

Post mortem changes in Ca²⁺ transporting proteins of sarcoplasmic reticulum in dependence on malignant hyperthermia status in pigs.

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Meat quality of pigs is dependent on biochemical and biophysical processes in the time course post mortem (p.m.) and is associated with the intracellular Ca²⁺ homeostasis. However, there is little known about changes in the Ca²⁺ transporting proteins controlling the Ca²⁺ uptake of sarcoplasmic reticulum (SR) in the time course p.m. In this study changes in the Ca²⁺ transporting proteins were investigated in homogenates of longissimus muscles of 4 malignant hyperthermia susceptible (MHS) and 6 malignant hyperthermia resistant (MHR) Pietrain pigs. Muscle samples were obtained at different time intervals: biopsy 2 h prior slaughtering and from the carcass immediately after exsanguination (0 h), 45 min, 4 h, and 22 h p.m. The SR Ca²⁺ uptake rate was measured immediately after homogenization with closed calcium release channel (CRC), with opened CRC and without manipulation of CRC. Additionally the SR Ca²⁺ ATPase activity was determined. The results show: (i) The ability of SR to sequester Ca²⁺ declined to about 60% in the first 45 min p.m. in MHS samples irrespective of CRC state, whereas in MHR samples this decline was about 5%; (ii) Ca²⁺ uptake and Ca²⁺ ATPase activity were not different between the biopsy and 0 h samples, i.e. the stress of slaughter was of no immediate influence; (iii) The Ca²⁺ ATPase activity of the SR declined at about the same rate as the Ca²⁺ uptake in both MHS and MHR pig samples in the course of time p.m.; (iv) In samples, taken immediately after exsanguination, the Ca²⁺ ATPase activity of MHS pigs was higher than that of MHR pigs. However, in samples taken 4 h p.m. Ca²⁺ ATPase activity of MHS pigs has declined to about 30% of the value at 0 h; (v) The CRC can be closed and opened in all samples up to 22 h p.m. and seems to be fully functional at all sampling times; (vi) The CRC of MHS pigs is almost fully open, whereas the CRC of MHR pigs is only partially open at all sampling times; (vii) The permeability of the SR membrane to Ca²⁺ (determined as the ratio of SR Ca²⁺ ATPase with and without ionophore A23187) is the same in both MHS and MHR and did not change with ongoing time; (viii) No uncoupling of uptake from ATP hydrolysis occurred up to 4 h p.m., but the coupling differed between MHS and MHR for all time intervals with lower values for MHS pigs. The results suggest that the decreasing Ca²⁺ uptake rate of homogenates, sampled at different times p.m., is essentially caused by changes in the Ca²⁺ pump and not by changes in the CRC or an increased phospholipid membrane permeability to Ca²⁺.