Original Research Article

EVALUATION OF INTRAVITREAL BEVACIZUMAB FOR ITS SYSTEMIC SIDE EFFECT THROMBOSIS AFTER CHRONIC ADMINISTRATION

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ABSTRACT:

Bevacizumab targets Vascular endothelial growth factor-A (VEGF-A). Bevacizumab specifically binds to the VEGF-A protein, thereby inhibiting the process of angiogenesis. Thrombosis and hypertension are the major systemic side effects of bevacizumab. As thrombosis and hypertension are the major systemic side effects of bevacizumab, whether this drug could predispose a patient to thrombosis or not after intravitreal administration, as it is absorbed even through intravitreal administration. We determined the safety of the drug. This study was conducted at Al Ibrahim eye hospital for 3 months. The drug was intravitreally administered by Professor Dr. P.S. Mahar. For this 10 patients were administered three doses of intravitreal bevacizumab at monthly interval and followed for chronic effects of the drug. Blood samples were taken to determine fibrinogen level, platelet count, prothrombin time (PT), activated partial thromboplastin time (APTT) and sodium level by kit method. Blood pressure was also monitored of all the patients before and after the drug administration. There has been a significant decrease seen in fibrinogen level. Non-significant rise in the PT. Platelet counts decrease insignificantly. Slight increase is noted in sodium level. Slight increase is noted in diastolic blood pressure whereas systolic blood pressure is insignificantly increased. Thus, results of our study indicate that there may be a bleeding tendency after bevacizumab so careful monitoring is required in patients receiving this drug, as well as monitoring of blood pressure is required.

KEYWORDS: Bevacizumab, thrombosis, intravitreal.

INTRODUCTION:

Bevacizumab is a humanized monoclonal antibody that targets Vascular endothelial growth factor-A (VEGF-A), an isoform of Vascular endothelial growth factor (VEGF) that is responsible for stimulating abnormal blood vessel growth and blood vessel leakage in diseases like diabetic retinopathy and retinal vein occlusion. Its role in the cause and progression of choroidal neovascularization in neovascular age-related macular degeneration has become increasingly important, hence by specifically blocking VEGF, there is reduction in pathological angiogenesis.

Four major steps of angiogenesis have been recognized; namely vasodilation and hyper permeability, vessel destabilization and matrix degradation, endothelial cell proliferation and migration, lumen formation and vessel stabilization. Angiogenesis is a complex biological process involving a delicate balance and
interplay between a variety of molecular angiogenic and angiostatic factors. The VEGF-A is believed to be prime regulator of angiogenesis and takes part in all four stages of angiogenesis [1]. Intravitreal bevacizumab has been described in some recent articles for treatment of proliferative diabetic retinopathy complicated by vitreous hemorrhage [2], macular edema in central retinal vein occlusion, and neovascular age-related macular degeneration [3]. Invivo studies revealed that in human nonproliferative diabetic retinopathy there was increased expression of VEGF-A [4]. Upregulation of VEGFR-1, VEGFR-2 and VEGFR-3 was also observed [4,5].

The role of VEGF-A in the development of choroidal neovascularization has been established. Increased VEGF-A expression was noted in surgically excised choroidal neovascularization, retinal pigment epithelium and vitreous of age-related macular degeneration [6].

The permeability inducing actions of VEGF appear to induce the development of macular edema. VEGF may cause macular edema that accompanies diabetic retinopathy, retinal vein occlusion and uveitis. Human eyes with macular edema secondary to uveitis, retinal vein accession and diabetic retinopathy have shown increased retinal levels of VEGF [7]. When VEGF was blocked, blood-retinal barrier breakdown could be both prevented and reversed through VEGF inhibition.

Bevacizumab (Avastin) is full length, recombinant humanized Anti-VEGF monoclonal antibody that binds to all isoforms of VEGF-A. It binds directly to the VEGF ligand (which is expressed by both normal and tumor cells) to prevent its interaction with receptors on the surface of endothelial cells, thereby inhibiting the biologic activity of VEGF as observed in invitro and invivo assay systems [8].

Angiogenesis is an essential process in tumor development [9,10]. The VEGF ligand is the predominant regulator of tumor angiogenesis [10]. Avastin directly targets the VEGF ligand to specifically inhibit angiogenesis [11]. Maintaining VEGF ligand inhibition may prevent tumor vessel regrowth over time [12,13].

It is the FDA’s approved Anti-VEGF agent in the treatment of patients with metastatic colorectal cancer [14], lung cancer, metastatic renal cell cancer FDA, 2009), glioblastoma multiforme [15]. It has shown promising result, through intravitreal route in the treatment of neovascular ocular diseases like choroidal neovascularization, macular edema and diabetic retinopathy [16,17].

As thrombosis and hypertension are the major systemic side effects of bevacizumab whether this drug could predispose a patient to thrombosis or not after intravitreal administration, as it is absorbed even through intravitreal administration. We determined the safety of the drug.

Since no work has been carried out before on this issue in our local setup, this study would be important in decision making regarding the safety of intravitreal bevacizumab in choroidal and retinal neovascular disorders.

MATERIAL AND METHODS:

In order to evaluate the systemic side effects of intravitreal bevacizumab total 10 patients were selected. Patients were selected from Al Ibrahim eye hospital, meeting the inclusion and exclusion criteria receiving Bevacizumab (Avastin) injection. The purpose, procedure, risks and benefits of the study was explained to the patients. Informed consent was taken. In operation theater before injecting the intravitreal bevacizumab, eye ball was anesthetized with topical proparacaine drops sterilized with povidone iodine 5%.
Inclusion criteria:
Patients having neovascularization in retina and choroid of eye like diabetic retinopathy and age related macular degeneration diseases including both males and females having age above 40 years. Hypertensive and non hypertensive, diabetic and non diabetic patients were included.

Exclusion criteria:
Patients having any hematological disorders like thrombosis. History of previous Bevacizumab injection.

Drug:
Patients received the drug Bevacizumab. This drug was intravitreally administered by Professor Dr. P.S. Mahar. The dose of intravitreal bevacizumab administered to the patient is 1.25mg in 0.05ml with 1ml syringe.

Experimental protocol:
These 10 patients were administered three doses of intravitreal bevacizumab at monthly interval and followed for chronic effects of drug. Blood samples were taken before and after one week of drug administration to determine fibrinogen level, platelet count, prothrombin time (PT), activated partial thromboplastin time (APTT) and sodium level by kit method. Blood pressure was also monitored of all the patients before and after the drug administration.

Sample collection:
Blood samples were collected in three types of tubes:
BD vacutainer EDTA.K2 tubes for collection of pure blood to study hematological parameter like platelet count.
Bio vac 3.2% sodium citrate tubes for collection of plasma to study coagulation parameters like PT and APTT and fibrinogen level. Prothrombin time and activated partial prothrombin time was checked manually using standard reagent kits of DiagnosticaStago France.
B-ject Gel clot activator vaccum tube for collection of serum to study sodium level.
After the collection of samples 3 ml of blood n 3.2% sodium citrate tubes, plasma was separated out and 4 ml of blood in Gel tubes, Serum was separated out, by centrifuging the blood samples in 800 centrifuge machine (china) at 4000 RPM for 5 minutes. The separated plasma and serum was stored in 2-8°C and within3hours all the coagulation and electrolyte estimation were performed.
The estimation of platelet was directly done on automated MS4E Vet Hematology analyzer model# 3MSR0214 by MeletSchloesing Switzerland.
After separating the plasma the fibrinogen level was measured by humaclot duo (coagulation analyzer, model# 18650) (Human Germany) using standard reagent kits of Human Germany. Hemostat fibrinogen (manual and automated determination of plasma fibrinogen) was used[18].
Neoplastin CI Plus was used for the determination of prothrombin time [19]. C.K.Prest was used for the determination of Kaolin-activated partial thromboplastin time [20, 21].
After separating the serum, the sodium level was measured by Microlab 300 semi-automated clinical chemistry analyzer by Merck Netherlands.
RESULT

**GRAPH-1:** Effect of Bevacizumab on fibrinogen level

**GRAPH-2:** Effect of Bevacizumab on prothrombin time

**GRAPH-3:** Effect of Bevacizumab on activated partial prothrombin time

**GRAPH-4:** Effect of Bevacizumab on platelet count
GRAPH-5: Effect of Bevacizumab on systolic blood pressure

Comparison of pre and post systolic blood pressure after chronic dosing of bevacizumab

GRAPH-6: Effect of Bevacizumab on diastolic blood pressure

Comparison of pre and post diastolic blood pressure after chronic dosing of bevacizumab

GRAPH-7: Effect of Bevacizumab on sodium level

Comparison of pre and post sodium level after chronic dosing of bevacizumab
DISCUSSION

Scutz in the study reported that Bevacizumab treatment is associated with significant increase in the risk of arterial thrombosis[22]. In the present study, determination of fibrinogen level, platelet count, prothrombin time (PT), activated partial thromboplastin time (APTT) and sodium level was conducted by kit method. Blood pressure was also monitored of all the patients before and after the drug administration.

There has been significant decrease in the fibrinogen levels seen after chronic administration. Fibrinogen is a key protein in the coagulation pathway, interacting in multiple processes of platelet aggregation, clot formation and wound healing, and contributing to the final step of the coagulation cascade[23]. Low level of fibrinogen may lead to predisposition to bleeding. Thus if a patient receiving intravitrealbevacizumab have any history of thrombosis should receive chronic dosing of intravitrealbevacizumab.

There has been some rise in the PT but it is non-significant. Decreased level of fibrinogen and platelet count may be causing this rise of PT. The results of our study indicate that there is slight risk of bleeding after using intravitrealbevacizumab in neovascularization of choroid and retina. APTT is non-significantly decreased after the injection in chronic stages.

Platelet counts decrease after the injection. Platelets are meant for maintaining hemostasis. They help in wound healing and prevent bleeding. One physiological function of platelets may be to act as scavengers of circulating VEGF to restrict angiogenic activity to sites of wound healing. This may also be occurring in the pathological situation of neovascularization [24].

Slight increase is observed in sodium levels after the injection. Slight increase is noted in diastolic blood pressure. Insignificant increase is seen in systolic blood pressure. This increase in blood pressure may be associated with elevated level of sodium. Raiser et al, study supports our results, reporting that there is a risk of dysregualtion of blood pressure or persistence of hypertension in hypertensive patients after intravitrealbevacizumab injections [25].

Mourad et al.,(2008) reported that bevacizumab treatment resulted in endothelial dysfunction and capillary rarefaction; both changes are closely associated and could be responsible for the rise in blood pressure observed in most patients [26].

The results of our study indicate that there may be bleeding tendency after bevacizumab because fibrinogen levels and platelet counts have decreased after chronic treatment, so careful monitoring is required in patients receiving this drug, as well as monitoring of blood pressure is required. In our study intravitreal administration of bevacizumab produced a slight increase in systolic and diastolic blood pressure. This work could be further extended to reassess the results with an increased sample size and also determine the effects in complicated patients having secondary problems.

REFERENCES: