Imatinib Sets Pericyte Mosaic in the Retina

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Abstract: The nervous system demands an adequate oxygen and metabolite exchange, making pericytes (PCs), the only vasoactive cells on the capillaries, essential to neural function. Loss of PCs is a hallmark of multiple diseases, including diabetes, Alzheimer's, amyotrophic lateral sclerosis (ALS) and Parkinson's. Platelet-Derived Growth Factor Receptors (PDGFRs) have been shown to be critical to the PC function and survival. However, how PDGFR-mediated PC activity affects vascular homeostasis is not fully understood. Here, we tested the hypothesis that imatinib, a chemotherapeutic agent and a potent PDGFR inhibitor, alters the PC distribution and thus induces vascular atrophy. We performed a morphometric analysis of the vascular elements in sham control and imatinib-treated NG2-DsRed mice. Vascular morphology and the integrity of the blood-retina barrier (BRB) were evaluated using blood albumin labeling. We found that imatinib decreased the number of PCs and blood vessel (BV) coverage in all retinal vascular layers, this was accompanied by a shrinkage of BV diameters. Surprisingly, the total length of capillaries was not altered, suggesting a preferential effect of imatinib on PCs. Furthermore, the blood-retina barrier disruption was not evident. In conclusion, our data suggest that imatinib could help in treating neurovascular diseases and serve as a model for PC loss, without BRB disruption.

Keywords: pericyte; retina; imatinib; Gleevec, PDGFR, neurovascular

1. Introduction

The Platelet-Derived Growth Factor Receptor-Beta (PDGFR-β) is amongst the major markers expressed by pericytes (PCs) found on the capillaries and responsible for metabolism and oxygen supply. Together with the PDGF-BB ligand, it promotes vascular maturation and stabilization by recruiting PCs and their progenitors [1, 2].

Although their role has been controversial for a long time [3], it is now clear that PCs are able to regulate blood flow in the nervous tissue as part of the neurovascular unit (NVU) (Fig. 1c) [4-8]. They express alpha-smooth muscle actin, making them capable of active vasoconstriction [9], functionally shown by Ivanova et al. 2017 [6] before. They also have a crucial role in many diseases, such as diabetic retinopathy [6, 10, 11]. In PC-deficient mice, hypoxia was reported in the hippocampus, as well as inflammation, and learning and memory impairment [12]. Embryonal ablation of either PDGF-BB or PDGFR-β induces microvascular leakage and hemorrhage and is therefore lethal [13, 14]. Interestingly, however, they are only essential in the formation of new blood vessels (BV) [15, 16], but not in adult blood-retina barrier (BRB) function in the retina [17]. PCs

regulate blood-brain barrier (BBB) in development [17, 18] while in the adult CNS, PC-deficient BVs become dilated without BBB impairment [19, 20].

PCs express PDGFR-β, endothelial cells (EC) secrete PDGF-BB, which is an important relationship to recruit PCs to BVs during development [21]. PDGFR-β is present on PCs, whereas mature ECs express PECAM1 (CD31) and both labels are restricted to the vasculature. In addition, PCs and smooth muscle cells (SMCs) express neural-glial marker-2 (NG2, or Chondroitin sulfate proteoglycan 4 (Fig. 1a). NG2 is a more suitable marker for PC morphological assessment and counting, as the somata of the cells are evenly visible and can be found in the inner retina in three separate layers [6, 7, 22] (Fig. 1a-b, 2).

Imatinib has been potentially used as selective cancer treatment acting through multiple signaling pathways [23-25]. It is a BCR-ABL tyrosine kinase inhibitor and platelet-derived growth factor receptor (PDGFR) and KIT inhibitor, thus induces apoptosis by inhibiting PDGFs but not insulin-induced PI 3-kinase/Akt survival signaling [26]. Imatinib has also been shown to impair the migration of PCs and fibroblasts in vitro [27].

Pharmacologic intervention with PDGFR-β inhibitors, such as imatinib, has shown significant effects in attenuating angiogenesis by reducing PC density. In addition, imatinib has been shown to successfully inhibit retinal neovascularization in oxygen-induced retinopathy in development by reducing the number of vascular cells, including PCs [24, 28]. Contrary, targeting PDGFR by imatinib administration in the adult retina has not been extensively evaluated as a potential model for pathological PC loss. Here we show the decrease of vascular areal densities, possibly as a result of PC depletion following either local or systemic treatment of imatinib in the adult mouse retina.

In most previous studies imatinib was injected intraperitoneally, ignoring the possible adverse effects of systemic treatment [29] (Fig. 1d). Although, such adverse effects directly linking PC depletion to a decrease in BVs density and retinal degeneration have not been proven [30, 31]. Optionally, a local, intravitreal treatment could be used to treat vascular related diseases in the retina as it has been shown to be most effective in many cases in treating diabetic retinopathy, AMD and retinal vein occlusion [32]. In summary, inhibition of PDGFs/PDGFRs might be a promising method for aimed PC elimination without BBB disruption. Our findings also suggest that local imatinib administration could be considered as an efficient treatment modality.
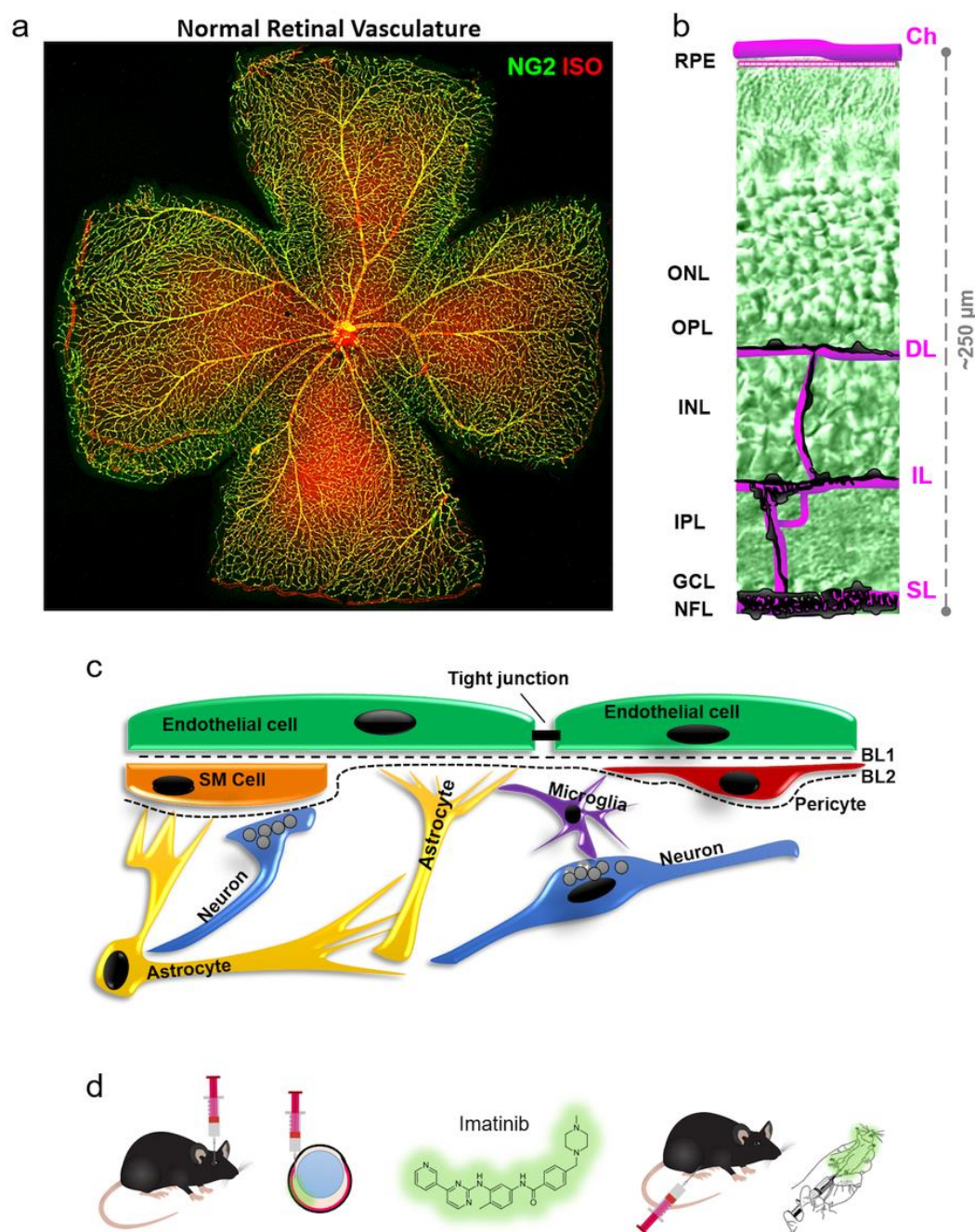
Figure 1. The essential role of PCs in the retina and in the neural tissue. (a) PCs (NG2 – green) and retinal vasculature in the mouse retina (IL, Isolectin GS-IB4 – red). (b) Layers of the retinal vasculature: choroid superficial, intermediate, deep: Ch, SL, IL, DL (c) Location of PCs in the NVU imatinib treatment schemes. (d) Local (intravitreal –IV, left) and global (intraperitoneal – IP, right) treatment with imatinib.

2. Results

The retinal vasculature supplies the retina with an adequate amount of oxygen and metabolites (Fig. 1a, b), thereby proving essential for retinal survival. We utilized two treatment options, systemic, intraperitoneal, injections and local, intravitreal, injections of imatinib to evaluate the effect of PDGF-inhibition (Fig. 1d). According to our hypothesis, imatinib treatment acts on PCs in the adult retina and thus a decrease in PC number accompanied by a decrease of vascular density and

interference in the NVU (Fig. 1c) function is expected that leads to various forms of retinal pathologies.
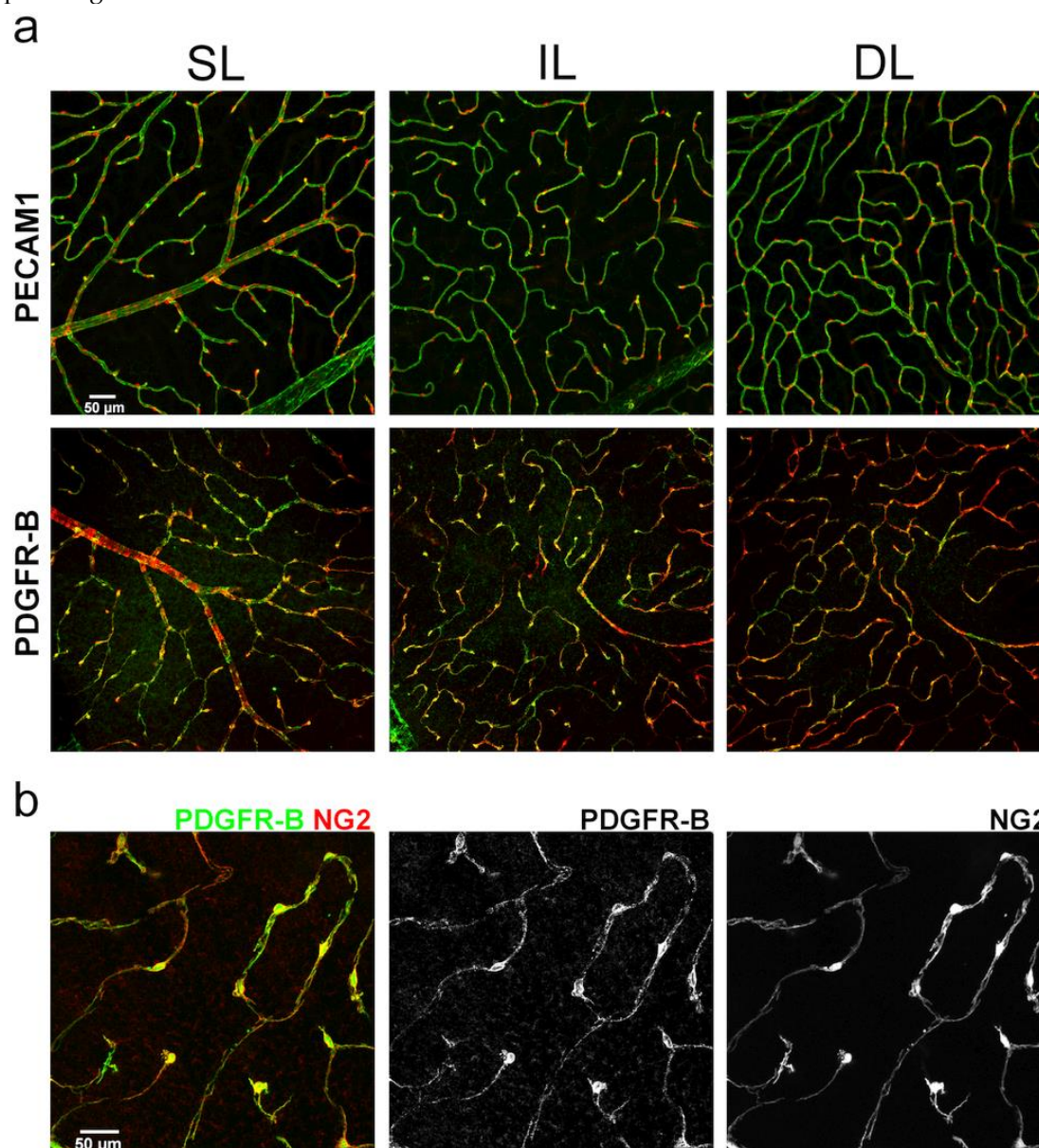
Figure 2. Pattern of endothelial PECAM1 (CD31) and PC PDGFR-β and NG2 in the mouse retina shows strong colocalization of PDGFR-β and NG2 in each layer of the retinal vasculature. NG2 is strictly expressed by mural cells attached to the PECAM1+ vasculature showed by confocal images of NG2-DsRed (red) and PECAM1/PDGFR-β (green) labeling (area 637x637 um) in mouse retinas from each (superficial-, intermediate-, deep-: SL, IL, DL) layer. PECAM1 labels endothelial cells, while PDGFR-β is expressed by NG2+ PCs and smooth muscle cells. It recruits vascularization and stabilizes the mature vasculature. All samples were controls.

2.1. PDGFR-β expression in the mouse retina shows colocalization with NG2

We performed PDGFR-β and PECAM1 immunohistochemical fluorescent labeling (n=3 retinas) in control retinas to show their colocalization with NG2. PCs express PDGFRβ and NG2 in a colocalized manner. PECAM1 is expressed by endothelial cells similar to normal adult retinas. We also tested our sham-control samples for any evident errors caused by PBS injection. We found no vascular aberrations in our sham control samples in PECAM1, PDGFRβ expression (Fig. 2).
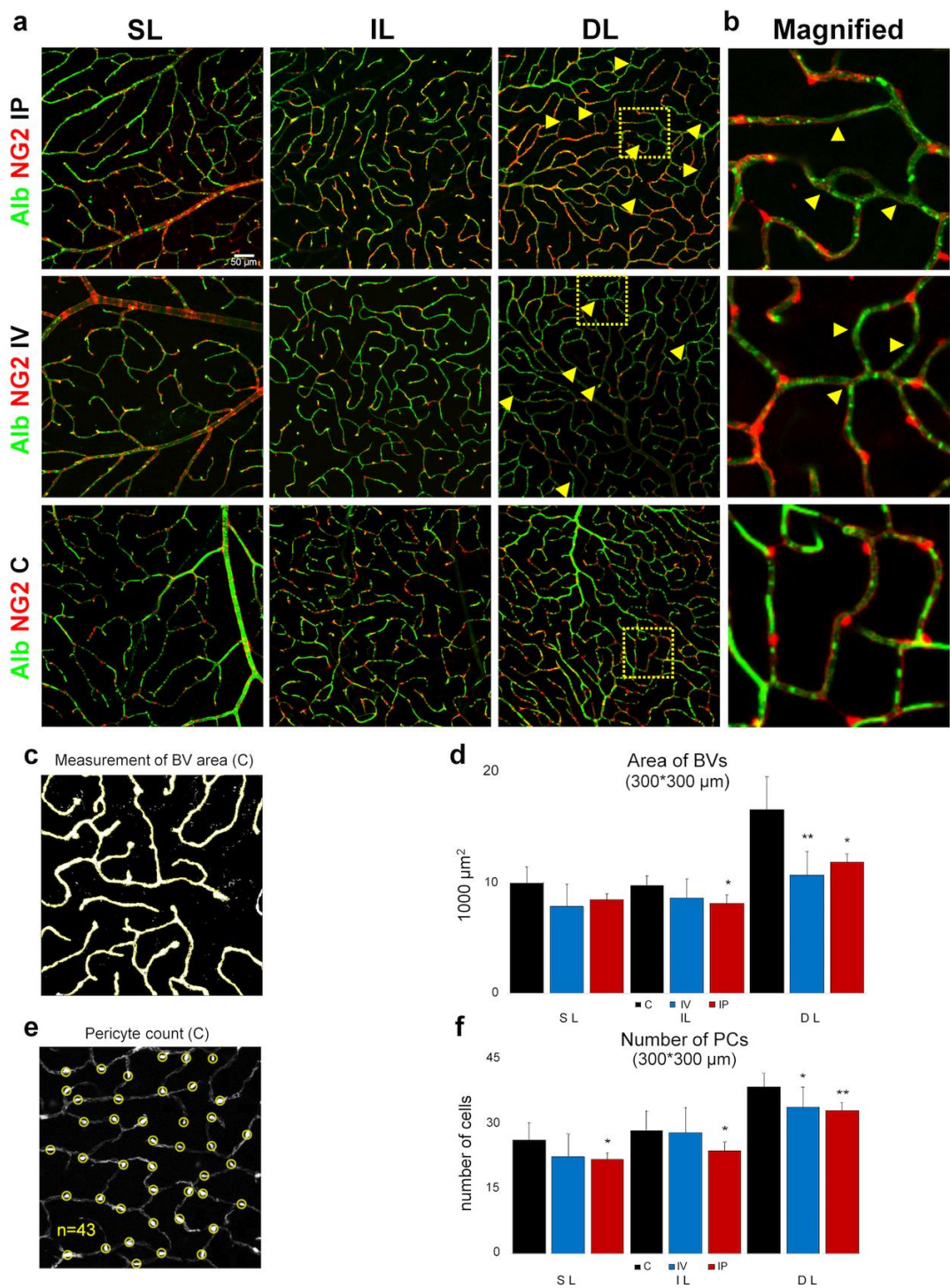
Figure 3. Reduction of PC count and the areal densities of blood vessels following imatinib treatment (a) Confocal images of NG2-DsRed (red) and Albumin (green) labeling (637*637 µm) in mouse retinas from each layer (superficial-, intermediate-, deep-: SL, IL, DL). Measurements were made from these original images, layer-by-layer, in a 300*300 µm area in sham-injected, intravitreally injected (2 µl, 8 µg/µl, IV) and intraperitoneally injected (100mg/kg/d, 2 days imatinib, IP) retinas. PC-loss is indicated with yellow arrows; we could not see PC loss in the sham-injected controls. Areas indicated by yellow squares are magnified and shown on panel b). We found no obvious sign of any extravascular albumin label in our imatinib treated specimen (nor our controls), suggesting that the brain-retina barrier (BRB) remained intact in retinas of these mice. n=13 (C 5, IV 3, IP 5). An example for (c) determining BV areas and (e) PC numbers. (d) BV areas and PC numbers (f) showing

a statistically significant decrease in the IL and DL and suggesting a non-significant decrease in the SL and in the IL of IV animals.

2.2. Imatinib decreases blood vessel coverage

First, we used albumin labeling to evaluate the effect of imatinib on the vascular densities in each retinal layer. It was not evident if imatinib induced any change in the BV coverage when albumin labeled confocal images were compared. To also obtain a quantitative comparison we performed a measurement of areal coverage of BVs in all layers of sham controls, IV and IP treated mouse retinas. Based on this comparison our data suggest that imatinib treatment induced a decrease in BV coverage in all layers for both treatment modalities. This decrement was evident and statistically significant in the deep layer and the intermediate layer of the IP injected animals (tested for n= 4 mice for C, 3 IP, 3 IV; ANOVA; Tukey's post hoc; $p < 0.05$) (Fig. 3c, d).

2.3. Imatinib treatment decreases the number of PCs

PCs are essential for blood-vessel formation [4] therefore we tested if the above-observed decrease of BV areas (Fig. 3d) was accompanied by a corresponding change in the number of PCs (Fig. 3f) following imatinib treatment. Our data suggest that both treatment modalities resulted in a decrease in PC coverage in all three vascular layers, however these changes were not statistically significant in the case of the SL and IL layers with IP treatment. This decrease was pronounced in all three layers of the IP injected mice and in the deep layer in the IV treated animals ($p < 0.5$) (Fig. 3f). This means a 14.7, 17.07% decrease in the SL, in the IV and IP; a 1.83 and 16.59 % decrease in the IL and 12.15 and 14,01% decrease in the IV, IP in the DL of treated animals (Table 1). This thus means that imatinib treatment results in a significant decrease of the PC numbers in all layers in case of IP treatment and a milder effect can be observed in the case of the local treatment in all layers (with a significant change in the DL).

Table 1. Layer-specific PC counts in the experimental specimens

		PC numbers			
Treatment	**Position**	**Mean**	**SD**	**Dev. from control [%]**	**N**
	DL	38.375	3.114		8
C	IL	28.375	4.502		8
	SL	26.125	3.980		8
	DL	33.000	1.732	-14.01	3
IP	IL	23.667	2.082	-16.59	3
	SL	21.667	1.528	-17.07	3
	DL	33.714	4.572	-12.15	7
IV	IL	27.857	5.669	-1.83	7
	SL	22.286	5.251	-14.7	7

*C: Sham-treated control, IP: intraperitoneal- and IV: intravitreal imatinib treated

To test whether there was a significant correlation between the change in PC number and the decrease in capillary surface area, using R (R Core Team 2013) we performed a Spearman's rank correlation test ($p < 0.01$, rho=0.7743385) and also fitted a Generalized Linear Model to our dataset, setting the capillary layer and the treatment type as factors (PC number - Area: p=0.039199). The results suggest a strong correlation between the PC number and capillary area.

2.4. Imatinib treatment does not change the length of capillaries but PC coverage and capillary diameter

Having proved that imatinib treatment caused PC loss (Fig. 3f), the next step was to determine whether the loss of PCs had an effect on BVs. However, the total lengths in all layers remained unchanged in any of the treatment modalities (Fig. 4a, b). Considering the PC-loss in both the IP and IV injected mice it was not surprising that the BV length vs. PC number ratio positively changed after both of the treatments in the IL and DL and in the SL of the IP injected animals. This change was not significant in the SL of the IV injected mouse (Fig. 4c).

However, since BV area coverage decreased, while the capillary length remained constant, we had to test where the decrease in the area comes from. We randomly tested capillary diameters in our dataset to show whether imatinib injection has an effect. In the case of both types of imatinib treatment modalities capillary diameters decreased significantly in the IL and DL measured next to PC somas (Fig. 4d, e), making this decrease in diameter responsible for the decrease in BV area coverage.
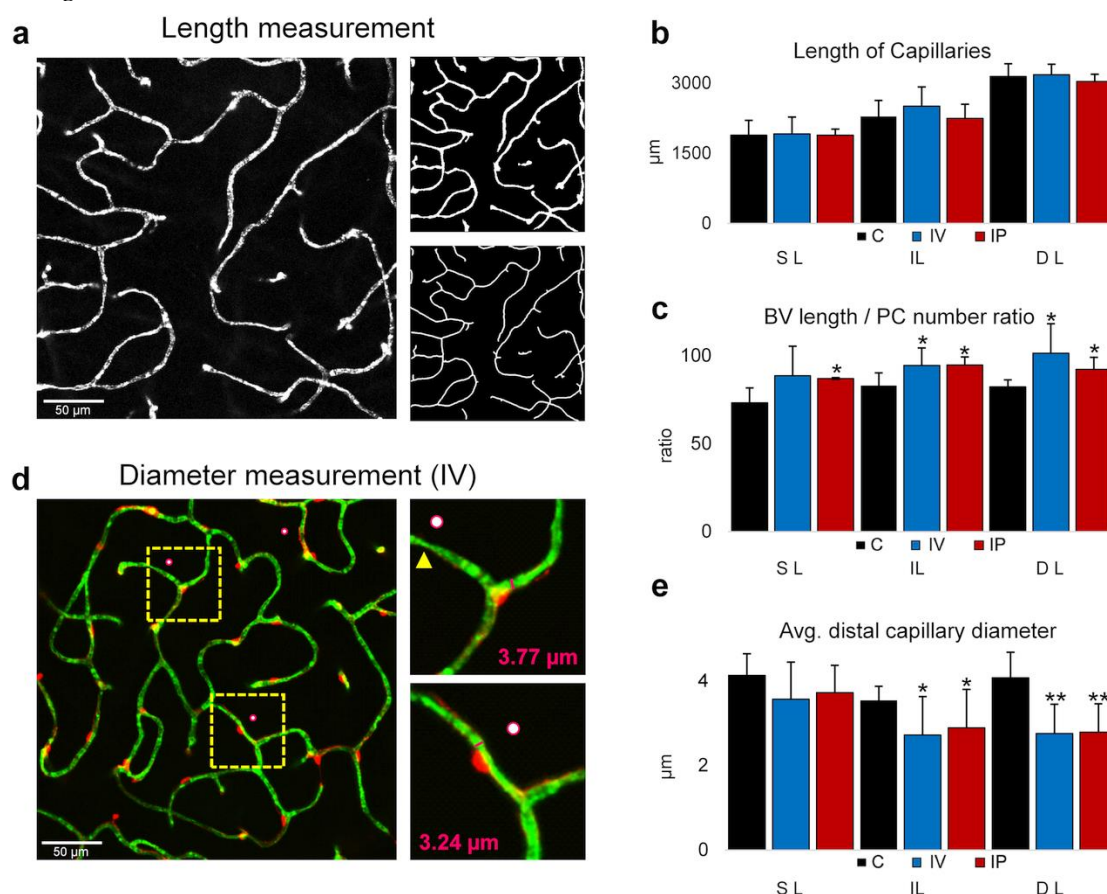
Figure 4. Reduction of PC number is not accompanied by BV length shrinkage following imatinib treatment (a) Illustrated step-by-step method for BV length measurement: normalized grayscale image of albumin IHC-labels merged from n=5 optical slides (top); binary image (right top); the skeleton of BVs used to measure length (right bottom). (b) Comparison of blood vessel lengths in retinal layers in Control (C), Intravitreally injected (2μl, 8μg/μl imatinib, IV) and intraperitoneally injected (100mg/kg/d, 2 days imatinib, IP) retinas. (c) Comparison of BV length / PC number ratios on the same dataset suggests that PCs fill out the gaps after PC loss and protect the underlying vasculature. Measurements on capillary diameters (d) show a reduction in the IL and DL (e). Measurements were made layer-by-layer in a 0.3*0.3 mm area from a mid-central region. C, IV, IP n=4, 4, 3. Error bars are ± SD. * P < 0.05; ** P < 0.01.

2.5. Imatinib did not induce BRB impairment.

In the albumin labeled samples, no albumin leakage into the extracellular space was detected, which would have indicated a BRB impairment (Fig. 3a). This thus confirms that the BRB was intact

in both sham control and imatinib treated animals and further suggests that imatinib treatment caused milder degeneration, without BRB breach in the retinal tissue as generally observed in other degenerative processes [33].

Notably, IP treated mice had an average of 11.15% decrease (p=0.065) in body weight 7 days after the first injection, suggesting a global negative effect of imatinib treatment. Nonetheless, we could not detect any characteristic change in the IV and the Control animal's weight.

3. Discussion

Several conditions and diseases, such as diabetes mellitus, hypertension, age-related macular degeneration (AMD) are associated with negative effects on the development and structure of the retinal vasculature [34, 35]. Hypertension, for example, causes an increased contractile potential of the vessel walls and later PC loss [36]. Hyperglycemia in diabetes mellitus can also lead to microvascular diseases, including diabetic retinopathy [37].

It remains a great challenge to define the key vascular components of diabetic retinopathy, however, our present knowledge and recent discoveries in this field showed that PCs are crucial in BV formation and maintaining its structure [37]. Diabetic retinopathy is characterized by several morphological changes, including loss of PCs, thickening of the basement membrane, increased vascular permeability, vascular occlusion, and microaneurysm. Therefore, it can be concluded, that PC depletion is one of the major hallmarks of Diabetic retinopathy [15]. Although, intraocular anti-VEGF injection and laser treatments can help regress these abnormal new vessels but they both can be detrimental or inadequate options for long term treatment [38]. Imatinib, as a potent PDGFR-, tyrosine kinase- and KIT inhibitor has been successfully used in treating cancer, and thus could provide an alternative treatment strategy in neurovascular diseases. It has also been shown previously that using imatinib as a PDGFR inhibitor during retinal development after OIR (oxygen-induced retinal) neovascularization successfully ameliorated retinal degeneration in the developing retina [24]. In this study, we found that the number of PCs were significantly depleted in all three layers of the retinal BVs in the healthy, adult retina (in a fixed area, not along BVs) after IP treatment. The deep layer was the most affected, out of the three layers, in both utilized treatment options. In addition, the decreased PC number was accompanied by a reduced capillary area and this correlation first suggested a strong association or even a causative link between these imatinib induced effects. However, we found no observable imatinib induced a change in the length of BVs. As PCs line up along capillaries their numbers are expected to follow the length rather than the summated area of the capillary system. Our observations therefore suggested a PC rearrangement. This hypothesis was supported by the imatinib induced increase of the BV area/ PC number ratio in both treatment modalities. This finding suggests that each retained PC covers a lengthened BV section to compensate for the imatinib induced PC loss. While BV areas actively change in live animals it is uncertain if the observed BV area reduction in the post-fixed state represents a corresponding BV area decrease in the live animal. Assuming that the constricted state is preserved in the fixed tissue the detected decrease in BV areas suggests that PCs constrict after imatinib treatment. Besides the above imatinib induced changes layer specific differences in PC loss was also observed (Fig 4; Table 1), which can be explained by the efficacy of local versus systemic treatment modalities. The DL is more affected, while the IL and SL is less affected by IV treatment. This may suggest that the DL is more sensitive to imatinib treatment, despite the greater distance and presumably lover local dose in IV, whereas the highest dose occurs at the level of the nearest vascular layer, the SL. On the other hand, there might be another path for IV fluid through the anterior chamber of the eye to affect the DL of retinal vasculature.

Inhibition of PDGFs/PDGFRs might be a promising therapy to suppress pathological angiogenesis. To test for imatinib induced side effects, we measured the bodyweight of the mice before and after the treatment and labeled the retinas with albumin, to detect any BRB breakdown.

In the retina albumin IHC has been extensively used by multiple groups to label BRB breach [33, 39, 40]. Armulik et al. showed that imatinib abolishes the accumulation of intravenously injected

tracers (Evans Blue - albumin; Cadaverin-Alexa 555, MW=~1 kDa) in the brain of PC-deficient mutants [17].

Our results show no hemorrhage and BRB breakdown after imatinib-treatment either in local- or in systemic treatment tests (Fig. 3), while, in parallel, the same albumin IHC adequately labels patches of extravasal albumin in the RD10 mouse retina (Sup Fig. 1). This evidence suggests that short exposure to imatinib (1*2 µl (8 µg/µl) IV; 2*100 mg/kg/day IP) already has an effect on PC density and vascular area, but not on the length and BRB integrity. This observation is consistent with other studies suggesting that the PDGF-BB/PDGFR-β signaling is not required to maintain EC–PC interaction for BRB integrity during adulthood, although it is indispensable in formation and maturation of BRB through active recruitment of PCs onto the growing retinal vessels and also could be related to the suggested BRB protective effect of imatinib [17] and elimination of PCs will eventually lead to loss of function in capillaries, thus possibly having a major effect on retinal health and resulting in further vascular atrophy.

A weight drop occurred in the IP injected, but not in the IV injected animals after treatment, suggesting an adverse systemic effect of imatinib that can be avoided with local administration. These findings, therefore, suggest that local imatinib administration could be considered as an additional treatment modality. On the other hand, the significant weight-loss occurred after the systemic treatment points towards negative side effects of the systemic imatinib administration.

In human chemotherapy, imatinib dose varies between 100-600 mg daily [41]. The highest clinical dose equals 8-12 mg/kg/day (depending on individual weight), whereas our experimental animals were treated with 100 mg/kg/day for only 2 days. Interestingly, however, this applied much higher dose was not disruptive on the BRB integrity but decreased the PC numbers in all layers. This observation seemingly contradicts a previous study of Armulik et al. [17], where no change was detected in the BV coverage and PC numbers after imatinib treatment. However, in that study, a shorter survival-time (48 h) was applied after the first treatment, whereas we waited 7 days to provide enough time to any possible rearrangement. This difference in the applied methodologies can easily explain the observed discrepancies. Additionally, local imatinib treatment may also have a somewhat different effect on retinal BV formation and maintenance. Our local IV treatment (16 µg imatinib / eye) equals a ~10-fold higher local dose in the eye, compared to systemic (IP) treatment, still having less negative systemic effect, since the dosage if the IV injection projected to the whole body weight is at least ~1500-fold lower if we take the eye/whole body weight ratio into account.

If used for a longer period, imatinib could cause retinal pathologies in chemotherapy [30, 31, 42]. This long term imatinib effect on vasculature may be explained by the loss of the PC dynamic interactions thus the ability to sustain retinal metabolic needs in some conditions [6].

Also, undeniably, imatinib treatment is not specific for PC elimination, since it is not exclusively specific for PDGFR-β, it could serve as a good model for PC dropout, having a role in multiple diseases. To focus its effect and to avoid possible side effects, our results suggest a local treatment on BVs if possible. Since imatinib has been approved for medical use, there are fewer obstacles ahead to benefit from its effect.

Finally, as we have recently shown, PCs form an essential 3D mosaic across the retina to provide a propagating vasomotor response in capillaries [22]. The signal for this propagation travels across gap junctions between neighboring PCs, thereby precisely supporting metabolic needs in the nervous tissue [6, 7, 22]. Any loss in this 3D-PC structure contributes to the loss of this essential function. We theorize that the PC mosaic can adapt to such changes to restore the function by filling out the gaps along the vasculature primarily resulted from the loss of some PCs. The given PC-loss, without a negative change in BV length, suggests that PCs may extend their processes and even move to rebuild the mosaic in order to protect BVs from degeneration. The decrease in capillary areas may come from either a decrease in their length or a decrease in diameter. We found that the diameters indeed shrunk after treatments. Although this shows a controversy with other findings [17], it could be explained by the longer recovery time (7 days). After our treatments the stress that might be still present caused by PC-mosaic reorganization. Although this theory needs to be further tested, it also serves as a direct explanation and a future path for PC research.

4. Materials and Methods

4.1. Animals & imatinib treatment

Animal handling, housing, and experimental procedures were reviewed and approved by the ethical committee of the University of Pécs (BA02/2000-6/2006; BA/35/51-42/2016) and in compliance with protocols approved by the Institutional Animal Care and Use Committee of Weill Cornell Medicine. All animals were treated in accordance with the ARVO Statement for the Use of Animals in Ophthalmic and Vision Research and with the National Institutes of Health Guide for the care and use of laboratory animals. All efforts were made to minimize pain and discomfort during the experiments and all procedures were done by obeying the 3R law.

We used multiple routes of imatinib administration to evaluate the effects of global, and local use of imatinib. Three mouse lines: C57BL/6J mice (The Jackson Laboratory, stock #000664, RRID: IMSR_JAX:000664), NG2-DsRed mice (The Jackson Laboratory, Tg(Cspg4-DsRed.T1)1Akik/J, stock #008241, RRID: IMSR_JAX:008241), RD10 (The Jackson Laboratory, stock #004297, RRID: MGI_3581193, see only in Suppl. Fig 1.) were used in this study. All animals were healthy, adult (1-6 mo.), males. Animals were injected either with imatinib (Methanesulfonate Salt, LC Laboratories, Cat. I-5508) or filtered phosphate-buffered saline (PBS; 0.1 M, pH = 7.3) as follows:

A: Imatinib intraperitoneal injection (IP) 100 mg/kg/day imatinib (dosage according to Raimondi 2014) in filtered PBS for 2 days, on n=5 mice.

B: Imatinib intravitreal injection (IV) 2 µl (8µg/µl) in filtered PBS on n=3 mice.

C: A/B equal volume injected from filtered PBS both IP and IV on n=5 mice; used as a sham control.

4.2. Immunohistochemistry

Eyecups of mice were fixed with 4% carbodiimide and 0.25% PFA in 0.1 M phosphate saline (PBS, pH 7.3) for 15 min at room temperature. After fixation, the samples were washed in PBS, and the retinas were dissected. Retinal whole mounts were blocked for 8-14 h in CTA (PBS solution containing 5% Chemiblocker (membrane-blocking agent, Millipore), 0.5% Triton X-100, and 0.05% sodium azide, Sigma). Primary antibodies were diluted in the same solution and applied for 72 h, followed by incubation for 48 h in the appropriate secondary antibody, conjugated to Alexa-488 (1:1000; green fluorescence, Invitrogen), Alexa-568 (1:1000; red fluorescence, Invitrogen), Alexa-633 (1:500; far-red fluorescence, Invitrogen). In multi labeling experiments, whole mounts were incubated in a mixture of primary antibodies, followed by a mixture of secondary antibodies. All steps were performed at room temperature. After staining, the retinal pieces were flat-mounted on a slide, ganglion cell layer up, and coverslipped using Vectashield mounting medium (H-1000, Vector Laboratories). The coverslip was sealed with nail polish using small pieces of a broken glass coverslip (number 1 size) as spacers. The primary antibodies used in this study were the following: rabbit anti-NG2 coupled to Cy3 fluorescent label (NG2, 1:500, EMD Millipore, AB5320C3), goat anti-mouse albumin (albumin, 1:1000, Bethyl Laboratories, A90-234A), goat anti-mouse PECAM1 (alternatively CD31; 1:5000, R&D Systems, AF3628) goat anti-mouse PDGFR-β (1:5000, R&D Systems, AF1042). BVs were visualized by Isolectin coupled to Alexa-488 fluorescent label (1:500, Invitrogen, I21411) to show the whole retinal vasculature with secondary antibodies (Fig 1). Retinal samples were imaged under a Nikon Eclipse Ti-U confocal microscope. For PC-evaluation retinas were imaged with 20x objective and areas of 300 x 300 µm² were selected for measurements (5 IP animals, 10 areas; 3 IV animals, 6 areas; 5 control animals, 8 areas). The samples were imaged under identical acquisition conditions, including laser intensity, photomultiplier amplification, and Z-stack step size.

4.3. Image analysis

All images were processed and analyzed using FIJI software (ImageJ, NIH; [43]). For measurements, we used areas of 300 x 300 µm2, merged (from 5-8 optical slices) images. BVs, labeled by Albumin, were traced with thresholding in FIJI and their total area was measured. NG2-positive PC

numbers were counted along capillaries, taking into account the local BV diameter and cell morphology, using the cell counter plugin in FIJI. In total, we evaluated 1565 individual PCs from all sample sets.

Albumin labeling was also used to evaluate BV length. After initial thresholding and global processing, images were converted to binary black and white. We used the dilate and erode binary function subsequently to smooth local unevenness at BV edges. Skeletonized BVs were measured using the analyze skeleton function in FIJI. Results were transferred to Excel for further processing.

Distal capillaries were specified by counting the 2nd or 3rd bifurcation from the arteriole and considering the diameter of BV and PC morphology (bump on the log) specific for distal capillaries (Fig 4). The diameter evaluation was made on composite images. First, a custom-made ImageJ script was used to generate random points on the images. We measured the diameter with the FIJI line tool on the nearest capillary from this point, next to the PC soma (Fig 4d).

4.4. Statistical analysis

For statistical analysis, Excel (Microsoft Corp., Redmond, USA), JASP (JASP Team, 2019 JASP, V 0.11.1), and R (R Core Team 2013) were used. An independent t-test was used to determine the BV/PC number ratio significance. For multiple comparisons, analysis of variance (ANOVA) with post-hoc Tukey's test or repeated-measures ANOVA was used with Pearson's correlation matrix. The number of samples (n) indicates the number of mice in the group or else indicated otherwise locally. The data is presented as mean ± SD unless otherwise indicated.

Author Contributions: conceptualization, T.KÖ. and B.S.; methodology, T.KÖ., E.I.; validation, T.KÖ., B.S.; formal analysis, T.KÖ., G.S., A.T.; investigation, T.KÖ.; resources, B.S.; data curation, T.KÖ., G.S., A.T.; writing—original draft preparation, T.KÖ.; writing—review and editing, B.S., B.V., E.I., A.T., G.S., T.KÖ.; visualization, T.KÖ.; supervision, B.S.; funding acquisition, B.S., B.V.

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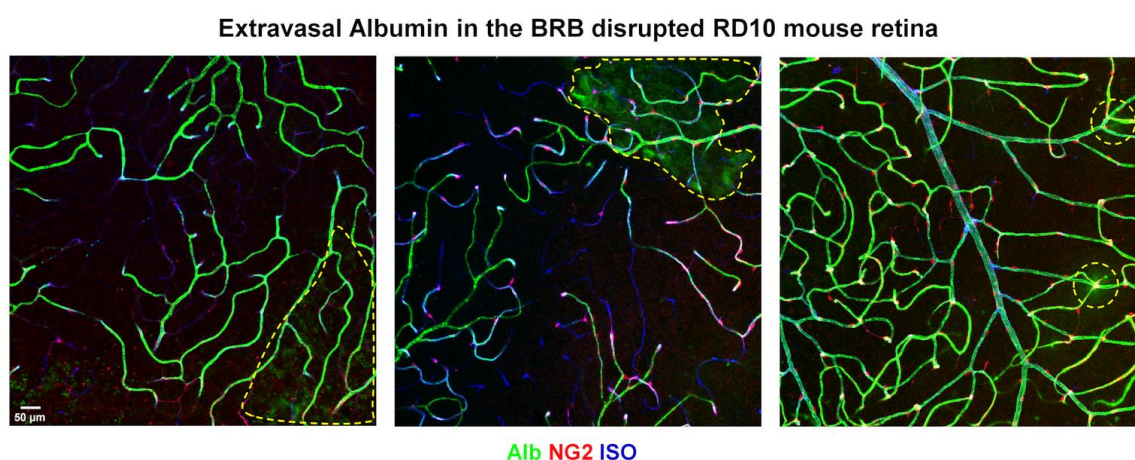
Conflicts of Interest: The authors declare no conflict of interest.

Supplementary Materials: [Supplementary materials can be found at www.mdpi.com/xxx/s1.]

Extravasal Albumin in the BRB disrupted RD10 mouse retina

Alb NG2 ISO

Supplementary figure 1. Albumin detection in the BRB disrupted RD10 mouse retina using the same IHC-method serves as a positive control for albumin extravasation (encircled by yellow, dashed lines) not detectable in our sampleset. Alb – albumin; NG2 – Neural glial factor-2; ISO – Isolectin (from Griffonia simplicifolia - IB4)

Abbreviations

ALS	Amyotrophic lateral sclerosis
AMD	Age-related macular degeneration
BBB	Blood-brain barrier
BRB	Blood-retina barrier
BV	Blood Vessel
C	Control
Ch	Choroid
CNS	Central nervous system
Cspg4	Chondroitin sulfate proteoglycan 4
DL	Deep Layer
EC	Endothelial cell
GCL	Ganglion cell layer
GS-IB4	Griffonia simplicifolia - IB4
IHC	Immunohistochemistry
IL	Intermediate Layer
INL	Inner nuclear layer
IP	Intraperitoneal
IPL	Inner plexiform layer
IV	Intravitreal
NFL	Nerve fiber layer
NG2	Neuron-glial antigen 2
NVU	Neurovascular Unit
OIR	oxygen-induced retinal
ONL	Outer nuclear layer
OPL	Outer plexiform layer
PBS	Phosphate-buffered saline
PC	Pericytes
PDGF	Platelet-derived growth factor
PDGFR	Platelet-derived growth factor receptor
PDGFR-β	Platelet-Derived Growth Factor Receptor-Beta
PECAM1	Platelet and endothelial cell adhesion molecule 1
RPE	Retinal pigment epithelium
SD	Standard deviation
SL	Superficial Layer
SMC	Smooth muscle cells

VEGF   Vascular endothelial growth factor

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