

Informed Consent for Photodynamic Therapy Treatment

You have been given information about your condition and the recommended surgical, medical or diagnostic procedure(s) to be used. This consent form is designed to provide a written confirmation of such discussions by recording some of the more significant medical information given to you. It is intended to make you better informed so that you may give or withhold your consent to the proposed procedure(s).

Nadeem N. Vaidya, M.D.

Condition: The doctor above has explained to me that the following condition(s) exist in my case:

- Exudative Age Related Macular Degeneration
- Presumed Ocular Histoplasmosis Syndrome
- Idiopathic Choroidal Neovascularization
- Myopic Degeneration
- Central Serous Retinopathy

Proposed Procedure(s): I understand that the procedure(s) proposed for evaluating and treating my condition is Photodynamic Therapy with Verteporfin of the:

- OD (right eye)
- OS (left eye)

Risks/Benefits of Proposed Procedure(s):

a. Just as there may be benefits to the procedure(s) proposed, I also understand that medical and surgical procedures involve risks. These risks include allergic reaction, bleeding, blood clots, infections, adverse side effects of drugs, blindness, and even loss of bodily function or life, as well as risks of transfusion reactions and the transmission of infectious disease, including Hepatitis and Acquired Immune Deficiency Syndrome, from the administration of blood and/or blood components.

b. I also realize that there are particular risks associated with the procedure(s) proposed for me and that these risks include, but are not limited to: *Pain, Inflammation, Infection, Local Reactions at the Site of the Injection, Extravasation of Photosensitive Dye, Skin Necrosis, Severe Sunburn, Temporary Vision Changes, Loss of Vision, and Loss of Eye.*

Cautions: I also understand that if I have *Porphyria, Severe Liver Disease or are allergic to Porphyrin* that I must inform my physician. Also I understand if I take Calcium Channel Blockers, or Polymyxin B that these medications may increase the concentration of the verteporfin dye. Additionally Tetracycline, Sulfonylurea, and diuretics may increase the risk of skin reactions. And β -carotene, Ethanol and Mannitol reduce the activity of verteporfin.

Complications; Unforeseen Conditions; Results: I am aware that in the practice of medicine, other unexpected risks or complications not discussed may occur. I also understand that during the course of the proposed procedure(s) unforeseen conditions may be revealed requiring the performance of additional procedures, and I authorize such procedures to be performed. I further acknowledge that no guarantees or promises have been made to me concerning the results of any procedure or treatment.

Acknowledgments: The available alternatives, some of which include: *no treatment, intra/periocular injections, or incisional surgery*, the potential benefits and risks of the proposed procedure(s), and the likely result without such treatment: *loss of vision*, have been explained to me. I understand what has been discussed with me as well as the contents of this consent form, and have been given the opportunity to ask questions and have received satisfactory answers. I understand that *I must avoid excessive sun exposure or bright light during the first 24 (Twenty Four) hours following treatment.*

Consent to Procedure(s) and Treatment: Having read this form and talked with the physicians, my signature below acknowledges that: I voluntarily give my authorization and consent to the performance of the procedure(s) described above (including the administration of blood and disposal of tissue) by my physician and/or his/her associates assisted by hospital personnel and other trained persons as well as the presence of observers.

Patient or Guardian _____ Date _____