Contribution to the Solution of the Question of Lympho-venous Anastomoses in Heart of Dog

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Summary

The efferent lymphatic vessels of the heart were ligated in 35 dogs and afterwards the systems of lymphatic and blood vessels were filled with various media by injection. Twelve of these dogs were sacrificed 30 min. after ligature and the remaining 23 dogs were left to survive for two to 132 days.

After acute ligature of lymph routes, no lymphatic anastomoses could be found. In the protracted experiment, lympho-lymphatic anastomoses had developed between the interrupted trunks and very thin ones at the sites of adhesions between the epicardium and pericardium by the seventh to 14th day. In five out of 23 dogs, lymphovenous anastomoses of a calibre of 40-60 µm. were found between the seventh and 28th day after ligature. After one month, these LVA could no longer be found. Both forms of anastomoses, i.e. lympho-lymphatic and lymphovenous, may be present at the same time. The mutual relationship between these forms was discussed in the paper.

Most papers on lymphatic vessels of the heart deal with the occurrence of lymphatic vessels in the heart of man and various mammals and the relations between the lymphatic networks of the myocardium, the endocardium and the subepicardial networks (1, 4, 6, 8, 9, 10, 14, 17). The lymphatic vessels of the cardiac valves are subject of number of papers (2, 3, 13, 17). Another group of authors have dealt with the changes in lymphatic vessels under pathological conditions in man and in experiments on animals (4, 5, 10, 13-15, 18-21, 24, 27).

In the last decade attention has been concentrated on the evidence of lymphovenous anastomoses (LVA) in the heart. The opinions as to their existence differ: Some authors deny the presence of LVA under normal conditions (Johnson et Blake, in man, 1968). Another group of authors confirm their presence. Among them are Vajda et al. (1972) who described numerous LVA in the epicardium of man and hog; in hog, Johnson et Blake (1968) also described LVA. A third group of authors found LVA in the heart under altered circulatory conditions: Zerbino et Gavriš (1970) showed that LVA usually existed in obstruction of lymph in the canine heart. Because of the contradictions in these opinions, the authors of this communication have asked themselves the question, as to whether or not and under what conditions do LVA develop in the heart of dog.

Material and Method

In 35 dogs patent blue was injected after thoracotomy carried out under general anaesthesia in order to demonstrate the course of the lymphatic vessels of the heart. Then the main lymphatic trunks in the cardiac atria, the aorta and the truncus pulmonalis (Fig 1) were ligated and severed. The dogs were divided into two groups: one of the acute and the other of the chronic experiment. The animals of the acute group (dogs 1-12) were sacrificed 30 min. after ligation of the lymphatic vessels. Immediately after death, a mixture of India ink and 4% gelatine was injected into the lymphatic vessels demonstrated by patent blue. The lymphatic vessels were filled with a pressure of 30-100 mm H₂O; when pressure rose to more than 200 mm H₂O, damage to lymph vessels and extravasation occurred. Approximately in one half of the heart specimens, this injection was supplemented by an intravenous injection of 6% gelatine containing a pigment dye. In the chronic experiment (dogs 13-35) the animals were left to survive ligation of lymphatic trunks for 2 to 132 days. After certain intervals some of the animals were
penetration of gelatine plus India ink into the venous bed was never observed after direct injection of the mixture into the subepicardial lymphatic vessels. Neither did it happen in the reversed procedure, i.e. after injection of coronary arteries and veins; the mixture did not penetrate into the lymphatic vessels, although the capillary network was well filled. From these findings, the following conclusion may be drawn: In normal heart no lymphovenous anastomoses are present. In direct injection of lymphatic vessels, the injected substance spreads from the apex towards the base of the heart; in retrograde injection, the lymphatic vessels are not filled at all because of the well functioning valves, and if the pressure increases, the wall bursts and extravasation takes place.

In the second group of 23 dogs surviving ligation of the cardiac lymphatic trunks for 2-132 days, the following changes were observed: The subepicardial lymphatic vessels were dilated, varicose and tortuous. By the second or third day even some of the side branches of main trunks were partly filled after retrograde injection. This collateral filling of the lymphatic network was the rule in all cases observed. The shunts between lymphatic trunks and their branches were made visible (between the anterior and posterior left and those of the right truncus). The collateral between the right and left truncus lymphaticus, running across the origin of the truncus pulmonalis (Fig. 1) was obviously thickened. Between the

**Observations**

In the group of acute experiments (12 dogs)
seventh and 14th day lympho-lymphatic anastomoses started to develop between the regions below and above the ligation, first as a network of thin, later of thick vessels (Between the aorta and the truncus pulmonalis (Fig. 2). Slight diffuse permeation of patent blue was also observed from the seventh day after operation in the disrupted adhesions of the pericardium with the epicardium. This also suggested that new lymphatic shunts had developed. Lympho-lymphatic anastomoses linked up with lymphatic vessels running towards the anterior mediastinal and tracheobronchial nodes. In five heart specimens (7, 7, 12, 14, 28 days after ligation) shunts between lymphatic vessels and a vein were observed. Their presence was already signified
in direct injection of lymphatic vessels by the permeation of the injected mass into the vein. In clarified specimens direct anastomoses between lymphatic vessels and veins were found, branching off the lymphatic vessels as an arch opening into a vein running deeper (Figs. 3 and 4). Some of the arched anastomoses split up into smaller branches, approximately 20 µm in calibre, most of them running straight for a distance of 40 to 60 µm. LVA were most frequently found on the anterior aspects of ventricles. Lymphovenous anastomoses appeared in the heart specimens seven to 14 days after ligation of lymphatic trunks, in one case they persisted up to 28 days. In cases, lasting more than one month, lymphovenous anastomoses were no longer found; the number of anastomoses was limited in each specimen (see discussion). At the same time, in three out of the five specimens referred to above, lympho-lymphatic anastomoses were found on the seventh, 14th and 28th day after ligation of lymphatic trunks. Appraisal of all chronic experiments revealed that lympho-lymphatic anastomoses had not developed but in two heart specimens (7th and 12th day) and were only found in cases in which lymphovenous anastomoses had developed.

Discussion

As has been shown by the papers of Johnson et Blake (68) and Vajda et al (72), LVA are found in the normal heart of hog. The opinions of these authors differ in the question as to whether or nor LVA exist in man, the former denied their existence, while Vajda et al. described them as frequent structures, 100-500 and even 700 µm. thick. When comparing the findings in the two papers with those of the present investigation in acutely stenosed lymphatic vessels of the canine heart, it may be said that in the normal heart of dog there are no LVA. Zerbino et Gavris (70) reported that they always found lymphovenous anastomoses in the canine heart after ligation of lymphatic vessels. Evidence of this is also furnished by the retrograde permeation of the injected substance from the veins into the lymphatic bed. Data referring to the duration of development and persistence of LVA have not been reported on, nor has there been made any mention in the paper of the development of lympho-lymphatic anastomoses. It emerges from the experiments of the present investigation that LVA were found in five out of 23 dogs (21,7%) in chronic occlusion of lymph routes. They were found from the seventh day after ligation together with developing lympho-lymphatic anastomoses. No LVA were found by retrograde injection of the lymphatic network from the venous bed. Because of the calibres of LVA it must be assumed that the technical conditions did not permit to determine their exact number. In clarified specimens, approximately six anastomoses were found in the subepicardial network of the ventricles. The authors do not agree with the opinions of Zerbino et Gavris who described LVA in the canine heart as multiple and common structures. Both Zerbino et Gavris and Vajda et al. never reported on the number of anastomoses. When observing the injected specimens, the error arising from the close neighbourhood of lymphatic vessels and veins, crossing each other frequently, should be avoided, and only by meticulous appraisal after dissection from all sides can it be ascertained whether or not these are LVA.

In agreement with other authors (Patek, 1939; Bradham, 1970 and Suschko, 1950) the authors of this communication do not consider evidence of LVA furnished by injection into the interstice as convincing, because puncture and pressure during injection lead to damage to the walls of blood vessels and permit the substance to escape into the veins.

It should be assumed that development of LVA is indirectly dependent on the development of lympho-lymphatic anastomoses. If lympho-lymphatic anastomoses have developed by the seventh day, a substituting drainage of lymph and the interstitial fluid via the LVA is not required and LVA do not develop. If, however, the drainage of lymph via the lympho-lymphatic anastomoses is insufficient, both forms of lymph drainage may exist simultaneously. This is borne out by the cases of the three out of five heart specimens with LVA referred to above. Rusznyak et al. (1957) produced the clinical picture of myocardial infarction in dog by ligation of the cardiac
lymphatic trunks; within five days the pathological signs receded in the ECG. This recession is, according to these authors, due to the development of lympho-lymphatic anastomoses. Vajda et al., on the other hand, explained the normalization of the ECG by the presence of permanent LVA. According to the opinion of the authors of this communication, the picture of myocardial infarction should never develop in the presence of permanent and thick LVA (as described by Vajda et al.) after occlusion of the lymphatic trunks. It may be assumed that development of lympho-lymphatic anastomoses only takes place under possible simultaneous development of lympho-venous anastomoses during the week in which the ECG changes recede.

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