

## 286. VARIANTS OF FORMATION OF SUPERFICIAL AND DEEP PALMAR ARCS

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**Introduction:** Our work with title "Desvoltage variants of superficial and deep palmar arcs" was performed at Department Topographic Anatomy and Operative Surgery with scientific coordinator Bedencova O.E.

**The objective:** To study variants of formation of superficial and deep palmar arcs on cadaveric material.

**Materials & Methods:** We used surgical instruments: scalpel, tweezers; 10 corpses at the Department Topographic Anatomy and Operative Surgery; camera.

**Results:** We identified 2 types of superficial arcs (arcus radioulnaris; arcus ulnaris) and 2 types of deep arcs (deep arc formed due anastomosis between the deep palmar branch of the radial artery and the deep branch of the ulnar artery; deep arc formed due anastomosis between the deep palmar branch of the radial artery with the the upper and lower of the deep branch of ulnar artery)

**Conclusion:** We have been revealed the most common desvoltage variant of arcs - arcus radioulnaris, which consists 20%.

## 287. THE GENETIC ASPECTS OF HYPERCHOLESTEROLEMIA

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**Introduction:** Hypercholesterolemia appears when a person swallows a big quantity of cholesterol, or it is synthesised in excess by hepatocytes. A high level of cholesterol increases the cardiovascular risk, and the incidence of myocardial and cerebral stroke. Hypercholesterolemia and overweight are caused by increased saturated fats consumption at persons with genetic idiosyncrasy and are supported by sedentariness. There is a increased production of LDL, and the genetic component is unlikely to be monogenic. The genes which are involved in appearance of hypercholesterolemia are: LDLR gene - protein that encodes LDL receptor on the hepatocytes; APOB gene - is the main component of chylomicrons and LDL, its disorder causes conformational changes of binding with LDL receptor; LDLRAP1 gene – gene that codifies pockets on the hepatocyte's membrane and PCSK9 - controls the number of LDL receptors. The most of all (60-75%) have dose-gene effect. The most common genetic disease that causes hypercholesterolemia is familial hypercholesterolemia, an autosomal dominant