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The CD47-SIRP α immune checkpoint

Meike E.W. Logtenberg¹, Ferenc A. Scheeren², Ton N. Schumacher^{1,3,4}

¹Division of Molecular Oncology and Immunology, Oncode Institute, The Netherlands Cancer Institute, Amsterdam, The Netherlands ²Department of Medical Oncology, Leiden University Medical Center (LUMC), Leiden, Netherlands ³Department of Immunohematology and Bloodtransfusion, Leiden University Medical Center, Leiden, The Netherlands

Abstract

The cytotoxic activity of myeloid cells is regulated by a balance of signals that are transmitted through inhibitory and activating receptors. The Cluster of Differentiation 47 (CD47) protein, expressed on both healthy and cancer cells, plays a pivotal role in this balance by delivering a “don’t eat me signal” upon binding to the Signal-regulatory protein alpha (SIRP α) receptor on myeloid cells. In this Perspective, we describe the current state of knowledge on the role of the CD47-SIRP α axis in physiological tissue homeostasis and as a promising therapeutic target in, amongst others, oncology, fibrotic diseases, atherosclerosis and stem cell therapies. In addition, we highlight where additional insight will be beneficial to allow optimal exploitation of this myeloid cell checkpoint as a target in human disease.

Introduction

The immune system has evolved to detect and destroy target cells that are diseased, such as pathogen-infected or cancer cells that would potentially be harmful when not inactivated. In addition, the immune system plays a central role in tissue homeostasis by removing unnecessary cells or, in some cases, by modifying their interactions with surrounding cells. Target cell clearance or modification mediated by myeloid effector cells, including macrophages, neutrophils and microglia, is regulated by receptors that either transmit activating or inhibitory signals upon ligand binding. The latter class of inhibitory receptors, also known as immune checkpoints, includes a series of immunoreceptor tyrosine-based inhibitory (ITIM)- and immunoreceptor tyrosine-based switch motif (ITSM)-containing receptors such as SIRP α , Leukocyte Immunoglobulin-like Receptor B1 (LILRB1), Siglec-10 and Programmed-cell Death Protein 1 (PD-1), which can bind CD47, MHC class I, CD24 and Programmed Death Ligand-1 (PD-L1) on target cells, respectively (Barkal et al., 2019; 2017; Gordon et al., 2017; Oldenborg et al., 2000).

⁴To whom correspondence should be addressed at t.schumacher@nki.nl.

Author contributions

M.E.W.L. researched data for the article, M.E.W.L., F.A.S and T.N.S jointly discussed data and co-wrote the article.

Competing interests

M.E.W.L., F.A.S., and T.N.S. are inventors of IP related to blockade of the CD47-SIRP α axis, T.N.S. is advisor to and shareholder in Scenic Biotech.

Work over the past decades has demonstrated that the CD47-SIRP α interaction regulates the outcome of myeloid cell-target cell crosstalk in a number of homeostatic processes. In particular, signaling through the CD47-SIRP α axis has been shown to influence erythrocyte, platelet and hematopoietic stem cell (HSC) maintenance, and to regulate synaptic pruning during neuronal development (Jaiswal et al., 2009; Lehrman et al., 2018; Lutz and Bogdanova, 2013; Yamao et al., 2002). Next to an important role in these homeostatic processes, CD47 expression on cancer cells has been shown to inhibit myeloid cell-mediated elimination in a manner that bears a superficial resemblance to the inhibition of intratumoral T cell activity by the PD-1/PD-L1 immune cell checkpoint (Sun et al., 2018).

Based on its role as a regulator of tumor cell fate, a series of molecules that block the CD47-SIRP α axis is currently in clinical development in oncological indications, with an encouraging clinical signal in some settings. In addition, a number of preclinical studies have provided compelling evidence for the potential value of the targeting of this axis in stem cell transplantation, and possibly also diseases such as atherosclerosis and fibrosis (Kojima et al., 2016; Wernig et al., 2017)

In this Perspective, we discuss the structural and functional properties of the CD47-SIRP α axis and highlight the deficits in our understanding of the, possibly context-dependent, mechanisms that regulate CD47-SIRP α signaling strength and that can be expected to influence the activity and toxicity of therapeutic targeting.

Molecular wiring of the CD47-SIRP α axis

The CD47 protein (also known as IAP, MER6 or OA3) consists of a single extracellular V-set IgSF domain, a presenilin domain with five membrane-spanning sections and a short cytoplasmic domain that is subject to alternative splicing, thereby giving rise to four isoforms (Campbell et al., 1992; Mushegian, 2002; Reinhold et al., 1995). In addition to binding SIRP α and its family member SIRP γ , CD47 also interacts with integrins and secreted glycoprotein extracellular matrix protein thrombospondin-1 (TSP-1) (reviewed in (Brown and Frazier, 2001) and (Oldenborg, 2013)). In contrast to the more restricted expression pattern of SIRP α (see below), CD47 is broadly expressed across different cell types in the body, with varying expression of the CD47 protein isoforms in different tissues (Reinhold et al., 1995).

The SIRP family of proteins consists of five members; SIRP α , SIRP β 1, SIRP β 2, SIRP γ and SIRP δ . Two of these, SIRP α (also known as PTPNS1, SHPS1, CD172A and P84) and SIRP γ (also known as CD172b), are known to bind CD47. SIRP α contains an extracellular region with three immunoglobulin superfamily (IgSF) domains, including a NH₂-terminal ligand binding V-domain. Allelic variants with polymorphisms in the ligand binding domain have been reported in the African, Japanese, Chinese and Caucasian populations, three of which (SIRP α V1, SIRP α V2 and SIRP α V8) are the most prominent haplotypes among the human population, jointly covering approximately 90% (Takenaka et al., 2007; Voets et al., 2019). The observation of significant polymorphism, including alleles that are substantially divergent in the ligand binding domain, is suggestive of evolutionary pressure. However, the observations that the polymorphic residues in the NH₂-terminal domain of SIRP α are distant

from the CD47 binding site, and do not influence CD47 binding capacity for the at least 5 variants for which this was tested, argues against selective pressure at the level of CD47-SIRP α signaling strength (Hatherley et al., 2014). The intracellular region of SIRP α contains both ITIM and ITSM motifs that are essential for inhibitory activity of the receptor (Fujioka et al., 1996; Kharitononkov et al., 1997). The extracellular domain of SIRP γ is similar to SIRP α , but binds to CD47 with a ten-fold lower affinity (K_D ~23 μ M for SIRP γ versus ~2 μ M for SIRP α) (Brooke et al., 2004). The cytoplasmic domain of SIRP γ consists of only 4 amino acids without an apparent signaling function (Brooke et al., 2004; Ichigotani et al., 2000).

SIRP α is expressed on all myeloid cell types, including monocytes, macrophages, neutrophils, a subset of dendritic cells and microglia (Adams et al., 1998; Seiffert et al., 1999; Veillette et al., 1998). In addition, SIRP α expression has also been observed in brain tissue (Uhlén et al., 2015) (<http://www.proteinatlas.org>), and on a subset of CD8⁺ T cells during chronic infection (Myers et al., 2019). Signaling through ITIM- or ITSM-containing receptors such as SIRP α serves to counteract the cellular activation that occurs when an activating – frequently an ITAM-containing – receptor is triggered. In other words, in the absence of such an activating signal, triggering of ITIM/ITSM receptors is considered a null event. Inhibition of signaling through activating immune receptors by SIRP α engagement requires the phosphorylation of the tyrosine residues within the cytoplasmic ITIM and ITSM sequences, resulting in the recruitment and activation of the SH2-domain-containing protein tyrosine phosphatases SHP-1 and SHP-2 (Fujioka et al., 1996; Kharitononkov et al., 1997). In addition, some evidence has been obtained for association of SIRP α with the inhibitory protein kinase Csk and, to a lesser extent, the Grb-2 adaptor molecule (Veillette et al., 1998). Although our understanding of the direct downstream targets of the SHP-1 and -2 phosphatases upon SIRP α signaling is likely to be incomplete, deactivation of motor protein myosin IIA, and decreased cytotoxic activity of myeloid effector cells, have been shown as downstream consequences of SIRP α ligation (Tsai and Discher, 2008). Over the past years, SIRP α signaling has been shown to counteract signals that myeloid cells receive through a variety of activating membrane receptors, including 1) Fc receptors (FcRs) that bind the Fc domain of target-opsonizing antibodies, 2) complement receptors, and 3) the lipoprotein-related protein (LRP) that binds calreticulin (CRT) (Chao et al., 2010b; Oldenborg et al., 2001). As a side note, the fact that FcRs are one class of activating receptors of which the activity is inhibited by SIRP α ligation implies that studies that utilize CD47 blocking molecules with intact FcR binding capacity should be interpreted with caution, as the intervention simultaneously removes an inhibitory signal and introduces an activating signal (see further below) (Ingram et al., 2017; Weiskopf et al., 2013).

Cell types under control of the CD47-SIRP α axis

Early work has provided compelling evidence that the CD47-SIRP α axis can regulate the activity of a variety of myeloid cell types. For example, a role for macrophages in tumor control has been shown with anti-CD47 mouse IgG1 blocking antibodies in various NSG cancer models that lack components of the adaptive immune system (Chao et al., 2010a; Majeti et al., 2009). In the case of neutrophils, antibody-dependent cellular cytotoxicity (ADCC) towards target cells has been shown to be enhanced upon pathway blockade *in vitro*

(Zhao et al., 2011). In addition, in a human Fc alpha receptor (FcαR) transgenic mouse model, the tumor cell killing that was induced by IgA1 anti-Her2 antibodies and that was boosted by pathway inhibition was primarily dependent on neutrophils (Logtenberg et al., 2019).

Over the past years, a growing body of evidence has emerged that suggests that the adaptive immune system, and in particular CD8+ T cells, can also contribute to the tumor control that is seen upon SIRPα-CD47 pathway inhibition. This observation may either be explained by a direct influence of the SIRPα-CD47 pathway on T cell activity, or by an indirect mechanism in which pathway blockade influences the capacity of myeloid cells to boost CD8+ T cell reactivity. In an adoptive transfer model, ovalbumin (OVA)-specific CD8+ T cells displayed enhanced killing capacity towards OVA-positive targets upon injection of macrophages that had been co-cultured with antigen-expressing target cells in the presence of anti-CD47 IgG (Tseng et al., 2013). These data suggest that the increased tumor control that is seen upon *in vivo* CD47 blockade may also in part be explained by improved APC function. In another study, enhanced antigen-presentation function of DCs, rather than macrophages, was proposed to be required for the observed anti-tumor effects of low dose intratumoral anti-CD47 treatment in syngeneic mouse models, as depletion of either CD11c+ cells or CD8+ T cells severely impaired the therapeutic effect (X. Liu et al., 2015). In a study by Sockolosky et al., anti-PD-L1 treatment synergized with antibodies targeting CD47 and the mouse melanoma antigen tyrosinase-related protein 1 (TRP-1), an observation that was also interpreted as evidence for a dominant role of T cells upon CD47 blockade, possibly via increased DC acquisition of CD47-opsonized tumor cells and enhanced antigen presentation (Sockolosky et al., 2016). As a side note, PD-1 is expressed in myeloid cells in tumor-bearing mice, and the observed synergy between PD-L1 and CD47 blockade may in theory also be explained by a role of PD-1 as an inhibitory receptor in myeloid cells, as shown in recent mouse experiments (Strauss et al., 2020). Finally, work by Li et al. has demonstrated that vaccination with CD47-deficient tumor cells induced expansion of CD11c+ DC, and that therapy-induced tumor control was abrogated upon CD11c+ cell depletion (Li et al., 2020).

While the observations in the above studies are likely to be explained by enhanced antigen-presentation upon pathway blockade, and hence do not provide substantial evidence for a direct effect of this pathway on T cell function, several other studies do make a somewhat stronger case for such a model. First, in a pooled genetic screen, CD47 deletion was shown to increase the sensitivity of melanoma cells to GVAX/ anti-PD-1 combination immunotherapy, and this effect was dependent on T cells (Manguso et al., 2017). It may be argued that this sensitizing effect of CD47 loss could reflect increased activation of myeloid cells and a resulting boosting of T cell activity. However, this would then have to assume a highly localized regulation of T cell activity by myeloid cells, as selective loss of CD47 negative cells in a sea of CD47 positive cells would otherwise not be expected. As a second 'myeloid centric' explanation, one could propose that while T cell activity is critical for tumor control, ultimate tumor cell killing may not be due to T cell activity but due to myeloid cell effector function. As an alternative explanation for the observed effect of CD47 loss in this study it may be proposed that the CD47-SIRPα axis may not only regulate T cell function by influencing APCs but can also influence T cell activity directly. Some support

for such a model comes from the observation that SIRP α is induced on a subset of virus-specific CD8⁺ T cells in mouse models of chronic viral infection and, while restricted to a smaller T cell subset, parallels PD-1 expression (Myers et al., 2019).

Regulation of tissue homeostasis and tissue remodeling by the CD47-SIRP α axis

The CD47-SIRP α axis regulates homeostatic processes by controlling myeloid cell-mediated removal of ageing cells, erythrocytes, HSCs and neuronal synapses (Figure 1). The first evidence demonstrating a protective role for CD47 came from *in vivo* experiments that showed rapid elimination of CD47-deficient erythrocytes by splenic macrophages when transferred into wild-type recipients (Oldenborg et al., 2000). Furthermore, increased clearance of CD47-deficient erythrocytes was abrogated in *SIRPA*-mutant mice lacking most of the SIRP α intracellular signaling domain (Ishikawa-Sekigami, 2006), thereby implying a role for both components of the CD47-SIRP α axis in the process. Over the past years, a number of factors have been identified that may influence the susceptibility of erythrocytes to macrophage-mediated elimination. Work by Khandelwal et al. has demonstrated that aged erythrocytes show a reduced CD47 expression (Khandelwal et al., 2007). In addition, increased display of a number of activating signals, such as increased phosphatidylserine (PS) exposure and auto-antibody binding to band 3 (Boas et al., 1998; Kay, 1984; Lutz and Bogdanova, 2013), might also contribute to the enhanced clearance of aged erythrocytes. The latter type of auto-antibody binding might also be of relevance in Autoimmune Hemolytic Anemia (AIHA), a syndrome characterized by accelerated clearance of erythrocytes (Barcellini, 2015), and spontaneous AIHA in nonobese diabetic (NOD) mice is substantially accelerated by CD47 deficiency (Oldenborg et al., 2002; 2001). In line with these data, SIRP α signaling has been shown to inhibit Fc γ - or complement receptor-dependent clearance of antibody- and complement-opsonized erythrocytes by macrophages (Oldenborg et al., 2001).

CD47 levels also play a role in the regulation of platelet turnover (Olsson et al., 2005) and in regulating the fate of circulating hematopoietic stem cells. Circulating human HSCs have been documented to express higher CD47 levels as compared to bone-marrow resident HSCs, and CD47-deficient c-Kit⁺ HSCs are phagocytosed more efficiently by macrophages *in vitro*. Furthermore, in a parabiosis model in which the circulatory systems of wild-type and CD47-deficient mice were surgically joined, bone marrow HSC chimerism was observed in the CD47 deficient animals but not in the wild-type animals, suggesting that CD47-deficient HSCs are cleared from the blood compartment *in vivo* (Jaiswal et al., 2009). In patients with Hemophagocytic Lymphohistiocytosis (HLH), a syndrome characterized by increased phagocytosis of HSCs by bone marrow macrophages, CD47 expression on the CD34⁺CD38⁻ HSC fraction is decreased relative to healthy subjects, reaching levels that are insufficient to prevent macrophage-mediated phagocytosis (Kuriyama et al., 2012). Activating signals on HSCs from healthy subjects or patients with HLH that promote this phagocytic process have not been identified to date, although CRT does not appear to be involved (Kuriyama et al., 2012).

As a final example of the role of CD47 as a physiological regulator of cell survival and cellular interactions, expression of CD47 has been shown to protect neuronal synapses from elimination during “synaptic pruning”, a developmental process in which microglia remove synaptic elements to form an organized brain circuitry (Lehrman et al., 2018; Paolicelli et al., 2011). Complement components such as C1q and C3 localize to developing synapses to promote the microglial engulfment of synaptic inputs that is necessary for refinement, in a microglial C3 receptor (CR3 or Mac-1) dependent manner (Schafer et al., 2012; Stevens et al., 2007). Microscopic analysis of coronal sections during the peak of synaptic pruning has demonstrated that CD47 protein is enriched in the dorsal geniculate nucleus (dLGN) of the thalamus and that SIRP α is highly expressed by microglia. Notably, CD47- and SIRP α -deficient mice display increased microglial engulfment of retinal ganglion cell (RGC) inputs and enhanced synaptic pruning, leading to a sustained reduction in the number of synapses. Interestingly, microglia are known to preferentially engulf less active synaptic inputs, and the observed preferential localization of CD47 to active inputs suggests a mechanism in which local inhibitory ligand expression is used to convey activity-related information to microglia (Lehrman et al., 2018).

Next to its role in tissue homeostasis, there is emerging evidence for a role of the CD47-SIRP α axis in tissue remodeling in fibrotic diseases (Wernig et al., 2017), and in atherosclerosis (Kojima et al., 2016). The fibrotic response is a normal component of tissue repair processes, however this response can lead to life-threatening conditions when uncontrolled, such as in idiopathic pulmonary fibrosis (IPF), primary myelofibrosis and scleroderma. In lung samples obtained from patients with idiopathic lung fibrosis, CD47 was highly expressed on fibroblasts, whereas calreticulin was found to be expressed on macrophages and a subset of bronchoepithelial cells. Furthermore, fibrosis was shown to be reduced by anti-CD47 treatment in mouse models of fibrotic disease (Wernig et al., 2017). In the development of atherosclerosis, impaired cell clearance leads to the formation of pathogenic plaques that are composed of diseased vascular and apoptotic cells. As such, atherosclerotic lesions are at risk of rupture and obstruction, increasing the possibility of myocardial infarction and stroke (Kasikara et al., 2018; Schrijvers et al., 2005). CD47 was found to be upregulated in human atherosclerotic plaques as compared to healthy vascular tissue, possibly due to the enhanced TNF α levels in these lesions, and treatment with CD47 blocking antibodies was shown to reduce atherosclerosis formation *in vivo* (Kojima et al., 2016). In both these studies, CD47 blockade involved the use of antibodies with Fc γ -receptor activating capacity, and in the work by Kojima et al., increased erythrocyte clearance and compensatory reticulocytosis was observed. In view of the known side-effects of FcR binding anti-CD47 antibodies, this appears to be an issue that deserves further attention.

Regulation of cancer cell fate by the CD47-SIRP α axis

CD47 expression is frequently observed on cancer cells in both hematological and solid malignancies, including non-Hodgkin lymphoma (NHL), acute myeloid leukemia (AML), glioblastoma, ovarian, breast, colon, bladder, hepatocellular, and prostate cancer (Chao et al., 2010a; Majeti et al., 2009; Willingham et al., 2012). While one study has reported copy number alterations in the *CD47* gene in 5% (15 of 316) of patients with ovarian serous

cystadenocarcinoma (Brightwell et al., 2016), in most human tumors CD47 levels are only modestly increased relative to surrounding healthy tissue (Willingham et al., 2012; Zhao et al., 2011). The lack of frequent overexpression of CD47 on tumor cells could either imply that the strength of myeloid cell inhibition is not boosted by a further increase in CD47 expression, or that the axis is not a substantial modifier of cancer cell fate during natural disease progression, an issue that requires further study.

As will be discussed in the next section, the CD47-SIRP α axis does form a critical regulator of tumor cell fate at the moment tumor cells are decorated with activating signals as a consequence of therapeutic intervention. In addition, cellular changes associated with malignant transformation may in some cases by themselves already lead to the expression of ligands for activating immune receptors by cancer cells. Only in such situations, where an endogenous activating signal is present, single agent blockade of CD47 is likely to suffice to induce tumor cell killing, and hence it is important to understand the nature and prevalence of such activating signals (Figure 1). As a first example of such an endogenous activating signal, CRT, the ligand of the myeloid cell receptor LRP-1, is present at high levels on several human cancers, and blockade of CRT binding to LRP can prevent the *in vitro* phagocytosis of certain cancer cell lines that is induced by anti-CD47 antibodies (Chao et al., 2010b). Of note, the CRT pool at the tumor cell surface need not originate from an endogenous CRT pool that translocates to the cell surface (Garg et al., 2012). Specifically, CRT has been shown to be secreted by macrophages in the tumor microenvironment, and secreted CRT can bind cancer cells that display specific patterns of membrane glycans (Feng et al., 2018). A similar binding of extracellular CRT has been described as a mechanism behind the clearance of ageing neutrophils (Feng et al., 2018). Next to CRT, myeloid cells may recognize other activating signals on cancer cells. Specifically, on cancer cell lines and hematopoietic cells that were phagocytosed equally efficient by wild-type and LRP-1-deficient macrophages upon CD47 blockade, the interaction between myeloid cell-expressed signaling lymphocytic activation molecule family 7 (SLAMF7; also known as CRACC, CS1 and CD319) and myeloid cell-expressed Mac-1 was shown to provide an activating signal (Chen et al., 2017). As SLAMF7 is a homotypic receptor, it was proposed that binding of myeloid cell-expressed SLAMF7 to SLAMF7 on cancer cells would be required for cancer cell recognition and phagocytosis, however this model has recently been contradicted (He et al., 2019). Additional evidence for the presence of endogenous activating signals on cancer cells comes from the increase in *in vitro* phagocytosis of cancer cell lines that is induced by anti-CD47 Fab fragments (Chao et al., 2010a). It is noted though that in these experiments, phagocytosis was significantly enhanced when anti-CD47 treatment was combined with Rituximab, suggesting that the endogenous activating signal is weak relative to the activating signal induced by antibody opsonization. In line with the notion that endogenous activating signals are frequently insufficient to allow significant CD47 blockade monotherapy activity, treatment with anti-CD47 F(ab')₂ fragments only increased neutrophil-mediated *in vitro* killing of tumor cells in the presence of anti-Her2/Neu antibody (Zhao et al., 2011). Furthermore, treatment with a high affinity SIRP α monomer only slowed lymphoma growth slightly and had no effect on breast cancer growth *in vivo*, whereas the combination with rituximab (in the case of lymphoma) or trastuzumab (in the case of breast cancer) resulted in a major reduction in tumor growth (Weiskopf et al., 2013). By the same token, even

localized secretion of CD47 blocking agents that lacked Fc domains was shown not to affect growth of melanoma tumors *in vivo* (Ingram et al., 2017), and genetic deletion of CD47 only minimally affected tumor growth in a B16 melanoma model (Barkal et al., 2017). Finally, also treatment with anti-SIRP α antibodies has been shown to be insufficient to induce myeloid cell-dependent tumor cell killing (Ring et al., 2017; Sim et al., 2019). Collectively, these data make a rather compelling case that the fraction of human tumors that intrinsically carry a sufficiently strong activating signal is limited, or that additional inhibitory signals are present on these cells that prevent the activity of single agent SIRP α -CD47 blockade. Future studies in this area would benefit substantially from efforts that would allow one to not only measure CD47 levels, but to also measure the presence of activating signals. Much like the presence of antigen can be used to predict the efficiency of T cell recognition at the moment inhibitory receptors are blocked, the quantification of molecules and/or tumor cell states that report on the presence of activating myeloid cell ligands could be used to improve our capacity to predict the consequence of CD47 pathway inhibition.

Regulation of CD47-SIRP α signaling strength

The importance of the CD47-SIRP α checkpoint in tissue homeostasis and remodeling, and its potential value in cancer therapy, are both evident. In view of this, it is surprising that relatively little is known on the parameters that regulate the strength of this signaling axis. As a general thought, an inhibitor of immune cell activation that would provide the same negative signal under all circumstances would not seem to provide any means of regulation, and simply making the activating receptor less sensitive would appear a more straightforward solution. For this reason, the presence of inhibitory receptor systems is likely to indicate a need to provide contextual information, and two conceptually different classes of such regulators may be distinguished. 1) In a “differential threshold system” a given target cell type or tissue may always have the same expression level of an inhibitory receptor ligand, and a given immune cell type may always have the same level of expression and signaling capacity of the corresponding inhibitory receptor. In such a setting, the same activating signal will lead to a differential response when present at a site that either shows low or high expression of the inhibitory receptor ligand. By the same token, the activating signal present on a given target cell type may lead to differential activation of distinct immune cell types, depending on their inhibitory receptor expression. In other words, in this setting, the expression level of inhibitory receptors and their ligands does not show dynamic regulation, but rather serve as a code to control the outcome of different cellular interactions. 2) On the contrary, in a “dynamic system”, inhibitory receptor-ligand interactions are not primarily used to regulate which immune cell types respond to which targets, but rather to have the same immune cell type respond differently, depending on environmental cues such as inflammation. A well-understood example of the latter type of systems is the PD-1/PD-L1 signaling axis that controls T cell activity (Sun et al., 2018), and, based on more recent data, in some cases also myeloid cell activity (Strauss et al., 2020). Specifically, expression of the PD-1 immune checkpoint on T cells is induced by T cell receptor triggering, thereby leading to a lowering of T cell sensitivity under conditions of chronic antigen encounter. Similarly, PD-L1 expression on tumor cells and other target cells can be increased many-fold by the T cell effector molecule IFN γ , thereby also creating a negative feedback loop by receptor

regulation on the target cell side. What is the evidence for a role of the CD47-SIRP α axis as either a “differential threshold regulator”, or as a “dynamic regulator” of immune cell activity?

In at least some situations, such as during mobilization of human HSCs where CD47 expression levels are increased approximately 4-fold, environmental regulation of axis activity appears to take place. Notably, this modest change in CD47 expression level does seem physiologically relevant, as HSCs from mice that lack a single CD47 allele, and that show a 2-fold reduction in CD47 expression levels, display a significant increase in susceptibility to macrophage-mediated phagocytosis *in vivo* (Jaiswal et al., 2009). Dynamic regulation of CD47-SIRP α signaling strength also appears relevant for synaptic pruning in brain development (Lehrman et al., 2018), where increased localization of CD47 to active synapses, thereby preventing engulfment by microglial cells, has been shown.

With respect to the mechanistic basis for alterations in CD47 expression or localization, *in vitro* studies have provided evidence that in a subset of cell lines, and most prominently in the MCF7 breast cancer cell line, CD47 expression levels are boosted through TNF α - and NF- κ B signaling, and the NF- κ B transcription factor was shown to directly bind to a super enhancer (SE) site near the *CD47* gene that promotes *CD47* gene transcription (Betancur et al., 2017). In addition, blockade of TNF α signaling was both shown to reduce CD47 expression and to induce increased phagocytosis of MCF7 cancer cells. Furthermore, tumor cells with active constituent enhancers in SEs near the *CD47* gene had increased CD47 levels compared to tumor cells lacking these (Betancur et al., 2017). Interestingly, cytokine stimulation may in certain settings also reduce CD47 expression levels, as observed in HLH patients. Incubation of CD34⁺CD38⁻ HSCs with a cytokine cocktail containing IL-6, M-CSF, IFN γ and TNF α at concentrations similar to those observed in patient serum led to a modest decrease in CD47 expression, to approximately 75% of control values, whereas CD47 levels on CD34⁺CD38⁺ progenitor or unfractionated mature cells were unaffected (Kuriyama et al., 2012). Similarly, TNF α was shown to induce a slight decrease in CD47 levels on malignant T cells (but not non-transformed T cells) from patients with Sézary syndrome, whereas CD47 levels were increased upon incubation with IL-13, IL-7 and IL-4 (Johnson et al., 2019). Finally, CD47 expression levels in cancer cells have also been shown to be regulated by miR-133a (Suzuki et al., 2012), miR-192 (Yang et al., 2015), the *MYC* oncogene (Casey et al., 2016), by HIF-1 in a subset of breast cancer cells when exposed to hypoxic conditions (Zhang et al., 2015), and by ERK signaling via nuclear respiratory factor 1 (NRF-1) in melanoma cells (F. Liu et al., 2017).

Other putative mechanisms that may control pathway strength by regulation of CD47 levels or function include alternative polyadenylation (ApA) of CD47 transcripts and posttranslational modification of the CD47 protein. With respect to the former, the transport of CD47 protein to the membrane is regulated by the length of the 3' untranslated region (3'UTR) of mRNA transcripts. Specifically, ApA leads to the generation of two CD47 mRNA isoforms that either contain a short or long 3'UTR. The long 3'UTR is rich in potential binding sites for the HuR RNA-binding protein that has been shown to mediate CD47 protein localization to the plasma membrane via SET and active RAC1. Unable to bind these proteins, the CD47 protein that is translated from the short 3'UTR version is

translocated into the endoplasmic reticulum (Berkovits and Mayr, 2015). Cells that express the CD47 protein translated from the short 3'UTR mRNA display increased susceptibility to macrophage-mediated phagocytosis even when total cellular CD47 levels are the same. The ability to create functionally distinct pools of CD47 protein by spatial separation may perhaps play a role in the CD47-mediated regulation of synaptic pruning, where local activity of the CD47-SIRP α axis in the cell membrane is thought to be important. Next to the regulation of CD47 protein localization, the activity of the CD47 protein has also been shown to depend on post-translational modification. Specifically, the amino-terminus of the CD47 protein contains a pyroglutamate (pGlu) residue that is essential to create a high affinity SIRP α binding site, and this modification has been shown to depend on the glutaminyl-peptide cyclotransferase-like (QPCTL) protein (Logtenberg et al., 2019). In line with the role of the CD47 pGlu in SIRP α binding, both genetic deletion of QPCTL and pharmacological inhibition of QPCTL suppresses inhibitory signaling via the CD47-SIRP α axis. At present, no evidence has been obtained indicating that regulation of either QPCTL, or of enzymes that may remove pGlu residues, is used as a physiological mechanism to tune pathway strength. Nevertheless, pharmacological manipulation of QPCTL activity using small molecule inhibitors forms a conceptually attractive strategy to manipulate pathway strength for therapeutic purposes.

With respect to the regulation of SIRP α levels on myeloid cells, stimulation of macrophages with the TLR ligands LPS and poly(I:C) has been shown to induce SIRP α downregulation (Dong et al., 2008; Kong et al., 2007). Furthermore, work by Zen et al. has provided evidence that inflammation can also reduce SIRP α signaling in polymorphonuclear (PMN) cells by inducing cleavage of the intracellular domain of the receptor (Zen et al., 2013). SIRP α proteins isolated from PMNs, but not monocytes, from donors with various inflammatory conditions consisted of a mixture of the full-length and a truncated protein, the latter lacking the ITIM/ITSM-containing cytoplasmic domains. Furthermore, the cleavage of SIRP α intracellular domains correlated with IL-17 levels in a dextran sulphate sodium (DSS)-induced colitis model, and could be blocked by anti-IL-17 antibody (Zen et al., 2013). As a separate form of regulation of SIRP α activity by proteolytic cleavage, the generation of a soluble extracellular domain of SIRP α following exposure of monocytic THP-1 cells to LPS and TNF α has been reported (Londino et al., 2015). By the same token, the generation of soluble extracellular SIRP α has also been reported in synapses in response to neuronal activity, presumably contributing to presynaptic maturation (Toth et al., 2013).

To conclude, while our understanding of the regulation of CD47-SIRP α axis signaling strength is presently limited, evidence indicating spatial or temporal control of CD47 and/or SIRP α expression level or function is starting to emerge. In addition, it is noted that regulation of signalling strength need not necessarily occur through changes in either receptor expression level or localization but could, for instance, also be achieved by altered intracellular signal transduction, and a further effort to understand potential regulation at this level would appear warranted.

Therapeutic manipulation of the CD47-SIRP α axis

Inhibition of the CD47-SIRP α axis can be considered attractive in all settings where clearance of intended target cells can be promoted while clearance of other vital cell types is avoided (Figure 2). At present, 3 classes of inhibitors of the CD47-SIRP α signaling pathway that are in varying stages of development may be distinguished: Molecules that inhibit pathway activity by blocking the CD47 molecules on target cells, molecules that inhibit pathway activity by blocking the SIRP α molecules on immune effector cells, and inhibitors of the QPCTL enzyme that is required for CD47 maturation. With respect to the first class of molecules, which includes not only anti-CD47 antibodies but also SIRP α -Fc fusions, it is important to distinguish molecules that do and do not lead to substantial FcR triggering. Specifically, the clinical value of CD47 binding molecules that retain substantial FcR activating capacity (e.g. human IgG1) is likely to be capped by on-target – off-tumor toxicity, such as depletion of erythrocytes (Ingram et al., 2017). Indeed, many of the efforts in the clinical development of anti-CD47 antibodies currently focus on molecules with decreased Fc γ R-binding capacity, such as IgG4, although it is noted that most of these IgG4 CD47-targeting molecules still induce substantial anemia in non-human primates or in cancer patients (Advani et al., 2018; Weiskopf et al., 2013). Interestingly, TTI-621, a protein consisting of the CD47-binding domain of SIRP α fused to the human IgG1 Fc domain appears to only minimally bind human erythrocytes and induce hemagglutination *in vitro*, in spite of the presence of the activating Fc domain (Petrova et al., 2017). This reduced toxicity may be due to the less effective clustering of CD47 on erythrocytes because of the stringent association with the cytoskeleton (Mouro-Chanteloup et al., 2003; Subramanian et al., 2006). To what extent the differential dependency on receptor clustering of this molecule would also influence the occurrence of any (other) major on-target off-tumor toxicities requires further understanding. However, clinical benefit with a decrease in dominant malignant T cells has been reported in four out of five patients with Sézary syndrome (Johnson et al., 2019). In more recent work, agents that simultaneously target CD47 and tumor-specific antigens have been tested in preclinical and clinical studies, with the aim to restrict CD47 blockade to the tumor antigen-positive cell population (Dheilly et al., 2017; Fischer et al., 2015; Ma et al., 2020; Piccione et al., 2015; 2016; Ponce et al., 2017). As these molecules contain active Fc domains, successful use of such bispecifics will require minimal binding to non-malignant CD47-positive cells, something that may be achieved by lowering the affinity of the CD47 targeting arm (Buatois et al., 2018; Piccione et al., 2016). Whether such molecules that carry a low affinity CD47 targeting arm allow sufficient blockade of CD47 molecules in the clinic remains to be established. In addition, this approach is expected not to be effective for tumor antigens that are expressed at low levels relative to CD47.

Biologicals that target SIRP α and small molecule inhibitors of QPCTL are currently being pursued in parallel to CD47 targeting agents. *In vitro* and *in vivo* data suggest that, much like Fc-null CD47 blocking antibodies, SIRP α -blocking agents can effectively induce anti-tumor control when used in combination with tumor-opsonizing antibodies (Ho et al., 2015; Ring et al., 2017; Voets et al., 2019; Zhao et al., 2011). In addition, as SIRP α is only expressed on a restricted number of cell types, targeting of the CD47-SIRP α axis with

SIRP α -blocking agents does not suffer from the antigen sink issue that has complicated the use of CD47 targeting molecules. With respect to the targeting of the CD47-SIRP α axis via the CD47 modifier QPCTL, at present *in vivo* data on small molecule inhibition of this enzyme are lacking. However, small molecule inhibition of the pathway may be attractive because of the expected high tissue penetrance of small molecule inhibitors and potential for oral bioavailability. Furthermore, CD47 molecules that newly arrive at the cell surface upon QPCTL inhibition lack pyroglutamate modification and, unlike antagonistic molecules that bind to CD47 or SIRP α , competition by the natural binding partners in the tumor microenvironment will thus not occur.

As discussed above, inhibition of the CD47-SIRP α axis will in many cases require simultaneous provision of an activating signal that marks a target cell population for destruction. Semi-selective marking of tumor cells may be achieved with antibodies against proteins such as CD20 and EGFR that opsonize the intended target cell population. However, sufficiently tumor-selective antibodies are lacking for most human malignancies. As a second strategy that does not rely on such opsonizing antibodies, activating signals may potentially be induced by cytotoxic therapies such as chemotherapy and low-dose radiation, with the implicit assumption that induction of stress signals would be biased towards the (tumor) cells that require clearing. The fact that CRT can be translocated to the cell surface upon induction of immunogenic cancer cells death by some chemotherapies (anthracyclines) and radiotherapies (reviewed in (Galluzzi et al., 2017)) provides some evidence for this model.

As a third strategy to achieve a sufficient level of selectivity, strategies that can provide either spatial or temporal control over CD47 pathway blockade could be considered. Spatial separation may, for instance, be attempted using technologies that allow local uncapping of antibodies in the tumor microenvironment (reviewed in (Autio et al., 2019)). Conceivably, uncapping of anti-CD47 or anti-SIRP α agents could be used to bias pathway inhibition to that site, while in the case of CD47 blockade also remedying the previously described antigen sink issue. Alternatively, selective recognition of cells at the tumor site may theoretically be achieved through the use of capped opsonizing antibodies. With respect to the use of temporal control of pathway inhibition to achieve an increased level of specificity, two strategies have been developed (Figure 2). As a first strategy, in order to ameliorate the erythrocyte depletion seen upon CD47 blockade, a priming strategy has been developed in which patients are first treated at a low dose, in order to remove the ageing erythrocytes that are more prone to display prophagocytic signals, thereby inducing compensatory hematopoiesis (Advani et al., 2018). Second, as shown by Chhabra et al, temporal control over pathway inhibition may also be used to facilitate cell transplants. The transplantation of allogeneic hematopoietic stem cells that is used to treat hematologic malignancies requires the elimination of host HSCs. To avoid the toxicities associated with the currently used host conditioning regimens, a strategy has been developed in which *in vivo* HSC depletion is achieved by co-administration of CD47 antagonists with anti-c-Kit antibodies that opsonize HSCs and their downstream progenitors (Chhabra et al., 2016; George et al., 2019). The resulting depletion of host HSCs then allows engraftment of donor HSCs, provided obviously that HSC opsonization/ CD47-SIRP α pathway inhibition no longer occurs at that point. Optimal temporal control over pathway inhibition may be achieved using antibodies

that display a short *in vivo* half-life or, perhaps preferably, with small molecule QPCTL inhibitors, as such inhibitors will not affect the preformed CD47 molecules on incoming cells. In future work, the concept of enhancing cell engraftment by transient targeting of an endogenous cell pool may conceivably be extended towards other (stem) cell therapies.

Clinical data that evaluate the potential of CD47 blockade have started to emerge. Evaluation of the combination of CD47 blockade with anti-CD20 treatment has yielded particularly promising data in patients with anti-CD20 refractory diffuse large B-cell lymphoma (DLBCL) and follicular lymphoma, with a 50% objective response rate, and a 36% complete response rate (Advani et al., 2018). In addition, early results suggest that the combination of CD47 blockade with azacitidine leads to significant activity in myelodysplastic syndrome (Chao et al., 2020). Next to the use of CD47 pathway inhibitors in oncology, cellular therapies, and possibly fibrosis and atherosclerosis, interventions that enhance pathway activity may also be of value in certain settings. Specifically, overexpression of CD47 on induced pluripotent stem cells (iPSCs), in combination with MHC class I and class II gene inactivation, was shown to render cells “hypoimmunogenic”, thereby enabling cell transplantation across MHC barriers (Deuse et al., 2019). These data provide the first tantalizing evidence indicating that enhancement of the CD47-SIRP α axis is likely to have biomedical value, and use in e.g. off-the-shelf CAR T cells would seem of potential interest.

Concluding remarks

The CD47-SIRP α axis is a key regulator of cell fate in a number of conditions and hence an attractive therapeutic target. To optimally exploit this axis as therapeutic target, it will be valuable to increase our understanding on the nature of the activating signals that are counterbalanced by the CD47-SIRP α axis in different cell systems. In addition, it will be important to determine by which mechanisms the strength of the SIRP α -mediated inhibitory signal is regulated. Finally, it will be critical to design therapeutic strategies that modify the relative strength of activating signals and the SIRP α -mediated inhibitory signal in either a time-dependent, location-specific, or cell type-specific manner. Strategies that manipulate the CD47-SIRP α axis are starting to show an intriguing clinical signal, and an improved understanding how to steer the balance between this axis and the activating signals that cells receive is likely to increase the scope of clinical application.

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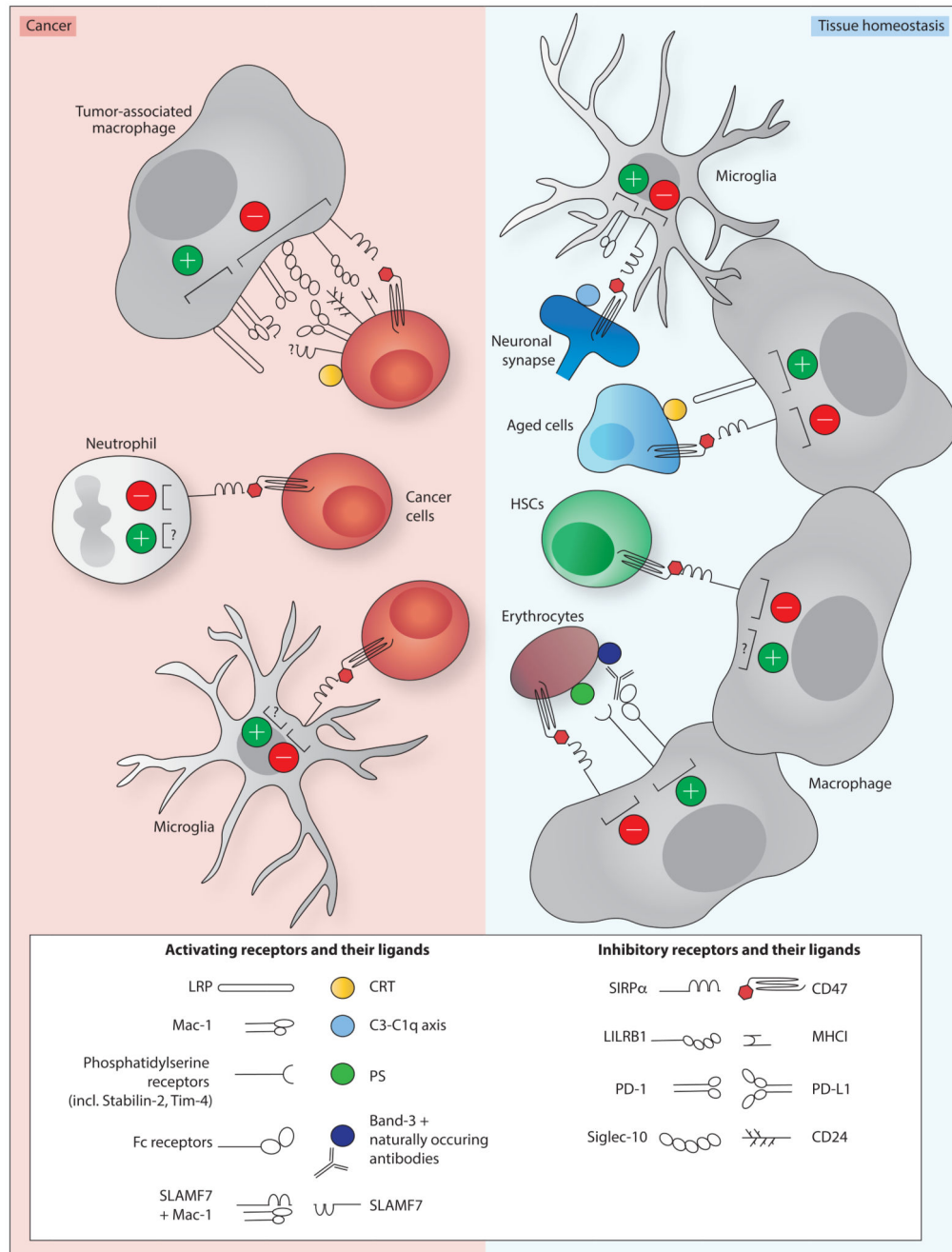


Figure 1. Activating receptors on myeloid cells that are influenced by the CD47-SIRP α axis in tissue homeostasis and in cancer

Macrophage-, neutrophil- and microglia-mediated cytotoxicity towards cancer cells (left panel) is inhibited by CD47-SIRP α signalling. Macrophage-mediated cytotoxicity has also been shown to be negatively influenced by additional inhibitory receptor-ligand interactions, including leukocyte immunoglobulin-like receptor 1 (LILRB1)/ major histocompatibility complex I (MHC-I) (Barkal et al. 2017), programmed cell death 1 (PD-1)/ programmed cell death 1 ligand 1 (PD-L1) (Gordon et al. 2017), and Siglec-10/CD24 (Barkal et al. 2019). The interaction between prolow-density lipoprotein receptor-related protein (LRP) and

calreticulin (CRT) (Chao et al. 2010) has been identified as activating signal in myeloid-mediated cytotoxicity towards cancer cells.

In tissue homeostasis (right panel), synaptic pruning by microglia, and removal of aged cells, erythrocytes and hematopoietic stem cells (HSCs) by macrophages has been shown to be inhibited by CD47-SIRP α signalling (Boas et al. 1998; Kay et al. 1994; Lutz & Bogdanova 2013; Oldenborg et al. 2000; Jaiswal et al. 2009; Lehrman et al. 2018; Feng et al. 2018). C1q or C3 localization to neuronal synapses enhances microglia-dependent synaptic pruning by binding Mac-1 (Schafer et al. 2012; Stevens et al. 2007). CRT binding to LRP on macrophages stimulates the removal of ageing neutrophils (Feng et al. 2018). Binding of naturally occurring antibodies to band-3 and increased phosphatidylserine (PS) exposure may enhance macrophage-mediated removal of ageing erythrocytes (Boas et al. 1998; Kay et al. 1994; Lutz & Bogdanova 2013). Activating signals on HSCs remain unidentified.

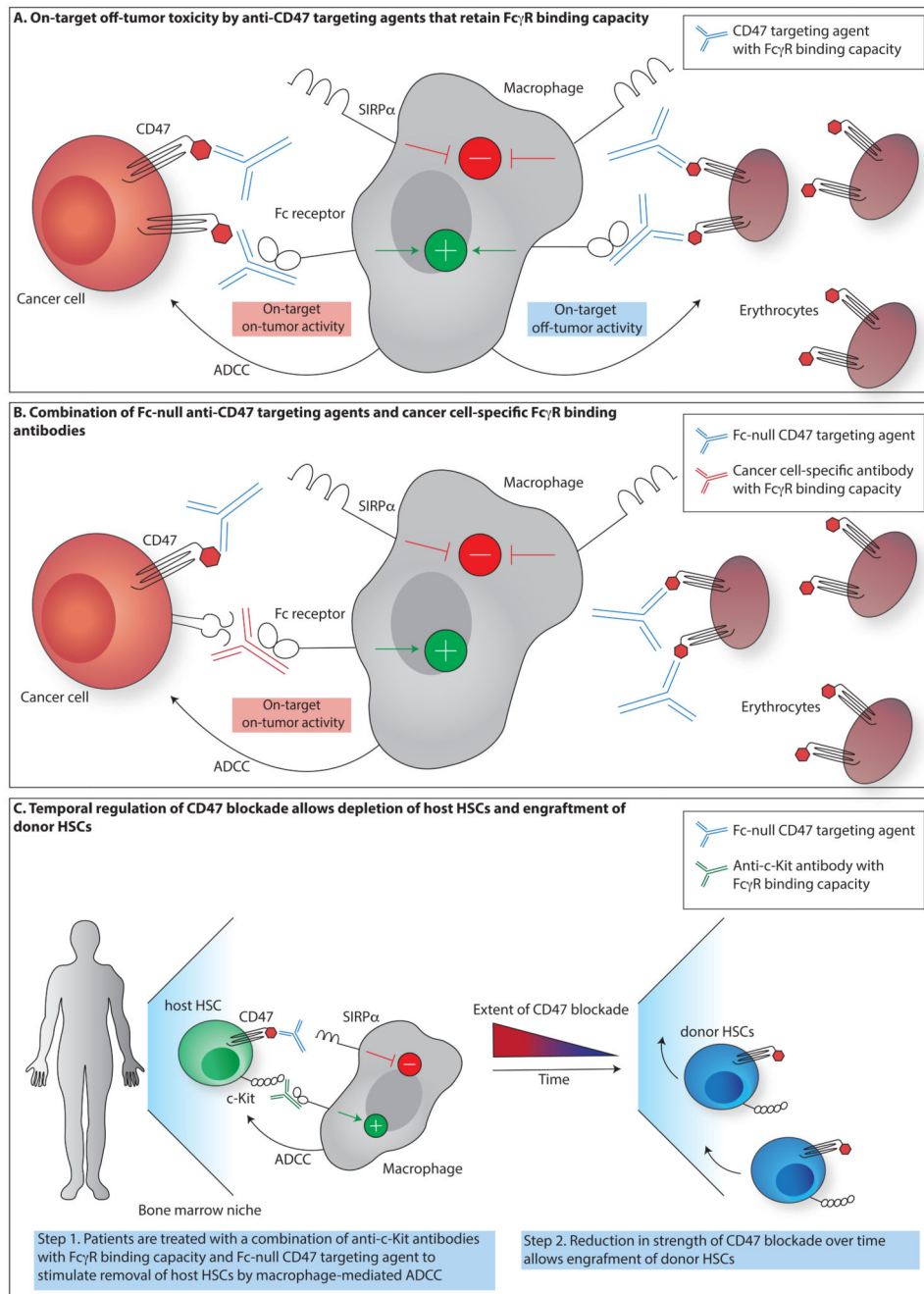


Figure 2. Strategies to exploit CD47 blockade.

A. Blocking CD47 targeting agents that retain Fc γ R binding capacity prevent inhibitory signalling through SIRP α , but simultaneously decorate CD47⁺ cells with an activating signal, thereby inducing antibody-dependent cytotoxicity (ADCC) by, for example, macrophages. Use of such agents can result in on-target off-tumor toxicity towards healthy CD47-expressing cells, such as erythrocytes. B. Fc-null CD47 targeting agents, such as the high-affinity SIRP α -Fc fusion protein ALX148 that contains an inactive Fc domain, prevent inhibitory signalling through SIRP α , but without the simultaneous delivery of an activating

signal. Combination of such CD47 blocking agents with cancer cell-specific antibodies that retain Fc γ R binding capacity forms a strategy to direct ADCC activity towards cancer cells while minimizing on-target off-tumor toxicity. C. Selective depletion of defined cell pools may be achieved by temporal control over CD47 blockade. For example, as shown by Chhabra et al., recipient hematopoietic stem cells (HSCs) can be targeted for removal by the combination of blocking agents targeting CD47 and opsonizing anti-c-Kit antibodies (Chhabra et al. 2016). Transplantation of donor HSCs at a time when the strength of CD47 inhibition has diminished then allows donor HSC engraftment.