

DEPARTMENT OF HRALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administrat 9200 Corporate Boulevard Rodevilla MD 20850

JAN 20 1999

Herbert J. Nevyas, M.D. Nevyes Eye Associates Delaware Valley Laser Surgery Institute 333 City Line Avenue Bala Cynwyd, PA 19004

G970088/S15 Re:

Sullivan Excimer Laser System (Nevvas Model)

Indications for Use: LASIK (Laser-Assisted In Sim Kerstomileusis) to correct myopia of -0.5 to -15 Diopters (D) with up to -7 D of astigmatism for protocol NEV-97-001 Myopia; and, LASIK retreatment to correct myopia and myopic astigmatism of eyes treated with this laser prior to IDE approval

Dated: January 5, 1999 Received: January 6, 1999 HCFA Category:

Next Annual Report Due: August 7, 1999

Dear Dr. Nevyas:

The Food and Drug Administration (FDA) has reviewed the supplement to your investigational device exemptions (IDE) application providing validation data for the contrast sensitivity study. You have corrected the deficiency cited in our September 24, 1998 conditional approval letter. Your application is approved, and you may continue your investigation at the institution enrolled in your investigation where you have obtained institutional review board (IRB) approval. Your investigation is limited to one institution and 1015 subjects (2030 eyes); 990 subjects (1980 eyes) for myopia (-0.5 to -15 D with up to -7 D astigmatism); and, 25 subjects (50 eyes) for enhancements of subjects treated prior to IDE approval (-0.5 to -15 D myopia with up to -7 D astigmatism).

Please be aware of the following:

In Table 1-1, the data appear to be quite scattered, with some subjects actually increasing in sensitivity during glare (e.g., see BC & CB at 3 cycles per degree (CPD)), while others are severely compromised (see ZM). In order to reduce variability in the data in the contrast sensitivity study, the person administering the test should have experience in this test and the subjects should be well trained prior to testing.