Csf1r-mApple Transgene Expression and Ligand Binding In Vivo Reveal Dynamics of CSF1R Expression within the Mononuclear Phagocyte System

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Csf1r-mApple Transgene Expression and Ligand Binding
In Vivo Reveal Dynamics of CSF1R Expression within the
Mononuclear Phagocyte System

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CSF1 is the primary growth factor controlling macrophage numbers, but whether expression of the CSF1 receptor differs between
discrete populations of mononuclear phagocytes remains unclear. We have generated a Csf1r-mApple transgenic fluorescent
reporter mouse that, in combination with lineage tracing, Alexa Fluor 647-labeled CSF1-Fc and CSF1, and a modified ΔCsf1r–
enhanced cyan fluorescent protein (ECFP) transgene that lacks a 150 bp segment of the distal promoter, we have used to dissect
the differentiation and CSF1 responsiveness of mononuclear phagocyte populations in situ. Consistent with previous Csf1r-driven
reporter lines, Csf1r-mApple was expressed in blood monocytes and at higher levels in tissue macrophages, and was readily
detectable in whole mounts or with multiphoton microscopy. In the liver and peritoneal cavity, uptake of labeled Csf1r largely
reflected transgene expression, with greater receptor activity in mature macrophages than monocytes and tissue-specific expres-
sion in conventional dendritic cells. However, CSF1 uptake also differed between subsets of monocytes and discrete populations
of tissue macrophages, which in macophages correlated with their level of dependence on CSF1 receptor signaling for survival
rather than degree of transgene expression. A double ΔCsf1r-ECFP-Csf1r-mApple transgenic mouse distinguished subpopulations
of microglia in the brain, and permitted imaging of interstitial macrophages distinct from alveolar macrophages, and pulmonary
monocytes and conventional dendritic cells. The Csf1r-mApple mice and fluorescently labeled Csf1r will be valuable resources for
the study of macrophage and CSF1 biology, which are compatible with existing EGFP-based reporter lines. The Journal of
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Abbreviations used in this article: AF647, Alexa Fluor 647; BM, bone marrow; cDC, conventional dendritic cell; DC, dendritic cell; ECFP, enhanced cyan fluorescent protein; Fli3, Fms-like tyrosine kinase 3; Flt3L, Flt3 ligand; KC, Kupffer cell; MFI, median fluorescence intensity; MHCI, MHC class I; MPS, mononuclear phagocyte system; pDC, plasmacytoid DC; WT, wild type.

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Csf1r mRNA than monocyte-derived APC (15, 22) and have been considered Csf1R independent (1). The levels of surface Csf1R largely distinguish Flt3L-dependent cDC2 from short-lived monocyte-derived CD11c<sup>−</sup> MHCII<sup>+</sup> cells in the serous cavities (23, 24). However, Csf1R expression from spleen express high levels of both Csf1r and Flt3 mRNA (www.biogps.org) and their numbers are controlled by CSF1 in vivo (25). Therefore, it remains unclear whether there is a genuine dichotomy between Csf1r and Flt3-dependent myeloid APC.

Csf1R on macrophages is continuously removed from the cell surface by endocytosis and degraded following ligand binding. For that reason, the detection of Csf1R protein by immunohistochemistry or flow cytometry does not provide a clear indication of functional expression. To identify Csf1r-expressing cells in situ, regulatory elements of the murine Csf1r locus, including a 150 bp segment of the distal promoter, were used to produce Csf1r-EGFP reporter mice (26). The same promoter construct was used to drive constitutive (27) and inducible cre-recombinase to support macrophage-specific conditional mutations (28) as well as lineage tracing (29), and these tools have been widely distributed among the research community. However, new resources are required to verify with single-cell resolution the extent to which Csf1r transgene expression reflects that of functional Csf1R protein.

In addition to aiding our understanding of the regulation of myeloid cells, visualization of Csf1r gene and protein expression may also be useful to study cell interactions in vivo due to the lack of tools to identify discrete MPS populations during multicellular imaging. A binary enhanced cyan fluorescent protein (ECPF) reporter (ΔCsf1r-Gal4VP16/UA5-EGFP) transgene with a 150 bp segment of the distal Csf1r promoter deleted, termed ΔCsf1r-ECFP, has provided a novel tool to support in vivo imaging of monocyte trafficking (30, 31), because expression was lost from the large majority of tissue macrophages but remained in blood monocytes, microglia, Langerhans cells, and cDC2 (32). In particular, dual reporter mice, such as those generated by crossing the Cx3cr1-EGFP and Ccr2-RFP mice, have been valuable tools for visualizing monocyte subsets and their differentiation in the brain and liver (33), findings that would not have been obtainable using single reporter mice. However, many other macrophage and nonmacrophage reporter genes use EGFP, rendering the original Csf1r-EGFP transgene of limited use for this purpose. Thus, additional monocyte/macrophage reporter mice that are compatible with existing EGFP-based reporters are needed.

Hence, we have created new tools and assays to image and assess Csf1r gene and protein expression that can be combined conveniently with common fluorophores, EGFP transgenes, and the ΔCsf1r-ECFP transgene for use in imaging and flow cytometry. In particular, we characterize a new Csf1r-mApple line expressing the red reporter gene mApple under the same promoter used in the Csf1r-EGFP reporter, and apply this in combination with the ΔCsf1r-ECFP transgene, lineage tracing, and labeled CSF1-Fc and CSF1 proteins to distinguish different cellular compartments within the MPS, and to dissect the homeostatic roles of CSF1.

Materials and Methods
Plasmid constructs
The 7.2 kb Csf1r reporter construct previously used to generate the Csf1r-EGFP mice (26) was digested with Apal and Sall (NEB) to remove EGFp before gel purification using the QiAquick gel extraction kit (Qiagen). Overhangs were removed with Mungbean nuclease (NEB) and DNA was purified using QIAGEN MinElute columns (Qiagen), then dephosphorylated using thermostable alkaline phosphatase (Promega). A construct encoding the fluorescent protein Csf1r-mApple (34) was digested with SmaI and AlIII, similarly purified, and overhangs removed before both constructs were precipitated with EtOH/NaOAc and then ligated with T4 ligase (NEB) at 16°C overnight. The resulting Csf1r-mApple construct was transformed into DH5α competent cells. The Csf1r-rtTA-M2 construct utilizing the same 7.2 kb mouse Csf1r promoter region was used previously to generate Csf1r-rtTA transgenic mice (35) For generation of transgenic mice, plasmid backbones were removed by digestion with DrdI/PvuI (Csf1r-mApple, NEB) and Sall/MluI (Csf1r-rtTA, Promega/NEB) and then gel-purified using a QAquick gel extraction kit. DNA was then further purified using AMPure XP beads (Agencourt) according to the instructions.

Generation of transgenic mice and animal maintenance
Animal experiments were permitted under license by the U.K. Home Office, and were approved by the University of Edinburgh Animal Welfare and Ethical Review Body. All mice including wild-type (WT) C57BL/6JolaHsd CD45.2<sup>+</sup>; congenic CD45.1<sup>+</sup>CD45.2<sup>+</sup>, Csf1r-EGFP (26), ΔCsf1r-Gal4VP16/UA5-ECFP (36), and Ccr2−/− (37) lines were bred and housed in specific-pathogen free facilities at the University of Edinburgh. Csf1r-mApple/Csf1r-rtTA transgenic mice were generated at the University of Edinburgh’s Central Biological Services Transgenic Core facility by microinjection of transgenes into the pronuclei of fertilized oocytes from C57BL/JolaHsd mice. The integration of the transgenes was determined by PCR analysis of genomic DNA isolated from ear biopsy using primers that amplified a 565 bp product between the c-fms promoter and rtTA gene, using primers 5′-TTT CAG AAC AGC AGC AGA-3′ (forward) and 5′-CTG TTC TCT CCA TAC GCA GC-3′ (reverse), and 5′-CCT ACA TGT GTG CCT AAG GA-3′ (forward) and 5′-CCT ACA TGT GTG CCT AAG GA-3′ (reverse), and amplification temperatures of 35 cycles of 94 at m, 55, and 72°C, after an initial denaturing step of 94°C for 5 min. Expression of Csf1r-mApple was verified by screening 10 μl blood for the presence of Csf1r-mApple transgene expression. One founder positive for both transgenes transmitted the transgenes to progeny and established the Csf1r-mApple/Csf1r-rtTA line (referred to as Csf1r-mApple). The Csf1r-mApple line was maintained by breeding to C57BL/6JolaHsd mice, or where specified, bred to the ΔCsf1r-ECFP line, for which subsequent analysis was performed on F1 progeny. For maintenance of the Csf1r-rtTA transgenic line, animals were initially identified by PCR analysis of genomic DNA and flow-cytometric assessment of the presence of Csf1r-mApple in blood cells, and subsequently by flow cytometry alone. For identification of myeloid populations replenished by CCR2-dependent bone marrow (BM) precursors, tissue-protected BM chimeric mice were generated as previously described (23). Briefly, anesthetized C57BL/6J CD45.1<sup>+</sup>CD45.2<sup>+</sup> congenic mice were exposed to a single dose of 9.5 Gy γ-irradiation, while all but the hind legs and lower abdomen were protected by a 2 inch lead shield. Animals were subsequently given 2–5 × 10<sup>5</sup> BM cells from CD45.2<sup>+</sup> C57BL/6Jd mouse or Ccr2−/− mice by i.v. injection before being left for 8 wk prior to analysis of chimerism in the tissue compartments. All experiments were performed with age- and sex-matched littermate control mice and approved by the University of Edinburgh Animal Welfare and Ethical Review Body under license granted by the U.K. Home Office.

Tissue digestion and FACS analysis
Unless otherwise stated, mice were cycled by a rising concentration of CO<sub>2</sub>. Then 100 μl of blood was collected by cardiac puncture into EDTA tubes. The peritoneal cavity was lavaged with RPMI 1640 containing 2 mM EDTA or 45 mM HEPES (Invitrogen). Cadavers were subsequently perfused and lung and liver removed and chopped finely, and digested in prewarmed collagenase mixture (0.625 mg ml<sup>−1</sup> collagenase D (Roche), 0.5% Papain (2.1 collagenase), 0.5% Dispase (Roche)), 1% Bio-Lytic (BioTechnologies, Invitrogen), and 30 U ml<sup>−1</sup> DNase (Roche Diagnostics) in RPMI 1640 for 22 and 45 min respectively in a shaking incubator at 37°C before being passed through a 100 μm filter. Lung preparations were washed in PBS containing 2 mM EDTA (Life Technologies, Invitrogen) and 0.5% BSA (Sigma-Aldrich), termed FACS buffer, followed by centrifugation at 300 g for 5 min, whereas liver preparations were washed in 50 ml then 30 ml of ice-cold RPMI 1640 followed by centrifugation at 300 g for 5 min. Erythrocytes in tissue and blood were lysed using RBC lysis buffer from Sigma-Aldrich or BioLegend, respectively. All cells were maintained on ice until further use. Cellular content of the preparations was assessed by cell counting using a CASY TT counter (Roche). Equal numbers of cells or equivalent volumes of blood were stained with Zombie Aqua viability dye (Invitrogen) blocked with 0.025% anti-CD16/32 (BD), 1:10 heat-inactivated mouse serum (Invitrogen), and then surface stained with a combination of Abs in FACS buffer. The following Abs were used: F4/80 (BM8), Siglec-F (E50-2440), Siglec-F (ES22-10D8), Ly6C (HK1.4), CD11b (M1/70), CD11c (N418), MHCII
CSF1R-dependent uptake of CSF1 AF647 or CSF1-Fc AF647 is presented as ing the Fortessa and rat anti-mouse CD45 (BioLegend). Flow cytometry was acquired using FlowJo V9 software. CD45+ cells were identified as live single cells by excluding 7AAD+ or Zombie Aqua+ cells and using forward scatter area versus forward scatter height characteristics. Cells positive for CD19, CD3, Ly6G, and SiglecF, or CD19, CD3, and Ly6G were referred to as Lineage+ and were excluded prior to analysis of liver, blood and cavity cells, or lung cells, respectively, as shown in the respective figures.

For the processing of brain tissue, double transgenic mice were perfused transcardially with physiological saline and brains were removed for regional dissection into cerebellum, cortex, hippocampus, and striatum. Mixed brain cell homogenates were prepared as described (32). The single-cell suspension of each region was incubated with 1 μg ml−1 anti-CD16/32 and subsequently stained with rat anti-mouse/human CD11b (M1/70) and rat anti-mouse CD45 (BioLegend). Flow cytometry was acquired using the Fortessa x20 (Becton Dickinson) and resulting data were analyzed using FlowJo V10 software.

Inhibition of CSF1R signaling
The CSF1R kinase inhibitor GW2580 (LC Laboratories) was suspended in 0.5% hydroxypropylmethylcellulose and 0.1% Tween 20 using a Teflon glass homogenizer. Diluent control or 160 mg/kg GW2580 was administered daily for 4 d by oral gavage before mice were culled on day 5.

Alexa Fluor 647-labeled CSF1 and anti-CSF1R mAb
Preservative-free sterile anti-CSF1 mAb (clone AF598) was purchased from Bioserv (Sheffield, U.K.). Porcine CSF1 and CSF1-Fc was prepared as described previously (12). CSF1 and CSF1-Fc were conjugated to Alexa Fluor 647 (AF647) using the AF647 Microscale Protein labeling kit from Thermo Fisher Scientific according to manufacturer’s instructions, and sodium azide subsequently removed using 7k MWCO Pierce polyacrylamide spin desalting columns (Thermo Fisher Scientific). Mice were injected i.v. with 0.5 mg anti-CSF1R mAb or PBS vehicle control, followed by 5 μg CSF1-FcAF647 or PBS vehicle i.v. 10 min later. After a further 10 min, 60 μl of blood was removed by tail venipuncture, with the animals then immediately culled by cervical dislocation, and tissues perfused with PBS through the inferior vena cava. For study of CSF1 uptake in the peritoneal cavity, mice were injected i.p. with or without 0.5 mg AF598 followed by 0.5 μg CSF1-mApple or PBS vehicle 2 min later, and then culled 10 min later by exposure to increasing levels of CO2. The degree of CSF1-dependent uptake of CSF1AF647 or CSF1-FcAF647 is presented as the Δ median fluorescence intensity (MFI) calculated as the MFI for individual samples from mice given labelled CSF1 minus the average MFI from all samples pretreated with AF598.

Imaging of tissues and cells
Whole-mount imaging of freshly isolated tissues from transgenic and WT littermate control mice aged 12–15 wk was performed using a Zeiss AxioZoom.V16 fluorescence microscope. Immediately after excision, tissues were kept at 4°C and protected from light. The fluorescent signal was acquired at 500–550 and 590–650 nm for EGFP and Csfr1-mApple, respectively. Acquisition of tissue background signal was performed by imaging WT tissue with the filter used for detection of the Csfr1-mApple protein.

Ex vivo confocal imaging of tissues
Male transgenic or WT male littermates were anesthetized, as per regulations, and intravenously injected in the tail vein with 5 μg/g of weight of Lectin-1 [from Grifonia (Bandeiraea) simplicifolia] tagged with FITC (Vector Labs). After 10 min, mice were perfused transcardially with HBSS (Thermo Fisher Scientific), at a rate of 10 ml per min, and the left lobe of the liver was excised. Lungs were inflated with a solution containing 1% low melting-point agarose (Sigma-Aldrich) and, upon agarose solidification, the left lung was excised. Detection of functional CSF1R in lung myeloid subsets was performed by administering 5 μg CSF1-FcAF647 and 5 μg/g of weight of Lectin-1 i.v. injection. Mice were perfused with HBSS, prior to lung excision and inflation with agarose, as described. After dissection, liver and lung were placed on coverslip-bottom chambers and covered with a sufficient volume of HBSS to prevent the surface of tissues from drying. Chambers were kept on ice and protected from light until tissues were imaged on a Zeiss LSM 710 microscope. Laser wave-lengths for ECFP, FITC, and mApple were 405, 488, and 543 nm, respectively. Fluorescence acquisition for ECFP, FITC, and mApple signals in liver and lung was 400–480, 525–600, and 602–758 nm, respectively. Acquisition settings for lung tissue treated with CSF1-FcAF647 were 400–479, 525–583, 593–651, and 651–755 nm for ECFP, FITC, mApple, and AF647, respectively. Postprocessing of images was performed by adjusting the black/white thresholds in the software ZEN 2012 (blue edition) developed by Carl Zeiss as follows: ECFP: 0–175, FITC: 0–100, mApple: 0–75, AF647: 0–200.

Statistics
Statistical tests detailed in the figure legends were performed using GraphPad Prism 6. Where necessary, data were log-transformed to achieve equal variance.

Results
Generation of Csfr1-mApple mice
C57BL/6 mouse embryos were comicro-injected with a construct containing the 7.2 kb Csfr1 promoter region used to create the Csfr1-EGFP mice (26) upstream of mApple, along with a construct encoding the reverse tetracycline inducible transactivator rtTA-m2 under control of the same promoter (Csfr1-rtTA), previously used to generate a Csfr1-driven Tet-on system (35). mApple was used because it is brighter than its parent mCherry, refractory to photobleaching (38), suffers little from background autofluorescence, and previously enabled whole-mount imaging of the avian response to CSF1 in Csfr1-mApple reporter chickens (39). A single founder positive by PCR for both transgenes and for mApple protein in blood cells by flow cytometry was mated with a WT C57BL/6 mouse to establish the Csfr1-mApple line. PCR analysis across 77 mice revealed that Csfr1-mApple and Csfr1-rtTA transgenes were exclusively co-inherited, suggesting cotaggregation (data not shown). PCR and flow cytometry analysis of blood demonstrated the Csfr1-mApple transgene to be inherited at a frequency of 44.0% (n = 207). The utility of the cotaggregated Tet-on cassette is under investigation and is not considered further in this study but preliminary data demonstrate rtTA-m2 mRNA is expressed in peritoneal cells (data not shown).

Comparison of Csfr1-EGFP and Csfr1-mApple expression across tissue
In whole-mount fluorescence microscopy of live organs from Csfr1-mApple mice expression patterns of mApple recapitulated EGFP in Csfr1-EGFP transgenic mice (Fig. 1A–F). Large stellate mApple+ cells were observed throughout the liver, lung, epidermis, and cardiac muscle. Both transgenic strains highlighted the abundant macrophage populations of the intestinal lamina propria (Fig. 1E), and the red pulp of spleen (Fig. 1F) (26). Background fluorescence in littermate control mice was negligible (Fig. 1A–F, left panel). The Csfr1-EGFP and ΔCsfr1-ECFP transgenes have been used extensively for in vivo imaging with multiphoton and spinning disc microscopes (e.g., Refs. 20, 30, 31, 40–43), providing high-resolution analysis of macrophage motility and the extent of their ramified processes. Multiphoton imaging of whole mounts of the muscularis externa of the intestine demonstrated the high signal-to-noise ratio obtainable with the Csfr1-mApple reporter (Fig. 1G), enabling visualization of the regular network of microglial-like macrophages in this site (44, 45). Furthermore, the impact of exogenous CSF1-Fc, which regulates the function of these cells (44), could be directly visualized as an increase in cell size.

Csfr1-mApple expression by blood myeloid cells
To determine efficiency, reliability, and specificity of transgene expression, flow cytometry was performed on the blood of a
The level of Csf1r-mApple expression distinguishes monocytes, macrophages, and cDC in different tissues

To determine if transgene expression distinguished cDC and macrophages across multiple tissues, we first confirmed the identity of marker-defined MPS populations before surveying transgene expression in Csf1r-mApple mice. In the peritoneal cavity, we have demonstrated that recruited monocytes continuously replenish rare short-lived F4/80lo MHCII+ macrophages that include both CD11c+ and CD11c− cells (23), although only slowly replacing the more abundant F4/80hi resident macrophages of embryonic origin (23). Both CD11c+ and CD11c− short-lived and F4/80hi peritoneal macrophage populations express detectable surface CSF1R. In contrast, Flt3-dependent cavity cDC of non-monocyte BM origin also express CD11c+ and MHCII+ and can be found among F4/80lo− cells, but can be distinguished as CSF1R− (23, 24, 48, 49). Based upon this published gating strategy and previously assigned ontogenies (Fig. 3A) (23), Csf1r-mApple was detected in Ly6C+ monocytes, all macrophage populations, and in CD11b+ cDC1 and CD11b+ cDC2 (Fig. 3B, 3C) (24). There was a progressive increase in Csf1r-mApple intensity between Ly6C+ monocytes, CD11c-defined subsets of short-lived F4/80lo MHCII+ macrophages, and long-lived F4/80hi macrophages (Fig. 3D), consistent with the linear developmental relationship between these populations and monocytes in adult mice (23, 48, 50). Csf1r-mApple fluorescence in both CD11b+ cDC1 and CD11b+ cDC2 was lower than in monocytes (Fig. 3D), consistent with the lack of surface CSF1R (Fig. 3A). EGFP expression in Csf1r-EGFP mice replicated this pattern (Fig. 3E, 3F).

In the lung, alveolar macrophages are readily identified based upon high levels of CD11c and SiglecF (Fig. 4A) (51, 52). Interstitial cells are more heterogeneous. Some MHCII+ cells with varying levels of CD11c have been defined as macrophages based upon their expression of the Fc receptor CD64 and CSF1R dependence (51, 53), which contrasts the Flt3 dependence of CD64+ interstitial cDC2 (15). To verify that CD64 expression distinguishes pulmonary interstitial macrophages from CD11b+ cDC2, we assessed the turnover kinetics of CD64-defined MHCII+ cells and their dependence on CCR2, an established method for determining the likely monocyte dependence of tissue MPS cells (23, 54). We used a BM chimeric system in which WT mice were irradiated with organs of interest shielded to prevent irradiation-induced injury and reconstituted with congenic WT or Ccr2−/− BM. This approach results in stable nonhost chimera in blood leukocytes of ∼30% in recipients of WT BM (23, 55) (Fig. 4B, short-dashed line) and allows the turnover kinetics of tissue MPS cells to be assessed. Importantly, in recipients of Ccr2−/− BM chimerism in monocytes (Fig. 4B, long-dashed line) but not other circulating leukocytes is largely abolished (23). Notably, putative CD64+ CD11b+ MHCII+ cDC2 were completely replaced within 8 wk, consistent with the short half-life of DCs (14). This occurred in a completely CCR2-independent manner, with identical chimerism in recipients of WT and Ccr2−/− BM (Fig. 4B). In contrast, relatively few CD64+ MHCII+ cells were replaced over 8 wk, although this was completely dependent on CCR2, suggesting slow replenishment from monocytes. Thus, consistent with previous work (15), CD64 accurately defines distinct CD11b+ MPS populations. Alveolar macrophages showed no evidence of chimerism (Fig. 4B), consistent with self-maintenance (52, 56, 57). Surprisingly, replenishment of cells...
defined as cDC1 was also dependent upon CCR2. However, these cells express CCR2 within the lung environment (58) and thus may require this receptor for tissue retention. Based upon the verified ontogenies, Ly6C² monocytes, CD64⁺ interstitial macrophages, and alveolar macrophages in the lungs were all Csf1r⁻⁻ (Fig. 4C, 4D), but expression increased progressively between monocytes and mature macrophages (Fig. 4C, 4E). Both cDC populations also expressed Csf1r⁻⁻⁻, but at lower levels than monocytes (Fig. 4C–E).

In the liver, the largest phagocyte population is the Kupffer cells (KC), but a minority CD11b⁺F4/80⁺ BM-derived population may include monocytes, cDC2, and possibly F4/80⁻⁻⁻⁻⁻⁻ BM-derived macrophages (14, 29, 57). KCs [F4/80⁻⁻⁻⁻⁻⁻CD11b⁻ (29, 59, 60)] (Fig. 5A) exhibited uniformly high expression of Csf1r⁻⁻⁻, but expression increased progressively between monocytes and mature macrophages (Fig. 4C, 4E). Both cDC populations also expressed Csf1r⁻⁻⁻, but at lower levels than monocytes (Fig. 4C–E).

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Detection of functional CSF1R using fluorescent CSF1-Fc
CSF1r mRNA may be posttranscriptionally regulated (62) and the protein may be cleaved from the cell surface in response to TLR signals (63). To assess functional CSF1r expression, we investigated the ability of MPS cells to take up labeled pig CSF1-Fc fusion protein, which produces a large increase in tissue macrophage populations when injected into mice (12) or pigs (64). CSF1-Fc conjugated with AF647 (CSF1-FcAF647) was found to bind specifically to monocytes in vitro (65). CSF1-FcAF647 was injected intravenously 10 min before mice were sacrificed. In the liver, the uptake of CSF1-FcAF647 was detected in KC, monocytes, and cDC2, but not cDC1, pDC (Fig. 6A, 6B), or neutrophils (data not shown). Within cDC2, CSF1-FcAF647 binding was prevalent in CD11c⁺⁺ cells (Fig. 6C) precluding any possible confusion with the CD11c⁺⁺ MHCII⁺ subcapsular macrophages described recently (42). In the lung, the majority of Ly6C⁺ monocytes and interstitial macrophages bound CSF1-FcAF647, whereas both cDC populations were negative (Fig. 6A, 6B). Uptake of labeled CSF1-Fc by myeloid populations was reduced or abolished by the anti-CSF1r Ab, AFS98 (Fig. 6A, 6B), a weak inhibitor of receptor-ligand binding (66), and an identical labeling profile was observed following injection of a non–Fc-fused AF647-labeled porcine CSF1-Fc (CSF1-FcAF647) (data not shown). No detectable CSF1-FcAF647 was bound by alveolar macrophages (Fig. 6A, 6B) and these cells also failed to bind appreciable levels in vitro (data not shown). Labeled anti-CD45 Ab was able to access all other myeloid populations in lung (data not shown) and liver (Fig. 6D), suggesting a lack of CSF1-FcAF647 or CSF1-FcAF647 uptake by certain cDC reflects an absence of surface CSF1R expression rather than the inaccessibility of the sites they may occupy (31, 53).

Consistent with previous population-level data on CSF1 clearance (68), KC bound the highest level of labeled CSF1 per cell in a receptor-dependent manner (Fig. 6B), and considerably more per
cell than blood monocytes (Fig. 6E). In turn, Ly6C+ monocytes in lung and liver were more intensely labeled than interstitial lung macrophages or liver CD11b+ cDC2 and liver Ly6C-2 monocytes (Fig. 6B), although distinct intravascular versus parenchymal locations of these cells (31, 53) means exposure to circulating CSF1 cannot be controlled in this comparison. Of note, Ly6C+ blood monocytes, identified independently of CSF1R expression (Supplemental Fig. 1A), were more intensely labeled than the Ly6C-2 subset (Fig. 6E), despite equivalent surface expression of CSF1R (Fig. 6F) (69, 70). Thus, novel differences in capacity to bind CSF1 were revealed using this ligand-binding approach.

The relationship between Csf1r transgene and CSF1R protein expression in the peritoneal cavity was also examined in the peritoneal cavity following i.p. injection. Monocytes and CD11c- cDC2 and liver Ly6C- macrophages were identified independently of Csf1r expression (Supplemental Fig. 1A), were more intensely labeled than the Ly6C- subset (Fig. 6E), despite equivalent surface expression of CSF1R (Fig. 6F) (69, 70). Thus, novel differences in capacity to bind CSF1 were revealed using this ligand-binding approach.

The relationship between Csf1r-mApple activity and CSF1R-mediated ligand uptake was also examined in the peritoneal cavity following i.p. injection. Monocytes and CD11c- and CD11c- F4/80C CD226+ macrophages (49, 50) were identified as described in Supplemental Fig. 1B, avoiding the use of Abs to CSF1R. Neither Csf1r-mApple-10 cDC population bound appreciable levels of CSF1AF647, whereas all three macrophage populations had higher levels of receptor-dependent uptake of CSF1AF647 than Ly6C- cavity monocytes (Fig. 7A). Surprisingly, the CD11c- F4/80C macrophages exhibited the greatest uptake. These differences were not explained by differential levels of receptor-independent macropinocytosis as uptake of injected OVA–Texas Red was largely equivalent between populations (Fig. 7B). Anti-CSF1R mAb inhibited CSF1 uptake to a similar degree in each population, with a reduction between 63 and 75%. The difference in receptor activity appeared to have functional significance, as treatment of mice daily for 4 d with a CSF1R kinase inhibitor, GW2580 (71), which has been shown to inhibit proliferation of microglia (41) and pleural macrophages (11), partly depleted the CD11c- subset of F4/80C MHCII+ peritoneal macrophages alone (Fig. 7C, left graph), despite inhibiting proliferation (as evidenced by Ki67 staining) of all macrophage populations (Fig. 7C, right graph).

Csf1r-mApple;ΔCsf1r-ECFP mice allow in situ imaging of distinct mononuclear phagocytes

The ΔCsf1r-ECFP transgene was crossed previously to the Cx3cr1+/EGFP or Itgax (CD11c)-EYFP mouse to distinguish pulmonary monocytes from other myeloid lung populations (31). To determine the utility of the Csf1r-mApple mouse to facilitate in vivo imaging of different myeloid populations, we crossed it to

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**FIGURE 3.** Csf1r transgene and CSF1R protein expression in the peritoneal cavity. (A) Flow cytometric strategy to identify peritoneal cavity myeloid cells as recently described (23). (B-D) Expression of Csf1r-mApple in peritoneal cavity myeloid populations determined by flow cytometry, showing a representative WT littermate control (left) and Csf1r-mApple (right) mouse (B), and graphs depicting the mean frequency of Csf1r-mApple+ cells in each population (C) and geometric mean fluorescence intensity (GeoMFI) of Csf1r-mApple for different peritoneal leukocytes gated on Csf1r-mApple+ cells (D). (E and F) Expression of Csf1r-EGFP in peritoneal cavity myeloid populations from Csf1r-EGFP mice, showing a representative flow cytometric overlay (E) and graphs depicting the frequency and GeoMFI of Csf1r-EGFP+ cells in each population (F) across multiple mice. Representative data from one of three experiments (B–D) or a single experiment (E and F). Data are presented as mean ± SD of four mice (C, D, and F).
the ΔCsf1r-ECFP line (32, 36). As in the intestine (32), the majority of cDC1, cDC2, and Ly6C⁺ and Ly6C⁻ monocytes in the liver expressed high levels of ECFP, whereas pDC expressed intermediate levels. Neutrophils, eosinophils, and lymphocytes were negative (data not shown) as were F4/80hi KC (Supplemental Fig. 2A). All ECFP⁺ cells expressed intermediate levels of mApple (Supplemental Fig. 2B, cyan gate), whereas all mApple hi cells were ECFP⁻ and represent KC (Supplemental Fig. 2B, red gate). We combined the two Csf1r reporters with detection of endothelial cells by injection of FITC-labeled Lectin I. In confocal images the mApple⁺ cells were almost completely restricted to the liver sinusoids (e.g., pink boxes), consistent with KC (Fig. 8). In contrast, ECFP⁺mApple⁺ double-positive cells were rarely detected (e.g., white box), despite the presence of numerous ECFP⁺ mApple⁻ cells (e.g., yellow box) (Fig. 8). These data suggest the intermediate levels of Csf1r-mApple expressed in ECFP⁺ cDC and monocytes (Fig. 5B, Supplemental Fig. 2A) are apparently below the threshold of detection of confocal imaging. This conclusion was supported by only weak detection of mApple expression when peripheral blood was imaged using identical microscope settings (Supplemental Fig. 3A). The Csf1r-ECFP⁺ cells in the liver (e.g., yellow box) were mainly detected outside the sinusoids and likely include the subcapsular liver macrophages that also express the ΔCsf1r-ECFP transgene (42). The high level of mApple expression in KC therefore allows imaging of these cells without detection of monocytes and other mApple⁺ cells in the liver.

In the lung, interstitial macrophages and DC were ECFP negative (Supplemental Fig. 2C) whereas the majority of alveolar macrophages and Ly6C⁺ monocytes expressed ECFP, as reported previously (31, 32). Combined, all ECFP⁺ cells were mApple⁺ and represent KC (Supplemental Fig. 2D, red gate). In confocal images of transverse lung sections, parenchymal populations broadly divided into rounded ECFP⁺mApple⁺ cells (Fig. 9, yellow box) consistent with alveolar macrophages or interstitial migratory monocytes (31, 67) and elongated stellar-shaped ECFP⁻ mApple⁺ cells (Fig. 9, white box). Injection of CSF1-Fc AF647 into the double transgenic mice selectively labeled the extravascular interstitial ECFP⁻ mApple⁺ cells (Supplemental Fig. 4, white box), visible as punctate staining indicative of internalization of labeled ligand,
FIGURE 5. Csf1r-mApple transgene expression in the liver. (A) Flow cytometric strategy to identify liver leukocytes. (B) Expression of Csf1r-mApple in liver leukocytes with histograms from a representative Csf1r-mApple (right) and WT littermate control (left) mouse. (C) Frequency of cells expressing Csf1r-mApple and (D) geometric mean fluorescence intensity (GeoMFI) of Csf1r-mApple+ cells for different liver leukocytes. (E) Frequency of donor cells within each hepatic population from tissue protected BM chimeric mice 8 wk after receiving BM from WT (white) or Ccr2−/− (black) mice. Mean donor chimerism for blood Ly6C+ monocytes is presented as short- or long-dashed lines for recipients of WT or Ccr2−/− BM, respectively. (F) Representative histograms showing forward scatter area (FSC-A) characteristics and CD26, CD11c, CD64, and PDCA1 expression (tinted) overlayed with FMO controls (open) for liver leukocytes and blood monocytes. (B–D) Representative data from one of three or (E) two experiments, with data presented as mean ± SD of four mice (C and D) or mean ± SEM of five (E) mice. The asterisk (*) indicates significant differences using multiple t tests corrected for multiple comparisons using the Holm–Sidak method.
and confirmed them to be interstitial macrophages rather than ECFP mApple\(^{2}\) pulmonary DC. Many extravascular ECFP mApple\(^{+}\) cells also took up CSF1-Fc AF647 (Supplemental Fig. 4), most likely migratory monocytes identified previously by live imaging (30, 31). In contrast, the most frequent cells observed within the pulmonary capillaries were ECFP\(^{2}\)mApple\(^{+}\) (Fig. 9D, cyan box) and failed to label with injected CSF1-FcAF647 (Supplemental Fig. 4), consistent with pulmonary neutrophils (72). Consistent with this, imaging of blood cells at identical power settings confirmed strong detection of mApple in ECFP\(^{+}\) monocytes and ECFP\(^{-}\) neutrophils (Supplemental Fig. 3B).

**Heterogeneous expression of Csf1r reporter genes in the brain**

Macrophages in the mouse embryo are ECFP positive in the \(\Delta\)Csf1r-ECFP line from their earliest appearance in the yolk sac (32). In addition to alveolar macrophages, one of the few locations in adults in which transgene expression is retained is in microglia.
Grabert et al. (73) reported differences in microglia numbers and gene expression profiles in different mouse brain regions, and changes in gene expression with age. We used the \( \text{Csf1r}^\text{mApple} \) cross to further dissect microglial heterogeneity in different brain regions. CD45 -CD11b+ classical microglia were uniformly strongly positive for mApple (Fig. 10A). Like blood monocytes and alveolar macrophages, microglia in cortex, hippocampus, and striatum were largely positive for ECFP, but in the cerebellum the percentage was much lower (37.4%, Fig. 10B). The level of ECFP in these cells was also lower, and lacked a clear peak, reminiscent of the ECFP profiles of macrophages and DC in the gut (32). The brain also contains a separable CD11b+, CD45 hi macrophage-like microglial population, a subset of which occupies perivascular locations and expresses higher levels of \( \text{Csf1r} \) than monocytes (74). By contrast to the classical microglia and blood monocytes, the CD11b+CD45 hi cells were >50% ECFP negative in all brain regions (Fig. 10B).

**Discussion**

We have developed a novel \( \text{Csf1r}^\text{mApple} \) reporter line. The \( \text{Csf1r} \) promoter construct used has been remarkably consistent in generating location and copy-number–independent expression of transgenes (75), further confirmed by the comparable pattern of \( \text{Csf1r}^\text{mApple} \) and \( \text{Csf1r}^\text{EGFP} \) transgene expression. Expression of \( \text{Csf1r}^\text{mApple} \) had no impact on numbers of tissue macrophages or circulating blood leukocytes (data not shown). With optimal microscope settings, the distinct profile of transgene expression across subsets of MPS cells allowed exclusive detection of mApple hi cells. When combined with the \( \text{Csf1r}^\text{ECFP} \) reporter gene, which selectively labels subsets of \( \text{Csf1r} \)-positive cells, \( \text{Csf1r} \)-Fc labeled with AF647, and FITC-labeled Lectin, we could identify and image lung interstitial macrophages and liver KC, and distinguish them from other myeloid cells. Despite high levels of \( \text{Csf1r}^\text{mApple} \) and \( \text{Csf1r}^\text{EGFP} \) (46) transgene expression, neutrophils are identifiable by injection of labeled Abs to Ly6G, a molecule with a negligible role in neutrophil trafficking or function (76, 77). Thus, there are numerous possibilities to produce live images of macrophage behavior and heterogeneity, particularly by combining with other established EGFP-based reporter mice.

In common with the \( \text{Csf1r}^\text{EGFP} \) reporter (78), \( \text{Csf1r}^\text{mApple} \) expression was uniformly higher in resident macrophages compared with monocytes, a difference reflected in the ability of at...
least liver and cavity macrophages to capture more CSF1 on a per-cell basis than monocytes in vivo. Consistently, peritoneal macrophages compete effectively for available CSF1 in mixed culture with proliferating BM-derived macrophages (79). The rapid uptake of CSF1 AF647 by KC is consistent with their role in regulating the circulating CSF1 concentration (68). Hence, upregulation of CSF1R expression may be a general feature of macrophage differentiation that allows them to compete for or control bioavailable CSF1. The apparent inability of alveolar macrophages to capture CSF1 is a notable departure from this tenet. However, alveolar macrophages are unaffected in adult CSF1-deficient op/op mice (80) and their replenishment from BM following irradiation is largely independent of CSF1R (81). Thus, our data are consistent with a lack of role for CSF1 in maintenance of the alveolar macrophage niche. However, it remains unclear why both alveolar macrophages and granulocytes express high levels of Csf1r transgene despite lacking a surface receptor. In the blood, Ly6C⁺ monocytes more readily took up CSF1 than their Ly6C⁻ progeny, a feature consistent with the suggestion that Ly6C⁺ monocytes regulate the availability of CSF1, thereby controlling the lifespan of the Ly6C⁻ population (57). Interestingly, higher consumption of CSF1 by classical monocytes is also evident in PBMCs from Csf1r-EGFP transgenic sheep (65), suggesting this feature is conserved across species.

Intensity of fluorescence in Csf1r-mApple mice also largely distinguished long-lived tissue-resident macrophages (KC, alveolar macrophages, and F4/80hi peritoneal macrophages) from those of more recent monocyte-origin (F4/80lo resident peritoneal and lung interstitial macrophages) but was not correlated with the ability to take up labeled CSF1. In the peritoneal cavity, the receptor activity was greatest in F4/80loCD11c² cells. Notably, these cells were selectively depleted following treatment with the CSF1R kinase inhibitor GW2580. Dynamics of loss of labeled histone 2B-GFP from peritoneal F4/80lo macrophages places the half-life for replenishment of both CD11c⁺ and CD11c⁻ subsets from monocytes at around 2 wk (23), much longer than the 4 d treatment regimen in this study. Hence, selective loss of F4/80lo CD11c⁻ cells likely results from reduced survival or retention in

**FIGURE 8.** Csf1r-mApple and ΔCsf1r-ECFP transgenes allow imaging of distinct lineages of hepatic myeloid cells. Confocal image of the surface of the left lobe of the liver of a ΔCsf1r-ECFP (A), Csf1r-mApple (B), WT (C), and Csf1r-mApple/ΔCsf1r-ECFP (D) mouse imaged ex vivo. FITC-Lectin I was injected i.v. to reveal liver sinusoidal endothelium. Scale bars represent 20 μm (A–C) or 50 μm (D).

**FIGURE 9.** Csf1r-mApple and ΔCsf1r-ECFP transgenes allow imaging of distinct lineages of pulmonary myeloid cells. Confocal image of a transverse section of lung from a ΔCsf1r-ECFP (A), Csf1r-mApple (B), WT (C), and Csf1r-mApple/ΔCsf1r-ECFP (D) mouse imaged ex vivo. FITC-Lectin was injected i.v. to reveal pulmonary vasculature. Scale bars in all panels represent 50 μm.
and pulmonary CCR2-independent cDC from CCR2-dependent reporter also provided a useful marker delineating peritoneal cavity rather than a failure of monocytes to differentiate and replenish these cells. Both F4/80<sup>lo</sup> and F4/80<sup>hi</sup> populations of peritoneal macrophages are rapidly lost upon Ab-mediated neutralization of CSF1 (24), suggesting only partial blockade of CSF1R signaling occurred with the oral inhibitor used in this study. Higher levels of CSF1R signaling are generally required for proliferation than survival of macrophages (8) and hence the uniform inhibition of Ki67 expression observed across the peritoneal macrophage compartment is consistent with a reduction of proliferation than survival of macrophages (8) and hence the uniform inhibition of Ki67 expression observed across the peritoneal macrophage compartment is consistent with a reduction of the functional diversity of classical macrophages in different organs (51, 88–90), APC differentiation is likely also organ specific. Because CSF1 drives a largely immunoregulatory program (91), the responsiveness of cDC2 to CSF1 may underlie the relatively weak APC activity in liver (92) and contribute to a tolerogenic environment in the liver (93). Similarly, competition of CSF1R<sup>+</sup> cDC2 together with KC and classical patrolling monocytes (94) for available CSF1 could contribute to a tolerogenic environment in the liver (93). Similarly, competition of CSF1R<sup>+</sup> cDC2 together with KC and classical patrolling monocytes (94) for available CSF1 could provide an explanation for the relative absence of hepatic monocyte-derived MHCI<sup>+</sup> APC.

In adult mice, labeling of cDC and macrophages in the ΔCsfr<sup>-</sup>-EGFP reporter is tissue specific (32). Using the ΔCsfr<sup>-</sup>-EGFP transgene, we highlighted the utility of the ΔCsfr<sup>-</sup>-EGFP reporter in new tools to dissect the differentiation or the proximity of individual cells to the tissue-specific factors that control its expression.

In overview, the Csfr<sup>-</sup>-mApple mouse recapitulates the expression profile of the widely used Csfr<sup>-</sup>-EGFP reporter. In combination with other reporters, and labeled CSF1, the Csfr<sup>-</sup>-mApple mouse provides a new tool to dissect the differentiation and function of the heterogeneous populations of mouse tissue mononuclear phagocytes and the homeostatic roles of CSF1. How different mononuclear phagocytes regulate CSF1R activity remains an important question given the continued interest in macrophages as possible vehicles for delivery of gene therapies and as targets of therapeutics.

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