunnett's test. The significance cut-off was p<0.05. All analyses were carried out with GraphPad Prism version 6.0c (GraphPad Software, San Diego California, USA). 136

137

#### 138 **RESULTS**

139 The effect of controlled isovolumetric respiratory efforts was tested after 30 min in supine position.

An example of the ensuing changes in IVC CSA is shown in Fig. 4, for a representative subject. Coherent tracings of CSA and alveolar pressure are plotted during expiratory (A) and thoracic and diaphragmatic inspiratory efforts (B, C). It can be observed that the maneuver produced immediate changes in both average size and pulsatility of IVC strictly related in their time course to changes in alveolar pressure. Furthermore, larger effects appear to be produced by diaphragmatic than thoracic inspiratory efforts at -5 mmHg, while even smaller effects are produced by the expiratory effort at +5mmHg (C).

#### 147 **Response to inspiratory efforts**

On average, the inspiratory maneuvers induced progressive reduction in CSA with decreasing alveolar pressures (p<0.001), the effect being significantly larger with diaphragmatic than thoracic efforts (p<0.05) (Fig. 5). In particular, the average CSA was  $2.88\pm0.43$  cm<sup>2</sup> in basal conditions (Fig. 5, dashed line), decreased to  $2.0\pm0.4$  cm<sup>2</sup> (n.s.) and to  $0.53\pm0.20$  cm<sup>2</sup> (p<0.01) during thoracic and diaphragmatic inspiratory efforts at -15 mmHg, respectively. (Fig. 5). Dunnett's test indicates that while thoracic inspiration did not provoke any statistically significant change from basal condition, diaphragmatic inspiration induced significant changes at all pressure levels.

In relative terms, the decrease of CSA ranged from 39±11% to 80±10% in diaphragmatic and up to
33±12% in thoracic maneuvers.

158 In some subjects, collapse of the IVC up to complete occlusion was observed during the maneuver.

159 This occurred more frequently at increasing negative pressures and during diaphragmatic efforts

160 (number of subjects: 1 at -5 mmHg; 5 at -15 mmHg).

## 161 **Response to expiratory efforts**

- 162 Expiratory efforts induced a progressive decrease in the IVC CSA with increasing positive alveolar
- 163 pressure (p< 0.05, Fig. 6). In relative terms, the CSA decreased by  $15 \pm 6\%$  (n.s., at +5mmHg) and

164 by 55±15% (at +15 mmHg, p<0.01).

165 In 3 subjects, complete collapse of the IVC was observed at +15 mmHg.

## 166 **Changes occurring in the stabilization phase**

Here we discuss the possible blood volume changes occurring during the first 30 min in the supine position (stabilization phase) on the different variables measured, a complete set of recording being achieved in 8 subjects (in two subjects some of the recordings were excluded from the analysis due to delays in following the protocol).

- 171 During the stabilization phase, the IVC CSA (measured in resting conditions at functional residual
- 172 capacity) was dependent on time, increasing from  $2.37\pm0.2$  cm<sup>2</sup> (t0) to  $2.90\pm0.21$  cm<sup>2</sup> (t30, p<0.01)
- 173 (see Fig. 7A), where t0 and t30 refer to the conditions at the beginning of the protocol and after 30
- 174 min, respectively, as indicated in Fig. 2. Assessment of CSA at the end of the protocol

175  $(2.89\pm0.2\text{cm}^2 \text{ at about 45 min})$  revealed that no further changes occurred during the last part of the 176 protocol.

177 The CCI exhibited a specular trend: from  $0.22\pm0.01 \text{ cm}^2$  (t0) to  $0.19\pm0.01 \text{ cm}^2$  (t30, p<0.05, Fig. 178 7B).

The response to respiratory efforts also exhibited a decreasing trend during the first 30 minutes (Fig. 7D). IVC collapse produced by a 5-mmHg diaphragmatic inspiratory effort decreased from  $60\pm12\%$  at t0 baseline to  $40\pm11\%$  at t30. A similar trend was exhibited by the response to thoracic inspiratory efforts (-5 mmHg, Fig. 7E), decreasing from  $20\pm15\%$  at t0 to  $1\pm10\%$  at t30, and expiratory efforts (+5 mmHg, Fig. 7F), decreasing from  $23\pm15\%$  (t0) to  $15\pm5\%$  (t30), although not reaching statistical significance.

185 The standard cross-sectional CI, measured manually during spontaneous breathing, confirmed the 186 same trend decreasing from  $0.41\pm0.19$  to  $0.31\pm0.13$  (p<0.05) (Fig. 7C).

187

## 188 **DISCUSSION**

189 With this study we show that controlled isovolumetric respiratory efforts produce consistent 190 changes in IVC CSA. During inspiratory efforts, these effects are strongly dependent on whether a 191 thoracic or diaphragmatic effort is made. In particular, diaphragmatic inspiratory efforts produced 192 the largest decreases in CSA (averaging across subjects, from 30% to 80% of basal condition, at -5 193 mmHg and -15 mmHg, respectively), while changes during thoracic inspiration were less than a 194 half. Expiratory efforts also decreased CSA (up to 65% at +15 mmHg). Preliminary results collected 195 during the stabilization phase in the supine position indicated that the response to standardized 196 respiratory efforts, along with other parameters including the IVC CSA and the cardiac and 197 respiratory CI, are potential indicators of changes occurring in the vascular volume.

In this study we adopted an approach that excludes most of the confounding factors affecting the
CI, the classical index of IVC collapsibility. 1) *Breath-to breath variability*, i.e., the amplitude of

200 the respiratory movements from breath to breath. Tobin et al (7) observed a coefficient of variation 201 in breath-to-breath tidal volume of 30% and 44% and in respiratory frequency of 20 and 28% in 202 young and old subjects, respectively. Since amplitude and speed of respiratory movements directly 203 affect intra-thoracic and abdominal pressures, their variability is expected to directly translate into 204 CI variability. Variability of spontaneous breathing was prevented in this study in which IVC size 205 changes were assessed in response to standardized respiratory maneuvers performed at constant 206 lung volume (functional residual capacity). 2) Thoracic/diaphragmatic breathing. Variability of the 207 respiratory pattern also concerns the relative proportion of thoracic vs. diaphragmatic activation (in 208 the inspiratory phase), which also exhibits breath-to-breath variability (CoV= 22-31%) (7) as well 209 as dependence on gender and age (9, 10). Although it is well known that thoracic and diaphragmatic 210 breathing affect abdominal pressure differently (15) and thus also the CI (6), it is impossible to 211 control for this confounder in patients because some self-consciousness and training are required for 212 thoracic or diaphragmatic respiration to be correctly performed. Preliminary training was necessary 213 for the healthy subjects of this study. Possibly, because of this difficulty, the issue is generally 214 overlooked and its implications ignored in the interpretation of the CI. 3) Respiratory movements of 215 the IVC. Longitudinal displacement of the IVC in the order of 2 cm in cranio-caudal direction has 216 been shown (11) and this may introduce an error of up to 30% in the estimate of the CI, depending 217 on the shape of the IVC (12). This error affects transversal as well as M-mode longitudinal 218 measurements, although the latter were recently indicated as the most sensitive indicators of 219 changes in volume status (16). These artifacts were prevented in the present study, because all 220 measurements were taken in static conditions (short apnea). 4) Non circular IVC cross-sectional 221 shape. The CI and other similar indices of collapse are always computed on the basis of maximum 222 and minimum IVC diameters, which is a misrepresentation, because the IVC usually presents a non-223 circular cross-sectional shape. Thus, the choice of a given "diameter" is arbitrary and its temporal 224 changes may not be representative of the behavior of the whole vessel. In fact, in hypovolemic 225 patients undergoing fluid replacement, Murphy et al (13) showed that the minor axis exhibited a

five-fold increase as compared to a 5% increase of the major axis (of IVC imaged in transversal section), which they called anisotropic behavior of the IVC. This problem was prevented here by assessing changes in the cross-sectional area of IVC rather than in a single, arbitrarily chosen diameter, following the approach proposed by Nakamura et al. (14).

230

# 231 IVC changes in thoracic vs. diaphragmatic inspiratory efforts.

232 IVC size varies in response to changes in transmural pressure (i.e., the difference between internal 233 and external pressures), according to its compliance (defined as the incremental variation of vessel 234 volume induced by a change of transmural pressure). Transmural pressure may change due to 235 changes in 1) internal pressure (i.e., central venous pressure, CVP) which directly depends on 236 changes in intra-thoracic pressure and 2) external (abdominal) pressure which may increase due to 237 contraction of the diaphragm (e.g., during diaphragmatic inspiration or inspiratory efforts) or of 238 abdominal expiratory muscles (e.g., during forced expiration or expiratory efforts). In this study, the 239 subjects engaged in isovolumetric respiratory efforts, attaining selected positive and negative levels 240 of alveolar pressure that, in the absence of airflow, could be measured at the mouth-piece. Since all 241 maneuvers were performed at the same lung volume (FRC), we can assume unchanged 242 transpulmonary pressure, which implies that changes in alveolar pressure produced equivalent 243 changes in intrathoracic pressure, affecting the CVP.

244 In light of these considerations, the results can be interpreted as follows. Diaphragmatic inspiratory 245 efforts produced both a reduction of blood pressure in the IVC (by decreased intrathoracic pressure) 246 and an increase of abdominal pressure (by diaphragm contraction). These effects concurred to 247 markedly decrease IVC transmural pressure, resulting in the observed marked decrease in IVC 248 CSA. The latter was roughly proportional to the intensity of the effort performed (Fig. 5, black 249 columns). Conversely, thoracic inspiratory efforts only affected intrathoracic pressure, with virtually 250 no effect on abdominal pressure, resulting in comparably lower reductions in IVC CSA. These were 251 negligible at -5 mmHg and did not further decrease for -10 to -15 mmHg of alveolar pressure (Fig. 5, white columns). This suggests that lowering intrathoracic pressure has a limited collapsing effecton the IVC, compared to increasing abdominal pressure.

254 This interpretation fits with other data in the literature. The thoracic/diaphragmatic respiratory 255 pattern also affects venous return, which is impaired by increased abdominal pressure. In fact, Miller et al. (15) elegantly showed that abdominal pressure increases (+ 6 cmH  $_2^{O}$ ) and venous 256 return (observed at the femoral vein) is arrested during the inspiratory phase of diaphragmatic 257 258 breathing, the same effects being produced by manually compressing the abdomen. On the contrary, 259 thoracic inspiration facilitates venous return (15). Gutzeit et al. (17) recently reported that forced 260 inspiration ("suction against resistance" at -20 mmHg) decreased venous return from inferior with 261 respect to superior vena cava. Although they did not control for thoraco/diaphragmatic inspiratory 262 patterns, it is likely that the diaphragm was activated to some extent, thus impairing venous return 263 in the IVC. In uncontrolled breathing at increasing inspiratory effort (0, -5, -10 mmHg) Gignon et 264 al. (18) showed that the increased CI was highly correlated with diaphragm displacement. To our knowledge, specific thoraco/diaphragmatic effects on IVC size were only investigated by Kimura et 265 266 al. (6), who reported that diaphragmatic breathing is associated with increased IVC excursions and 267 CI, with respect to thoracic breathing at comparable tidal volumes. Based on a different approach 268 (isovolumetric efforts at comparable levels of blood volume and alveolar pressure), we show here 269 that increasing inspiratory diaphragmatic efforts progressively decreased IVC size down to 270 complete collapse in 5/10 subjects. Notably, intrathoracic pressure changes during spontaneous breathing are in the order of 2-3 cm H  $_{2}^{0}$ , well below 5 mmHg. On this basis, the data presented 271 272 here support the notion that abdominal pressure is a major determinant of IVC size and further 273 emphasize that an uncontrolled breathing pattern may confound volume status assessment based on 274 measured CI (6, 18).

275

#### 277 IVC changes during expiratory efforts

278 Attaining a positive alveolar pressure at constant lung volume results in equivalent increase in 279 intrathoracic pressure, which in turn should increase blood pressure in the IVC and increase its size. 280 However, the expiratory effort requires contraction of both thoracic and diaphragmatic expiratory 281 muscles and results in increased abdominal pressure (19), which per se produces the opposite effect 282 on IVC transmural pressure and size. On this basis, predicting the outcome on IVC size is not trivial 283 and may depend on both lung volume and how the maneuver is actually performed (20). Grant et al. 284 (20) observed reduction of the IVC in 100% of males and 50% of females performing a Valsalva 285 maneuver. In a recent study, IVC deformation by the Valsalva maneuver was reinvestigated with 286 computed tomography (21) and a systematic decrease of IVC size was reported (to 22% of resting 287 CSA, on average). However in none of these studies was the maneuver controlled or standardized in 288 terms of exerted pressure. The authors did not measure abdominal pressure simultaneously with 289 IVC size. Since it averaged about 80 mmHg, the expiratory effort was presumably close to maximal 290 in the second study (21). In this study, rather low pressure values (5, 10 and 15 mmHg) were 291 attained at the mouthpiece, as compared to the 40 mmHg commonly employed for the Valsalva 292 maneuver (22), but a progressive and significant decrease in IVC size was observed, again 293 indicating that abdominal pressure prevails on intra-thoracic pressure.

294

## 295 Detection of volume changes during the stabilization phase

A fluid shift from the extravascular to the intravascular compartment is known to occur when switching from the orthostatic to the clinostatic position (23). The increase in plasma volume, often assessed on the basis of the accompanying decrease in protein concentration and hematocrit (so called postural pseudoanemia (24)), was reported to be in the order of 8% and 12% after 15 and 60 min from changing posture (23). Other studies have substantially confirmed these figures, indicating volume shifts of about 400 ml within 30 min from postural change, from lying to standing or vice versa (24, 25). Such increase in plasma volume presumably occurred in the present 303 study, during the stabilization phase. In agreement with this hypothesis, a significant increase in 304 IVC CSA was detected, along with other effects indicating decreased IVC compliance, such as the 305 decrease in cardiac and respiratory CI and a trend to decrease in the IVC CSA response to 306 inspiratory efforts. These results emphasize the importance of allowing for a stabilization phase in 307 the supine position before starting experimental protocols and that failing to do so may introduce 308 large errors in the measurement of IVC variables. The response to a standardized respiratory 309 maneuver has the potential to reveal changes in blood volume; however further studies and larger 310 population samples are needed to characterize the sensitivity of the different parameters to the 311 actual changes in blood volume occurring in this and other conditions.

312

#### 313 Limitations

Although maneuvers were performed at constant lung volume, some movement of the diaphragm could still occur at the onset of inspiratory and expiratory efforts (although not during measurements), resulting in longitudinal displacements of the IVC that may not have been adequately compensated for by the operator. In addition, some residual movements of the diaphragm during thoracic inspiration, or of the thorax during abdominal inspiration, may have occurred. This, however, did not prevent detection of large differences in the IVC responses to the two breathing patterns.

The subjects involved were selected for echogenicity and for their ability to correctly perform the respiratory maneuvers required by the protocol. They were partly recruited among amateur swimmers, often characterized by a particularly large IVC (26), and this may account for the relatively high average IVC CSA and low CI observed.

325 A further limitation concerns the applicability of this approach to the clinical setting. The 326 implementation of these respiratory maneuvers required well-trained subjects and may be 327 unfeasible with poorly collaborative patients.

328

## 329 Conclusions

330 We describe for the first time IVC responses to positive and negative changes in alveolar pressure, 331 producing equivalent changes in intrathoracic pressure. The results indicate that diaphragmatic 332 activation, affecting abdominal pressure, is a major determinant of IVC size and suggest that 333 individual variability in the thoracic/diaphragmatic respiratory patterns may account for the large 334 variability normally observed in the respiratory CI. Implementation of isovolumetric controlled maneuvers proved effective in probing IVC compliance in the absence of the confounding effects 335 336 otherwise introduced by spontaneous breathing and potentially adequate to detect changes in 337 volume status. Further studies are required to test this possibility and to make the procedure 338 compatible with the clinical setting.

# 340 ACKNOWLEDGMENTS

341 This work was supported with grants from University of Torino.

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417 FIGURES

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Figure 1: Visual feedback. At the bottom, the visual feedback of alveolar pressure (black) and the target level to be reached (grey). At the top, the trigger signal fed to the echograph for synchronization. Dashed lines indicate the 4-s intervals used in the analysis, identifying "baseline" and "effect".

Figure 2. Experimental protocol. Bars of different height indicate expiratory (positive, grey) and inspiratory (negative) diaphragmatic (black) and thoracic (white) efforts of different magnitudes (5, 10 and 15 mmHg). Squared brackets indicate whether maneuvers were used to characterize the effects of respiratory efforts on IVC or to detect possible changes in blood volume over the first 30 min in the supine position.

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430 Figure 3. Representation of the automated detection of the IVC wall.

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Figure 4. IVC response to respiratory efforts, in a representative subject. Recordings of the crosssectional area of the inferior vena cava (IVC CSA, upper trace) in response to controlled changes in alveolar pressure (lower trace) are shown in an expiratory effort at +5 mmHg (A) and a thoracic (B) and adiaphragmatic (C) inspiratory effort at -5 mmH. Note that different effects are produced in terms of changes in CSA and pulsatility by the different maneuvers.

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Figure 5. Average IVC response to inspiratory efforts. Cross sectional area of inferior vena cava (IVC-CSA) during diaphragmatic (black) and thoracic (white) inspiratory efforts at different pressure levels. The dashed horizontal line represents the average basal value (pre-maneuver) of IVC-CSA and the grey band and error bars represents standard error. Symbols on single columns indicate significant difference from baseline. \*) p<0.05; #) p<0.01.

<sup>423</sup> 

444	Figure 6. Average IVC response to inspiratory efforts. Cross sectional area of inferior vena cava
445	(IVC-CSA) during the expiratory efforts at different pressure levels. The dashed horizontal line
446	represents the average basal (pre-maneuver). Other notations as in Fig. 5.
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448	Figure 7. Changes observed during the first 30 min in supine position. A) Cross sectional area of
449	inferior vena cava (IVC-CSA). B) Cardiac caval index (CCI). C) Respiratory caval index assessed
450	by the echographer. Response to diaphragmatic (D) and thoracic (E) inspiratory efforts at -5 mmHg
451	and to expiratory efforts (+5 mmHg, F), expressed in terms of % change in CSA with respect to
452	baseline; *) significantly different from t0, p<0.05. #) significantly different from t0, p<0.01.
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FIG.1



FIG.2













FIG.5



FIG.6







