Effect of atorvastatin on hard exudates in type 2 diabetic patients with normal lipid profile

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Abstract

Purpose: To evaluate the effect of Atorvastatin on hard exudates in diabetic retinopathy in type 2 diabetic patients with normal lipid profile.

Materials and Methods: Prospective study of 50 patients with clinically significant macular edema involving centre was done over 2 years (2015-16). We excluded patients with type 1 diabetes mellitus, patients with nephropathy/ heart disease, dyslipidemic patients, patients with laser treatment within 3 months of enrollment, patients with proliferative diabetic retinopathy requiring laser treatment. Detailed history was taken and thorough examination was done with +90 D lens for size and extent of clinically significant macular edema and grading of hard exudates. OCT was done for maximum retinal thickness. The outcome measures were visual acuity, change in grade of hard exudates and maximum retinal thickness. Change in maximum retinal thickness by 30% was considered as significant reduction of macular edema. Patient had minimum follow up of 6 months.

Results: Of 50 patients 34 were male and 16 female. Mean follow up period was 10.5±4.0 months. Baseline mean visual acuity was 0.73±0.4 log MAR and baseline mean MRT was 360.03± 30.94 μ. Patients with grade 3 hard exudates were 6, with grade 4 hard exudates were 28 and with grade 5 hard exudates were 16 at baseline. Final mean visual acuity was 0.76±0.4 log MAR. Mean MRT at final visit was 284.6± 48.3 μ. 34 patients (68%) showed improvement by one grade and 2 patients showed improvement by two grade. The grade of hard exudates remained unchanged in 12 patients (24%) and deteriorated in 2 patients. **Conclusion:** Atorvastatin leads to significant improvement in grade of hard exudates in clinically significant macular edema and decrease in maximum retinal thickness on OCT. However visual acuity status was not significant over short term follow up.

Keywords: Atorvastatin, Clinically significant macular edema, Hard exudates.

Introduction

Diabetic retinopathy (DR) & diabetic macular edema are the leading cause of blindness in working age population in both developed and developing countries. According to a study done by Sankara Nethralaya in 2009, the prevalence of diabetic retinopathy in an urban Indian population with diabetes mellitus was 18%. Hyperlipidemia has been linked to the presence of retinal hard exudates in patients with DR and increases the risk of visual loss by 50%. The Early Treatment Diabetic Retinopathy Study (ETDRS) data suggested that lipid-lowering therapy may reduce hard exudates and microaneurysms & helped in preserving vision in patients with clinically significant macular edema (CSME).

Lipid lowering drugs like statins are the standard of care for CSME with dyslipidemia. Statins are 3-hydroxy-3-methylglutaryl coenzyme A (HMG CoA) reductase inhibitors and they also have anti thrombotic, anti inflammatory and anti proliferative properties. Clinical studies have shown beneficial effect of statins on DR. In a study by Gordon et al pravastatin improved the appearance of hard exudates and microaneurysms. Another study reported improvement of visual acuity in diabetic patients with hypercholesterolemia using simvastatin. Later studies showed atorvastatin reduce the severity of hard exudates and microaneurysms in patients with diabetic macular edema. The role of statins in improvement of endothelial function and

ocular blood flow has been emphasized. Atorvastatin improved retinal blood flow velocities in patients with background or proliferative DR.⁹

It was reported that intensive glycemic control and combination treatment of dyslipidemia reduced the rate of progression of DR. ¹⁰ The mechanism, however, seemed to be related to intraretinal lipid transportation rather than serum lipid levels. However, to our knowledge, no study has evaluated the role of lipid-lowering drugs in the management of clinically significant macular edema with normal lipid profile. In the present study, we evaluated the role of Atorvastatin in Clinically Significant Macular Edema (CSME) in type 2 diabetic patients with normal lipid profile.

Materials and Methods

This was a prospective randomised study including 50 metabolically stable diabetic (34 males & 16 females) with CSME involving macular centre (2015-2016). We included eyes with non-proliferative diabetic retinopathy (NPDR) with CSME that presented to retina clinic of our institute which is a tertiary eye care centre. The study was approved by our institute ethical committee. The study included type 2 diabetic patients with CSME with normal lipid profile i.e. total cholesterol < 190mg%, LDL < 115mg%, HDL > 40mg% and serum triglycerides < 180mg% was taken as standard. We excluded patients with type 1 diabetes mellitus, patients with nephropathy/ heart disease,

patients who had been subjected to pars plana vitrectomy for any reason, dyslipidemic patients, patients with laser treatment within 3 months of enrolment, patients with proliferative diabetic retinopathy requiring laser treatment.

Detailed history and thorough ocular examination including visual acuity by Snellen's chart, intraocular pressure by Goldmann applanation tonometer, slit lamp biomicroscopy, fundus evaluation was done with +90 D to look for the extent & location of the hard exudates. Macular OCT was also done for size and extent of CSME. Grading of hard exudates was done as per ETDRS Report No.10.¹¹ Atorvastatin in dosage of 10-20 mg was started in consultation with internist. Patient had follow up at 1 month, at 3 months, at 6 months and thereafter. Patient had a minimum follow up of 6 months.

Results

The study included 50 patients of which 34 were male and 16 were female with type 2 diabetes with normal lipid profile. The Mean age \pm SD was 54.55 ± 7.7 years. Mean follow up period was 10.5 ± 4.0 months.

The outcome was evaluated in terms of visual acuity, change in grade of hard exudate and change in maximum retinal thickness. Visual acuity outcome was

regarded as improvement (two lines gain in VA), stabilization (within two lines of initial VA) or deterioration (loss of two lines). Hard exudates status was regarded as improved, stable or worsened depending on any change in grade of hard exudates. Change in maximum retinal thickness by 30% was considered as significant reduction of macular edema.

Baseline mean visual acuity was 0.73 ± 0.4 log MAR and baseline mean MRT was $360.03\pm30.94~\mu$. Final mean visual acuity was 0.76 ± 0.4 log MAR and mean MRT at final visit was $284.6\pm48.3~\mu$ (Table1). The change in visual acuity from baseline was not statistically significant as p value is 0.92 which is >0.05. The change in maximum retinal thickness from baseline was statistically significant (p value <0.001)

Patients with grade 3 hard exudates were 6 (12%), with grade 4 hard exudates were 28 (56%) and with grade 5 hard exudates were 16 (32%) at baseline (Table 2). 34 patients (68%) showed improvement by one grade (Fig. 1 & 2) and 2 patients showed improvement by two grade. The grade of hard exudates remained unchanged in 12 patients (24%) and deteriorated in 2 patients (Table 3).

Table 1: Showing baseline & at final visit change of VA and maximum retinal thickness

	Baseline	Final Visit	P value
VA (logMAR)	0.73±0.4	0.76±0.4	>0.05
MRT (µm)	360.03± 30.94	284.6 ± 48.3	< 0.001

Table 2: Showing grading of hard exudates at baseline

Grading of Hard Exudates	Baseline (n)		
Grade 3	6 (12%)		
Grade 4	28 (56%)		
Grade 5	16 (32%)		

Table 3: Showing status of hard exudates at final visit

Status of Hard Exudates	Final visit (n)
Improved	36
Stabilized	12
Worsened	2

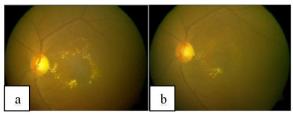


Fig.1: Fundus photograph L/E of patient after 20 weeks atorvastatin therapy showing reduction in the macular edema as well as improvement in grade of hard exudates from grade 4 to grade 3.

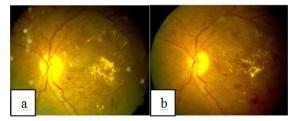


Fig. 2: Fundus photograph L/E of patient showing reduction in the macular edema as well as improvement in grade of hard exudates from grade 5 to grade 4 after 4 months of oral atorvastatin

Discussion

HMG Co A reductase inhibitors have been established as safe and efficacious drugs in management of diabetic retinopathy with dyslipidemia. Atorvastatin has beneficial effect in decreasing hard exudates severity and clinically significant macular edema in diabetics with dyslipidemia. ^{12,13} The present study was to evaluate the role of atorvastatin in diabetics with normal lipid profile. Lipid exudation at the macula in diabetic patients is the result of increased vascular permeability and dysfunctional outer bloodretina barrier and retinal pigment epithelium. ¹⁴ The Action to Control Cardiovascular Risk in Diabetes

(ACCORD) eye studies demonstrated that intensive control of dyslipidemia using combined simvastatin–fenofibrate treatment can significantly (40%) delay the progression of DR in patients with Type 2 diabetes. ¹⁵

A study by Gupta et al reported dramatic regression of hard exudates after correction of dyslipidemia in diabetics with severe hard exudates over a mean follow up of 18 weeks. Thus, lipid-lowering drugs preserve vision and reduce the risk of hard exudates in clinically-significant macular edema (CSME) in diabetics with an abnormal lipid profile. But their role in reducing CSME in diabetics with a normal lipid profile is not yet known. Our study showed that atorvastatin leads to significant decrease in grade of hard exudates & also decreases maximum retinal thickness in type 2 diabetic patients with normal lipid profile.

Another study by Narang et al reported that atorvastatin does not affect treatment outcome in CSME over a short term follow-up. Results were based on visual acuity only. OCT and hard exudate grading was not done in that study as in our study. ¹⁶

Michael Cusick et al showed histopathological evidence of regression of hard exudates and macular edema after reduction of elevated serum lipid levels in dyslipidemic patients. Now-a-days trends are shifting towards use of lipid lowering drugs as primary preventive measure for coronary artery disease in type II diabetics with normal lipid profile. Due to the histopathological similarity between diabetic retinopathy and coronary artery disease, atorvastatin could also have a role to play in diabetic retinopathy with normal lipid profile.

The results of our study suggest that lipid-lowering therapy in patients with diabetes may be a useful adjunct in the management of diabetic macular edema with severe hard exudates even in patients with normal lipid profile.

Conclusion

Atorvastatin leads to significant improvement in grade of hard exudates in clinically significant macular edema and decrease in maximum retinal thickness on OCT. However visual acuity status was not significant over short term follow up.

Conflict of Interest: None

Source of Funding & Support: Nil

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