Short communication

Immunohistochemical investigation of cells expressing CD21, membrane IgM, CD32 and a follicular dendritic cell marker in the lymphoid tissues of neonatal calves

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Activation of B lymphocytes in the presence of passive maternal antibodies depends on expression of CD21, membrane IgM and CD32. On colligation with IgM, CD32 inhibits activation whereas CD21 enhances it. Recently, we assessed expression of CD21 and CD32 on IgM+ cells from lymphoid tissues of newborn calves by flow cytometry, but this approach does not provide information about spatial distribution within lymphoid compartments. Therefore, histologic sections of lymphoid tissues from newborn and 7-month-old calves were examined using an immunoperoxidase technique. In all calves, CD21 and IgM stained cells were collocated in the cortex and paracortex of the retropharyngeal lymph node, in the marginal zone of the spleen and in lymphoid aggregates of palatine tonsils. Most CD32+ cells were in the mantle zone of lymphoid follicles in 7-month-old calves, whereas only weak staining was observed in newborns. A few CD32+ cells were also observed in the paracortex at both ages. Absence of CD32+ cells in the center of follicles suggests that IgM+CD32− cells observed previously by flow cytometry were from germinal centers. Overall, there were few organized lymphoid aggregates within lymphoid tissues of newborn calves, and follicular dendritic cells were virtually undetectable. Their absence may be an important limitation for neonatal immunization.
bound to antigen (Dempsey et al., 1996; Lyubchenko et al., 2005). However, various soluble complement components including C3 (which on breakdown forms C3d), are low in neonates (reviewed by Firth et al., 2005).

CD32 (FcγRII) is an IgG Fc binding receptor expressed by B cells, neutrophils and monocytes, that binds IgG within immune complexes (Nimmerjahn and Ravetch, 2008). The b isoform (FcγRIIb) of CD32 expressed by B cells has an immunoreceptor tyrosine-based inhibitory motif (ITIM) in the cytoplasmic tail. Colligation of mIgM and CD32 by antigen–antibody complexes suppresses anti-motif (ITIM) in the cytoplasmic tail. Colligation of mIgM and CD32 by antigen–antibody complexes suppresses antibody responses by increasing the threshold for activation. We have shown that the majority of IgM+ B lymphocytes in the blood is potential for colligation of mIgM with both CD21 and CD32 on B cells by complement bound antigen–antibody complexes. Thus, the relative proportion of C3d and IgG in complexes may control B cell activation or inhibition by regulating the threshold of activation. We have shown that the majority of IgM+ B lymphocytes in the blood of calves express both CD21 and CD32 (Chattha et al., 2010a). In the spleen and lymph nodes of newborn and 7-week-old calves almost all IgM+ B cells express CD21, whereas only 75% to 83% of IgM+ B cells express CD32, suggesting a possible bias toward activation (Chattha et al., 2010b). However, receptor expression was investigated by flow cytometry, which does not provide information about the location of receptor-expressing cells in the lymphoid tissues. The location of CD21, IgM and CD32 expressing cells can have important implications for immune responsiveness. Therefore, we compared the distribution of cells expressing these receptors in the lymphoid tissues of newborn and 7-month-old calves (as representative of adults) by immunohistochemistry.

Follicular dendritic cells (FDCs) also play an important role in antigen presentation for induction of antibody responses. Complement and Fc receptors, including CD32, on FDCs trap the antigen in its complexed form (antigen–antibody or antigen–complement or antigen–antibody–complement). FDCs display antigen, help in affinity maturation and selection of activated B lymphocytes, and contribute towards formation of germinal centers (GCs), and generation of memory and plasma cells (Tew et al., 2001). The low number of mature FDCs in the lymphoid tissues of neonatal mice correlates with low serum IgG, the small number of antibody secreting cells and low number of GCs in lymphoid tissues (Pihlgren et al., 2003). Limitations of B cells and various accessory cells including FDCs in neonates may partly be responsible for observed poor antibody responses (Marshall-Clarke et al., 2000; Siegrist and Aspinall, 2009). Recently, a new monoclonal antibody FDC-B1 was produced, which is specific for a 28-kDa (as measured by western blotting) surface antigen (postulated to be a glycoprotein) of bovine FDCs (Melot et al., 2004). It was used in this study to analyse the FDC network in neonatal calves since the number of FDCs in lymphoid tissues could be an important limiting factor for induction of antibody responses.

2. Materials and methods

All experimental procedures were approved by the University of Guelph Animal Care Committee and were conducted according to the guidelines of the Canadian Council for Animal Care.

2.1. Lymphoid tissue samples

Spleen, retropharyngeal lymph node (RPLN) and palatine tonsils were collected at necropsy from 4 clinically healthy Holstein bull calves. The two newborn calves (2 and 5 days of age) were born at the Elora Dairy Research Station, whereas the two 7-month-old calves were born off site but were raised at the Ponsonby Research Station; these two research facilities are operated coordinately by the University of Guelph. Calves received at least four litres of colostrum within 16 h of birth. Calves were humanely euthanized by intravenous injection of Euthansol (Intervet/Schering-Plough Animal Health, Kirkland, Quebec). 5 mm × 5 mm × 5 mm sections of tissue were harvested and embedded in OCT compound (Tissue-Tek, Sakura Finetek, Torrance, CA), then snap frozen by dipping in liquid nitrogen for 15–20 s. Frozen tissue samples were kept at –80°C until processed.

2.2. Immunohistochemistry staining

5 μm thick serial sections of lymphoid tissues were cut using a cryostat (Leica CM3050 S) and mounted on charged glass slides (Fisher Scientific, Pittsburgh, PA). Cryostat sections were fixed in pre-cooled acetone (–20°C) for 10 min at 4°C, then air-dried at room temperature (RT). Slide-mounted tissue sections were rehydrated in phosphate buffered saline (pH = 7.4) and pre-treated with 0.03% hydrogen peroxide (Sigma–Aldrich) for 10 min to block endogenous tissue peroxidases. Slides were washed twice with PBS for 5 min, then blocked with serum-free protein blocking solution (Dako, Carpenteria, CA) for 15–20 min at RT. Sequential tissue sections were stained

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<td>Monoclonal antibodies (mAb) used in immunohistochemistry.</td>
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<td>(Clone)</td>
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<td>CCG36 (mouse)</td>
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<td>CC21 (mouse)</td>
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<td>BM-23 (mouse)</td>
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<td>FDC-B1 (mouse)</td>
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separately with appropriate dilutions of mouse monoclonal antibodies (mAb) directed against bovine CD21, IgM, CD32 and the 28-kDa surface antigen of bovine FDC (referred to as anti-FDC) (Table 1). Lectin from *Arachis hypogaea* (peanut agglutinin [Sigma], working dilution 1:400) was used to detect GCs (Coico et al., 1983). To stain for CD21, IgM or GCs, slides were incubated with the working dilution of mAbs or peanut agglutinin for 1 h at RT, whereas for CD32 and FDC, slides were incubated overnight at 4 °C. A goat anti-mouse/anti-rabbit immunoglobulin (EnVision™+ dual link) polymer detection system (Dako) was used as the secondary antibody with Nova Red (SK 4800, Vector labs, Peterborough, UK) as chromogen. Sections were counter-stained with Harris haematoxylin for 5 min, followed by destaining in 1% acid alcohol (1% hydrochloric acid in 70% alcohol) and weak ammonia solution. Cover slips were applied using glycerol-mounting medium (Dako) and slides were kept at RT until examined by light microscopy (Leica DM RA2).

3. Results

3.1. Newborn calves

The cortex of the RPLN was small in size and contained few organized primary follicles. Most of the CD21+ cells were in the subcapsular region and cortex; only a few cells were present in the paracortex and medulla (Fig. 1A). IgM and CD21 positive cells exhibited the same staining patterns and were located in the same regions (Fig. 1B). CD32+ cells were scarce and detected primarily in the paracortex and medulla. These cells could be lymphocytes, macrophages or dendritic cells (Fig. 1C). There was minimal CD32 staining in the cortex.

In the spleen, cells demonstrating CD21 and IgM immunoreactivity were present primarily in the marginal zone of the white pulp (Fig. 2A, B). There were no recognizable GCs or lymphoid follicles around the periarteriolar lymphoid sheath (PALS) (Fig. 2A, B). A few IgM+ cells were scattered in the red pulp of spleen consistent
with the staining pattern and morphology expected for plasma cells (Fig. 2C). Subjectively, staining for IgM and CD21 was less extensive in newborns compared to older calves.

In palatine tonsils, as in lymph nodes, CD21 and IgM positive cells had identical staining patterns and were present in the sub-epithelium, as scattered cells or organized lymphoid follicles (Fig. 3A, B). A few IgM+ cells were also

![Fig. 2. Spleen of newborn (A, B, C) and 7-month-old calves (D, E) stained with monoclonal antibodies for CD21 (A, D) and IgM (B, C, E). Arrows indicates plasma cells in the red pulp (C). Figures A and B are serial sections from spleen of a newborn calf and show similar distribution of staining for CD21+ and IgM+ cells. Figures D and E are serial sections from spleen of a 7-month-old calf and show similar distribution for CD21+ and IgM+ cells. Magnification is indicated by the bar in each image. PALS = periarteriolar lymphoid sheath, MZ = marginal zone, RP = red pulp, WP = white pulp, Fo = follicle.](image)

![Fig. 3. Palatine tonsils of newborn (A, B, C) and 7-month-old calves (D, E) stained with monoclonal antibodies for CD21 (A, D), IgM (B, E) and CD32 (C). Figures A, B and C are serial sections from tonsil of a newborn calf and show similar distribution for CD21+, IgM+ cells and minimal expression of CD32. Figures D and E are serial sections from tonsil of a 7-month-old calf and show similar distribution for CD21+ and IgM+ cells. Magnification is indicated by the bar in each image. SG = salivary gland, CE = crypt epithelium, Fo = follicle, IFZ = interfollicular zone.](image)
detected within crypt epithelium (Fig. 3B). CD32 staining was weak, staining only a few cells in the lymphoid follicles and showed some overlap with CD21 and IgM staining (Fig. 3C).

FDCs could not be detected in RPLN, spleen or palatine tonsils of newborn calves (Fig. 4A, D).

3.2. 7-Month-old calves

In RPLN, CD21 and IgM positive cells showed overlapping patterns of staining in the mantle zone of the follicles and in the paracortical area (Fig. 1D, E). CD21+ cells were also present within the follicles suggesting expression on centroblasts (rapidly proliferating lymphocytes in the dark zone of GC), centrocytes (lymphocytes in the light zone of GC with low rates of proliferation) and FDCs (Fig. 1F). CD32 was primarily detected in the corona of follicles, interfollicular area and paracortex whereas almost no staining was detected in the GCs suggesting absence of this receptor on centroblasts and centrocytes (Fig. 1G).

In spleen, CD21+ and IgM+ cells showed overlapping staining patterns in the follicles around the PALS and in the marginal zone of the white pulp (Fig. 2D, E). CD32 staining overlapped with CD21 except for minimal staining in the GCs (data not shown).

In palatine tonsils, CD21 and IgM staining was detected in the mantle zone and GCs of lymphoid follicles, and in the interfollicular zone (Fig. 3D, E). PNA was used to identify GCs and the staining patterns were compared to those for other antibodies to confirm the presence of stained cells in GCs (Fig. 4C). FDCs in a characteristic mesh-like network were detected in lymphoid follicles of retropharyngeal lymph nodes and spleen and in lymphoid aggregates in palatine tonsils (Fig. 4B, E, F, G). FDCs were present primarily in the light zone, with few in the dark zones of GCs. Cells in these regions stained with CD21 mAb as well, suggesting co-expression of this receptor (Figs. 1F, 2D, 3D). Cells in the FDC regions also stained with anti-IgM mAb suggesting display of IgM complexes on FDCs or their interaction with IgM+ B cells (data not shown).

4. Discussion

This study was completed to determine the spatial distribution of CD21, IgM and CD32 expressing cells in the lymphoid tissues and is supplementary to an earlier...
investigation showing expression of CD21 and CD32 on IgM+ cells in lymphoid tissues of calves by flow cytometry (Chattha et al., 2010b). In addition, this is the first study to investigate the appearance of FDCs with age in cattle. FDCs are important for antibody responses and their absence may result in limited responses in newborns (Pihlgren et al., 2003).

Our previous study using flow cytometry showed that greater than 90% of IgM+ cells in the spleen and lymph nodes of newborn calves express CD21. The present study shows a major overlap in immunostaining patterns for CD21 and IgM in these tissues in both newborns and 7-month-old calves. Because CD21 is the receptor for complement C3d, these findings suggest that B cells in the lymphoid tissues of newborns can be activated by complement-containing antigen or adjuvants (Pihlgren et al., 2004). A recent study has shown a previously unsuspected role for CD21 on B cells in carrying and transferring antigen–antibody complexes from subcapsular sinus macrophages to the FDCs in the lymph nodes (Phan et al., 2007). The absence of FDCs in neonates may be a major limitation for employment of this mechanism in the generation of antibody responses.

The presence of CD21 expressing lymphocytes in the marginal zone of the spleen in newborn calves suggests that expression of this receptor may not be a limiting factor for T independent responses as observed in human neonates and infants (Timens et al., 1989). However, the overall reduced number of B cells expressing CD21 and IgM, in this age group may impose constraints on B cell responses and delay the kinetics of antibody production. Since antigenic stimulation was minimal in the animals used in this study, the lack of organized GCs in the spleen of newborn calves was not surprising. In rats B cell follicles are not apparent histologically before 2 weeks of age and GCs appear by 3–4 weeks of age (Dijkstra and Dopp, 1983), whereas GCs are not observed in human neonates until several months after birth (Namikawa et al., 1986).

Similar to human lymphoid tissues, immunostaining for CD32 was limited mainly to the mantle zone of the follicles suggesting that not all the IgM+ cells express CD32 (Tuijman et al., 1993). This finding correlates with flow cytometry data showing that only 75% to 83% of IgM+ cells express CD32 in newborn and older calves (Chattha et al., 2010b). The reduced expression of CD32 on centroblasts and centrocytes has been demonstrated by flow cytometry in the human tonsil GCs (Macardle et al., 2002). It has been hypothesized that reduced expression of CD32 may play an important role in selection and affinity maturation of B lymphocytes in GC reactions (Macardle et al., 2002). In the 7-month-old calves we observed minimal immunostaining for CD32 in the region of FDCs in the GC suggesting low-level expression of this receptor on FDCs. This receptor has been suggested to play a role in displaying immune complexes (Tew et al., 1997).

The absence of follicular dendritic cells in newborn calves is an important finding and may limit early GC reactions, including generation of antibody secreting cells and memory cells. In mice, adult equivalent patterns for FDCs appear by 3 weeks of age (Pihlgren et al., 2003) whereas mature FDC networks appear by 2 months of age in humans (Timens et al., 1989). Investigations in neonatal mice have shown that FDC precursors are unresponsive to B cell associated factors such as TNF-α, LT-α and LT-β, which are considered essential for maturation of FDC networks in adults (Fu and Chaplin, 1999; Pihlgren et al., 2003). The appearance of mature FDC networks in the lymph nodes of 7-week-old calves (data not shown) suggests appearance of these cells somewhere between a week and 2 months of age. Thus, early immunization may result in limited B cell responses in neonatal calves, whereas vaccination after this time is more likely to be efficacious.

The marker detected by FDC-B1 mAb is not known. There is a possibility that mature FDCs may exist in newborn calves with minimal or no expression of the particular antigen recognized by FDC-B1 mAb. However, we could not identify CD21+ cells with FDC morphology, indirectly suggesting that FDC were absent from the lymphoid tissues. Likewise the failure to observe organized follicles in lymphoid tissues can also be attributed to absence of mature FDCs in newborn calves. Interestingly, regions with FDCs also exhibited staining with anti-IgM mAb in 7-month-old calves (data not shown) indicating that IgM-antigen or IgM-complement-antigen complexes could be retained by FDCs. This retention may occur through either a complement receptor or the suggested presence of an Fc alpha/mu receptor (Fccα/μR) in cattle similar to humans (Kikuno et al., 2007).

In brief, the co-location of IgM+ and CD21+ cells is consistent with observations from our flow cytometry study, which found that the majority of the IgM+ cells in the lymphoid tissues of newborns express the activating receptor CD21, but a lower proportion express the inhibitory receptor CD32. Therefore, C3d tagged antigens might be effective in driving antibody responses in the neonate despite the presence of maternal antibodies. B cells with minimal expression or absence of CD32 occur in GCs. However, the low number of B lymphocytes and the minimal number of FDCs in the lymphoid tissues are important limitations for neonatal antibody responses following immunization. Discovering means to accelerate FDC maturation in newborns will be an important milestone on the road to effective neonatal vaccines.

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References


