Expression of Death Cellular Receptors FAS/CD95 and DR4 During Porcine Placentation

Expresión de los Receptores de Muerte Celular FAS/CD95 y DR4 Durante la Placentación Porcina

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SUMMARY: Apoptosis is a permanent and dynamic physiological process by which an organism eliminates the undesirable cells without causing an inflammatory response. The objective of this work was to study the expression of FAS, DR4 and other members of the TNF-R1 superfamily extrinsic route apoptotic receptors the DNA fragmentation and the cellular apoptosis in placental samples at the early, mid and late pregnancy on ± 30, ± 55 and ± 114 gestational days, respectively. We used placental histological sections of samples fixed in buffered saline formaldehyde. Immunohistochemical techniques were performed to detect the apoptotic receptors, whereas the DNA fragmentation was detected by TUNEL reaction and apoptotic cellular ultrastructure was detected by TEM conventional techniques. Apoptosis related receptors were immunolocalized in the early pig gestation and correlated with apoptosis, suggesting a role in the cellular remodelling of the placenta. At gestation day 55, apoptosis might be correlated to FAS route, but not by DR4-mediating pathway. At the end of gestation, increased apoptosis and both receptors markers were detected showing cellular death due to the extrinsic route through FAS and DR4 receptors. In conclusion, the immunolocalization of FAS and TNF-R1 receptors along the pig placental development correlates with TUNEL reaction and with apoptotic ultrastructure observed by TEM and seems to occur through different pathways along gestation.

KEY WORDS: FAS/CD95; DR4 and TNF-R1 family; Apoptosis receptors; Pig placenta.

INTRODUCTION

The apoptosis or programmed cellular death is an essential, permanent, dynamic and interactive biological process by which an organism eliminates the undesirable cells without causing an inflammatory response. The sequence of characteristic apoptotic processes is regulated by the interrelation of pro and anti apoptotic mechanisms, so the cellular death can be inhibited, balanced or stimulated (King & Cidlowski, 1998; Ameisne, 2002; Jersak & Bischof, 2002; Adams, 2003; Van Gurp et al., 2003; Fink & Cookson, 2005).

The programmed cellular death constitutes a key factor in the development of different physiological processes such as the placenta, an essential reproductive phenomenon in most mammals. It can be induced through intrinsic or mitochondrial route or by extrinsic route that begins by binding of ligands to membrane receptors. The best characterized cellular death receptors are FAS/CD95, DR4 and TNF-R1 (Yang & Korsmeryer, 1996; Barnhart et al., 2003; Inoue et al., 2003; Lavrik et al., 2005).

Early gestation in pigs, is characterized by rapid development and growth of the uterus and the embryos, resulting in the formation of an epitheliochorial, diffuse, noninvasive, folded and adequate placenta. The placenta plays a fundamental role during the gestation, allowing the implantation and maintenance of the pregnancy, and also determining the survival of the piglets (Amoroso, 1952; Dantzer, 1985; Dantzer & Leiser, 1993; Leiser & Kaufmann, 1994; Van der Lende & Van Rens, 2003).

One of the mechanisms of tissues remodelling regulation in both humans and animals is apoptosis (Hardy et al., 2001). Preliminary reports on remodelling of porcine placental cells have been done in our laboratory through morphologic and immunohistochemical detection of
apoptosis, in an attempt to characterize this process during porcine placentation (Merkis et al., 2007; Cristofolini et al., 2007).

In pigs, early embryonic mortality without specific-infectious or toxic cause ranges between 30 to 40%, most of them occurring before Day 30 of gestation (Lawrence, 1993; Pope, 1994). However, in our geographical area, Rio Cuarto (Argentina), this percentage can reach values up to 50-52% (Bosch et al., 2001). Therefore, determination of the basic mechanisms that trigger cell placental death or survival through key molecules identification during porcine gestation, will allow deepening the knowledge on physiological mechanisms that contributes for a successful pregnancy. This knowledge might be increased to improve the survival of the embryos/fetus and to improve the pig reproductive health, collaborating with the high productive and valuable pig industry of our region.

The aim of this work was to study the expression of FAS, DR4 and other members of the TNF-R1 superfamily extrinsic route apoptotic receptors by immunohistochemistry, DNA fragmentation by TUNEL and the apoptotic cellular ultrastructure by TEM in placental samples from early (±30 days), mid (±55 days) and late gestation (±114 days), in order to analyze the apoptotic process associated with porcine placentation.

MATERIAL AND METHOD

Mixed breed swines from a pig breeding establishment of Rio Cuarto city, Cordoba, Argentina (33.11°S; 64.3°O) were used. According to the ante-mortem and postmortem examinations they were considered free of disease. Of these bristles, a total of 15 placentas were processed: ±30 days of gestation (n=5), ±55 days (n=5) and upon maturity (approximately 114 days of pregnancy) (n=5). In all cases the reproductive tract was removed immediately after slaughter, washed with saline solution of Hank’s (SSH) containing sodic penicillin G, streptomycine sulphate and fungizone (Gibco, Grand Island, NY USA) and was maintained at 4°C until processing in the laboratory. Palpation was made to detect the location of the embryos or fetuses. The uterine horns were opened carefully and longitudinally with an incision on the anti-mesometrial edge to observe the implantation site and to gather samples of mesometrial endometrial and fetal placental tissues. Portions of approximately 6 mm each were taken from 5 placentas of every gestational period, fixed in 4 per cent (v/v) buffered-saline formaldehyde, pH 7.2-7.4 at 4°C and embedded in paraffin. From these paraffin-embedded tissues, histological sections of ±4 μm were obtained. The gestational age of the placentas was determined according to the crown-rump length of the embryos and/or fetuses obtained of each gestating bristle (Marrable, 1971).

Immunohistochemical techniques were performed using Santa Cruz, Inc. commercial antibodies: FAS (B-10), that recognizes FAS receptor and other members of the TNF R-1 superfamily, FAS (C-20), that recognizes only FAS receptor and DR4 that binds DR4 receptors, biotinylated secondary antibodies pool, streptavidin conjugated to horseradish peroxidase (LSAB®+Systems HRP, Dako Cytomation) and 3,3-diaminobenzidine chromogen solution (Liquid DAB+Substrate Chromogen System, Dako Cytomation). The sections were counterstained with Mayer’s haematoxylin. The results were expressed as semiquantitative values, determining: (-): negative, (+): weak, (++): moderate and (+++): strong immunostaining. For every gestational period the distribution of the immunolabelling intensity was determined by means of the High Score value (Selam et al., 2001). Simultaneously, immunohistochemical negative controls were carried out.

The detection of the DNA fragmentation was made by means of TUNEL technique (TdT-mediated-dUtp Nick End Labeling) using commercial equipment ApopTag® (Chemicon International). The samples were observed in a light microscope Axiophot (Carl Zeiss) and the images acquired realized with a digital camera Powershot G6, 7.1 megapixels (Canon INC, Japan). The results were expressed as qualitative.

For electron microscopy the placental samples were fixed in 2.5 % glutaraldehyde in 0.2 M-s-collidine pH 7.4, post-fixed in 1% osmium tetroxide in 0.2 M-s-collidine pH 7.4, dehydrated in increasing concentrations of acetone and embedded in EMBed 812 resin. This ultra-thin sections (60 nm) were cut and placed on cupper grids, counterstained in uranyl acetate and lead citrate. The sections were examined in transmission electronic microscope Elmiskop 101 (Siemens, Germany).

RESULTS

At 30 days of pregnancy, no immunolabelling for FAS B-10 was seen in the villi, blood vessels and glands, whereas in placental connective tissue weak labelling was observed. Immunoreactivity of FAS C-20 was found moderate in the endometrial connective tissue and weak in the mesenchyme, while no labelling was seen in the villi, glands and blood vessels. Villi, blood vessels and glands were not reactive to...
membrane receptor DR4 but its expression was moderate in endometrial and mesenchymal connective tissues.

At 55 days of pregnancy moderate immunolabelling for FAS B-10 was seen in the villi and endometrial connective tissue, weak expression in blood vessels and moderate labelling in glands. Similarly, FAS C-20 expression was moderate in the villi and in endometrial connective tissue. In this gestational period no expression of the DR4 was observed neither in the villi nor in maternal blood vessels, but weak and moderate staining was seen in mesenchyme and endometrial connective tissue, respectively.

At late pregnancy, immunoreaction detected weak labelling for FAS B-10 in mesenchymal tissue but no staining in villi and blood vessel. FAS C-20 expression was found negative in villi, moderate in mesenchymal connective tissue and weak in blood vessels. For DR4 receptors, the immunolabelling was negative in villi and blood vessels, and moderate in mesenchymal tissues.

Immunohistochemical negative controls showed no labelling expression in the studied tissues. At the early gestation in the chorionic villi the distribution of the immunolabelling intensity determined by means of the High Score value revealed low immunolabelling of DR4 receptors.
and high expression of FAS membrane receptors and others belonging to the TNF R-1 superfamily (Fig. 1). At 55 days, the distribution of the immunolabelling intensity of the three apoptosis receptors diminished, predominating the expression of FAS receptors. At late pregnancy a remarkable increase in the expression of the apoptosis receptors was observed, especially for DR4 receptors.

**DISCUSSION**

Our results indicate that at the beginning of the porcine gestation the cellular remodelling of the placenta may be occurring through FAS membrane receptor and other members of the TNF-R1 superfamily. The HScore percentages of this receptors in the first gestational period correlate to the apoptosis levels by way of FAS comparable with those observed in organs in situation of immunological privilege as it is the placenta.

During porcine gestation a correct metabolic exchange between maternal and fetal systems is critical to meeting the nutritional demands of the conceptus. The adequate remodelling of placental cell through apoptotic processes is essential for the maintenance of homeostasis of this transitional organ with advancing gestation (Dantzer & Leiser; Finch et al., 2004). The post-implantational period is when an important angiogenesis takes place, which is necessary to obtain the suitable transport of nutrients to the embryo/fetus (Finch et al., 2004; Merkis et al., 2006). In humans and pigs, low weight at birth is the most important factor of neonatal morbidity and mortality; in pigs the faults that occur at 30 days of pregnancy would directly influence

RESUMEN: La apoptosis es un proceso fisiológico, dinámico y permanente a través del cual un organismo elimina células indeseables sin provocar una respuesta inflamatoria. El objetivo del presente trabajo fue estudiar la expresión de los receptores de la vía extrínseca de apoptosis, FAS, DR4 y otros miembros de la super familia TNF-R1, la fragmentación del ADN y la apoptosis celular a través de TEM, en muestras placentarias del inicio, la mitad y el final de la gestación, hacia el día ± 30, ± 55 y ± 114 de preñez, respectivamente. Se realizaron cortes histológicos de las muestras placentarias fijadas en formol tamponado. Para la detección de los receptores de apoptosis se realizaron técnicas inmunohistoquímicas, para el estudio de la fragmentación del ADN se utilizó el ensayo TUNEL y para el análisis de la ultraestructura celular apoptótica la técnica convencional de TEM. La inmunolocalización de los receptores de muerte celular al inicio de la preñez porcina sugiere el rol de la apoptosis en la remodelación celular placentaria. Hacia el día 55 de preñez, la apoptosis detectada ocurriría únicamente a través de la vía del receptor FAS, no del receptor DR4. Al final de la gestación, se detectó un incremento de la apoptosis y la expresión de ambos receptores, indicando que la muerte celular a través de la vía de señalización extrínseca estaría inducida por los receptores FAS y DR4. En conclusión, la inmunolocalización de los receptores FAS y otros miembros del TNF-R1, los resultados de TUNEL y la ultraestructura celular apoptótica observada en la placencía porcina, indican que la apoptosis detectada ocurre por diferentes vías de inducción a lo largo de la gestación.

PALABRAS CLAVE: FAS/CD95; DR4 y superfamilia TNF-R1; Receptores de apoptosis; Placenta porcina.
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