A Cyclic Peptide Epitope of an Under-Explored VEGF-B Loop 1 Demonstrated In Vivo Anti-Angiogenic and Anti-Tumor Activities

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Pathological angiogenesis is mainly initiated by the binding of abnormal expressed vascular endothelial growth factors (VEGFs) to their receptors (VEGFRs). Blocking the VEGF/VEGFR interaction is a clinically proven treatment in cancer. Our previous work by epitope scan had identified cyclic peptides, mimicking the loop 1 of VEGF-A, VEGF-B and placental growth factor (PlGF), inhibited effectively the VEGF/VEGFR interaction in ELISA. We described here the docking study of these peptides on VEGFR1 to identify their binding sites. The cellular anti-angiogenic activities were examined by inhibition of VEGF-A induced cell proliferation, migration and tube formation in human umbilical vein endothelial cells (HUVECs). The ability of these peptides to inhibit MAPK/ERK1/2 signaling pathway was examined as well. On chick embryo chorioallantoic membrane (CAM) model, a cyclic peptide named B-cL1 with most potent in vitro activity showed important in vivo anti-angiogenic effect. Finally, B-cL1 inhibited VEGF induced human gastric cancer SGC-7901 cells proliferation. It showed anti-tumoral effect on SGC-7901 xenografted BALB/c nude mouse model. The cyclic peptides B-cL1 constitutes an anti-angiogenic peptide drug lead for the design of new and more potent VEGFR antagonists in the treatment of angiogenesis related diseases.

Keywords: VEGF, VEGFR, anti-angiogenic, loop mimetics, cyclic peptides

INTRODUCTION

Angiogenesis (formation of new blood vessels) plays a crucial role during the development of certain diseases, such as cancer and age-related macular degeneration (Carmeliet, 2003; Chung and Ferrara, 2011). For example, angiogenesis provides nutrients and oxygen supply for tumoral cells proliferation as well as for tumoral cells metastasis (Hanahan and Folkman, 1996). Anti-angiogenic therapy consists in inhibition of angiogenesis to cut down the diffusion of nutrients in pathological tissues, which is a clinically proven target therapy to treat angiogenesis related diseases (Potente et al., 2011; Apte et al., 2019; Eelen et al., 2020). The principle of anti-angiogenic therapy is to inhibit the interaction of pro-angiogenic factors with their receptors, such as the interaction of vascular endothelial growth factors (VEGF) and their receptors (VEGFR), or to inhibit...
tyrosine kinase activity of VEGFRs and the downstream signal transduction pathways (Ferrara et al., 2003; Vasudev and Reynolds, 2014). Thus, anti-angiogenic drugs research has been mostly focused on the inhibition of VEGF/VEGFR interaction or the disruption of VEGF downstream signal transduction.

Currently, three major types of anti-angiogenic drugs: antibodies, nucleotide aptamers and tyrosine kinase inhibitors, have been extensively used in clinical practice and achieved great benefits in the treatment of age-related macular degeneration (Ng et al., 2006) and several type of cancers, such as non-small cell lung cancer, gastric cancer and metastatic colorectal cancer (Carmeliet and Jain, 2011; Moserle et al., 2014). Tyrosine kinase inhibitors (sunitinib, sorafenib, axitinib or pazopanib) are mostly small molecules, they inhibit angiogenesis by blocking the kinase activity of intracellular domains of VEGFRs (Wilhelm et al., 2006; Faivre et al., 2007; Van Geel et al., 2012). They are oral administration drugs with good bioavailability but have low specificity and thus side effects (Aparicio-Gallego et al., 2011; Sun et al., 2013). Patients treated by tyrosine kinase inhibitors develop frequently drug resistance (Morais, 2014; Cabral et al., 2020). Antibodies, act as antagonists of VEGF or VEGFR, indirectly inhibiting intracellular protein kinase activity and downstream signaling pathway. They exert anti-angiogenesis activities by blocking the interaction between VEGF and VEGFR. For example, bevacizumab targets the VEGF and ramucirumab targets the VEGFR2, both blockade VEGF-VEGFR interaction driven angiogenesis (Kong et al., 2017). Antibodies have high specificity, but also high production cost and high pharmacokinetic variability (Paci et al., 2020). Meanwhile, recent research showed that peptides, especially cyclic peptides, which have the similar action mode as antibodies, showed high target specificity as well as good bioavailability and metabolic stability (Fosgerau and Hoffmann, 2015; Henninot et al., 2018; Choi and Joo, 2020). Thus, development of synthetic peptide as antagonists of the VEGF/VEGFR interaction become an attractive research topic.

Two major strategies are involved to develop peptide antagonists of the VEGF/VEGFR interaction: rational design or random screen of peptide libraries. Rational design of peptide antagonists is based on the complex structures of VEGF/VEGFR. VEGFs, especially VEGF-A, was reported as a key pro-angiogenic factor, which binds to membrane receptors including VEGFR1, VEGFR2, and NRP1 to exert the angiogenesis activity (Ferrara et al., 2003; Shibuya and Claesson-Welsh, 2006; Shibuya, 2014). VEGFR1 and VEGFR2 consist of seven extracellular immunoglobulin (Ig)-like domains (D1−D7), the deletion analysis showed that VEGF-A bends both on the second (D2) and the third domains (D3) of VEGFR1 (Wiesmann et al., 1997). The first structurally determined interaction between VEGF-A and VEGFRs was a complex between VEGF-A dimer and unique D2 of VEGFR1 (PDB: 1FLT) (Wiesmann et al., 1997). Numerous peptide inhibitors of VEGFRs have been designed based on this complex (Goncalves et al., 2007a; Basile et al., 2011; Garcia-Aranda et al., 2013; Wang et al., 2014). Later, more complete structures of interactions between VEGF-A and VEGFRs have been resolved, for example, a complex of VEGF-A with VEGFR2 D2-D3 (PDB: 3V2A) (Brozzo et al., 2012) and a complex of VEGF-A with VEGFR1 D1-D6 (PDB: 5T89) (Markovic-Mueller et al., 2017). These complexes showed that VEGF-A binds also partly to the third domain (D3) of VEGFR1 and VEGFR2. For example, the loop 1 of VEGF-A mainly binds to VEGFR1 D3 (Figure 1). Meanwhile, many other interaction structures of different growth factors with VEGFRs have been reported, but only with D2 of VEGFR1, such as placental growth factor (PlGF) dimer with D2 of VEGFR1 (PDB code 1VR6) (Christinger et al., 2004) and VEGF-B dimer with D2 of VEGFR1 (PDB: 2XAC) (Iyer et al., 2010). All these structure data promoted the rational design of peptide inhibitors targeting VEGF or VEGFRs for the purpose of blocking VEGF/VEGFR interaction to inhibit angiogenesis (Basile et al., 2011; Garcia-Aranda et al., 2013; Assareh et al., 2019), including our works (Goncalves et al., 2007a; Gautier et al., 2010; Wang et al., 2014; Reille-Seroussi et al., 2015; Wang et al., 2017; Wang et al., 2019).

We have recently reported the design of three cyclic peptides to mimic Loop 1 of VEGF-A, VEGF-B, and PlGF, which bind mainly to D3 of VEGFR1 and VEGFR2 according to the crystal structure data (Wang et al., 2017). These cyclic peptides showed competitive binding with VEGF to VEGFR1 and a primary tube formation inhibition in human umbilical vein endothelial cells (HUVECs) (Wang et al., 2017). In this study, we performed docking and binding energy calculation of these three cyclic peptides (named A-cL1, B-cL1, and P-cL1) with VEGFR1 to investigate the structure-activity relationship. We also examined the in vitro activities by evaluation of their inhibitory activity in the VEGF-A induced HUVECs proliferation, migration and tube formation by inhibiting MAPK/ERK1/2 signaling pathway. Peptide B-cL1, which showed the most important in vitro anti-angiogenic activity, inhibits human gastric cancer SGC-7901 cells proliferation. At last, B-cL1 was evaluated on chick embryo chorioallantoic membrane (CAM) model and human gastric cancer SGC-7901 cells xenographed BALB/c nude mouse model for its in vivo anti-angiogenic and antitumoral activity.

RESULTS

Docking With VEGFR1

VEGF-A, VEGF-B, and PlGF are high sequence homology ligands of VEGFR1, they bind similarly to VEGFR1 according to the structure data (PDB: 1FLT, 2XAC, and 1VR6). It has been then demonstrated that Loop 1 of VEGF-A binds mainly to D3 of VEGFR1 (Markovic-Mueller et al., 2017) and VEGFR2 (Brozzo et al., 2012). In our previous study, we designed peptides mimicking different binding epitopes of VEGF-A, VEGF-B, and PlGF (Wang et al., 2017). Cyclic peptides derived from Loop 1 showed very interesting inhibition activity to disrupt the interaction of VEGF-A/VEGFR1 in a VEGF-A/VEGFR1 interaction-based ELISA assay (Table 1). Although most of reported inhibitors target VEGFR1 D2, Wiesmann et al had shown by domain deletion analysis that the deletion of D3 from VEGFR1 D1-D2-D3 induce a loss of more than 20-fold
VEGF-A binding affinity (Wiesmann et al., 1997). In our reported study, cyclic peptides derived from Loop 1: A-cL1, B-cL1, and P-cL1 showed higher activity than those derived from the helix α1 or the loop 2 or the loop 3. We thus focus this study on these three peptides (sequences in Table 2, formulas in Supplementary Scheme S1).

Peptides A-cL1, B-cL1, and P-cL1 were respectively designed from Loop 1 of VEGF-A, VEGF-B, and PlGF, which were supposed to bind mainly to the domain 3 (D3) of VEGFR1. The docking results showed that A-cL1 and P-cL1 had similar interaction mode with VEGFR1, mainly binding to VEGFR1 D3 (Figures 2A,C). However, B-cL1, which had larger cycle, binds differently to VEGFR1. B-cL1 binds to the interface of VEGFR1 D2 and D3 (Figure 2B). The ten highest score peptide structures were superimposed in the binding with VEGFR1 (Supplementary Figures S1–S3). The major interactions between A-cL1, B-cL1, P-cL1, and VEGFR1 were analyzed according to the docking (Table 2).

### Inhibition of HUVECs Proliferation, Migration and Tube Formation

VEGFR1 and VEGFR2 have very similar structures. The crystal structure of VEGF-A/VEGFR1 D1-D6 (PDB: 5T89) and the crystal structure of VEGF-A/VEGFR2 D2-D3 (3V2A) showed the same binding epitopes on VEGF-A, which suggests that the epitope mimic peptides of VEGF-A are normally able to bind both VEGFR1 and VEGFR2. An in vitro inhibitor selection assay was previously developed in our laboratory based on the interaction of VEGF-A/VEGFR1 (Goncalves et al., 2007b). Inhibitors selected from this assay that are able to inhibit the interaction of VEGF-A/VEGFR1, should probably inhibit as well...

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**TABLE 1 |** Peptide ID, original Loop 1 sequence and derived cyclic peptide sequence. IC50 were determined in an ELISA based on the VEGF-A/VEGFR1 interaction.

| Peptide ID | Original Loop 1 sequence | Cyclic peptide sequence | IC50 (μM) determined by ELISA [37] |
|------------|---------------------------|-------------------------|-----------------------------------|
| A-cL1      | (VEGF-A) FQEYPDEIEYIK50   | [CQEYPDEIEYIC]K         | 50.4 ± 11.5                       |
| B-cL1      | (VEGF-B) LTVELMGTVAKQLVPS20 Ac-[CTVELMGTVAKQLVPC] | 10.4 ± 2.8               |
| P-cL1      | (PlGF) VSEYPSEVEHMFS26   | [CSEYPSEVEHMC]S         | 56.0 ± 11.4                       |

**TABLE 2 |** The interaction analysis of peptide A-cL1, B-cL1, and P-cL1 with VEGFR1 according to the docking.

| Type of interaction | A-cL1 [CQEYPDEIEYIC]K13 | B-cL1 Ac-[CTVELMGTVAKQLVPC]s | P-cL1 [CSEYPSEVEHMC]S13 |
|---------------------|---------------------------|-------------------------------|------------------------|
| Hydrogen bonds      | P9 (O) with ARG280 (HN); E9 (O) with GLN263 (HNE2) | None | P9 (O) with ARG280 (HN); E9 (O) with GLN263 (HNE2) |
| Hydrophobic effects | I8 with VAL278, PHE292; I11 with HIS223; ARG261; K13 with ILE145, HIS223 | V6 with ILE145; A10 with HIS223; K11 with ARG281 | V6 with VAL278; ARG280; PHE292 |

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**FIGURE 1 |** Interaction between VEGF-A with VEGFR1 D1 to D6. The complex structure of VEGF-A dimer with VEGFR1-D1-D6 was adopted from PDB ST89 (Markovic-Mueller et al., 2017). The VEGF-A/VEGFR binding regions were framed in blue dash lines. Loop 1 of VEGF-A, circled in red dash lines, binds mainly to VEGFR1-D3.
as the interaction of VEGF-A/VEGFR2 and their downstream signaling. The investigation of A-cL1, B-cL1, and P-cL1 induced anti-angiogenic effects was performed by cell proliferation, cell migration and tube formation in HUVECs.

Inhibition of HUVECs Proliferation

As shown in Figure 3, additional VEGF-A (0.2 μg/ml) in culture medium stimulated HUVECs proliferation (control group). Bevacizumab (Avastin®), a monoclonal antibody drug targeting VEGF-A to block the interaction of VEGF-A with VEGFRs, was tested as positive control. At 6.5 μM (1 mg/ml), bevacizumab resulted in strong inhibition of VEGF-A induced HUVECs proliferation. Peptide A-cL1, B-cL1, and P-cL1 showed a dose-dependent inhibition of HUVECs proliferation at five concentrations (0.2, 1, 5, 25, and 125 μM) (Figure 3). A-cL1 and B-cL1 showed better inhibition of HUVECs proliferation than P-cL1. They showed similar inhibition activity at 25 μM as bevacizumab at 6.5 μM, while B-cL1 exhibited more effective inhibition at 0.2 μM than A-cL1. All the three peptides showed slight cytotoxicity at 125 μM.

Inhibition of HUVECs Migration

The inhibition of VEGF-A stimulated cell migration by A-cL1, B-cL1, and P-cL1 was evaluated by a wound healing assay.
FIGURE 4 | A-cL1, B-cL1, and P-cL1 inhibit VEGF-A (0.2 μg/ml) stimulated HUVECs migration. (A) Representative images of scratch wound-healing assay. After creation of scratches, HUVECs were cultured in 2% serum medium containing 0.2 μg/ml VEGF-A, treated with serum-free medium (control group, blue), or with 6.5 μM bevacizumab (positive control, red), or with 1 μM, 5 and 25 μM of A-cL1, B-cL1 or P-cL1. Images were taken before the treatment (T0h), after 6 h treatment (T6h) and 12 h treatment (T12h). Scale bar, 100 μm. (B) Quantitative analysis of the cell migration (%) by measuring wound closure area after 6 h treatment using ImageJ. (C) Quantitative analysis of the cell migration (%) by measuring wound closure area after 12 h treatment using ImageJ. The relative cell migration (%) were analyzed with GraphPad Prism 8 (mean ± SEM of three independent experiments), compared to the control group in one-way ANOVA statistical analysis. *p < 0.05, **p < 0.01, ***p < 0.001, ****p < 0.0001, “ns” meaning non-significant.

FIGURE 5 | A-cL1, B-cL1, and P-cL1 inhibit VEGF-A stimulated tube formation in HUVECs. (A) Representative images of the capillary tube formation assay. After seeded on Matrigel® HUVECs were cultured in 2% serum medium containing 0.2 μg/ml VEGF-A, treated with serum-free medium (control group), or with bevacizumab (positive control), or with 1 μM, 5 and 25 μM of A-cL1, B-cL1 or P-cL1. Images were taken after 6 h treatment. (B) Quantitative analysis of the tube formation by counting the capillary number. Data are presented as mean ± SEM of three experiments, statistics analysis using GraphPad Prism 8, compared to the control group in one-way ANOVA. ***p < 0.001, ****p < 0.0001.
After stimulation with VEGF-A (0.2 μg/ml), HUVECs were able to migrate through the scratched area, and completely fill the scratched area after 12 h (Figure 4, control group). Bevacizumab (6.5 μM) resulted in efficient inhibition of HUVECs migration. After treatment with A-cL1, B-cL1, and P-cL1, HUVECs migration was inhibited. B-cL1 and A-cL1 induced dose-dependent inhibition of HUVECs migration. B-cL1 was able to inhibit strongly HUVECs migration at 5 μM, less than 10% of migration were observed at both 6 and 12 h. A-cL1 inhibited partially HUVECs migration at 5 μM, but it could also completely inhibit HUVECs migration at 25 μM. P-cL1 showed a week inhibition of HUVECs migration at 6 h even at 25 μM, HUVECs filled up the scratched area after 12 h treatment of P-cL1 at the three concentrations.

Inhibition of HUVECs Tube Formation
Capillary-like tube formation is one of the most important pseudo angiogenic test using HUVECs. In this test, two-dimensional tube formation was stimulated by additional VEGF-A (0.2 μg/ml) on Matrigel®. A-cL1, B-cL1, and P-cL1 were tested at three concentrations (1 μM, 5 and 25 μM) for their ability to inhibit VEGF-A stimulated tube formation in HUVECs. The total numbers of formed tubes were measured (Figure 5). All the three peptides showed a dose-dependent inhibition of tube formation in HUVECs. B-cL1 was able to inhibit significantly HUVECs tube formation at 25 μM, and it had an equal inhibition activity at 5 μM as bevacizumab (6.5 μM). A-cL1 also exhibited a good inhibition of VEGF-A stimulated tube formation, it had a similar inhibition activity at 25 μM as bevacizumab (6.5 μM). P-cL1 inhibited VEGF-A stimulated tube formation in HUVECs in a dose-dependent manner with a weaker activity than A-cL1 and B-cL1, short capillary-like structures were still observed after treatment with P-cL1 at 25 μM.

These results showed that peptides A-cL1, B-cL1, and P-cL1, derived from epitope Loop 1 of VEGF-A, -B, and PI GF can inhibit VEGF-A/VEGFRs interaction induced migration and tube formation in HUVECs. B-cL1 showed the highest inhibitory activity, while P-cL1 showed the lowest activity among the three peptides. The results are in agreement with peptides IC50 determined in a VEGF/VEGFR1 interaction-based ELISA assay (Wang et al., 2017) (Table 1).

Inhibition of Intracellular Signaling Pathways Associated With VEGF
A-cL1, B-cL1, and P-cL1 having different VEGFR1 binding affinity showed dose-dependent different inhibitory effects in HUVECs migration and tube formation. In order to verify whether the different activity of A-cL1, B-cL1, and P-cL1 could result in different intensity of downstream signal transduction, we performed western blot analysis of the p-ERK1/2 level in the MAPK/ERK1/2 signaling pathway, which is a proven signal pathway induced by the interaction of VEGF/VEGFRs (Shibuya, 2011). After HUVECs incubation with A-cL1, B-cL1, and P-cL1 at different concentrations (1 μM, 5 and 25 μM) in the presence of VEGF-A (0.2 μg/ml), the p-ERK1/2 level was determined in cell lysate. As shown in Figure 6, bevacizumab (6.5 μM) had the most important inhibition of p-ERK1/2 formation. A-cL1 and B-cL1 significantly decreased the level of p-ERK1/2 in a dose-dependent manner, while P-cL1 showed a limited decrease of p-ERK1/2 formation compared to the control.

B-cL1 Inhibits Angiogenesis on Chick Embryo Chorioallantoic Membrane Model
Among the three peptides, B-cL1 showed the most important in vitro anti-angiogenic activity. In order to further study the in vivo anti-angiogenic activity of B-cL1, we performed an evaluation in a chick embryo chorioallantoic membrane (CAM) model, which is a common physiological and pathological angiogenesis study model in vivo (Rezzola et al., 2020). As shown in Figure 7A, on day 9, capillaries are well formed on the surface of the chick embryo chorionic epithelium. After treatment with different concentrations of B-cL1 (1 μM, 5
and 25 μM) for 48 h, formation of new capillaries was inhibited compared with control treated by PBS (Figure 7A). At 1 μM of B-cL1, short capillaries formation was still observed. However, the increase in thickness of blood vessels was inhibited. At 5 μM of B-cL1, both the formation of new capillaries and the development of blood vessels thickness were inhibited. At 25 μM of B-cL1, destruction of pre-existing capillaries was observed. AngioTool (Zudaire et al., 2011), a software for quantitative analysis of angiogenesis, which allows to quantify the percentage of capillary area in total area (Klotz et al., 2015; Sawaguchi et al., 2017), was used to analyze the increasing or decreasing vessels area after treatment. The results showed that B-cL1 was able to inhibit new capillaries formation in a dose-dependent manner (Figure 7B).

**B-cL1 Inhibits Human Gastric Cancer SGC-7901 Cell Proliferation and Tumor Growth on Human Gastric Cancer Xenografted BALB/c Nude Mouse Model**

Gastric cancer is one of the most common malignancy that threaten the health of human beings, and the human gastric cancer SGC-7901 cell line is widely used in anti-tumoral studies in cellular level and on xenografted mouse model (Lv et al., 2019). This kind of tumor is largely explored in clinical trial by anti-angiogenic approach, including anti-VEGF, which have shown some benefits in second line treatment (Park et al., 2015). Consequently, anti-tumoral activity of B-cL1 was evaluated by VEGF-induced (50 ng/ml) human gastric cancer SGC-7901 cell proliferation and on BALB/c mice subcutaneous xenograft of human gastric cancer SGC-7901 cells.

B-cL1 showed a dose-dependent inhibition of human gastric cancer SGC-7901 cell proliferation at five concentrations (0.2, 1, 5, 25 and 125 μM), no cytotoxicity was observed at 125 μM (Figure 8A). It showed similar anti-proliferation activity at 25 μM as bevacizumab at 6.5 μM. Meanwhile, B-cL1 (5 mg/kg/day and 10 mg/kg/day) and bevacizumab (5 mg/kg, only once) were intravenously administrated for 2 weeks on BALB/c nude mice subcutaneous xenograft of SGC-7901 cells, starting when the tumor volume arrived at 150–300 mm³. Tumor volume and mice body weight were measured every 2 days. After 14 days of treatment, the mice were euthanized using CO₂ followed by cervical dislocation to ensure death and the tumors were separated (Figure 8B). The results showed that B-cL1 (5 mg/kg/day) reduced 60% of tumor weight, 53% of tumor volume, compared to control group; B-cL1 (10 mg/kg/day) showed 62% reduction of tumor weight, 51% reduction of tumor volume, compared to control group. As a positive control, bevacizumab (5 mg/kg) reduced 57% of tumor weight and 57% of tumor volume with only one administration (Figures 8C,D). It is worth to indicate that no mortality of mice was observed during the 2 weeks of treatment, and the body weights of mice was increased reasonably (Figure 8E), which suggest weak toxicity of B-cL1 for an administration up to 10 mg/kg/day.

**DISCUSSION**

Blocking VEGF/VEGFRs interaction is a strategy to treat angiogenesis related diseases. Antibody drugs, such as bevacizumab, ramucirumab, ranibizumab, and fusion protein, such as aflibercept, can bind either VEGF or VEGFRs to inhibit VEGF/VEGFRs interaction (Lazzeri et al., 2015; Eelen et al., 2020). They have been approved in clinical treatment of multitype of cancers or neovascular (wet) age-related macular degeneration (AMD). Current research of peptide-based inhibitors for VEGF/VEGFRs interaction mainly start with rational design or random screen of peptide libraries targeting mainly the D2 of VEGFRs or the VEGF (Basile et al., 2011; Garcia-Aranda et al., 2013; De Rosa et al., 2014; Bayo-Puxan et al., 2016). They have shown ability to inhibit VEGF/VEGFRs interaction, and some of them showed in vitro activities. However, their in vivo activities were less reported. Previously, we have designed several series of peptides mimicking different VEGFRs binding epitopes (helix and loops) of VEGF-A, VEGF-B, and PIGF (Goncalves et al., 2007a; Gautier et al., 2010; Wang et al., 2014; Wang et al., 2017; Wang et al., 2019). Three peptides...
respectively mimicking Loop 1 of VEGF-A, VEGF-B or PlGF showed the most important inhibition of VEGF-A/VEGFRs interaction in an in vitro competition assay (Wang et al., 2017). Here, we performed VEGF-A induced cell migration and tube formation assays in HUVECs, followed by western blot analysis of VEGFRs downstream signaling studying ERK activation. We also performed the evaluation of physiological anti-angiogenesis activity in chick embryo chorioallantoic membrane (CAM) model. Finally, we evaluated the inhibition of human gastric cancer SGC-7901 cells proliferation and tumor growth activity in SGC-7901 cells subcutaneous xenograft model on BALB/c nude mice. All these experiments allowed us to investigate the in vitro and in vivo anti-angiogenesis activity as well as anti-tumor activity of this series of cyclic peptides derived from under-explored Loop 1.

The docking results identified the different binding sites on VEGFR1 of A-cL1, B-cL1, and P-cL1. A-cL1 and P-cL1 bind similarly as the original Loop 1 structure of VEGF-A and PlGF to the domain 3 (D3) of VEGFR1. However, B-cL1 with a larger cycle binds differently. It binds to the interface of VEGFR1 D2-D3 with higher flexibility (Supplementary Figure S2). VEGF-A was reported to bind tightly to VEGFR1 D1-D7. The deletion of D1 or D4-D7 had slight influence on VEGF binding affinity, the deletion of D3 could cause a 25-fold diminution of binding affinity. However, D2 stays as the major binding domain of VEGF-A (Wiesmann et al., 1997). A-cL1 and P-cL1 binding majorly to D3 had higher IC₅₀ (50 and 56 µM respectively), whereas B-cL1 binding to both D2 and D3 had lower IC₅₀ (10 µM) (Table 1) (Wang et al., 2017). The docking results showed that targeting only D3 is insufficient to inhibit VEGF/VEGFR interaction.

![Figure 8](image-url)
The *in vitro* anti-angiogenic activities of A-cL1, B-cL1, and P-cL1 were evaluated in VEGF-induced HUVECs proliferation, migration and tube formation. The results showed that their different inhibitory activity was correlated to their IC\(_{50}\) values of VEGF-A/VEGFR1 interaction inhibition (*Table 1*). B-cL1, having the lowest IC\(_{50}\) value (10 \(\mu\)M) measured in a VEGF/VEGFR1 interaction-based ELISA assay, demonstrated the most potent inhibition in VEGF-induced HUVECs proliferation, migration and tube formation. Moreover, B-cL1 showed the most potent inhibitory effect on VEGF-induced VEGFRs signaling through MAPK/ERK1/2 pathways. These results showed that B-cL1, designed from Loop 1 of VEGF-B, had very important *in vitro* anti-angiogenic activity by binding to VEGFRs to inhibit VEGF-A/VEGFRs interaction, thus inhibiting VEGFRs signal transport.

The *in vivo* evaluation of B-cL1 in CAM model showed that B-cL1 was able to inhibit physiological angiogenesis on the surface of chorionic epithelium during the chick embryo development at 5 \(\mu\)M, and more significantly at 25 \(\mu\)M.

The anti-tumor activities of B-cL1 were then studied. B-cL1 was able to inhibit *in vitro* in a dose-dependent way VEGF-induced human gastric cancer SGC-7901 cells proliferation. In SGC-7901 cells subcutaneous xenograft BALB/c mouse model, intravenous administration of bevacizumab (5 \(mg/kg\), just once injection for 2 weeks) significantly inhibit tumor growth. Bevacizumab, as recommended, was injected by intravenous every 14 days at the dose of 5 \(mg/kg\) (Garcia et al., 2020). However, peptides are commonly considered with poor *in vivo* stability (Diao and Meibohm, 2013; Henninot et al., 2018). Thus, we performed *in vivo* evaluation of cyclic peptide B-cL1, at doses 5 \(mg/kg\) and 10 \(mg/kg\)/day. The results showed similar tumor growth inhibition as bevacizumab at 5 \(mg/kg\). No obvious toxicity of B-cL1 was observed up to 10 \(mg/kg\)/day. Interestingly, non-significant difference in tumor growth inhibition was observed for administration of 5 \(mg/kg\)/day and 10 \(mg/kg\)/day of B-cL1, which suggests that 5 \(mg/kg\)/day of B-cL1 can induce maximal tumor growth inhibition. The tumor growth inhibition observed *in vivo* is probably correlated with the inhibition of ERK1/2 phosphorylation observed *in vitro* on HUVE cells. Which may be the result of the inhibition of VEGF-VEGFR interaction. Recently, the group of S. M. Asghari has designed a series of peptides derived from VEGF-A and VEGF-B epitopes binding to D2 of VEGFR. The peptides named VGB (Sadremontaz et al., 2018), VGB1 (Assareh et al., 2019), VGB4 (Farzaneh Behelgardi et al., 2018), and VGB3 (Sadremontaz et al., 2020) were tested in murine mammary carcinoma tumor model (MCT) where BALB/c mice were implanted with murine breast cancer 4T1 cells. In VEGFR1-dependent MCT model, the peptides exhibited inhibition of tumor growth at doses of 5 \(mg/kg\), with peptide VGB3 displaying effects at a very low dose of 0.2 \(mg/kg\). Therefore, investigation of the peptide B-cL1 on tumor models characterized by various expression levels of VEGFR1 and VEGFR2 will be of interest to determine its anti-angiogenesis activity profile more accurately. Otherwise, our results, coupled with those of the group of S. M. Asghari, show a relative stability of cyclic peptides in physiological medium.

Peptide antagonists reported till now are all D2 targeting molecules except this series of cyclic peptides A-cL1, B-cL1, and P-cL1. In this study, we determined by molecular docking that B-cL1 binds D2 and D3 simultaneously, but A-cL1 and P-cL1 bind mainly to D3. We confirmed that B-cL1 has higher anti-angiogenic activities than A-cL1 and P-cL1. The results open a new way to develop VEGFR inhibitors. Indeed, the conception of Affibrecpt, consisting of VEGFR1-D2 and VEGFR2-D3 fused to human IgG1 Fc portion (Lazzeri et al., 2015), supports the importance of targeting both D2 and D3 of VEGFR in the research of VEGFR inhibitors.

### MATERIALS AND METHODS

#### Peptide Inhibitors

The design and synthesis of peptides derived from different epitopes of VEGFR1 binding sites on VEGF-A, VEGF-B, and PlGF have been reported in our previous work (Wang et al., 2017). Briefly, to mimic the loop structure, the original peptide sequences have been modified and then cyclized through disulfide bonds. To facilitate the reading, we rename these peptides (peptide 14, 18, and 19 in previous work) as described in the *Table 1*.

#### Docking With VEGFR1

Protein Preparation Wizard Workflow program (Gayatri et al., 2016) provided in Maestro 9.0 was used to build the structures of peptides A-cL1, B-cL1, and P-cL1 based on the original crystal structures of VEGF-A, VEGF-B, and PlGF with VEGFR1-D2 (PDB code: 1FLT, 2XAC, 1RV6). Peptides were built by mutating underlined residues to cysteine residues and formed disulfide bond on the side chains (*Table 1*). They were then aligned and docked near the binding pocket of Loop1 on VEGFR1 (using complex structure of VEGF-A with VEGFR1 D1-D6, PDB: 5T89) in Pymol to generate the initial PDB complex file of A-cL1/5T89, B-cL1/5T89 and P-cL1/5T89. Afterwards, the peptide-protein complexes were adapted to optimize the side chains, in order to eliminate internal collision that are not related to intermolecular interactions in Rosetta FlexPepDock program (Raveh et al., 2010). At last, the docking was performed with optimized structures of the three peptides on Rosetta online server (http://flexpepdock.furmanlab.cs.huji.ac.il/index.php) (Raveh et al., 2010; London et al., 2011). Ten structure models of each peptide with highest score were obtained among 200 generated structures (*Supplementary Figures S1–S3*) and were output for binding analysis.

#### Cell Culture

Human umbilical vein endothelial cells (HUVECs), of passage 3 to 6, were purchased from Shanghai Fuheng Biotechnology Co. LTD. and cultured in endothelial cell culture medium (ECM: Gibco, Life Technologies, United States), with 10% fetal bovine serum (FBS: Gibco, Life Technologies, United States), 100 U/mL penicillin and 100 mg/ml streptomycin in an incubator at 37°C and containing 5% carbon dioxide (CO\(_2\)). Human gastric cancer SGC-7901 cells, from Cell Bank of the Chinese Academy of
incubated at 37°C for 6 h. Resulting tube networks were observed by Cell Counting Kit-8 (CCK-8, Sigma, United States) assays. The absorbance was measured at 450 nm with an enzyme-linked immunosorbent assay reader (AMR-100, ALLSHENG, Hangzhou, China). The experiment was repeated three times.

**VEGF-Induced HUVECs Proliferation Assay**
HUVECs (3 × 10⁴) were seeded on the 96-well plate per well in ECM containing 5% FBS, and incubated overnight at 37°C, with 5% CO₂. Then the medium was removed. New serum-free medium containing 0.2 μg/ml VEGF-A (R&D Systems, United Kingdom) with different concentrations of peptides (0.2, 1, 5, 25 and 125 μM), bevacizumab (Roche, Switzerland) at 6.5 μM (1 mg/ml) or control group (serum-free medium) were added and incubated for another 48 h (6 wells/concentration/group). The effects of proliferation were quantified by Cell Counting Kit-8 (CCK-8, Sigma, United States) assays. The absorbance was measured at 450 nm with an enzyme-linked immunoassay reader (AMR-100, ALLSHENG, Hangzhou, China). The experiment was repeated three times.

**VEGF-Induced Wound Healing Assay in HUVECs**
The six-well plate was coated with HUVECs and incubated with ECM containing 10% FBS until the confluence of monolayer cells reached 100%. A scratching wound was created with 10 μl pipette tip. Then the medium was removed. New medium containing only 2% serum and 0.2 μg/ml (5 nM) VEGF-A (R&D Systems, United Kingdom) with different concentrations of peptides (1 μM, 5 and 25 μM), bevacizumab (1 mg/ml, 6.5 μM) or 2% serum medium (control group) were added and incubated at 37°C, with 5% CO₂ for another 12 h. Cell migration was observed and photographed at 6 and 12 h under a microscope (TE-2000U, NIKON, Japan). The results were calculated from three independent experiments with four replicates. Wound area (the open area of the scratch) was quantified using ImageJ, and the percentage of wound closure were calculated: [1 - (wound area at 6 h or 12 h/wound area at 0 h)] × 100%.

**HUVECs Tube Formation Assay**
Growth factor-reduced basal membrane extract (BD Biocoat™, 356,230, United States) was melted overnight in refrigerator at 4°C, and precoated on the 96-well plate. The 96-well plate was then incubated at 37°C for 1 h until Matrigel solidified. 5 × 10⁴ HUVECs/well, pre-treated with different concentrations of peptides (1 μM, 5, and 25 μM), bevacizumab (6.5 μM) or 2% serum medium (control group) were added to the plate and incubated at 37°C for 6 h. Resulting tube networks were observed and photographed by using the microscope (TE-2000U, NIKON, Japan). The numbers of cell branches were quantified in a blind manner. The results were calculated from three independent experiments with four replicates.

**Western Blot Analysis**
HUVECs treated with different concentrations of peptides (1 μM, 5, and 25 μM), bevacizumab (6.5 μM) or serum-free medium (control group) were lysed with RIPA lysis buffer (Beyotime, Shanghai, China) containing protease and phosphatase inhibitors (Sigma, United States). After centrifugation at 4°C, protein concentration was determined by bicinchoninic assay (BCA) (Beyotime, Shanghai, China). Polyacrylamide gel electrophoresis was performed with the same amount of protein of different groups. The protein on the gel was then transferred to the PVDF membrane (Millipore, United States), and the membrane was blocked with skimmed milk powder in Tris-buffered saline (TBS, 5%) for 2 h. Later, the membrane was washed with TBST (1 × Tris Buffered saline, 0.1%Tween20), and incubated with the primary antibody (anti-phospho ERK1/2, anti-ERK1/2, and anti-GAPDH as control, Cell Signaling Technology, United States) and TBST overnight at 4°C. The membrane was washed 3 times with TBST, and then incubated with horseradish peroxidase-conjugated goat anti-mouse IgG secondary antibody and goat anti-rabbit IgG secondary antibody for 1 h at room temperature. At last, proteins were analyzed by ECL reagent (WesternBright ECL, United States) following the manufacturer’s protocol, after 3 times washing with TBST. The experiment was repeated three times. Protein expression was quantified by ImageJ.

**In Vivo Anti-Angiogenic Evaluation of B-cL1 Using the Chick Chorioallantoic Membrane Assay**
A chick chorioallantoic membrane (CAM) assay was carried out to determine the in vivo anti-angiogenic activity of peptide B-cL1. Two-days old fertilized specific pathogen free (SPF) eggs (Ningbo Chunpai Agricultural Science and Technology Co. LTD., Ningbo, China) were incubated at 37.5°C, 60–70% relative humidity for 1 week. Blood vessels in SPF eggs were observed within lighting. Then a window of about 2 cm² was opened on the eggs and a sterilized silica gel ring (1.5 cm inner diameter) was placed onto the CAM. 20 μL of different concentration of B-cL1 (1 μM, 5, and 25 μM in PBS) or PBS (control group) were added inside the silica gel ring (10 eggs per group). The window was recovered with sterilized tape, and the eggs were incubated for another 48 h. The sterilized tape was removed, and the area of capillary blood vessels photographed under a stereomicroscope (Soptop, Shanghai, China). The experiment was repeated three times, and the percentage of capillary area in total analyzed area was calculated using AngioTool (Zudaire et al., 2011). The percentage of increasing/decreasing capillary area was then calculated: percentage of capillary area at 48 h - percentage of capillary area at 0 h.

**VEGF-Induced Human Gastric Cancer SGC-7901 Cells Proliferation Assay**
Human gastric cancer SGC-7901 cells (2 × 10⁴) were seeded on the 96-well plate per well in RPMI 1640 medium containing 2% FBS, and incubated overnight at 37°C, with 5% CO₂. Then the medium was removed. New serum-free medium containing 50 ng/ml VEGF-A (R&D Systems, United Kingdom) with different concentrations of peptides (0.2, 1, 5, 25, and 125 μM),
bevacizumab (Roche, Switzerland) at 6.5 μM (1 mg/ml) or control group (serum-free medium) were added and incubated for another 48 h (6 wells/concentration/group). The effects of proliferation were quantified by Cell Counting Kit-8 (CCK-8, Sigma, United States) assays. The absorbance was measured at 450 nm with an enzyme-linked immunoassay reader (AMR-100, ALLSHENG, Hangzhou, China). The experiment was repeated three times.

**In Vivo Antitumor Study of B-cL1 on Xenografted Mouse Model**

The evaluation of antitumoral activity of B-cL1 was performed in Laboratory of Experimental Animal Science, Hangzhou Normal University (Hangzhou, China) maintained under standardized environmental conditions, with approved protocols by the Institutional Animal Care and Use Committee (IACUC) of Hangzhou Normal University. Human gastric cancer SGC-7901 cells (1 x 10⁶ cells/500 μL) from Cell Bank of the Chinese Academy of Sciences (Shanghai, China) were injected to the right flanks of 8 weeks old female BALB/c mice. When tumor size grown to 100–300 mm³, BALB/c mice were randomized to groups (n = 6). B-cL1 (5 mg/kg/day and 10 mg/kg/day) and PBS (control group) were intravenously administrated for 2 weeks, bevacizumab (5 mg/kg) was intravenously administrated only once during the 2 weeks as positive control. The tumor volume was measured every 2 days by a digital Vernier caliper, using the following formula: \( v = a^2 \times b \times 0.52 \) (where a is the shortest diameter of tumor and b is the longest diameter of tumor). After 14 days of treatment, the mice were euthanized using CO₂ followed by cervical dislocation to ensure death. Then, the tumors were separated, photographed and weighed on balance (OHAUS Adventure, United States).

**Statistical Analysis**

Data are expressed as the arithmetic mean ± SEM of at least three different experiments using the GraphPad Prism software version 8.00 (San Diego, United States). The statistical significance of results was evaluated by one-way ANOVA, with probability values *\( p < 0.05 \), **\( p < 0.01 \), ***\( p < 0.001 \), ****\( p < 0.0001 \), being considered as significant, “ns” meaning non-significant.

**CONCLUSION**

Our designed peptide B-cL1, mimicking Loop 1 of VEGF-B, showed significant anti-angiogenic and anti-tumor activities both in vitro and in vivo. B-cL1 can thus be considered as a peptide lead for anti-angiogenic drug research. The results in this study suggest targeting simultaneously D2 and D3 domains of VEGFR as a new concept to develop potent VEGFRs antagonists. The optimization of B-cL1 to achieve better dual binding peptides are under investigation.

Anti-VEGF therapeutic agents like antibodies have been generally used in combination with a cytotoxic agent. Recent studies demonstrate their beneficial use in combination with programmed cell death protein 1 (PD-1)/programmed cell death ligand 1 (PD-L1) interaction inhibitors in various solid tumor types (Gao and Yang, 2020). New peptides with different pharmacokinetic properties as compared to antibodies will be explored as potential drugs in combination with such inhibitors.

**DATA AVAILABILITY STATEMENT**

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

**ETHICS STATEMENT**

The animal study was reviewed and approved by Zhejiang Sci-tech University and Institutional Animal Care and Use Committee (IACUC) of Hangzhou Normal University.

**AUTHOR CONTRIBUTIONS**

Conceptualization, LW and W-QL; peptides preparation, LW, SB, and W-QL; docking study, LZ, FY, and JJ; in vitro and in vivo assays, LW, MX, and HH; writing, original draft preparation, LW, MX, HH, LZ, FY, JJ, HF, GC, and JC; writing, review and editing, LW, ZL, SB, MV, and W-QL; funding acquisition, LW and MV. All authors have read and agreed to the published version of the manuscript.

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**SUPPLEMENTARY MATERIAL**

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fphar.2021.734544/full#supplementary-material
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