Association of anatomical defects of the inferior vena cava and gonadal veins with pelvic varicose veins

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ARTICLE INFO

Article history:
Submitted: 30. 3. 2021
Accepted: 6. 6. 2021
Available online: 30. 11. 2021

SOUHRN

Znalost klinických a anatomických odchylek pánevního žilního systému hraje významnou úlohu v diagnos-tice a použití katetrizačních metod u pánevních varíků. Nejvýznamnější charakteristiky gonadálních žil jsou definovány flebografickým vyšetřením dolní duté žily a pánevních žil. Aby se zabránilo případným kompli-kacím, je při endovaskulárních intervencích nutno brát tyto charakteristiky v úvahu.

ARTICLE INFO

Knowledge of the clinical and anatomical variations of the pelvic venous system plays a significant role in the diagnosis and transcatheter methods of treatment of pelvic varicose veins. The most important characteristics of gonadal veins are identified by phlebography of the inferior vena cava and pelvic veins. These characteristics should be taken into account during endovascular interventions in order to avoid possible complications.

Pelvic varicose veins (PVV) is a disease characterized by dilatation of ovarian veins and pelvic venous plexuses. Nonetheless, when there is retrograde blood flow, ovarian venous insufficiency can be encountered in patients with normal venous diameter (4–5 mm). According to various authors, the incidence of PVV among females is approximately 30% which shows the gravity of this medical and social problem. Per several studies, 10–70% of PVV patients also develop varicose veins of the lower extremities:1 in 10–62% of cases, the cause of recurrent lower limbs varicosities is pelvic-perineal reflux.2,3

The diagnosis and treatment of PVV remains the focus of attention among phlebologists and vascular surgeons, because there are still no generally accepted recommendations that dictate treatment algorithms for patients with unusual venous anatomies as well as clear indications and contraindications for surgical cure.4 Phlebography is currently the standard diagnostic method for PVV. In the normal pelvic venous system, the right and left common iliac veins empty into the inferior vena cava (IVC) at the level of the lower edge of the L5.3

Typically, inferior vena cava (IVC) is valveless, retroperi-toneal, and on the right of the abdominal aorta. The ascending lumbar vein which joins the paired lumbar veins empties directly into IVC. The longer left ovarian vein (LOV) usually drains into the left renal vein, while the shorter right ovarian vein (ROV) on the other hand empties directly into IVC.5,6

Normally, the LOV forms anastomoses with the inferior mesenteric vein, which in 49% of women drain into the
middle segment of the LOV. However, the ROV is more variable: it anastomosizes with the superior mesenteric vein in 60% of cases, drains into the infrarenal segment of IVC, or in 30% of cases, empties into the confluence of the right renal vein (RRV) and the IVC. In 10% of cases, it drains directly into the intrarenal branches of the right renal vein.7

The first pelvic phlebography ever performed was by Guillaume and Beau in 1954. It showed tortuous and dilated gonadal veins without pelvic congestion syndrome (PCS). The anatomy of the female pelvic venous system differs significantly from person-to-person and is quite complex – especially with respect to the various venous plexuses surrounding the vagina and uterus.8 Whilst the cystic plexus has valves, the uterine and the pampiniform plexuses are valveless.9

In 6–15% of women, the left gonadal veins are valveless. Meanwhile, in 41% of patients, congenital gonadal venous insufficiency occurs on the left and on the right in 46%.10 As a rule, in most patients, the development of PCS is facilitated by both congenital defects of the tributaries of the IVC and the normal anatomical ovarian blood flow.11

Per the results of many studies, 17% of people have abnormal renal veins,12,13 0.5–30% of whom have multiple renal veins (most often of the right side).12 In 26–29% of these people, the renal veins are double, whereas triple renal veins occur in 5–9.7% of them.14 Anomalies of LRV have been found in only 5% of observed patients.12 A good example is a case of four right renal veins with multiple renal arteries.15

Apart from the above-described defects in the development of renal veins, other structural anomalies include circumaortic LRV, which has quite a rare incidence (0.3–5% of cases).16 Observations by N. Drabe et al. (2001) showed that the prevalence of retroaortic LRV was 0.8–7%.17 According to M. Izuimiya et al. (1997), retroaortic LRV develops when the ventral arch of the aortic collar regresses, whilst the dorsal arch persists to form LRV. The ventral portion develops between supracardinal veins, and the posterior portion originates from the supracardinal venous anastomosis.19 The clinical manifestations of the retroaortic position of LRV are impaired venous outflow and development of renal venous congestion, resulting in ovarian varicose veins.20

From a surgical point of view, in order to avoid recurrence of PVV, it is very important to take into account the possibility of 1–2 lower lumbar veins or double gonadal veins draining into the LRV. This anatomical variation occurs in 37–65% of females.19 Moreover, for a successful phlebography, it is necessary to know all the possible anatomical defects of LRV drainage into the IVC. Anomalies of LVR are classified according to:

1. **Number of veins**
   - I) presence of accessory renal veins;
   - II) double left renal vein:
     - a) with drainage of adrenal vein into the upper left renal vein;
     - b) with drainage of adrenal vein into the lower left renal vein;
   - III) incomplete bifurcation of left renal vein;
   - IV) multiple left renal veins.

2. **Retroaortic left renal vein**

3. **Extracaval drainage**
   - I) into the iliac vein;
   - II) into the hemiazygos vein;
   - III) into the iliac and hemiazygos veins (after splitting);
   - IV) into the precaval venous plexus.

4. **Vertebral level**
   - I) high – above the L1;
   - II) low – below L2.

5. **Venous pathologies**
   - I) stenosis;
   - II) presence of valves;
   - III) other.

Furthermore, J.K. Nam et al. (2010) described 4 anatomical variations of drainage of a retroaortic LRV into IVC (Fig. 1).25 Though most of the anatomical variations of pelvic veins have no functional significance, they can pose difficulties during femoral vessels catheterization or pelvic surgeries.21

Fig. 1 – Possible forms of retroaortic left renal vein drainage: type 1 – directly into the inferior vena cava (typical at L1–L2); type 2 – inferiorly into the inferior vena cava; type 3 – circumaortically into the inferior vena cava; type 4 – into the left common iliac vein.

Mack Shin et al. (2014) in a retrospective study to determine the prevalence of congenital defects of iliac veins reviewed the multidetector computed tomography of 2,488 patients. They presented their findings in a schematic drawing (Fig. 2). A normal IVC is right-sided, i.e., it lies to the right of the aorta. A left-sided IVC is a congenital defect whereby IVC is located on the left of aorta with LRV crossing aorta anteriorly to join RRV and the right-sided suprarenal segment of IVC (Fig. 3).22 The prevalence of a left-sided IVC is 0.2–0.5%.23

A double IVC is a rare occurrence, found in 0.6–2.3% of study subjects.24,25 M. Itoh et al. (2001) explained that a double IVC forms when left supracardinal veins persist (Fig. 4).26 Also, aplasia of the hepatic segment of IVC and its communication with the azygos vein is seen in 0.6% of instances.26 This defect of IVC deserves much attention since under unfavourable conditions, (trauma, surgery, or a history of infection) it can be complicated by deep vein thrombosis and chronic venous insufficiency.27
Case report. Right ovarian vein dilatation, most probably due to inferior vena cava transposition

A 27-year-old nulligravid patient presents with dyspareunia, dysmenorrhea, and pain in the right iliac region. She had these symptoms for over 3 years. The pain intensified during physical activities and was refractory to phlebotonics. She is on hormonal replacement therapy (YAZ). Her previous medical history includes autoimmune thyroiditis.

An MRI revealed a left-sided infrarenal segment of IVC, signs of stenosis in the middle and proximal segments of the right common iliac vein, and dilated right ovarian vein (12 mm), ascending lumbar vein (10–12 mm), parametric venous plexus (12 mm), arcuate vein (9 mm). The IVC above its bifurcation was 20 × 16.5 mm (Fig. 4). She was diagnosed with pelvic varicose veins and pelvic congestion syndrome. A phlebography of IVC and pelvic veins by femoral access with a 5F Cobra 1 catheter (Cook Medical, Bloomington, IN, USA) revealed the same findings. A superselective embolization of the right ovarian vein with a Penumbra Coil 35 was performed (Fig. 5).
700 Association of anatomical defects of IVC and gonadal veins with PVV hypertension, which culminates in the development of ovarian varicose veins.\textsuperscript{20,28} Also, anaplasia of the hepatic segment of IVC can cause symptoms of peripheral venous thrombosis and chronic venous insufficiency, which in 50\% of instances are bilateral.\textsuperscript{29}

The high prevalence of multiple right renal veins is associated with the peculiarities of development of the abdominal right and left venous system. Whereas RRV develops from multiple mesonephros veins, LRV originates from left subcardinal and supracardinal venous anastomosis.\textsuperscript{30} Variations of IVC and its tributaries development can have serious clinical effects (from PVV to gonadal and iliac venous thrombosis), and so, should be taken into consideration during interventional procedures. Careful interpretation of results of non-invasive diagnostic methods could help prevent serious mistakes in patient management.

Reference

Conclusion
Therefore, congenital defects of IVC and its tributaries is a risk factor for the development of chronic pelvic venous insufficiency. A retroaortic LRV may present clinically as obstructed venous outflow and congestive renal venous hypertension, which culminates in the development of ovarian varicose veins.\textsuperscript{20,28} Also, anaplasia of the hepatic segment of IVC can cause symptoms of peripheral venous thrombosis and chronic venous insufficiency, which in 50\% of instances are bilateral.\textsuperscript{29}

The high prevalence of multiple right renal veins is associated with the peculiarities of development of the abdominal right and left venous system. Whereas RRV develops from multiple mesonephros veins, LRV originates from left subcardinal and supracardinal venous anastomosis.\textsuperscript{30} Variations of IVC and its tributaries development can have serious clinical effects (from PVV to gonadal and iliac venous thrombosis), and so, should be taken into consideration during interventional procedures. Careful interpretation of results of non-invasive diagnostic methods could help prevent serious mistakes in patient management.
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