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A Csf1r-EGFP Transgene Provides a Novel Marker for Monocyte Subsets in Sheep

Clare Pridans,* Gemma M. Davis,* Kristin A. Sauter,* Zofia M. Lisowski,* Yolanda Corripio-Miyar,*† Anna Raper,* Lucas Lefevre,* Rachel Young,* Mary E. McCulloch,* Simon Lillico,* Elspeth Milne,† Bruce Whitelaw,* and David A. Hume*

Expression of Csf1r in adults is restricted to cells of the macrophage lineage. Transgenic reporters based upon the Csf1r locus require inclusion of the highly conserved Fms-intronic regulatory element for expression. We have created Csf1r-EGFP transgenic sheep via lentiviral transgenesis of a construct containing elements of the mouse Fms-intronic regulatory element and Csf1r promoter. Committed bone marrow macrophage precursors and blood monocytes express EGFP in these animals. Sheep monocytes were divided into three populations, similar to classical, intermediate, and nonclassical monocytes in humans, based upon CD14 and CD16 expression. All expressed EGFP, with increased levels in the nonclassical subset. Because Csf1r expression coincides with the earliest commitment to the macrophage lineage, Csf1r-EGFP bone marrow provides a tool for studying the earliest events in myelopoiesis using the sheep as a model. The Journal of Immunology, 2016, 197: 000–000.

The development of macrophages requires signaling from the CSF1 receptor (CSF1R) initiated by one of two ligands, CSF1 and IL-34 (1, 2). Expression of Csf1r mRNA in adult mice is restricted to cells of the macrophage lineage (3). The lineage-restricted expression of Csf1r provides the basis for the development of macrophage-specific transgenes. MacGreen (Csf1r-EGFP) mice were created previously by placing EGFP expression under the control of a 7.2-kb region of the murine Csf1r proximal promoter. Expression of the transgene is dependent on the inclusion of the highly conserved Fms-intronic regulatory element (FIRE), located in the second intron of Csf1r (4). MacGreen mice express EGFP in the same locations as the endogenous gene and have been widely used to visualize macrophages (5–8). Embryonic expression of the MacGreen transgene is restricted to macrophages and trophoblasts (4), and expression of the latter relies on a conserved 150-bp region of the Csf1r promoter. The trophoblast promoter region was omitted from the construct used in the generation of the MacBlue (Csf1r-Gal4VP16/ UAS-ECFP) transgenic mice (9). In these mice, ECFP is absent from osteoclasts and the majority of tissue macrophages, but expression is retained in microglia and Langerhans cells (10). MacGreen and MacBlue mice were created by standard pronuclear injection techniques, a relatively inefficient gene delivery tool when compared with lentiviral-based methods (11). To increase the efficiency of macrophage-specific transgene delivery, we created a lentiviral vector containing elements of murine Csf1r (12). The Csf1r promoter (0.5-kb including the start codon) was fused to the coding sequence of EGFP and cloned upstream of the intronic region containing FIRE. The lentivirus (Csf1r-EGFP-FIRE) was capable of directing EGFP expression in mouse, rat, human, pig, cow, sheep, and even chicken macrophages in vitro. This remarkable cross-species reactivity led us to produce Csf1r-EGFP sheep via injection of lentivirus into the perivitelline space of fertilized embryos. This is a highly efficient method to generate transgenic sheep (13). In addition to being economically important, sheep are increasingly being used as models for human diseases such as Huntington disease (14) and Batten disease (15). They themselves are susceptible to a variety of diseases such as paratuberculosis or Johne disease where macrophages are known to play a role (16). Although ovine macrophages have been studied in response to infection (17, 18), little is known about the characteristics of their monocytic precursors. In contrast, monocyte subpopulations have been well described in human, mouse, and pig (reviewed in Ref. 19). In this article, we describe the generation and characterization of Csf1r-EGFP sheep. Monocytes and their precursors express EGFP, yet expression of the transgene is down-regulated in differentiated macrophages. These sheep provide a useful tool for studies involving subpopulations of monocytes and their precursors.

Materials and Methods

Animals

Approval was obtained from The Roslin Institute’s and the University of Edinburgh’s Protocols and Ethics Committees. The experiments were carried out under the authority of a U.K. Home Office Project License under the
regulations of the Animals (Scientific Procedures) Act 1986. Sheep and mice were euthanized via captive bolt or CO₂ asphyxiation, respectively.

Constricts and lentivirus preparation

Constricts and preparation of lentivirus (Csf1r-EGFP-FIRE) was performed as described by Pridans et al. (12). The viral titer was 2.2 × 10⁷ transducing U/ml as assayed by endpoint dilution on the D-17 cell line.

Embryo manipulation and generation of transgenic sheep

Zygotes were obtained via methods described by Ritchie et al. (20) using abattoir-derived oocytes and frozen sperm from a Shetland ram. The lentivirus was injected into the perivitelline space of zygotes that were developed in vitro until the blastocyst stage (days 6–7), at which point they were transferred into the uterine horn of recipient ewes. The integration of the transgene in lambs was investigated by PCR analysis of ear biopsy DNA, amplifying EGFP using primers 5′-GCAAGCTTCTCTCCGTCGCGCATGCC-3′ (forward) and 5′-CGGGATCTTGAAGTTGACCTTGGCGC-3′ (reverse). Plasmid DNA (Csf1r-EGFP-FIRE) and genomic DNA from a wild type sheep were used as positive and negative controls, respectively.

Isolation of PBMCs from sheep

Bags containing citrate phosphate dextrose adenine (Infusion Concepts, Sowerby Bridge, U.K.) were used to collect blood from live animals <1 y of age. Blood was layered onto an equal volume of Lymphoprep (Axis-Shield, Oslo, Norway) and centrifuged at 1200 × g for 25 min with no brake. The PBMC layer was washed in an equal volume of PBS containing 2% FCS (FACS buffer). Any contaminating RBCs were removed by resuspending the pellet in 1 ml RBC lysis buffer (BioLegend, London, U.K.) and immediately topping up to 50 ml with FACS buffer. After centrifugation (400 × g, 5 min) cells were washed once with buffer.

Bone marrow and alveolar macrophage isolation from sheep

Bone marrow (BM) and alveolar macrophages (AMs) were isolated as described by Kapetanovic et al. (21), except RBC lysis buffer (BioLegend) was used and the medium was RPMI 1640 supplemented with 20% sheep serum, 100 U/ml penicillin, 100 µg/ml streptomycin, and 1 mM glucose (Glu-taMAX (Life Technologies, Paisley, U.K.).

In vitro differentiation of sheep macrophages

Freshly isolated BM and PBMCs were cultured on bacteriological plates at a density of 2 × 10⁵ and 3 × 10⁵ cells/cm², respectively. Complete medium was supplemented with 10% FCS (100 ng/ml recombinant human CSF1 (rhCSF1); a gift from Chiron, Emeryville, CA). Fresh media containing rhCSF1 was added on day 4 and cells analyzed on day 7.

Phagocytosis assays using macrophages and whole blood from sheep

Freshly isolated PBMCs and BM from Csf1r-EGFP sheep were cultured in a two-well Lab-Tek chamber slides (Nunc) at a density of 5 × 10⁵ cells/cm² as described earlier. Zymosan (Saccharomyces cerevisiae) BioParticles labeled with Alexa Fluor (AF) 594 (Thermo Fisher Scientific, Waltham, MA) were added at 8 × 10⁵ particles/well and incubated for 2 h at 37°C. After addition of ice-cold PBS, cells were washed four times with PBS and fixed in 4% paraformaldehyde for 20 min at room temperature. Cells were washed in PBS and viewed by confocal microscopy. For comparison of phagocytic activity between wild type and Csf1r-EGFP sheep, whole blood was collected in EDTA tubes and assays performed based on a modified protocol from Bicker et al. (22). pHred Red Escherichia coli BioParticles (Thermo Fisher) were added to an equal volume of whole blood and centrifuged at 34 × g, 37°C for 1 h. A control blood sample containing the BioParticles was left on ice during this time. Samples were incubated on ice for 10 min and then stained with mouse anti-human CD14 AF647 (clone TUK4, 1:20; BD Serotec, Kidlington, U.K.) for 1 h on ice. Blood was then prepared for flow cytometry using DAKO Uti-lyse erythrocyte lysing solution (Dako) using Zombie Violet (BioLegend) to exclude dead cells. CSF1-Fc (6) was labeled with an AF647 labeling kit (Thermo Fisher Scientific). Analysis was performed with FlowJo software (FlowJo, Ashland, OR).

Cell sorting and cytospins

Freshly isolated BM and PBMCs were used for sorting EGFP populations on a FACS Aria III cell sorter (BD). SYTOX Blue (Life Technologies) was used to exclude dead cells. Cells were stained with Leishman’s stain and mounted with DPX mounting media.

cDNA synthesis and PCR

Total RNA from sheep BM-derived macrophages (BMDMs) was isolated using TRIzol (Invitrogen), and the aqueous phase was purified with an RNasey Mini kit (Qiagen). cDNA was prepared as in Pridans et al. (12). For standard PCR, EGFP and ovine Csf1r were amplified with Invitrogen Taq polymerase using the following oligonucleotides: EGFP 5′-CCACAAGGTTCAGGTTGGTCTC-3′ and 5′-CTTGGACAGCTGCTTGGACATC-3′; Csf1r 5′-AGTCTTGACCTGAACTCAACTG-3′ and 5′-GGGGTGTAGCTTGAGGTGGTGTAT-3′. For quantitative PCR, cDNA was amplified with Power SYBR Green PCR Master Mix using the 7500 fast Real Time PCR system (Applied Biosystems, Thermo Fisher Scientific). The oligonucleotides used were: Hpr 5′-GACACTTGGAGAACAGATGCA-3′ and 5′-GCTTCTTTCTACCGAAGCT-3′; Csf1r 5′-TTGGTAGTACCTTCCATCGATC-3′ and 5′-CTTGTGATTCGCAACCATTGC-3′. Primer efficiency was validated with a standard curve of four serial dilution points (efficiency ranging between −3.28 and −3.38), and tests for nonamplification of genomic DNA were carried out systematically. Data were normalized according to the 2-ΔCq model (24).

Western blot

Radioimmunoprecipitation assay lysates (50 µl of Tris pH 7.0, 150 mM of NaCl, 0.1% SDS, 1% Igepal CA-630, 1.27 µM of sodium deoxycholate) were prepared from CSF1R-starved BMDMs (1 ml per 5 × 10⁵ cells) and EGFP RAW264.7 cells (12). Samples were mixed with 4× loading dye (0.25 M Tris-HCl pH 6.8, 8% SDS, 30% glyceral, 0.02% bromophenol blue, 10% 2-ME), heated for 5 min at 95°C, run on a 4–12% gradient SDS-PAGE gel, and transferred onto a polyvinyliden difluoride membrane as per Bio-Rad instructions. The membrane was blocked using a GPP Tag mAb (Life Technologies) and β-actin Ab (C4; Santa Cruz Biotechnology). Secondary Abs were anti-rabbit and -mouse IgH RHP (Cell Signaling Technology).

Results

Production and screening of Csf1r-EGFP transgenic sheep

Our recent in vitro study of a lentiviral vector containing control elements of murine Csf1r revealed macrophage-specific gene reporter expression in multiple species including sheep (12). Zygotes were injected with lentivirus (Csf1r-EGFP-FIRE [12]) and developed to blastocysts in vitro to create germline transgenic Csf1r-EGFP sheep. Eight recipient ewes were transplanted with either two or three blastocysts, which resulted in three pregnancies. Four founder lambs (two male and two female) were born and all were positive for the transgene (EGFP) via confocal microscopy and PCR (Fig. 1A). The LPS receptor CD14 is a monocyte marker (25) and was used to initially screen the founders by flow cytometry. Two reporter gene expression patterns were observed: all CD14+ monocytes expressed EGFP in two of the lambs (Fig. 1Bi), whereas the other two lambs had a population of CD14+EGFP- monocytes
Of the 13 lambs born, 10 were Blackface ewes. Seven lambs were born to the CD14+EGFP+ founder males (as shown in Fig. 1B) were bred to wild type did not express CD14, which is discussed later (Fig. 1B). The two macrophage growth factor, CSF1 (1, 2). Monocytes in MacGreen Differentiation of these subsets appears to be controlled by the markers including CD14, CD16, CX3CR1, and CD163 (29–31).

Peripheral blood monocytes in mice, humans, and pigs can be analyzed by flow cytometry. To exclude autofluorescent eosinophils (27) and other granulocytes, we gated the marrow population on lymphocytes/monocytes based on forward light scatter (FSC) and side scatter (SSC) profiles (Fig. 2A). The small mononuclear population had an average of 28.5% EGFP+ cells (n = 5) and could be divided into EGFPHi and EGFPlo populations. There was an average of 1.7% EGFPlo cells that were CD16+CD172a+ and expressed an increasing level of CD14. The EGFPlo cells were more heterogeneous. There was a wider range of CD14 expression than the EGFPhi cells, and only a small percentage (<10%) also expressed CD16 and CD172a (Fig. 2B). Leishman staining of sorted EGFP+ cells revealed that the majority of cells were monocytoid. The large blast cells were characteristic of myeloblasts and monoblasts (28) (Fig. 2C).

Analysis of PBMCs from Csf1r-EGFP sheep

Peripheral blood monocytes in mice, humans, and pigs can be divided into functional subsets based upon expression of surface markers including CD14, CD16, CX3CR1, and CD163 (29–31). Differentiation of these subsets appears to be controlled by the macrophage growth factor, CSF1 (1, 2). Monocytes in MacGreen mice are EGFP+ (4). To analyze EGFP expression in monocytes from the sheep, blood samples were taken from live animals and PBMCs purified using a density gradient. Sheep are known to have a lower percentage of blood monocytes compared with humans or mice (32), and an average of 4.2% of PBMCs expressed EGFP (range 1.2–9.9%). EGFPlo PBMCs could be divided into EGFPhi and EGFPlo populations (Fig. 3A). The EGFPhi cells were CD14+CD16+++, characteristic of human classical and intermediate monocytes (33) whereas the EGFPlo cells displayed the nonclassical monocyte phenotype (CD14+ CD16++). Both populations expressed CD172a (Fig. 3B), and FSC could be used to separate the EGFPlo cells into two distinct populations. Nonclassical monocytes are known to be smaller than classical monocytes (34), and Leishman staining on sorted cells revealed that both subsets presented monocyte morphology (Fig. 3C). CD16+ human monocytes express higher levels of Csf1r mRNA than CD16− monocytes (35), and this could be reflected in the increase in EGFP expression in CD16+ monocytes from Csf1r-EGFP sheep.

Downregulation of the EGFP transgene in macrophages of sheep

Treatment of BM cells or blood monocytes from other species with CSF1 can promote maturation into macrophages in vitro (4, 21, 36). To analyze EGFP expression in macrophages from Csf1r-EGFP sheep, we isolated AMs via lung lavage and PBMCs/BM were differentiated in vitro in the presence of rhCSF1. The resulting cell populations were analyzed for surface expression of CD14, CD16, and CD172a. All three cell populations retained high levels of surface CD14, but they differed in expression of CD16 and CD172a. AMs were the only cell type to express CD16, and BMDMs expressed very low levels of CD172a (Fig. 4). In all of these populations of cells, detectable expression of the EGFP transgene was extinguished. Hence the transgene provides a novel marker for blood monocytes that is lost from macrophages.

To determine when EGFP expression was lost during differentiation, we cultured BM cells in rhCSF1 and analyzed EGFP expression via flow cytometry (Fig. 5A). After 24 h in culture with rhCSF1, the EGFP+ population was no longer visible and 50% of the cells expressed low levels of EGFP. This pattern of expression continued until day 4, when there was a slight increase in the granularity of the cells. After 7 d in culture, once the BM had differentiated into macrophages, EGFP expression was no longer detected. Expression of Csf1r mRNA is downregulated by CSF1 (37), and macrophages express lower levels of Csf1r compared.
with monocytes (38). Because the levels of EGFP in Csf1r-EGFP sheep are very low, downregulation of Csf1r during differentiation would likely result in loss of detectable EGFP. Indeed, comparison of EGFP in PBMCs from Csf1r-EGFP sheep and mice demonstrates the sheep express 1 log-fold less EGFP (Fig. 5B) and still express EGFP mRNA in BMDMs (Fig. 5C). Mature macrophages in Csf1r-EGFP mice are easily detected via FACS and microscopy (4). Notwithstanding the lower expression, EGFP could still be detected in the sheep macrophages via Western blot (Fig. 5D). As we have previously shown that Csf1r-EGFP lentivirus is capable

FIGURE 2. BM of Csf1r-EGFP sheep contains a heterogeneous EGFP+ population of cells. BM from Csf1r-EGFP sheep was isolated from ribs and analyzed via flow cytometry. (A) Gating strategy. Dead cells were excluded with propidium iodide staining and granulocytes excluded via FSC/SSC. EGFPlo and EGFPhi were selected once quadrants were set with negative control BM. (B) Analysis of CD14, CD16, and CD172a expression in GFPlo and GFFhi cells. Quadrants were set using isotype controls for each population. (C) Leishman-stained EGFP+ cells (combined EGFPlo and EGFPhi). Scale bar, 10 µm. Dot plots and Leishman staining of EGFP+ cells representative of five and two sheep, respectively. Leishman staining was representative of 15 images per animal.

FIGURE 3. Monocytes are EGFP+ in Csf1r-EGFP sheep. Peripheral blood monocytes were isolated from the blood of live animals and analyzed via flow cytometry. (A) EGFP+ gating strategy. (B) Expression of CD14, CD16, and CD172a in EGFPlo and EGFPhi PBMCs. Quadrants were set using isotype controls for each population. (C) Leishman-stained EGFP+ cells. Scale bars, 10 µm. Dot plots and Leishman staining of EGFP+ cells are representative of nine and three sheep, respectively. Leishman staining was representative of 10 images per animal.
of transducing primary sheep macrophages in vitro to generate an EGFP+ population (12), producing $Csf1r$-EGFP sheep with a higher titer of lentivirus may result in higher levels of EGFP+ in macrophages.

Functional analyses of monocytes and macrophages from $Csf1r$-EGFP sheep

MacGreen ($Csf1r$-EGFP) mice express the transgene in the same locations as the endogenous gene (4). To verify whether expression of the transgene in sheep was restricted to cells expressing $Csf1r$, we used fluorescently labeled CSF1-Fc (6) to detect $Csf1r$ in whole blood via flow cytometry. Both EGFPlo and EGFPhi PBMCs bound the labeled CSF1-Fc, whereas the EGFPhi PBMCs did not (Fig. 6A). Surprisingly, the granulocyte population showed low levels of CSF1-Fc binding. Murine granulocytes from the MacGreen mouse express $Csf1r$ mRNA and are EGFP+, yet they do not express $Csf1r$ on the cell surface (39). There is evidence that human granulocytes express functional $Csf1r$, albeit at lower levels than CD14+ monocytes (40). We examined EGFP expression in sheep granulocytes. Consistent with the level of CSF1-Fc binding, and identical to the levels of $Csf1r$ in humans, this population expressed low levels of EGFP (Fig. 6A). Expression of endogenous $Csf1r$ was not affected by the transgene; the level of mRNA detected by quantitative PCR in BMDMs from wild type and transgenic sheep was not significantly different (Fig. 6B).

BM and PBMCs from $Csf1r$-EGFP sheep differentiated into macrophages in the presence of rhCSF1 (Figs. 4, 5). Phagocytosis assays were performed to further investigate whether the transgene had any effect on monocyte and macrophage function in $Csf1r$-EGFP sheep. Both monocyte-derived macrophages (MDMs) and BMDMs phagocytosed Zymosan A $S. cerevisiae$ BioParticles within 2 h (Fig. 6C), highlighting that they were indeed functional macrophages. To compare the function of monocytes from wild type and $Csf1r$-EGFP sheep, we performed phagocytosis assays on whole blood. There was no difference in the capability of CD14+ monocytes to phagocytose $E. coli$ (Fig. 6D), suggesting that the transgene also had no effect on the function of monocytes.

The development of mature APCs relies on the expression of surface MHC class II molecules. We assessed the expression in wild type and transgenic monocytes via flow cytometry and found
that both groups had CD14+ monocytes that expressed MHC class II (Fig. 6E). In humans, the highest expression levels of MHC class II are found on CD16+ monocytes (30). As in humans, all monocytes express high levels of MHC class II and it is unaffected in the Csf1r-EGFP sheep. Again, in common with humans, the CD16+ monocytes expressed higher levels than the CD14++ subset.

**Discussion**

CSF1 controls the proliferation, differentiation, and survival of monocytes and macrophages and their BM progenitors (41), and the receptor, CSF1R, provides a marker for cells of the mononuclear phagocyte lineage. MacGreen (Csf1r-EGFP) mice express the transgene in the same locations as the endogenous gene, and expression is dependent on FIRE (4). FIRE acts as an antisense promoter in macrophages (42) and is remarkably conserved from humans to reptiles (43). It is even more highly conserved than the proximal promoter. We have recently described a lentivirus containing control elements of murine Csf1r capable of driving transgene expression in macrophages from multiple species, including sheep, in vitro (12). This lentivirus was used in the generation of Csf1r-EGFP transgenic sheep. Alignment of the Ensembl predicted sheep Csf1r gene with the murine promoter, and FIRE reveals 50 and 80% identity, respectively (data not shown). All of the myeloid-expressed transcription factor binding sites contained within FIRE (44) are conserved between mouse and sheep. Our data show that the apparent conservation reflects function in that the mouse elements were sufficient to drive EGFP expression in monocytes from all four Csf1r-EGFP founder sheep.

The Csf1r promoter region in mice contains two separate promoters to drive expression in macrophages and placental trophoblast cells (45). In humans, the trophoblast promoter is located 26 kb upstream of CSF1R (46). When the human trophoblast promoter sequence is aligned against the sheep genome (Oar_v3.1/oviAri3) using BLAST-like alignment tool, the highest scoring match is located at the 3' end of Pdgfb. This suggests the ovine trophoblast promoter, as in humans, is located at least 20 kb upstream of Csf1r. Deletion of the trophoblast promoter in the MacBlue mice abolishes transgene expression in the majority of tissue macrophages (10). Hence the lack of EGFP expression in AMs from the transgenic sheep may be caused by the lack of the trophoblast promoter in the construct used to generate these animals. We have previously shown that sheep macrophages are EGFP+ after incubation with Csf1r-EGFP lentivirus, which used polybrene to increase transduction efficiency (12). The low levels of EGFP expression in Csf1r-EGFP sheep could likely be increased by the use of a higher titer lentivirus (and hence more lentiviral insertion events) or by use of the ovine instead of murine Csf1r in the lentiviral construct. However, it is also possible that the reduced expression level in sheep versus mouse is associated with either species- or lentiviral-specific methylation of the transgene. A study of EGFP+ sheep generated by lentiviral injection of zygotes using a ubiquitous promoter indicated increased methylation patterns correlated with lower EGFP intensity (47).

Monocytes are a population of leukocytes that can be functionally characterized by their ability to phagocytose, produce cytokines, and present Ag. They make up 5–10% of the PBMCs in humans and show both antigenic and morphological (size, granularity, and nuclear morphology) heterogeneity. Their initial identification was based on expression of CD14; however, variation in surface expression of Ags has led to the description of various subsets (reviewed in Ref. 48). Human and bovine monocytes can be divided into subpopulations based on surface expression of
CD14/CD16, whereas porcine and murine monocytes can be identified by CD14/CD163 and Ly6C/CX3CR1, respectively (reviewed in Ref. 19). The majority of knowledge about monocyte development in the BM stems from work performed in mice (reviewed in Ref. 49). Monocytes develop from hemopoietic stem cells in the BM via a series of progenitors such as the common myeloid progenitor, monocyte-macrophage dendritic cell progenitor, and the common monocyte progenitor, which all express CD14/CD16, whereas porcine and murine monocytes can be identified by CD14/CD163 and Ly6C/CX3CR1, respectively (reviewed in Ref. 19). The majority of knowledge about monocyte development in the BM stems from work performed in mice (reviewed in Ref. 49). Monocytes develop from hemopoietic stem cells in the BM via a series of progenitors such as the common myeloid progenitor, monocyte-macrophage dendritic cell progenitor, and the common monocyte progenitor, which all express Csf1r (50–53). These progenitors can be identified as Lin– and by differential expression of CD117 (c-kit), CX3CR1, Ly6C, and CD31. In humans, the common myeloid progenitor can be distinguished from other progenitors based on expression of CD38, CD45RA, FLT3, CD7, and CD10 (54). In-depth studies of monocyte development in larger animals have been hampered by the lack of species-specific Abs. In pigs, the Swine Workshop Cluster molecules were originally used to identify committed monocyte precursors in the BM. Swine Workshop Cluster 3 was identified as the earliest marker of myeloid cell development (55) and was later identified as CD172a (56). In cows and sheep, CD34+ progenitor cells have been identified (57, 58), yet specific analysis of monocyte progenitors has not been performed.

As noted earlier, the expression of the EGFP transgene in the sheep is downregulated by CSF1 in vitro and in tissue macrophages in vivo. Tissue macrophages depend upon CSF1 for their differentiation, and they are rapidly depleted after treatment of mice with anti-CSF1R Ab (5). The downregulation of the reporter gene by CSF1 probably reflects direct actions on FIRE. FIRE enhancer activity is acutely regulated by transcription factor Runx1, which is expressed at high levels in progenitors and acutely downregulated by CSF1 (59). There is an emerging view that most tissue macrophages are replaced by self-renewal, rather than replacement from the blood monocyte pool (60, 61). Like the MacBlue transgene in mice, which is effectively monocyte specific (62, 63), the Csf1r-EGFP lentiviral transgene could provide a useful marker to monitor monocyte extravasation and trafficking in tissues, and might be applied to other species.

Mice and humans differ immunologically, and the mouse has limitations as a model (64–66). Macrophages from the pig more closely resemble those of humans in terms of their response to bacterial LPS (21) or sheep in terms of pattern recognition receptor expression (67). Hence larger animal models are likely to represent a better model of human macrophage development. Phenotypic analysis of macrophage development in sheep has not been described. Instead, studies have focused on the response of monocytes and macrophages to infection. We have shown that macrophages from Csf1r-EGFP sheep are functional and comparable with their wild type counterparts, and that the transgene had no effect on expression of the endogenous gene. Hence, as expression of Csf1r is one of the earliest markers of macrophage lineage commitment (51), BM from Csf1r-EGFP sheep could be a valuable resource to study the earliest events in myelopoiesis in an underused species.

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Disclosures
The authors have no financial conflicts of interest.
References


