Mechanical Activation of Hypoxia-Inducible Factor 1α Drives Endothelial Dysfunction at Atheroprone Sites

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Objective—Atherosclerosis develops near branches and bends of arteries that are exposed to low shear stress (mechanical drag). These sites are characterized by excessive endothelial cell (EC) proliferation and inflammation that promote lesion initiation. The transcription factor HIF1α (hypoxia-inducible factor 1α) is canonically activated by hypoxia and has a role in plaque neovascularization. We studied the influence of shear stress on HIF1α activation and the contribution of this noncanonical pathway to lesion initiation.

Approach and Results—Quantitative polymerase chain reaction and en face staining revealed that HIF1α was expressed preferentially at low shear stress regions of porcine and murine arteries. Low shear stress induced HIF1α in cultured EC in the presence of atmospheric oxygen. The mechanism involves the transcription factor nuclear factor-κB that induced HIF1α transcripts and induction of the deubiquitinating enzyme Cezanne that stabilized HIF1α protein. Gene silencing revealed that HIF1α enhanced proliferation and inflammatory activation in EC exposed to low shear stress via induction of glycolysis enzymes. We validated this observation by imposing low shear stress in murine carotid arteries (partial ligation) that upregulated the expression of HIF1α, glycolysis enzymes, and inflammatory genes and enhanced EC proliferation. EC-specific genetic deletion of HIF1α in hypercholesterolemic apolipoprotein E–deficient mice reduced inflammation and endothelial proliferation in partially ligated arteries, indicating that HIF1α drives inflammation and vascular dysfunction at low shear stress regions.

Conclusions—Mechanical low shear stress activates HIF1α at atheroprote regions of arteries via nuclear factor-κB and Cezanne. HIF1α promotes atherosclerosis initiation at these sites by inducing excessive EC proliferation and inflammation via the induction of glycolysis enzymes. (Arterioscler Thromb Vasc Biol. 2017;37:00-00. DOI: 10.1161/ATVBAHA.117.309249)

Key Words: apolipoproteins E ■ atherosclerosis ■ endothelial cells ■ glycolysis ■ hypoxia-inducible factor 1

Although it is associated with risk factors that act systemically (eg, hypercholesterolemia, smoking, age), atherosclerosis is a focal disease that develops preferentially near branches and bends of arteries.1 The physiology of endothelial cells (EC) varies considerably according to their location in the arterial tree. EC at regions that are predisposed to lesion formation (atheroprone) are characterized by relatively high rates of proliferation8,9 and proinflammatory activation.8–10 These features, which we refer to as endothelial dysfunction, promote early atherogenesis by increasing the accessibility of the vessel wall to leukocytes and its permeability to cholesterol-rich lipoproteins10 that are key drivers of atherogenesis. By contrast, EC at atheroprotected sites remain quiescent and are resistant to inflammatory activation. The spatial localization of EC phenotypes and atherosclerosis is tightly linked to local hemodynamics. Blood flow generates a frictional force at the endothelial surface called wall shear stress that varies in magnitude and direction according to vascular anatomy. Atheroprotected regions of arteries with relatively uniform geometry are exposed to high time-averaged shear stress that is unidirectional, whereas atheroprone sites near branches and bends are exposed to complex flow patterns generating low time-averaged shear stress that varies in direction (eg, oscillatory bidirectional and...
The application of flow to cultured cells has revealed a causal relationship between flow and EC physiology. Atheroprotective flow patterns induce numerous coding and noncoding RNAs that induce quiescence and reduce inflammation, whereas atheroprotive flow activates multiple signaling pathways and transcription factors that promote EC dysfunction and inflammation.12

The transcription factor HIF1α (hypoxia-inducible factor 1α) is a central regulator of cellular responses to hypoxia. In cells exposed to physiological levels of oxygen (normoxia), HIF1α is modified with hydroxyl groups by prolyl hydroxylase domain (PHD) enzymes, thus targeting it for rapid ubiquitination and degradation.13–16 However, this oxygen-dependent process is inactivated in conditions of hypoxia, leading to HIF1α accumulation. HIF1α plays an essential role during angiogenesis by inducing vascular endothelial growth factor and other growth factors, thereby enhancing perfusion of ischemic tissues to restore oxygenation. Of note, this process also requires HIF1α-dependent interaction of multiple glycolytic enzymes, thus causing a metabolic switch that allows ATP and macromolecules to be generated by glycolysis under anerobic conditions.19,21 HIF1α is expressed in advanced atherosclerotic plaques,22 and a recent study demonstrated that genetic deletion of HIF1α from EC reduced experimental atherosclerosis in a murine model.20 However, the potential role of HIF1α in focal EC dysfunction and inflammation linked to early atherogenesis has not been studied.

Here, we show for the first time that HIF1α can be activated in EC by mechanical low shear stress, leading to enrichment of HIF1α expression at atheroprotective regions of arteries. HIF1α is upregulated via a dual mechanism involving transcriptional activation by nuclear factor (NF)-κB and stabilization via the ubiquitination switching that allows ATP and macromolecules to be generated by glycolysis under anerobic conditions. Enriched expression of HIF1α at the low shear stress region was active at the shear stress site (aligned nuclei; Figure I in the online-only Data Supplement). It was concluded that HIF1α isolated from the low shear stress region had enriched expression of HIF1α and several downstream targets, including the glycolysis regulator 6-phosphofructo-2-kinase/fructose-2,6-bisphosphatase 3 (PFKFB3), glycolysis enzymes (hexokinase 2 [HK2], enolase 2), and glucose transporters (glucose transporter 1, glucose transporter 3; Figure 1A).

We also performed en face staining of the murine aortic endothelium to quantify the expression of HIF1α at sites that are known to be exposed to low (inner curvature of arch) or high (curvature) shear stress.11 It demonstrated that HIF1α protein was expressed at higher levels at a low shear compared with a high shear stress site (Figure 1B, top). Tiling of multiple fields of view revealed a sharp delineation in HIF1α expression, which was observed in EC exposed to low shear stress (note nonaligned nuclei) but not in EC exposed to high shear stress (aligned nuclei; Figure 1 in the online-only Data Supplement). It was concluded that HIF1α was active at the low shear region as a portion of the cellular pool localized to the nucleus (Figure 1B, top, arrows); in addition, the expression of HK2 and enolase 2 target molecules was also enriched at the low shear site (Figure 1B, center and bottom). The influence of atherogenesis on HIF1α expression was studied using ApoE−/− mice exposed to a high-fat diet for 6 weeks. En face staining revealed that HIF1α was expressed in EC overlying plaques and that the level of expression at the low shear region was similar in wild-type and ApoE−/− mice (Figure II in the online-only Data Supplement). Thus, we conclude that HIF1α
Figure 1. Enrichment of HIF1α (hypoxia-inducible factor 1α) and glycolytic enzymes at atheroprone sites. The expression of HIF1α and several of its target genes was quantified at low wall shear stress (WSS; inner curvature) and high WSS (outer curvature) regions of the porcine aorta by quantitative reverse transcription polymerase chain reaction (A) and en face staining of the mouse aorta (B). Mean levels±SEM are shown for n=5 pigs and n=3 mice. In representative images, endothelial cells were identified by costaining with anti-CD31 antibodies conjugated to fluorescein isothiocyanate (green). Cell nuclei were identified using TOPRO3 (blue). Arrows indicate nuclear HIF1α. Differences between means were analyzed using a paired t test. ENO2 indicates enolase 2; GLUT, glucose transporter; HK2, hexokinase 2; and PFKFB3, 6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3.
and downstream glycolysis genes are expressed preferentially at low shear atheroprone sites and that HIF1α expression is maintained during early atherogenesis.

**Low Shear Stress Induces HIF1α in Conditions of Atmospheric Oxygen**

Given its expression at atheroprone sites, we hypothesized that HIF1α is regulated by shear stress. In preliminary studies, we validated the detection of HIF1α by Western blotting by demonstrating that anti-HIF1α antibodies recognize a single band (at ≈120 kDa) in cells treated with the PHD inhibitor dimethylxalylglycine and that this band was suppressed by small interfering RNA sequences designed to target HIF1α (with no effect on HIF2α; Figure III in the online-only Data Supplement). The potential relationship between shear stress and HIF1α was investigated using cultured EC exposed to flow in the presence of atmospheric oxygen. Two complimentary systems were used: an orbital system that generates regions of lower shear stress (5 dyn/cm²) with variation in direction at the center and higher unidirectional shear stress (11 dyn/cm²) at the periphery and a parallel plate system that was used to generate unidirectional shear stress of 4 or 13 dyn/cm². Although these shear stress magnitudes are within the physiological range, they are referred to as low (4–5 dyn/cm²) and high (11–13 dyn/cm²) shear stress, respectively, for the sake of brevity. We have previously validated both the orbital and parallel plate systems; for example, high shear stress reduces apoptosis by inducing antiapoptotic genes. First, we demonstrated using systems; for example, high shear stress reduces apoptosis by inducing antiapoptotic genes.24 First, we demonstrated using

**NF-κB and Cezanne-Dependent Mechanism Induce and Stabilize HIF1α Under Low Shear Stress**

The mechanism linking low shear stress to HIF1α upregulation was investigated. qRT-PCR revealed that HIF1α mRNA levels were significantly elevated in human umbilical vein EC exposed to low or low oscillatory shear stress using the orbital (Figure 3A) or parallel plate (Figure 3B) systems compared with cells exposed to high shear stress or static conditions, indicating that low shear stress induces HIF1α at the transcript level. Similarly, HIF1α mRNA levels were elevated in human coronary artery EC exposed to low shear stress compared with cells exposed to high shear stress or static conditions (Figure V in the online-only Data Supplement). We hypothesized that flow regulates HIF1α mRNA via NF-κB, a transcription factor that can induce HIF1α in response to inflammatory signaling.10,31 In support of this, DNA-binding enzyme-linked immunosorbent assay revealed that low shear stress activated RelA and p50 subunits in cultured EC (Figure 3C). A direct link between NF-κB and HIF1α expression was established using an expression plasmid containing IκBα, which inhibited NF-κB activity in transfected EC (Figure VI in the online-only Data Supplement). Overexpression of IκBα reduced HIF1α expression in EC exposed to low shear stress, whereas an empty control plasmid had no effect (Figure 3D), indicating that NF-κB positively regulated HIF1α under low shear conditions. Similarly, silencing of RelA NF-κB subunits reduced HIF1α expression in EC exposed to low shear (Figure 3E).

![Figure 2](image-url)

**Figure 2.** Low shear stress induced HIF1α (hypoxia-inducible factor 1α). A, Human umbilical vein endothelial cells (HUVEC) were exposed to orbital flow to generate low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS) or were maintained under static conditions. After 72 h, they were exposed to dimethylxalylglycine (DMOG) for 4 h or remained untreated. B, Alternatively, HUVEC were exposed to high (13 dyn/cm²) or low oscillatory shear stress (4 dyn/cm²; 0.5 Hz) WSS for 72 h using a parallel plate system. A and B, The expression levels of HIF1α were assessed by Western blotting using specific antibodies, and anti-Calnexin or anti-PDHX (pyruvate dehydrogenase complex component X) antibodies were used to control for total protein levels. Representative blots are shown. Bands were quantified by densitometry. Data were pooled from 3 independent experiments, and mean HIF1α expression±SEM is shown. Differences between means were analyzed using a paired t test or 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons.
Finally, chromatin immunoprecipitation studies demonstrated that HIF1α promoter sequences coprecipitated with anti-histone H3 antibodies (positive control) and anti-RelA antibodies but did not coprecipitate with control IgG in EC exposed to orbital flow (Figure 3F), indicating that RelA interacts with the HIF1α promoter under low shear stress conditions. Collectively, these observations indicate that low shear stress induces HIF1α mRNA via an NF-κB–dependent process.
We reasoned that low shear stress must also prevent HIF1α degradation to enhance its expression at the protein level. To identify the mechanism, we examined the effects of flow on the expression of PHD proteins and Von Hippel Lindau (VHL) E3 ubiquitin ligase which destabilize HIF1α by modifying it with hydroxyl groups and ubiquitin, respectively. Western blotting revealed that PHD1, PHD3, and VHL were not suppressed by low shear stress (data not shown), indicating that HIF1α accumulation under these conditions is not mediated by reduced hydroxylation or ubiquitination. However, HIF1α can be stabilized by the deubiquitinating enzyme Cezanne that rescues it from degradation. The expression of Cezanne in cultured EC exposed to flow was studied by Western blotting. Antibodies against the N-terminal portion generated bands at \( \approx 95 \) and \( \approx 105 \) kDa (Figure 4), which were confirmed to result from Cezanne expression by gene silencing (Figure VII in the online-only Data Supplement). It was concluded that Cezanne is expressed at significantly higher levels in EC exposed to low shear stress compared with cells exposed to high shear stress or static conditions (Figure 4A and 4B). Similarly, qRT-PCR demonstrated that Cezanne mRNA levels were enhanced under low compared with high shear stress (Figure 4C and 4D). These observations were validated by qRT-PCR analysis of EC isolated from regions of the porcine aortic arch (Figure 5A) and en face staining of the murine aortic endothelium (Figure 5B; Figure I in the online-only Data Supplement), which demonstrated that Cezanne protein was expressed at higher levels at low shear stress compared with high shear stress sites.

To assess whether Cezanne is responsible for stabilizing HIF1α under low shear stress, we transfected cells with a dominant negative form that is mutated at the catalytic cysteine (GFP-Cezanne Cys/Ser). Immunofluorescent staining revealed

![Figure 4. Low shear stress induced Cezanne. A and C, Human umbilical vein endothelial cells (HUVEC) were exposed to orbital flow to generate low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS) or were maintained under static conditions. B and D, Alternatively, HUVEC were exposed to high (13 dyn/cm²) or low (4 dyn/cm²) WSS for 72 h using a parallel plate system. A and B, The expression levels of Cezanne were assessed by Western blotting using specific antibodies, and anti-PDHX (pyruvate dehydrogenase complex component X) antibodies were used to control for total protein levels. Representative blots are shown. Bands were quantified by densitometry. Data were pooled from 3 independent experiments, and mean Cezanne expression±SEM is shown. C and D, The expression levels of Cezanne mRNA were assessed by quantitative reverse transcription polymerase chain reaction. Data were pooled from 3 independent experiments, and mean expression±SEM is shown. Differences between means were analyzed using a 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons (A) or a paired t test (B–D).](http://atvb.ahajournals.org/Downloaded from)
that the expression of HIF1α was reduced by expression of GFP-Cezanne Cys/Ser and increased by GFP-Cezanne (wild type) compared with control GFP-transfected cells (Figure 6A), indicating that Cezanne activity is required to enhance HIF1α protein expression under low shear stress conditions. The effects of Cezanne on HIF1α ubiquitination were assessed by Western blotting of HIF1α immunoprecipitates generated from EC exposed to orbital flow. The abundance of high molecular weight forms of HIF1α and coprecipitating polyubiquitin was enhanced by expression of GFP-Cezanne Cys/Ser compared with expression of GFP-Cezanne or GFP alone (Figure 6B). These data are consistent with the hypothesis that Cezanne can target HIF1α for deubiquitination in EC exposed to low shear stress. Consistent with these observations, silencing of Cezanne significantly reduced HIF1α expression at the protein level (Figure 6C) but not at the mRNA level (Figure VIII in the online-only Data Supplement) in EC exposed to low shear stress. In conclusion, HIF1α is upregulated by low shear stress via dual processes that induce it at a transcriptional level via NF-κB and enhance it at the protein level via Cezanne.

HIF1α Promotes Glycolysis Under Low Shear Conditions

HIF1α drives angiogenesis by inducing a glycolytic switch,17–21 and we wondered whether this process can also be induced by low shear stress. qRT-PCR and Western blotting revealed that several glycolysis regulators, including HK2, enolase 2, and PFKFB3, were induced in cultured EC by the application of low shear stress (Figure 7A and 7C). Their expression was reduced by silencing of HIF1α (Figure 7B and 7C) or by silencing of Cezanne (Figure VIII in the online-only Data Supplement) and enhanced in cells exposed to dimethyloxyallylglycine (Figure IX in the online-only Data Supplement), indicating that Cezanne-HIF1α signaling activates glycolysis genes in EC exposed to low shear stress.

To study the influence of shear stress on glycolysis directly, we applied flow to cultured EC for 72 hours and then monitored extracellular acidification rate using a Seahorse XF analyzer.29 In the presence of glucose, glycolytic extracellular acidification rate was elevated in EC exposed to low compared with high shear stress (Figure 7D). In addition, glycolytic capacity, which is the ability of the glycolytic pathway to upregulate in time of energy need, was assessed after the addition of oligomycin that is a specific inhibitor of the mitochondrial ATP synthase. Glycolytic capacity was also higher in cells exposed to low shear stress compared with those cultured under high shear (Figure 7D). As a control, the glucose analogue 2-deoxyglucose (an inhibitor of glycolysis) was used to inhibit glycolysis and establish the nonglycolytic extracellular acidification rate. Collectively, these data indicate that low shear stress drives glycolysis via HIF1α-dependent induction of glycolytic enzymes.

HIF1α-Dependent Glycolysis Enhanced Proliferation and Inflammation in EC Exposed to Low Shear Stress

We next investigated whether HIF1α-dependent glycolysis influences EC proliferation and inflammation because these processes are involved in the initiation of atherosclerosis at low shear
Figure 6. Cezanne stabilizes HIF1α (hypoxia-inducible factor 1α) expression under low shear stress conditions via deubiquitination. A and B, Human umbilical vein endothelial cells (HUVEC) were transfected using pGFP-Cezanne, pGFP-Cezanne Cys/Ser, or with pGFP alone as a control. Cells were subsequently exposed to low wall shear stress (WSS; 5 dyn/cm²) using the orbital system for 72 h. A, The expression levels of HIF1α were assessed by immunofluorescent staining. Representative images and quantification of HIF1α expression (mean±SEM) are shown. B, Cells were treated with MG132 (proteasome inhibitor; 50 μmol/L) and bafilomycin (lysosome inhibitor; 100 nmol/L) for the final 4 h of orbiting. HIF1α immunoprecipitates were tested by Western blotting using antiubiquitin (top) or anti-HIF1α (bottom) antibodies. Representative blots are shown. Bands were quantified by densitometry. Data were pooled from 3 independent experiments, and mean levels±SEM are shown. B and C, Differences between means were analyzed using 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons (A and B) or a paired t test (C). DAPI indicates 4′,6-diamidino-2-phenylindole.
Figure 7. Low shear stress enhanced glycolysis via HIF1α (hypoxia-inducible factor 1α)-dependent induction of glycolytic enzymes. A. Human umbilical vein endothelial cells (HUVEC) were exposed to orbital flow to generate low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS). After 72 h, the expression levels of hexokinase 2 (HK2), enolase 2 (ENO2), glucose transporter (GLUT) 1, GLUT3, and PFKFB3 mRNA were assessed by quantitative reverse transcription polymerase chain reaction (qRT-PCR).

B. and C. HUVEC were transfected with small interfering RNA (siRNA) targeting HIF1α or with scrambled sequences. Cells were subsequently exposed to orbital flow to generate low WSS for 72 h. B, The expression levels of HIF1α, HK2, ENO2, and 6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 (PFKFB3) (Continued)

D. Glycolytic capacity was measured over time in response to low and high WSS conditions. The addition of oligomycin and 2DG inhibited glycolytic activity, illustrating the role of HIF1α in regulating glycolysis.
regions. This was studied using cultured cells exposed to flow in vitro and 2 different measures of proliferation (PCNA [proliferating cell nuclear antigen] and Ki67 staining). Gene silencing of HIF1α significantly reduced proliferation of EC exposed to low shear but did not alter EC exposed to high shear conditions (Figure 8A; Figure X in the online-only Data Supplement), indicating that low shear stress induces EC proliferation via HIF1α. The contribution of enhanced glycolysis to proliferation was assessed using 2 deoxyglucose or 3-(3-Pyridinyl)-1-(4-pyridinyl)-2-propen-1-one (an inhibitor of PFKFB3). Pretreatment of EC with either compound significantly reduced EC proliferation under low shear stress conditions (Figure 8B; Figure XI in the online-only Data Supplement). Moreover, the expression of inflammatory molecules in EC exposed to low shear stress was significantly reduced by silencing of Cezanne (Figure VIII in the online-only Data Supplement) or pretreatment of cells with 2DG (Figure 8C). Pretreatment of EC with 2DG did not alter the expression of Cezanne or HIF1α (Figure XII in the online-only Data Supplement), suggesting that glycolysis does not feedback to control Cezanne-HIF1α signaling.

Figure 8. Low shear stress enhanced endothelial proliferation and inflammation via glycolysis. A, Human umbilical vein endothelial cells (HUVEC) were transfected with small interfering RNA (siRNA) targeting HIF1α (hypoxia-inducible factor 1α) or with scrambled sequences. After 24 h, cells were exposed to orbital flow to generate low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS). B, HUVEC were exposed to orbital flow to generate low WSS in the presence of 2 deoxyglucose (2DG; 5 mmol/L) or 3-(3-Pyridinyl)-1-(4-pyridinyl)-2-propen-1-one (3PO; 10 μmol/L) or dimethyl sulfoxide (DMSO) vehicle as a control. A and B, After 72 h, EC proliferation was quantified by immunofluorescent staining using anti-PCNA (proliferating cell nuclear antigen) antibodies and costaining using 4’6-diamidino-2-phenylindole (DAPI). Representative images are shown. The % PCNA-positive cells were calculated for multiple fields of view. C, HUVEC were exposed to orbital flow to generate low WSS in the presence of 2DG (5 mmol/L) or DMSO vehicle for 72 h. They were exposed to tumor necrosis factor α (10 ng/mL) for the final 4 h. The expression levels of VCAM-1 and PDHX (pyruvate dehydrogenase complex component X) were assessed by Western blotting using specific antibodies. Representative images are shown. The % PCNA-positive cells were calculated for multiple fields of view. A–C, Data were generated from 3 independent experiments, and differences between means were analyzed using a paired t test.
Collectively, these data indicate that low shear stress enhances EC proliferation and inflammatory activation via HIF1α-dependent induction of glycolysis enzymes.

**HIF1α Promotes EC Dysfunction at Low Shear Stress Regions of Arteries In Vivo**

We validated our in vitro observations by imposing low shear stress in murine carotid arteries using the partial ligation model that promotes arterial wall remodeling and atherosclerosis in hypercholesterolemic mice. It should be noted that low shear stress is not sufficient to drive vascular inflammation per se, but it primes ECs for enhanced responses to inflammatory stimuli. Because atherosclerosis is a lipid-driven disease, we analyzed the function of HIF1α in partially ligated carotid arteries using hypercholesterolemic mice (ApoE−/− mice exposed to a high-fat diet for 6 weeks). We previously demonstrated by morphometry analysis that promotes arterial wall remodeling and atherosclerosis in hypercholesterolemic mice.

*Figure 9.* HIF1α (hypoxia-inducible factor 1α) induces glycolytic enzymes and inflammatory transcripts in arterial endothelium exposed to low shear in vivo. HIF1α−/− or HIF1α+/+ mice were subjected to partial ligation of the left carotid artery (LCA), whereas the right carotid artery (RCA) was sham-operated as a baseline control (RCA; unligated). After surgery, mice were exposed to a Western diet for 6 wk. The expressions of (A) regulators of glycolysis (hexokinase 2 [HK2], enolase 2 [ENO2], glucose transporter [GLUT] 1, 6-phosphofructo-2-kinase/fructose-2,6-biphosphatase 3 [PFKFB3]) or (B) inflammatory molecules (E-selectin, ICAM-1 [intercellular adhesion molecule 1], VCAM-1 [vascular cell adhesion molecule 1], MCP-1 [monocyte chemotactic protein 1]) were assessed by quantitative reverse transcription polymerase chain reaction. Data were pooled from n=4 mice per group. Mean gene expression±SEM in the LCA is presented. Differences between mean LCA values were analyzed using an unpaired t test. Because differences between LCA and RCA values were not tested statistically, we have presented RCA values as a baseline comparator (normalized to 1; dotted line).

|          | HK2       | E-selectin | ENO2      | ICAM-1    | GLUT1    | PFKFB3    | MCP1      |
|----------|-----------|------------|-----------|-----------|----------|-----------|-----------|
| **A**    |           |            |           |           |          |           |           |
| Relative mRNA levels | 10        | 15         | 8         | 15        | 10       | 15        | 5         |
| RCA      |           |            |           |           |          |           |           |
| **B**    |           |            |           |           |          |           |           |
| Relative mRNA levels | 5         | 25         | 6         | 25        | 5        | 20        | 2         |
| RCA      |           |            |           |           |          |           |           |

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of cross-sections and microcomputed tomography that lesion area in partially ligated carotid arteries was reduced by >50% and that lumen volume was increased by ≈30% in HIF1αCre;K0 ApoE−/− mice compared with HIF1αCre;K0 ApoE−/− mice. Correspondingly, it was demonstrated by morphometry of partially ligated carotid arteries that HIF1αCre;K0 ApoE−/− mice displayed a trend toward reduced total wall area compared with HIF1αCre;K0 ApoE−/− mice (HIF1αCre;K0, 39,660±9961 μm²; HIF1αCre;K0, 50,495±8423 μm²; mean±SEM; n=5; P<0.09), whereas medial thickness was unaltered by HIF1α deletion from EC (HIF1αCre;K0, 26854±3794 μm²; HIF1αCre;K0, 24730±1097 μm²; P=0.19).

It was demonstrated by qRT-PCR (Figure 9A) and immunohistochemistry (Figure XIII in the online-only Data Supplement) that partially ligated left carotid arteries (LCA) of ApoE−/− mice displayed enhanced expression of several regulators of glycolysis compared with sham right carotid arteries. The expression of inflammatory molecules was also enhanced in partially ligated LCA compared with sham-operated right carotid arteries (Figure 9B; Figure XIII in the online-only Data Supplement). The function of HIF1α in low shear-driven arterial inflammation and dysfunction was studied by inducible deletion of HIF1α from EC. First, it was demonstrated that the expression of HIF1α was significantly reduced in HIF1αCre;K0 ApoE−/− mice compared with HIF1αCre;K0 ApoE−/− mice at both the mRNA and protein levels (Figure XIV in the online-only Data Supplement). It was subsequently demonstrated by qRT-PCR that the expression of glycolysis regulators (Figure 9A; HK2, enolase 2, glucose transporter 1, PFKFB3) and inflammatory molecules (Figure 9B; E-selectin, ICAM-1 [intercellular adhesion molecule 1], VCAM-1 [vascular cell adhesion molecule 1], MCP-1 [monocyte chemotactic protein 1]) in partially ligated LCA was significantly reduced in ApoE−/− mice that lacked endothelial HIF1α (compare HIF1αCre;K0 with HIF1αCre;K0). This concurred with immunostaining that demonstrated that the expression of proteins that regulate glycolysis (HK2, PFKFB3; Figure 10A) or inflammation (ICAM-1, VCAM-1; Figure 10B) was significantly reduced in partially ligated LCA by endothelial deletion of HIF1α. In parallel studies, we assessed the rate of EC proliferation in partially ligated LCA and demonstrated that this was significantly reduced by genetic deletion of HIF1α (Figure 10C). These data indicate that exposure of arteries to low shear stress leads to the activation of HIF1α driving glycolysis, inflammation, and EC proliferation.

In summary, mechanical shear stress induces NF-κB and Cezanne-dependent upregulation of HIF1α at regions of arteries that are prone to atherosclerosis. HIF1α contributes to lesion initiation at atheroprone sites by activating glycolysis that enhances inflammation and EC proliferation. These data identify the Cezanne-HIF1α-glycolysis pathway as a novel mechanism contributing to lesion initiation at low shear stress regions of arteries.

**Discussion**

Although HIF1α is canonically activated in response to hypoxia, here we show for the first time that mechanical low shear stress can activate HIF1α under normoxic conditions. The underlying molecular mechanism for low shear-induced HIF1α involves the transcription factor NF-κB that is known to be activated at low shear atheroprone regions. This finding is consistent with previous observations that NF-κB can induce transcription of the HIF1α gene in other contexts. We reasoned that a second mechanism should be involved because transcriptional activation of the HIF1α gene per se is unlikely to enhance protein levels in the face of PHD- and VHL-mediated degradations. We observed abundant expression of PHD and VHL in EC exposed to low shear stress, suggesting that HIF1α is hydroxylated and ubiquitinated in these conditions. We, therefore, analyzed the expression of Cezanne, a deubiquitinating enzyme that can rescue HIF1α from VHL-mediated degradation by cleaving ubiquitin from it. Cezanne was enhanced under low shear stress conditions in both in vitro and in vivo models, and manipulation of its activity or expression revealed that Cezanne enhances HIF1α expression at the protein level. Our collective data suggest a model where low shear stress enhances HIF1α via dual processes: NF-κB-dependent induction of HIF1α mRNA and subsequent Cezanne-dependent stabilization of HIF1α protein via ubiquitin editing. Our study also indicates that a functional interplay exists between EC mechanoresponses and hypoxia because exposure to low shear stress enhanced HIF1α activation in response to hypoxic signaling. Thus, EC at atheroprone sites are primed for responses to localized hypoxia, which may be caused by secondary flows that convect oxygen away from the arterial wall or from consumption of oxygen by metabolically active macrophages.

The Cezanne protein was initially described by this laboratory as a negative regulator of NF-κB activation downstream from tumor necrosis factor or interleukin-1 receptor signaling. Thus, it was surprising to observe that Cezanne and NF-κB can co-operate to enhance HIF1α expression in low shear conditions. However, our subsequent studies revealed that silencing of Cezanne had modest effects on canonical NF-κB activation (RelA Ser536 phosphorylation) and did not alter non-canonical NF-κB (measured by p100 processing to p52) in EC exposed to low shear stress (data not shown). Thus, we suggest that cross-talk between Cezanne and NF-κB may differ according to physiological context. Cezanne can regulate NF-κB downstream from tumor necrosis factor α or interleukin-1 receptors but not in response to low shear stress. Consistent with this, tumor necrosis factor α and interleukin-1 receptors signal to NF-κB via the generation of polyubiquitin chains that can be dismantled by Cezanne, whereas shear stress controls NF-κB via alternative pathways that may be insensitive to Cezanne. It is also interesting to compare our observations with those of Passerini et al who found using microarrays that Cezanne mRNA was enriched in the descending thoracic aorta (a high shear stress site) compared with the inner curvature of the aortic arch (low shear stress site). Further work is required to assess whether Cezanne regulation by shear stress differs between the aortic arch and thoracic descending aorta.

Although it is well established that HIF1α is expressed in mature atherosclerotic plaques, its role in atherogenesis remains a topic of intense study. Recent work revealed that oral administration of a PHD2 inhibitor reduced lesion formation in mice by lowering serum cholesterol levels, indicating that systemic activation of HIF1α plays an atheroprotective role via modulation of lipid metabolism. However, conditional deletion or enforced expression of HIF1α in specific cell types revealed that the effects of HIF1α on atherogenesis are complex and vary according to cellular context. For example, expression of HIF1α in vascular smooth muscle cells or macrophages promoted inflammation in atherosclerosis, whereas HIF1α activity in CD11c-positive...
antigen-presenting cells protected arteries from lesion formation by reducing T-cell infiltrates. Of particular note, a recent study revealed that HIF1α expression in EC promoted lesion formation by enhancing expression of proinflammatory microRNA-19a. Previous studies demonstrated that low shear stress promotes atherosclerosis initiation by inducing high rates of EC proliferation.
that increase the permeability of arteries to cholesterol-containing lipoproteins\textsuperscript{10} and by activating inflammatory pathways.\textsuperscript{2,3} Our findings illuminate the underlying molecular mechanism by demonstrating that HIF1\(\alpha\) drives inflammation and proliferation in low shear stress conditions. Interestingly, genetic deletion of HIF1\(\alpha\) did not completely restore the expression of glycolytic enzymes or inflammatory molecules to the baseline value. This implies that although HIF1\(\alpha\) is required for the full induction of glycolytic enzymes and inflammatory molecules in response to low shear stress, other molecules also contribute to this response independently from HIF1\(\alpha\). Thus, therapeutic targeting of HIF1\(\alpha\) would be expected to dampen inflammation of arteries but not prevent it completely. Several signaling molecules including P53\textsuperscript{34} and JNK\textsuperscript{1}\textsuperscript{-}(c-Jun N-terminal kinase) have been implicated in enhanced EC turnover at low shear stress sites, and future studies should determine whether these pathways cross-talk with HIF1\(\alpha\) in atheroprotected endothelium.

Our studies revealed that HIF1\(\alpha\) enhances EC proliferation at low shear stress sites by inducing glycolytic enzymes to upregulate glycolysis. However, a recent study demonstrated that high shear stress reduces the rate of glycolysis in atheroprotected EC by activating the transcription factor KLF2 (Kruppel like factor 2) for transcriptional repression of PFKFB3.\textsuperscript{44} Thus, EC metabolism is regulated by high and low shear stress that trigger opposing signaling pathways; high shear reduces glycolysis via KLF2, whereas low shear enhances it via HIF1\(\alpha\). It is well established that HIF1\(\alpha\)-dependent glycolysis plays an essential role during angiogenesis because it allows rapid ATP generation under hypoxic conditions and produces intermediates for macromolecule synthesis, thus promoting EC proliferation and migration.\textsuperscript{19,20} This pathway is important in tumor vascularization, and clinical trials are underway to test the ability of glycolysis inhibitors to treat cancer.\textsuperscript{20} Now, we show for the first time that this pathway is activated in adult arteries, specifically at atheroprotected sites where it contributes to lesion initiation by promoting excessive EC proliferation and inflammation. It is likely that glycolysis also contributes to the progression of atherosclerosis, and it has recently been linked to inflammation in coronary artery disease.\textsuperscript{27}

In summary, we demonstrate for the first time that HIF1\(\alpha\) can be activated mechanistically by low shear stress. This non-canonical pathway, which requires activation of NF-\(\kappa\)B and Cezanne, promotes HIF1\(\alpha\) accumulation at atheroprotected sites leading to excessive EC proliferation and focal inflammation. Thus, the mechanically activated Cezanne-HIF1\(\alpha\) axis contributes to the initiation of lesions at branches and bends and may provide a novel therapeutic target to promote vascular function.

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Disclosures

None.

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Highlights

- HIF1α (hypoxia-inducible factor 1α) can be activated by mechanical shearing of arterial endothelial cells in the presence of oxygen, a noncanonical mechanism that promotes HIF1α accumulation at atheroprotective sites.
- The underlying mechanism involves dual processes; nuclear factor-kB–dependent induction of HIF1α mRNA and stabilization of HIF1α protein by the ubiquitin-editing enzyme Ceazanne.
- HIF1α promotes atherogenic processes at low shear stress regions by inducing excessive endothelial proliferation and inflammation via upregulation of glycolysis.
- Thus, noncanonical mechanical activation of HIF1α plays an important role in focal endothelial dysfunction and has the potential to be targeted therapeutically to enhance vascular function.
Mechanical Activation of Hypoxia-Inducible Factor 1α Drives Endothelial Dysfunction at Atheroprone Sites

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MATERIALS AND METHODS

Isolation of EC from porcine aortae. Pig aortas from 4-6 month old animals (weight approximately 80kg) were obtained immediately after slaughter from a local abattoir. They were cut longitudinally along the outer curvature to expose the lumen. EC exposed to high (outer curvature) or low (inner curvature) wall shear stress (WSS) were harvested using collagenase (1 mg/ml for 10 minutes at room temperature) prior to gentle scraping.

Mouse lines. Mice were housed under specific-pathogen free conditions. All animal experiments were reviewed and approved by the local authorities in accordance with German animal protection law. HIF1α was deleted from endothelial cells (EC) of ApoE^-/- mice by crossing Hif1α^floxflox ApoE^- mice VE-Cad-Cre-ER^T2 mice which express Tamoxifen-activated Cre under control of the VE-cadherin promoter as described1. Experiments were carried out using female mice because vascular physiology varies between sexes in mice with females having more pronounced and less variable pathophysiological features2. Thus female VE-Cad-Cre-ER^T2/ Hif1α^floxflox/ ApoE^-/- conditional knockout mice (called HIF1α^EC-cKO) and VE-Cad-Cre-ER^T2/ Hif1α^+/+ ApoE^-/- (HIF1α^+/+) mice were treated with tamoxifen (2 mg/20 g body weight) dissolved in neutral oil via intraperitoneal (i.p.) injections for 5 consecutive days. After one week, mice aged 6-8 weeks were anesthetized (ketamine (80 mg/kg, i.p.), xylazine (10 mg/kg, i.p.)) and the left carotid artery was partially ligated by closing the external, internal and the occipital artery restricting blood outflow to the superior thyroid artery only as described3. The right carotid artery was sham-operated as a control. Mice were fed a high fat diet (0.15% cholesterol) for 6 weeks.

Staining of murine endothelium. The expression levels of specific proteins were assessed in EC at regions of the inner curvature (susceptible site) and outer curvature (protected site) of murine aortae by en face staining. Animals were killed by I.P injection of pentobarbital or by isoflurane overdose. Aortae were perfused in situ with PBS (at a pressure of approximately 100 mm Hg) and then perfusion-fixed with 4% Paraformaldehyde prior to harvesting. Fixed aortae were tested by immunostaining using specific primary antibodies and Alexafluor568-conjugated secondary antibodies (red). EC were identified by co-staining using anti-CD31 antibodies conjugated to the fluorophore FITC (green). Nuclei were identified using a DNA-binding probe with far-red emission (To-Pro-3). Stained vessels were mounted prior to visualization of endothelial surfaces en face using confocal laser-scanning microscopy (Zeiss LSM510 NLO inverted microscope). Isotype-matched monoclonal antibodies raised against irrelevant antigens or pre-immune rabbit sera were used as experimental controls for specific staining. The expression of particular proteins at each site was assessed by quantification of fluorescence intensity for multiple cells (at least 50 per site) using Image J (1.49p) and calculation of mean fluorescence intensities with standard error of the mean.

Consecutive sections at an interval of 120 μm made from experimental carotid arteries were immunostained using specific primary antibodies. Non-specific IgG served as a negative control. Primary antibodies were detected using horse radish peroxidase-conjugated secondary antibodies and 3,3-diaminobenzidine (DAB; brown) or NovaRed (red) substrates (Vector Laboratories, Burlingame, CA, USA).
Sections were counterstained using haematoxylin and eosin. The extent of EC staining in cross-sections was scored by 3 researchers blinded to the experimental conditions. Lesions were identified in carotid arteries by elastic van Gieson staining of serial sections (4–5 µm thick) at an interval of 100 µm, and the total vessel area and medial area were calculated by morphometry using Image J software.

**EC culture and exposure to WSS.** HUVEC were isolated using collagenase digestion. Cells were cultured on 1% gelatin and maintained in M199 growth medium supplemented with foetal bovine serum (20%), L-glutamine (4 mmol/L), endothelial cell growth supplement (30 µg/ml), penicillin (100 U/ml), streptomycin (100 µg/ml) and heparin (10 IU/ml). Human coronary artery EC (HCAEC) were purchased from PromoCell and cultured according to the manufacturer’s recommendations. EC at passage 3–5 were cultured until confluent in 6 well plates and exposed to flow using an orbital shaking platform (PSU-10i; Grant Instruments) housed inside a cell culture incubator. The radius of orbit of the orbital shaker was 10 mm and the rotation rate was set to 210 rpm. This motion caused swirling of the culture medium over the cell surface generating low WSS (approximately 5 dynes/cm²) with varied directionality at the centre and high (approximately 11 dynes/cm²) WSS with uniform direction at the periphery. Alternatively, HUVEC were cultured on Ibidi® gelatin-coated µ-Slides (Ibidi GmbH) until they reached confluency. Flowing medium was then applied using the Ibidi® pump system to generate low (4 dyn/cm²), low oscillatory (+/- 4 dyn/cm², 0.5 Hz) or high (13 dyn/cm²) WSS. The slides and pump apparatus were enclosed in a cell culture incubator warmed to 37°C.

**Gene silencing and overexpression.** Cell cultures were transfected with siRNA sequences that are known to silence HIF1α (Dharmacon SMARTpool: ON-TARGETplus L-004018-00-0005, Dharmacon Individual: ON-TARGETplus J-004018-08-0002), RelA (Dharmacon SMARTpool: ON-TARGETplus L-003533-00-0005) or Cezanne (Dharmacon siGENOME D-008670-03) using the Lipofectamine® RNAiMAX transfection system (13778-150, Invitrogen) following the manufacturer’s instructions. Alternatively, cells were transfected with the Neon™ Transfection system (Invitrogen; 1200 volts, 40 ms, 1 pulse). Final siRNA concentration used was 25nM for lipofectamine4 and 50 mm by electroporation. After knockdown, cells were then incubated in complete M199 growth medium for 2 h before exposure to flow. Non-targeting scrambled sequences were used as a control (D-001810-01-50 ON-TARGETplus Non targeting siRNA#1, Dharmacon). HUVEC were transfected with expression vectors containing IkBα (pCMV-IkBα, GFP-Cezanne or GFP-Cezanne Cys/Ser (catalytically inactive))5 using Lipofectamine (ThermoFisher).

**Comparative real time PCR.** RNA was extracted using the RNeasy Mini Kit (74104, Qiagen) and reverse transcribed into cDNA using the iScript cDNA synthesis kit (1708891, Bio-Rad). The levels of human, porcine or murine transcripts were assessed using quantitative real time PCR (qRT-PCR) using gene-specific primers (Supplementary Table 1). Reactions were prepared using SsoAdvanced universal SYBR®Green supermix (172-5271, Bio-rad) and following the manufacturer’s instructions, and were performed in triplicate. Relative gene expression was calculated by comparing the number of thermal cycles that were
necessary to generate threshold amounts of product. Fold changes were calculated using the ΔΔCt method. Data were pooled from at least three independent experiments and mean values were calculated with SEM.

**Immunoprecipitation of HIF1α.** Cells were lysed using ice-cold lysis buffer (50 mM Tris (pH 7.4), 250 mM NaCl, 0.3% Triton X-100, 1mM EDTA, 10µM MG132) supplemented with protease inhibitor cocktail (Roche, Basel, Switzerland). Lysates were subjected to three freeze-thaw cycles before immunoprecipitation using anti-HIF1α antibodies. Beads were then washed extensively using lysis buffer.

**Western blotting** Total cell lysates were isolated using lysis buffer (containing 2% SDS, 10% Glycerol and 5% β-mercaptoethanol). Western blotting was carried out using specific antibodies against HIF1α, HIF2α, Cezanne, IkBa, HK2, ENO2, PFKFB3, VCAM-1, PDHX, Calnexin (Supplementary Table 2) and horse radish peroxidase-conjugated secondary antibodies obtained commercially from Dako and chemiluminescent detection was carried out using ECL Prime® (GE Healthcare).

**Chromatin immunoprecipitation.** A commercial kit was used (Cell Signaling Technology). HUVECs were fixed using formaldehyde and nuclei were purified and subjected to sonication. For immunoprecipitation reactions, the nuclear lysates were incubated overnight at 4°C with protein G magnetic beads coated with either anti-RelA (sc-372; Santa Cruz) or with anti-histone H3 antibodies or isotype-matched IgG control (both from Cell Signaling Technology). Immunocomplexes were washed, and co-precipitating DNA fragments were quantified by real-time PCR using specific DNA primers (Supplementary Table 1).

**Immunofluorescent staining of cultured EC.** The expression levels of proteins were assessed by immunostaining using specific antibodies followed by widefield fluorescence microscopy (LeicaDMI4000B). HUVEC were fixed with Paraformaldehyde (4%) and permeabilised with Triton X-100 (0.1%). Following blocking with goat serum for 30 min monolayers were incubated for 16 h with primary antibodies against HIF1α, PCNA (proliferation marker) or Ki67 (proliferation marker) and AlexaFluor488- or Alexafluor568-conjugated secondary antibodies. Nuclei were identified using the DNA-binding probe DAPI (Sigma). Image analysis was performed using Image J software (1.49p) to calculate average fluorescence. Isotype controls or omission of the primary antibody was used to control for non-specific staining.

**Glycolysis assay.** HUVEC were plated in a Cell Tak-coated Seahorse cell culture plate in XF assay media pH 7.4 (Seahorse Bioscience) supplemented with 2 mM glutamine and 25 mM glucose. Extracellular acidification rate (ECAR) was measured basally prior to addition of oligomycin to assess glycolytic capacity. After reading basic ECAR and maximum ECAR induced by oligomycin, glycolysis was suppressed via addition of 2 deoxyglucose (2DG) thus allowing measurement of non-glycolytic ECAR as described6. Glycolytic ECAR was calculated by taking the ECAR value prior to oligomycin injection and subtracting the ECAR value in the presence of 2DG. Glycolytic capacity was calculated by subtracting ECAR in the presence of oligomycin from ECAR in the presence of 2DG.
Statistics. Differences between samples were analysed using an unpaired or paired Student’s t-test or ANOVA with the Bonferonni correction for multiple pairwise comparisons.
Supplementary Table 1  
PCR primers.

|           | Forward 5'-3' | Reverse 5'-3' |
|-----------|---------------|---------------|
| **Mouse** |               |               |
| Mouse HK2 | TCGCATAATGATGCCTGCTTA | TCTGAGAGACGCATGTGGTAG |
| Mouse ENO2| GAGAAGATTGGCGGCGAGAG | CGGAAGACCTTGGCGAGCT |
| MouseGLUT1| GGATCACTGAGTTGGCTTA  | AGCGGTGTTCCATGGTTTA |
| Mouse PFKFB3 | GAAGCTGACTCGCTACCTCA | TGTACTGCTCTACAGGCTCA |
| Mouse MCP1 | CTTGAGCTACAGGTTGGCTTA | CAGCTCTTGGGACACCCGT |
| Mouse ICAM1 | CCGCTGTCAGCGGCTTGGCTTA | GAACATTGCTACAGGCTCAGG |
| Mouse VCAM1 | TGGGAGCCTCAACCGGTACCT | AGCAATCGTTTGTATCCAGGG |
| Mouse E-selectin | GAATGCCTCAGGCTTCCTTCTCTC | GAGCTCAGGGGCTATTGTA |
| **Pig**   |               |               |
| Pig HIF1α | TGTTACTCAGAGATGATGGAC | CATTCCCTCATGGTGACAG |
| Pig HK2   | GAACAGGAGAGTTGGCTTA  | CTTTCTGAGTTCCTTTGCT |
| Pig ENO2  | GCCCTGCTCTGCTGGAAC   | ATTGCCGACACACTGTTA |
| Pig GLUT1 | GGCATCAACGGTGTTTTCAT | AGACTCAGCGACCGGACG |
| Pig GLUT3 | TGCTCTGAGAGGCGATCAA  | GACACAGGCGACTGTAAG |
| Pig PFKFB3 | CTGACCCGCTACCTCAAAGC | AGGAGCTGACTGCTTTGAG |
| **HUMAN** |               |               |
| Human HIF1α | TTGTTCTCACAGATGATGGAC | CTCAATTTCTCATGGTCACAG |
| Human HK2  | TCACGGAGACTCCAATACG | GCTCAAAGCCCTTTCTCCAT |
| Human ENO2 | CCAGCTGAGTTGGGATGCTTA | TCATGATGCTGCGAGGCT |
| Human GLUT1 | TCCTCCGCTGCTGTTGCTA | CAGCGTCTCCTCCACAG |
| Human GLUT3 | CTCTCGCTCAACGCCCTGTG | CCCTAAAGCAAGCCCAAG |
| Human PFKFB3 | CCGTTGAGACTGACGCAGA | GACAGGCTGCTGCGGAGG |
| Human Cezanne | GAGTTGAGAAAGGACGCTT | CACTCTGCTGCTGCTTTA |
| Human HIF1A promoter | CTTCCGAGAAAGGCGAGAAGT | GTGCTTCGCTGCTGTTTAGCG |
## Supplementary Table 2  Antibodies: suppliers and concentrations used.

| Antibody | Company | Use | Dilution Ratio | Final Concentration |
|----------|---------|-----|----------------|---------------------|
| HIF1α    | Mouse monoclonal Anti-Human HIF-1α, 610959, BD Biosciences | WB | 1/500 | 5μg/ml |
| HIF1α    | Rabbit monoclonal antibody to HIF1α, ab51608, Abcam | IF | 1/100 | 10μg/ml |
| VCAM1    | Rabbit monoclonal anti-VCAM1 antibody, Ab134047, Abcam | WB | 1/5000 | 80ng/ml |
|          |                             | ICC | 1/200 | 2μg/ml |
| PCNA     | Rabbit polyclonal anti- PCNA antibody, Ab18197, Abcam | IF | 1/100 | 2μg/ml |
| PFKFB3   | Rabbit monoclonal anti- PFKFB3 antibody, Ab181861, Abcam | WB | 1/3000 | 0.2μg/ml |
|          |                             | ICC | 1/100 | 5μg/ml |
| ENO2     | Rabbit polyclonal NSE/ENO2 Antibody [NB100-91898], Novus | WB | 1/2000 | 0.5μg/ml |
| HK2      | Rabbit polyclonal Hexokinase II Antibody [NBP2-16814], Novus | WB | 1/5000 | 0.2μg/ml |
|          |                             | IF | 1/200 | 5μg/ml |
| HIF2α    | Abcam, Rabbit polyclonal anti-HIF-2-alpha antibody, Ab199 | WB | 1/1000 | 1μg/ml |
| ICAM1    | Rabbit polyclonal anti-ICAM1 antibody, 10020-I-AP, ProteinTech | ICC | 1/250 | 5μg/ml |
| PDHX     | Rabbit polyclonal E3BP(PDHX) antibody, sc-98751, Santa Cruz | WB | 1/5000 | 40ng/ml |
| Calnexin | Mouse monoclonal Anti-calnexin, 610524, BD Biosciences | WB | 1/5000 | 50ng/ml |
| Cezanne  | Rabbit polyclonal to OTUD7B, 16605-I-AP, ProteinTech | WB | 1/3000 | 50ng/ml |
|          |                             | IF | 1/100 | 2.5μg/ml |
| Ki67     | Rabbit polyclonal to Ki67, ab15580, Abcam | IF | 1/100 | 5μg/ml |

WB, Western blotting; IF, immunofluorescence; ICC, immunocytochemistry.
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Supplementary Figure I. Low magnification images of en face staining of HIF1α and Cezanne in the murine aorta.

The expression of HIF1α (A) and Cezanne (B) was assessed at low shear stress (inner curvature) and high shear stress (outer curvature) regions of the murine aorta by en face staining. EC were identified by co-staining with anti-CD31 antibodies conjugated to FITC (green). Cell nuclei were identified using TOPRO3 (blue). Regions exposed to high and low shear stress were identified by anatomical landmarks and confirmed by assessment of nuclei which are aligned specifically under high shear stress. Images representing tiling of multiple fields of view are presented. Note the distinct difference in HIF1α and Cezanne expression between high and low shear stress regions (delineated by a broken white line). Scale bar, 20 µm.
Supplementary Figure II. HIF1α is expressed by endothelial cells overlying plaques at low shear stress regions.

ApoE<sup>-/-</sup> mice aged 8 weeks were exposed to a high fat diet for 6 weeks (n=4). C57BL/6 (wild-type) mice aged 14 weeks were also studied (n=4). The expression of HIF1α was quantified at a low wall shear stress region (inner curvature) by en face staining. EC were identified by co-staining with anti-CD31 antibodies conjugated to FITC (green). Cell nuclei were identified using TOPRO3 (blue). A representative image of HIF1α expression in EC overlying a plaque in the ApoE<sup>-/-</sup> model is shown (left). Mean levels of HIF1α expression (+/- SEM) were calculated for ApoE<sup>-/-</sup> and wild-type mice (right). Differences between means were analysed using a paired t-test. n.s., not significant.
Supplementary Figure III. Validation of HIF1α siRNA.
HUVEC were transfected with siRNA targeting HIF1α or with scrambled sequences. After 24 h, cells were exposed to DMOG for 4 h. The expression levels of HIF1α and HIF2α were assessed by Western blotting using specific antibodies, and anti-Calnexin antibodies were used to control for total protein levels. Representative blots are shown.
Supplementary Figure IV. Endothelial cells exposed to low shear stress in atmospheric oxygen were not hypoxic. HUVEC were exposed to orbital flow to generate low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS) either within a hypoxic chamber (1% O₂) or under atmospheric levels O₂. After 72 h, the cells were treated with Pimonidazole (60 ng/ml) for 90 min before fixation with 4% PFA and staining using rabbit polyclonal anti-pimonidazole antibodies. Hypoxic cells were assessed by Pimonidazole staining (red). Cell nuclei were identified using DAPI (blue).
Supplementary Figure V. Low shear stress induced HIF1α mRNA in coronary artery EC.

Human coronary artery EC were exposed to low (5 dyn/cm²) or high (11 dyn/cm²) wall shear stress (WSS) for 72 h using an orbital system or were exposed to static conditions. HIF1α mRNA was quantified by qRT-PCR. Data were pooled from three different donors and mean HIF1α expression levels +/- SEM are shown. Differences between means were analysed using a 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons.
Supplementary Figure VI. Validation of enforced IκBα expression. HUVEC were transfected with pCMV-IκBα to inhibit NF-κB or with an empty plasmid as a control. After 24 h, the expression levels of IκBα and VCAM-1 were assessed by Western blotting using specific antibodies, and anti-Calnexin antibodies were used to control for total protein levels. Representative blots are shown.
Supplementary Figure VII. Validation of antibodies used to detect Cezanne by Western blotting.

HUVEC were transfected with siRNA targeting Cezanne or with scrambled sequences. Cells were then exposed to orbital flow to generate low WSS (5 dyn/cm²). After 72 h, the expression levels of Cezanne were assessed by Western blotting using specific antibodies, and anti-PDHX antibodies were used to control for total protein levels. Representative blots are shown.
Supplementary Figure VIII. Cezanne positively regulates the expression of glycolysis genes and inflammatory molecules in EC exposed to low shear stress.

HUVEC were transfected with siRNA targeting Cezanne or with scrambled sequences. Cells were subsequently exposed to orbital flow to generate low WSS (5 dyn/cm²) for 72 h. The expression levels of Cezanne, HIF1α (A) or HK2, ENO2, PFKFB3, VCAM-1, ICAM-1, E-selectin, IL-8 and MCP1 (B) mRNA were assessed by qRT-PCR. Data were generated from three independent experiments and differences between means were analysed using a paired t-test.
Supplementary Figure IX. Low shear stress primed EC for hypoxic signaling.

HUVEC were exposed to orbital flow to generate low (5 dyn/cm$^2$) or high (11 dyn/cm$^2$) wall shear stress (WSS). After 72h, cells were treated with DMOG for 4 h. The expression levels of HK2, ENO2 and PFKFB3 were assessed by Western blotting using specific antibodies, and anti-Calnexin antibodies were used to control for total protein levels. (A) Representative blots are shown. (B) Bands were quantified by densitometry. Data were pooled from three independent experiments and mean expression +/- SEM is shown. Differences between means were analysed using a 2-way ANOVA with the Bonferroni correction for multiple pairwise comparisons.
Supplementary Figure X. Low shear stress enhanced endothelial proliferation (Ki67 staining) via HIF1α.

HUVEC were transfected with siRNA targeting HIF1α or with scrambled sequences. After 24 h, cells were exposed to orbital flow to generate low (5 dyn/cm²) or high (5 dyn/cm²) shear stress. After 72 h, EC proliferation was quantified by immunofluorescent staining using anti-Ki67 antibodies and co-staining using DAPI. Representative images are shown. The % Ki67-positive cells were calculated for multiple fields of view. Data were pooled from three independent experiments and mean expression +/- SEM is shown. Differences between means were analysed using a 2-way ANOVA with the Bonferroni correction for multiple pairwise comparisons.
Supplementary Figure XI. Low shear stress enhanced endothelial proliferation (Ki67 staining) via glycolysis.

HUVEC were exposed to orbital flow to generate low shear stress (5 dyn/cm²) in the presence of 2DG (5 mM) or 3PO (10 µM) or DMSO vehicle as a control. After 72 h, EC proliferation was quantified by immunofluorescent staining using anti-Ki67 antibodies and co-staining using DAPI. Representative images are shown. The % Ki67-positive cells were calculated for multiple fields of view. Data were pooled from three independent experiments and mean expression +/- SEM is shown. Differences between means were analysed using a 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons.
Supplementary Figure XII. FDG treatment did not alter the expression of Cezanne or HIF1α in EC.
HUVEC were exposed to FDG for 24 h or 72 h or remained untreated as a control. The expression levels of HIF1α and Cezanne were assessed by Western blotting using specific antibodies, and anti-PDHX antibodies were used to control for total protein levels. Representative blots are shown. Bands were quantified by densitometry. Data were pooled from three independent experiments and mean levels +/- SEM are shown. Differences between means were analysed using 1-way ANOVA with the Bonferroni correction for multiple pairwise comparisons. n.s., not significant.
Supplementary Figure XIII. Glycolytic enzymes, inflammatory proteins and proliferation were induced by low shear in vivo. ApoE<sup>−/−</sup> mice were subjected to partial ligation of the left carotid artery (LCA). After surgery, mice were exposed to a Western diet for 6 weeks. Immunostaining was performed to assess the expression of the glycolysis regulator HK2, the inflammatory proteins ICAM-1 and VCAM-1, and the proliferation marker Ki67 in cross-sections using DAB (brown) or NovaRed (red) substrates. n=4 mice per group were studied. Representative images are shown. Histology scores of positive endothelial staining were pooled from 3 independent evaluations and mean values +/- SEM are shown. Differences between means were analysed using a Mann Whitney test.
Supplementary Figure XIV. Validation of inducible HIF1α genetic deletion from EC.

HIF1α\textsuperscript{EC-\textit{cKO}} or HIF1α\textsuperscript{+/+} mice were subjected to partial ligation of the left carotid artery. After surgery, mice were exposed to a Western diet for 6 weeks. (A) The expression of HIF1α was assessed by qRT-PCR. Data were pooled from n=4 mice per group. Differences between means were analysed using an unpaired t-test. (B) Immunostaining was performed to assess the expression of HIF1α protein in cross-sections using NovaRed (red) substrate. n=5 mice per group were studied. Representative images are shown with arrowheads indicating EC that stained positive and high magnification insets. Histology scores of positive endothelial staining were pooled and mean values +/- SEM are shown. Differences between means were analysed using a Mann Whitney test.