**Myb-Independent Macrophages: A Family of Cells That Develops with Their Tissue of Residence and Is Involved in Its Homeostasis**

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In most metazoans, all tissues contain phagocytes “in residence,” generally termed “macrophages” in vertebrates. In contrast to myeloid cells produced continuously by the bone marrow (BM), tissue-resident macrophages develop during embryogenesis together with their tissue of residence, and persist in adulthood, independently of hematopoietic stem cells and the transcription factor *Myb*. They therefore represent an independent lineage from blood monocytes, dendritic cells, and monocytes/macrophages that are recruited to tissues during inflammation. Tissue-resident macrophage functions are yet to be completely defined. They all share the ability to scavenge toxic compounds, lipids, microorganisms, and dead cells and contribute to tissue remodeling, via phagocytosis and the production of growth factors. In contrast, the production of inflammatory mediators seems to be more associated with BM-derived cells. Tissue-resident macrophages and BM-derived myeloid cells thus differ in developmental origin and functions; the term “macrophages” could be reserved for *Myb*-independent-resident macrophages to avoid confusion. A genetic and molecular dissection of resident macrophage functions will reveal their roles in tissue metabolism and the maintenance of homeostasis independently of the extravasation of inflammatory leukocytes, and in the control of the recruitment of BM-derived cells in overt inflammation.

Elie Metchnikoff studied the development and functions of “phagocytes,” cells capable of engulfing particles, which are observed through all phyla, from protozoan to mammals. In triploblastic organisms (having three germinal layers: ecto-, meso-, and endoderm), phagocytes are mesodermal cells that have retained the ability to engulf foreign bodies, microorganisms, and neighboring cells. They contribute to tissue renewal and maintenance (as shown for Echinoderm larvae [Metchnikoff 1884b]) and respond to wounding and pathogens (Metchnikoff 1884b). In the tree of life, the tissue phagocytes are observed before the emergence of the vascular system. In animals with a vascular system, Metchnikoff suggested that such “extra-vascular” or “connective tissue” phagocytes, which he termed “Macrophages,” participate in tissue remodeling and in the recruitment of the vascular leukocytes in response to tissue damage (Metchnikoff 1884a).

Since then, many attempts were made to characterize their origin and their relationship with blood leukocytes. The most prevalent model, developed in the 1970s, integrated macrophages into the concept of the “mononuclear phagocyte system” (MPS), based on (1) common morphological criteria, (2) common functions—professional phagocytes that attach firmly to glass surface, and (3) the hypothesis that they share a common origin in the bone marrow (BM), from hematopoietic stem and progenitor cells (HSPCs) (van Furth and Cohn 1968; van Furth et al. 1972). The MPS thus proposed that myeloid precursors from the BM differentiate into monocytes that circulate in the blood and populate tissues as macrophages in the steady state and during inflammation. In accordance with this model, circulating monocytes can extravasate from the blood and give rise to inflammatory macrophages in response to inflammatory cues and other signals, such as tumor progression and wound healing (Geissmann et al. 2003; Serbina and Pamer 2006). The classical dendritic cell (DC) lineage (Meredith et al. 2012), initially described by Ralph Steinman, also shares with monocytes a common BM progenitor that circulates in the blood and seed lymphoid tissues (Fogg et al. 2006; Liu et al. 2007; Onai et al. 2007). The MPS model allowed progress into the study of the function of macrophages and DCs, using monocyte-derived cells as a model.

However, a large number of observations are not in accordance with the MPS model in regard to macrophages. A population of macrophages appears in the embryo before hematopoietic stem cells (HSCs) are detected (Sorokin et al. 1992). During embryogenesis, macrophages in human, mice, and zebrafish were shown to develop by passing the monocyte stage (Naito et al. 1996; Herbomel et al. 1999; Naito 2008). Tissue-resident macrophages are not affected in monocytopenic mice (Yamada et al. 1990; Ajami et al. 1999; Naito et al. 1996) and do not exchange between parabiotic mice (Mered et al. 2002; Ajami et al. 2007). After irradiation and BM graft, tissue macrophages are only partially “replaced,” and this depends on the level of tissue damage, rather than the BM replacement (Mildner et al. 2007). In addition, several populations of tissue-resident macrophages, such as Langerhans cells in the epidermis, Kupffer cells in the liver, microglia in the central nervous...
system, and macrophages from the peritoneum and pleura, proliferate in situ in steady state and in inflammation (Sawyer et al. 1982; Miyachi and Hashimoto 1987; Yamada et al. 1990; Chorro et al. 2009; Davies et al. 2011; Jenkins et al. 2011). Pulse-labeling at E7.25–7.5 (before hematopoietic stem cell emergence) in Runx1-MER-iCre-MER; R26LSL-YFP embryos results in the labeling of the microglia in adults (Ginhoux et al. 2010).

Collectively, these observations suggested that tissue macrophages might not meet the criteria to be included in the MPS. Indeed, they develop from embryonic progenitors distinct from HSPCs, persist in adult independently of the transcription factor Myb (Schulz et al. 2012), and renew independently of BM-derived cells and monocytes (Schulz et al. 2012; Hashimoto et al. 2013; Yona et al. 2013). We here review experimental evidence investigating the origin of macrophages and discuss the existence and consequences of two models of differentiation of myeloid cells that coexist through embryonic development and adulthood.

**EXPERIMENTAL EVIDENCE FOR A Myb-INDEPENDENT HSC-INDEPENDENT LINEAGE OF TISSUE MACROPHAGES**

**Myb-Independent Tissue Macrophages Self-Renew Independently from the BM**

Gene invalidation in the mouse has allowed the identification of genes that are required for the generation and/or maintenance of progenitors and hematopoietic stem cells. Among these genes, the transcription factor Myb is required for development of HSPCs and all HSPC-derived monocytes, macrophages, and DCs (Mucenski et al. 1991). Myb-deficient mice present a phenotype reminiscent of that of Runx1-deficient mice, but do not present hemorrhage and die later with anemia between E15 and E17 (Mucenski et al. 1991; Mukoyama et al. 1999; Sumner et al. 2000; Schulz et al. 2012). Although Myb-deficient AGM and fetal liver (FL) do not present HSC activity, macrophages are present in normal numbers in E10.5 yolk sac (YS) (Schulz et al. 2012), suggesting that Myb is dispensable for YS hematopoiesis (Sumner et al. 2000; Schulz et al. 2012). At later stages of embryonic development, two populations of tissue myeloid cells can be defined on the expression of F4/80 and CD11b. As expected for HSC-derived myeloid cells, CD11b^{bright} myeloid cells in all organs and CD117/c-Kit^{+} progenitors in the FL were absent in Myb-deficient animals. However, tissue macrophages, defined as F4/80^{bright}, were present in all organs examined, such as the liver, skin, brain, spleen, pancreas, lungs, and kidneys (Schulz et al. 2012).

In adult mice, the loss of Myb by conditional deletion also causes failure of hematopoiesis (Lieu and Reddy 2009; Schulz et al. 2012). Mx1-Cre mice allow a widespread expression of Cre in all cell types upon Poly(I:C) administration. Targeted disruption of the Myb gene in Cdh5.2; Mx1Cre; Myb^{flox/flox} mice leads to a rapid depletion of the HSC pool and of blood monocytes and granulocytes, and a syngeneic Cdh5.1 Myb^{-/-} BM engrafts without irradiation. Monocytes and granulocytes, as well as CD11b^{bright} myeloid cells in the spleen, liver, kidney, and pancreas, are completely replaced by donor BM-derived cells, whereas microglia, epidermal LCs, Kupffer cells, and a large proportion of F4/80^{bright} in other tissues remain of the host, Myb-deficient, origin 3 mo after transplantation of a wild-type BM (Schulz et al. 2012). Two independent groups have confirmed that tissue-resident macrophages self-renew independently from monocytes and BM progenitors using expression of Cre driven by CX3CR1 (Yona et al. 2013) or by following repopulation of myeloid cells and tissue-resident macrophages after genetic depletion (Hashimoto et al. 2013). Previous studies had shown that depletion of blood monocytes by strontium-89 does not affect the number of resident peritoneal and alveolar macrophages (Sawyer et al. 1982) and Kupffer cells (Yamada et al. 1990) nor their proliferation capacity (Sawyer et al. 1982; Yamada et al. 1990).

Therefore, tissue “resident” macrophages develop in the absence of Myb and HSPCs and persist in adult tissues independently of HSPCs (Fig. 1).

**Myb-Independent Macrophages Originate from a CSF1-R Progenitor Generated in the YS, Are Largely FLT3-Independent, and Are Capable of Proliferation**

In an unsupervised hierarchical clustering analysis of gene expression arrays from E10.5 YS macrophages, E16.5 F4/80^{bright} macrophages, and E16.5 CD11b^{bright} myeloid cells from Myb wild-type (WT) and Myb-deficient embryos, F4/80^{bright} tissue macrophages clustered with E10.5 YS macrophages. The Csfr1 receptor (Csfr1r) was part of the Myb-independent F4/80^{bright} signature, whereas Flt3, the FLT3L receptor, was part of the Myb-dependent F4/80^{bright} signature (Schulz et al. 2012). Therefore, Csfr1 and Flt3 were used to investigate the persistence of YS-derived macrophages into adulthood using fate-mapping strategies.

Early expression of Csfr1 in YS precursors and the availability of a tamoxifen-dependent Csfr1-CRE mouse line (Csfr1-MER-iCre-MER) (Qian et al. 2011) allow us to identify in adults the progeny of CSF1R^{+} cells labeled during early development. In utero administration of TAM in E8.5 Csfr1-MER-iCre-MER; R26LSL-YFP embryos results in YFP expression by adult Myb-independent macrophages, whereas Myb-dependent HSC-derived cells such as blood leukocytes are not labeled (Schulz et al. 2012). Thus, E8.5 CSF1R^{+} precursors give rise to tissue macrophages, such as microglia, Langerhans cells, Kupffer cells, and macrophages in the spleen, lung, pancreas, and kidneys that persist in adults (Fig. 1). They are most likely generated in the YS, because they are labeled before the HSC emergence from the aorto-gonado-mesonephros (AGM). Whether these progenitors differentiate in situ in the YS or whether they migrate into the FL, where they amplify and differentiate before seeding other tissues, is still an open question (discussed below).

Flt3 is expressed on multipotent hematopoietic progenitors (Buza-Vidas et al. 2011), MDPs (Auffray et al.
The progeny of FLT3\(^+\) progenitors can be detected by expression of YFP in the F1 progeny of Flt3\(^{-}\)-Cre\(^+\)/C2R26LSL-YFP mice (Srinivas et al. 2001; Benz et al. 2008). YFP expression is overall restricted to blood leukocytes and to tissue Myb-dependent CD11b\(^{high}\) myeloid cells (Schulz et al. 2012), indicating that the development of Myb-independent macrophages occurs largely independently of FLT3\(^+\) precursors (Fig. 1).

Persistence of Myb-independent macrophages in adulthood could be explained by cell proliferation. Kupffer cells filled with phagocytized beads are present 3 mo after intravenous administration of beads and mitotic bead-filled Kupffer cells can also be detected (Bouwens et al. 1986), suggesting that Kupffer cells have a long life span and can proliferate even after phagocytosis. In humans, epidermal Langerhans cells were also shown to remain of donor (limb) origin 10 yr after limb graft (Kanitakis et al. 2004, 2011). This is compatible with results from several recent studies that have re-investigated the proliferation capacities of resident macrophages and confirmed earlier observations that microglia, epidermal LCs, and macrophages in the liver, peritoneum, and pleura can proliferate during inflammation and in the steady state (Bouwens et al. 2009), cDC progenitor (Onai et al. 2007), and common lymphoid progenitors in the BM (Buza-Vidas et al. 2011). The progeny of FLT3\(^+\) progenitors can be detected by expression of YFP in the F1 progeny of Flt3\(^{-}\)-meriCre; Rosa-YFP F1 embryos labels macrophages/macrophage precursors (green circles) presumably in the YS. These macrophages/macrophage precursors continue to be generated during development (dashed circles) in the YS or the FL. Progeny of E8.5–E9.5 CSF1R\(^+\) cells in adults include liver Kupffer cells, epidermal Langerhans cells, brain microglia, and F4/80\(^{bright}\) macrophages in other tissues. A second wave of hematopoiesis within the embryo proper gives rise to Myb-dependent FLT3\(^+\) HSPCs that expand in the fetal liver and can be fate mapped using Flt3-Cre mice (orange circle). Their myeloid progeny in adults includes monocytes, classical dendritic cells, and PDCs, neutrophils, mast cells, and osteoclasts.

The Case of the BM Radiation Chimera

BM transplantation results in a very high level of chimerism for granulocytes, monocytes, classical DCs, and plasmacytoid DCs, indicating that they are short-lived, being replaced continually from BM precursors (Fogg et al. 2006; Onai et al. 2007; Auffray et al. 2009; Liu et al. 2009) (reviewed in Geissmann et al. 2010). It also leads to a partial, relatively inefficient, and variable level of chimerism of tissue macrophages. The latter seems to be dependent on the local tissue damage and/or macrophage death induced by the irradiation protocol, and the presence of a T-cell allogeneic response by donor cells. Irradiation and/or conditioning therapies induce cell death, tissue damage, and alteration of the basement mem-

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**Figure 1.** Differentiation of myeloid cells. Model for the development of the two myeloid lineages: the Myb independent (blue) and HSC dependent (orange). The origin of HSCs and yolk sac (YS) progenitors from hemangioblasts or hemogenic endothelium is still a source of controversy (as reviewed in Ueno and Weissman 2010). During embryonic development, hematopoiesis in the YS comprises EMPs, macrophage-restricted progenitors (MP), and multipotent progenitors (MPP) (Palis et al. 2001; Bertrand et al. 2005). Pulse labeling at E8.5 using TAM (green arrow) in Csf1r-meriCre; Rosa-YFP F1 embryos labels macrophages/macrophage precursors (green circles) presumably in the YS. These macrophages/macrophage precursors continue to be generated during development (dashed circles) in the YS or the FL. Progeny of E8.5–E9.5 CSF1R\(^+\) cells in adults include liver Kupffer cells, epidermal Langerhans cells, brain microglia, and F4/80\(^{bright}\) macrophages in other tissues. A second wave of hematopoiesis within the embryo proper gives rise to Myb-dependent FLT3\(^+\) HSPCs that expand in the fetal liver and can be fate mapped using Flt3-Cre mice (orange circle). Their myeloid progeny in adults includes monocytes, classical dendritic cells, and PDCs, neutrophils, mast cells, and osteoclasts.
brane—for example disruption of the blood–brain barrier—and promote inflammation (Diserbo et al. 2002; Kaya et al. 2004; Barcellos-Hoff et al. 2005; Kierdorf et al. 2013b). This is responsible for the recruitment of donor blood leukocytes in host tissues, akin to some inflammatory processes. For example, in a model of encephalitis where microglial cells are not killed by irradiation, monocytes are transiently recruited but do not ultimately contribute to the resident microglial pool and vanish, whereas resident microglia proliferate and persist following remission (Ajami et al. 2011). In BM radiation chimera, local brain irradiation is indeed required for the monocyte recruitment into the parenchyma (Mildner et al. 2007). In mice irradiated with a head protection and receiving a syngeneic BM transplant, monocytes fail to enter the brain parenchyma and microglia remains entirely of the host origin (Mildner et al. 2007). Similarly, Kupffer cells and epidermal Langerhans cells can remain of the host origin for extended periods of time after syngeneic BM transplantation (Katz et al. 1979; Perreault et al. 1984; Merad et al. 2002; Ajami et al. 2007).

Therefore, the irradiation chimera represents a useful tool to manipulate BM-derived leukocytes, but may not inform on the biology of tissue-resident macrophages—first because they are not of BM origin and also because irradiation induces cell death among tissue cells and their associated macrophages, creates inflammation, and recruits inflammatory cells. Whether recruited BM cells may functionally replace the resident macrophages when the latter are partially killed by irradiation is an open debate in the field and is briefly discussed below. Para-biosis studies have been very useful to investigate the contribution of BM to tissue macrophages and exposed some experimental issues associated with irradiation (Merad et al. 2002; Ajami et al. 2007; Liu et al. 2007). However, para-biosis studies are considered unethical in several countries. BM transplantation following genetic depletion of hematopoietic stem cells by conditional inactivation of Myb (Schulz et al. 2012) may be a useful tool in this regard.

**HSCS-DERIVED Myb-DEPENDENT MELOID CELLS: MONOCYTES AND DENDRITIC CELLS**

In contrast to resident tissue macrophages, adoptive transfer experiments and analysis of syngeneic BM transplantation have clearly shown that blood leukocytes, the classical DC lineage (Meredith et al. 2012), and PDCs develop from BM-derived HSCs. Extensive work has clarified the relationship between DC and monocyte lineages and identified their respective precursors in the blood and BM in the steady state (Fogg et al. 2006; Liu et al. 2007, 2009; Onai et al. 2007). Monocytes, DCs, and granulocytes develop from a common myeloid progenitor (CMP) (Akashi et al. 2000). A monocyte/macrophage and DC precursor (MDP) (Fogg et al. 2006) gives rise to monocytes and to the common DC precursor (CDP) (Onai et al. 2007; Liu et al. 2009). Of note, the development of monocytes and, to some extent, DCs also depends on the growth factor receptor CSF1R (Sasmono et al. 2003; MacDonald et al. 2005), and Csf1r deficiency led to a reduced number of circulating monocytes (Dai et al. 2002). Flt3- and Flt3L-deficient mice have shown their critical role in the differentiation of cDCs and PDCs (McKenna et al. 2000; Waskow et al. 2008) but appeared to be dispensable for the development of monocytes.

**Monocytes as Effector Cells**

Monocytes are a heterogeneous population in regard to shape, size, and function (reviewed in Grage-Griebenow et al. 2001). In mice, at least two monocyte subsets have been identified. The Ly6C+ “inflammatory” subset of blood monocytes, an equivalent to human CD14+ monocytes (Geissmann et al. 2003; Cros et al. 2010), extravasate in infected and inflamed tissues where they produce reactive oxygen species (ROS), iNOS, and cytokines and contribute to T-cell activation or polarization (Serbina et al. 2008; Soudja et al. 2012). CCR2-mediated signaling is important for the release of Ly6C+ monocytes from the BM into the blood and their recruitment to inflammation sites (Serbina and Pamer 2006; Tsou et al. 2007).

The Ly6Cint subset of monocytes and their human putative counterparts (CD14dim) (Cros et al. 2010) patrol the vasculature (Auffray et al. 2007), survey the endothelium, scavenge intraluminal debris, and can orchestrate the neutrophil-mediated necrosis of endothelial cells in response to a danger signal (Carlin et al. 2013). They are also equipped with a full set of Fc receptors involved in the uptake of immune complexes (Biburger et al. 2011).

Instead of being intermediates between the BM and tissue macrophages, Ly6C+ and Ly6Cint blood monocytes rather appear to be bona fide effector cells in inflamed tissues and within the vasculature, respectively.

**Dendritic Cells**

DCs are specialized in processing and presenting antigens to elicit specific T-cell effector functions. They are found in both lymphoid and nonlymphoid organs. Because many macrophage subsets express class II antigen and because blood monocytes can stimulate T cells in vitro, the distinction between DCs and macrophage is sometimes blurred. However, as outlined above, the classical DC lineage and plasmacytoid DCs are BM-derived cells with a short half-life (Liu et al. 2009; Meredith et al. 2012).

Recent studies on human immunodeficiencies have contributed to the understanding and the identification of transcription factors critical for their development. Patients carrying IRF8 and GATA-2 mutations present severe immunodeficiencies and a complete lack of circulating monocytes and DCs, although tissue macrophages appear to be normal (Dickinson et al. 2011; Hambleton et al. 2011).

One particular case is that of epidermal Langherans cells, which are currently classified as DCs, but which do not appear to be important for T-cell responses, develop from Myb-independent embryonic progenitors, and self-renew in adults (Merad et al. 2002; Chorro and Geissmann 2010).
THE ORIGINS OF Myb-INDEPENDENT MACROPHAGES AND Myb-DEPENDENT MYELOID CELLS

Different Developmental Waves of Progenitors with Different Hematopoietic/Myeloid Potential

During mouse hematopoiesis, the simplest hypothesis is that Myb-independent macrophages originate from extra-embryonic progenitors and/or macrophages at E8.5 and the requirement for Myb for HSCs maintenance (Mucenski et al. 1991) suggest that the progenitors that give rise to the Myb-independent lineage are generated in the YS. In support, it has been suggested that microglia arise from EMP progenitors in the YS (Kierdorf et al. 2013a). However, it is important to note that YS-generated progenitors seed the FL (Palis et al. 2001) as early as E9 (Kieusseian et al. 2012), which is also the site where Myb-dependent HSCs proliferate and differentiate from E11. Therefore, the FL probably represents a niche where the progenitors of Myb-independent macrophages and Myb-dependent HSCs probably coexist and expand, although a fate-mapping analysis of YS and HSC-derived hematopoietic cells within the FL has not yet been published. Work is needed to identify the progenitors from which Myb-independent macrophages develop, their differentiation potential, and the anatomical niches where they expand in vivo.

Although the simplest hypothesis is that Myb-independent macrophages originate from extra-YS hematopoietic progenitors distinct from intra-embryonic hematopoietic stem cells, the intra-embryonic sites not only generate HSCs or imHSCs, but also release multipotent progenitors without LTR activity into the circulation. P-SP and, later, the AGM region are sources of such intra-embryonic multi-potent hematopoietic progenitors at E8.5 (Godin et al. 1993, 1995; Medvinsky et al. 1993; Muller et al. 1994). Reciprocally, there is still a controversy on the origin of HSCs, and the restriction of YS progenitors to primitive erythromyelopoiesis is debated (Ueno and Weissman 2010). Transplantation of E9 YS cells into the YS cavities of synchronic embryos allows the detection of donor-derived spleen colonies in secondary transplant recipient (Weissman et al. 1978; Ueno and Weissman 2010). In addition, E9.0 YS cells repopulated erythroid, lymphoid, and myeloid lineages long-term upon transplantation into the YS.
newborn recipient animals (Yoder and Hiatt 1997). High proliferative hematopoietic precursors have also been reported in the YS before they can be detected in the blood or the embryo proper (Palis et al. 2001). Endothelial cells from E9.5 YS were shown to generate all blood cell types, including lymphocytes, suggesting that the YS endothelium may be a source of multipotent HSCs (Nishikawa et al. 1998).

It is of note that the blood circulation between the YS and the embryo is established very early at the eight-somite stage (E8–8.25) and therefore precursors can pass through the circulation from one site to the other, hampering efforts to study independently both pools of progenitors. In organotypic cultures of YS and P-Sp harvested before circulation, the progenitors have different potentials (i.e., YS progenitors have no lymphoid potential). Shortly after the start of circulation, P-Sp precursors can seed the YS, and cells of both origins would colonize the FL (Cumano et al. 1996). In accordance, mutant mice without circulation between both sites, because of lack of blood vessel formation (VE-Cadherin-deficient embryos), present erythroid and myeloid but no lymphoid potential within the YS (Rampon and Huber 2003). However, in apparent contradiction with these reports, hematopoietic progenitors are decreased at E9.5 in the embryo, but unaffected in the YS of Ncx1−/− mice, which die at E10, and have blood vessels but no cardiac contraction (Lux et al. 2008).

In summary, although available data suggest that Myb-independent, HSC-independent macrophage originate from extraembryonic progenitors, their intraembryonic origin cannot be excluded. It is therefore important to keep in mind that FL hematopoiesis should not be equated to fetal HSC-derived hematopoiesis and a fortiori to adult BM hematopoiesis, and that YS hematopoiesis cannot be opposed to FL hematopoiesis. Fate-mapping strategies or transplantation experiments are useful to assess the potential of such progenitors but do not allow drawing quantitative conclusions. Identifying novel transcription factors regulating Myb-independent myelopoiesis would help to investigate in vivo the development and biology of these cells.

BM-Derived Myeloid Cells and Myb-Independent Macrophages Have Different Functions

Myb-independent macrophages and Myb-dependent myeloid cells may mediate distinct roles in response to tissue damage and infection (Fig. 2). In contrast to monocytes recruited to the inflamed brain, which produce inflammatory mediators and appear to correlate with disease progression and poor outcome (Ajami et al. 2011), microglia neither activate NF-κB signaling nor secrete inflammatory mediators in exacerbated tissue inflammation. Rather, they seem to protect against neuronal death (Saijo et al. 2009). Similarly, lung macrophages have been proposed to respond to infection-induced tissue damage rather than to the infection itself (Jamieson et al. 2013). Kupffer cells, the Myb-independent macrophages of the liver, accumulate in necrotic areas of liver failure patients and may release anti-inflammatory mediators, whereas recruited monocytes may exacerbate tissue inflammation (Antoniades et al. 2012). Compatible with this hypothesis...
is the observation that in patients undergoing BMT, irradiation leads to the slow “replacement” of tissue macrophages by monocyte-derived cells in the lung (Thomas et al. 1976), whereas in the alveoli, the repopulating cells have a different morphology and fail to respond to a Candida challenge (Winston et al. 1982), suggesting that tissue-resident alveolar macrophages and BM-derived cells have different functions.

They may also have different responses in situation of metabolic imbalance. Kupffer cells in the liver (Bieghs et al. 2010) and peritoneal macrophages (Li et al. 2004) uptake lipid and become “foamy” in a high-fat diet. This is also the case of BM-derived monocytes/macrophages, at least in vitro. However, although lipid uptake by BM-derived monocytes was shown to activate NF-κB signaling possibly through toll-like receptors (TLRs), resulting in the secretion of proinflammatory mediators such as TNF, IL-1β, Cxcl10, and CXCL9 (Moore and Tabas 2011; Biswas and Mantovani 2012), peritoneum F4/80bright macrophages in contrast did not activate TLR-4 and NF-κB signaling and did not secrete proinflammatory molecules in response to a high-fat diet (Spann et al. 2012). This suggests that Myb-independent macrophages and BM-derived myeloid cells may present distinct responses to the same metabolic stimulus, here an excess of tissue lipids. This is in accordance with the proposition by R. Medzhitov that tissue-resident macrophages are responsible for an adaptive response to tissue stress or malfunction—which he termed parainflammation—intermediate between the basal homeostatic state and a classic inflammatory response where leukocyte extravasate (Medzhitov 2008).

Because Myb-independent macrophages develop within their tissue of residence and accompany it through development and adult life, they may play a role in morphogenesis and the maintenance of homeostasis. Microglia play an important role in the homeostasis of the surrounding neurons, by monitoring and pruning synapses (Wake et al. 2009; Paolicelli et al. 2011), phagocytosing dead cells during development and adulthood, and responding to local brain injury (Davalos et al. 2005; Nimmerjahn et al. 2005). As professional scavengers, macrophages could be seen as a sensor of the metabolic state of the tissue and its variations and provide growth factors accordingly and other mediators to tissue cells (Fig. 2).

Of note, whether the “pro-homeostatic” responses of macrophages are ultimately beneficial or detrimental in situation of chronic stress, such as, for example, a prolonged lipid-rich diet, remains to be investigated (Medzhitov 2008).

CONCLUSIONS

Macrophages and DCs are present in all tissues and are critical effectors and regulators of immune responses. The data reviewed here challenge the classical view that the MPS is a population of BM-derived cells that develops from HSCs along distinct differentiation pathways in response to internal and external cues. In contrast, we define a lineage of Myb-independent macrophages, alongside BM-derived inflammatory cells, which constitute tissue-resident macrophages and may contribute to tissue homeostasis and instruct tissue response to stress or malfunction. It will thus be important to characterize the molecular and cellular mechanisms that allow the development and maintenance of the macrophage networks, mediate their activation, and underlie their functions.

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E.G.P. wrote the first draft and E.G.P. and F.G. discussed, revised, and edited the manuscript.

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REFERENCES

Ajami B, Bennett JL, Krieger C, Tetzlaff W, Rossi FM. 2007. Local self-renewal can sustain CNS microglia maintenance and function throughout adult life. Nat Neurosci 10: 1538–1543.

Ajami B, Bennett JL, Krieger C, McNagny KM, Rossi FM. 2011. Infiltrating monocytes trigger EAE progression, but do not contribute to the resident microglia pool. Nat Neurosci 14: 1142–1149.

Akashi K, Traver D, Miyamoto T, Weissman IL. 2000. A clonogenic common myeloid progenitor that gives rise to all myeloid lineages. Nature 404: 193–197.

Antoniades CG, Quaglia A, Taams LS, Mitry RR, Hussain M, Abeles R, Possamai LA, Bruce M, McPhail M, Starling C, et al. 2012. Source and characterization of hepatic macrophages in acetaminophen-induced acute liver failure in humans. Hepatology 56: 735–746.

Auffray C, Fogg D, Garfa M, Elain G, Join-Lambert O, Kayal S, Sarnacki S, Cumano A, Lauvau G, Geissmann F. 2007. Monocyte-derived cells in the lung (Thomas et al. 2010), and peritoneal macrophages (Li et al. 2004) as professional scavengers, macrophages could be seen as a sensor of the metabolic state of the tissue and its variations and provide growth factors accordingly and other mediators to tissue cells (Fig. 2).

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REFERENCES

Ajami B, Bennett JL, Krieger C, Tetzlaff W, Rossi FM. 2007. Local self-renewal can sustain CNS microglia maintenance and function throughout adult life. Nat Neurosci 10: 1538–1543.

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Antoniades CG, Quaglia A, Taams LS, Mitry RR, Hussain M, Abeles R, Possamai LA, Bruce M, McPhail M, Starling C, et al. 2012. Source and characterization of hepatic macrophages in acetaminophen-induced acute liver failure in humans. Hepatology 56: 735–746.

Auffray C, Fogg D, Garfa M, Elain G, Join-Lambert O, Kayal S, Sarnacki S, Cumano A, Lauvau G, Geissmann F. 2007. Monocyte-derived cells in the lung (Thomas et al. 2010), and peritoneal macrophages (Li et al. 2004) as professional scavengers, macrophages could be seen as a sensor of the metabolic state of the tissue and its variations and provide growth factors accordingly and other mediators to tissue cells (Fig. 2).

Of note, whether the “pro-homeostatic” responses of macrophages are ultimately beneficial or detrimental in situation of chronic stress, such as, for example, a prolonged lipid-rich diet, remains to be investigated (Medzhitov 2008).

CONCLUSIONS

Macrophages and DCs are present in all tissues and are critical effectors and regulators of immune responses. The data reviewed here challenge the classical view that the MPS is a population of BM-derived cells that develops from HSCs along distinct differentiation pathways in response to internal and external cues. In contrast, we define a lineage of Myb-independent macrophages, alongside BM-derived inflammatory cells, which constitute tissue-resident macrophages and may contribute to tissue homeostasis and instruct tissue response to stress or malfunction. It will thus be important to characterize the molecular and cellular mechanisms that allow the development and maintenance of the macrophage networks, mediate their activation, and underlie their functions.

AUTHORS’ CONTRIBUTIONS

E.G.P. wrote the first draft and E.G.P. and F.G. discussed, revised, and edited the manuscript.

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Chorro L, Geissmann F. 2010. Development and homeostasis of resident myeloid cells: The case of the Langerhans cell. Trends Immunol 31: 438–445.

Chorro L, Sarde A, Li M, Woollard KJ, Chambon P, Malissen B, Dai XM, Ryan GR, Hapel AJ, Dominguez MG, Russell RG, Cumano A, Ferraz JC, Klaine M, Di Santo JP, Godin I. 2001. Expansion of the epidermal LC network. 2009. Langerhans cell (LC) proliferation mediates neonatal inflammation. New Engl J Med 365: 127–138.

Carlin LM, Stamatiades EG, Auffray C, Hanna RN, Glover L, Godin IE, Garcia-Porrero JA, Coutinho A, Dieterlen-Lievre F. 2003. Blood monocytes of the mouse colony-stimulating factor 1 receptor gene results in osteopetrosis, mononuclear phagocyte deficiency, in vivo. Blood 101: 375–386.

Clemens RK, Lutz M, Ferrer A, Naus D, Jockusch BM. 2011. Exome sequencing identifies GATA-2 mutation as the cause of dendritic cell, monocyte, B and NK lymphoid defects. Science 334: 792–804.

Cotta-Ramusino C, Hansel D, Dejana E, Baldassarre M. 2002. Blood–brain barrier permeability after injury. Nat Neurosci 5: 105–112.

Davalois D, Grutzendler J, Yang G, Kim JV, Zuo Y, Jung S, Littman DR, Dustin ML, Gran WB. 2005. ATP mediates rapid microglial response to local brain injury in vivo. Nat Neurosci 8: 752–758.

Davies LC, Rosas M, Smith PJ, Fraser DJ, Jones SA, Taylor PR. 2011. A quantifiable proliferative burst of tissue macrophages restores homeostatic macrophage populations after acute inflammation. Nat Immunol 12: 213–216.

Delassus S, Cumano A. 1993. Para-aortic splanchnopleura from early caudal intraembryonic splanchnopleura. J Immunol 151: 2481–2493.

Disease maps: Observations made on a composite tissue allograft. PloS One 8: e58544.

Ding W, Kettman R, Nucifora G, Shi Y, Wang B, Pannaraj PS, et al. 2013. Conditional Nr4a1−/−dependent Ly6Clow monocytes monitor postcoitus. Hepatology 57: 139–153.

Disease maps: Observations made on a composite tissue allograft. PloS One 8: e58544.

Ding W, Kettman R, Nucifora G, Shi Y, Wang B, Pannaraj PS, et al. 2013. Conditional Nr4a1−/−dependent Ly6Clow monocytes monitor postcoitus. Hepatology 57: 139–153.
impaired proliferation and accelerated differentiation. Proc Natl Acad Sci 106: 21689–21694.

Liu K, Wiklund U, Yao K, Tish K, Nussenzweig M. 2007. Origin of dendritic cells in peripheral lymphoid organs of mice. Nat Immunol 8: 578–583.

Liu K, Victoria GD, Schwickert TA, Guermonprez P, Meredith MM, Yao K, Chu FF, Randolph GJ, Rudensky AY, Nussenzweig M. 2009. In vivo analysis of dendritic cell development and homeostasis. Science 324: 392–397.

Liu CT, Yashihoro M, McGrath K, Coyne SJ, Palis J, Yoder MC. 2008. All primitive and definitive hematopoietic progenitor cells emerging before E10 in the mouse embryo are products of the yolk sac. Blood 111: 3435–3438.

MacDonald KP, Rowe V, Bofinger HM, Thomas R, Sasmono T, Hume DA, Hill GR. 2005. The colony-stimulating factor 1 receptor is expressed on dendritic cells during differentiation and regulates their expansion. J Immunol 175: 1399–1405.

McKenna HJ, Stocking KL, Miller RE, Brasel K, De Smedt T, Liu K, Waskow C, Liu X, Yao K, Hoh J, Nussenzweig M. 2007. Microglia in the adult brain arise from Ly-6ChiCCR2+ monocytes emerging before E10 in the mouse embryo. Nature 445: 428–435.

Mack M, Heikenwalder M, Bruck W, Priller J, Prinz M, Hara T, Watanabe T. 1999. Hematopoietic cells in cultures of murine embryonic aorta-gonad-mesonephros region are induced by stromal-89. Lab Invest 81: 527–530.

Mack M, Heikenwalder M, Bruck W, Priller J, Prinz M. 2007. Microglia in the adult brain arise from Ly-6ChiCCR2+ monocytes emerging before E10 in the mouse embryo. Nature 454: 21689–21694.

Mack M, Heikenwalder M, Bruck W, Priller J, Prinz M. 2007. Identification of clonogenic common Flt3+ M-CSFR+ plasmacytoid and conventional dendritic cell progenitors in mouse bone marrow. Nat Immunol 8: 1207–1216.

Paolicelli RC, Bolosac G, Pagani F, Maggi L, Scianni M, Panzaneli P, Giustetto M, Ferreira TA, Guiducci E, Dumas L, et al. 2011. Synaptic pruning by microglia is necessary for normal brain development. Science 333: 1456–1458.

Perreault C, Pelletier M, Landry D, Gyger M. 1984. Study of Langerhans cells after allogeneic bone marrow transplantation. Blood 63: 807–811.

Pforte A, Gerth C, Voss A, Beer B, Haussinger K, Jutting U, Burger G, Ziegler-Heitbrock HW. 1993. Proliferating alveolar macrophages in BAL and lung function changes in interstitial lung disease. Eur Respir J 6: 951–955.

Qian BZ, Li J, Zhang H, Kitamura T, Zhang J, Campion LR, Kaiser EA, Snyder LA, Pollard JW. 2011. CCL2 recruits inflammatory monocytes to facilitate breast-tumor metastasis. Nature 475: 222–225.

Rae F, Woods K, Sasmono T, Campanale N, Taylor D, Ovchinnikov DA, Grimmond SM, Hume DA, Ricardo SD, Little MH. 2007. Characterisation and trophic functions of murine embryonic macrophages based upon the use of a Csf1r-EGFP transgene reporter. Dev Biol 308: 232–246.

Rampon C, Huber P. 2003. Multilineage hematopoietic progenitor activity generated autonomously in the mouse yolk sac: Analysis using angiogenesis-defective embryos. Int J Dev Biol 47: 273–280.

Sasmono T, RT, Oecandy D, Pollard JW, Tong W, Pavli P, Wainwright BJ, Ostrowski MC, Himes SR, Hume DA. 2003. A macrophage colony-stimulating factor receptor-green fluorescent protein transgene is expressed throughout the mononuclear phagocyte system of the mouse. Blood 101: 1155–1163.

Sawyer RT, Strausbauch PH, Volkman A. 2006. Monocyte emigration from bone marrow by chemokine receptor CCR2. Anatom Rec 258: 1127–1132.

Spann NJ, Garmire LX, McDonald JD, Myers DS, Milne SB, Shibata N, Reichart D, Fox JN, Shaked I, Heudobler D, et al. 2015. In vivo analysis of dendritic cell development and homeostasis. Science 350: 1314–1318.
2012. Regulated accumulation of desmosterol integrates macrophage lipid metabolism and inflammatory responses. Cell 151: 138–152.

Srinivas S, Watanabe T, Lin CS, William CM, Tanabe Y, Jessell TM, Costantini F. 2001. Cre reporter strains produced by targeted insertion of EYFP and ECFP into the ROSA26 locus. BMC Dev Biol 1: 4.

Sumner R, Crawford A, Mucenski M, Frampton J. 2000. Initiation of adult myelopoiesis can occur in the absence of c-Myb whereas subsequent development is strictly dependent on the transcription factor. Oncogene 19: 3335–3342.

Thomas ED, Ramberg RE, Sale GE, Sparkes RS, Golde DW. 1976. Direct evidence for a bone marrow origin of the alveolar macrophage in man. Science 192: 1016–1018.

Thorod P, Heldmann U, Gomes-Leal W, Gisler R, Darsalia V, Tancrea J, Nygren JM, Jacobsen SE, Kokaia Z, et al. 2009. Long-term accumulation of microglia with pro-neurogenic phenotype concomitant with persistent neurogenesis in adult subventricular zone after stroke. Glia 57: 835–849.

Tsou CL, Peters W, Si Y, Slaymaker S, Aslanian AM, Weissberg SP, Mack M, Charo IF. 2007. Critical roles for CCR2 and MCP-3 in monocyte mobilization from bone marrow and recruitment to inflammatory sites. J Clin Invest 117: 902–909.

Ueno H, Weissman IL. 2010. The origin and fate of yolk sac hematopoiesis: Application of chimera analyses to developmental studies. Int J Dev Biol 54: 1019–1031.

van Furth R, Cohn ZA. 1968. The origin and kinetics of mononuclear phagocytes. J Exp Med 128: 415–435.

van Furth R, Cohn ZA, Hirsch JG, Humphrey JH, Spector WG, Langevoort HL. 1972. The mononuclear phagocyte system: A new classification of macrophages, monocytes, and their precursor cells. Bull World Health Organ 46: 845–852.

Wake H, Moorhouse AJ, Jinno S, Kohsaka S, Nabekura J. 2009. Resting microglia directly monitor the functional state of synapses in vivo and determine the fate of ischemic terminals. J Neurosci 29: 3974–3980.

Waskow C, Liu K, Darrasse-Jeze G, Guermonprez P, Ginhoux F, Meral M, Shengelia T, Yao K, Nussenzweig M. 2008. The receptor tyrosine kinase Flt3 is required for dendritic cell development in peripheral lymphoid tissues. Nat Immunol 9: 676–683.

Weissman IL, Warnke R, Butcher EC, Rouse R, Levy R. 1978. The lymphoid system. Its normal architecture and the potential for understanding the system through the study of lymphoproliferative diseases. Hum Pathol 9: 25–45.

Winston DJ, Territo MC, Ho WG, Miller MJ, Gale RP, Golde DW. 1982. Alveolar macrophage dysfunction in human bone marrow transplant recipients. Am J Med 73: 859–866.

Yamada M, Naito M, Takahashi K. 1990. Kupffer cell proliferation and glucan-induced granuloma formation in mice depleted of blood monocytes by strontium-89. J Leukocyte Biol 47: 195–205.

Yoder MC, Hiatt K. 1997. Engraftment of embryonic hematopoietic cells in conditioned newborn recipients. Blood 89: 2176–2183.

Yona S, Kim KW, Wolf Y, Mildner A, Varol D, Breker M, Strauss-Ayali D, Vinokov S, Guilliams M, Misharin A, et al. 2013. Fate mapping reveals origins and dynamics of monocytes and tissue macrophages under homeostasis. Immunity 38: 79–91.

Zovein AC, Turlo KA, Ponec RM, Lynch MR, Chen KC, Hofmann JJ, Cox TC, Gasson JC, Iruela-Arispe ML. 2010. Vascular remodeling of the vitelline artery initiates extravascular emergence of hematopoietic clusters. Blood 116: 3435–3444.