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# COX-2-Derived Prostanoids and Oxidative Stress Additionally Reduce Endothelium-Mediated Relaxation in Old Type 2 Diabetic Rats

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## Abstract

Endothelial dysfunction in resistance arteries alters end organ perfusion in type 2 diabetes. Superoxides and cyclooxygenase-2 (COX-2) derivatives have been shown separately to alter endothelium-mediated relaxation in aging and diabetes but their role in the alteration of vascular tone in old diabetic subjects is not clear, especially in resistance arteries. Consequently, we investigated the role of superoxide and COX-2-derivatives on endothelium-dependent relaxation in 3 and 12 month-old Zucker diabetic fatty (ZDF) and lean (LZ) rats. Mesenteric resistance arteries were isolated and vascular tone was investigated using wire-myography. Endothelium (acetylcholine)-dependent relaxation was lower in ZDF than in LZ rats (60 versus 84% maximal relaxation in young rats and 41 versus 69% in old rats). Blocking NO production with L-NAME was less efficient in old than in young rats. L-NAME had no effect in old ZDF rats although eNOS expression level in old ZDF rats was similar to that in old LZ rats. Superoxide level and NADPH-oxidase subunits (p67phox and gp91phox) expression level were greater in ZDF than in LZ rats and were further increased by aging in ZDF rats. In young ZDF rats reducing superoxide level with tempol restored acetylcholine-dependent relaxation to the level of LZ rats. In old ZDF rats tempol improved acetylcholine-dependent relaxation without increasing it to the level of LZ rats. COX-2 (immunolabelling and Western-blot) was present in arteries of ZDF rats and absent in LZ rats. In old ZDF rats arterial COX-2 level was higher than in young ZDF rats. COX-2 blockade with NS398 restored in part acetylcholine-dependent relaxation in arteries of old ZDF rats and the combination of tempol and NS398 fully restored relaxation in control (LZ rats) level. Accordingly, superoxide production and COX-2 derivatives together reduced endothelium-dependent relaxation in old ZDF rats whereas superoxides alone attenuated relaxation in young ZDF or old LZ rats.

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## Introduction

Type 2 diabetes is the most frequently encountered metabolic disorder, currently affecting 5% to 10% of most populations, and the incidence continues to grow among developing countries [1]. Associated with obesity, type 2 diabetes is characterized by an insulin resistance inducing metabolic changes such as hyperinsulinemia, hyperglycaemia, dyslipidaemia and hypertension, all leading to increased cardiovascular risk [2]. The morbidity and mortality associated with type 2 diabetes are essentially related to the vascular lesions that develop over time [3–5]. Microcirculation is mainly involved, and consequently vital organs such as the brain, heart, kidneys and the limbs are progressively damaged. While the consequences of type 2 diabetes on macrocirculation have been extensively studied [6,7], much less is known about its effects on microcirculation, especially in old subjects. Aging is associated with a rising incidence of hypertension, obesity and type 2 diabetes; unfortunately the 3 risk factors are usually associated [8–10]. Therefore the aim of the present study was to investigate

the effect of type 2 diabetes on the endothelium in old rats. Certainly, type 2 diabetes is usually detected late after developing, usually during a medical checkup or when adverse effects induce alerting disorders such as retinal or kidney disorders. Consequently, investigating endothelium-mediated relaxation in older diabetic rats is a key issue.

Type 2 diabetes is associated with an increased reactive oxygen species (ROS) production [11–13], which may reduce endothelium-dependent dilation [14,15]. Similarly, aging is associated with a change in endothelium-dependent dilation with a reduction in NO-dependent dilation in both large and small arteries [16]. The effect of aging is less obvious in small resistance arteries [17,18]. In resistance arteries ROS may also play a role in the reduction in NO-dependent dilation observed in aging [16]. Furthermore, alteration of the endothelium may also involve an inflammatory process [19]. Consequently, type 2 diabetes associated with aging is likely to induce greater damage to the endothelium. Conse-

quently, we investigated the effect of type 2 diabetes on endothelium-dependent relaxation in young and old rats.

Using Zucker Diabetic Fatty (ZDF) rats, we found that ROS and COX-2 vasoconstrictor derivatives additionally reduced endothelium-dependent relaxation in old diabetic rats whereas ROS alone are involved in the decrease in endothelium-dependent relaxation in young ZDF or in old LZ rats.

## Methods

### Animals

Adult male Zucker fatty diabetic (ZDF) and lean Zucker (LZ) rats, 12–14 and 50–52 week-old, were purchased from Charles River (L'Arbresles, France). Rats were anesthetized (Isoflurane, 2.5%) and the right femoral artery was catheterized for blood pressure measurement [20]. Animals were then sacrificed by CO<sub>2</sub> inhalation, the gut excised and mesenteric arteries gently dissected. From each rat, several arterial segments (second order mesenteric arteries similar in size) were isolated and used for pharmacology,

**Table 1.** Rat body weight, glycaemia and mean arterial blood pressure (MAP) were measured in 3- and 12-month old LZ and ZDF rats.

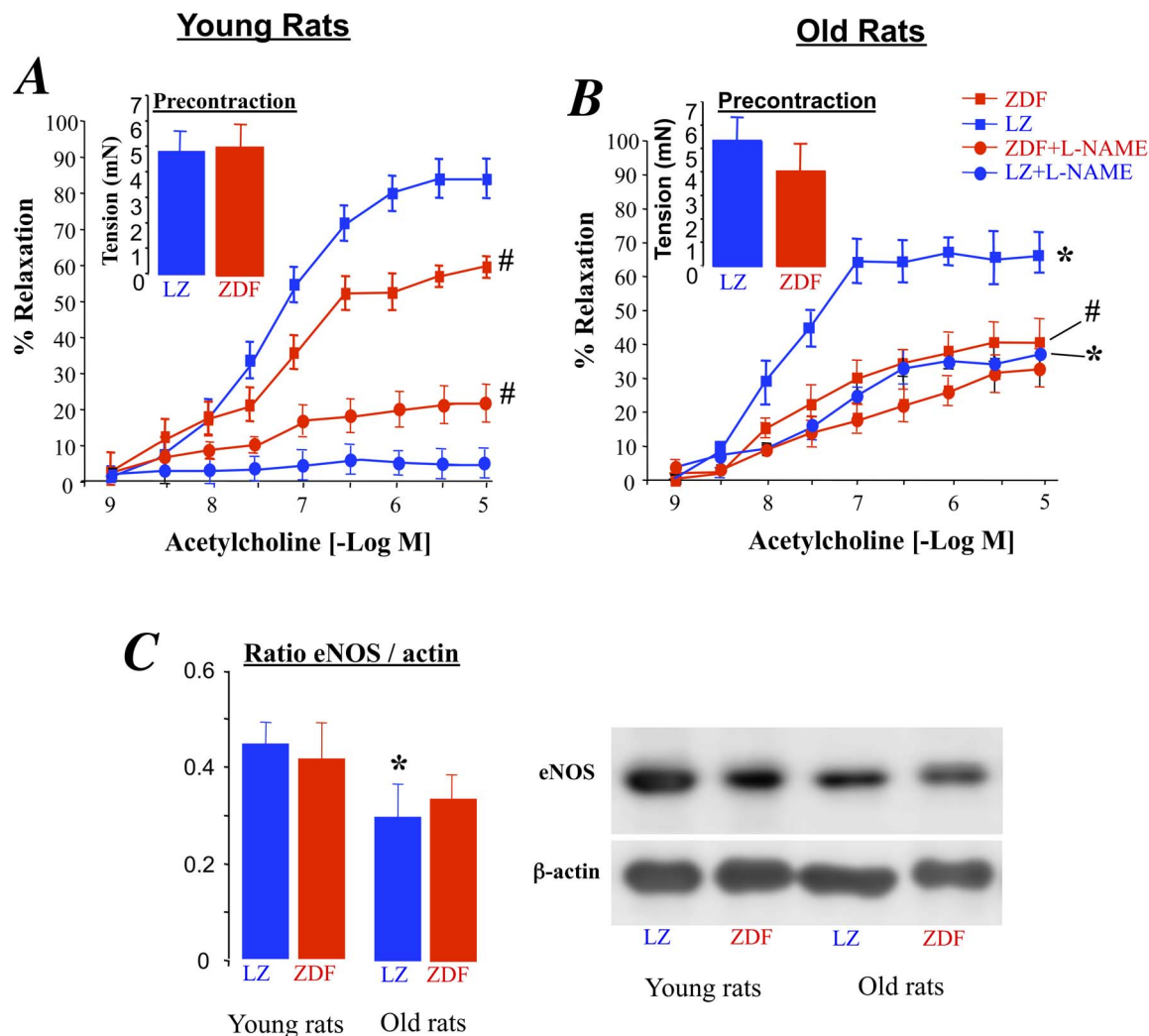
	3-month old rats		12-month old rats	
	LZ rats	ZDF rats	LZ rats	ZDF rats
Body weight (g)	325±11	441±17*	567±28#	382±10*#
Glycaemia (mg/dL)	112±13	302±23*	125±11	421±37*#
MAP (mmHg)	98±4	108±6	112±5	132±6*#

Mean±SEM is presented (n=12 rats per group).

\*P<0.05, ZDF versus LZ of the same age.

#P<0.05, effect of age within each group.

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**Figure 1. Acetylcholine-induced relaxation in young (A) and old (B) LZ and ZDF rats.** Relaxation was obtained before and after NO synthesis inhibition with the NO synthesis blocker L-NAME. The inserts show the precontraction applied before adding acetylcholine. Beta-actin and eNOS expression levels were measured using Western-Blot analysis (typical blots on the right side). Data is given as a ratio eNOS/beta-actin (C). Mean ± SEM is presented (n=10 per group). #P<0.01, ZDF versus LZ. \*P<0.01, old versus young rats. doi:10.1371/journal.pone.0068217.g001

biochemical and immuno-histological analyses. Before sacrifice, a sample of blood was collected for analyses [21].

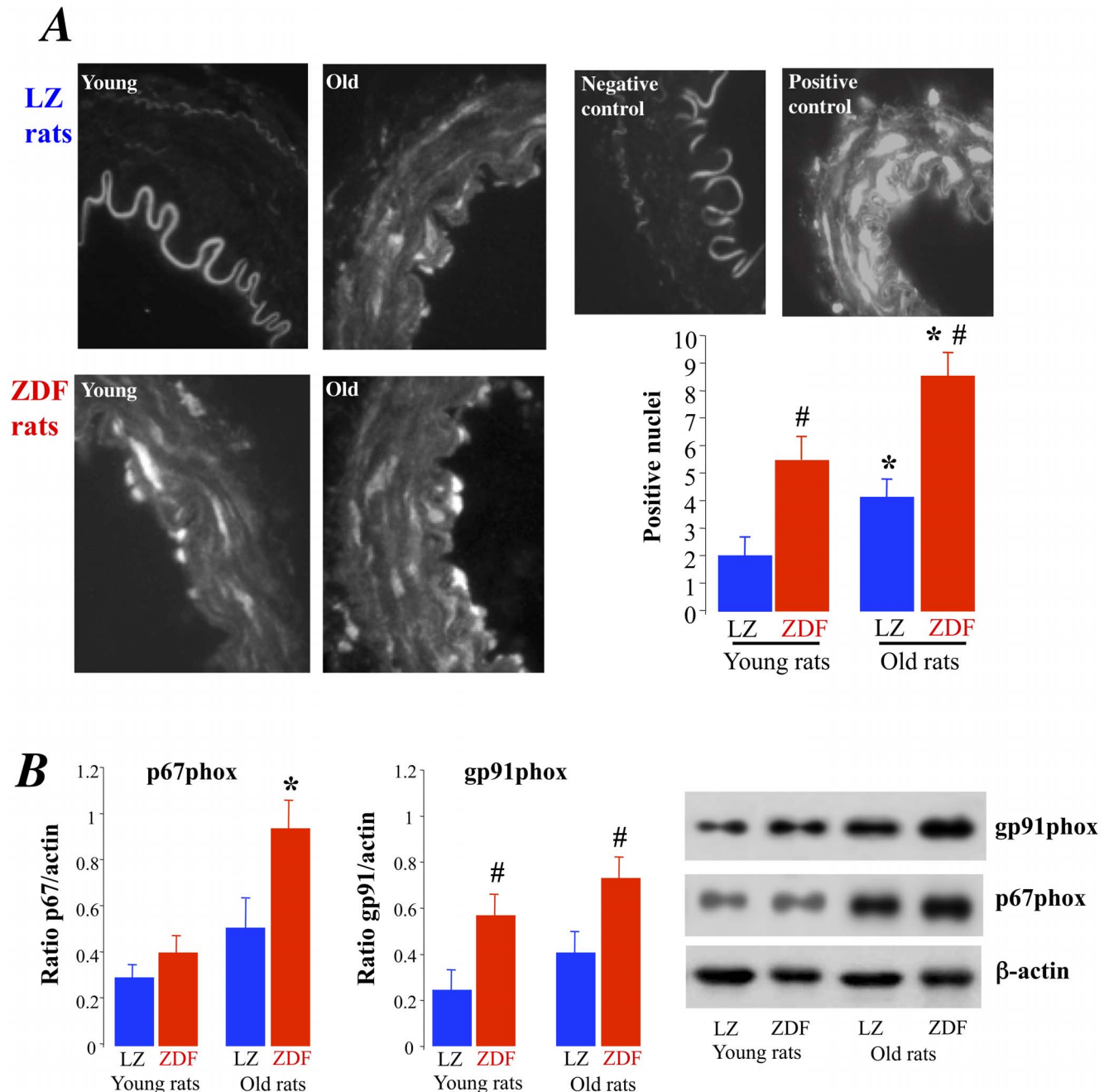
In a separated series of experiments 12-month-old male LZ and ZDF rats were treated with the COX2 inhibitor Celecoxib (25 mg/kg/day, forced feeding once a day, 21 days) or with the combination of Celecoxib (25 mg/kg/day) plus 1-Oxyl-2, 2, 6, 6-tetramethyl-4-hydroxypiperidine (tempol, forced feeding once a day, 10 mg/kg per day, 21 days) [12,22].

The procedure followed in the care and euthanasia of the study animals complied with the European Community Standards on the Care and Use of Laboratory Animals (Ministère de

l'Agriculture, France, authorization No. 6422). The protocol was approved by the Committee on the Ethics of Animal Experiments of the "Pays de la Loire" Region ("Comité d'éthique en Expérimentation Animale", CEEA, permit # CEEA PdL 2008.10).

#### Pharmacological Profile of Isolated Mesenteric Arteries

Segments of mesenteric arteries (2 mm long) were mounted on a wire-myograph (DMT, Aarhus, DK) [23]. 2 tungsten wires (40  $\mu$ m diameter) were inserted into the lumen of the arteries and respectively fixed to a force transducer and a micrometer. Arteries



**Figure 2. Measurement of the reactive oxygen species level in mesenteric resistance arteries isolated from LZ and ZDF rats using dihydroethydim microfluorography (A).** The expression level of the NADP(H)-oxidase subunits gp91 and p67 (B) was measured using Western-Blot analysis. Mean  $\pm$  SEM is presented (n = 10 per group). #P<0.01, ZDF versus LZ. \*P<0.01, old versus young rats. doi:10.1371/journal.pone.0068217.g002

were bathed in a physiological salt solution (PSS) of the following composition (mmol/L): 130.0, NaCl; 15.0, NaHCO<sub>3</sub>; 3.7, KCl; 1.6, CaCl<sub>2</sub>; 1.2, MgSO<sub>4</sub> and 11.0, glucose (pH 7.4, PO<sub>2</sub> 160 mm Hg, and PCO<sub>2</sub> 37 mm Hg) [24]. Wall tension was applied as described previously [25]. Artery viability was tested using a potassium rich PSS (80 mmol/L). Cumulative concentration-response curve to phenylephrine (1 nmol/L to 10  $\mu$ mol/L) was then performed [26]. After washout the arteries were precontracted with phenylephrine to a level approximately equivalent to 50% of the maximal response. Cumulative concentration-response curve to acetylcholine (ACh, 0.001 to 3  $\mu$ mol/L) was then performed in the presence or absence of N (omega)-nitro-L-arginin methyl ester (L-NAME, 100  $\mu$ mol/L, 20 min), tempol (10  $\mu$ mol/L, 20 min) or N-[2-(cyclohexyloxy)-4-nitrophenyl]-methanesulfonamide (NS398, 10  $\mu$ mol/L, 20 min) for 20 min [27].

### Western Blot Analysis

Other segments of arteries were homogenized and proteins (25  $\mu$ g total protein from each sample) were separated by SDS-PAGE using a 4% stacking gel followed by a 10% running gel. Proteins were detected with specific antibodies (Transduction Laboratories, eNOS 1:1000, Cav-1 1:4000, p67 and gp91 and actin 1:1000 in T-TBS-BSA 5%). COX-2 polyclonal antibodies (1:1000) were obtained from Santa Cruz Biotechnology. Protein expression was visualized using the ECL-Plus Chemiluminescence kit (Amersham) [28].

### Detection of Reactive Oxygen Species (ROS) Using Confocal Microscopy

Other arterial segments were embedded vertically in Tissue-tek (Sakura) and frozen in isopentane. ROS detection was performed on transverse cross sections 7  $\mu$ m thick incubated with dihydroethidine (DHE) as previously described [29]. DHE, in the presence of superoxide, is briefly oxidized to fluorescent ethidium bromide. Ethidium bromide is trapped by intercalation with DNA, and the number of fluorescent nuclei indicates the relative level of superoxide production. In negative control experiments arterial sections were incubated with Tempol before DHE staining [29]. Positive control experiments were performed using MRA isolated from a rat treated for 6 hours with bacterial lipopolysaccharide. Positive staining was visualized using confocal microscopy and QED-image software (Solamere Technology, Salt Lake City, UT) [30]. Image analysis was performed using Histolab (Microvision, France) [31].

### Immunohistological Analysis of COX-2

Immunostaining of COX-2 was performed as previously described [12] segments of mesenteric resistance arteries were mounted in embedding medium (Tissue-Tek, Miles, Inc), frozen in isopentane, pre-cooled in liquid nitrogen and stored at  $-80^{\circ}$ C. Transverse cross sections (7  $\mu$ m thick) were incubated with primary goat anti-COX-2 polyclonal antibodies (Santa Cruz Biotechnology, 1:100) followed by the fluorescent secondary antibody (1:200). In negative control experiments the primary antibody was omitted. Positive control experiments were performed using MRA isolated from a rat treated for 6 hours with bacterial lipopolysaccharide. Positive staining was visualized as described above [32].

### Statistical Analysis

Results are expressed as means  $\pm$  SEM. Significance of the difference between arteries was determined by ANOVA (1-factor

ANOVA or ANOVA for consecutive measurements, when appropriate). Means were compared by paired t-test or by the Bonferroni test for multigroup comparisons. Values of  $p < 0.05$  were considered to be significant.

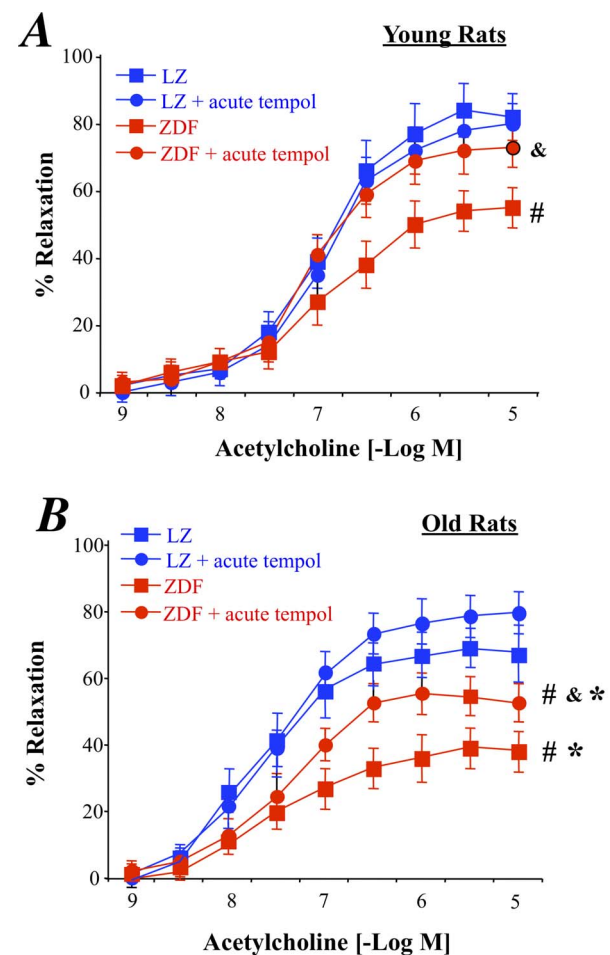
## Results

### Animals Parameters

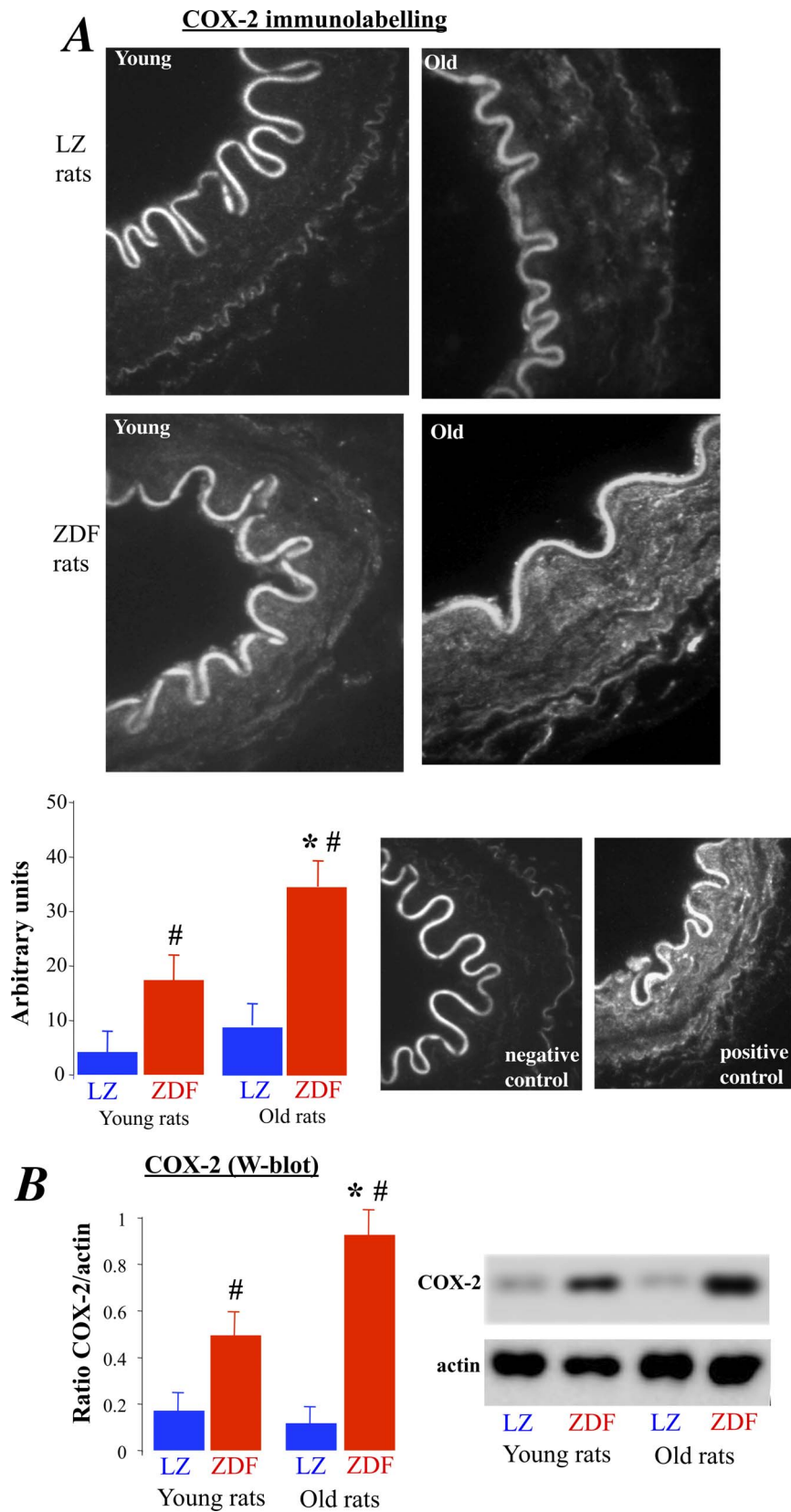
Rat body weight, greater in young ZDF rats than in young LZ rats decreased with age in ZDF rats (table 1), in agreement with previous studies [33,34]. Nevertheless, blood glucose in ZDF rats was significantly higher than in LZ rats and was greater in old ZDF rats. ZDF rats were slightly, but significantly hypertensive (table 1).

### Endothelium-dependent Relaxation of Mesenteric Resistance Arteries

Acetylcholine-dependent relaxation was reduced in ZDF rats compared to lean rats (figure 1A–B). Furthermore, aging per se was also associated with a significant reduction in acetylcholine-mediated relaxation in both LZ and ZDF rats (figure 1A–B). The



**Figure 3. Effect of the acute treatment of mesenteric resistance arteries with the antioxidant tempol on acetylcholine-induced relaxation in mesenteric resistance arteries isolated from young (A) and old (B) LZ and ZDF rats.** Mean  $\pm$  SEM is presented (n = 10 per group). # $P < 0.01$ , ZDF versus LZ. \* $P < 0.01$ , old versus young rats. & $P < 0.05$ , effect of tempol on acetylcholine-induced relaxation. doi:10.1371/journal.pone.0068217.g003



**Figure 4. Measurement of COX-2 expression level in mesenteric resistance arteries isolated from young and old LZ and ZDF rats using immunolabelling (A) and Western-blot analysis (B).** Mean  $\pm$  SEM is presented (n = 10 per group). <sup>#</sup>P<0.01, ZDF versus LZ. <sup>\*</sup>P<0.01, old versus young rats.

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reduction in relaxation due to ageing was greater in ZDF rats than in LZ rats ( $47 \pm 6\%$  in old rats versus  $28 \pm 4\%$  in young rats,  $P < 0.01$ ), showing that the combination of diabetes and aging further impaired endothelium-dependent relaxation. Precontraction prior to acetylcholine-mediated relaxation was similar in the different groups (inserts in figure 1 A and B).

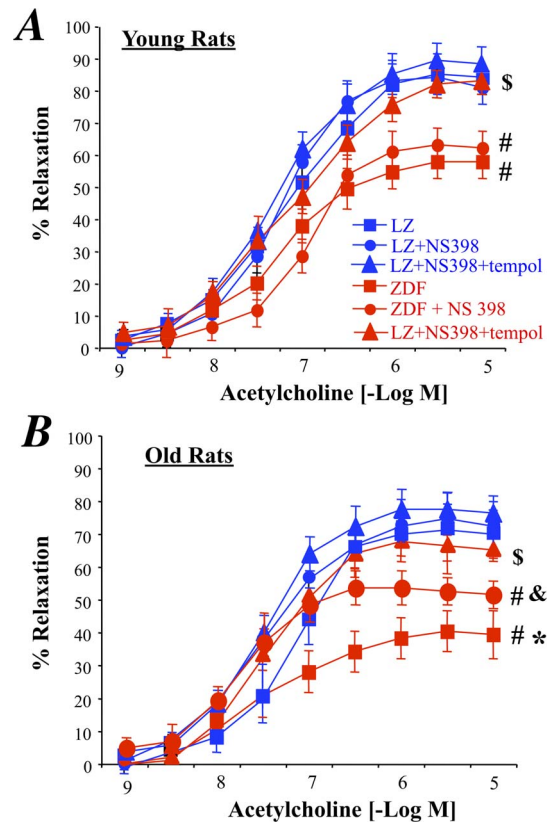
In young LZ rats, L-NAME suppressed acetylcholine-mediated relaxation whereas it was inhibited, but not suppressed, in old LZ rats (figure 1B). In ZDF rats, L-NAME significantly reduced acetylcholine-induced relaxation by  $61 \pm 8\%$  in young rats (figure 1A) and by only  $24 \pm 4\%$  in old rats (figure 1B,  $P < 0.01$  versus young ZDF rats). Accordingly the inhibitory effect of L-NAME was greatly reduced in old ZDF rats. Furthermore, eNOS expression level (relative to beta-actin) was lower in older rats than in young rats, although no difference between ZDF and LZ rats was found in the mesenteric resistance arteries (figure 1C). Finally, eNOS expression level, reduced in old versus young rats, was not altered by type 2 diabetes.

### Reactive Oxygen Species Detection and Effect on Endothelium-dependent Relaxation

As excessive oxidative stress has been shown to reduce endothelium-mediated dilation in diabetes, we investigated the ROS level in mesenteric arteries was analyzed using dihydroethidine (DHE) microfluorography. We found that DHE staining was higher in ZDF than in LZ rats whatever the age. In addition, in older animals the ROS level was higher than in younger animals in both LZ and ZDF rats (figure 2A). The increased DHE staining was associated with a greater expression level of the NAD(P)H-oxidase subunit p67phox in old ZDF rats than in the other groups (figure 2B) whereas the gp91phox expression level was significantly higher in both young and old ZDF rats compared to age-matched LZ rats (figure 2B). Furthermore, tempol, acutely added to the bath of isolated mesenteric arteries improved acetylcholine-induced relaxation in both young and old ZDF rats without affecting relaxation in LZ rats (figure 3). In young ZDF rats, acetylcholine-induced relaxation in the presence of tempol was equivalent to the relaxation found in LZ rats (figure 3A). On the other hand, although tempol significantly improved relaxation in old ZDF rats, it did not fully restore relaxation to the level found in LZ rats (figure 3B) suggesting that another vasoconstrictor agent prevented complete relaxation.

### COX-2 Detection and Effect on Endothelium-dependent Relaxation

COX-2 was expressed in ZDF rats and not in LZ animals (figure 4A and B). In addition, in old rats COX-2 expression level was significantly greater than in young ZDF rats. In order to determine the role of COX-2 in the endothelium dysfunction, we tested the effect of the COX-2 inhibitor NS398 on acetylcholine-induced relaxation in the arteries of ZDF rats (figure 5). First, NS398 had no significant effect on acetylcholine-induced relaxation in young and old LZ rats. Similarly, NS398 did not affect acetylcholine-induced relaxation in young ZDF rats (figure 5A). In young ZDF rats, NS398 and tempol improved relaxation so that it was no different than LZ rats. By contrast with young ZDF rats, NS398 in old ZDF rats significantly improved acetylcholine-induced relaxation (figure 5B), although relaxation in the presence of NS 398 remained significantly lower than in LZ rats. The addition of tempol to NS398 fully improved acetylcholine-induced relaxation in old ZDF rats, so that acetylcholine-induced relaxation in the presence of NS398 and tempol was no different than in LZ rats. Accordingly, in old ZDF COX-2 derived



**Figure 5. Effect of an acute treatment of mesenteric resistance arteries with the COX-2 inhibitor NS398 or NS398 and the antioxidant tempol on acetylcholine-induced relaxation in young (A) and old (B) LZ and ZDF rats.** Mean  $\pm$  SEM is presented ( $n = 10$  per group). # $P < 0.01$ , ZDF versus LZ. \* $P < 0.01$ , old versus young rats.  $^{\&}$  $P < 0.05$ , effect of NS398 alone on acetylcholine-induced relaxation.  $^{\$}$  $P < 0.05$ , effect of NS398+tempol on acetylcholine-induced relaxation.

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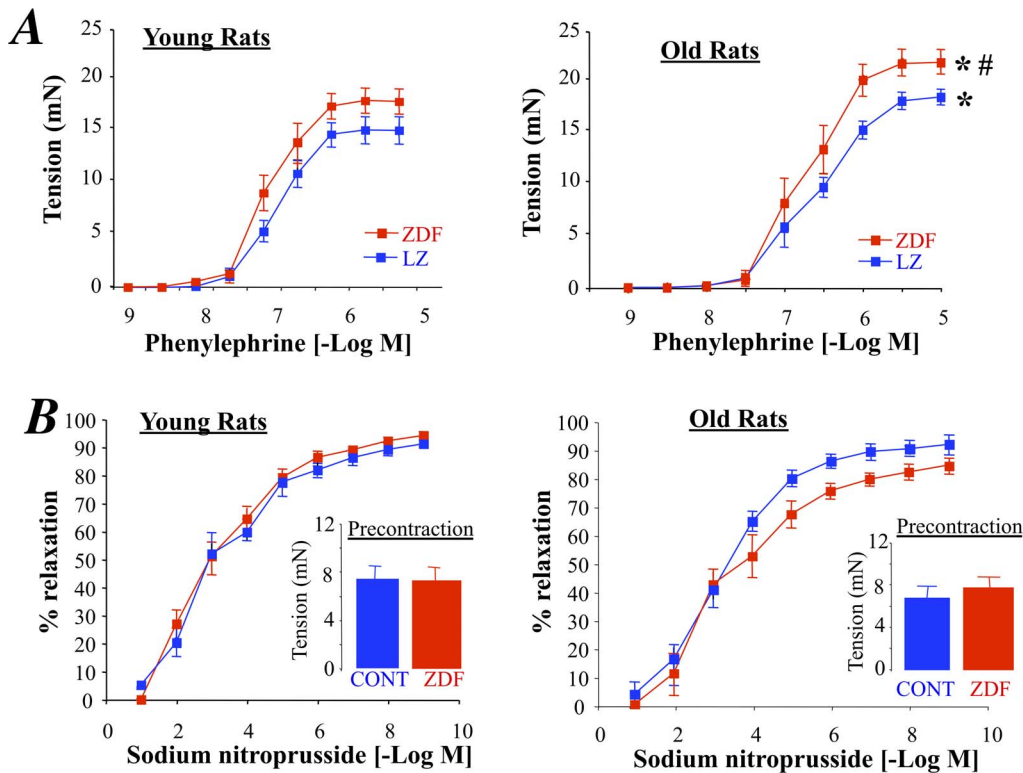
vasoconstrictor prostanoids was associated with superoxide in order to further reduce endothelium-dependent relaxation.

### Phenylephrine-mediated Contraction and Endothelium-independent Relaxation

Finally, in old rats phenylephrine-mediated contraction was greater than in young rats (figure 6A). The contraction was also higher in ZDF than in LZ rats, although the difference only reached significance in old rats (figure 6A). On the other hand, sodium nitroprusside, which induced endothelium-independent relaxation, was not affected by aging or by diabetes (figure 6B).

### In vivo Blockade of COX-2

In old ZDF rats treated chronically with the COX-2 inhibitor celecoxib, acetylcholine-induced relaxation was improved whereas celecoxib did not affect relaxation in old LZ rats (figure 7A). Nevertheless, acetylcholine-induced relaxation in old ZDF rats treated with celecoxib did not reach the level of relaxation obtained in LZ rats (figure 7A). In old ZDF rats treated chronically with the celecoxib plus the antioxidant tempol, acetylcholine-induced relaxation was improved and reached the level obtained in LZ rats (figure 7B). The treatment with celecoxib plus tempol did not affect relaxation in old LZ rats (figure 7B).



**Figure 6. Concentration-response curves to phenylephrine (A) and sodium nitroprusside (B) performed in isolated arteries from young and old LZ and ZDF rats.** The inserts in panels B represent the precontraction applied to arteries before sodium nitroprusside-mediated relaxation. Mean  $\pm$  SEM is presented (n = 10 per group). \* $P < 0.05$ , old versus young rats # $P < 0.05$ , ZDF versus LZ rats. doi:10.1371/journal.pone.0068217.g006

## Discussion

In this study we found that endothelium-dependent relaxation was reduced in young rats suffering from type 2 diabetes due to excessive ROS production, whereas in one year-old diabetic rats COX-2 derivatives in association with an excessive ROS production reduced endothelium dependent relaxation.

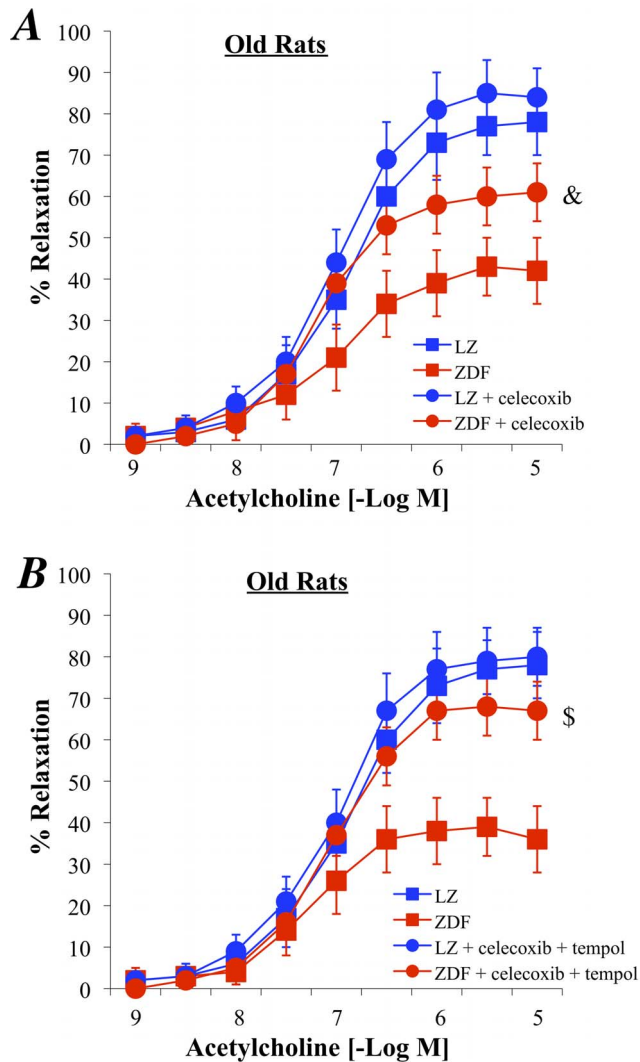
First, we found that endothelium-mediated dilation, evidenced by acetylcholine-dependent relaxation was reduced in ZDF rats compared to lean rats. This finding is in agreement with previous studies showing that type 2 diabetes is associated with endothelial dysfunction in rats [35] as well as in humans [36]. Furthermore, aging per se was associated with a significant reduction in acetylcholine-mediated relaxation in both LZ and ZDF rats, in agreement with previous studies in large conduit arteries [37] and in small resistance arteries [12,35,38]. Also in agreement with these previous works, we found that the combination of diabetes and aging further impaired endothelium-dependent relaxation.

As endothelium-mediated relaxation in mesenteric arteries relies greatly on the production of NO [39–42], concentration-response curves to acetylcholine were repeated after incubation arterial with L-NAME. Although L-NAME significantly reduced relaxation in all groups, the inhibitory effect of L-NAME was greatly reduced in old ZDF rats suggesting a reduced involvement of NO in the relaxation. Thus, in order to find the origin of the excessive endothelial dysfunction found in old ZDF rats we measured eNOS expression level in the mesenteric artery in the 4 study groups. Nevertheless, we found no difference between ZDF and LZ rats. This observation is in agreement with previous studies of resistance arteries [38,43–45] and of large arteries [46] showing a reduced NO-dependent relaxation in diabetic rats. Similarly, NO-mediated

dilation is also reduced in humans [11,47,48]. Surprisingly, eNOS expression level, reduced in old versus young rats, was not altered by type 2 diabetes. This is in agreement with previous studies performed on mesenteric arteries isolated from diabetic db/db mice, another model of type 2 diabetes [49]. On the other hand, in streptozotocine-induced diabetes, eNOS expression decreases together with a reduced NO production [50]. In human subcutaneous arterioles, obesity associated with diabetes or not, was accompanied by a reduction in eNOS expression level and endothelium-mediated dilation [51]. Indeed, it is most likely that NO bioavailability is reduced in diabetes mainly due to excessive oxidative stress. Dimerization of the enzyme is a key step in the production of NO [13,52] and eNOS dimerization is strongly reduced in ZDF rats' mesenteric arteries [53]. The authors have shown that NO-mediated dilation was reduced due to increased oxidative stress leading to eNOS uncoupling rather than to a reduction in expression level.

As excessive oxidative stress has a major role in the endothelium dysfunction observed in diabetes, we investigated the role of superoxide in acetylcholine-mediated relaxation. Certainly, oxidative stress has been shown to play a major role in the reduction in endothelium-mediated relaxation in diabetes in most vascular territories [54] including the mesenteric vasculature [12,49]. A similar superoxide effect has been shown in the metabolic syndrome [10,21,22,55,56]. In the present study the ROS level in mesenteric arteries was analyzed using dihydroethidine (DHE) microfluorography as previously shown [21,22,57]. We have previously validated this approach in the detection of ROS in isolated mesenteric arteries. Certainly, we obtained similar results with DHE staining as with rhodamin-staining, 3-nitrotyrosine





**Figure 7. Effect of a chronic treatment of old LZ and ZDF rats with celecoxib (A) or with the combination of celecoxib and tempol (B) on acetylcholine-induced relaxation of isolated mesenteric resistance arteries.** Mean  $\pm$  SEM is presented ( $n=8$  per group). <sup>a</sup> $P<0.05$ , effect of celecoxib alone on acetylcholine-induced relaxation. <sup>b</sup> $P<0.05$ , effect of celecoxib+tempol on acetylcholine-induced relaxation.

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labeling, NADP(H)Oxidase subunits level measurement and O<sub>2</sub>-quantification using electron paramagnetic resonance [29]. In the present study, DHE staining was larger in ZDF than in LZ rats whatever the age and it was higher in old than in young animals in both LZ and ZDF rats. The increased DHE staining was associated with a greater expression level of the NAD(P)H-oxidase subunits in old ZDF rats. The occurrence of an excessive oxidative stress reducing endothelium (NO)-dependent relaxation was confirmed using the acute effect of the antioxidant tempol as previously shown and validated [21,29,54]. This test is indeed a functional proof of the effect of ROS on endothelium-mediated relaxation. Acute tempol improved acetylcholine-induced relaxation in both young and old ZDF rats without affecting relaxation in LZ rats whereas in young ZDF rats, acetylcholine-induced relaxation in the presence of tempol was equivalent to the relaxation found in LZ rats. This observation shows that ROS reduced endothelium (NO)-mediated relaxation in both young

and old ZDF rats. This is in agreement with previous studies showing that ROS reduces endothelium-mediated relaxation in ZDF rats as well as in other animal models of type 2 diabetes [15,45,53,58] and in human endothelial cells [59]. Nevertheless, in old ZDF rats, tempol did not fully restore relaxation to the level found in LZ rats suggesting that another vasoconstrictor agent was involved.

In order to better understand this endothelial dysfunction found in old ZDF rats, we investigated the involvement of COX-2 in acetylcholine-induced relaxation. Indeed, COX-2 is expressed in resistance arteries [22,60,61] as well as in large vessels [62] in animal models of ageing. In these studies, COX-2 expression was associated with the production of vasoconstrictor prostanoids. In the present study, COX-2 was detected using immunostaining and Western-Blot in ZDF rats and not in LZ animals. Furthermore, in old ZDF rats COX-2 expression was significantly larger than in young ZDF rats. In agreement with this observation, we found that NS398 had no effect on acetylcholine-induced relaxation in young and old LZ rats as well as in young ZDF rats. By contrast, NS398 in old ZDF rats significantly improved acetylcholine-induced relaxation, although relaxation in the presence of NS 398 remained significantly lower than in LZ rats. The addition of tempol to NS398 fully improved acetylcholine-induced relaxation in old ZDF rats, so that acetylcholine-induced relaxation in the presence of NS398 and tempol was no different than in LZ rats. Accordingly, in old ZDF COX-2 derived vasoconstrictor prostanoids was associated with superoxide in order to further reduce endothelium-dependent relaxation. As COX-2 derived vasoconstrictors were involved in old ZDF rats which also have a greater oxidative stress a possible explanation could be that ROS per se activate COX-2, at least in part. Indeed, a recent study has clearly shown that oxidative stress has a key role in the increased COX-2 activity observed in renal arteries of renovascular hypertensive rats. Indeed, this later study also shows that COX-2-derived PGF<sub>2 $\alpha$</sub>  has an major role in mediating endothelial dysfunction [63]. The involvement of ROS and COX-2 derivatives was further confirmed using old ZDF rats treated chronically with the COX-2 inhibitor celecoxib alone or in combination with tempol. Endothelium-mediated relaxation was fully restored to control level only when the 2 substances were given together to the rats. Nevertheless, it should be noted that the use of COX-2 inhibitors is not recommended as these drugs possess deleterious side effects including higher risk of myocardial infarction [64].

Finally, in old rats phenylephrine-mediated contraction was greater than in young rats. Furthermore, phenylephrine-mediated contraction was also higher in ZDF than in LZ rats. These observations are in agreement with previous studies, which have shown that both aging [37,65] and diabetes [12,66] are associated with arterial wall hypertrophy and consequently with hypercontractility.

## Conclusion

We found that long-term exposure to diabetes increased endothelium damages and further reduced endothelium-mediated relaxation in mesenteric resistance arteries. In these arteries COX-2 derivatives in addition to ROS contributed to severely reduce endothelium-dependent dilation. This finding is of importance as it shows that deleterious pathways may add one to each other when diabetes evolves over time. Certainly, the present experimental conditions most probably represent a more common pathological condition as diabetes is usually diagnosed several months or years after the onset of the metabolic disorders leading to hyperglycemia. Consequently, the present study reinforces the

assumption that not only oxidative stress but also inflammation contribute, most probably synergistically, to the vascular disorders occurring in type 2 diabetes.

## References

- Wild S, Roglic G, Green A, Sicree R, King H (2004) Global prevalence of diabetes: estimates for the year 2000 and projections for 2030. *Diabetes Care* 27: 1047–1053.
- Eckel RH, Grundy SM, Zimmet PZ (2005) The metabolic syndrome. *Lancet* 365: 1415–1428.
- Adams KF, Schatzkin A, Harris TB, Kipnis V, Mouw T, et al. (2006) Overweight, obesity, and mortality in a large prospective cohort of persons 50 to 71 years old. *N Engl J Med* 355: 763–778.
- Ford ES (2005) Risks for all-cause mortality, cardiovascular disease, and diabetes associated with the metabolic syndrome: a summary of the evidence. *Diabetes Care* 28: 1769–1778.
- Mokdad AH, Bowman BA, Ford ES, Vinicor F, Marks JS, et al. (2001) The continuing epidemics of obesity and diabetes in the United States. *Jama* 286: 1195–1200.
- Pieper GM, Siebeneich W (1997) Diabetes-induced endothelial dysfunction is prevented by long-term treatment with the modified iron chelator, hydroxyethyl starch conjugated-deferoxamine. *J Cardiovasc Pharmacol* 30: 734–738.
- Winer N, Sowers JR (2007) Diabetes and arterial stiffening. *Adv Cardiol* 44: 245–251.
- Grundey SM (2008) Metabolic syndrome pandemic. *Arterioscler Thromb Vasc Biol* 28: 629–636.
- Barton M (2010) Obesity and aging: determinants of endothelial cell dysfunction and atherosclerosis. *Pflugers Arch* 460: 825–837.
- Vessieres E, Freidja ML, Loufrani L, Fassot C, Henrion D (2012) Flow (shear stress)-mediated remodeling of resistance arteries in diabetes. *Vascul Pharmacol* 57: 173–178.
- Tousoulis D, Kampoli AM, Stefanadis C (2012) Diabetes mellitus and vascular endothelial dysfunction: current perspectives. *Curr Vasc Pharmacol* 10: 19–32.
- Retailleau K, Belin de Chantemele EJ, Chanoine S, Guihot AL, Vessieres E, et al. (2010) Reactive oxygen species and cyclooxygenase 2-derived thromboxane A2 reduce angiotensin II type 2 receptor vasorelaxation in diabetic rat resistance arteries. *Hypertension* 55: 339–344.
- Sharma A, Bernatchez PN, de Haan JB (2012) Targeting endothelial dysfunction in vascular complications associated with diabetes. *Int J Vasc Med* 2012: 750126.
- Portik-Dobos V, Anstadt MP, Hutchinson J, Bannan M, Ergul A (2002) Evidence for a matrix metalloproteinase induction/activation system in arterial vasculature and decreased synthesis and activity in diabetes. *Diabetes* 51: 3063–3068.
- Ding H, Aljofan M, Triggle CR (2007) Oxidative stress and increased eNOS and NADPH oxidase expression in mouse microvessel endothelial cells. *J Cell Physiol* 212: 682–689.
- Toda N (2012) Age-related changes in endothelial function and blood flow regulation. *Pharmacol Ther* 133: 159–176.
- Dumont O, Pinaud F, Guihot AL, Baufreton C, Loufrani L, et al. (2008) Alteration in flow (shear stress)-induced remodeling in rat resistance arteries with aging: improvement by a treatment with hydralazine. *Cardiovasc Res* 77: 600–608.
- Pinaud F, Bocquet A, Dumont O, Retailleau K, Baufreton C, et al. (2007) Paradoxical role of angiotensin II type 2 receptors in resistance arteries of old rats. *Hypertension* 50: 96–102.
- Kampoli AM, Tousoulis D, Briasoulis A, Latsios G, Papageorgiou N, et al. (2011) Potential pathogenic inflammatory mechanisms of endothelial dysfunction induced by type 2 diabetes mellitus. *Curr Pharm Des* 17: 4147–4158.
- Driss A, Devaux C, Henrion D, Duriez M, Thuillez C, et al. (2000) Hemodynamic stresses induce endothelial dysfunction and remodeling of pulmonary artery in experimental compensated heart failure. *Circulation* 101: 2764–2770.
- Bouvet C, Belin de Chantemele E, Guihot AL, Vessieres E, Bocquet A, et al. (2007) Flow-induced remodeling in resistance arteries from obese Zucker rats is associated with endothelial dysfunction. *Hypertension* 50: 248–254.
- Vessieres E, Belin de Chantemele EJ, Toutain B, Guihot AL, Jardel A, et al. (2010) Cyclooxygenase-2 inhibition restored endothelium-mediated relaxation in old obese Zucker rat mesenteric arteries. *Front Physiol* 1: 145.
- Dowell EJ, Henrion D, Benessiano J, Poitevin P, Levy B (1996) Chronic infusion of low-dose angiotensin II potentiates the adrenergic response in vivo. *J Hypertens* 14: 177–182.
- Henrion D, Laher I, Laporte R, Bevan JA (1992) Further evidence from an elastic artery that angiotensin II amplifies noradrenaline-induced contraction through activation of protein kinase C. *Eur J Pharmacol* 224: 13–20.
- Mulvany MJ, Halpern W (1977) Contractile properties of small arterial resistance vessels in spontaneously hypertensive and normotensive rats. *Circ Res* 41: 19–26.
- Henrion D, Laher I, Bevan JA (1992) Intraluminal flow increases vascular tone and  $^{45}\text{Ca}^{2+}$  influx in the rabbit facial vein. *Circ Res* 71: 339–345.
- Bolla M, Matrougui K, Loufrani L, Maclouf J, Levy B, et al. (2002) p38 mitogen-activated protein kinase activation is required for thromboxane-induced contraction in perfused and pressurized rat mesenteric resistance arteries. *J Vasc Res* 39: 353–360.
- Loufrani L, Retailleau K, Bocquet A, Dumont O, Danker K, et al. (2008) Key role of  $\alpha(1)\beta(1)$ -integrin in the activation of PI3-kinase-Akt by flow (shear stress) in resistance arteries. *Am J Physiol Heart Circ Physiol* 294: H1906–1913.
- Cousin M, Custaud MA, Baron-Menguy C, Toutain B, Dumont O, et al. (2010) Role of angiotensin II in the remodeling induced by a chronic increase in flow in rat mesenteric resistance arteries. *Hypertension* 55: 109–115.
- Freidja ML, Toutain B, Caillon A, Desquiret V, Lambert D, et al. (2011) Heme oxygenase 1 is differentially involved in blood flow-dependent arterial remodeling: role of inflammation, oxidative stress, and nitric oxide. *Hypertension* 58: 225–231.
- Baron-Menguy C, Bocquet A, Guihot AL, Chappard D, Amiot MJ, et al. (2007) Effects of red wine polyphenols on postischemic neovascularization model in rats: low doses are proangiogenic, high doses anti-angiogenic. *FASEB J* 21: 3511–3521.
- Tarhouni K, Guihot AL, Freidja ML, Toutain B, Henrion B, et al. (2013) Key role of estrogens and endothelial estrogen receptor alpha in blood flow-mediated remodeling of resistance arteries. *Arterioscler Thromb Vasc Biol* 33: 605–611.
- Doisaki M, Katano Y, Nakano I, Hirooka Y, Itoh A, et al. (2010) Regulation of hepatic branched-chain alpha-keto acid dehydrogenase kinase in a rat model for type 2 diabetes mellitus at different stages of the disease. *Biochem Biophys Res Commun* 393: 303–307.
- Howarth FC, Al Kitibi MK, Hameed RS, Adeghate E (2011) Pancreatic peptides in young and elderly Zucker type 2 diabetic fatty rats. *JOP* 12: 567–573.
- Oltman CL, Richou LL, Davidson EP, Coppey LJ, Lund DD, et al. (2006) Progression of Coronary and Mesenteric Vascular Dysfunction in Zucker Obese and Zucker Diabetic Fatty Rats. *Am J Physiol Heart Circ Physiol*.
- Naka KK, Papathanassiou K, Bechlioulis A, Kazakos N, Pappas K, et al. (2012) Determinants of vascular function in patients with type 2 diabetes. *Cardiovasc Diabetol* 11: 127.
- Najjar SS, Scuteri A, Lakatta EG (2005) Arterial aging: is it an immutable cardiovascular risk factor? *Hypertension* 46: 454–462.
- Freidja ML, Tarhouni K, Toutain B, Fassot C, Loufrani L, et al. (2012) The AGE-breaker ALT-711 restores high blood flow-dependent remodeling in mesenteric resistance arteries in a rat model of type 2 diabetes. *Diabetes* 61: 1562–1572.
- Dowell EJ, Henrion D, Duriez M, Michel JB (1996) Vascular reactivity in mesenteric resistance arteries following chronic nitric oxide synthase inhibition in Wistar rats. *Br J Pharmacol* 117: 341–346.
- Loufrani L, Levy BI, Henrion D (2002) Defect in microvascular adaptation to chronic changes in blood flow in mice lacking the gene encoding for dystrophin. *Circ Res* 91: 1183–1189.
- Henrion D, Dechaux E, Dowell EJ, Maclour J, Samuel JL, et al. (1997) Alteration of flow-induced dilatation in mesenteric resistance arteries of L-NAME treated rats and its partial association with induction of cyclo-oxygenase-2. *Br J Pharmacol* 121: 83–90.
- Matrougui K, Levy BI, Henrion D (2000) Tissue angiotensin II and endothelin-1 modulate differently the response to flow in mesenteric resistance arteries of normotensive and spontaneously hypertensive rats. *Br J Pharmacol* 130: 521–526.
- Matsumoto T, Ishida K, Nakayama N, Taguchi K, Kobayashi T, et al. (2010) Mechanisms underlying the losartan treatment-induced improvement in the endothelial dysfunction seen in mesenteric arteries from type 2 diabetic rats. *Pharmacol Res* 62: 271–281.
- Belin de Chantemele EJ, Vessieres E, Guihot AL, Toutain B, Maquignau M, et al. (2009) Type 2 diabetes severely impairs structural and functional adaptation of rat resistance arteries to chronic changes in blood flow. *Cardiovasc Res* 81: 788–796.
- Leo CH, Hart JL, Woodman OL (2011) 3',4'-Dihydroxyflavonol reduces superoxide and improves nitric oxide function in diabetic rat mesenteric arteries. *PLoS One* 6: e20813.
- Sena CM, Matafome P, Louro T, Nunes E, Fernandes R, et al. (2011) Metformin restores endothelial function in aorta of diabetic rats. *Br J Pharmacol* 163: 424–437.
- Laight DW, Carrier MJ, Anggard EE (2000) Antioxidants, diabetes and endothelial dysfunction. *Cardiovasc Res* 47: 457–464.
- Hile C, Veves A (2003) Diabetic neuropathy and microcirculation. *Curr Diab Rep* 3: 446–451.

## Author Contributions

Conceived and designed the experiments: DH LL. Performed the experiments: EV ALG BT MM. Analyzed the data: DH EV ALG BT LL. Contributed reagents/materials/analysis tools: EV. Wrote the paper: DH LL CF.

49. Huang A, Yang YM, Feher A, Bagi Z, Kaley G, et al. (2012) Exacerbation of endothelial dysfunction during the progression of diabetes: role of oxidative stress. *Am J Physiol Regul Integr Comp Physiol* 302: R674–681.
50. El-Remessy AB, Tawfik HE, Matragoon S, Pillai B, Caldwell RB, et al. (2010) Peroxynitrite mediates diabetes-induced endothelial dysfunction: possible role of Rho kinase activation. *Exp Diabetes Res* 2010: 247861.
51. Georgescu A, Popov D, Constantin A, Nemezc M, Alexandru N, et al. (2011) Dysfunction of human subcutaneous fat arterioles in obesity alone or obesity associated with Type 2 diabetes. *Clin Sci (Lond)* 120: 463–472.
52. Forstermann U, Munzel T (2006) Endothelial nitric oxide synthase in vascular disease: from marvel to menace. *Circulation* 113: 1708–1714.
53. Leo CH, Hart JL, Woodman OL (2011) Impairment of both nitric oxide-mediated and EDHF-type relaxation in small mesenteric arteries from rats with streptozotocin-induced diabetes. *Br J Pharmacol* 162: 365–377.
54. Oniki H, Goto K, Fujii K, Kansui Y, Murakami N, et al. (2012) Effects of the Superoxide Dismutase Mimetic Tempol on Impaired Endothelium-Dependent and Endothelium-Independent Relaxations in Type II Diabetic Rats. *Clin Exp Hypertens*.
55. Munzel T, Gori T, Bruno RM, Taddei S (2010) Is oxidative stress a therapeutic target in cardiovascular disease? *Eur Heart J* 31: 2741–2748.
56. Vessieres E, Belin de Chantemele EJ, Guihot AL, Jardel A, Toutain B, et al. (2013) Cyclooxygenase-2-derived prostanoids reduce inward arterial remodeling induced by blood flow reduction in old obese Zucker rat mesenteric arteries. *Vascul Pharmacol* 58: 356–362.
57. Belin de Chantemele EJ, Vessieres E, Dumont O, Guihot AL, Toutain B, et al. (2009) Reactive oxygen species are necessary for high flow (shear stress)-induced diameter enlargement of rat resistance arteries. *Microcirculation* 16: 391–402.
58. Tian XY, Wong WT, Xu A, Lu Y, Zhang Y, et al. (2012) Uncoupling protein-2 protects endothelial function in diet-induced obese mice. *Circ Res* 110: 1211–1216.
59. Cosentino F, Hishikawa K, Katusic ZS, Luscher TF (1997) High glucose increases nitric oxide synthase expression and superoxide anion generation in human aortic endothelial cells. *Circulation* 96: 25–28.
60. Gendron ME, Thorin-Trescases N, Villeneuve L, Thorin E (2007) Aging associated with mild dyslipidemia reveals that COX-2 preserves dilation despite endothelial dysfunction. *Am J Physiol Heart Circ Physiol* 292: H451–458.
61. Belin de Chantemele EJ, Vessieres E, Guihot AL, Toutain B, Loufrani L, et al. (2010) Cyclooxygenase-2 preserves flow-mediated remodelling in old obese Zucker rat mesenteric arteries. *Cardiovasc Res* 86: 516–525.
62. Wong SL, Leung FP, Lau CW, Au CL, Yung LM, et al. (2009) Cyclooxygenase-2-derived prostaglandin F2alpha mediates endothelium-dependent contractions in the aortae of hamsters with increased impact during aging. *Circ Res* 104: 228–235.
63. Tian XY, Wong WT, Leung FP, Zhang Y, Wang YX, et al. (2012) Oxidative stress-dependent cyclooxygenase-2-derived prostaglandin f(2alpha) impairs endothelial function in renovascular hypertensive rats. *Antioxid Redox Signal* 16: 363–373.
64. Garcia Rodriguez LA, Gonzalez-Perez A, Bueno H, Hwa J (2011) NSAID use selectively increases the risk of non-fatal myocardial infarction: a systematic review of randomised trials and observational studies. *PLoS One* 6: e16780.
65. Dumont O, Loufrani L, Henrion D (2007) Key role of the NO-pathway and matrix metalloprotease-9 in high blood flow-induced remodeling of rat resistance arteries. *Arterioscler Thromb Vasc Biol* 27: 317–324.
66. Prevost G, Bulckaen H, Gaxatte C, Boulanger E, Beraud G, et al. (2011) Structural modifications in the arterial wall during physiological aging and as a result of diabetes mellitus in a mouse model: are the changes comparable? *Diabetes Metab* 37: 106–111.