#### How I write a scientific paper:

Selling your data with power writing.

by

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A talk presented locally, invited at other University and at National Scientific meetings.

- I. What is a paper?
- II. Parts
  - A. Title
  - B. Abstract
  - C. Introduction
  - D. Materials and Methods
  - E. Results (& Discussion)
  - F. (Discussion)
  - G. Acknowledgments
  - H. References
  - I. Tables
  - J. Figures and Captions
- III. Where do I Begin?
  - A. Data, then
  - B. Everywhere
- IV. Data
  - A. The heart of the work!
  - B. What is your point? or, What hypothesis was tested?
- V. The parts in detail
  - J. Figures and Captions They should tell the story; Think of final size in journal Caption should allow figure to stand alone
  - I. Tables
  - A. Title
    - Primary key words
  - B. Abstract Secondary key words Packed with concrete information
  - C. Introduction Only information that points to your point, hypothesis
  - D. Materials and Methods Can another person repeat exactly what you've done?
  - E. Results (& Discussion) Lead reader to your point, interpretation, conclusions
  - F. (Discussion)
  - G. Acknowledgments Support, Funding reports.

- H. References All as appropriate
- VI. The Writing
  - A. Write to the reader's expectations!
  - B. Write with emphasis.
  - C. Write for mind control
  - D. Write to persuade the reader.
- VII. How?
  - A. At the document level
    - 1. Issue, topic
    - 2. persuasive discussion
    - 3. Coda, finale, conclusion
    - 4. Chaining
  - B. At the paragraph level
    - 1. Issue, topic
      - 2. persuasive discussion
      - 3. Coda, finale, conclusion
    - 4. Chaining
  - C. At the sentence level
    - 1. Bold, with emphasis on point
    - 2. Simple and direct
    - 3. Chaining
      - a. old-p --> new-1 (Issue)
      - b. old-1 --> new-2
      - c. old-2 --> new-3
      - d. old-3 --> new-4
      - e. old-4 --> new-5
      - Χ. ...
      - z. Coda, finale, conclusion, emphasis
- VII. Concrete Examples
  - 1. Jurkiewicz BA; Buettner GR.(1996) Photochem. Photobiol. **64:**918-922
  - Kelley EE; Buettner GR; Burns CP. (1997) Photochem. Photobiol. 65:576-580.
- VIII. Finis

# Types of Writing Creative Persuasive, narrative writing

Who is the audience?

- Technical Experts?
- Knowledgeable in area?
- Broader Area?
- All Scientists?
- All Readers?

3

# I. What is a paper?

# A means to publish data.

# DATA

#### Interpretation

**Definition:** An acceptable primary scientific publication must be the first disclosure containing sufficient information to enable peers:

- (1) to assess observations,
- (2) to repeat experiments, and
- (3) to evaluate intellectual processes;

moreover, it must be susceptible to sensory perception, essentially permanent, available to the scientific community without restriction, and available for regular screening by one or more of the major recognized secondary services (*e.g.* currently, Biological Abstracts, Chemical Abstracts, Index Medicus, Excerpta Medica, Bibliography of Agriculture, etc., in the United States and similar facilities in other countries).<sup>1,2</sup>

<sup>&</sup>lt;sup>1</sup> Council of Biology Editors. (1968) Proposed definition of a primary publication. *Newsletter, Council of Biology Editors.* November, p. 1-2.

<sup>&</sup>lt;sup>2</sup> Day, Robert A. (1983) *How to Write and Publish a Scientific Paper*. iSi Press, Philadelphia, PA. p. 2.

# II. Parts

- A. Title
- **B.** Abstract
- C. Introduction
- **D. Materials and Methods**
- E. Results (& Discussion)
- F. (Discussion)
- G. Acknowledgments
- H. References
- I. Tables
- J. Figures and Captions

# III. Where do I Begin?

# A. Data, then

## **B. Everywhere**

# IV. Data

# A. The heart of

# the work!

# B. What is your point? or, What hypothesis was tested? What problem was addressed

# **J. Figures and Captions**

- They should tell the story; Story board
- 2. Think of final size in journal See Kelley figures.
- 3. Caption should allow Figure to stand alone.
- 4. Informative rather than descriptive title.

Examples:

# Kelley et al. Figure 1

Jurkiewicz Figures 1 & 3

Figure 4



Figure 5. PhGPx inhibits <sup>1</sup>O<sub>2</sub>-induced lipid-derived radical generation.<sup>3</sup>

A. Cells were pretreated with 6 μg/mL Photofrin for 24 h in full media. After 5-min light exposure, cells were incubated for 6 h in the dark. Radical formation was then assessed using EPR spin trapping. Each spectrum represents the signal-averaged result of 45 scans. The height of the low-field line of the DMPO/lipid-derived radical adduct was used as a measure of radical yield. Inset: An example X-band EPR spectrum of the DMPO adduct (a<sup>N</sup> = 15.2 G, a<sup>H</sup> = 10.2 G in ethyl acetate) formed from MCF-7 cells. Control cells were incubated with Photofrin but not exposed to light. The yield of DMPO adduct in the control samples was below the limit of detection (data not shown).

(Mean  $\pm$  SE, n = 3, \* p< 0.05, \*\* p< 0.005 compared to Wt.)

B. Lipid radical formation correlates inversely with PhGPx activity. Data are derived from Figures 3A and 5A.

Figure 5

<sup>&</sup>lt;sup>3</sup> Wang HP, et al. (2001) Free Radic Biol Med. in press.)

The University of Iowa, Free Radical and Radiation Biology Program

## I. Tables

#### Same as Figures, make them easy to understand.

#### UV Light Increases Iron in Skh-1 Mouse Skin<sup>4</sup>

Skin Sample	[Fe(III)Desferal]/m M	
Non UV Exposure	$5.2 \pm 0.4$	
UV Exposed (16 weeks)	10.0 ± 2.3	
(16 weeks)	0.1	

n = 3; mean ± SEM; p ~ 0.1

Cell Line	Median Diameter	<b>SD</b> <sup>b,c</sup>	Range <sup>b</sup>	Range <sup>d</sup>
	nm	± mm	nim	nn
CEM	9 <sup>a</sup>	1.8	5.2 - 14.1	8 - 12
HEL	12.1 <sup>b</sup>	1.3	9.1 - 15.8	10 - 18
HL-60	10 <sup>a</sup>	1.4	6.1 – 13.6	8 - 12
K-562	14.3 <sup>b</sup>	1.6	9.3 – 18.7	10 -20
KG-1	10.3 <sup>b</sup>	1.4	6.1 – 13.3	8 - 18
L1210	10.4 <sup>b</sup>	1.1	7.5 – 13.3	8 - 10
Molt-4	11.4 <sup>b</sup>	1.3	8.5 - 15.5	10 - 12
THP-1	12.5 <sup>b</sup>	1.4	9.2 – 16.6	8 - 12
U-937	13.0 <sup>a</sup>	1.8	9.1 - 18.3	8 - 18

#### Table 1 Size of nine leukemia cell lines<sup>5</sup>

<sup>a</sup> These diameters were determined using conventional light microscopy and were used as standards for the flow cytometric determinations.

- <sup>b</sup> These values were determined from the FSC measured by flow cytometry, using the equation: cell diameter = (0.0893•FSC - 0.677) μm.
- <sup>c</sup> The standard deviation (SD) of the cell diameter distribution was determined from the gaussian FSC distribution. Using the equation in <sup>b</sup>, (FSC<sub>median</sub> ± SD<sub>FSC</sub>) was converted to cell diameters. The SD in μm is then estimated as the range of these two values/2.
- <sup>d</sup> These values were determined using a micrometer and conventional light microscopy.

#### Table 1

<sup>5</sup> Schafer FQ. (2000) partially published.

9

- 1. Make a list of the Primary key words
- Construct an informative title that contains these key words (or the most important of them).
- 3. The title will most likely have a verb in it.
- 4. If appropriate, the title should state the main conclusion of the paper.
- 5. Use mainstream words.
- Let it age; constantly rethink, especially as you are putting on the final touches to the manuscript.

# B. Abstract

Secondary key words

Packed with concrete information

If appropriate:

Introduction

Hypothesis/Problem

**Data/Facts** 

Conclusion

< 250 words

Example: See Jurkiewicz

12

# C. Introduction

Present information that points to:

your point,

your hypothesis,

your problem; i.e.

The reason for the work.

- But also think about educating the reader so they can understand where you are going and how you are getting there.
- At the end is often a general statement of the findings.

Example: See Jurkiewicz or Kelley

### **D.** Materials and Methods

Can another person, competent in this field, repeat exactly what you've done?

E. Results (& Discussion)

Lead reader to: your point, and your interpretation, and your conclusions

- F. (Discussion)
- G. Acknowledgments Support, Funding reports.

# H. References All as appropriate

# VI. The Writing

# A. Write to the reader's expectations!

- B. Write with emphasis.
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# VII. How?

- A. At the document level
  - 1. Issue, topic
  - 2. Persuasive discussion
  - 3. Coda, finale, conclusion

# 4. Chaining, *i.e.* everything flows

# B. At the paragraph level

- 1. Issue, topic
- 2. Persuasive discussion
- 3. Coda, finale, conclusion
- 4. Chaining, *i.e.* everything flows

Examples:

# J10, J12

K12, K13, K14

# C. At the sentence level

- 1. Bold, with emphasis on point
- 2. Simple and direct
- 3. Chaining
  - a. old-p --> new-1 (Issue)
  - b. old-1 --> new-2
  - c. old-2 --> new-3
  - d. old-3 --> new-4
  - e. old-4 --> new-5
  - X. ...
  - z. Coda, finale, conclusion,

# Always ask what should be the emphasis of a sentence.

Examples: K1, Chaining J16, Chaining and emphasis