NON-SPECIFIC MORPHOLOGICAL CHANGES IN THE ORGANS OF
CHICKENS INFECTED EXPERIMENTALLY WITH MAREK'S
DISEASE VIRUSES

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Abstract


Description is given of hitherto unnoticed pathological changes in the organs of Hybro, White Hisex, F1 inbred line CB×IA and Brown Leghorn chickens after experimental infection with oncogenic strains (GA, field isolates VUB-83 and RB-1B) and non-oncogenic strains (field isolates M and K) of Marek's disease virus (MDV). The chickens were infected on the 2nd post-hatching day and sacrificed at weekly intervals up to the 5th week p. i. Pathomorphological examination revealed, in addition to MD-specific and cytolytic morphological changes in the organs, the presence of syncytial formations, granulomas, "Hassall's corpuscle-like" structures, dystrophic changes in follicular epithelium-associated (FAE) cells and changes in the corticomedullary membrane of the bursa of Fabricius follicles, hypertrophy and hyperplasia of Hassall's corpuscles, hyperplasia of myoid elements in the thymus, spheroids in peripheral nerves and glomerulopathy.

The changes observed in the lymphoid organs are regarded as a morphological manifestation of local cellular defence, and the changes in the nerves and kidney as an autoimmune process and, consequently, as non-specific of MD. Their description contributes to a better understanding of the complex pathogenesis of Marek's disease of poultry.

Marek's disease, non-specific morphological changes, lymphoid organs, cellular immunity

A characteristic feature of Marek's disease of poultry (MD) as of many other herpes virus-produced diseases is a differing type of virus-cell interaction, ranging from fully productive and semi-productive infection to non-productive neoplastic and non-productive latent infection (Payne 1985). However, the final outcome of infection with Marek's disease virus (MVD) is strongly affected by pathogenicity of the viral strain, genotype of the host and its immune status. The main pathological changes occurring in the organs during the complex pathogenesis of MD produced by different MDV strains are well-known (Calnek 1986).

Lymphoid infiltration and proliferation, and development of lymphomas are regarded as specific of MD (Payne and Biggs 1967). That MDV is lymphotropic appears also from cytolytic changes with a secondary inflammatory reaction in lymphoid organs, particularly in the bursa of Fabricius (BF), which we do not regard as specific of MD but as a major criterion of the pathogenicity of MVD (Halouzka and Jurajda 1991).

The cell structure of the BF corresponds in essence to the germinal centres of mammalian lymphoreticular tissues and performs the function of a peripheral lymphoid organ (Toivanen et al. 1987). It can therefore be assumed that the morphological changes arising in lymphoid follicles of the bursa after antigenic stimulation are also a manifestation of local immune response.

Our experimental studies of the pathogenesis of MD revealed, in addition to specific changes in various tissues and regressive changes in the lymphoid organs of poultry, also some other changes that have not been generally described in the pathogenesis of MD. The object of the present study is to describe these unusual changes designated by us as non-specific of MD and point out their formal and causal pathogenesis.
Materials and Methods

There oncogenic MDV strains were used. Prototype strain Georgia (GA) (Eidson and Schmittle 1968) was administered in doses of approximately $1.6 \times 10^3$ and $1.6 \times 10^4$ PFU to 1-old meat-type Hybro chicks, layer-type White Hisex chicks and to F₁ inbred line CB×IA chicks that are more resistant to MD than Brown Leghorns (BrL) (Jurajda et al. 1989). A highly oncogenic MDV strain VUB-83 (Jurajda and Halouzka 1989) was administered to 1-day old Hybro, White Hisex and BrL chicks in the same doses as MDV strain GA. An oncogenic MDV strain RB-1B (Jurajda and Halouzka 1990) was administered to BrL chicks in doses of approximately $10^2$, $10^3$ and $10^4$ PFU in 0.2 ml inoculum volumes.

The non-oncogenic MDV strains used in our study were field isolates M and K (Jurajda and Halouzka 1991). They were administered to 2-day old BrL chicks in doses of approximately $10^2$, $10^3$ and $10^4$ PFU in 0.2 ml inoculum volumes (Halouzka and Jurajda 1991).

Ten chickens of each infected group were sacrificed, together with 10 controls, 1, 2, 3 and 5 weeks p. i. All of them were subjected to pathomorphological examination. Samples for histopathological examination were taken from lymphoid organs, skin, peripheral nerves, gonads, liver and kidney. They were fixed in 10% aqueous solution of neutral formaldehyde and stained with haematoxylin and eosin, Giemsa stain and luxol blue. Cytolytic changes in the lymphoid organs were assessed quantitatively using immunosuppression (Si) degrees of 0 to 4 (Halouzka and Jurajda 1991).

Results

Unusual morphological changes were observed in the BF, thymus, skin, peripheral nerves and kidney of experimental chickens. The frequency of these changes varied from experiment to experiment and in dependence on the post-infection period.

Bursa of Fabricius

In the germinal centres of lymphoid follicles classified as Si degrees 2 to 3, syncytial formations (1 to 3 in number), arisen by fusion of 2 to 5 dentritic reticular cells, were observed (Fig. 1). Some of them corresponded to the morphological characteristics of large polykaryons, others were reminiscent Hassall's corpuscles of the thymus. They were found most frequently in Hybro chickens, CB×IA hybrids and White Hisex chickens 1 and 2 weeks after infection with strain GA and in BrL chickens 3 weeks after infection with strain M and considerably less frequently 1 week after infection with strain K. In the experiments with strain VUB-83 they were observed only sporadically 1 week p. i. in Hybro and White Hisex chickens. In all experiments they were found very rarely also in non-infected controls (in a total of 5 control birds).

In experiments with strain K some syncytial formations gave rise to structures morphologically identical with minute granulomas showing central markedly eosinophilic necrosis rimmed with cells with signs of epitheloid elements.

Vacuolar and dystrophic changes to necrosis of follicular epithelium-associated (FAE) cells constituting the surface epithelium of the villi were an accompanying phenomenon of cytolytic changes in the follicular medulla from the 3rd week p. i. in all experiments (Fig. 2). Sporadically, their protrusion to desquamation to the cavity of the organ was observed.

Distortion and thickening of the corticomedullary membrane of the follicles and transition of flat epithelial cells to cubic ones was seen in all cases classified as Si degree 2 to 4, particularly where marked lymphocyte depletion of the follicle medulla was observed. These changes were conspicuous mainly in BrL chickens infected with MDV strains VUB-83, GA and K.
Fig. 1. Production of syncytial structures in the germinal centres of lymphoid follicles in the bursa of Fabricius. HE $\times$ 640.

**Thymus**

Hyperplasia and hypertrophy of Hassall’s corpuscles due to increased hyalination of epithelial cells, frequently in the form of concentric hyaline layers, accompanied the regressive changes in the thymus (Fig. 3). With the higher Si degrees this phenomenon was more pronounced and the centres of Hassall’s corpuscles showed cavities containing chromatin granules and residues of cellular bodies. These corpuscles were rimmed with granulocytes. The incidence of altered Hassall’s corpuscles was associated with an increase of myoid elements and with the development of germinal lymphoid centres in the medulla. Although these changes were observed in all experiments, they were most frequent 1 week after infection with strains GA and VUB-83 in Hybro and White Hisex chickens and 2 and 3 weeks after infection with strains M and K in BrL chickens.

**Skin**

In BrL chickens infected with strain K feather-pulp changes were regular findings in addition to well-known skin lesions. A conspicuous finding was an enlarged capillary network and mixed cellulization of the thin mesenchymal reticulum of the medulla round the veins. The medulla itself was markedly eosinophilic and showed dissociation of the structure. These changes affected more chickens and at a higher intensity at the end of the experiment, being particularly intense in the group given the high virus dose.
Peripheral Nerves

Multiple spheroid eosinophilic to amphophilic formations in neuraxons (Fig. 4) accompanied by granulation of myelin sheaths of the surrounding neurits were frequent and marked findings in RB-1B-infected BrL chickens from the 3rd post-infection week. These lesions were accompanied neither by interneuritic nor perineural lymphoid infiltration nor by any damage to Schwann cells.

Kidney

Two BrL chickens infected with strain GA showed proliferation of mesangial cells, thickening of the basement membranes and swelling of the endothelia of capillaries in some glomeruli 5 weeks p. i. (Fig. 5). These changes were accompanied by MD-specific pleomorphic infiltration of the intertubular tissue.

Discussion

Syncytial formations, also referred to as polykaryons or “Hassall’s corpuscle—like” structures, in the medulla of lymphoid follicles of the BF have not been described in the literature on the pathogenesis of MD before. However, avian germinal centres of the BF follicles are known to contain, besides T- and B-lymphocytes, also macrophages and dentritic reticular cells (Jeuring et al. 1989).
Moreover, the avian follicular medulla contains another two cell types that have not been identified in mammals, namely follicular epithelial cells between the medullary and cortical parts of the follicles and "Hassall's corpuscle-like" structures consisting of epithelial cells (Naukkarinen and Sorvari 1982). Structures in the germinal centres of mammalian lymphoreticular tissues identical with the syncytial structures found by us in the follicular medulla of the BF have been described at these locations after antigenic stimulation. Macrophages are transformed to cells epitheloid in appearance and fused (Syrjanen 1982). In our view, the syncytia, or polykaryons, in the BF follicular medulla are therefore a morphological manifestation of cellular immunity after antigenic stimulation with MDV virus as is the increase in, and enlargement of, "Hassall's corpuscle-like" structures.

In the light of our observations the formation of granulomas in the BF can be regarded as an enhanced immune response to stimulation with MDV. From the viewpoint of general pathology epitheloid granulomas represent T-cell-mediated type of immunity and are a favourable reaction of the body (Bednář 1982). To a certain extent, they can be therefore regarded as morphological manifestation of the relation between the pathogenicity of MDV strain K and the defence potential of BrL chickens. In other experiments the granulomatous type of reaction was not observed.

FAE cells are of mesenchymal origin and have, similarly to mammalian M-cells (Lupetti et al. 1983a, b; Toivanen et al. 1987), an important function in antigen transport. Dystrophic changes of FAE cells after infection with MDV
are apparently a result of the interaction between the virus and these cells, similarly to cytolytic changes in other elements of mesenchymal origin.

The incidence of pathologically altered Hassall’s corpuscles of the thymus correlated with the production of granulomas in the BF. Avian thymus has also the properties of a peripheral lymphoid organ, and reticular epithelium has the ability of phagocytosis (Janković et al. 1972 as cited by Benda 1990). Cytolytic infection results in the proliferation of reticular epithelium in the medulla (Payne 1985) and this proliferation contributes to the intensification of the production of Hassall’s corpuscles and of subsequent changes which we regard as a manifestation of local cellular immunity. No explanation can be offered for the increase of myoid elements.

The pulpitis observed in the skin was a frequent finding in our experiments with isolate K. Payne (1985) in his study on MD has mentioned the occurrence of occasional lymphoid infiltrates in feather pulp without specifying their character. In our view, the pulpitis of the skin in strain K-infected chickens is the result of viraemia and part of MD-specific changes.

The pathogenesis of spheroids observed in the peripheral nerves is difficult to interpret; whether it is the case of primary or secondary degeneration cannot be decided without knowing the status of the respective ganglion cells. Nevertheless, some writers have described degenerative changes of heavily infiltrated nerves in MD as Waller’s type of degeneration (Lampert et al. 1977). From some other studies, however, it appears that primary demyelination is caused by an immunopathological process and evidence has been obtained to indicate that

Fig. 4. Multiple spheroids in neuraxons of the n. ischiadicus. HE × 640.
idiopathic polyneuritis is a spontaneous autoimmune disease of poultry (Pepose et al. 1981). In our view, a similar process cannot be excluded in our experiments with MDV strain RB-1B.

The cause of renal changes cannot be detected with the methods used in our study. An autoimmune character of similar changes has been demonstrated (Ley and Yamamoto 1979, Winter and Majid 1984). The description by Pradham et al. (1988) of immune complex-mediated glomerulopathy in MD 10 weeks after infection with MDV is morphologically identical with our findings.

Comparison of the morphological observations made in our experimental studies on the pathogenesis of MD with the corresponding results reported by other writers allows us to conclude that the changes in the BF and thymus are a manifestation of local cellular defence of the body and are not specific of MD. The changes in the nerves and kidney suggest their autoimmune character. The identification of the pathomorphological changes observed in the organs contributes to a better understanding of the complex pathogenesis of MD.

Nespecifické morfologické změny v orgánech kuřat po experimentální infekci viry Markovy nemoci

Práce pojednává o dosud nepozorovaných patologických změnách v orgánech experimentálních kuřat genotypů Hybro. Hisex bílý, inbredních linii F1 CB × IA a Brown Leghorn po infekci onkogenními kmeny (GA, terénní izoláty VUB-83,
RB—1B) a neonkogenními kmeny (terénní izoláty M a K) MDV. Kuřata byla infikována 2. den věku a patomorfologicky vyšetřována v týdenních intervalech do 5. týdne p. i. Kromě MD-specifických a cytologických morfologických změn v orgáncích, byly pozorovány syncytialní útvary, granulomy, „Hassal’s corpuscle-like” struktury, dystrofické změny FAE buněk a změny kortikomedulární membrány ve folikulech Fabriciovy burzy, hypertrofie a hyperplazie Hassalových tělísek a hyperplasie myoidních elementů v thymu, sféroidy v periferních nervech a glomerulopathie.

Pozorované změny v lymfatických orgáncích jsou považovány za morfologický projev místní celulární obrany a změny v nervech a leduváčkách za autoimunitní proces a tudíž nespecifické pro MD. Jejich popis přispívá k objasnění složité patogeneze Markovy nemoci drůbeže.

Неспецифические морфологические изменения органов цыплят после экспериментальной инфекции вирусами болезни Марека

В работе дается изложение до сих пор не наблюдаемых патологических изменений органов экспериментальных цыплят генотипов Гибро, Гисекс бельй, инбредных линий F1 СВх IA и Браун Легхорн после инфекции онкогенными штаммами (GA, изолями VUB-83, RB-1B) и неонкогенными штаммами (полученными в полевых условиях изолями М и К) MDV. Цыплята инфицировали в возрасте двое суток и патоморфологически исследовали с недельным интервалом до 5 недели после инфекции. Помимо MD-специфических и цитологических морфологических изменений органов наблюдали синцитиальные формирования, грануломы «Hassal’s corpuscle-like» структуры, дистрофические изменения клеток FAE и изменения кортика-медуллярной мембраны в фолликулах фабрициевой сумки, гипертрофию и гиперплазию телца Гассалла и гиперплазию миоидных элементов тимуса, сфероиды в периферических нервах и глюмерулопатию.

Наблюдаемые изменения лимфатических органов считали морфологическим проявлением местной целлулярной защиты и изменения нервов и почек - автоиммунным процессом и, следовательно неспецифическими для болезни Марека. Их описание вносит вклад в выяснение сложного патогенезиса болезни Марека птицы.

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