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Regulation of metabolic supply and demand during B cell activation and subsequent differentiation

Takeshi Egawa¹ and Deepta Bhattacharya²

¹Department of Pathology and Immunology, Washington University School of Medicine, Saint Louis, MO 63110, U.S.A.

²Department of Immunobiology, University of Arizona College of Medicine, Tucson, AZ 85724, U.S.A.

Abstract

B cell activation and differentiation are associated with marked changes in proliferative and effector functions. Each stage of B cell differentiation thus has unique metabolic demands. New studies have provided insight on how nutrient uptake and usage by B cells are regulated by B cell receptor signals, autophagy, mammalian target of rapamycin, and transcriptional control of transporters and rate-limiting enzymes. A recurring theme is that these pathways play distinct roles ranging from survival to antibody production, depending on the B cell fate. We review recently published data that define how these pathways control metabolic flux in B cells, with a particular emphasis on genetic and *in vivo* evidence. We further discuss how lessons from T cells can guide future directions.

Introduction:

The primary role of B cells is to produce antibodies against infectious pathogens. To accomplish this seemingly simplistic goal, a highly regulated process is initiated to clonally expand relevant B cells, followed by a massive contraction to narrow the response to the minimal set of cells required to maintain protection against re-infection. In a conventional T cell-dependent (TD) response, naïve B cells are initially activated through cognate interactions between antigen and the B cell receptor (BCR). This triggers activation signals, internalization and processing of antigen to display to cognate CD4⁺ T cells, and co-stimulation through CD40:CD40L interactions. After a period of limited clonal expansion, B cells then differentiate either into plasma cells (PCs), germinal center (GC) B cells, or directly into IgM⁺ memory B cells [1]. PCs formed early in the response are localized to the extrafollicular regions and mostly live for a period of a few days [2]. GC B cells engage

Address correspondence to: Deepta Bhattacharya, deeptab@email.arizona.edu, Ph: 520-626-8088.

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multiple rounds of rapid proliferation coupled with isotype class switching, somatic hypermutation (SHM) of their immunoglobulin genes, and selection by CD4⁺ follicular helper T cells for high affinity variants [3]. Rare memory B cells and PCs emerge from the GC through selection processes that remain incompletely understood. Together, these cell types maintain long-term immunity against the offending pathogen and its relatives, sometimes for life [4]. The metabolic requirements of each of these stages differ [5,6]. We discuss below a number of recent studies that help explain how stage-specific metabolic requirements are communicated to B cells, and then how the resultant nutrients are used to support function.

Metabolic demands of initial B cell activation

Because of the difficulties in following the very early stages of B cell activation *in vivo*, most work in this area has come from *in vitro* studies. As much of the historical work has been reviewed recently [5], we focus instead on studies published within the past few years. Naïve B cells minimally proliferate and thus have relatively few bioenergetic requirements [7], such that lineage-determining transcription factors such as Pax5 and Ikaros suppress glucose uptake and ribonucleotide synthesis [8•,9]. BCR-activated B cells must thus markedly change their metabolic profiles to expand, proliferate, and generate ATP, much of which is driven by enhanced glucose uptake through the transporter GLUT1 [10–12]. BCR-induced metabolic changes are dependent on PKC β , which is required for the metabolic switch to glycolysis and for downstream mTOR activation, mitochondrial function, and heme biosynthesis [11,13].

As with T cells, antigen receptor-stimulated B cells use second signals such as Toll-like receptor- or CD40-engagement to become fully activated, thereby reducing the chances of autoimmunity. In the absence of these secondary signals, B cells accumulate mitochondrial calcium and produce excessive amounts of superoxides, eventually leading to activation-induced cell death [14]. Costimulatory signals such as TLR activation enhance uptake of glucose and glutamine, which are used as carbon sources for the pentose phosphate pathway to support proliferation, *de novo* lipogenesis to expand the endoplasmic reticulum [15], and for mitochondrial respiration and anaplerosis [16]. Nutrient uptake and the extent of B cell activation is tempered by the expression of the let-7 microRNA cluster, which suppresses glucose and glutamine transporter expression. In the absence of let-7, excessive antibodies are produced in response to T-independent antigens *in vivo*. These studies collectively provide a valuable baseline that begins to explain how B cells integrate antigen-specific signals with metabolic programs to support subsequent differentiation (Figure 1).

Germinal center metabolism:

Following initial T-dependent activation, antigen-specific B cells and CD4⁺ T cells form GCs, which are specialized anatomical structures located within the centers of B cell follicles. GCs are divided into two distinct areas, light- and dark-zones (LZ and DZ), in which B cells exist in distinct proliferative and metabolic states [17,18]. In the LZ, B cells are mostly in a non-proliferative state where they compete with one another for CD4⁺ T cell help by presenting peptide on MHC class II. Only a small fraction of LZ B cells are selected

by T cells to enact a proliferative program as they migrate to the DZ to divide for several days. These proliferating B cells are the templates for AID-mediated SHM, and are thus critical for the emergence of protective antibodies with affinity-enhancing mutations. This proliferative program is enabled by an anabolic response driven by the transcription factor c-MYC, which not only regulates cell cycle progression but also promotes expression of transporters of nutrients and enzymes for glycolysis and glutaminolysis [19–21]. GC B cells require both BCR and CD40 signals for c-MYC induction, thus enforcing a stringent selection checkpoint dependent on CD4⁺ T cell help [22]. As a counterexample, either BCR or CD40 engagement is sufficient for naive B cells to induce c-MYC and proliferation. Despite the absolute requirement for c-MYC in GC B cells, its expression is restricted to a small fraction of LZ B cells and becomes undetectable in proliferating DZ B cells [17,20,21]. This paradox is explained by a MYC-induced transcription factor AP4, which maintains many c-MYC target genes, including nutrient transporters and metabolic enzymes after downregulation of c-MYC [23•]. Activity of mTORC1 correlates well with engagement of the MYC-AP4 pathway. mTORC1 is activated upon selection of GC B cells in the LZ and is required for cell growth, sustained expansion, and dwelling of selected B cells in the DZ [24••]. Thus, c-MYC-dependent and mTORC1-centered metabolic programs, both of which are activated by T cell help, independently or synergistically support durable proliferation of DZ B cells.

While cell-intrinsic programs triggered in selected GC B cells by T cell-derived help enact metabolic pathways to meet the demands of DZ B cell proliferation, environmental factors specific to the GC microenvironment also modulate GC B cell differentiation. One such example is restriction of mTORC1 activation and AID expression in LZ B cells by hypoxia [25]. Hypoxic conditions in the LZ induce elevated activity of HIF transcription factors. HIFs in turn inhibit activity of mTORC1 and expression of AID, and thereby restrict proliferation of GC B cells and isotype class switching [25]. As a result, overexpression of HIFs through genetic deletion of their inhibitor VHL leads to diminished GC responses. Given that oxygen availability is normally controlled by proximity to blood vessels, it is unclear how hypoxia is preferentially induced in LZ vs. DZ cells, or whether the reciprocal deletion of HIF transcription factors would be sufficient to cause exaggerated GC responses. The enzyme glycogen synthase kinase 3 (GSK3) is also important for GC B cell responses. Through restriction of c-MYC, GSK3 protects proliferating B cells from deprivation of nutrients, such as glucose and amino acids [26]. Since GCs contain a high density of activated cells with elevated glycolytic and mTORC1 activity, GC B cells may have adapted or been desensitized to nutrient deprivation in the microenvironment through GSK3-mediated pathways.

The engagement of autophagy is another notable metabolic program in GC B cells. Compared to resting B cells, autophagy is upregulated in GC B cells, which is consistent with the demand for nutrients to support rapid proliferation [27]. Interestingly, B cells lacking WIPI2, a component in the autophagy pathway, differentiate preferentially to ASCs at the expense of GC B cells, and this altered B cell differentiation is associated with a specific increase in non-canonical autophagy and elevated mitochondrial activity. Although the detailed mechanism is unknown, a shift between distinct autophagy pathways or

autophagy-independent function of WIPI2, may control the checkpoint between GC B cells and PCs. These recent findings are collectively summarized in Figure 2.

Metabolic programs to support plasma cells and memory B cells:

The major goal of the GC reaction is to output affinity-matured memory B cells and antibody-secreting PCs, the principal effectors within the B cell lineage. A model of LPS-induced differentiation *in vivo* has shown that PCs are formed by a series of transcriptional changes downstream of the canonical PC transcription factor BLIMP1 and EZH2 [28–30]. These transcriptional changes are division-dependent and related to ER stress genes, oxidative metabolism enzymes, and amino acid transporters, thereby leading to mTORC1 activation and enhanced antibody secretion [31,32•]. Other studies have shown that autophagy is also activated and required to regulate ER stress responses, antibody secretion, and energy metabolism [33].

In general, PCs formed early in the response tend to be relatively short-lived, while those formed later from the GC reaction are long-lived [34•]. Yet throughout the immune response, PCs of varying lifespans are generated, demonstrating a far greater complexity belied by the simple short- vs. long-lived designation [35]. Indeed, different vaccines and infections lead to a wide range of durations of antibody production [4], though what distinguishes the underlying PCs remains a mystery. Moreover, while it has generally been assumed that T cell-independent responses fail to generate long-term immunity, many studies have conclusively demonstrated that this is not the case. In fact, through activation of B-1 cells, T-independent responses robustly yield persistent PCs that constitutively secrete antigen-specific IgM [36–39]. Recent studies have demonstrated that PC subsets of varying lifespans and antibody secretion rates can be distinguished based on uptake of a fluorescent glucose analog, surface expression of amino acid transporters, and autophagic markers [35,40•]. PCs with relatively long lifespans import more glucose and predominantly use it to glycosylate antibodies [40•], consistent with their single-minded focus on antibody production. Upon bioenergetic shortages, this glucose can be diverted away from glycosylation to glycolysis and the resultant pyruvate transported into the mitochondria to restore homeostasis [40•]. Similarly, long-lived PCs use glutamine both as a building block for antibodies and for mitochondrial respiration and anaplerosis [35]. These data suggest a link between immunoglobulin production and bioenergetic metabolism, and indeed, long-lived PCs secrete antibodies at a greater rate than do their short-lived counterparts [35]. Yet surprisingly, very few ER stress responses or transcriptional changes are associated with these profound metabolic differences [35,41]. These data demonstrate that there is no obligatory link between transcriptional regulation and metabolism, and some caution is warranted when such assumptions are made (Figure 3).

Metabolic support of memory B cells and recall responses:

Unlike their memory T cell counterparts, which have been the subject of many metabolic studies, only a few reports have described properties of memory B cell metabolism. An elegant set of studies have shown that memory B cells lacking the autophagy gene ATG7 form normally, but fail to persist after immunizations with haptens or infections with

influenza [42,43]. As a result, recall responses are severely diminished. ATG7 protects memory B cells against an unusual form of death related to oxidative stress rather than conventional apoptosis or RIPK1-driven necrosis. Another recent study showed that upon stimulation with TLR ligands and IFN α , human IgD⁺ memory B cells rapidly become glycolytic, activate mTORC1, and differentiate into plasmablasts to a greater extent than do naïve B cells [44]. Whether these differences between memory and naïve B cells reflect an intrinsically distinct metabolic wiring or simply the magnitude of TLR signaling remains to be determined.

Future directions: Lessons from T cells

Studies of B cell metabolism lag behind those of their T cell cousins. Yet this provides opportunities for analogous concepts to be adopted and tested while avoiding some of the pitfalls already encountered by other immunometabolism studies. Some of these issues are technical. Inhibitors such as etomoxir, which antagonizes long-chain fatty acid oxidation, are plagued by off-target effects [45], resulting in inconsistencies between genetic and pharmacological studies [46,47]. Yet relying on genetic studies alone can also be problematic, as deletion of genes such as Cpt1a can be compensated by the paralogs Cpt1b and Cpt1c, leading to incomplete ablation of fatty acid oxidation [46]. Choosing non-redundant genes such as Cpt2 that are essential to a pathway to perform genetic studies [48], and carefully titrating pharmacological inhibitors are the best ways to avoid such issues as the B cell immunometabolism field moves forward.

A key biological question that has emerged in the T cell field is whether metabolic pathways are supportive of a given lineage, as opposed to instructive, which implies the suppression of alternate fates. In most contexts, the evidence suggests that metabolic programs play supportive roles. For example, memory T cells normally depend on fatty acid oxidation and spare respiratory capacity for formation and function [47], but functional cells can also be generated by maintaining glycolysis after the effector stage [49]. Thus, a specific metabolic pathway is required for neither enforcing memory T cell fate nor for extinguishing the effector lineage. In most cases we suspect a supportive role will be the dominant metabolic mechanism in B cells as well. For example, although mTORC1 activation assists in the expression of the germinal center-determining transcription factor BCL6 [50], this pathway is activated in plasma cells as well [32]. Nonetheless, if anything is to be gleaned from recent years of immunometabolism research, it is to expect the unexpected.

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Highlights:

- -Nutrient uptake by naïve B cells is limited by Pax5 and Ikaros
- -Antigen and second signals promote B cell nutrient uptake and metabolic rewiring
- -Germinal center B cells use c-myc and mTOR for their proliferative burst
- -Plasma cell longevity and antibody secretion are defined by metabolic pathways

Figure 1: Metabolic changes following initial B cell activation.

Resting naïve B cells import relatively little glucose, in part due to repression by the transcription factors Pax5 and Ikaros. After ligation of the BCR and co-ligation of either CD40 or Toll-like receptors (TLR), an influx of glucose and glutamine is triggered. Accompanying this elevated nutrient import, the size of the cell expands, mitochondrial mass increases, and transcriptional changes are enacted to support metabolic function.

Figure 2: Metabolic and transcriptional regulation of germinal center B cell response.

B cell clones that express BCRs with higher affinity than neighboring clones are selected by follicular helper T cells. The selected clones upregulate transcription factors c-MYC and AP4 and activate mTORC1 to support their rapid proliferation, which may be modulated by local environmental factors, such as hypoxia or nutrient deprivation.

Figure 3: Metabolic, but not transcriptional changes distinguish short-from long-lived plasma cells.

Short-lived plasma cells import less of a fluorescent glucose analog and amino acids than do long-lived plasma cells. However, these cell types are highly similar in ER stress responses and transcriptional profiles. Adapted from [35].