Cardiovascular Health: The Role of Endothelial Calcineurin Signaling

Highlights

- Endothelial calcineurin is dispensable for physiological and tumor angiogenesis.
- Endothelial calcineurin stabilizes vessels and restrains metastatic outgrowth.
- Calcineurin/NFAT target BMP2 induces differentiation of metastatic cells.

Authors

Stefanie Hendrikx, Sanja Coso, Borja Prat-Luri, ..., Holger Gerhardt, Mauro Delorenzi, Tatiana V. Petrova

Correspondence
tatiana.petrova@unil.ch

In Brief

Hendrikx et al. show that endothelial calcineurin signaling is dispensable for physiological and tumor angiogenesis. Instead, it promotes vascular stabilization and, in cancer, restrains metastatic outgrowth. Immunosuppressive therapy with calcineurin inhibitors thus also directly affects the endothelium, which may contribute to aggressive cancer progression in organ transplant recipients.
Endothelial Calcineurin Signaling Restrains Metastatic Outgrowth by Regulating Bmp2

Stefanie Hendriks, Sanja Coso, Borja Prat-Luri, Laureline Wetterwald, Amélie Sabine, Claudio A. Franco, Sina Nassiri, Nadine Zangger, Holger Gerhardt, Mauro Delorenzi, and Tatiana V. Petrova

INTRODUCTION

The formation of new blood vessels, or angiogenesis, is a hallmark of many human cancers that supports tumor growth and metastasis (De Palma et al., 2017). Signaling via vascular endothelial growth factors (VEGFs) and their endothelial receptor tyrosine kinases (VEGFRs) are essential for vessel growth and remodeling. Blocking VEGF-A/VEGFR-2 interaction or downstream signaling pathways such as phosphoinositide 3-kinase (PI3K), the mammalian target of rapamycin (mTOR), and Akt interfere with angiogenesis and vascular maintenance in many pathological and physiological situations (De Palma et al., 2017; Graupera et al., 2008; Kerr et al., 2016). VEGFR-2 activation also triggers the Ca²⁺/phospholipase-C gamma cascade, which activates the Ca²⁺-dependent serine/threonine phosphatase calcineurin. Calcineurin dephosphorylates the nuclear factor of activated T cells (NFAT), leading to its nuclear entry and transcriptional activity. Among the 5 members of the NFAT family, NFAT1–NFAT4 are regulated by calcineurin. They were first described in T cells; however, it is now evident that they are expressed by numerous cell types and have a wide range of cellular functions (Mancini and Toker, 2009).

In vitro studies of the calcineurin/NFAT pathway suggested that it is a key downstream mediator of VEGFA-VEGFR2 responses, such as endothelial cell (EC) migration and proliferation (Minami et al., 2004; Schweighofer et al., 2009; Zaichuk et al., 2004). However, the in vivo role of the calcineurin/NFAT signaling pathway in the tumor vasculature is not fully understood. Studies of Down syndrome critical region-1 (DSCR1), a direct target and endogenous inhibitor of calcineurin/NFAT, showed that Dscr1 overexpression suppresses calcineurin signaling, inhibits tumor angiogenesis, and prevents tumor growth (Baek et al., 2009). Dscr1 germine deletion reduced tumor growth as a result of overactive calcineurin signaling, leading to EC apoptosis (Ryeom et al., 2008). In addition, Dscr1−/− mice develop more lung metastases due to the increased production of angiopoietin-2, which promotes vascular permeability and facilitates tumor cell extravasation (Minami et al., 2013).

Here, we directly investigated the role of calcineurin signaling in tumor angiogenesis, growth, and metastasis, using a model with...
Figure 1. Endothelial Calcineurin Signaling Controls Vessel Regression in the Postnatal Retina

(A) Scheme for tamoxifen injection and retina collection of pups.
(B) Representative images of a retina leaflet from Cnb1 WT and Cnb1 Δc nec pups. Staining for collagen IV (green), Icam2 (red), and IB4 (blue).
(C) Vascular radial expansion is impaired in Cnb1 Δc nec retinas. ***p = 0.0001; n = 6 Cnb1 WT; n = 6 Cnb1 Δc nec.
(D) Vessel density measured by IB4 is reduced in Cnb1 Δc nec retinas. *p = 0.0347; n = 6 Cnb1 WT; n = 6 Cnb1 Δc nec.
(E) Representative images of empty sleeves from Cnb1 WT and Cnb1 Δc nec retinas. Staining for Icam2 (green) and collagen IV (red). Arrowheads indicate empty sleeves.
(F) The number of empty sleeves is increased in Cnb1 Δc nec retinas. Quantification of the number of Icam2−/ collagen IV+ area normalized to the total vascularized area. *p = 0.0273; n = 6 Cnb1 WT; n = 6 Cnb1 Δc nec.
(G) Increased association of pericytes with empty collagen sleeves in Cnb1 Δc nec retina. Immunofluorescent staining for pericyte marker CD13 (gray), CD31 (green), and collagen IV (red). Arrowheads indicate a CD13− empty sleeve in Cnb1 WT and a CD13+ empty sleeve in Cnb1 Δc nec.
(H) Quantification of total pericyte coverage in Cnb1 WT and Cnb1 Δc nec pups. n = 3 Cnb1 WT; n = 3 Cnb1 Δc nec.

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Endothelial calcineurin signaling does not contribute to sprouting angiogenesis, but rather is implicated in vessel stabilization.

**Endothelial Calcineurin Signaling Is Dispensable for Tumor Angiogenesis and Tumor Growth**

To study the role of calcineurin in pathological angiogenesis, we injected syngeneic melanoma B16F10 cells subcutaneously in Cnb1\(^{WT}\) and Cnb1\(^{Abec}\) mice (Figure 2A). The growth of primary B16F10 tumors was minimally decreased in Cnb1\(^{Abec}\) animals, and we observed similar results in the Lewis lung carcinoma model (Figure 2B and data not shown). The quantification of tumor volume over time and tumor weight after sacrifice showed a non-significant tendency toward reduced primary tumor size in Cnb1\(^{Abec}\) mice (Figures 2C and 2D). EGFP was uniformly high in tumor ECs, indicating high activity of the Pdgfb-iCreERT2-Ires-EGFP transgene (Claxton et al., 2008), and analysis of sorted tumor ECs confirmed the significantly reduced expression of both Cnb1 and the calcineurin/NFAT target gene Dscr1 (Figures S2A–S2C). Further analysis of the tumor vasculature did not reveal differences in vascular density or vessel pericycle coverage between Cnb1\(^{WT}\) and Cnb1\(^{Abec}\) tumors (Figures 2E–2H), indicating that calcineurin is dispensable for tumor vessel expansion and maturation.

The loss of calcineurin increased the number of empty sleeves in the retinal vasculature (Figure 1F); therefore, we analyzed collagen IV distribution in B16F10 tumors. We observed no change in the overall collagen IV staining (Figures 2I and 2J). However, when analyzing the avascular collagen IV, we noticed an increase in the collagen IV+ empty sleeves in Cnb1\(^{Abec}\) tumors (Figures 2I and 2K). Our results indicate that similar to the retinal vasculature, calcineurin is dispensable for the tumor angiogenic response but fine-tunes the tumor vessel stability, and its inactivation has only a marginal impact on primary tumor growth.

**Calcineurin Deletion in the Endothelium Increases Cancer Cell Metastasis in the Lung**

To study the role of endothelial calcineurin in a metastatic context, we injected intravenously (i.v.) B16F10 cells (Figure 3A). We analyzed whether the early steps of metastasis are affected, and 5 days after the i.v. injection, there was no detectable difference in the number of lung metastases between Cnb1\(^{WT}\) and Cnb1\(^{Abec}\) mice (Figures 3B and 3C). However, we observed significantly more metastases in Cnb1\(^{Abec}\) lungs compared to wild-type lungs 12 days after cancer cell injection (Figures 3D and 3E). The increased metastasis in Cnb1\(^{Abec}\) lungs was also apparent at 19 days post-injection (Figures 3F and 3G), but the difference between the 2 genotypes was less prominent. Similarly, i.v.-injected MC38-GFP colorectal cancer cells generated significantly more metastases in Cnb1\(^{Abec}\) mice (Figures S3A and S3B). Our results thus indicate that endothelial calcineurin signaling restrains the growth of lung metastasis.

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1. CD13 mean fluorescence intensity (MFI) is similar between Cnb1\(^{WT}\) and Cnb1\(^{Abec}\) pups. \(n = 3\) Cnb1\(^{WT}\); \(n = 3\) Cnb1\(^{Abec}\).
2. \(\text{CD13}^{\text{Abec}}\) retinas have an increased number of CD13+ pericytes in the empty sleeves area. \(p = 0.0346; \ n = 3\) Cnb1\(^{WT}\); \(n = 3\) Cnb1\(^{Abec}\).
3. ns, not significant; error bars represent means ± SDs. Two-tailed Student’s t test was performed on all of the data represented. Scale bars: (B) 200 \(\mu m\); (G and J) 50 \(\mu m\).
Figure 2. Endothelial Calcineurin Signaling Does Not Affect Primary Tumor Growth and Angiogenesis

(A) Scheme for tamoxifen and subcutaneous B16F10-luc cell injections.

(B) Representative pictures of tumors ex vivo from Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> mice.

(C) Tumor growth is not affected in Cnb1<sup>ΔΔc</sup> mice; n = 4 per genotype.

(D) Tumor weight is not affected in Cnb1<sup>ΔΔc</sup> mice; n = 4 per genotype.

(E) Comparable primary tumor blood vascular density in Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> mice. Staining for Vegfr2 (red) and DNA (blue) of primary tumors.

(F) Quantification of the blood vessel density in Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> tumors. Vegfr2 area was normalized to the total DAPI area and plotted relative to wild-type (WT); n = 5 per genotype.

(G) Representative images of pericytes in Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> B16F10 primary tumors. Staining for NG2 (red), CD31 (green), and DNA (blue).

(H) Pericyte (NG2) area normalized to the CD31 area is comparable in Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> tumors; n = 4 per genotype.

(I) More avascular collagen IV is present in Cnb1<sup>WT</sup> tumors compared to Cnb1<sup>ΔΔc</sup> tumors. Immunofluorescent staining for collagen IV (red) and CD31 (green) in Cnb1<sup>WT</sup> and Cnb1<sup>ΔΔc</sup> tumors. Arrowheads indicate empty sleeves.

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Immune cells are major players in metastasis initiation and progression. Natural killer (NK) cells and CD8+ T cells can eradicate metastatic cancer cells, whereas macrophages and neutrophils enhance metastasis formation (Kitamura et al., 2015). Therefore, we analyzed major immune cell populations in Cnb1WT and Cnb1bec mice after i.v. injection of B16F10 cells. We did not observe significant changes in B cells, neutrophils, monocytes and macrophages, or NK cells between Cnb1WT and Cnb1bec mice. Although there was some increase in naive T cell infiltration in Cnb1bec animals, this did not translate into significant changes in activated and memory T cells (Figures S4A–S4G). Thus, the increased metastasis phenotype observed in Cnb1bec mice is likely not initiated by immune cells.

Calcineurin Deletion in the Endothelium Increases the Outgrowth Potential of Cancer Cells in Lungs

To evaluate whether calcineurin affects the survival of cancer cells in the circulation or their extravasation, we analyzed the fate of fluorescently labeled B16F10 cells in the lungs 24 h after i.v. injection. A similar number of cancer cells were present in the lungs of wild-type and Cnb1bec mice (Figures S4H and S4I). Thus, endothelial calcineurin/NFAT signaling does not play a major role in early metastatic dissemination.

To study whether this pathway affects the outgrowth of cancer cells in the lung microenvironment, we injected Cnb1WT and Cnb1bec mice i.v. with B16F10 cells first and deleted calcineurin 5 days later (Figure 4A), when micrometastases were already established. Mice were sacrificed 12 days after tumor cell injection, and thus 7 days after calcineurin inactivation. As expected, there was no difference in the total number of metastatic foci between Cnb1WT and Cnb1bec mice. However, we observed a significantly increased number of large colonies (>0.5 mm) in Cnb1bec lungs compared to Cnb1WT lungs (Figures 4B–4D), indicating that endothelial calcineurin restrains metastatic outgrowth.

The increased difference in metastatic foci was observed at day 12 (Figures 3D and 3E), but it was reduced at a later stage (Figures 3F and 3G). These results suggest that calcineurin is important during the initial metastatic outgrowth, but contributes less after metastatic lesions have reached a certain size. To study this question, we induced calcineurin deletion 9 days after the i.v. injection of cancer cells (Figure 4E), and the metastases were quantified at day 16. In this setting, we did not observe changes in the number or size of the metastatic foci between Cnb1WT and Cnb1bec mice (Figures 4F–4H). We thus conclude that the endothelium of Cnb1bec mice promotes the outgrowth of cancer cells during a restricted period.

The increased outgrowth at early stages could be due to the fact that in macrometastases, proportionally fewer tumor cells are located in the vicinity of blood vessels (Figure S5A). The growth of cancer cells in macrometastases thus may be less sensitive to the endothelial-derived paracrine factors. We evaluated whether calcineurin loss affects the proliferation of cancer cells in macrometastases by analyzing EDU incorporation. We did not observe any differences in the proliferation of cancer cells in macroscopic lesions of Cnb1WT or Cnb1bec lungs when analyzing lungs 19 days post-injection of cancer cells (Figures S5B and S5C). However, in the 12-day outgrowth experiments, we observed an increase in the EdU+ metastatic area in Cnb1bec lungs compared to wild-type lungs (Figures 5A and 5B). Further analysis using GFP-labeled B16F10 cells confirmed the increased proliferation of cancer cells 12 days post-injection in Cnb1bec lungs (Figures 5C and 5D). We also evaluated whether endothelial NFAT signaling is influenced by the presence of metastasis and whether such activation is organ specific. The expression of the direct NFAT target gene Dscr1 was not affected in the lung endothelium by the presence of cancer cells (Figure S5D). However, we observed that the basal level of NFAT signaling in the lung ECs was significantly higher in comparison to ECs from the gut and skin (Figure S5E). Our data thus reveal tissue-specific differences in the levels of endothelial calcineurin activation and show that its loss promotes the proliferation of lung micrometastases.

Calcineurin Signaling Regulates Bmp2 Expression in Lung Endothelial Cells

To mechanistically unravel the contribution of endothelial calcineurin signaling to cancer cell outgrowth, we transduced human umbilical vein ECs (HUVECs) with lentiviruses encoding a constitutively active form of NFATc1 (caNFATc1). The mutation of serines in the SRR and SPXX repeat motifs of the NFATc1 regulatory domain prevents the phosphorylation of NFATc1 and imposes its nuclear localization (Monticelli and Rao, 2002). As expected, HUVECs transduced with caNFATc1 had increased nuclear NFATc1 levels compared to GFP controls (Figures S6A and S6B). Similarly, a lower-motility NFATc1 band was observed in lysates from caNFATc1-HUVECs, confirming the loss of phosphorylation (Figure S6C). There was a 10-fold upregulation of the NFAT target Dscr1 in caNFATc1-HUVECs, thereby confirming the overactivation of NFAT signaling (Figure 6A). To identify potential paracrine factors that affect metastatic outgrowth, we analyzed the expression of cell surface or secreted molecules that were previously shown to be regulated in HUVECs by calcineurin/NFAT signaling (Suehiro et al., 2014; Figure S6D). We observed a significantly increased expression of Tissue Factor and BMP2, but not of other target genes such as VCAM1, ANG2, or ESELECTIN or another member of the BMP family, BMP4 (Figure 6A).

To study whether the target genes identified by our in vitro analyses are also regulated by calcineurin signaling in vivo, we sorted lung blood ECs from wild-type and Cnb1bec mice i.v. injected with B16F10 cells (Figures S6E–S6H). As expected, we observed a strong decrease in Cnb1 in ECs from Cnb1bec mice.

(J) Overall collagen IV staining in Cnb1WT and Cnb1bec tumors is unchanged; n = 5 Cnb1WT; n = 4 Cnb1bec; (K) Avascular collagen IV is increased in Cnb1bec tumors compared to WT tumors. Quantification of the avascular (CD31−) collagen IV area or empty sleeves, plotted relative to WT. *p = 0.0399; n = 5 Cnb1WT; n = 4 Cnb1bec; ns, not significant; error bars represent means ± SDs. Two-tailed Student’s t test was performed on all of the data represented. Scale bars: (B) 1 mm, (E) 50 μm, (G and I) 100 μm.
Figure 3. Endothelial Calcineurin Signaling Restrains Lung Metastasis

(A) Scheme for tamoxifen and B16F10 cell injections.
(B) Representative images of lungs 5 days after B16F10 i.v. injection.
(C) The number of B16F10 colonies in lungs is not affected in Cnb1^Δbec^ mice 5 days after i.v. injection; n = 7 per genotype. Data combined from 2 independent experiments.
(D) Representative images of lungs 12 days after B16F10 i.v. injection.
(E) Cnb1^Δbec^ lungs have more B16F10 colonies 12 days after tumor cell injection. **p = 0.0066; n = 9 per genotype. Data combined from 2 independent experiments.
(F) Representative images of Cnb1^Δbec^ and Cnb1^WT^ lungs 19 days after B16F10 i.v. injection.
(G) Cnb1^Δbec^ lungs have more B16F10 colonies 19 days after tumor cell injection. *p = 0.0212; n = 3 Cnb1^WT^; n = 5 Cnb1^Δbec^.
ns, not significant; error bars represent means ± SDs. Two-tailed Student’s t test was performed on all of the data represented.
Scale bars: (B, D, and F) 1 mm.
Figure 4. Endothelial Calcineurin Deletion Increases the Outgrowth of Metastatic Cancer Cells
(A) Scheme for tamoxifen and intravenous (i.v.) B16F10 injection for early outgrowth experiments.
(B) Cnb1<sup>ΔABC</sup> lungs have significantly larger (>0.5 mm) colonies compared to Cnb1<sup>WT</sup> lungs 12 days after i.v. injection and tamoxifen administration at day 5. **p = 0.0037; n = 4 Cnb1<sup>WT</sup>; n = 5 Cnb1<sup>ΔABC</sup>.
(C) Representative pictures of lungs from Cnb1<sup>WT</sup> and Cnb1<sup>ΔABC</sup> mice from the 12-day outgrowth experiment.
(D) The number of lung B16F10 colonies is not affected in Cnb1<sup>ΔABC</sup> mice 12 days after i.v. injection and tamoxifen administration at day 5; n = 4 Cnb1<sup>WT</sup>; n = 5 Cnb1<sup>ΔABC</sup>.
(E) Scheme for tamoxifen and intravenous B16F10 injections for late outgrowth experiments.
(F) Similar size B16F10 colonies are present in Cnb1<sup>WT</sup> and Cnb1<sup>ΔABC</sup> lungs 16 days after i.v. injection and tamoxifen administration at day 9; n = 5 Cnb1<sup>WT</sup>; n = 4 Cnb1<sup>ΔABC</sup>.
(G) Representative pictures of Cnb1<sup>WT</sup> and Cnb1<sup>ΔABC</sup> lungs from the 16-day outgrowth experiment.
(H) The number of lung B16F10 colonies is not affected in Cnb1<sup>ΔABC</sup> mice 16 days after i.v. injection and tamoxifen administration at day 9; n = 5 Cnb1<sup>WT</sup>; n = 4 Cnb1<sup>ΔABC</sup>.
ns, not significant; error bars represent means ± SDs. Two-tailed Student’s t test was performed on all of the data represented.
Scale bars: (C and G) 1 mm.
mice (Figure 6B). Calcineurin inactivation significantly decreased the endothelial expression of Bmp2, while the expression of other previously published target genes such as TissueF, Angpt2, or Thbs1 was not modified (Figure 6B).

**Bmp2 Inhibits Tumor Cell Growth in 3D**

Lung BMP signaling maintains cancer cells in a dormant state by preventing their self-renewal (Gao et al., 2012). Based on this finding and our own observations, we selected BMP2 as a potentially important calcineurin target during tumor progression. BMP2 treatment strongly activated Smad-1/5 phosphorylation in B16F10 cells (Figure 6C), which was inhibited by dorsomorphin homolog 1 (DMH1), a small molecule antagonist targeting the activation of BMP type I receptors (Hao et al., 2010). However, BMP2 did not affect melanoma cell growth in a 2-dimensional (2D) monolayer (Figure 6D). As metastases grow in a 3D environment, we studied whether BMP2 affects the ability of B16F10-luc cells, tagged with firefly luciferase to form spheroids in Matrigel. In 3D conditions, the addition of BMP2 significantly decreased luciferase signal and spheroid growth (Figures 6E–6G). BMP2 treatment significantly reduced the number of large 3D colonies, whereas the number of small colonies was increased, indicating that BMP signaling inhibits spheroid outgrowth (Figure 6G). The inhibition of spheroid
Figure 6. Calcineurin Regulates Bmp2 Expression in the Endothelium, a Potent Inhibitor of Tumor Cell Growth
(A) caNFATc1 induces the expression of DSRC1, BMP2, and TISSUEF in HUVECs. The qRT-PCR analyses of the indicated genes; n = 3, *p = 0.0091 DSRC1, "p = 0.0257 BMP2, and "p = 0.0318 TISSUEF.

(B) Loss of calcineurin reduces the expression of Cnb1, Dscr1, and Bmp2 in the BEC-2 population of lung ECs. *p = 0.0316, "p = 0.0029; n = 3 Cnb1 WT-GFP; n=4 Cnb1 D

(C) BMP2 induces the phosphorylation of Smad-1/5 in B16F10 cells. Cells were treated with 100 ng/mL BMP2 in the presence or absence of 1 or 3 μM DMH1.

(D) BMP2 does not affect B16F10 cell growth in 2D. Cells were treated with vehicle or 100 ng/mL BMP2 for 24 and 48 h and analyzed by CyQuant DNA-based assay.

(E) BMP2 reduces B16F10 cell growth in 3D. Representative images of B16F10 spheroids treated with 100 ng/mL BMP2, BMP2 + 3 μM DMH1, or vehicle.

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expansion by BMP2 was rescued by the combined treatment with DMH1 (Figures 6E, 6H, and 6I). Similar to B16F10 cells, BMP2 treatment activated Smad-1/5 signaling and reduced the 3D growth of MC38 cells, which was rescued by DMH1 (Figures S7A–S7C).

**BMP2 Signaling Promotes Melanoma Cell Differentiation**

To further evaluate the pathways affected by BMP2 in melanoma cells, we compared the transcriptomes of control and BMP2-treated B16F10 3D spheroids. The expression of 54 and 72 genes was significantly up- or downregulated in response to BMP2 (false discovery rate [FDR] <0.05, fold change [FC] >2). As expected, BMP2 target genes ID2, SMAD6, and ATOH8 were significantly induced (Figure 7A; Koch et al., 2016). BMP2 strongly repressed the nuclear factor κB (NF-κB) pathway, previously implicated in melanoma cell survival, invasion, and metastasis (Figure 7B; Amiri and Richmond, 2005). In agreement with reduced spheroid growth, BMP2 downregulated the expression of genes involved in mitosis, targets of CMYC and E2F transcription factors, and genes associated with the mTOR complex 1 pathway (Figure 7B). BMP2 treatment also repressed gene signatures associated with hypoxia and glycolysis and oxidative phosphorylation, indicating overall reduced metabolic activity (Figure 7B). We observed significantly increased expression of genes important for melanogenesis, such as Serpin F1, TYR, GPR114, Mlh, SLC45a2, and MITF (Figure 7A; Chen et al., 2016; Fernández-Barral et al., 2014; Kondo and Hearing, 2011). Analysis of Gene Ontology (GO) terms for biological processes revealed that 5 of 9 GO terms related to pigmentation, lung metastases had reduced tyrosinase activity (Figures 7E and 7F). Our data indicate that BMP2 signaling inhibits the 3D growth of cancer cells by promoting differentiation along melanocytic lineage and reducing metabolic activity and cell-cycle progression.

**Endothelial BMP2 Regulates Retinal Angiogenesis and Metastatic Colonization**

To study the role of endothelial-derived BMP2 in vivo we generated Bmp2flox/flox; Pdgfb-iCreERT2 mice (Bmp2−/−) and analyzed retinal angiogenesis and metastatic colonization. In postnatal retina, vascular density but not radial expansion was significantly reduced upon endothelial inactivation of BMP2, and there was a tendency toward an increased number of collagen IV⁺ empty sleeves (Figures 7G–7J). Most important, in lung colonization experiments, we observed that the loss of endothelial BMP2 enhanced the formation of metastases (Figures 7K and 7L). The effect of endothelial Bmp2 loss on metastatic outgrowth was less pronounced compared to calcineurin inactivation (Figures 3D and 3E). These in vivo results indicate that BMP2 is an important effector molecule acting downstream of calcineurin in ECs, although additional calcineurin targets are also likely involved.

**DISCUSSION**

In this work, we used an EC-specific calcineurin deletion model to investigate the role of this signaling pathway in physiological and pathological angiogenesis and tumor metastasis. We report that angiogenic sprouting and primary tumor growth are not significantly affected by the absence of calcineurin in ECs. However, both in physiological and pathological situations, calcineurin modulates vascular regression. Most important, our study shows that the ablation of calcineurin in the endothelium promotes metastatic outgrowth.

Earlier work suggested that calcineurin has an essential role in sprouting angiogenesis in vitro and in vivo (Hernández et al., 2001; Mena et al., 2014). However, during embryogenesis, calcineurin is crucial for the formation of heart valves and coronary vessels and for the maturation of lymphatic vessels, but it is dispensable for the growth of other vascular beds (Bushid et al., 2003; de la Pompa et al., 1998; Normén et al., 2009; Sabine et al., 2012; Zeini et al., 2009). Here, in a model of physiological retinal angiogenesis, we show that calcineurin is dispensable for EC proliferation and sprouting. In contrast, vessel regression was affected, indicating that endothelial calcineurin signaling is involved in vessel stabilization and remodeling. In addition to the VEGF-A/VEGFR2 pathway, calcineurin signaling is involved in vessel stabilization and remodeling.

Error bars represent means ± SDs. Two-tailed Student’s t test was performed on all of the data represented. Scale bars: (E) 200 μm, (F and H) 2 mm.
Figure 7. BMP2 Regulates the Growth and Differentiation of Melanoma Cells and Retinal Angiogenesis

(A) Heatmap of BMP2 targets and melanogenesis-related genes induced by BMP2 in B16F10 cells.

(B) Gene set enrichment analysis (GSEA) waterfall plot for the hallmark gene sets significantly affected by BMP2 in B16F10 3D spheroids; upregulated (pink) and downregulated (blue) pathways with adjusted p < 0.05.

(C) GO biological processes “Pigmentation” and “Response to Bmp” are significantly enriched in BMP2-treated B16F10 spheroids.

(D) Signature of BMP2-treated B16F10 spheroids is enriched in human differentiated melanomas.

(E) Representative pictures of l-DOPA staining in Cnb1WT and Cnb1Dbec lungs 12 days post-i.v. injection.

(F) Tyrosinase activity is reduced in B16F10 metastases in Cnb1Dbec lungs 12 days post-i.v. injection; n = 5 Cnb1WT; n = 5 Cnb1Dbec.

(G) Representative images of a P5 retina leaflet from BMP2 WT or BMP2Dbec mice, staining for CD31 (green).

(H) Reduced vascular density in P5 BMP2Dbec retina; ***p = 0.0008, n = 6 BMP2WT; n = 8 BMP2Dbec.

(I) Radial expansion is similar in BMP2WT and BMP2Dbec retinas; n = 4 BMP2WT; n = 6 BMP2Dbec.

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regulates mural cell behavior. It is not yet clear whether pericytes from remodeling vessels remain in the empty sleeves and undergo apoptosis or migrate onto the surviving vessel (Korn and Augustin, 2015). Our findings indicate that pericytes remain in the empty sleeves and that upon calcineurin deletion, these pericytes would have better survival rates.

Pharmacological calcineurin inhibition decreased tumor angiogenesis and tumor growth in several studies (Courtwright et al., 2009; Medyouf et al., 2007). However, in our model, despite the efficient deletion of Cnb1 and the inhibition of calcineurin/NFAT signaling in the tumor endothelium, vascular density was not significantly affected, whereas vascular stabilization was decreased. The tumor phenotype of Cnb1<sup>Abec</sup> mice is thus also rather similar to that of mice with deficient endothelial non-canonical WNT signaling, in which a defective vessel stabilization and only a minor effect on primary tumor growth have been reported (Scholz et al., 2016). It is conceivable that the pharmacological inhibition of calcineurin blunts tumor growth in cancers, in which calcineurin/NFAT drives malignant cell proliferation, such as in T and B cell lymphomas (Medyouf et al., 2007), or alternatively, when the tumor vasculature is especially sensitive to the loss of non-canonical WNT signaling.

A more intriguing part of our study was that endothelial calcineurin restricts the outgrowth of metastases. This may explain in part the clinical observations that solid organ transplant recipients are both more susceptible to develop de novo cancer and have a more aggressive course of the disease in comparison to the general population (Chapman et al., 2013; Sherston et al., 2014). Studies of Dscr1<sup>−/−</sup> mice with hyperactive NFAT signaling proposed that the increased production of endothelial Ang2 drives cancer cell extravasation by increasing lung vascular permeability (Minami et al., 2013). In contrast, in mice with endothelial calcineurin inactivation, we did not observe significant changes in endothelial Angpt2 expression or metastatic cancer cell behavior up to 12 days after intravenous injection, which argues against the importance of calcineurin during the early stages of metastasis. Rather, we observed an increased outgrowth of metastatic colonies in Cnb1<sup>Abec</sup> mice during a critical time window, after which metastatic growth becomes independent of endothelium-derived paracrine signals.

Mechanistically, we propose that intact calcineurin signaling in lung ECs induces the expression of BMP2, which restrains the outgrowth of micrometastases by promoting cell differentiation. Calcineurin activation induces BMP2 in ECs in vitro and in vivo, and BMP2 treatment potently suppresses the 3D growth of cancer cells. Furthermore, the analyses of melanoma cell transcriptomes revealed that together with reducing metabolic activity and cell-cycle progression, BMP2 directed differentiation along the melanocytic lineage. In addition, metastases in Cnb1<sup>Abec</sup> lungs had lower tyrosinase activity, confirming the reduced differentiation of B16F10 cells when Bmp2 expression downstream of calcineurin is impaired. Most important, direct in vivo inactivation of endothelial Bmp2 increased the metastatic potential of cancer cells, although to a lesser extent than calcineurin deletion. We have not confirmed whether endothelial Bmp2 deletion directly affects the differentiation of B16F10 in vivo. However, comparison of the Bmp2-dependent transcriptional signature of B16F10 cells with human melanomas (Tsui et al., 2017) showed that it correlates best with a melanocytic differentiated subtype. Calcineurin activates all 4 NFAT isoforms (Mancini and Toker, 2009), and previous studies suggested the roles both for NFATc1 and NFATc3 in vascular remodeling and angiogenesis (Bushid et al., 2003; Scholz et al., 2016). The question of which NFAT isoform relays calcineurin activation in the endothelium during metastatic outgrowth thus merits future investigation.

From the clinical point of view, the use of calcineurin inhibitors for immunosuppressive therapy in solid organ transplant recipients is strongly associated with increased de novo malignancies, including melanomas, and especially with a more aggressive course of disease (Chapman et al., 2013; Sherston et al., 2014; Tremblay et al., 2002). Given the important roles of calcineurin/NFAT in immune cells, impaired anti-tumor immunity is conceivably the main culprit leading to the development of cancers in transplant recipients. In addition, our data argue that the loss of endothelial calcineurin/NFAT signaling directly contributes to the progression of at least some cancer types because of the loss of paracrine inhibitory communication between endothelial and cancer cells.

In conclusion, we demonstrate the role of calcineurin in the postnatal vasculature using an inducible endothelial-specific deletion model. In both physiological retinal and pathological tumor vasculatures, calcineurin is dispensable for sprouting angiogenesis and is mostly implicated in vessel stabilization, which does not affect primary tumor growth. EC calcineurin activation restrains the outgrowth of metastases, suggesting that post-transplantation immunosuppressive therapy also directly targets the endothelium, which is crucial for initial metastasis formation, and is independent of immune cells.

**STAR★METHODS**

Detailed methods are provided in the online version of this paper and include the following:

- **KEY RESOURCES TABLE**
- **CONTACT FOR REAGENT AND RESOURCE SHARING**
- **EXPERIMENTAL MODEL AND SUBJECT DETAILS**
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(1) Quantification of empty sleeves in Bmp2<sup>WT</sup> and Bmp2<sup>Abec</sup> retinas. The number of CD31<sup>+</sup>, collagen IV<sup>+</sup> areas was normalized to the total vascularized area; n = 6 Bmp2<sup>WT</sup>; n = 8 Bmp2<sup>Abec</sup>.

(2) Representative images of Bmp2<sup>WT</sup> and Bmp2<sup>Abec</sup> lungs 12 days after B16F10 injection.

(3) Quantification of lung metastases in Bmp2<sup>WT</sup> and Bmp2<sup>Abec</sup> 12 days post-i.v. injection. *p = 0.0113, n = 13 Bmp2<sup>WT</sup>; n = 10 Bmp2<sup>Abec</sup>. Data combined from 3 independent experiments.

Error bars represent means ± SDs. Two-tailed Student’s t test was performed on (F), (H–J), and (L). Scale bars: (E) 100 μm, (G) 500 μm, (K) 1 mm.
We thank J. Schlom and L. Borsig for providing MC38 and MC38-GFP cells, M. Acknowledgments

SUPPLEMENTAL INFORMATION Supplemental Information includes seven figures and two tables and can be found with this article online at https://doi.org/10.1016/j.celrep.2019.01.016.

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AUTHOR CONTRIBUTIONS

S.H., S.C., and T.V.P. designed the study. S.H. performed the experiments, analyzed and interpreted the data, and wrote the manuscript. S.C., B.P.-L., and L.W. performed the experiments and analyzed and interpreted the data. A.S., C.F., and H.G. performed the experiments and analyzed and interpreted the data. N.Z., and L.W. performed the experiments and analyzed and interpreted the data, and wrote the manuscript. All of the authors revised the manuscript critically.

DECLARATION OF INTERESTS

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# STAR METHODS

## KEY RESOURCES TABLE

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Further information and requests for resources and reagents should be directed to and will be fulfilled by the Lead Contact, Tatiana V. Petrova (Tatiana.petrova@unil.ch).

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Animal models

Experiments were approved by the Animal Ethics Committee of the Canton of Vaud. Ppp3r1fl/fl, BMP2fl/fl and Pdgfb-iCreERT2 mice (Claxton et al., 2008; Ma and Martin, 2005; Zeng et al., 2001) were on C57Bl6J background. Adult (5–10 weeks old) female mice were injected i.p. with 50 μg/g mouse of tamoxifen in sunflower seed oil (Sigma). Unless indicated otherwise, tamoxifen was administered on day −2 and 0 of cancer cell injection and twice weekly thereafter. Deletion efficiency was confirmed by qRT-PCR of sorted lung and tumor ECs. Pups were injected at P1 and P3 with 20 μl of 2 mg/ml tamoxifen.
**Cell culture**

Cells were cultured in a sterile incubator at 37°C and 5% CO₂. HUVECs (pooled donor, Lonza) were cultured in endothelial growth medium with full supplements (EBM-2, Lonza) and were used up to passage 6. MC38, B16F10-luc and 293T were cultured in DMEM complete (10% FBS, 1% P/S), for B16F10-luc cells zeocin 0.2 mg/ml was added to the culture medium.

**METHOD DETAILS**

**Tumor models**

For subcutaneous tumor experiments 5x10⁵ B16F10-luc cells (B16F10-luc-G5, Perkin Elmer) in 100 μl PBS were injected in the right flank of the mice. When tumors reached 1 cm³ mice were sacrificed and perfused intracardially with PBS and 2% PFA. 1 hour prior to sacrifice mice were injected i.p. with 5 μg/g EdU (Life Technologies). Tumors were fixed in 4% PFA for 4 hours at 4°C, incubated for 12 hours in 30% sucrose and embedded in OCT. For the lung colonization experiments 5x10⁵ B16F10-luciferase cells in 200 μl PBS were injected into tail vein, mice were sacrificed after 1, 5, 12 and 19 days and perfused intracardially with PBS and 2% PFA. Lungs were fixed in 4% PFA for 4 hours, left in 30% sucrose for 12 hours, OCT inflated via the trachea and embedded in OCT. In outgrowth experiments the first dose of tamoxifen was administered 5 or 9 days after cancer cell injection, and every two days thereafter. For the 24h tumor cell extravasation assay B16F10-luc were labeled with 15 μm CMTMR (Thermo Fisher) by incubating for 1h at 37°C in serum free medium.

**Protein extraction and immunoblotting**

Cells were lysed in RIPA buffer (50 mM Tris pH 8, 150 mM NaCl, 1% Triton X-100, 0.5% sodium deoxycholate, and 0.1% sodium dodecyl sulfate) supplemented with phosphatase (PhosSTOP, Roche) and protease (Complete, Roche) inhibitors. Lysates were cleared by centrifugation and supernatants were loaded on 10% SDS-PAGE gels, transferred on nitrocellulose membranes and blocked for 1h at RT in 5% BSA. Membranes were incubated overnight at 4°C with primary antibodies and for 1h at RT with secondary antibodies conjugated to horseradish peroxidase (Dako). Blots were developed using Super Signal West Femto (Thermo Scientific) with a CCD camera (Fusion, Vilber). Details of the antibodies used are provided in the Key Resources Table.

**Plasmids and transfections**

caNFAcTc1 construct Addgene plasmid #11102 was subcloned into the pSD44 lentiviral backbone. pSD44-GFP was used as control. Lentiviral particles were prepared by transfecting HEK293T cells with the pSD44 vector of interest and packaging plasmids pSD11 and pSD16. 48h after transfection supernatants were collected and concentrated. For infection 3.5x10⁵ HUVECs were plated in a 10cm dish and infected the day after in the presence of polybrene (8 μg/ml) with either GFP or caNFATc1 lentiviruses. The medium was changed 24h later and another 24h later cells were selected with 0.5 μg/ml puromycine for 48 hours.

For the generation of the B16F10-GFP cell line B16F10 were seeded in a 6 well plate and infected the day after with GFP lentivirus. 48h after transfection selection was started with 0.6 μg/ml puromycine, which remained in the culture medium. Single clones were selected to choose cells with the brightest GFP signal.

**2D Proliferation analysis**

For analysis of cell growth, cells were seeded in 96 well plates at a density of 1000 cells/well, 100 ng/ml rBMP2 (R&D) or vehicle treatment was added for 24 or 48 hours. Live cell number was determined using CyQuant NF DNA-based assay (Invitrogen).

**Spheroid formation**

500 B16F10 or 300 MC38 cells were plated in 50 μl of 7 mg/ml Matrigel (Corning) and cultured for approximately 10 days, depending on the experiment. 100 ng/ml of mouse rBMP2 (R&D) was added in the presence or absence of 3 μM DMH1 (Tocris) in DMEM complete medium and was changed every other day. At the end of the experiment, the whole well was imaged with a Leica M205FA stereomicroscope, using Leica camera DFC300FXR2 and LAS AF6000 software. For quantification, 3 images per well were taken with a Leica DMI3000 microscope, using a DFC3000G camera and LAS X software. B16F10-luc spheroids were imaged with the Xenogen IVIS Lumina II system after supplementation with 0.3 mg/ml luciferin (Biosynth) and analysis was performed with Living Image software (Perkin Elmer).

**Retina preparation**

Pups were injected at P1 and P3 with 20 μl of 2 mg/ml tamoxifen in sunflower seed oil. Pups were sacrificed at day 5 and eye balls were fixed in 4% PFA for 4 hours at 4°C. Retinas were dissected and blocked with blocking buffer (0.5% BSA, 5% donkey serum, 0.3% Triton X-100, 0.1% NaCl) for 6h at 4°C. Primary antibodies were diluted in blocking buffer and incubated overnight at 4°C. Subsequently, retinas were incubated with fluorophore-conjugated Alexa secondary antibodies (Thermo Fisher) in blocking buffer overnight at 4°C and mounted with mounting medium (Fluoromount-G, Invitrogen). Details of the antibodies used are provided in the Key Resources Table.
Cell staining

Cells were grown on glass coverslips, washed once with cold PBS (with calcium chloride and magnesium chloride), fixed for 10 minutes in 4% PFA in PBS, permeabilized for 10 minutes with 0.1% Triton X-100 in PBS, blocked for 30 minutes with blocking buffer (0.5% BSA, 5% donkey serum, 0.3% Triton X-100, 0.1% NaN₃) and incubated with primary antibodies overnight at 4°C in blocking buffer. The following day, slides were washed 3x with 0.3% Triton-X PBS for 10 minutes and fluorophore-conjugated Alexa secondary antibodies (Thermo Fisher) were added in blocking buffer for 1h at RT. Slides were washed 3x 10 minutes with 0.3% Triton-X PBS and once with PBS before mounting.

Immunofluorescent and histochemical staining

8 μm OCT sections were thawed for 20 minutes at RT, fixed for 10 minutes in 4% PFA and washed with 0.3% Triton-X PBS. After blocking for 30 minutes with blocking buffer (0.5% BSA, 5% donkey serum, 0.3% Triton X-100, 0.1% NaN₃) slides were incubated with primary antibodies overnight at 4°C in blocking buffer. On day 2, slides were washed 3x with 0.3% Triton-X PBS for 10 minutes and fluorophore-conjugated Alexa secondary antibodies (Thermo Fisher) were added in blocking buffer for 1h at RT. Slides were washed 3x 10 minutes with 0.3% Triton-X PBS and once with PBS before mounting.

For the analysis of tyrosinase activity lung cryostat sections were incubated with 5 mM of L-3,4-dihydroxyphenylalanine (L-DOPA, Sigma-Aldrich) in 0.1 nM sodium phosphate buffer at pH 7.4 for 2 h at 37°C; controls were incubated with the buffer alone. The sections were fixed in 4% PFA for 15 min at RT, counterstained with Mayer’s hematoxilin and mounted.

qRT-PCR

RNA was isolated using the QIAGEN RNeasy Plus Mini Kit. Reverse transcription was performed using Transcriptor First Strand cDNA Synthesis Kit (Roche). Alternatively, RNA from FACS samples was isolated using the QIAGEN RNeasy Plus Micro Kit and mRNA was amplified using the Ovation RNA-seq system V2 (Nugen). StepOnePlus (Applied Biosystems) and SYBR Green PCR Master Mix (Kapa Biosystems/Bioline) were used for qRT-PCR analyses. Data was normalized to GAPDH (HUVECs) or B2m (FACS samples) using the comparative Ct (ΔΔCt) method. Sequences of PCR primers are provided in Table S2.

RNA-sequencing

RNA quality was assessed on a Fragment Analyzer (Advanced Analytical Technologies, Inc., Ankeny, IA, USA) and all RNAs had a RQN between 9.4 and 10.

RNA-seq libraries were prepared using 500 ng of total RNA and the Illumina TruSeq Stranded mRNA reagents (Illumina; San Diego, California, USA). Cluster generation was performed with the resulting libraries with the Illumina TruSeq SR Cluster Kit v4 reagents and sequenced on the Illumina HiSeq 2500. Sequencing data were demultiplexed using the bcl2fastq Conversion Software (v. 2.20, Illumina; San Diego, California, USA).

Endothelial cell sorting

For sorting of endothelial cells we used the Pdgfb-iCreERT2GFP (Cnb1WT-GFP) as controls, which express EGFP in the Pdgfb-iCreERT2 construct (Claxton et al., 2008)

For isolation of tumor endothelial cells Cnb1WT-GFP and Cnb1WT-GFP mice were injected i.v. with 5x10⁵ B16F10-luc cells and tamoxifen was administered. 7 days after cancer cell injection mice were sacrificed. Following 30 min incubation with 10% collagenase digestion mix at 37°C, the tumor digest was filtered through 100 μm and 40 μm cell strainers to obtain single cell suspension. Cell were stained with antibodies listed in the Key Resources Table. Endothelial cells were selected as the CD45-GFP+CD31+ population.

Intestinal LECs were isolated from adult small intestine using cell sorting as described previously (Bernier-Latmani et al., 2015). The intestine was dissected and flushed with ice-cold PBS. Peyers patches were removed and the intestine was cut into 1 cm pieces, which were put in a 10 mM EDTA solution with agitation at 37°C for 30 minutes to remove epithelial cells. The remaining tissue was then digested with Collagenase IV (3 mg/ml) in complete DMEM (GIBCO; ThermoFisher Scientific) containing CaCl₂ (2 mM) and 50 μg/ml DNaseI (Roche) in 0.1% BSA. Lungs were cut in pieces and incubated with 0.1% collagenase A (Roche) and 50 μg/ml DNase (Roche) in 0.1% BSA. Lungs were cut in pieces and incubated in digestion mix at 37°C for 45 minutes while continuously rotating at 150 rpm. Digested tissue was filtered through 100 μm and 40 μm cell strainers to obtain single cell suspension. 20 million cells were stained with antibodies listed in Key Resources Table. All FACS sorting was performed on a BD FACS Aria II (BD Bioscience).
Initially, we aimed to isolate lung BECs as Pdpn-CD31+GFP+ and lymphatic endothelial cells (LECs) as Pdpn+CD31+GFP- populations. Unexpectedly, we identified a third major Pdpn+CD31+GFP+ endothelial cell population, which we refer to here as BEC-2 (Figure S6E). qRT-PCR analysis of BEC, BEC-2 and LEC populations demonstrated that all of them expressed the pan-endothelial markers CD31 and VE-cadherin (Figure S6F). BEC and BEC-2 were furthermore enriched in the blood endothelial-specific marker Vegfr1 (Jurisic et al., 2012) (Figure S6G). In contrast lymphatic markers Proxl, Vegfr3, Itga9, Nrp2 and Pdpn were highly expressed in LECs (Figure S6H). BEC-2 population showed no expression of lymphatic markers and more strikingly, although the cells were sorted based on the cell surface Pdpn+ expression, they did not contain Pdpn mRNA (Figure S6H). We thus propose that due to the particular organization of the lung capillary vasculature, the endothelial cells, located in close proximity of the podoplanin-expressing epithelial cells in order to allow gas exchange, acquire membrane of podoplanin positive cells during tissue digestion. The phenomena of cell membrane transfer was described before in subcapsular sinus macrophages (Gray et al., 2012). However, these cells that appear as Pdpn+ BECs (BEC-2) in our FACS analyses, will have not undergone any RNA transfer as they are devoid of Pdpn mRNA.

**Immune cell analysis**

For immune cell analysis we used Cnb1WT and Cnb1Δbec animals injected with 5x10⁵ B16F10-luc cells. 5 or 7 days after cancer cell injection, mice were anesthetized and perfused intracardially with PBS. Lungs were cut in pieces and digested in digestion mix (1mg/ml Collagenase type IV in DMEM) at 37°C for 45 minutes while continuously rotating at 150 rpm. Digested tissue was filtered through a 70 μm cell strainer to obtain single cell suspension. Cells were then stained for flow cytometry with anti-CD45-FITC, anti-CD3ε-e660, anti-CD8α-PerCP-Cy5.5, anti-CD4-A700, anti-CD44-Pe-Cy7, anti-CD62LPe and anti-B220-Texas Red (=lymphoid panel) or with anti-CD45-FITC, anti-CD11b-eFluor780, anti-Gr1-Alexa647, anti-Ly6C-PerCP-Cy5.5, anti-NK1.1-Pee, and anti-B220-Texas Red (=innate immune cell panel). Data were acquired on LSR II (BD Bioscience) and analyzed using FlowJo software (BD Bioscience). The following populations were analyzed: B cells (CD45+ B220+), effector memory (CD62L-CD44+), central memory B cells (CD62L+CD44+), naive (CD62L+CD44-) and CD4+ and CD8+ T cells, monocytes (Cd11b+Ly6C+), natural killer (Cd11b+/NK1.1+) and neutrophils (Cd11b+GR1+).

**QUANTIFICATION AND STATISTICAL ANALYSIS**

**Image analysis**

Retinas were imaged with an upright Zeiss Axio Imager Z1 using the tile scan and stitching function, for general overview pictures and to quantify the vessel branching and radial expansion. For vessel regression pictures were obtained with an inverted Zeiss LSM 880 confocal microscope. Retina analysis was performed with the Fiji software. Vessel regression analyses were performed by counting the collagen IV positive Icam2 negative structures and normalizing the counts to the total Icam2 positive area. Radial expansion was measured as the distance between the vascular front and the central optic nerve. Arterial and venous vessel branching was counted as the number of vessels branching off from the big central artery of vein visible per leaflet. Overall pericyte coverage was determined by measuring the total CD13 stained surface, further collagen IV positive, CD31 negative areas were selected and checked for CD13 positive staining to observe pericytes in empty sleeves. Proliferation of endothelial cells was measured in regions containing the vascular sprouting front by dividing the total number of ERG+ endothelial cell nuclei by the number of positive for ERG/EdU cells in 3–5 20x objective images. Apoptosis was quantified as the number of cleaved caspase-3+ events in CD31+ vessels divided by the total CD31 vessel area, and given as percentage of control.

For whole section analysis slides were imaged with the upright Zeiss Axio Imager Z1 using the tile scan and stitching function or either with the slide scanner Axio Scan.Z1. Confocal images were captured with an inverted Zeiss LSM 880 microscope. Area measurements were carried out using ImageJ software. Tumor vessel density was quantified as the CD31 area normalized to the total DAPI tumoral area. Empty sleeves in primary tumors were defined as avascular, CD31 negative regions, that were positive for collagen IV and 5 images per tumor were analyzed. For proliferation in lung metastasis, EdU was normalized to the GFP positive tumoral area. Empty sleeves in primary tumors were defined as avascular, CD31 negative regions, that were positive for collagen IV and 5 images per tumor were analyzed. For proliferation in lung metastasis, EdU was normalized to the GFP positive tumoral area.

**Statistical analysis**

Pairwise comparisons were performed by two-tailed Student’s t test. The MC38 3D spheroid growth *in vitro* and the mRNA expression of LEC, BEC and BEC-2 for their specific markers was analyzed by two-way ANOVA. Statistical details of experiments can be found in the figure legends.

**Public dataset analysis**

Affymetrix Human Genome U133 Plus 2.0 Array gene expression profiling dataset GSE49426 (Suehiro et al., 2014) from HUVECs treated with 50ng/ml VEGFA in the presence or absence of 1µM Cyclosporin A or HUVECs with adenovirally overexpressed caNATC1 was downloaded and normalized with RMA (affy R package). Differential expression was computed with limma. In the absence of replicates, cutoffs based on log fold change (LFC) were decided to define differentially expressed genes: LFC > 0.5 defined upregulation by VEGFA in the absence of Cyclosporin A, LFC > 0.5 defined upregulation by VEGFA in the presence of...
Cyclosporin A, LFC > 2.5 defined upregulation by caNFATc1. Genes induced by VEGFA without Cyclosporin A and induced by caNFATc1, but not induced by VEGFA in the presence of Cyclosporin A were selected and their expression is shown in a heatmap.

**RNA-seq analysis**

Mouse target transcript sequences were obtained from ENSEMBLE (GRCm38.p6), and the abundances of transcripts were quantified using Kallisto (0.44.0) with default parameters (Bray et al., 2016). Kallisto’s transcript-level estimates were then summarized at the gene-level using tximport (1.8.0) from Bioconductor (Soneson et al., 2015). Raw and processed data are accessible from Gene Expression Omnibus (GEO) under accession number GSE117074.

Differential expression analysis was performed using DESeq2 (1.20.0) from Bioconductor (Love et al., 2014), with FDR < 0.05 and FoldChange > 2 as significance criteria. For gene set enrichment analysis (GSEA), human orthologs of mouse genes were obtained using biomaRt (2.36.1) (Durinck et al., 2005, 2009). GSEA enrichment scores were then computed using fgsea (1.6.0) from Bioconductor (Sergushichev, 2016), with gene sets obtained from MSigDB database (Liberzon et al., 2015; Subramanian et al., 2005), and DESeq2’s LOG2FC as gene ranking metric.

We defined the Bmp2-treated mouse melanoma signature as the aggregated list of up- and downregulated genes obtained from DESeq2. To compare the enrichment of the Bmp2-treated mouse melanoma signature across human melanoma subtypes, processed gene expression data of metastatic skin cutaneous melanoma (SKCM) from TCGA was obtained from the cBioPortal (Cerami et al., 2012; Gao et al., 2013). Expression data was log transformed with offset = 1, and standardized across samples. Sample labels assigning each tumor to one of the four differentiation subtypes (melanocytic, transitory, neural crest-like, and undifferentiated) were directly obtained from (Tsoi et al., 2018). An enrichment score was then computed for the Bmp2-treated mouse melanoma signature per tumor, using a weighted sum of standardized expression values, where the weight was set to +1 for upregulated genes and −1 for downregulated genes.

**DATA AND SOFTWARE AVAILABILITY**

The RNA-seq data are deposited into Gene Expression Omnibus under accession number GEO: GSE111783.