REVIEW

Sixty Years of Heart Research in the Institute of Physiology of the Czech Academy of Sciences

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Summary

In 2023 six decades have elapsed since the first experimental work on the heart muscle was published, in which a member of the Institute of Physiology of the Czech Academy of Sciences participated as an author; Professor Otakar Poupa was the founder and protagonist of this research domain. Sixty years – more than half of the century – is certainly significant enough anniversary that is worth looking back and reflecting on what was achieved during sometimes very complicated periods of life. It represents the history of an entire generation of experimental cardiologists; it is possible to learn from its successes and mistakes. The objective of this review is to succinctly illuminate the scientific trajectory of an experimental cardiological department over a 60-year span, from its inaugural publication to the present. The old truth – *historia magistra vitae* – is still valid.

Key words

Heart • Adaptation • Development • Hypoxia • Protection

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Fig. 1. Participants of the Symposium on Scientific Basis for the Practice of Cardiology", organized by Department of Developmental

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Cardiology in Prague, 2010. Introduction

Cardiovascular diseases represent the most serious health disorders in the contemporary era, accounting for over 50% of total mortality in the developed countries. A singular illness, ischemic heart disease, is responsible for half of this grim statistic. It is, therefore, understandable that the interest of both experimental and clinical cardiologists is driven by the effort to positively influence this unfavourable situation. Moreover, clinicalepidemiological studies have clearly shown that the risk factors of serious cardiovascular diseases manifest already during the early phases of ontogenetic development. Hence, ischemic heart disease and atherosclerosis are thus no longer diseases of the fifth and higher decades of life, necessitating a shift in experimental studies on the pathogenetic mechanisms of these disturbances to the early phases of ontogenetic development. Moreover, sex differences in the sensitivity to ischemia-reperfusion injury are so obvious that they should be taken into consideration in both experimental and clinical cardiology. The importance of the developmental and sex approach for experimental and clinical cardiology is thus indisputable. And, finally, cardiology is the scientific discipline where the close cooperation between the theoretical and clinical cardiologists has the longstanding tradition, spanning from the molecular level to the patient's bed.

The aim of the present survey is to demonstrate briefly the real scientific life of one experimental cardiological department over the 60-year period, from its inaugural experimental work to the present time. It is necessary to emphasize that the program of the cardiovascular research in the Institute of Physiology of the Czech Academy of Sciences was based on all the above presumptions.

Establishment of the Prague school of experimental cardiology

In 2023, six decades have elapsed since the first experimental work on the heart muscle was published, in which a member of the Institute of Physiology participated as an author; he was the founder and protagonist of this research domain, professor Otakar Poupa [1]. This luminary scientist, possessing charisma and renaissance qualities, came with his "Laboratory for Physiology and Pathophysiology of Metabolism" to the Institute of Physiology from the then Institute for Nutrition Research. In a short time, within the newly established department, he founded a cardiological laboratory, which worked closely with the Institute of Pathological Physiology of the Faculty of Pediatrics (today the 2nd Faculty of Medicine of the Charles University), of which he concurrently assumed external leadership. Shortly, he managed to create very unique example of cooperation between the Academy of Sciences and the Faculty of Medicine, both in the field of experimental research and in teaching the medical students. He developed broad-based research on the phylogenetic and ontogenetic development of the heart muscle, with special attention to the needs of clinical cardiology, particularly pediatric cardiology. This orientation attracted a large number of young adepts of science and became the basis of the Prague School of Adaptive and Developmental Cardiology, successfully continuing the Czech tradition in evolutionary medicine, pioneered by Eduard Babák. There were published numerous still cited original results about the development of cardiac muscle during phylogeny and ontogeny [2-8], the cardioprotective effect of adaptation to chronic hypoxia [9,10] and factors influencing the extent of experimental cardiac necrosis, induced by high doses of isoproterenol [11-13]. Studies on increasing cardiac tolerance to oxygen deprivation in animals adapted to chronic hypoxia were the first published experimental results in this field. Poupa's group closely cooperated with the excellent cardioembryologist Zdeněk Rychter, author of the first work on experimental heart defects [14], one of the founders of Czech experimental embryology. In this connection it is necessary to mention, that even a modern cardioprotective phenomenon - preconditioning - has its roots in Poupa's department. Already in 1966, twenty years before the originally recognized discovery of this phenomenon, Poupa and his colleagues published a paper in which they showed that repeated administration of small doses of isoprenaline significantly reduced the extent of heart muscle damage, induced by subsequent high dose of this catecholamine [15]. Unfortunately, this work was published in the local journal, and the described effect was not given a commercially successful name. However, persisting interest of world cardiologists in this originally forgotten publication remains a source of pride for Czech cardiologists. The successes of Poupa's Prague school soon crossed the borders of the country. By the late 1960s, Poupa in collaboration with US scientists Bing and Bajusz conceived the idea to create an international scientific that would bring together experimental society cardiologists from all over the world. This was indeed successful, and the International Society for Heart

Research (ISHR), as it is now called, represents the only world society of its kind. Naturally, Poupa became a member of the editorial board of the official journal of this society, the Journal of Molecular and Cellular Cardiology, which is still one of the prestigious world-renowned periodicals. The fruitful years culminated in the time of the Prague Spring during which Poupa's merits were recognized with a state prize.

Development after 1968

The Soviet occupation in August 1968, which started the so-called normalization era, adversely affected the development of the entire Czech science including experimental cardiology for many years. Poupa, as one of the authors of the memorable manifesto "Two Thousands Words", chose emigration in September 1968, setting in motion the gradual departure of other Prague school members. The significant limitation of foreign contacts gradually led to the interruption of promising development, to professional isolation and a substantial lag behind global scientific advancements.

Experimental cardiologists in the Institute of Physiology (Faltova, Ostadal, Pelouch, Prochazka, later Kolar, Ostadalova, Papousek and Rychter) perpetuated Poupa's legacy, establishing the Department of Developmental Cardiology (further Department). Their main task became the study of the structural, functional and metabolic properties of the developing cardiac muscle, crucial for understanding the ontogenetic changes of cardiac resistance to oxygen deprivation [16-19]. Interestingly, it was observed that fetal spongious musculature persists in children with congenital heart disease [20]. Special attention was paid to the possibilities of protective influence on the myocardium, especially by the adaptation to chronic hypoxia [21-25]. Moreover, for the first time sex difference in cardiac sensitivity to hypoxia was described [26]. This orientation led to the establishment of very effective cooperation both with other laboratories in the Institute of Physiology as well as with cardiologists addressing cardiopulmonary diseases (Institute for Clinical and Experimental Medicine - IKEM Prague, led by Jiří Widimský) [27-32] and clinical department of pediatric cardiology (Children's Cardiac Center Prague-Motol, under the leadership of Milan Samanek) [33-39]. Simultaneously, at the Institute of Physiology, Krecek's department continued the research activities in the field of ontogenesis: developmental aspects of the pathogenesis of hypertension were very

successfully addressed by Jelinek, Kunes and Zicha [40,41]. However, the history of this research would deserve a separate treatise.

Period after November 1989

Only a greatest optimist could assume that the possibility of verifying dreams and reality in the open field of world science is not a utopia. The events of November 1989 ushered in a transformative era, offering Czech science the invaluable gift of global reintegration. First came the altruistic invitations from foreign colleagues but at the same time become clear that excuses for past unhappy years would not be enough. Of course, all of this also applies to experimental cardiology. Where possible, old contacts were established, long-standing literary acquaintances were personified, and seminal results timidly found their way to scientific meetings and prestigious journals. The explosion of foreign sojourns of young researchers began to bear fruit, and the intellectual and methodological background gradually improved. In this context, it should be stressed that molecular cardiology, without which we can no longer imagine current experimental research, began to be developed with great delay. The grant system established a competitive approach to financial resources and undoubtedly contributed to the improvement of the scientific quality.

The advantage of our experimental cardiology was that it entered the last 30 years organizationally prepared. The Committee of Experimental Cardiology (KEK), with the foundation of which (1973 by Braveny and Ostadal) the cardiologists from the Institute of Physiology are intrinsically connected, represented a unique national scientific society, even on a world scale, with its philosophy, organization and scientific activity. This strategic advantage prompted ISHR to entrust KEK with organizing the 1995 World Congress in Prague. The president was Pavel Braveny from Brno, the secretary general Ostadal and the entire team of the Department participated in the organization. With the passage of time, it can be emphasized that the first meeting of ISHR organized east of Alps, with more than 1200 participants, became a real culmination of the KEK's activities to date. It was also possible to present adequately the traditional issues of the Institute of Physiology, i.e. cardiac development and adaptation.

The scientific research of the Department during this period was concentrated on the question, how to increase cardiac tolerance to oxygen deficiency. Focused investigation probed the molecular and cellular mechanisms involved in the protection of the ischemic myocardium [42-56] and the analyses of the cardiac resistance during early phases of ontogenetic development [57-62]. Other studies have investigated e.g. the effect of increased pressure on neonatal heart growth [63-65] or right ventricular function in hypoxic pulmonary hypertension [43, 66-68]. The Department seized opportunities offered both for substantional improvements of methodical equipment as well as for significant expansion of contacts with the top foreign laboratories. Very fruitful was the intensive cooperation with the groups at University of Manitoba, Winnipeg [69-71], University of Ottawa [72-78], Max-Planck Institute in Bad Nauheim [79], Free University Berlin [80-82], Catholic University of Louvain in Brussels [83-85], University of Strathclyde, Glasgow [86], INSERM Paris [87,88], Institute for Heart Research, Bratislava [89,90], and others. The representatives of the Department became members of the committees of the international scientific societies, such as ISHR and the International Academy of Cardiovascular International activities of the Department Sciences. continued in the organization of several scientific meetings, such as Czech-French-Slovak Symposium on Basic Cardiology (1994), The Developing Heart (2000), Mendel symposium I on Genes and the Heart (2003), Mendel Symposium II (2008) and Symposium on Scientific Basis for the Practice of Cardiology (2010). The interest of many pre- and post-graduate students, both from the Faculty of Science and from the Faculty of Medicine was gratifying. Intensive cooperation has developed with domestic experimental laboratories of the Faculty of Science or the 2nd Faculty of Medicine and continued successful collaboration with the clinical institutions such as Children's Cardiac Center and IKEM.

Research Center for Cardiovascular Diseases

In 1999, the first representative research centers were established as part of a project of the Ministry of Education, Youth and Sports. An informal group of cardiology-oriented, internationally experienced and freely cooperating laboratories of the Institute of Physiology of the Czech Academy of Sciences (Department of Developmental Cardiology, Department of Hypertension), of the 2nd Faculty of Medicine (Departments of Physiology and Pathophysiology) and Center for Experimental Medicine of IKEM, seemed to be a suitable model for this purpose. "Center for

Cardiovascular Research" was acknowledged among the first, with Ostadal as the responsible researcher. The research concept of this Center aligned with cardiovascular research priorities in the European Union. The aim of the research activity was to clarify some of the molecular and cellular mechanisms involved in the development of ischemic heart disease and main risk factors, such as atherosclerosis and high blood pressure. The orientation on developmental approach in cardiovascular diseases was based on long-standing traditions of Czech cardiac research. At the end of the fiveyear period the evaluating council stated that the Center represents a research base that has no parallels in our country, leading to a resounding endorsement for another seven years. New competition rules included the close cooperation with clinical research facilities, Children's Cardiac Center in Prague-Motol and Departments of Cardiology and Cardiac Surgery of IKEM. The theoretical part was further significantly strengthened by a group from the Faculty of Science and two further departments of the Institute of Physiology, the newly established Department of Cardiac Morphogenesis and the Department of Biomaterials and Tissue Engineering. The Center gradually ceased to be a formal grouping of individual workplaces but became a virtual institute. New laboratories were created and methodical approaches and technical equipment increased significantly. The main output of the Center's activities were high-quality original results published in renowned journals. The Center was very successful in attracting young researchers: a number of future cardiologists and cardiac surgeons completed their scientific training in the experimental and clinical research. The number of full-time researchers was in average 109, 30 of them were under 35 years of age. During existence of the Center, 25 postgraduate students defended their PhD thesis. Regrettably, the dissolution of this long-established team of experimental and clinical researchers interrupted the well-started and effective cooperation; the successful system of scientific centers officially terminated in 2012.

Cardiovascular research in the last decade

Although the cessation of the Centre's activities led to the end of joint funding, the scientific cooperation of the Department with several partners, in particular the Faculty of Science, continued and developed further on a bilateral basis. In the framework of this collaboration, a number of studies have recently been published that have

further characterized the differences in cardioprotective mechanisms induced by various regimens of continuous and intermittent chronic hypoxia [91-97]. Together with the Center for Experimental Medicine of IKEM, we have also addressed questions concerning the influence of comorbidities, especially various forms of systemic hypertension, on myocardial ischemic tolerance and new possibilities of therapeutic pharmacological interventions [98-102]. The merger with the Department of Cardiac Morphogenesis has led to the expansion of the studied topics to include the development of the structure and function of the cardiac conduction system and to the enrichment of the methodological tools [103-106]. Special attention was paid to the developmental and sex differences in cardiac tolerance to ischemia/reperfusion injury and the possible role of mitochondria in this process [107-116]. A newly established international collaboration with the Medical College of Wisconsin, Milwaukee, has yielded new findings with translational potential on the beneficial effects of eicosanoids on cardiac injury by ischemia and on the development of post-ischemic heart failure [117-120]. The traditional focus of the Department's research programme on the cardiovascular effects of chronic hypoxia has prompted joint projects with the Laboratory of Molecular Pathogenetics, Institute of Biotechnology, investigating the role of the transcription factor HIF-1a in the mechanism of ischemic tolerance and in the pathogenesis of diabetic cardiomyopathy and heart failure [121-124]. Last but not least, we are involved with colleagues from Comenius University in Bratislava in research on mechanisms of new forms of cell death in myocardial infarction and heart failure [125-127]. Recently, the Department (newly Laboratory) has also shown promising developments in advanced molecular biology methods and their use in the study of epigenetic RNA regulatory mechanisms involved in the ontogenetic development of the heart and in the pathogenesis of heart disease [128-131]. Number of young enthusiastic researchers, modern methodologies and attractive scientific programme represent promising perspectives for future.

The scientific contribution of experimental cardiologists from the Institute of Physiology

To evaluate the contribution of scientific work is always tricky; it depends on many points of view. On the first place is of course the originality of the published results, but it is also necessary to asses to whom the results will help, what are the perspectives of their further use and how they have been accepted by the world scientific community, i.e. how often they were cited. The harshest critic is time, which will test the results and show "evergreen" ones. For the sake of completeness, we would like to summarize at least some of them.

- studies on the normal phylogenetic and ontogenetic development of the cardiac muscle, myocardial blood supply and conduction system;
- persistence of the fetal avascular spongious myocardium, supplied by diffusion from the ventricular cavity in patients with pulmonary stenosis;
- cardiac adaptation to pressure overload during early postnatal development;
- the first observation in experimental cardiology, demonstrating that the female heart is more tolerant to hypoxia than the male heart;
- developmental and sex differences in cardiac tolerance to oxygen deprivation; the role of mitochondria, and protective strategies for the immature myocardium;
- metabolic adaptation to chronic hypoxia in children with congenital heart disease;
- intermittent chronic hypoxia-induced right ventricular hypertrophy and pulmonary hypertension; possibilities of pharmacological interventions;
- studies on the effect of perinatal hypoxia on the sex-dependent hypoxic tolerance of the adult myocardium;
- molecular mechanisms of the long-lasting cardioprotective effect of adaptation to chronic hypoxia, regular exercise and other adaptive interventions;
- altered cardiac ischemic tolerance associated with various forms of systemic hypertension; possibilities of genetic and pharmacological interventions;
- pathogenetic mechanisms of diabetic cardiomyopathy; ischemic tolerance of diabetic heart;
- progression of post-ischemic heart failure in hypertensive animals; novel experimental therapy;
- role of epitranscriptomic regulatory mechanisms in heart physiology and pathophysiology.

Conclusion

60 years – more than half of the century – is certainly significant enough anniversary that it is worth looking back and reflecting on what has been achieved. It represents the history of an entire generation of experimental cardiologists; it is possible to learn from its successes and mistakes; and that was the main purpose of this historical reflection. Indeed, the old truth – *historia magister vitae* – is still valid.

Conflict of Interest

There is no conflict of interest.

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Fig. 2. Participants of the "Mendel symposium II: Genes and the Heart", organized by Department of Developmental Cardiology in Liblice, 2008.

References

- 1. Rakusan K, Poupa O. Changes in the diffusion distance in the rat heart muscle during development. Physiol Bohemoslov 1963;12:220-227.
- 2. Rakusan K, Poupa O. Development postnatal des vaisseaux terminaux et des fibres musculaires dans le coeur du rat. J Physiol Paris 1964;56:636.
- 3. Rakusan K, Jelinek J, Korecky B, Soukupova M, Poupa O. Postnatal development of muscle fibres and capillaries in the rat heart. Physiol Bohemoslov 1965;14:32-37.
- 4. Rakusan K, Poupa O. Developmental changes in the protein composition of heart muscle in the rat. Physiol Bohemoslov 1966;15:132-136.
- 5. Dbaly J, Ostadal B, Rychter Z. Development of the coronary arteries in rat embryos. Acta anat (Basel) 1968;71:209-222. <u>https://doi.org/10.1159/000143186</u>

- Poupa O, Ostadal B. Experimental cardiomegalies and cardiomegalies"- in free-living animals. Ann NY Acad Sci 1969;156:445-468. <u>https://doi.org/10.1111/j.1749-6632.1969.tb16744.x</u>
- Poupa O, Rakusan K, Ostadal B: The effect of physical activity upon the heart of vertebrates. In: Medicine and Sport. Physical Activity and Aging. D BRUNNER, E JOKL (eds). Karger, New York, Basel, 1970, pp 202-233. <u>https://doi.org/10.1159/000387290</u>
- 8. Ostadal B, Rychter Z, Poupa O. Comparative aspects of the development of the terminal vascular bed in the myocardium. Physiol Bohemoslov 1970;19:1-7.
- 9. Poupa O, Rakusan K, Prochazka J, Krofta K. Resistance of the myocardium to hypoxia induced by anaemia of different types. Physiol Bohemoslov 1965;14:452-455.
- 10. Poupa O, Turek Z, Kalus M, Krofta K. Acute cardiac infarct like necrosis in high altitude adaptes rats. Physiol Bohemoslov 1965;14:542-545.
- 11. Poupa O, Turek Z, Pelouch V, Prochazka J, Krofta K. Increased resistance of the myocardium to anoxia in vitro after repeated application of isoprenalin. Physiol Bohemoslov 1965;14:536-541.
- 12. Poupa O, Prochazka J, Pelouch V. Effect of catecholamines on resistance of the myocardium to anoxia and on the heart glycogen concentration. Physiol Bohemoslov 1968;17:36-42.
- Ostadal B, Rychterova V, Poupa O. Isoproterenol-induced acute experimental cardiac necrosis in the turtle (Testudo Horsfieldi). Amer Heart J 1968;76:645-649. <u>https://doi.org/10.1016/0002-8703(68)90163-4</u>
- Rychter Z. Experimental morphology of the aortic arches and the heart loop in chick embryos. In: Advances in morphogenesis. Academic Press, New York/London, 1962, pp 333-371. <u>https://doi.org/10.1016/B978-1-4831-9949-8.50012-2</u>
- 15. Turek Z, Kalus M, Poupa O. The effect of isoprenaline pretreatment on the size of acute myocardial necrosis induced by the same drug. Physiol Bohemoslov 1966;15:353-356.
- Ostadal B, Schiebler TH. Kapillarentwicklung in Rattenherzen. Elektronenmikroskopische Untersuchungen. Z Anat Entwickl-Gesch 1971;133:288-304. <u>https://doi.org/10.1007/BF00519304</u>
- Bass A, Ostadal B, Pelouch V, Vitek V. Differences in weight parameters, myosin ATPase activity and the enzyme pattern of energy supplying metabolism between the compact and spongious cardiac musculature of carp (Cyprinus carpio) and turtle (Testudo Horsfieldi). Pflügers Arch 1973;343:65-77. <u>https://doi.org/10.1007/BF00586575</u>
- Ostadal B, Schiebler TH, Rychter Z: Relations between development of the capillary wall and myoarchitecture of the rat heart. In: Cell Impairment in Aging and Development. VJ CRISTOFALO, E HOLECKOVA (eds). Adv Exp Med Biol 1975;53:375-388. <u>https://doi.org/10.1007/978-1-4757-0731-1_33</u>
- Ostadal B, Rychter Z, Rychterova V. The action of isoproterenol on the chick embryo heart. J Mol Cell Cardiol 1976;8:533-544. <u>https://doi.org/10.1016/0022-2828(76)90054-7</u>
- 20. Dusek J, Ostadal B, Duskova M. Postnatal persistence of spongy myocardium with embryonic blood supply. Arch Path 1975; 99:312-317.
- McGrath J, Prochazka J, Pelouch V, Ostadal B. Physiological responses of rats to intermittent high-altitude stress: effects of age. J Appl Physiol 1973;34:289-293. <u>https://doi.org/10.1152/jappl.1973.34.3.289</u>
- 22. McGrath J, Ostadal B, Prochazka J, Wachtlova M, Rychterova V. Experimental cardiac necrosis in hypobaric and anemic hypoxia. J Appl Physiol 1975;39:205-208. <u>https://doi.org/10.1152/jappl.1975.39.2.205</u>
- Wachtlova M, Mares V, Ostadal B. DNA synthesis in the ventricular myocardium of young rats exposed to intermittent high altitude (IHA) hypoxia. An autoradiographic study. Virchows Arch B Cell Path 1977;24:335-342. https://doi.org/10.1007/BF02889289
- Ostadal B, Mirejovska E, Hurych J, Pelouch V, Procházka J. Effect of intermittent high altitude hypoxia on the synthesis of collagenous and non-collagenous proteins of the right and left ventricular myocardium. Cardiovasc Res 1978;12:303-308. <u>https://doi.org/10.1093/cvr/12.5.303</u>
- Jarkovska D, Ostadal B. Intermittent high altitude hypoxia-induced structural changes of the pulmonary myocardium in young mice. Virchows Arch (Cell Pathol) 1983;43:327-336. <u>https://doi.org/10.1007/BF02932965</u>
- 26. Ostadal B, Prochazka J, Pelouch V, Urbanova D, Widimsky J. Comparison of cardiopulmonary responses of male and female rats to intermittent high altitude hypoxia. Physiol Bohemoslov 1984;33:129-138.

- 27. Widimsky J, Urbanova D, Ressl J, Ostadal B, Pelouch V, Prochazka J. The effect of intermittent altitude hypoxia on the myocardium and lesser circulation in rat. Cardiovasc Res 1973;7:798-808. https://doi.org/10.1093/cvr/7.6.798
- 28. Widimsky J, Urbanova D, Ressl J, Ostadal B, Pelouch V, Prochazka J: Chronic pulmonary hypertension induced by intermittent high altitude exposure. In: Pulmonary Circulation. J WIDIMSKY, S KARGER (eds), Prog Resp Res, Basel, New York, 1975;9:121-125.
- Widimsky J, Ostadal B, Urbanova D, Ressl J, Prochazka J, Pelouch V. Intermittent high altitude hypoxia. Chest 1980;77:383-389. <u>https://doi.org/10.1378/chest.77.3.383</u>
- 30. Pelouch V, Ostadal B, Urbanova D, Prochazka J, Ressl J, Widimsky J. The effect of intermittent high altitude hypoxia on the structure and enzymatic activity of cardiac myosin. Physiol bohemoslov 1980;29:313-322.
- Pelouch V, Ostadal B, Prochazka J, Urbanova D, Widimsky J. Effect of high altitude hypoxia on the protein composition on the right ventricular myocardium. Progr Resp Res 1985;20:41-48. <u>https://doi.org/10.1159/000410421</u>
- 32. Kolar F, Ostadal B, Prochazka J, Pelouch V, Widimsky J. Comparison of cardiopulmonary response to intermittent high-altitude hypoxia in young and adult rats. Respiration 1989;56:57-62. <u>https://doi.org/10.1159/000195778</u>
- Skovranek J, Ostadal B, Pelouch V, Prochazka J. Ontogenetic differences in cardiac sensitivity to verapamil in rats. Pediatr Cardiol 1986;7:25-29. <u>https://doi.org/10.1007/BF02315478</u>
- Bass A, Samanek M, Ostadal B, Hucin B, Stejskalova M, Pelouch V. Differences between atrial and ventricular energy supplying enzymes in children. J Appl Cardiol 1988;3:397-405.
- Samanek M, Bass A, Ostadal B, Hucin B, Stejskalova M. Effect of hypoxaemia on enzymes supplying myocardial energy in children with congenital heart disease. Int J Cardiol 1989;25:265-270. <u>https://doi.org/10.1016/0167-5273(89)90216-7</u>
- 36. Samanek M, Bass A, Ostadal B, Hucin B: The importance of the energy-supplying metabolism of the heart in preand postoperative care of patiens with tetralogy of Fallot. In: *Perspectives in Paediatric Cardiology*. GK SRUPI, L PARENZANI, RH ANDERSON (eds), Future Publ Comp, Inc, Mount Kisco, New York, 1989, pp 152-154.
- Jarkovska D, Janatova T, Hruda J, Ostadal B, Samanek M. The physiological closure of ductus arteriosus in the rat. An ultrastructural study. Anat Embryol 1989;180:497-504. <u>https://doi.org/10.1007/BF00305125</u>
- Janatova T, Jarkovska D, Hruda J, Samanek M, Ostadal B. Effect of the administration of prostaglandins (PGE2) in the early postnatal period on closure of the ductus arteriosus in the laboratory rat. Physiol Bohemoslov 1989;38:201-206.
- Pelouch V, Milerova M, Ostadal B, Hucin B, Samanek M. Differences between atrial and ventricular protein profiling in children with congenital heart disease. Mol Cell Biochem 1995;147:43-49. <u>https://doi.org/10.1007/BF00944782</u>
- 40. Zicha J, Kunes J, Jelinek J. Experimental hypertension in young and adult animals. Hypertension 1986;8:1096-1104. https://doi.org/10.1161/01.HYP.8.12.1096
- Zicha J, Kunes J. Ontogenetic aspects of hypertension development: analysis in the rat. Physiol Rev 1999;79:1227-1282. <u>https://doi.org/10.1152/physrev.1999.79.4.1227</u>
- 42. Kolar F, Cole WC, Ostadal B, Dhalla NS. Transient inotropic effects of low extracellular sodium in perfused rat heart. Am J Physiol Heart Circ Physiol 1990;259:H712-H719. <u>https://doi.org/10.1152/ajpheart.1990.259.3.H712</u>
- 43. Kolar F, Ostadal B. Right ventricular function in rats with hypoxic pulmonary hypertension. Pflügers Arch 1991;419:121-126. <u>https://doi.org/10.1007/BF00372996</u>
- Asemu G, Papousek F, Ostadal B, Kolar F. Adaptation to high altitude hypoxia protects the rat heart against ischemia-induced arrhythmias. Involvement of mitochondrial KATP channel. J Mol Cell Cardiol 1999;31:1821-1831. <u>https://doi.org/10.1006/jmcc.1999.1013</u>
- 45. Szarszoi O, Asemu G, Vanecek J, Ostadal B, Kolar F. Effects of melatonin on ischemia and reperfusion injury of the rat heart. Cardiovasc Drugs Ther 2001;15:251-257. <u>https://doi.org/10.1023/A:1011920407691</u>
- 46. Neckar J, Papousek F, Novakova O, Ostadal B, Kolar F. Cardioprotective effects of chronic hypoxia and ischaemic preconditioning are not additive. Basic Res Cardiol 2002; 97:161-167. <u>https://doi.org/10.1007/s003950200007</u>

- Neckar J, Szarszoi O, Koten L, Papousek F, Ostadal B, Grover GJ, Kolar F. Effects of mitochondrial KATP modulators on cardioprotection induced by chronic high altitude hypoxia in rats. Cardiovasc Res 2002;55:567-575. <u>https://doi.org/10.1016/S0008-6363(02)00456-X</u>
- Kolar F, Ostadal B. Molecular mechanisms of cardiac protection by adaptation to chronic hypoxia. Physiol Res 2004;53:S3-S13. <u>https://doi.org/10.33549/physiolres.930000.53.S3</u>
- Neckar J, Markova I, Novak F, Novakova O, Szarszoi O, Ostadal B, Kolar F. Increased expression and altered subcellular distribution of PKC-delta in chronically hypoxic rat myocardium: involvement in cardioprotection. Am J Physiol Heart Circ Physiol 2005;288:H1566-H1572. <u>https://doi.org/10.1152/ajpheart.00586.2004</u>
- 50. Kolar F, Neckar J, Ostadal B. MCC-134, a blocker of mitochondrial and opener of sarcolemmal ATP-sensitive K+ channels, abrogates cardioprotective effects of chronic hypoxia. Physiol Res 2005;54:467-471. https://doi.org/10.33549/physiolres.930000.54.467
- Fitzpatrick CM, Shi Y, Hutchins WC, Su j, Gross GJ, Ostadal B, Tweddell JS, Baker JE. Cardioprotection in chronically hypoxic rabbits persists upon exposure to normoxia: Role of nitric oxide synthase and KATP channels. Am J Physiol Heart Circ Physiol 2005;288:H62-H68. <u>https://doi.org/10.1152/ajpheart.00701.2004</u>
- Kolar F, Jezkova J, Balkova P, Breh J, Neckar J, Novak F, Novakova O, Tomasova H, Srbova M, Ostadal B, Wilhelm J, Herget J. Role of oxidative stress in PKC-δ upregulation and cardioprotection induced by chronic intermittent hypoxia. Am J Physiol Heart Circ Physiol 2007;292:H224-H230. https://doi.org/10.1152/ajpheart.00689.2006
- 53. Hlavackova M, Neckar J, Jezkova J, Balkova P, Stankova B, Novakova O, Kolar F, Novak F. Dietary polyunsaturated fatty acids alter myocardial protein kinase C expression and affect cardioprotection induced by chronic hypoxia. Exp Biol Med 2007;232:823-832.
- 54. Hlavackova M, Kozichova K, Neckar J, Kolar F, Musters RJP, Novak F, Novakova O. Up-regulation and redistribution of protein kinase C-δ in chronically hypoxic heart. Mol Cell Biochem 2010;345:271-282. https://doi.org/10.1007/s11010-010-0581-8
- Borchert GH, Yang C-T, Kolar F. Mitochondrial BKCa channels contribute to protection of cardiomyocytes isolated from chronically hypoxic rats. Am J Physiol Heart Circ Physiol 2011; 300:H507-H513. https://doi.org/10.1152/ajpheart.00594.2010
- Snorek M, Hodyc D, Sedivy V, Durisova J, Skoumalova A, Wilhelm J, Neckar J, Kolar F, Herget J. Short-term fasting reduces the extent of myocardial infarction and incidence of reperfusion arrhythmias in rats. Physiol Res 2012;61:567-574. <u>https://doi.org/10.33549/physiolres.932338</u>
- 57. Kolar F, Seppet EK, Vetter R, Prochazka J, Grünermel J, Zilmer K, Ostadal B. Thyroid control of contractile function and calcium handling in neonatal rat heart. Pflügers Arch 1992;421:26-31. https://doi.org/10.1007/BF00374729
- Ostadalova I, Kolar F, Ostadal B, Rohlicek V, Rohlicek J, Prochazka J. Early postnatal development of contractile performance and responsiveness to Ca2+, verapamil and ryanodine in the isolated rat heart. J Mol Cell Cardiol 1993;25:733-740. <u>https://doi.org/10.1006/jmcc.1993.1085</u>
- Ostadalova I, Kolar F, Ostadal B. Inotropic effect of low extracellular sodium on perfused perinatal rat heart. Can J Physiol Pharmacol 1995;73:50-54. <u>https://doi.org/10.1139/y95-007</u>
- Kolar F, Papousek F, Pelouch V, Ostadal B, Rakusan K. Pressure overload induced in newborn rats: effects on left ventricular growth, morphology, and function. Pediatr Res 1998;43:521-526. <u>https://doi.org/10.1203/00006450-199804000-00014</u>
- 61. Ostadalova I, Ostadal B, Jarkovska D, Kolar F. Ischemic preconditioning in chronically hypoxic neonatal rat heart. Pediatr Res 2002;52:561-567. <u>https://doi.org/10.1203/00006450-200210000-00016</u>,
- 62. Rakusan K, Chvojkova Z, Oliviero P, Ostadalova I, Kolar F, Chassagne C, Samuel J-L, Ostadal B. ANG II type 1 receptor antagonist irbesartan inhibits coronary angiogenesis stimulated by chronic intermittent hypoxia in neonatal rats. Am J Physiol Heart Circ Physiol 2007;292:H1237-H1244. <u>https://doi.org/10.1152/ajpheart.00965.2006</u>
- Kolar F, Papousek F, Pelouch V, Ostadal B, Rakusan K. Pressure overload induced in newborn rats: Effects on left ventricular growth, morphology, and function. Pediatr Res 1998;43:521-526. <u>https://doi.org/10.1203/00006450-199804000-00014</u>

- 64. Sedmera D, Thompson RP, Kolar F. Effect of increased pressure loading on heart growth in neonatal rats. J Mol Cell Cardiol 2003;35:301-309. <u>https://doi.org/10.1016/S0022-2828(03)00011-7</u>
- 65. Novotny J, Hrbasova M, Kolar F, Svoboda P. Cardiomegaly induced by pressure overload in newborn rats is accompanied by altered expression of the long isoform of Gsα protein and deranged signaling of adenylyl cyclase. Mol Cell Biochem 2003;245:157-166. <u>https://doi.org/10.1023/A:1022828430565</u>
- Cihak R, Kolar F, Pelouch V, Prochazka J, Ostadal B, Widimsky J. Functional changes in the right and left ventricle during development of cardiac hypertrophy and after its regression. Cardiovasc Res 1992;26:845-850. <u>https://doi.org/10.1093/cvr/26.9.845</u>
- 67. Pelouch V, Kolar F, Ostadal B, Milerova M, Cihak R, Widimsky J. Regression of chronic hypoxia-induced pulmonary hypertension, right ventricular hypertrophy and fibrosis: Effect of enalapril. Cardiovasc Drugs Ther 1997;11:177-185. <u>https://doi.org/10.1023/A:1007788915732</u>
- Rivolta I, Lucchini V, Rocchetti M, Kolar F, Palazzo F, Zaza A, Miserocchi G. Interstitial pressure and lung oedema in chronic hypoxia. Eur Respir J 2011;37:943-949. <u>https://doi.org/10.1183/09031936.00066710</u>
- 69. Dhalla NS, Yates JC, Naimark B, Dhala KS, Beamish RE, Ostadal B: Cardiotoxicity of catecholamines and related agents. In: Cardiovascular Toxicology. D ACOSTA (ed), Raven Press, New York, 1992, pp 239-282.
- 70. Lee S L, Ostadalova I, Kolar F, Dhalla NS. Alterations in Ca2+-channels during the development of diabetic cardiomyopathy. Mol Cell Biochem 1992;109:173-179. <u>https://doi.org/10.1007/BF00229773</u>
- Ostadal B, Ostadalova I, Dhalla NS. Development of cardiac sensitivity to oxygen deficiency: comparative and ontogenetic aspects. Physiol Rev 1999;79:635-659. <u>https://doi.org/10.1152/physrev.1999.79.3.635</u>
- 72. Kolar F, MacNaughton C, Papousek F, Korecky B. Systolic mechanical performance of heterotopically transplanted hearts in rats treated with cyclosporin. Cardiovasc Res 1993;27:1244-1247. <u>https://doi.org/10.1093/cvr/27.7.1244</u>
- 73. Kolar F, MacNaughton C, Papousek F, Korecky B, Rakusan K. Changes in calcium handling in atrophic heterotopically isotransplanted rat hearts. Basic Res Cardiol 1995;90:475-481. <u>https://doi.org/10.1007/BF00788540</u>
- 74. Kolar F, Papousek F, MacNaughton C, Pelouch V, Milerova M, Korecky B. Myocardial fibrosis and right ventricular function of heterotopically transplanted hearts in rats treated with cyclosporin. Mol Cell Biochem 1996;163/164:253-260. <u>https://doi.org/10.1007/BF00408666</u>
- 75. Sladek T, Sladkova J, Kolar F, Papousek F, Cicutti N, Korecky B, Rakusan K. The effect of AT1 receptor antagonist on chronic cardiac response to coronary artery ligation in rats. Cardiovasc Res 1996;31:568-576. <u>https://doi.org/10.1016/0008-6363(95)00244-8</u>
- 76. Heron MI, Kolar F, Papousek F, Rakusan K. Early and late effect of neonatal hypo- and hyperthyroidism on coronary capillary geometry and long-term heart function in rat. Cardiovasc Res 1997; 33:230-240. <u>https://doi.org/10.1016/S0008-6363(96)00198-8</u>
- 77. Rakusan K, Cicutti N, Kolar F. Effect of anemia on cardiac function, microvascular structure, and capillary hematocrit in rat hearts. Am J Physiol Heart Circ Physiol 2001; 280:H1407-H1414. https://doi.org/10.1152/ajpheart.2001.280.3.H1407
- Rakusan K, Cicutti N, Kolar F. Cardiac function, microvascular structure and capillary hematocrit in hearts of polycythemic rats. Am J Physiol Heart Circ Physiol 2001; 281:H2425-H2431. <u>https://doi.org/10.1152/ajpheart.2001.281.6.H2425</u>
- 79. Deindl E, Kolar F, Neubauer E, Vogel S, Schaper W, Ostadal B. Effect of intermittent high altitude hypoxia on gene expression in rat heart and lung. Physiol Res 2003;52:147-157. <u>https://doi.org/10.33549/physiolres.930295</u>
- 80. Kolar F, Seppet E K, Vetter R, Prochazka J, Grünermel J, Zilmer K, Ostadal B. Thyroid control of contractile function and calcium handling in neonatal rat heart. Pflügers Arch 1992;421:26-31. https://doi.org/10.1007/BF00374729
- Vetter R, Studer R, Reinecke H, Kolar F, Ostadalova I, Drexler H. Reciprocal changes in the postnatal expression of the sarcolemmal Na+-Ca2+-exchanger and SERCA2 in rat heart. J Mol Cell Cardiol 1995;27:1689-1701. <u>https://doi.org/10.1016/S0022-2828(95)90788-2</u>
- Cernohorsky J, Kolar F, Pelouch V, Korecky B, Vetter R. Thyroid control of sarcolemmal Na⁺-Ca²⁺ exchanger and SR Ca²⁺-ATPase in developing rat heart. Am J Physiol Heart Circ Physiol 1998;275:H264-H273. https://doi.org/10.1152/ajpheart.1998.275.1.H264

- Wibo M, Kolar F, Zheng L, Godfraind T. Influence of thyroid status on postnatal maturation of calcium channels, beta-adrenoceptors and cation transport ATPases in rat ventricular tissue. J Mol Cell Cardiol 1995;27:1731-1743. <u>https://doi.org/10.1016/S0022-2828(95)90887-0</u>
- Zheng L, Wibo M, Kolar F, Godfraind T. Calcium channels and cation transport ATPases in cardiac hypertrophy induced by aortic constriction in newborn rats. Mol Cell Biochem 1996;163/164:23-29. https://doi.org/10.1007/BF00408637
- Wibo M, Feron O, Zheng L, Maleki M, Kolar F, Godfraind T. Thyroid status and postnatal changes in subsarcolemmal distribution and isoform expression of rat cardiac dihydropyridine receptors. Cardiovasc Res 1998;37:151-159. <u>https://doi.org/10.1016/S0008-6363(97)00228-9</u>
- Ostadalova I, Ostadal B, Kolar F, Parratt JR, Wilson S. Tolerance to ischaemia and ischaemic preconditioning in neonatal rat heart. J Mol Cell Cardiol 1998;30:857-865. <u>https://doi.org/10.1006/jmcc.1998.0653</u>
- Oliviero P, Chassagne C, Kolar F, Adamy C, Marotte F, Samuel J-L, Rappaport L, Ostadal B. Effect of pressure overload on angiotensin receptor expression in the rat heart during early postnatal life. J Mol Cell Cardiol 2000;32:1631-1645. <u>https://doi.org/10.1006/jmcc.2000.1198</u>
- Ratajczak P, Oliviero P, Marotte F, Kolar F, Ostadal B, Samuel J-L. Expression and localization of caveolins during postnatal development in rat heart: implication of thyroid hormone. J Appl Physiol 2005;99:244-251. https://doi.org/10.1152/japplphysiol.01292.2004
- Ravingerova T, Neckar J, Kolar F, Stetka R, Volkovova K, Ziegelhöffer A, Styk J. Ventricular arrhythmias following coronary artery occlusion in rats: is the diabetic heart less or more sensitive to ischaemia? Basic Res Cardiol 2001;96:160-168. <u>https://doi.org/10.1007/s003950170066</u>
- Ravingerova T, Neckar J, Kolar F. Ischemic tolerance of rat hearts in acute and chronic phases of experimental diabetes. Mol Cell Biochem 2003;249:167-174. <u>https://doi.org/10.1023/A:1024751109196</u>
- 91. Hrbasova M, Novotny J, Hejnova L, Kolar F, Neckar J, Svoboda P. Altered myocardial Gs protein and adenylyl cyclase signaling in rats exposed to chronic hypoxia and normoxic recovery. J Appl Physiol 2003;94:2423-2432. https://doi.org/10.1152/japplphysiol.00958.2002
- 92. Neckar J, Borchert GH, Hlouskova P, Micova P, Novakova O, Novak F, Hroch M, Papousek F, Ostadal B, Kolar F. Brief daily episode of normoxia inhibits cardioprotection conferred by chronic continuous hypoxia. Role of oxidative stress and BKCa channels. Curr Pharm Des 2013;19:6880-6889. https://doi.org/10.2174/138161281939131127115154
- Waskova-Arnostova P, Kasparova D, Elsnicova B, Novotny J, Neckar J, Kolar F, Zurmanova J. Chronic hypoxia enhances expression and activity of mitochondrial creatine kinase and hexokinase in the rat ventricular myocardium. Cell Physiol Biochem 2014;33:310-320. <u>https://doi.org/10.1159/000356671</u>
- 94. Chytilova A, Borchert GH, Mandikova-Alanova P, Hlavackova M, Kopkan L, Hye Khan A, Imig JD, Kolar F, Neckar J. Tumour necrosis factor α contributes to improved cardiac ischaemic tolerance in rats adapted to chronic continuous hypoxia. Acta Physiol 2015;214:97-108. <u>https://doi.org/10.1111/apha.12489</u>
- 95. Waskova-Arnostova P, Elsnicova B, Kasparova D, Hornikova D, Kolar F, Novotny J, Zurmanova J. Cardioprotective adaptation of rats to intermittent hypobaric hypoxia is accompanied by the increased association of hexokinase with mitochondria. J Appl Physiol 2015;119:1487-1493. https://doi.org/10.1152/japplphysiol.01035.2014
- 96. Alanova P, Chytilova A, Neckar J, Hrdlicka J, Micova P, Holzerova K, Hlavackova M, Machackova K, Papousek F, Vasinova J, Benak D, Novaková O, Kolar F. Myocardial ischemic tolerance in rats subjected to endurance exercise training during adaptation to chronic hypoxia. J Appl Physiol 2017;122:1452-1461. https://doi.org/10.1152/japplphysiol.00671.2016
- 97. Kohutova J, Elsnicova B, Holzerova K, Neckar J, Sebesta O, Jezkova J, Vecka M, Vebr P, Hornikova D, Szeiffova-Bacova B, Egan Benova T, Hlavackova M, Tribulova T, Kolar F, Novakova O, Zurmanova JM. Anti-arrhythmic cardiac phenotype elicited by chronic intermittent hypoxia is associated with alterations in connexin-43 expression, phosphorylation and distribution. Front Endocrinol 2019;9:789. <u>https://doi.org/10.3389/fendo.2018.00789</u>
- 98. Neckar J, Kopkan L, Huskova Z, Kolar F, Papousek F, Kramer HJ, Hwang SH, Hammock BD, Imig JD, Maly J, Netuka I, Ostadal B, Cervenka L. Inhibition of soluble epoxide hydrolase by cis-4-[4-(3-adamantan-1-yl-

ureido)cyclohexyl-oxy]benzoic acid exhibits antihypertensive and cardioprotective actions in transgenic rats with angiotensin II-dependent hypertension. Clin Sci 2012;122:513-525. <u>https://doi.org/10.1042/CS20110622</u>

- Neckar J, Svatonova A, Weissova R, Drahota Z, Zajickova P, Brabcova I, Kolar D, Alanova P, Vasinova J, Silhavy J, Hlavackova M, Tauchmannova K, Milerova M., Ostadal B, Cervenka L, Zurmanova J, Kalous M, Novakova O, Novotny J, Pravenec M, Kolar F. Selective replacement of mitochondrial DNA increases the cardioprotective effect of chronic continuous hypoxia in spontaneously hypertensive rats. Clin Sci 2017;131:865-881. https://doi.org/10.1042/CS20170083
- 100. Neckar J, Alanova P, Olejnickova V, Papousek F, Hejnova L, Silhavy J, Behuliak M, Bencze M, Hrdlicka J, Vecka M, Jarkovska D, Sviglerova J, Mistrova E, Stengl M, Novotny J, Ostadal B, Pravenec M, Kolar F. Excess ischemic tachyarrhythmias trigger protection against myocardial infarction in hypertensive rats. Clin Sci 2021;135:2143-2163. <u>https://doi.org/10.1042/CS20210648</u>
- 101. Nedvedova I, Kolar D, Elsnicova B, Hornikova D, Novotny J, Kalous M, Pravenec M, Neckar J, Kolar F, Zurmanova J. Mitochondrial genome modulates myocardial Akt/GLUT/HK salvage pathway in spontaneously hypertensive rats adapted to chronic hypoxia. Physiol Genomics 2018;50:532-541. https://doi.org/10.1152/physiolgenomics.00040.2017
- 102. Nedvedova I, Kolar D, Neckar J, Kalous M, Pravenec M, Silhavy J, Korenkova V, Kolar F, Zurmanova JM. Cardioprotective regimen of adaptation to chronic hypoxia diversely alters myocardial gene expression in SHR and SHR-mtBN conplastic rat strains. Front Endocrinol 2019;9:809. <u>https://doi.org/10.3389/fendo.2018.00809</u>
- 103. Kolesova H, Bartos M, Hsieh WC, Olejnickova V, Sedmera D. Novel approaches to study coronary vasculature development in mice. Dev Dyn 2018;247:1018-1027. <u>https://doi.org/10.1002/dvdy.24637</u>
- 104. Kvasilova A, Olejnickova V, Jensen B, Christoffels VM, Kolesova H, Sedmera D, Gregorovicova M. The formation of the atrioventricular conduction axis is linked in development to ventricular septation. J Exp Biol 2020;223:jeb229278. <u>https://doi.org/10.1242/jeb.229278</u>
- 105. Olejnickova V, Kocka M, Kvasilova A, Kolesova H, Dziacky A, Gidor T, Gidor L, Sankova B, Gregorovicova M, Gourdie RG, Sedmera D. Gap junctional communication via connexin43 between Purkinje fibers and working myocytes explains the epicardial activation pattern in the postnatal mouse left ventrikle. Int J Mol Sci 2021;22:2475. <u>https://doi.org/10.3390/ijms22052475</u>
- 106. Neffeova K, Olejnickova V, Nanka O, Kolesova H. Development and diseases of the coronary microvasculature and its communication with the myocardium. WIREs Mech Dis 2022;e1560. <u>https://doi.org/10.1002/wsbm.1560</u>
- 107. Skarka L, Bardova K, Brauner P, Flachs P, Jarkovska D, Kopecky J, Ostadal B. Expression of mitochondrial uncoupling protein 3 and adenine nucleotide translocase 1 genes in developing rat heart: putative involvement in control of mitochondrial membrane potential. J Mol Cell Cardiol 2003;35:321-330. <u>https://doi.org/10.1016/S0022-2828(03)00016-6</u>
- 108. Netuka I, Szarszoi O, Maly J, Besik J, Neckar J, Kolar F, Ostadalova I, Pirk J, Ostadal B. Effect of perinatal hypoxia on cardiac tolerance to acute ischaemia in adult male and female rats. Clin Exp Pharmacol Physiol 2006;33:714-719. <u>https://doi.org/10.1111/j.1440-1681.2006.04423.x</u>
- 109. Ostadal B, Netuka I, Maly J, Besik J, Ostadalova I. Gender differences in cardiac ischemic injury and protectionexperimental aspects. Exp Biol Med 2009;234:1011-1019. <u>https://doi.org/10.3181/0812-MR-362</u>
- 110. Netuka I, Szarszoi O, Maly J, Riha H, Turek D, Ostadalova I, Ostadal B. Late effect of early hypoxic disturbance in the rat heart: Gender differences. Physiol Res 2010;59:127- 31. <u>https://doi.org/10.33549/physiolres.931833</u>
- 111. Milerova M, Charvatova Z, Skarka L, Ostadalova I, Drahota Z, Fialova M, Ostadal B. Neonatal cardiac mitochondria and ischemia/reperfusion injury. Mol Cell Biochem 2010;335:147-153. <u>https://doi.org/10.1007/s11010-009-0251-x</u>
- 112. Ostadal B, Ostadal P. Sex-based differences in cardiac ischaemic injury and protection: therapeutic implications. Br J Pharmacol 2014;171:541-554. <u>https://doi.org/10.1111/bph.12270</u>
- 113. Ostadal P, Ostadal B. Women and the management of acute coronary syndrome. Can J Physiol Pharmacol 2012;90:1151-1159. <u>https://doi.org/10.1139/y2012-033</u>
- 114. Milerova M, Drahota Z, Chytilova A, Tauchmannova K, Houstek J, Ostadal B. Sex difference in the sensitivity of cardiac mitochondrial permeability transition pore to calcium load. Mol Cell Biochem 2016;412:147-154. <u>https://doi.org/10.1007/s11010-015-2619-4</u>

- 115. Ostadal B, Drahota Z, Houstek J, Milerova M, Ostadalova I, Hlavackova M, Kolar F. Developmental and sex differences in cardiac tolerance to ischemia-reperfusion injury: the role of mitochondria. Can J Physiol Pharmacol 2019;97:808-814. <u>https://doi.org/10.1139/cjpp-2019-0060</u>
- 116. Ostadal B, Ostadalova I, Szarszoi O, Netuka I, Olejnickova V, Hlavackova M. Sex-dependent effect of perinatal hypoxia on cardiac tolerance to oxygen deprivation in adults. Can J Physiol Pharmacol 2021;99:1-8. https://doi.org/10.1139/cjpp-2020-0310
- 117. Alanova P, Huskova Z, Kopkan L, Sporkova A, Jichova S, Neckar J, Imig JD, Klevstig M, Kolar F, Reddy NR, Falck JR, Sadowski J, Nishiyama A, Kramer HJ, Melenovsky V, Cervenkova L, Kujal P, Vernerova Z, Cervenka L. Orally active epoxyeicosatrienoic acid analog does not exhibit antihypertensive and reno- or cardioprotective actions in two-kidney, one-clip Goldblatt hypertensive rats. Vasc Pharmacol 2015;73:45-56. https://doi.org/10.1016/j.vph.2015.08.013
- 118. Neckar J, Hsu A, Md Hye Khan A, Gross G, Nithipatikom K, Cyprova M, Benak D, Hlavackova M, Sotaova-Kasparova D, Falck J, Sedmera D, Kolar F, Imig J. Infarct size-limiting effect of epoxyeicosatrienoic acid analog EET-B is mediated by hypoxia inducible factor-1α via down regulation of prolyl hydroxylase 3. Am J Physiol Heart Circ Physiol 2018;315:H1148-H1158. <u>https://doi.org/10.1152/ajpheart.00726.2017</u>
- 119. Neckar J, Md Hye Khan A, Gross GJ, Cyprova M, Hrdlicka J, Kvasilova A, Falck JR, Campbell WB, Sedlakova L, Skutova S, Olejnickova V, Gregorovicova M, Sedmera D, Kolar F, Imig JD. Epoxyeicosatrienoic acid analog EET-B attenuates post-myocardial infarction remodeling in spontaneously hypertensive rats. Clin Sci 2019;133:936-951. https://doi.org/10.1042/CS20180728
- 120. Hrdlicka J, Neckar J, Papousek F, Huskova Z, Kikerlova S, Vanourkova Z, Vernerova Z, Akat F, Vasinova J, Hammock BD, Hwang SH, Imig JD, Falck JR, Cervenka L, Kolar F. Epoxyeicosatrienoic acid-based therapy attenuates the progression of postischemic heart failure in normotensive Sprague-Dawley but not in hypertensive Ren-2 transgenic rats. Front Pharmacol 2019;10:159. <u>https://doi.org/10.3389/fphar.2019.00159</u>
- 121. Bohuslavova R, Kolar F, Kuthanova L, Neckar J, Tichopad A, Pavlinkova G. Gene expression profiling of sex differences in HIF1-dependent adaptive cardiac responses to chronic hypoxia. J Appl Physiol 2010;109:1195-1202. https://doi.org/10.1152/japplphysiol.00366.2010
- 122. Cerychova R, Bohuslavova R, Papousek F, Sedmera D, Abaffy P, Benes V, Kolar F, Pavlinkova G. Adverse effects of Hifla mutation and maternal diabetes on the offspring heart. Cardiovasc Diabetol 2018;17:68. https://doi.org/10.1186/s12933-018-0713-0
- 123. Bohuslavova R, Cerychova R, Papousek F, Olejnickova V, Bartos M, Gorlach A, Kolar F, Sedmera D, Semenza GL, Pavlinkova G. HIF-1 α is required for development of the sympathetic nervous system. Proc Natl Acad Sci USA 2019;116:13414-13423. <u>https://doi.org/10.1073/pnas.1903510116</u>
- 124. Hrabalova P, Bohuslavova R, Matejkova K, Papousek F, Sedmera D, Abaffy P, Kolar F, Pavlinkova G. Dysregulation of hypoxia-inducible factor 1α in the sympathetic nervous system accelerates diabetic cardiomyopathy. Cardiovasc Diabetol 2023;22:88. <u>https://doi.org/10.1186/s12933-023-01824-5</u>
- 125. Adameova A, Hrdlicka J, Szobi A, Farkasova V, Kopaskova K, Murarikova M, Neckar J, Kolar F, Ravingerova T, Dhalla NS. Evidence of necroptosis in hearts subjected to various forms of ischemic insults. Can J Physiol Pharmacol 2017;95:1163-1169. <u>https://doi.org/10.1139/cjpp-2016-0609</u>
- 126. Lichy M, Szobi A, Hrdlicka J, Horvath C, Kormanova V, Rajtik T, Neckar J, Kolar F, Adameova A. Different signaling in infarcted and non-infarcted areas of rat failing hearts: a role of necroptosis and inflammation. J Cell Mol Med 2019;23:6429-6441. <u>https://doi.org/10.1111/jcmm.14536</u>
- 127. Lichy M, Szobi A, Hrdlicka J, Neckar J, Kolar F, Adameova A. Programmed cell death in the left and right ventricle of the late phase of post-infarction heart failure. Int J Mol Sci 2020;21:7782. <u>https://doi.org/10.3390/ijms21207782</u>
- 128. Semenovykh D, Benak D, Holzerova K, Cerna B, Telensky P, Vavrikova T, Kolar F, Neckar J, Hlavackova M. Myocardial m6A regulators in postnatal development: effect of sex. Physiol Res 2022;71:877-882. <u>https://doi.org/10.33549/physiolres.934970</u>
- 129. Benak D, Kolar F, Zhang L, Devaux Y, Hlavackova M. RNA modification m6Am: the role in cardiac biology. Epigenetics 2023;18:2218771. <u>https://doi.org/10.1080/15592294.2023.2218771</u>
- Benak D, Benakova S, Plecita-Hlavata L, Hlavackova M. The role of m6A and m6Am RNA modifications in the pathogenesis of diabetes mellitus. Front Endocrinol 2023;14:1223583. <u>https://doi.org/10.3389/fendo.2023.1223583</u>

131. Benak D, Holzerova K, Hrdlicka J, Kolar F, Olsen M, Karelson M, Hlavackova M. Epitranscriptomic regulation in fasting hearts: implications for cardiac health. RNA Biol 2024;21:1-14. https://doi.org/10.1080/15476286.2024.2307732