Bevacizumab Reverses Need for Liver Transplantation in Hereditary Hemorrhagic Telangiectasia

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Abbreviations: A, ascites; HA, hepatic artery; HHT, hereditary hemorrhagic telangiectasia; LVED, left ventricular end-diastolic; LVES, left ventricular end-systolic; MR, magnetic resonance; PV, portal vein; VEGF, vascular endothelial growth factor.

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hemoptysis, whereas cerebral and spinal malformations may present with seizures, hemorrhage, headaches, and neurological deficits.

Currently, treatment of patients with hepatic HHT is primarily symptomatic without alteration of the underlying pathological neovascularization. Hepatic arterial embolization or occlusion is associated with significant morbidity and mortality and is not recommended outside palliative circumstances. Liver transplantation is curative but involves an 18% risk of acute mortality as well as problems of long-term immunosuppression. We present a case of a patient with severe HHT who responded dramatically to treatment with bevacizumab, a vascular endothelial growth factor (VEGF) antibody, which obviated the need for liver transplantation.

CASE REPORT

A 47-year-old woman was referred for assessment for liver transplantation due to HHT complicated by high-output cardiac failure, portal hypertension, cholestasis, and malnutrition. She had been diagnosed with HHT as a teenager and fulfilled consensus diagnostic criteria requiring the presence of epistaxis, telangiectasia on her lips and oral cavity, visceral involvement of her liver, and a family history. Her mother had died at the age of 56 of liver failure secondary to HHT, her brother and sister suffered epistaxis, and a cousin had received a liver transplant for HHT. After her diagnosis, she had received periodic iron infusions for anemia but had otherwise been well and was working full-time.

Over a 6-month period, she developed worsening high-output cardiac failure, ascites, and bilateral pleural effusions that became diuretic-resistant and required large-volume paracentesis. She had progressive lethargy and a reduction in her performance status such that she was unable to walk more than 30 m or stand for a prolonged length of time. This was associated with a progressive loss of weight and muscle mass, with her dry weight body mass index falling from 21.7 to 17.9 kg/m². Serum biochemistry demonstrated progressive cholestasis (Fig. 1). Liver magnetic resonance (MR) angiography confirmed an enormous liver (4807 mL) and massive arteriovenous malformations (Fig. 2A). Cardiac MR demonstrated severely dilated atria and ventricles and a high resting cardiac output (Table 1) despite beta-blockade. Pulmonary and intracranial vascular malformations were not present on computed tomography or MR scans. Endoscopy demonstrated

![Figure 1. Dramatic improvement of cholestatic liver enzymes with bevacizumab. Bilirubin is represented by a continuous line; alkaline phosphatase is represented by a broken line.](image1)

![Figure 2. Magnetic resonance scans with intravenous gadolinium before and 6 months after bevacizumab, demonstrating a marked reduction in hepatic size and vascularity and resolution of ascites. Abbreviations: A, ascites; HA, hepatic artery; PV, portal vein.](image2)

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grade II esophageal varices. Because of her poor cardiac status and malnutrition, a bridge therapy was felt necessary. After informed consent was obtained from the patient, 6 courses of bevacizumab (5 mg/kg) were given over a 3-month period.

An immediate improvement in her cholestatic liver enzymes was observed (Fig. 1), followed by a dramatic improvement in her clinical state. Over the course of 3 months, her cardiac failure, ascites, and pleural effusions resolved clinically and radiologically (Fig. 2B). This was associated with improvement in her nutritional status and a 10% increase in dry weight. By 6 months, all symptoms had resolved, and she was able to cease all medications and return to full-time work. She was subsequently delisted as a transplant candidate. Serial MR liver and cardiac scans (Table 1) confirmed a marked diminution of hepatic vascularity, a 2-fold reduction in liver volume, and normalization of cardiac output. Her previous grade II esophageal varices were markedly small and barely visible on follow-up endoscopy. She remains well 6 months after completing treatment.

**DISCUSSION**

VEGF is a key regulator of angiogenesis and is upregulated in a variety of disease states including malignancy, ocular disease, and inflammatory conditions. Vessels formed by VEGF are disorganized and tortuous, similar to those observed in patients with HHT. Not surprisingly, serum and tissue expression of VEGF is increased in patients with HHT and correlates with microvessel density. The genetic mutations of HHT affect endoglin (HHT type 1), activin receptor-like kinase 1 (HHT type 2), and the SMAD 4 gene. These genes are involved in the transforming growth factor-β signaling pathway, which is a potent stimulator of VEGF production. Thus, blocking VEGF represents a logical therapeutic target for HHT.

Bevacizumab is a humanized recombinant monoclonal antibody against VEGF. It induces the arrest of endothelial cell proliferation, thereby preventing vessel growth and causing the regression of existing vessels by increasing endothelial cell death. Bevacizumab is effective in diseases of abnormal angiogenesis such as non–small-cell lung cancer and metastatic colorectal cancer, where it improves survival. Furthermore, it improves visual acuity in age-related macular degeneration by reducing choroidal neovascularization.

When given in combination with pemetrexed for treatment of malignant mesothelioma, bevacizumab was noted to reduce the severity of epistaxis and transfusion requirements in 1 patient who had concomitant HHT. The case that we describe here is the first in which bevacizumab has been used to deliberately ameliorate life-threatening complications of HHT. Complete resolution of all symptomatic manifestations was observed and objectively quantified. Therapy with bevacizumab was well tolerated with minimal toxicity. In contrast, the current treatment options for advanced symptomatic hepatic HHT of hepatic arterial ligation or liver transplantation carry significant risk of morbidity and mortality.

In conclusion, bevacizumab induced a strong biological and clinical response in this patient with symptomatic hepatic HHT. Biological agents targeting angiogenic growth factors represent a novel form of treatment for patients with HHT with the potential to alleviate disease-related morbidity and mortality. The impressive response of this case mandates the further use of such agents in a clinical trial setting.

**REFERENCES**


**TABLE 1. Reduction in Liver Volume and Cardiac Output with Bevacizumab**

<table>
<thead>
<tr>
<th>Time</th>
<th>Courses of Bevacizumab (5 mg/kg)</th>
<th>Liver Volume (mL)</th>
<th>LVED Volume (mL)</th>
<th>LVES Volume (mL)</th>
<th>Cardiac Output (L/minute)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0 months</td>
<td>0 courses</td>
<td>4807</td>
<td>192</td>
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<td>3 months</td>
<td>6 courses</td>
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<td>2269</td>
<td>182</td>
<td>58</td>
<td>5.1</td>
</tr>
</tbody>
</table>

*Abbreviations: LVED, left ventricular end-diastolic; LVES, left ventricular end-systolic.*