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Topical Treatment With Brimonidine and Somatostatin Causes Retinal Vascular Dilation in Patients With Early Diabetic Retinopathy From the EUROCONDOR

Jakob Grauslund,1–3 Ulrik Frydkjaer-Olsen,1,2 Tunde Peto,2,4 Jimena Fernández-Carneado,5 Berta Ponsati,5 Cristina Hernández,6 José Cunha-Vaz,7 and Rafael Simó6; for the EUROCONDOR

1Department of Ophthalmology, Odense University Hospital, Odense, Denmark
2Department of Clinical Research, University of Southern Denmark, Odense, Denmark
3Steno Diabetes Center Odense, Odense University Hospital, Odense, Denmark
4Centre for Public Health, Institute of Clinical Sciences, Queen’s University Belfast, Belfast, United Kingdom
5BCN Peptides, Barcelona, Spain
6Diabetes and Metabolism Research Unit, Institut de Recerca Hospital Universitari Vall d’Hebron (VHIR), Barcelona, Spain
7Association for Innovation and Biomedical Research on Light and Image (AIBILI), and University of Coimbra, Coimbra, Portugal

PRACTICE POINTS

8–10 The concept has been supported by experimental data demonstrating a prevention of retinal neurodegeneration by topical administration of neuroprotective drugs.8–10 The European Consortium for the Early Treatment of Diabetic Retinopathy (EUROCONDOR) aimed to test the effect of topically administered brimonidine and somatostatin on retinal neurodysfunction in patients with type 2 diabetes with no or early DR.11 The study reported that in patients with preexisting retinal neurodysfunction, topical neuroprotection arrested further loss of retinal function as measured by multifocal electroretinography implicit time.12 It would also be important to investigate the structural vascular effects of topical neuroprotective treatment. However, as demonstrated by the Diabetes Control Complications Trial (DCCT), even a strong systemic intervention like strict glycemic regulation would take 3 years to affect the level of DR.13 Hence, more subtle preclinical endpoints for vascular dysfunction are needed. In this aspect, the retinal vascular calibers, which can be measured noninvasively, have been associated with14,15 and even predictive of sight-threatening DR.16 In order to explore the effect of topical neuroprotection on the retinal vasculature of patients with type 2 diabetes and early vis.

Keywords: diabetic retinopathy, topical, retinal vasculature, brimonidine, somatostatin

PURPOSE. Structural retinal microvascular changes have been identified as risk markers of diabetic retinopathy (DR). In order to estimate the retinal response of neuroprotective eye drops, we aimed to evaluate the effect of topical retinal neuroprotection on retinal microvascular changes in early DR.

METHODS. Patients with type 2 diabetes with no or early DR were randomized 1:1:1 to topical treatment with placebo, brimonidine, or somatostatin in a 96-week prospective, phase II to III, European multicenter trial. Retinal vascular calibers were measured semi-automatically in digital fundus images by certified graders at baseline and follow-up and summarized as central retinal arteriolar and venular equivalent (CRAE and CRVE).

RESULTS. Of 449 patients originally included, 297 completed the study with gradable retinal images. Median age and duration of diabetes was 64.5 and 9.9 years, and 65.7% were male. At baseline, Early Treatment Diabetic Retinopathy Study levels were 10 (no DR, 42.8%), 20 (minimal DR, 28.3%), and 35 (mild DR, 29.0%), and CRAE and CRVE did not differ between groups. As opposed to patients with no or minimal DR at baseline, patients with mild DR in the active groups developed a larger retinal arteriolar (brimonidine: +13.9 μm, P = 0.001; somatostatin: +7.2 μm, P = 0.006) and venular (brimonidine: +13.9 μm, P = 0.01; somatostatin: +14.3 μm, P = 0.0001) caliber in contrast to those in the placebo group.

CONCLUSIONS. Topical treatment with brimonidine and somatostatin causes retinal arteriolar and venular dilation in patients with type 2 diabetes and preexisting early DR. Upcoming studies should elaborate on the potential of these findings in arresting early DR.

Keywords: diabetic retinopathy, topical, retinal vasculature, brimonidine, somatostatin

Diabetic retinopathy (DR) is the most common complication in diabetes1 and among the leading cause of blindness among working age adults in the Western world.2 DR has traditionally been considered a microvascular disease, but recent evidence has implicated retinal neurodegeneration as an early event that may even precede vascular dysfunction.3,4 At present, retinal photocoagulation and intravitreal therapy with vascular endothelial growth factor inhibitors are considered the gold standard treatment in DR.5–7; but given the invasive nature of the treatments, these are only considered in patients with sight-threatening DR.8–10 The concept has been supported by experimental data demonstrating a prevention of retinal neurodegeneration by topical administration of neuroprotective drugs.8–10 The European Consortium for the Early Treatment of Diabetic Retinopathy (EUROCONDOR) aimed to test the effect of...
Topical Neuroprotective Treatment in Diabetes Patients

DR, the aim of the present study was to evaluate the result of 2 years of treatment with brimonidine and somatostatin on the retinal vascular calibers in a randomized, clinical trial.

METHODS

Study Subjects

The EUROCONDOR [NCT01726075] was a European multicenter, 96-week prospective, interventional, phase II to III, randomized controlled clinical trial that aimed to evaluate the effect of topical neuroprotection to arrest or prevent early retinal neurodegeneration in DR. In a study including 11 European centers, a total of 449 patients were recruited and randomized 1:1:1 to topical treatment twice daily (BID) with placebo, brimonidine tartrate 0.2%, or somatostatin 0.1%. Criteria of inclusion were: type 2 diabetes with no, minimal, or mild DR (Early Treatment Diabetic Retinopathy Study [ETDRS] levels 10, 20, or 35, respectively), duration of type 2 diabetes for at least 5 years, and age between 45 and 75 years. Exclusion criteria included previous retinal photocoagulation, retinal degenerative diseases (i.e., glaucoma), and refractive error or ±5 diopeters or more, hazy ocular media, inadequate pupil dilation, renal failure (creatinine >124 μmol/L) or HbA1c >10% (86 mmol/mol) in the previous 6 months.

For each patient, one eye was identified as the study eye by a central reading center (Coimbra Ophthalmology Reading Centre, Coimbra, Portugal), which also provided grading of the level of DR at baseline and follow-up.

The study was approved and funded by the European Commission Seventh Framework Program (Grant Agreement No. FP7-278040). At all centers, the study was conducted in accordance with the tenets of the Declaration of Helsinki with written informed consent from all patients.

Statistical Methods

Descriptive statistics were calculated for all parameters. Continuous data are presented as median (with range), and categoric data are given as percentage. Development in retinal vascular calibers and level of DR were given as differences between follow-up and baseline.

RESULTS

Among the 297 participants of the study, median age and duration of diabetes at baseline were 64.5 and 9.9 years, and 65.7% were male. Median levels of HbA1c, blood pressure, and body mass index were 7.0%, 135/78 mm Hg and 30.0 kg/m², respectively. Baseline ETDRS levels of DR were 10 (42.8%), 20 (28.3%), or 35 (29.0%), and CRAE and CRVE was 146.6 μm and 214.8 μm, respectively.

Number of patients in each treatment group were 108 (36.4%), 85 (28.6%), and 104 (35.0%) for placebo, brimonidine, and somatostatin, respectively. Among treatment groups, patients did not differ at baseline according to age, sex, duration of diabetes, HbA1c, blood pressure, body mass index, CRAE (146.7 μm, 147.1 μm, and 145.2 μm, P = 0.68) or CRVE (215.6 μm, 212.7 μm, and 215.7 μm, P = 0.89; Table 2). Likewise, there was no difference in CRAE according to baseline level of DR (level 10, 144.1 μm; level 20, 147.7 μm; level 35, 148.2 μm; P = 0.10), but there was a trend toward a higher CRVE for patients with increasing levels of DR (level 10, 213.7 μm; level 20, 214.4 μm; level 35, 216.1 μm; P = 0.056).

During 24 months of topical treatment, there was a median increase of CRAE and CRVE of 1.0 μm (P = 0.001) and 2.6 μm (P = 0.0002) in the overall population. Changes in retinal vascular calibers depended on baseline level of DR and treatment group (Table 3). Overall, patients with higher levels of DR at baseline developed a higher CRAE (level 10, −1.0 μm; level 20, −1.0 μm; level 35, +6.1 μm; P = 0.001; Fig. 1) and CRVE (level 10, +1.3 μm; level 20, +1.4 μm; level 35, +10.3 μm; P = 0.001; Fig. 2) at follow-up. Even though there was no general difference between treatment groups in the alteration of CRAE (placebo, +1.8 μm; brimonidine, +1.9 μm; somatostatin, −0.2 μm; P = 0.98) or CRVE (placebo, +3.7 μm; brimonidine, +1.3 μm; somatostatin, +2.4 μm; P = 0.78), it was demonstrated that the development of wider retinal vascular calibers in higher levels of DR depended on the treatment regimen; patients treated with placebo did not increase in CRAE (P = 0.17) or CRVE (P =...
0.56) with increasing baseline levels of DR. In contrast, there was a DR-dependent arteriolar widening in patients treated with somatostatin ($P = 0.01$), and a venular dilation in patients treated with brimonidine ($P = 0.048$) and somatostatin ($P = 0.01$).

At follow-up, level of DR improved, remained unchanged and worsened in 28.0% ($n = 83$), 65.7% ($n = 195$), and 6.4% ($n = 19$) patients, respectively. There were no differences in rates of improvement, unchanged conditions, and DR worsening between patients in the different treatment groups (placebo, 28.7% vs. 66.7% vs. 4.6%; brimonidine, 25.9% vs. 63.5% vs. 10.6%; somatostatin, 28.9% vs. 66.4% vs. 4.8%, $P = 0.47$). Of patients with a potential to improve or worsen more than one level of DR from baseline, this was only found in 23.3% (20 of 86 with ETDRS 35 at baseline) and 1.2% (2 of 127 with ETDRS 10 at baseline), respectively. There was no difference in CRAE ($P = 0.51$) or CRVE ($P = 0.75$) when patients were compared according to development in DR during the study.

### DISCUSSION

In a 96-week, prospective, randomized trial evaluating the retinal effect of topical neuroprotective treatment, we concluded that in patients with preexisting mild DR, brimonidine and somatostatin both induced retinal vascular dilation as opposed to placebo.

Our results are in contrast with the earlier conception that topical treatment is inadequate to induce a vascular retinal effect given the corneal barrier and the intraocular distance between the anterior and posterior part of the eye. However,
this concept has been challenged by animal studies as well as a 1-week trial by Tilma and Bek demonstrating retinal arteriolar narrowing in 22 patients with type 1 diabetes treated with topical latanoprost. In the present study, we demonstrate a long-term effect of the retinal vasculature by two independent neuroprotective drugs, even though the study period might not have been long enough to arrest or prevent development of DR. As indicated by the DCCT in type 1 diabetes, this would likely take at least 3 years, which was the reason that EUROCONDOR was designed to study surrogate markers of early vascular and neurogenic dysfunction.

Retinal vascular calibers have been identified as early biomarkers of retinal structural dysfunction in diabetes. In a 16-year prospective study of patients with type 1 diabetes, Broe et al. reported that a 10μm retinal arteriolar narrowing and venular dilation independently predicted long-term proliferative DR, nephropathy, and peripheral neuropathy with odds ratios of 1.4 to 3.0. Likewise, the Wisconsin Epidemiological Study of Diabetic Retinopathy found that a 10μm retinal venular (but not arteriolar) dilation in 4 years, associated with a higher risk of various DR-endpoints in the following 6 years.

### TABLE 3. Development in Retinal Vascular Caliber

<table>
<thead>
<tr>
<th>Level of DR</th>
<th>All (n = 297)</th>
<th>Somatostatin (n = 104)</th>
<th>Brimonidine (n = 85)</th>
<th>Placebo (n = 108)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Central Retinal Arteriolar Equivalent, μm</td>
<td>Placebo (n = 108)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>-0.5 (P = 0.07)</td>
<td>+0.6 (P = 0.30)</td>
<td>-0.6 (P = 0.70)</td>
<td>+0.3 (P = 0.70)</td>
</tr>
<tr>
<td>20</td>
<td>-0.5 (P = 0.1)</td>
<td>+0.3 (P = 0.20)</td>
<td>-0.5 (P = 0.10)</td>
<td>+0.2 (P = 0.20)</td>
</tr>
<tr>
<td>35</td>
<td>0.17 (P = 0.70)</td>
<td>+0.7 (P = 0.0001)</td>
<td>+0.6 (P = 0.0001)</td>
<td>+0.6 (P = 0.0001)</td>
</tr>
</tbody>
</table>

Data presented as median, and level of DR defined according to Early Treatment Diabetic Retinopathy Study scale:

- Placebo (n = 108)
- Somatostatin (n = 104)
- Brimonidine (n = 85)
- All (n = 297)

* Statistically significant (P < 0.05).
† Test for trend of development in central retinal arteriolar and venular equivalent across different levels of DR.
‡ P < 0.17
§ 0.07
¶ 0.01
\* 0.001

demonstrating retinal arteriolar narrowing in 22 patients with type 1 diabetes treated with topical latanoprost. In the present study, we demonstrate a long-term effect of the retinal vasculature by two independent neuroprotective drugs, even though the study period might not have been long enough to arrest or prevent development of DR. As indicated by the DCCT in type 1 diabetes, this would likely take at least 3 years, which was the reason that EUROCONDOR was designed to study surrogate markers of early vascular and neurogenic dysfunction.

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Both of these studies identify retinal vascular dilation as a risk factor for DR progression, which might seem in opposition with our findings of a retinal vascular dilation in response to neuroprotective treatment. However, comparisons are difficult given that most patients in the studies by Broe et al. and Klier et al.24 had more mild DR, whereas our results were found in patients with preexisting mild DR at baseline. In addition, Broe et al.16 only evaluated patients with type 1 diabetes as opposed to type 2 diabetes in the EUROCONDOR.

In the EUROCONDOR, we have previously reported that there was no association between baseline levels of DR and diabetes as opposed to type 2 diabetes in the EUROCONDOR.30 One of the main reasons of the lack of vasodilation is pericyte loss, which may lead to capillary dropout and diabetes can be caused by basement membrane thickening or addition.30 Noticeably, in the present study only patients with preexisting DR developed arteriolar and venular widening, and we demonstrate that this was influenced by the topically administered agent. This is in alignment with the principal findings of the EUROCONDOR which report that neuroprotective treatment was only effective in patients with preexisting retinal neurodysfunction.12 In particular, patients with mild NPDR in both neuroprotective groups had a retinal venular dilation of more than three times of that in the placebo group. This effect size is likely to be clinically relevant. To illustrate, the retinal venular dilation induced by topical neuroprotection in our patients with preexisting mild DR (+13.9–14.3 μm) has the same size as the difference in CRVE between an average 50- and 85-year-old person, in which the former would be 13 μm wider.25

There is no clear mechanistic explanation of the topically induced vascular dilation in patients with moderate NPDR, but it could potentially be given by compensatory increased blood flow to prevent peripheral nonperfusion which is strongly associated with DR progression26 and poor treatment outcome.27 Another potential explanation was proposed by Ludovico et al.28 that demonstrated retinal vasodilation in 50% of patients with early diabetes and suggested that a lack of vasodilation could be a harmful response. While hypoxia induces retinal vasodilation in healthy eyes,29 lack of dilation in diabetes can be caused by basement membrane thickening or pericyte loss, which may lead to capillary dropout and closure.30 One of the main reasons of the lack of vasodilation in nontreated diabetic eyes can be attributed to the impairment of the neurovascular unit, which prevents vasodilation, thus impeding to adapt to the higher metabolic demands that exist in the diabetic retina. Consequently, the neuroactive drugs used in EUROCONDOR may have a favorable effect by improving the neurovascular unit function. This effect results in vasodilation and, therefore, improved autoregulation and delayed capillary dropout. However, with the data at hand, we cannot conclude if treatment-induced retinal vascular dilation in mild DR is a specific consequence of topical neuroprotection with brimonidine and somatostatin, or if this would also be observed with other interventions.

It can be difficult to compare retinal vascular calibers between studies. For instance, Drobnjak et al.31 reported median CRAE and CRVE of 163.5 and 251.0 μm, which were substantially higher than the baseline finding of our study (146.6 and 214.8 μm).32 Unbalances might be explained by differences in ages, populations, methodology, and by the fact that diabetes was only present in a subset of patients in the study by Drobnjak et al.31

Strengths of the present study include the prospective, randomized design and the long follow-up with repeated measurements over time. On the other hand, limitations should be acknowledged. First, we did not include retinal flow measurements which could have provided additional information regarding retinal perfusion. Second, there was a higher than expected dropout which may limit the generalizability of the results. Third, retinal vascular calibers were measured by different graders at baseline and follow-up.

In conclusion, the present study demonstrated retinal vascular dilation induced by long-term topical neuroprotection in 297 patients with type 2 diabetes and mild DR. The concept that eye drops may induce retinal vascular changes would be appealing in order to prevent or delay DR in the early phases, and upcoming prospective studies would be needed to translate this into clinical care.

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References
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APPENDIX

Members of the EUROCONDOR

Scientific Institute San Raffaele, Milan, Italy: Francesco Bandello.

Association for Innovation and Biomedical Research on Light and Image, Coimbra, Portugal: José Cunha-Vaz.

Moorfields Eye Hospital, London, United Kingdom: Catherine Egan.

Vall d’Hebron University Hospital, Barcelona, Spain: José García-Arumí.

University of Aston, Birmingham, United Kingdom: Jonathan Gibson.

University of Liverpool, Liverpool, United Kingdom: Simon Harding.

International Diabetes Federation Europe, Brussels, Belgium: Sehnaz Karadeniz.

University of Ulm, Ulm, Germany: Gabriele Lang.

Hôpital Lariboisière-APHP, Paris, France: Pascale Massin.

University of Padova, Padova, Italy: Edoardo Midena.

BCN Peptides, Barcelona, Spain: Berta Ponsati.

University of Turin, Turin, Italy: Massimo Porta.

Cheltenham General Hospital, Cheltenham, United Kingdom: Peter Scanlon, Stephen Aldington.

Vall d’Hebron Research Institute, Barcelona, Spain: Rafael Simo, Cristina Hernandez.

University of Southern Denmark, Odense, Denmark: Jakob Grauslund, Ulfrik Frydkjaer-Olsen.