

1976 #8a

Indicate below the numbers and titles of sessions in which your abstract might be programmed (see Topic Category List); CAREFUL SELECTION IS CRITICALLY IMPORTANT.

1st = 031 Title Microcirculation  
2nd = 033 Title Peripheral Circulation

(See paragraph below for the number of photocopies to be submitted with abstracts.)

# Unified 1976 FASEB Abstract Form

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Mail to:  
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## PRESENTATION PREFERENCE

Preferred choice (CHECK ONE ONLY)

- ☐ Poster presentation  
☒ Slide presentation  
☐ Indifferent

16 mm. films (silent or optical sound) are permitted if essential to 10-minute session presentation.

MOVIE YES..... NO..X.....

Submit justification by letter to Society Office, with abstract.

## IMPORTANT:

See sample abstracts, typing and mailing instructions on reverse side; use enclosed Check List for preparation of abstract.

PULSATILITY OF MICROVASCULAR BLOOD VELOCITY. H.N. Mayrovitz\*, R.T. Tuma,\* and M.P. Wiedeman, Dept. of Physiology, Temple University School of Medicine, Phila., Pa. 19140

Red blood cell velocity has been measured in the microvasculature of the unanesthetized mammal, (little brown bat) to determine the magnitude of pulsatile effects produced by the heart. Velocity measurements were made in wing vessels belonging to five different branching orders (diameter ranges; 42-71, 18-64, 14-17, 5-7, and 3.2-6 microns) using a modification of the dual slit method and crosscorrelating the resultant photoptic signals. To evaluate the extent of damping of the pulsatile component, the data were analyzed in terms of the ratio ( $\gamma$ ) equal to the pulsatile velocity component to the average velocity at each branching order. The results show a trend for selective damping of the pulsatile component as evidenced by a decrease in  $\gamma$  with increasing branching order. A tendency for the damping to increase with heart rate was also found. In spite of the damping however,  $\gamma$  values as large as 0.3 are found in third and fourth order vessels. The finding of pulsatile component synchronous with the heart rate of this magnitude within the terminal branches of the microvasculature implies that microcirculatory analyses and data interpretation (e.g. shear stress, flow dynamics, vasomotion) in which blood velocity plays an essential role, must include the effect and significance of such pulsatility. (Supported in part by SCOR Thrombosis Grant #HL 14217).

All compounds that are designated by code or initial letters must be identified adequately in the abstract, e.g., MJ-1999: 4-(2-isopropylamino-1-hydroxyethyl) methanesulfonamide hydrochloride.

Each Abstract Form submitted MUST BE SIGNED by a member of the Society to which the abstract is sent.\*

Man P. Wiedeman  
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\*See the enclosed unified rules for eligibility of papers.

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