The photosensing processes in *Stentor coeruleus* and *Blepharisma japonicum* can be described within the photo-signal transduction cascade shown above, with stentorin and blepharismin serving as the photosensor molecules, respectively. In the photophobic responses of these ciliates, light signal (red wavelength light and intensity gradient) is perceived by the photosensor molecules localized in the pigment granule. A signal transduction then leads to the reversal of the direction of ciliary beating that is entailed in the stop-turning motion of the cell away from the source of light. In this paper we briefly describe the photosensory transduction which utilizes stentorin and blepharismin as the photosensor molecules.

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Period: February 1991 - February 1995

Title: Photo-Signal Transduction in Motile Ciliate Blepharisma

Contract or Grant Number: DAAL03-91-G-0061

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# Table of Contents

- Statement of the Problem Studied .................................................. 3  
- Summary of the Most Important Results ........................................... 3  
- List of Publications and Technical Reports ..................................... 5  
- Participating Scientific personnel ................................................... 6  
- Inventions ......................................................................................... 6
Statement of the Problem Studied:

**Title:** Photo-Signal Transduction in Motile Ciliate *Blepharisma*

The single cell ciliate, *Blepharisma japonicum*, is responsive to a sudden increase in light intensity and is capable of discriminating colors of different wavelengths ("color vision"). This remarkable single cell vision is mediated by the photosensor molecule called blepharismin. The objectives of the research conducted included:

1. Elucidation of the chemical structure of the photosensor molecule
2. Study of its photochemical mechanism of action
3. Exploration of the light signal transduction cascade triggered by the light excitation of the photosensor molecule

Summary of the Most Important Results:

During the three year period of research supported by the Army Research Office grant, the following important results have been obtained to solve the problem outlined above:

1. The chemical structures of the two forms of the *Blepharisma* photosensor molecule, red and blue blepharismin, have been elucidated by means of UV-visible spectroscopy, NMR, FTIR and mass spectrometry. We have also determined the structure of stentorin, the photosensor molecule in the closely related ciliate *Stentor coerules*. Both blepharimins and stentorin represent a structurally unique class (hypericin derivatives) of the biological light receptor molecules that have been added to the select few photosensors that are known up to this point (i.e., rhodopsin, phytochrome, and blue light cryptochromes, based on retinal, tetrapyrrole and flavin as the chromophore molecules, respectively).
(2) The primary photochemical mechanism of the ciliate photosensor molecules appears to be electron transfer coupled with proton release. This reaction occurs in a few picosecond time scale in both blepharismin and stentorin. The resulting intracellular pH change serves as an initial physiological signal.

(3) The light signals in terms of intensity gradient and different wavelength colors are transduced and amplified in the cells via a transducin-like G-protein coupled to a cGMP-dependent phosphodiesterase. Activation of the phosphodiesterase enzyme depolarizes the cell membrane by opening the voltage-dependent calcium channels. The calcium influx results in a transient reversal in ciliary stroke, thus eliciting the photomotile response in *Blepharisma japonicum* and *Stentor coeruleus*. 
List of all publications and technical reports published:


Participating Scientific personnel:

Pill-Soon Song, Principal Investigator (1991-1995)
Stanislaw Fabczak, Postdoctoral Fellow (1990-1992)
Hanna Fabczak, Postdoctoral Fellow (1990-1992)
Jae Seok Hyon, Postdoctoral Fellow (1991-1992)
Nengbing Tao (Ph.D. degree in 1994)
Renke Dai (Ph.D. degree in 1994)
Phun Bum Park (Ph.D. degree expected 1995)
Susanne Meza-Keuthen (M.S. degree in 1993)
Mark Looyenga (M.S. degree in 1994)
Gregg Timm, Undergraduate Assistant (1993)

Inventions:

No patents planned or pending