Optimized Multicolour Immunofluorescence Panel for Cattle B Cell

Phenotyping by an 8-Colour, 10-Parameter Panel

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Abstract:

This 8-colour, 10-parameter panel has been optimised to distinguish between functionally distinct subsets of cattle B cells in both fresh and cryopreserved peripheral blood mononuclear cells (PBMCs). Existing characterised antibodies against cell surface molecules (immunoglobulin light chain (S-Ig(L)), CD20, CD21, CD40, CD71 and CD138) enabled the discrimination of 24 unique populations within the B cell population. This allows the identification of five putative functionally distinct B cell subsets critical to infection and vaccination responses; 1) naïve B cells (B_{Naïve}), 2) regulatory B cells (B_{Reg}), 3) memory B cells (B_{Mem}), 4) plasmablasts (PB) and 5) plasma cells (PC). Although CD3 and CD8α can be included as an additional dump channel, it does not significantly improve the panel's ability to separate

“Classical” B cells. This panel will promote better characterisation and tracking of B cell responses in cattle as well as other bovid species as the reagents are likely to cross react.

Key terms:

Flow cytometry; Cattle; B cells; antibody secreting cells; naïve B cells; memory B cells; regulatory B cells; B cell subsets

Background:

As our knowledge of immune cell subsets and their functions increases, so does the need to identify and measure alterations in their phenotype and frequency. The mammalian B cell population consists of several functionally distinct subsets that together comprise the major mediator of humeral immunity (1,2). The development of naïve B cells ($B_{\text{Naïve}}$) is important for long term immune protection (3–5). Driving the development of antibody secreting cells (ASC) and memory B cells (B_{Mem}) is an essential requirement of many vaccines that elicit neutralizing antibody responses (6–8). Furthermore, these subsets are often the source of therapeutic antibody candidates (as vaccines or immunotherapies) against infectious diseases (6–8). Regulatory B cells (B_{Reg}) also play a vital role in suppressing infectious diseases (9,10). Consequently, the identification and relative quantification of B cell subsets is

a fundamental requirement when evaluating pathogen or vaccine induced immune responses and ultimately the development of better strategies to control diseases (1).

The capability to dissect B cell responses at high resolution is limited in many non-model species through a combination of limited reagents, lack of knowledge of species-specific B cell markers and standardised methods (11). This is certainly the case for cattle, a key food producing species and crucial for human nutrition globally, as a universal B cell lineage marker (i.e. CD19) and reagents against other well-known B cell subsets (e.g. IgD and CD38) are lacking (12). As technologies to design and deliver protective immunogens continue to emerge rapidly, it is essential to evaluate their applicability in other species as part of one health approaches. Consequently, a need to study cattle B cell responses and their maturation at a high resolution.

We have developed a flow cytometry panel using existing antibodies against cell surface markers based on knowledge in humans and mice (13). With no pan B cell markers known in cattle, such as CD19, CD72 or CD79 α , we separated B cells from other lymphocytes using CD14 (CCG33, (14)) to exclude the monocytes, CD40 (IL-A158, (15)) as a B cell lineage marker, and included previously described cattle B cell markers such as CD21 (CC21, (16,17)) and surface immunoglobulin light-chain (S-Ig(L), IL-A58, (16,18)) (2,19). Subsets within these populations were further dissected

using the activation and differentiation markers CD71 (IL-A165, (20)), CD20 (MEM-97, (21)) and CD138 (recombinant-F1.20/A, (Personal comm. Washington State University)).

Based on well characterised human and mouse B cell populations we hypothesise that these markers will identify five major subsets of B cells in cattle lymphocytes (Online Table 3): B_{Naïve}, B_{Mem}, B_{Reg}, plasmablasts (PB) and plasma cells (PC) (13). The panel further allows for more in-depth characterisation of cattle B cells into 24 phenotypically unique subsets, following the gating strategy set out in Fig. 1; however, the functional discrimination and therefore importance between these subsets remains to be determined.

Our gating strategy consists of plotting CD40 against CD14 to select “classical” (CD40⁺CD14⁻) B cells. Although CD3 and CD8α are often used as a dump channel to isolate cattle B cells, their inclusion did not significantly improve separation (Online Figure 8). After identifying B cells, S-Ig(L) was plotted against CD21, allowing discrimination of four putative cattle B cell populations: CD21-S-Ig(L)⁺ and CD21⁺S-Ig(L)⁻ single positive (SP), CD21⁺S-Ig(L)⁺ double positive (DP), and CD21-S-Ig(L)⁻ double negative (DN) cells (Fig. 1 A). Next, each population was further sub-divided by comparing CD71 against CD20 and sub-gated into CD71⁺CD20⁻ SP, CD71⁺CD20⁺ DP and CD71⁻ populations (Fig.1 B). Lastly, each of these sub-gates were divided as

either CD138⁺ or CD138⁻ (Fig.1 C), resulting in 24 minor subsets of cattle B cells. An important step while labelling the PBMCs is to first stain the cells with the CD20 antibody before adding any of the other antibodies in the panel (Online Figure 9).

Although this panel was developed and optimised on a BD LSRFortessa, it performed equally well using a BD Aria IIIU when sorting cattle B cell subsets for further molecular investigation. The panel also has potential to be adapted by moving the CD14 antibody into the Live/Dead channel (for example coupled to APC-Cy7), provided the monocytes do not need to be specifically analysed, or into an available violet channel, which will free the PE channel for an additional marker if needed (e.g. intra-cellular staining). The CD71 antibody in APC can also be moved into an available violet channel. Further potential to expand and tailor this panel includes the addition of a cattle cross-reactive human CD27 antibody, which could resolve the limitations in identifying B_{Mem} cells or the addition of an antibody against CD5, which is known to identify B1 cells in mice (22).

By identifying functional subsets of B cells this panel has the potential to dramatically improve our understanding of cattle immune responses to infection and vaccination, moving towards addressing some of the problems detailed in both Entrican *et al.* and Barroso *et al.*, e.g. the lack of reagents to study the developmental cascade of cattle B cells (11,12). Additionally, the panel allows the enrichment or

isolation of specific single B cells or their populations to further study function, specificity, and drive antibody discovery.

Similarity to published OMIPs:

None to date.

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B Cell phenotyping in Cattle

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Figure 1 Gating strategy of the Cattle B cell panel into 24 minor subsets. A) The gating strategy followed from all events to the S-Ig(L) vs CD21 populations. B) S-Ig(L) vs CD21 further sub-gated into CD71 vs CD20 subsets for each of the four major subsets. C) The 24 minor subsets identified as the CD138⁺ and CD138⁻ from the three previous subsets in B. The parent population for each of the 24 minor subsets are listed above each gated population.

Table 1 Summary table for Optimized Multicolour Immunofluorescence Panel

Purpose	B cell phenotyping
Species	Cattle
Cell type	Fresh or cryopreserved PBMC, bone marrow, lymph node and tonsillar mononuclear cells
Cross reference	None

Table 2 Reagents used for Optimized Multicolour Immunofluorescence Panel

Specificity	Clone	Isotype	Fluorochrome	Manufacturer (Cat. No.)	Purpose	Dilution
Ig(L)	IL-A58	α-Mo IgG2a	DyLight 405	1°Ab ^a ITBX ^b (ITB00514) Kit ^c : abcam (ab201798)	B Cell lineage, B cell subset	1:3000
CD20	MEM-97	α-Mo IgG1	PE-Cy7	1°Ab Thermo Fisher Scientific (MA1-19008) Kit: Bio-Rad (LNK112PECY7)	B Cell development	1:500
CD21	CC21	α-Mo IgG1	FITC	1°Ab Bio-Rad (MCA1424GA) Kit: Bio-Rad (LNK061F)	B Cell lineage, B cell subset	1:4000
CD40	IL-A158	α-Mo IgG1	PerCP-Cy5.5	1°Ab Bio-Rad (MCA2431GA) Kit: Bio-Rad (LNK142PERCPCY5.5)	Co-stimulatory, B Cell lineage marker	1:2000
CD71	IL-A165	α-Mo IgG1	APC	1°Ab ITBX (ITB00261) Kit: Bio-Rad (LNK032APC)	Activation marker, Activated B Cells	1:1000
CD138	r(F1.20/A)	α-Mo IgG1	PE-Texas Red	1°Ab WSU ^d Kit: Bio-Rad (LNK172PETR)	Antibody secreting cells (Plasma cell and Plasmablasts)	1:1000
CD14	CCG33	α-Mo IgG1	PE	1°Ab Bio-Rad (MCA2678GA) Kit: Bio-Rad (LNK022RPE)	dump, Monocyte lineage marker	1:8000
CD3	MM1A	α-Mo IgG1	APC-Cy7	1°Ab Bio-Rad (MCA6080) Kit: Bio-Rad (LNK132APCCY7)	dump, T Cell lineage marker	1:100
CD8α	CC63	α-Mo IgG2a	APC-Cy7	1°Ab Bio-Rad (MCA837GA) Kit: Bio-Rad (LNK132APCCY7)	dump, T Cells, NK Cells	1:500
Amine-reactive	LIVE/DEAD Near-IR		APC-Cy7	Thermo Fisher Scientific (L10119)	Viability	1:2000

^aPrimary antibody.

^bImmunological Toolbox (23)

^cCommercial conjugation kit from Bio-Rad LYNX range or abcam Lightning-Link range.

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