Complaint of Alleged Scientific Misconduct by
Professors Stanton A. Glantz and Lisa A. Bero
That Violates the UCSF Campus Code of Conduct
and Policy on Ethical Conduct of Research

James E. Enstrom, Ph.D., M.P.H.
University of California
Box 951772
Los Angeles, CA 90095-1772
jenstrom@ucla.edu

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Background

I am submitting this complaint because I have substantial evidence of alleged scientific misconduct by UCSF Professors Stanton A. Glantz (Glantz) and Lisa A. Bero (Bero). I allege that this evidence, in specific instances and in its totality, represents scientific misconduct as defined by the UCSF Campus Code of Conduct and the UCSF Office of Research policy on "Ethical Conduct of Research." The Code of Conduct states: "Misconduct or Misconduct in Science means fabrication, falsification, plagiarism, or other practices that seriously deviate from those that are commonly accepted within the scientific community for proposing, conducting, or reporting research." (http://ucsfhr.ucsf.edu/policies/files/finalcc.pdf). The policy on "Ethical Conduct of Research" states: "UCSF is committed to creating and maintaining an environment for research that conforms to the highest ethical principles, fosters intellectual honesty and integrity, and encourages scientific advances of the highest quality." (http://www.research.ucsf.edu/QG/orQgEth.asp).

This scientific misconduct has occurred during the past two years, during which time Glantz and Bero have repeatedly and maliciously attacked me and my epidemiologic research, using fabrication and falsification in many instances. In the interest of brevity, I have limited this submission to nine pages of text, which includes web links that document all the sources that I have used. If my allegations of misconduct are not clear from the details below, then I request the opportunity to make a personal presentation in order to fully explain them. If my allegations are sustained, then I request that appropriate action be taken regarding Glantz and Bero. Since much of the alleged misconduct is related to the selective use of tobacco industry documents by Glantz and Bero, I have used, wherever possible, my own links to documents from the UCSF Legacy Tobacco Documents Library (http://legacy.library.ucsf.edu). All of the links given below should be freely accessible via connection with the UC electronic library system.

This submission involves the very fundamental issues of academic freedom, scientific integrity, and professional conduct. These issues have gotten the attention of high officials at both the University of California and the National Institutes of Health. Some aspects are discussed in the February 2005 *Nature Medicine*, where Dr. Lawrence B. Coleman, Vice Provost for Research at the University of California, stated "Academic freedom has to be absolute or no one has it"

(http://www.nature.com/nm/journal/v11/n2/pdf/nm0205-106a.pdf) and in the March 2005 *Nature Medicine*, where NIH Director Elias Zerhouni "has called for an 'ethics summit,' and rules for scientists at outside institutions receiving NIH grants could be heavily scrutinized." (http://www.nature.com/nm/journal/v11/n3/pdf/nm0305-235.pdf).

I would like to begin with a few brief sentences about my background in order to assure you that I am a serious scientist with an important message. I have a Ph.D. from Stanford University, awarded in 1970, and my dissertation advisor is a Nobel Laureate. Also, I have postdoctoral certification in cancer epidemiology and a M.P.H. in epidemiology from UCLA, awarded in 1975 and 1976. Since 1976 I have been on the research faculty at the UCLA School of Public Health. I have had a long and successful career as an epidemiologist. I am a 24-year Fellow of the American College of Epidemiology and I am listed in *Who's Who in America* in recognition of my epidemiologic research.

During the first 33 years of my professional career no one ever once questioned my honesty or integrity as a scientist. However, that situation changed dramatically in May 2003 when I and my co-author, Dr. Geoffrey C. Kabat (Kabat), published a paper in the May 17, 2003 British Medical Journal (*BMJ*), "Environmental tobacco smoke and tobacco related mortality in a prospective study of Californians, 1960-98" (http://bmj.com/cgi/reprint/326/7398/1057.pdf), henceforth referred to as "my *BMJ* paper" or "my *BMJ* study." This paper describes the largest and most detailed epidemiologic study on environmental tobacco smoke (ETS) and tobaccorelated mortality ever published in a major medical journal and the second largest study ever published in terms of its statistical power. This study found no relationship between ETS and tobacco-related mortality and instantly became very controversial, as described in the following May 18, 2003 Sunday Telegraph (London) newspaper article:

http://www.telegraph.co.uk/news/main.jhtml?xml=/news/2003/05/18/nsmoke18.xml.

Since May 15, 2003 many false and misleading charges have been made against me and my research, primarily because the study was partially funded by the tobacco industry. These charges have damaged my professional reputation and my ability to publish in several journals that are now aware of the unjustified controversy surrounding me. However, in the two years since its publication, no errors have been identified in my *BMJ* paper, the alleged flaws in the study have not been substantiated with any actual evidence, and the *BMJ* editor has strongly defended his decision to publish the paper (http://bmj.com/cgi/reprint/327/7413/501).

As evidence that Glantz, or Glantz in collaboration with Bero, have engaged in an unprofessional, two-year campaign to discredit me and my research, I document below the initial aspect of their attack, plus two egregious aspects that have occurred since March 8, 2005. These aspects represent only a portion of their full attack, but they should provide sufficient evidence to document scientific misconduct.

Aspect 1) Early Statements by Glantz and Bero Meant to Discredit Me and My Research

On May 15, 2003, Glantz and Bero participated in a Miami press conference of "international experts" assembled to "debunk" my study before he could have possibly read it in any detail. These "experts" falsely claimed the paper said "Marry a smoker, get less cancer" and falsely

claimed it was a "tobacco industry study" (http://www.no-smoking.org/may03/05-15-03-4.html). It is not clear how these "experts" learned of the study, but they apparently violated the press embargo on the paper, which lasted until 12:01 AM May 16, 2003 UK time (or 7:01 PM May 15, 2003 Miami time). Glantz and Bero could not possibly have read the full version of the *BMJ* paper, which was first posted on bmj.com at this same time.

On May 16, 2003 Glantz told the San Francisco Chronicle: "... that because secondhand smoke was so common in the early years of the study, UCLA's research was fatally flawed and could only produce the kind of result the tobacco industry wanted... the British Medical Journal report was a textbook case of why UC researchers should not be allowed to accept funding from the tobacco industry.... It is an embarrassment that this came out of UCLA...." (http://www.sfgate.com/cgi-bin/article.cgi?f=/c/a/2003/05/16/MN259820.DTL).

On May 24, 2003 Glantz and Bero wrote a Rapid Response (electronic letter) to bmj.com entitled "Misleading the public about secondhand smoke . . . again." They stated "Enstrom and Kabat's study is the latest in a long string of studies supported by the tobacco industry to deny the evidence about secondhand smoke and confuse the public. . . . The Enstrom and Kabat study may be another example of the financial disclosure not fully describing the extent of involvement of the tobacco industry in the design, conduct and dissemination of the study. . . . By publishing Enstrom and Kabat's paper, the *BMJ* has helped the tobacco industry mislead the public about the harmful effects of secondhand smoke exposure. Only a retraction could stem some of the damages to public health goals that have already been inflicted by this paper." (http://bmj.bmjjournals.com/cgi/eletters/326/7398/1057#32596)

On June 20, 2003 comments involving Glantz and my *BMJ* study were made during the California Air Resources Board Scientific Review Panel meeting that was evaluating the CARB draft report on "Proposed Identification of Environmental Tobacco Smoke as a Toxic Air Contaminant." The exact comments are shown below, as taken directly from pages 84 and 85 of the meeting transcript (http://www.arb.ca.gov/srp/030620.pdf):

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PETERS SHORTHAND REPORTING CORPORATION (916) 362-2345 84 & 85
25 CHAIRPERSON FROINES: As a member of the UCLA

1 School of Public Health, I apologize.

2 (Laughter.)

3 PANEL MEMBER GLANTZ: You should.

4 (Laughter.)

5 PANEL MEMBER GLANTZ: We're doing a study of how

6 that paper came to pass. And it's going to get even more

7 unpleasant.

8 CHAIRPERSON FROINES: James Enstrom's paper --

9 PANEL MEMBER GLANTZ: -- that dreamt up by

10 Phillip (sic) Morris.

11 CHAIRPERSON FROINES: Go ahead.

12 PANEL MEMBER HAMMOND: How smoking doesn't cause

13 any lung cancer.
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On June 23, 2003 Glantz told UPI ".... As far as I know, there's no legitimate scientist in the world who doesn't think secondhand smoke causes lung cancer and heart disease. There are a number of people paid by cigarette companies to say that it doesn't.... I think it is shameful the British Medical Journal published that study...."

(http://www.hawaiireporter.com/story.aspx?e3a73f3e-4104-43ba-8ece-a7169b47149c).

On July 25, 2003 Glantz co-wrote an eight-page to Dr. Lawrence B. Coleman, UC Vice Provost for Research(http://www.ucsf.edu/senate/townhallmeeting/TobIndFundingColeman7-25-03.pdf). The following excerpt is taken from pages 3 and 4:

"A Recent Example: The UCLA Study of Secondhand Smoke

The most recent example of how the tobacco industry uses funding of university research as part of its for propaganda campaign is a May 17, 2003 study from UCLA on the health effects of secondhand smoke published in the *British Medical Journal*. This study concluded that secondhand smoke does not increase the risk of lung cancer or heart disease.

This paper has attracted a storm of scientific criticism, manifest as an unusually large number of criticisms through the *British Medical Journal*'s "rapid response" mechanism (available, along with the original paper, at http://bmj.com/cgi/eletters/326/7398/1057.) There is little possibility that it will be taken seriously in scientific circles. To this extent, the normal scientific process of open discussion and debate is functioning effectively.

The difficulty is that the discussion of this study is not simply occurring within the academic community. If it were, this paper would go down as one bit of poor research done at a university with a reputation for high quality scholarship that slipped into a good journal because of the foibles of the peer review process. Instead, the paper is already being exploited world-wide in the tobacco industry's public relations and political campaign against effective measures to protect the public from the toxins in secondhand smoke. Public health experts are doing what they can to set the record straight, but the fact remains that the multinational tobacco industry has a much larger megaphone.

Like others reaching similar conclusions, this project was supported by the tobacco industry as a 'special reviewed' project selected by industry lawyers and executives. (See the attached bibliography for more examples.) Rather than sharing the University's objective of supporting the best science as part of a larger search for truth, the tobacco industry has a long and well practiced program of using specially reviewed projects to stimulate and support just such studies as part of its propaganda and political activities.

The fact that a poorly conceived study funded by the tobacco industry reaches conclusions that, in contrast to the established literature, support the tobacco industry's agenda is not surprising. It is, however, unseemly and harms the University's reputation."

These are all libelous statements by the common definition of libel, "a written or oral defamatory statement or representation that conveys an unjustly unfavorable impression" (*Webster's New Collegiate Dictionary*). The full impact of these libelous statements can most accurately be seen when they are viewed as part of a larger campaign to "silence science" regarding my *BMJ* paper. This evidence is documented in the 19-page 2005 paper by two sociologists, Drs. Sheldon Ungar and Dennis Bray, entitled "Silencing science: partisanship and the career of a publication disputing the dangers of secondhand smoke" (http://pus.sagepub.com/cgi/reprint/14/1/5). Ungar and Bray described in detail the "efforts to prevent the making of specific scientific claims in any *or* all of the arenas in which these claims are typically reported or circulated" as they related to my *BMJ* paper.

Aspect 2) March 8, 2005 KQED Radio Forum involving Glantz

On Tuesday, March 8, 2005 at 9:00 AM Glantz participated in the San Francisco based KQED Radio *Forum* (http://www.kqed.org/epArchive/R503080900). The topic was "Funders and Academic Research: *Forum* assesses the controversy surrounding the relationship between funders and academic research." The host was Michael Krasny and the primary participants, in addition to Glantz, were Dr. Max Neiman, Chair of the system-wide University of California Committee on Research Policy, and Dr. Michael Kleinman, Adjunct Professor of Community and Environmental Medicine at UC Irvine. This program can be listened to in its entirety by clicking on the above link.

The program initially discusses Glantz's attempt to have the California Attorney General's Office launch a criminal investigation into an epidemiologic review article on ETS and SIDS written by Dr. Frank M. Sullivan, a retired Professor from University of London, who has had a long and distinguished career as a toxicologist in England. During this 52 minute program, the discussion of the "scandal" about me and my *BMJ* study occurred during the following time period (minutes:seconds): 16:57-19:15.

Four examples of libelous statements against me by Glantz, and the exact time at which they begin, are given below, along with my statement of the actual facts of the matter.

At 17:20 Glantz says the *BMJ* study "was not funded by the American Cancer Society" but "done with Philip Morris's money." Actually, the inflation-adjusted funding for the study, which began in 1959 and was published in 2003, came from the three primary sources: ~90% ACS, ~5% TRDRP, and ~5% CIAR (the 'tobacco money' portion). NO Philip Morris money was used for this study. Glantz, who could not precisely know the 44 year funding history of the study, simply made false statements about the funding of my study.

At 17:50 Glantz says I am "a damn fool" who was told by ACS that I "made inappropriate use of the data." Actually, my use of ACS data began in 1991 and I had the full cooperation of and long standing working relationships with Lawrence Garfinkel and Dr. Clark W. Heath, Jr., now retired ACS Vice Presidents for Epidemiology. I have been conducting important long-term epidemiologic research with the California portion of the CPS I cohort. My dealings with ACS epidemiologists date back to 1978, when I received all my research funding from ACS. I am the ONLY investigator outside of ACS who has ever been allowed to follow ACS subjects. This access was granted largely because of the high quality of my ACS-funded epidemiologic research, begun in 1973. The ACS epidemiologists who I worked with, Garfinkel and Heath, realized the great potential value of long-term follow-up of the CPS I cohort and they would not have given important confidential data to "a damn fool." Only in May 2003, when the BMJ paper was published, did the ACS (most specifically, Dr. Michael J. Thun) complain about my use of their data. I worked with Health on this study until 2001, when he was no longer able to continue because of his retirement. Heath was a co-author on the first version of the study that was submitted to and given serious consideration by the New England Journal of Medicine. I never worked with Thun on this study.

At 18:10 Glantz implies that I am "advocating a pro-tobacco position." Actually, I am a lifelong nonsmoker and have never advocated a pro-tobacco position in my entire 35-year career or in my entire life for that matter. As evidence I am not "pro-tobacco," I have spent much of my career documenting the health benefits of being a nonsmoker, as can be seen by reading my publications on Mormons, which date back to 1975 (http://legacy.library.ucsf.edu/tid/gei79c00). In 1999 I published two papers indicating active smoking may be more dangerous than generally believed because its impact on mortality was less reversible by cessation than generally believed (http://legacy.library.ucsf.edu/tid/tbf19c00) and http://legacy.library.ucsf.edu/tid/wve19c00). My findings regarding lung cancer and smoking cessation were largely confirmed in a Mayo Clinic study of Iowa women published the May 2003 *Journal of Clinical Oncology* (http://www.jco.org/cgi/reprint/21/5/921).

At 18:39 Glantz states "the science that the UCLA study did was crap." Actually, my study is the largest and most detailed epidemiologic study on secondhand smoke and mortality ever published in a major medical journal, the *BMJ*. It is by far the largest study on Californians. The paper was deemed to be scientifically sound and worthy of publication after being peer reviewed by two of the world's leading epidemiologists, Drs. Kenneth Rothman and George Davey-Smith. Rothman is the author of several major textbooks on epidemiology and founding editor of *Epidemiology* and Davey-Smith is co-editor of *International Journal of Epidemiology*. In the interest of transparency, the *BMJ* took the unusual step of posting the entire prepublication history of the paper online (http://bmj.bmjjournals.com/cgi/content/full/326/7398/1057/DC1).

There might be some logic to Glantz's attack if he had found a single error in my *BMJ* paper or had proved that the paper was "fatally flawed" because everyone alive in 1959 was equally exposed to ETS. But he has done neither of these things. The August 30, 2003 *BMJ* letter by me and Kabat clarified the findings of our 1999 follow-up survey, which clearly showed that 1959 spousal smoking history was a valid measure of relative ETS exposure, particularly for females (http://bmj.bmjjournals.com/cgi/content/full/327/7413/504). Also, I have evidence from three other independent surveys that spousal smoking status is a valid indicator of relative ETS exposure for subjects alive in the 1950s. Keep in mind, all subjects in the other US cohort studies were alive in 1959 and these studies are not considered to be "fatally flawed" by Glantz.

Aspect 3) April 2005 Tobacco Control paper by Bero and Glantz

Bero, Glantz, and M.-K. Hong are authors of a nine-page paper in the April 2005 issue of *Tobacco Control (TC)*, entitled "The limits of competing interest disclosures." It is now posted on the following UCSF web site: http://www.tobaccoscam.ucsf.edu/pdf/Enstrom-TC.pdf. This paper questions the veracity of the following 200+ word competing interest disclosure statement made at the end of my *BMJ* paper: "Funding: The American Cancer Society initiated CPS I in 1959, conducted follow up until 1972, and has maintained the original database. Extended follow up until 1997 was conducted at the University of California at Los Angeles with initial support from the Tobacco-Related Disease Research Program, a University of California research organisation funded by the Proposition 99 cigarette surtax (www.ucop.edu/srphome/trdrp). After continuing support from the Tobacco-Related Disease Research Program was denied, follow up through 1999 and data analysis were conducted at University of California at Los Angeles with support from the Center for Indoor Air Research, a 1988-99 research organisation that received funding primarily from US tobacco companies. Competing interests: In recent years JEE has received funds originating from the tobacco industry for his tobacco related epidemiological research because it has been impossible for him to obtain

equivalent funds from other sources. GCK never received funds originating from the tobacco industry until last year, when he conducted an epidemiological review for a law firm which has several tobacco companies as clients. He has served as a consultant to the University of California at Los Angeles for this paper. JEE and GCK have no other competing interests. They are both lifelong non-smokers whose primary interest is an accurate determination of the health effects of tobacco."

Any doubts that a reasonable person might have had regarding our competing interest disclosures were addressed in the August 30, 2003 *BMJ* letter by me and Kabat: "We want to make clear that the tobacco industry played no part in our paper other than providing the final portion of the funding. The tobacco industry never saw any version of our paper before it was published, never attempted to influence the writing of the paper in any way, and did not even know the paper was being published until it became public. In addition, we have never testified on behalf of the tobacco industry, never owned any stock in the tobacco industry, never been employees of the tobacco industry, and would never have accepted tobacco industry funds if there had been any other way to conduct this study." (http://bmj.bmjjournals.com/cgi/content/full/327/7413/504).

In spite of this clear and unequivocal statement, Bero and Glantz still went ahead and wrote the *TC* paper, which completely mischaracterized the relationships that Kabat and I have had with the tobacco industry. This paper is simply an *ad hominem* attack designed to impugn our scientific integrity and damage our professional reputations. It is clearly libelous by the common definition of libel, "a written or oral defamatory statement or representation that conveys an unjustly unfavorable impression" (*Webster's New Collegiate Dictionary*). In fact, I believe it contains the greatest amount of malicious libel ever published in a single peer-reviewed paper.

One particularly reprehensible example of the libel is Table 1 in the *TC* paper, which shows "Financial ties between Enstrom, Kabat, and the tobacco industry" dating back to 1975. Although I actually had no "financial ties" of any kind with the tobacco industry before July 1, 1992, Bero and Glantz listed six alleged ties under "Enstrom" in Table 1 dated from 1975 to 1991. To illustrate the maliciousness of this libel, I have examined their first entry in detail.

As my first alleged "financial tie," Bero and Glantz cited my 1975-76 correspondence with the Council for Tobacco Research (CTR), a research organization funded by the tobacco industry, regarding proposed epidemiologic research on Mormons (TC references 23-27). However, they failed to mention that the actual December 1975 grant application to CTR was submitted by Dr. Lester Breslow, then Dean and Professor at the UCLA School of Public Health (http://legacy.library.ucsf.edu/tid/bei79c00). Breslow, a world-renowned public health authority whose accomplishments are summarized in Who's Who in America, has been described as "Mr. Public Health" in a detailed October 13, 1997 Los Angeles Times newspaper article (http://www.pacificnet.net/jue/healthfitness/mrpublichealth.html). He was my mentor and the prinicipal investigator on several grant applications that we submitted to potential funding agencies beginning in 1974. Next, they failed to cite Breslow's July 6, 1976 letter withdrawing the unfunded CTR application once we had received funding for this Mormon research from the ACS (http://legacy.library.ucsf.edu/tid/sei79c00). Then, they failed to mention my 1978 CANCER publication on cancer mortality among active Mormons (http://legacy.library.ucsf.edu/tid/msd3aa00), which acknowledged on the first page the funding received from the ACS (Grant PDT-51).

Finally, Bero and Glantz failed to mention that Mormons are a religious group that advocates ABSTENTION from tobacco and that I was (and still am) studying them because their unusually

low cancer rates offer an excellent opportunity to better understand etiologic factors associated with the prevention of cancer. If Bero and Glantz had any interest in fairly and accurately portraying me and my epidemiologic research interests during the past 30 years they would have cited my initial findings on Mormons that appeared on the front page of the Washington Post on November 18, 1974 (http://legacy.library.ucsf.edu/tid/zci79c00).

This one example related to one line in Table 1 of the *TC* paper indicates how Bero and Glantz have selectively used tobacco industry documents in order to deliberately distort my career and my relationship with the tobacco industry. On its surface the *TC* paper purports to provide evidence of the inadequacy of the *BMJ*'s requirements concerning competing interest disclosures. However, beneath this veneer, the paper's true objective is to smear the reputations of two honest scientists who had the temerity to publish an influential paper reporting results which run counter to the firmly held beliefs of Bero and Glantz.

In comparison with our 107-word *BMJ* funding statement shown above, please note the 15-word *TC* funding statement: "Research support – California Tobacco-Related Disease Research Program grant 9RT0193 and National Cancer Institute grant CA-87472." Missing from this brief statement is any clarification that TRDRP Grant 9RT0193 is a \$442,670 award to Bero for "Analysis of tobacco industry documents on scientific research" (http://trdrp.org/research/PageInvestigator.asp?person_id=449). NCI Grant 5R01CA087472 is a multi-million dollar NIH grant awarded to Glantz for "Analysis of Tobacco Industry Documents" (http://www.crisp.cit.nih.gov), as part of a large NCI program on "Review and Analysis of Tobacco Industry Documents" (http://cancercontrol.cancer.gov/tcrb/grant_doc.asp). Thus, both Glantz and Bero have a direct financial interest in writing a paper designed to justify their examination and analysis of tobacco industry documents. But they failed to clearly and fully disclose/explain their "competing interest" in the *TC* paper, which is the very thing they alleged that Kabat and I did in the *BMJ* paper.

In addition, the "Declaration of Competing Interests" statement on the *TC* web site (http://tc.bmjjournals.com/misc/echoice3.shtml) fails to disclose that since 2001 *TC* Editor, Dr. Simon Chapman of Sydney, Australia has received NCI Grant 5R01CA087110 for the "Analysis of Tobacco Industry Documents--Asia/Australia" (http://www.crisp.cit.nih.gov). This grant is from the same NCI program as Glantz's grant. Thus, Chapman also has a direct financial interest in justifying the examination and analysis of tobacco industry documents. This "competing interest" may have influenced Chapman's decision to publish the *TC* paper.

Furthermore, Glantz, Bero, and *TC* Editor Chapman have ignored the NIH requirement that the following disclaimer be included in NIH-funded papers, such as, the *TC* paper: "Its contents are solely the responsibility of the authors and do not necessarily represent the official views of the [NCI]." This requirement is clearly stated in the "Rights in Data (Publication and Copyrighting)" section of the NIH Grants Policy Statement (12/03) (http://grants2.nih.gov/grants/policy/nihgps_2003/NIHGPS_Part7.htm).

The program announcement for the "Review and Analysis of Tobacco Industry Documents" program (http://grants1.nih.gov/grants/guide/pa-files/PAR-99-114.html) explains the purpose of this program. Based on my reading, this program was never intended for *ad hominem* attacks on

honest scientists. In conclusion, I believe that NIH funds have been inappropriately used for the writing of a *TC* paper that contains malicious libel and that has no direct connection to the mission of NIH. I find it particularly offensive that American taxpayers like myself, who expect NIH funds to be spent on finding ways to cure and prevent diseases, have to pay for the assassination of their own character. I have been able to get the attention of NIH Director Elias Zerhouni on this matter and hopefully it will be part of the "ethics summit" that he is proposing regarding NIH funded investigators, as mentioned in the March 2005 *Nature Medicine* article.

Two Specific Examples of Alleged Scientific Misconduct

To illustrate the alleged scientific misconduct in the clearest possible way, I have specifically identified two examples from the extensive evidence presented above. If necessary, further examples can be specifically identified from this evidence.

1) <u>Claim by Glantz that my BMJ study was "dreamt up by Philip Morris" and "done with Philip Morris's money"</u>

It is blatant fabrication and falsification for Glantz to claim that my BMJ study was "dreamt up by Philip Morris" and "done with Philip Morris's money." In actual fact, my BMJ study grew out of concerns that I had about the epidemiologic evidence on ETS and lung cancer in the 1992 EPA report, entitled "Respiratory Health Effects of Passive Smoking: Lung Cancer and Other Disorders" (http://legacy.library.ucsf.edu/tid/oys44f00). My concerns were expressed in a January 28, 1993 *Investor's Business Daily* newspaper article by Michael Fumento, entitled "Is EPA blowing its own smoke? How much science is behind its tobacco finding?" (http://legacy.library.ucsf.edu/tid/cap91a00). During 1993 I developed the initial research proposal for my BMJ study in conjunction with ACS epidemiologists Heath and Garfinkel. On February 18, 1994 I submitted this proposal, entitled "Tobacco-Related Disease Trends Among 118,000 Californians," to the University of California Tobacco-Related Disease Research Program (TRDRP) through the UCLA Office of Contract and Grant Administration. I still have full documentation regarding this proposal and Philip Morris had absolutely no involvement. Regarding the funding of my BMJ study, which was begun by the ACS in 1959, continued by me at UCLA beginning in 1991, and published in 2003, it came from three primary sources (in inflation-adjusted dollars): ~90% ACS, ~5% TRDRP, and ~5% CIAR (the 'tobacco money' portion). NO Philip Morris money was used for this study.

2) Claim by Bero and Glantz that "Financial ties between Enstrom, Kabat, and the tobacco industry" existed from 1975 to 1991

It is blatant fabrication and falsification for Bero and Glantz to represent in the *TC* paper that "Financial ties between Enstrom, Kabat, and the tobacco industry" existed from 1975 to 1991. In actual fact, as discussed in my detailed response above, I had no financial ties with the tobacco industry in 1975. Furthermore, I can fully document that I had no "financial ties" of any kind with the tobacco industry before July 1, 1992. Glantz and Bero listed six alleged ties under "Enstrom" in Table 1 of their *TC* paper that were dated before 1992, but they have not provided any evidence that I actually received money from the tobacco industry before 1992.

Epidemiologic Perspectives & Innovations



Analytic Perspective

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Defending legitimate epidemiologic research: combating Lysenko pseudoscience

James E Enstrom^{1,2}

Address: ¹University of California, Los Angeles, CA, USA and ²Scientific Integrity Institute, Los Angeles, CA, USA Email: James E Enstrom - jenstrom@ucla.edu

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Abstract

This analysis presents a detailed defense of my epidemiologic research in the May 17, 2003 *British Medical Journal* that found no significant relationship between environmental tobacco smoke (ETS) and tobacco-related mortality. In order to defend the honesty and scientific integrity of my research, I have identified and addressed in a detailed manner several unethical and erroneous attacks on this research. Specifically, I have demonstrated that this research is not "fatally flawed," that I have not made "inappropriate use" of the underlying database, and that my findings agree with other United States results on this relationship. My research suggests, contrary to popular claims, that there is not a causal relationship between ETS and mortality in the U.S. responsible for 50,000 excess annual deaths, but rather there is a weak and inconsistent relationship. The popular claims tend to damage the credibility of epidemiology.

In addition, I address the omission of my research from the 2006 Surgeon General's Report on Involuntary Smoking and the inclusion of it in a massive U.S. Department of Justice racketeering lawsuit. I refute erroneous statements made by powerful U.S. epidemiologists and activists about me and my research and I defend the funding used to conduct this research. Finally, I compare many aspect of ETS epidemiology in the U.S. with pseudoscience in the Soviet Union during the period of Trofim Denisovich Lysenko. Overall, this paper is intended to defend legitimate research against illegitimate criticism by those who have attempted to suppress and discredit it because it does not support their ideological and political agendas. Hopefully, this defense will help other scientists defend their legitimate research and combat "Lysenko pseudoscience."

Background

This analysis presents a detailed response to the extensive attacks that have been made on my epidemiologic research in the May 17, 2003 *British Medical Journal*, "Environmental tobacco smoke and tobacco related mortality in a prospective study of Californians during 1960–98" [1]. I seek to defend the honesty and scientific integrity of my research and I directly respond to my most powerful critics, who have attempted to suppress and discredit findings that do not support their ideological and political

agendas. To put a historical perspective on the tactics that have been used against me, I conclude by making an analogy with the pseudoscientific practices of Trofim Denisovich Lysenko [2]. Hopefully, my defense will encourage and/or help other honest scientists to defend their research against unwarranted and illegitimate criticism.

This analysis deals with several important elements of the attacks, with a primary focus on the epidemiologic issues involved. Additional elements of the attack are mentioned

briefly in this analysis and are presented in detail on my Scientific Integrity Institute website, under 'Research Defense' [3]. Being attacked for publishing unpopular scientific findings is not unique to me or my research. However, the nature and scope of the attacks to which I have been subjected is quite unusual and needs to be documented and addressed.

Being able to distinguish between real and implied scientific misconduct is important to the integrity of science in general and to the integrity of individual scientists in particular. Falsely accusing an honest scientist of scientific misconduct is just as wrong as scientific misconduct itself. Implying that an honest scientist has committed scientific misconduct because he has published unpopular findings or has used an unpopular funding source is wrong and falls under the category of "scientific McCarthyism" [4].

Analysis

Background on BMJ Paper

I begin with a presentation of the background necessary to understand the issues involved with the May 17, 2003 British Medical Journal (BMJ) paper that I wrote with Dr. Geoffrey C. Kabat [1]. This account primarily involves me and thus is written in the first person, but it also refers to Kabat where appropriate and not otherwise noted. Our paper found no relationship between environmental tobacco smoke (ETS) and tobacco-related mortality in a prospective study of Californians during 1960–1998, with some associations slightly below the null and some slightly above the null, but none statistically different from the null. It concluded, "The association between exposure to environmental tobacco smoke and coronary heart disease and lung cancer may be considerably weaker than generally believed." It is the largest (in terms of statistical power), most detailed (in terms of results presented), and most transparent (in terms of information about its conduct) epidemiologic paper on ETS and mortality ever published in a major medical journal.

The study is based on the California (CA) portion of the original 25-state Cancer Prevention Study (CPS I) [1]. CA CPS I was begun by the American Cancer Society (ACS) in 1959 and has been conducted at UCLA by me since 1991. Kabat and I are both well qualified epidemiologists who have had long and successful careers dating back to the 1970s, as can be confirmed by examining our epidemiologic publications on PubMed. Our paper was deemed to be scientifically sound and worthy of publication after being peer reviewed by two distinguished epidemiologists, a *BMJ* statistician, and a *BMJ* editorial committee. The details of the entire peer review process and the names of all the individuals involved in the review process are available online as the "Prepublication history" [5]. The paper was subjected to the same review process and selec-

tion criteria as other papers submitted to the *BMJ*, which publishes less than 10% of the total submissions it receives [6].

In the interest of transparency and full disclosure, the paper included the following detailed statements about the funding history of the study and the competing interests of the authors: "Funding: The American Cancer Society initiated CPS I in 1959, conducted follow up until 1972, and has maintained the original database. Extended follow up until 1997 was conducted at the University of California at Los Angeles with initial support from the Tobacco-Related Disease Research Program, a University of California research organisation funded by the Proposition 99 cigarette surtax. After continuing support from the Tobacco-Related Disease Research Program was denied, follow up through 1999 and data analysis were conducted at University of California at Los Angeles with support from the Center for Indoor Air Research, a 1988-99 research organisation that received funding primarily from US tobacco companies. Competing interests: In recent years JEE has received funds originating from the tobacco industry for his tobacco related epidemiological research because it has been impossible for him to obtain equivalent funds from other sources. GCK never received funds originating from the tobacco industry until last year, when he conducted an epidemiological review for a law firm which has several tobacco companies as clients. He has served as a consultant to the University of California at Los Angeles for this paper. JEE and GCK have no other competing interests. They are both lifelong nonsmokers whose primary interest is an accurate determination of the health effects of tobacco." [1].

Initial Attacks on BMJ paper

Even though our paper satisfied (and in many ways exceeded) the accepted standards of epidemiologic analysis and writing, it was immediately attacked by people who did not like the results we reported. Beginning in the days before May 17, 2003, our BMJ paper was subjected to a large-scale ad hominem attack. Since our honesty or scientific integrity had never previously been questioned, such an attack seemed to us to be quite implausible and indeed incredible. Based on what I have learned since May 2003, I describe the key elements of this attack in order to expose the tactics that have been used in an attempt to discredit and silence legitimate epidemiologic research. Additional details are presented on my Scientific Integrity Institute website [3]. The attack has been largely due to the fact that we published politically incorrect null findings from a long-term study primarily funded by the ACS, but completed with a research award to UCLA from the Center for Indoor Air Research (CIAR), a now-defunct tobacco-industry funded research organization.

On May 9, 2003 I learned that our paper was to be published in the May 17, 2003 BMJ and that an embargoed BMJ press release was to be issued on May 13, 2003. The strict publication/broadcast embargo regarding our paper was to last until 00:01 hours (UK time) on May 16, 2003, which was 19:01 (7:01 PM) EDT on May 15, 2003 in Florida and 16:01 (4:01 PM) PDT on May 15, 2003 in California. During this period, the ACS was informed of our forthcoming paper and the press embargo. The ACS then prepared its own press release entitled "American Cancer Society Condemns Tobacco Industry Study for Inaccurate Use of Data." The May 14, 2003 version of the ACS press release was inserted into a May 15, 2003 email message of Stanton A. Glantz, Ph.D., Professor of Medicine at the University of California, San Francisco (UCSF). Glantz send out this message worldwide to his UCSF listserv before the press embargo ended [7]. The official May 15, 2003 version of the ACS press release, which adhered to the press embargo, was issued in a separate PDF form [8]. Then it was permanently posted on the ACS web site in a slightly different format [9].

The instantaneous attack on our paper appears to have been a coordinated effort, primarily organized by the ACS and Glantz. Glantz is a well-known anti-smoking activist who has worked closely with the ACS for many years [10]. As part of this coordinated effort, Glantz organized a May 15, 2003 Miami, Florida press conference involving a panel of "international experts" in order to "debunk" our "Marry a Smoker, Get Less Cancer" study before the press embargo ended [11]. At the time of the ACS press release and the Miami press conference, neither the ACS, Glantz, or the other Miami "experts" had access to the full tenpage version of our paper, let alone time to read it and carefully analyze it. The full version of our paper was not posted on the BMJ website until the press embargo lifted at 7:01 PM EDT on May 15, 2003 [1]. The only version available when the embargoed BMJ press release was issued on May 13, 2003 was the abridged five-page paper that appears in the print version of the BMJ [12]. Obviously, these critics chose to hastily write a press release and hold a press conference based on limited information. They did not have the integrity or objectivity to read our full ten-page paper or to contact the authors before beginning their attack, which included erroneous claims about the paper's content and quality.

The ACS press release was authored by Michael J. Thun, M.D., ACS Vice President, Epidemiology and Surveillance Research, and Harmon J. Eyre, M.D., ACS Chief Medical Officer. This press release makes several entirely false statements about the study, such as:

1) "Tobacco Industry Study" was "Part of Organized Effort to Confuse Public About Secondhand Smoke"

- 2) "Society researchers repeatedly advised Dr. Enstrom that using CPS-I data to study the effects of secondhand smoke would lead to unreliable results"
- 3) "this study is neither reliable nor independent"
- 4) "The study suffers from a critical design flaw: the inability to distinguish people who were exposed to second-hand smoke from those who were not"
- 5) "exposure to secondhand smoke was so pervasive [in 1959] that virtually everyone was exposed to ETS, whether or not they were married to a smoker".

Further distracting from the actual content of the study and the legitimacy of the analysis, the press release added a number of out of context quotes from formerly confidential tobacco industry documents that had nothing to do with the conduct, analysis, or publication of the study. For the past several years these documents have been available online from the Legacy Tobacco Documents Library at UCSF [13], which was established by Glantz [14]. These documents are also available at other online tobacco document libraries [15]. As shown above, my tobacco industry funding and competing interests were clearly and accurately described in more than 200 words in the BMJ paper [1]. However, in order to raise doubts about my honesty and scientific integrity, the ACS made a great effort to locate and extract selective quotes from the professional correspondence I have had with the tobacco industry during my career. This ad hominem attack diverted attention from the paper itself and obscured its contribution to the body of epidemiologic evidence regarding the lethality of ETS.

A major element of the attack included the submission to the *BMJ* website of over 150 mostly negative electronic letters, known as "rapid responses" (rrs) [16]. The overall content and nature of these rrs was summarized by a *BMJ* associate editor in an August 30, 2003 letter [17]. Particularly troubling are May 19 and 20, 2003 rrs by Thun [18,19], a May 30, 2003 rr by Thun and 13 other members of the International Agency for Research on Cancer (IARC) Working Group on tobacco smoke [20], and a August 19, 2003 rr by Drs. Phillip S. Gardiner, Charles Gruder, and Francisco Buchting of the University of California Office of the President [21]. None of the authors of these criticisms ever contacted us for a clarification of any aspect of our *BMJ* paper or our contacts with the tobacco industry before posting their rrs.

Most of the press coverage of the study was muted or equivocal because of the issues raised by the ACS criticism of the paper. Typical of this type of newspaper coverage was the May 16, 2003 Los Angeles Times article on page A26, "Study Downplays the Health Risks From Secondhand Smoke." This article concludes with the following quote from Dr. Jonathan Samet, Professor and Chair of Epidemiology at the Johns Hopkins University Bloomberg School of Public Health: "We have one very flawed study that does not find an association. It flies in the face of so much evidence and so much scientific understanding that it just doesn't contribute." [22].

Supportive Commentary on the BMJ Paper

A supportive press account appeared in the May 18, 2003 Sunday Telegraph newspaper article, "Warning: the health police can seriously addle your brain," by Robert Matthews [23]. The article noted, "More than any other health debate, the question of whether smokers kill others as well as themselves is engulfed in a smog of political correctness and dubious science." Other supportive commentaries also appeared. Michael Fumento, a Senior Fellow at the Hudson Institute, wrote a September 11, 2003 syndicated column, "Second-hand Smoke is Harmful to Science" [24]. Elizabeth Whelan, Sc.D., President of the American Council on Science and Health (ACSH), wrote an August 13, 2004 ACSH column entitled "American Cancer Society a Danger to Science?" [25]. Michael Fitzpatrick, M.D., a general practice physician in London, wrote a November 15, 2004 Spiked commentary entitled "We have ways of making you stop smoking." [26]. These commentaries put our BMJ findings in context and described the excesses of the anti-smoking critics who attacked us.

Two sociologists, Drs. Sheldon Ungar and Dennis Bray, noticed the rrs and the other media coverage of my paper and described the phenomena that they observed in their own January 2005 paper [27]. They described in detail the "efforts to prevent the making of specific scientific claims in any or all of the arenas in which these claims are typically reported or circulated" as they related to my BMJ paper. Their "results suggest that the public consensus about the negative effects of passive smoke is so strong that it has become part of a regime of truth that cannot be intelligibly questioned." Given all the controversies involving other epidemiologic risk factors, such as, hormone replacement therapy, air pollution, and vitamin supplements, this state of affairs regarding ETS is quite amazing. Indeed, the evidence regarding the lethality of ETS is not "a regime of truth," but collection of weak results that have turned into a "causal" relationship by carefully chosen committees. As I will discuss later, the epidemiologic evidence on this subject has changed in recent years and needs to be completely and objectively reassessed in order to reach a valid conclusion.

Authors and Editor Defend the BMJ Paper

The attack described above was quite startling to me as someone whose honesty and scientific integrity had never been questioned during the 33-year period from July 1970, when I received my Ph.D. [28], until May 2003 [1]. It was also startling that the attack was initiated by the ACS, the very organization that had given me the original California Cancer Prevention Study (CA CPS I) data in 1991 upon which the *BMJ* study was based. Kabat and I dealt with some of the initial controversy by responding to specific criticisms in our August 30, 2003 *BMJ* letter [29] and in our January 31, 2004 *Lancet* letter [30]. In particular, in these letters we refuted the five false ACS statements shown above:

- 1) This was not a "Tobacco Industry Study," but rather a UCLA study conducted by two qualified epidemiologists with ACS cooperation up until publication of the *BMJ* paper. This was not "Part of Organized Effort to Confuse Public About Secondhand Smoke", but rather it was an accurate representation of the results of one study. The tobacco industry played no role in the conduct, writing, or publication of the paper, and did not even know it was being published until it appeared.
- 2) It is a complete fabrication that "Society researchers repeatedly advised Dr. Enstrom that using CPS-I data to study the effects of secondhand smoke would lead to unreliable results." Indeed, the ACS Vice President for Epidemiology prior to Thun worked closely with me on the overall CA CPS I follow-up study from 1991 until 2001 because he felt that this was a valuable project. He was a co-author on the first version of the ETS and mortality paper when it was submitted to the *New England Journal of Medicine* in 2001 and was co-author on my first publication based on the CA CPS I cohort, which dealt with smoking cessation and mortality trends [31]. He was not able to remain as co-author on the ETS and mortality paper after 2001 because of his retirement from the ACS and his growing distance from the project.
- 3) It is absolutely false that "this study is neither reliable nor independent." First, this study is just as reliable as other epidemiological studies that have been conducted in a similar manner Indeed, the *BMJ* peer review process found that the results of the study were sound and sufficiently reliable to be worthy of publication and the ACS has thus far identified no specific errors in the study. Second, the study was conducted independent of influence from both the ACS and the tobacco industry.
- 4) It is absolutely false that "The study suffers from a critical design flaw: the inability to distinguish people who were exposed to secondhand smoke from those who were not." This cohort study was done in the same way as the

other spousal smoking studies and our 1999 follow-up questionnaire survey results clearly showed that there were subjects who had varying degrees of exposure to ETS as shown in Tables 1 and 2 of the *BMJ* paper. This issue was clearly addressed in the *BMJ* paper in response to Thun's 1999 concerns about this issue [32].

5) It is absolutely false that "exposure to secondhand smoke was so pervasive [in 1959] that virtually everyone was exposed to ETS, whether or not they were married to a smoker." The results of the 1999 survey shown in Table 4 of the *BMJ* paper clearly showed that among never smokers married to never smokers as of 1959, 43.5% of males and 61.7% of females reported no regular exposure to cigarette smoke from others in work or daily life as of 1999.

Although the ACS disputes the validity of my 1999 survey, they have not conducted their own ETS exposure survey of the approximately 50 million Americans who were born before 1950 and who are currently alive. Such a survey would yield actual evidence as to whether or not all Americans alive during the 1950s and 1960s were equally exposed to ETS. The ACS cannot simply make an unsubstantiated claim that "virtually everyone was exposed to ETS" and expect this claim to negate all the evidence presented in my *BMJ* paper.

In addition to the published letters cited above, we submitted to the BMJ on June 30, 2003 Manuscript BMJ/ 2003/084269, a detailed commentary that vigorously defended specific aspects of our BMJ paper. We showed that there was, in fact, substantial agreement between our results regarding ETS and those of the ACS and pointed out inconsistencies in ACS findings that had not been previously noted. Unfortunately, on September 19, 2003 the BMJ declined to publish this commentary, which would have helped resolve the controversy that had erupted over our BMJ paper. We then spent over two years attempting to publish various portions of this commentary in other journals until we successfully published in 2006, as described in our January 24, 2006 rr to bmj.com [33]. Portions of Manuscript BMJ/2003/084269 are presented later in this paper and the entire manuscript is posted for historical reference [34].

In spite of the numerous attacks described above, the *BMJ* has stood behind the *BMJ* paper since its publication. For all of the vehemence of the rrs, only about 3% referred to actual data in the paper and none identified anything approaching scientific error or scientific fraud [16]. Indeed, our paper was ranked among the "Top tens from bmj.com" in 2003 [3531]. *BMJ* Editor Richard Smith strongly defended his decision to publish the paper on both May 18, 2003 [36] and August 30, 2003 [37]. Fur-

thermore, Smith again defended this decision in his 2006 book, *The Trouble with Medical Journals*, in which he stated "it would be antiscience to suppress systematically one source of research" [38]. To date, no impropriety, bias, or omission has been identified in the review process and no error in the results has been identified in the paper, not even by Thun, who is in a position to check our findings and to publish additional findings.

Support for the BMJ paper from Other Epidemiologic Research

To further document the validity of our BMJ findings, Kabat and I compared them with the other U.S. epidemiologic evidence on ETS and coronary heart disease (CHD), in our 2006 peer-reviewed meta-analysis of environmental tobacco smoke and CHD mortality in the United States [39]. This comprehensive meta-analysis focuses on the U.S. cohort studies of ETS and CHD death in never smokers. These cohort studies are all fairly similar in design; ETS exposure was approximated by spousal smoking; CHD death was the endpoint; and they constitute virtually all the U.S. evidence and the majority of the world-wide evidence. In contrast to the previous major meta-analyses on this topic, such as the one in 1999 by Thun [32], our analysis includes the results of our 2003 study and the 1995 study by LeVois and Layard based on CPS I data [40]. We have applied consistent criteria to the selection of results included in the analysis. The results are summarized in terms of overall relative risks and doseresponse relationships. In addition, available data on misclassification of ETS exposure, personal monitoring of actual ETS exposure, and dose-response data for active smoking are discussed in order to characterize the estimates of ETS exposure in epidemiologic studies.

Contrary to the claims of the ACS and other critics, our results do not differ in any material way from those of the other studies, particularly for females. A further example of the ACS misrepresentations on the ETS issue can be found in the following simple comparison of statements about the findings in their major 1982 Cancer Prevention Study (CPS II) cohort. In the May 15, 2003 ACS press release Harmon J. Eyre, MD, stated: "CPS-II is one of more than 50 studies now published that have shown nonsmokers married to smokers have an increased risk of lung cancer" [8,9]. But, the 1995 doctoral dissertation based on CPS II by Victor Cardenas, "Environmental tobacco smoke and lung cancer mortality in the American Cancer Society's Cancer Prevention Study II", was inconclusive [41]. The dissertation abstract states: "This study found no evidence of an association between self-reported ETS and lung cancer risk among nonsmokers. However, using spousal smoking habits to assess exposure, we found ETS is only weakly, and not statistically significantly, related to lung cancer risk among nonsmoking women in seven years of follow-up of the CPS II cohort." [41]. Even though our findings are entirely consistent with Cardenas' findings, Eyre impugned our study with his statement: "Bad science can haunt us for generations. And regrettably, if questionable studies make it to publication, the damage is done." [8,9].

Furthermore, we specifically refuted the unsubstantiated claim by Thun that our BMJ study is "fatally flawed because of misclassification of exposure" [42]. Thun implied that virtually everyone in the U.S. during the 1950s and 1960s was equally exposed to ETS because it was so pervasive. Results from four independent surveys, as well as our 1999 CA CPS I survey, show that Americans were not equally exposed to ETS. Additional surveys show that exposure to ETS comes primarily from spousal smoking, not public smoking, particularly for females. Indeed, there was a clear relationship between spousal smoking and self-reported ETS exposure among never smokers who lived a major portion of their life before the introduction of restrictions on public smoking in the 1970s. One of these surveys is contained in the 1995 Cardenas dissertation [41]. Although Thun served on the Cardenas dissertation committee, to my knowledge, he has never cited results from this dissertation.

We found that when all relevant studies are included in the meta-analysis and the results of the individual studies are appropriately combined, current or ever exposure to ETS, as approximated by spousal smoking, is associated with roughly a 5% increased risk of death from CHD in never smokers, not the widely cited 25% in the meta-analyses of Thun and others. Furthermore, we found no doseresponse relationship and no elevated risk associated with the highest level of ETS exposure in males or females.

Another paper which sheds light on the CPS II findings concerning ETS is a 1995 analysis which linked data on ambient air pollution from 151 U.S. metropolitan areas with mortality data from CPS II individuals who resided in those areas [43]. The results of this analysis showed that in never smokers there was a statistically significant association of all cause mortality with both sulfate and fine particle concentrations after controlling for covariates, including "hours per day of ETS exposure." The authors, one of whom was Thun, did not report the specific results for the confounding variable of ETS exposure. However, in order to resolve a major dispute over the validity of the results in this air pollution analysis [44], a reanalysis was conducted in 2000 by the Health Effects Institute (HEI) [45]. The Cox proportional hazards regression model (PHREG) results included in Appendix F of the resulting HEI Reanalysis Report make it clear that the independent variable "passive" (hours per day of ETS exposure) shows no association with mortality from lung cancer, cardiop-

ulmonary disease, or all causes in never smokers [46]. Results are shown as a relative risk (RR) and 95% confidence interval (CI). For lung cancer in CPS II, RR(passive) = 1.020 (0.938-1.110) for males, 1.004 (0.995-1.013) for females, and 1.005 (0.957-1.055) for both sexes. These relative risks agree well with those in my CA CPS I study, where RR(7 level index) = 0.88 (0.70-1.10) in males and RR(8 level index) = 0.97 (0.90-1.05) in females. For cardiopulmonary diseases in CPS II, RR(passive) = 1.004 (0.987–1.021) for males, 1.015 (1.000– 1.029) for females, and 1.010 (0.999-1.021) for both sexes. For all causes in CPS II, RR(passive) = 0.996 (0.984-1.009) for males, 1.004 (0.995-1.013) for females, and 1.001 (0.994-1.009) for both sexes. A key portion of the actual PHREG computer printout for these diseases for males, females, and both sexes has been assembled and posted [46]. The PHREG program used in the CPS II study [43,46] is the same as that used in the CA CPS I study [1].

My BMJ results for coronary heart disease are also consistent with those in the Western New York State study published in the October 9, 2006 Archives of Internal Medicine, which found "After adjustment for covariates, exposure to secondhand smoke [SHS] was not significantly associated with an increased risk of myocardial infarction [MI]" [47]. Furthermore, this study concluded "Exposure to SHS has declined sharply among nonsmokers in recent years. In the absence of high levels of recent exposure to SHS, cumulative lifetime exposure to SHS may not be as important a risk factor for MI as previously thought." This study was entirely independent of my study and was done without tobacco industry funding and came to the same conclusion with regard to heart disease. Finally, my BMJ results for lung cancer in the CA CPS I cohort are consistent with those of the original 1981 ACS analysis of the nationwide CPS I cohort [48]. This analysis examined lung cancer mortality during 1960-1972 and found "Compared with nonsmoking women married to nonsmoking husbands, nonsmokers married to smoking husbands showed very little, if any, increased risk of lung cancer." This analysis was entirely funded by and conducted by ACS and came to the same conclusion as my BMJ analysis.

Ongoing Misrepresentations Regarding ETS

Much of the evidence above is not being properly presented and there is misrepresentation of other evidence. For instance, serious misrepresentation of CPS II results is evident when one examines the 1997 Cardenas peerreviewed paper [49], which was based on the 1995 Cardenas dissertation [41]. Table 4 of the Cardenas paper presents exposure to spousal smoking among women by the husband's level of smoking, but is deceptively labeled. Women with the highest level of exposure, labeled "40+

cpd by spouse", have a RR of 1.9 (95% CI 1.0-3.6) and the P for dose-response trend is 0.03 (cpd = cigarettes per day). However, Table 38 of the Cardenas dissertation makes clear that the RR for spouses of current smokers of 40+ cpd is only 0.9 (95% CI 0.2-3.9) and the P for trend is 0.34. If it were not for Table 38 the reader would not know that Table 4 is based on the combination of current and former smokers. This combination of current and former smokers by cpd is highly unorthodox, has not been done in other ETS studies, and is not meaningful for assessing a trend based on current spousal smoking. The Cardenas dissertation makes it very clear that there is no dose-response relationship between spousal smoking and lung cancer in CPS II. Key sections of Cardenas' Tables 4 and 38 are shown side by side in Table 1 and they reveal a serious discrepancy in the presentation of the same data. Because Cardenas' Table 38 appears to present the underlying findings and because these findings contradict Eyre's statement above, the ACS should clarify this major discrepancy. However, no clarification has been made and only the positive dose-response relationship in Cardenas' Table 4 is ever cited [49].

For instance, Cardenas' Table 4 findings are now cited in the 2004 WHO IARC Monograph 83 "Tobacco Smoke and Involuntary Smoking" [50]. This major 1452-page report contains a review of the epidemiologic evidence on ETS and lung cancer on pages 1231–1271 [51]. The section "Exposure-response relationships" on page 1236 contains the statement "The study by Cardenas *et al.* (1997) also found a significant exposure-response relationship. When the husbands smoked 1–19, 20–39, \geq 40 cigarettes/day, the relative risks for women exposed to secondhand smoke were 1.1, 1.2, and 1.9 respectively (p value for trend test, 0.03)".

In addition, a January 2004 J Natl Cancer Inst (JNCI)summary of IARC Monograph 83 shows results for ≥40 cigarettes/day in Table 3 and it contains the erroneous value RR = 1.9 [52]. Obviously Thun, a member of the IARC Working Group for Monograph 83, did not notify the IARC Working Group about the 1995 Cardenas dissertation. This type of selective analysis and presentation of results has been termed "publication bias in situ" and it is often difficult to detect [53]. I was able to detect this irregularity only because I knew of the Cardenas dissertation. In other scientific fields, the type of data manipulation done in Cardenas' Table 4 would most likely be treated as a serious ethical violation. Also, it is noteworthy that 14 authors of the JNCI article signed an August 30, 2003 BMJ letter criticizing my BMJ paper, but then made no mention of my paper in their January 2004 JNCI article.

Continuing ACS Campaign to Discredit the BMJ Study

Although I have refuted the erroneous statements in their May 15, 2003 press release, the ACS has shown no interest in correcting the record with regard to me and my research. Their press release has been posted on up to 1,000 locations on the Internet during the past four years, based on Google searches of the phrase "American Cancer Society Condemns Tobacco Industry Study." It is still posted on many websites in addition to ACS's own website. Our *BMJ* and *Lancet* letters and our new meta-analysis defending the validity of our *BMJ* paper are being ignored by the ACS. Instead, the ACS and other activist organizations continue to post defamatory information about us and our research.

Our new meta-analysis shows that the relationship between ETS and CHD in U.S. never smokers is very weak (estimated relative risk of 1.05 with no dose-response relationship) [39]. Yet the ACS continues to state in their 2007 "Cancer Facts and Figures" that "ETS causes an estimated 35,000 deaths from heart disease in persons who are not current smokers" (page 36) [54]. The source the ACS uses for this CHD death estimate is a 1992 JAMA paper [55], even though more than 90% of the U.S. epidemiologic evidence has been published since 1992. Our new meta-analysis shows that the vast majority of the existing U.S. evidence originates from the ACS CPS I and CPS II cohorts, yet the ACS simply ignores or dismisses most of this evidence. The CPS I and CPS II evidence is summarized in Table 2, which is taken from Table 6 of our meta-analysis paper [39].

Continuing Glantz Campaign to Discredit Enstrom

Beginning with his activities at the time of the publication of our BMJ paper, Glantz has continually attacked me and my research, in spite of the fact that we are both established, long-term faculty members in the University of California system. Glantz is well-known as a long-time anti-smoking activist [10,56], whose ultimate goal is achieving a society free of smokers [57]. However, as a UC faculty member, he is supposed to adhere to the UCSF Campus Code of Conduct [58] and the UC Standards of Ethical Conduct [59]. For instance, the Code of Conduct states "Misconduct or Misconduct in Science means fabrication, falsification, plagiarism, or other practices that seriously deviate from those that are commonly accepted within the scientific community for proposing, conducting, or reporting research." The UC Standards of Ethical Conduct states "Members of the University community are expected to conduct themselves ethically, honestly, and with integrity in all dealings."

However, based on his clearly documented written and verbal attack on me, he has not adhered to these codes. Indeed, I have spent the past four years responding to his

http://www.epi-perspectives.com/content/4/1/11

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Table 1: Comparison of the CPS II dose-response results of 1995 Cardenas disseration and 1997 Cardenas paper: relative risk (RR & 95% CI) of lung cancer death by ETS exposure (spousal smoking) among female never smokers in CPS II. Definition in 1995 Cardenas dissertation [41]: 'Analyses restricted to nonsmoking spouses married to nonsmoking spouses and those married to cigarette smokers (and not other type of tobacco), with complete smoking data, married once at a time of interview, and with valid data on age at first marriage. Definition in 1997 Cardenas paper [49]: 'The referent group includes never-smoking women married to husbands who did not smoke during the marriage. The exposed categories are split into approximate tertiles, and are restricted to never-smokers married to cigarette smokers with complete smoking data, married once, and with valid information on age at marriage.'

	1995	Cardenas dissertation [41]			15	997 Cardenas paper [49]	1
Spousal smoking (cigarettes per day)	Deaths/Person-years	1982–89 CPS II Fully-adjusted RR (95% CI)	Deaths/Person-years	1982–89 CPS II Fully-adjusted RR (95% CI)	Cigarettes per day by spouse	Deaths/Person-years	1982–89 CPS II Fully-adjusted RR (95% CI)
	Table 38 as s	shown on page 117	Proper summ	nary of Table 38 data	Table 4: im	proper summary of Tab	le 38 data
Never	30/311,333	1.0	30/311,333	1.0	0 (never)	30/333,946	1.0
Former(I-I9)	4/61,677	0.6 (0.2–1.8)					
Former(20-39)	12/120,585	0.8 (0.4–1.7)					
Former(40+)	11/49,304	2.0 (1.0-4.0)					
Former – total			27/231,566	1.13 (0.72–1.78)			
Current(I-I9)	5/32,524	1.7 (0.7–4.4)	5/32,524	1.7 (0.7–4.4)	I–I9 (current or former)	9/83,074*	1.1 (0.5–2.2)
Current(20–39)	10/69,060	1.6 (0.8–3.4)	10/69,060	1.6 (0.8–3.4)	20–39 (current or former)	22/179,751*	1.2 (0.7–2.2)
Current(40+)	2/24,900	0.9 (0.2–3.9)	2/24,900	0.9 (0.2–3.9)	40+ (current or former)	13/71,618*	1.9 (1.0–3.6)
P test for trend for 'former'		P = 0.29					
P test for trend for 'current'		P = 0.34			P test for 'current	or former'	P = 0.03

^{*} Current and Former Combined

Table 2: Dose-response relationship between ETS exposure and CHD mortality. Relative risk of spousal smoking related to CHD deaths among never smokers in CA CPS I [1], CPS II [82], and CPS I [40] and in the summary RR of the three studies.

Spousal smoking	1960–98 CA CPS I Age-adjusted RR (95% CI)	I 982–89 CPS II Fully-adjusted RR (95% CI)	1960–72 CPS I Age-adjusted RR (95% CI)	Summmary'Age-adjusted' RR (95% CI)
	Enstrom [1] (extracted from Tables 7 & 8)	Steenland [82](extracted from Table 2)	LeVois [40] (extracted from Table 4)	Enstrom + Steenland + <u>LeVois</u>
<u>Males</u>				
Vever	1.00	1.00	1.00	1.00
ormer	0.94 (0.78–1.12)	0.96 (0.83–1.11)	0.95 (0.83-1.09)	0.95 (0.87–1.04)
current				
I-19 cigs/day	0.91* (0.78–1.06)	1.33 (1.09–1.61)	0.99 (0.89–1.09)	1.02 (0.94–1.10)
20 cigs/day	0.92 (0.74–1.15)	1.17 (0.92–1.48)		
20+ cigs/day			0.96* (0.83–1.11)	1.02 (0.92–1.12)
21+ cigs/day	1.20* (0.88–1.64)	1.09 (0.77–1.53)		
- - - - - - - - - - - - - - - - - - -				
Never	1.00	1.00	1.00	1.00
ormer	1.02 (0.93–1.11)	1.00 (0.88–1.13)	0.99 (0.93–1.05)	1.00 (0.95–1.05)
Current				
I-19 cigs/day	1.07* (0.96–1.19)	1.15 (0.90–1.48)	1.04 (0.97–1.12)	1.05 (0.99–1.12)
20 cigs/day	1.04 (0.92–1.16)	1.07 (0.83–1.40)		
20–39 cigs/day			1.06 (0.98–1.15)	1.04 (0.98–1.10)
21–39 cigs/day	0.95 (0.80–1.12)	0.99 (0.67–1.47)		
40+ cigs/day	0.83 (0.65–1.06)	1.04 (0.67–1.61)	0.95 (0.78–1.15)	0.92 (0.79–1.06)

^{*} indicates RR was based on combining other RRs

false and misleading statements and defending my honesty and scientific integrity. The full details of his campaign are too extensive to present here, but the selected examples below demonstrate the tactics that he used against me and the epidemiologic research that I have been conducting at UCLA.

On July 25, 2003 Neal L. Benowitz, MD, UCSF Professor of Medicine, and Glantz co-wrote an eight-page letter to the UC Vice Provost for Research Lawrence Coleman which attempts to make the case that acceptance of tobacco industry funding for research violates current Regents and University policy and should be ended [60]. On pages 3 and 4 of this letter they claim: "The most recent example of how the tobacco industry uses funding of university research as part of its for propaganda campaign is a May 17, 2003 study from UCLA on the health effects of secondhand smoke published in the British Medical Journal. . . . There is little possibility that it will be taken seriously in scientific circles. . . . this paper would go down as one bit of poor research done at a university with a reputation for high quality scholarship that slipped into a good journal because of the foibles of the peer review process."

On March 8, 2005 Glantz participated with other UC faculty members in a San Francisco based KQED radio program entitled "Funders and Academic Research: Forum assesses the controversy surrounding the relationship between funders and academic research," which can be listened to on the Internet and audio files [61]. During this program Glantz attempted to discredit well qualified scientists and their peer-reviewed research publications by inappropriately linking them to the tobacco industry. The "scandal" about me and my BMJ study was discussed during minutes 17-19 of this 52 minute program, when Glantz made several clearly false and inflammatory statements. First, Glantz claimed that the BMI study "was not funded by the American Cancer Society," but was "done with Philip Morris' money." Actually, the study was funded by ACS from 1959 to 1990, by the UC Tobacco-Related Disease Research Program from 1991 to 1997, and by the Center for Indoor Air Research (CIAR) from 1998 to 2003. Philip Morris provided no direct funding for this study and had no role in its conduct. Then, Glantz stated that I was "a damn fool" who was told by ACS that I "made inappropriate use of the data", an unsubstantiated claim made only after Glantz and ACS learned of my results. Then, Glantz implied that I was "advocating a protobacco position" when I have never done so. Finally, Glantz claimed "the science that the UCLA study did was crap", whereas it clearly conformed to the standards of epidemiologic research. These statements indicate the unprofessional approach used by Glantz to attack scientific findings with which he disagrees and to advocate positions that are not supported by the facts.

Glantz's arguments for banning tobacco industry funding of research at UC have been rejected in favor of academic freedom. The UC administration has expressed its strong support for academic freedom and UC Vice Provost for Research Coleman has stated "Academic freedom must be absolute or no one has it" [62]. On May 11, 2005 the UC Academic Senate adopted a strong Academic Senate Resolution on Research Funding Sources which clearly supports the right of individual UC faculty members to accept research support from any source, including the tobacco industry, as long as this funding adheres to University policy [63]. In spite of this strong faculty resolution, in September 2006 Glantz brought the issue of a ban on tobacco industry funding to the UC Regents, the governing body of the University [64]. Glantz cited my BMJ study as one rationale for such a ban in written documents [65] and in a January 18, 2007 presentation before the UC Regents [66]. The UC Regents requested advice on this issue from the UC Academic Senate, which spent several months carefully evaluating the matter [67]. My perspective, including a defense of my research, my funding, and my scientific integrity, was presented to the UC Academic Senate in April 2007 [68]. In May 2007 representatives of the UC Academic Senate voted almost unanimously (15 to 1 by the Academic Council and 44 to 5 by the Academic Assembly) in favor in academic freedom and against a proposed ban on tobacco industry funding advocated by Glantz [69-71].

One final example of Glantz's unprofessional treatment of my research is contained in his May 24, 2005 Circulation report, where he attempts to make the case that passive smoking has nearly the same impact as active smoking on cardiovascular effects [72]. In his meta-analysis of the relation between ETS and CHD, he found "The pooled relative risk computed with a random-effects model (computed with Stata Version 7) was 1.31 (95% CI, 1.21 to 1.41), similar to the estimates of earlier metaanalyses." To achieve this result, he omitted the two largest studies, which represent a major portion of the available evidence. My BMJ study, which began in 1960 [1], was omitted based on his unsubstantiated claim that it had "serious misclassification bias" and the 1995 study by LeVois and Layard, which also began in 1960 [40], was omitted without comment and was not even cited. However, Glantz included the other cohort studies which began in the 1960s and 1970s without any comment about their misclassification bias. Kabat and I fully addressed all these studies and the issue of misclassification bias in our 2006 meta-analysis [39]. Glantz's biased analysis regarding the relation between ETS and CHD is

evident when his 2-page 2005 meta-analysis [72] is compared with our 12-page 2006 meta-analysis [39].

Jonathan M. Samet, M.D., and the 2006 Surgeon General's Report

False and misleading statements about my research were also made by Jonathan M. Samet, M.D., M.S., who has played a prominent role in reviews of the epidemiologic evidence on ETS for over 20 years. First, Samet made a statement that neither he nor anyone else has substantiated in the May 16, 2003 Los Angeles Times, when he described my BMJ paper as "one very flawed study" that "just doesn't contribute" [22]. Then, he co-signed serious accusations about my research that appeared in a May 30, 2003 BMJ rapid response [20] and an August 30, 2003 BMJ letter [73]. These two items stated "Enstrom and Kabat's conclusions are not supported by the weak evidence that they offer, and although the accompanying editorial alluded to 'debate' and 'controversy', we judge the issue to be resolved scientifically, even though the 'debate' is cynically continued by the tobacco industry." To understand the outlandish nature of these accusations, recall that we used a large and highly respected dataset and accepted epidemiologic methods; we reported study details in the paper itself, in the "Prepublication History", and in our subsequent letters; we have supported our conclusions to a greater extent than can be found for any other study of ETS and mortality; our methods have never been substantively challenged; and our results are consistent with the entire body of U.S. evidence [39].

These statements from Samet might have been somewhat plausible if he had any evidence that there were errors in my 2003 paper or that I was "pro-tobacco" based on my research before 2003. But neither he nor other critics have made a plausible case for fundamental errors in my paper, and I have never been "pro-tobacco." Samet has been aware of my epidemiologic research since we both participated in the August 23-25, 1978 National Cancer Institute Workshop on "Populations at Low Risk of Cancer" held in Snowbird, Utah. The proceedings of the workshop, including the list of participants, were published in JNCI in November 1980 [74]. I gave three talks at this Workshop and two of them described the reduced cancer death rates among nonsmokers, one dealing with Mormons [75] and another dealing with a representative sample of U.S. nonsmokers [76]. Indeed, I have investigated the healthy lifestyles of Mormons and other nonsmokers during my entire epidemiologic career [77,78].

Further evidence of Samet's willingness to dismiss scientific evidence when it does not support his agenda appears in the June 27, 2006 release and publication of the 727-page Surgeon General's Report on "The Health Consequences of Involuntary Exposure to Tobacco Smoke" [79].

Samet was the Senior Scientific Editor of this report and the most influential epidemiologist involved with the report [80]. In addition, Glantz was a Contributing Editor and Thun was a Reviewer on this report. Although Samet, Thun, and Glantz were fully aware of the importance of my BMJ paper, as evidenced by their extensive efforts to discredit it, the paper was simply omitted from the Surgeon General's Report without comment. A search for "enstrom j" of the entire PDF version of the report [79], reveals that the only mention of the BMJ paper is in the Appendix on page 673, where it is listed as one of the papers not included in the report. Another search reveals that the BMJ paper was omitted without explanation from the database for the Report [81]. This database was prepared by Johns Hopkins University and the Centers for Disease Control and Prevention's Office on Smoking and Health. It includes "approximately 900 key articles regarding involuntary smoking and disease outcomes" and supposedly "reflects the most recent findings in the scientific literature."

In order to illustrate the selective and unscientific nature of this omission, I examined the references used in Chapters 1-10 of the Surgeon General's Report and the references in the Appendix that were not used. Of 38 total references from 2003, 33 were used in Chapters 1-10 and only 5 references, including the BMJ paper, were not used. Of 71 references from 2004, 53 were used and 18 were not used; of 39 references from 2005, 26 were used and 13 were not used; of 22 references from 2006, 7 were used and 15 were not used. In summary, the report used 119 references from 2003-2006, but omitted without comment the 2003 BMJ paper. The BMJ paper was the only U.S. study relating ETS to lung cancer and coronary heart disease that was omitted. Because of this omission, the Surgeon General's Report does not accurately reflect all the peer-reviewed epidemiologic evidence on the relation of ETS to lung cancer and coronary heart disease mortality in the U.S.

Chapter 7, page 423, reports: "This chapter considers the full body of evidence on secondhand smoke exposure and lung cancer published through 2002, the ending date for the systematic review of the epidemiologic studies." Based on comparing never smokers ever married to a smoker with never smokers never married to a smoker, a world wide relative risk (RR) of 1.21 (1.13–1.30) was reported on page 435. However, there is no reason for an ending date of 2002, given that other sections of the report cite results published during 2003–2006 (by my count 119 such publications are cited). It appears that the ending date of 2002 was intentionally selected in order to exclude my 2003 *BMJ* results. Consequently, the above worldwide RR is misleading because it does not reflect that fact that my results substantially weaken the U.S. evidence [1,29].

My own meta-analysis of all U.S. spousal smoking studies, yields a U.S. RR of 1.10 (1.00–1.21), which barely constitutes a relationship.

Chapter 7 contains this inaccurate statement on page 435: "There were no significant differences in the RR estimates by geographic area; the point estimate was 1.15 (95 percent CI, 1.04–1.26) for studies conducted in the United States and Canada, 1.16 (95 percent CI, 1.03–1.30) for studies conducted in Europe, and 1.43 (95 percent CI, 1.24–1.66) for studies conducted in Asia." Obviously, the RR = 1.43 for studies in Asia is statistically greater than the RR = 1.15 for studies in U.S. and Canada and the RR = 1.16 for studies in Europe. Indeed, there is substantial variation around the world and all these results cannot be accurately represented by a single RR of 1.21. This geographic variation should have been properly acknowledged in the Report.

Chapter 8 contains on page 521 selective criticism about and dismissal of the analysis by LeVois and Layard of ETS and CHD deaths in the ACS CPS I and CPS II studies [40]. This paper is important because of its size and statistical power, as discussed in our 2006 meta-analysis of ETS and CHD deaths in the U.S. [39]. One basis for the dismissal is the inaccurate statement, "The investigators did not distinguish between current exposures from spousal secondhand smoke and former exposures, nor did they separately report the effect of current spousal smoking on the risk of CHD." Table 4 of the LeVois and Layard paper clearly shows results for three levels of current ETS exposure for both males and females. Furthermore, Table 2 summarizes the dose-response relationship between ETS and CHD deaths based on the results from the three largest U.S. studies [1,40,82]. There is no meaningful difference in the results for these studies and no dose-response relationship in any of them.

Furthermore, note that the meta-analysis of ETS and CHD is summarized in Figure 8.1 on page 524. Since this figure only shows studies through 2001 it obviously omits the 2003 *BMJ* study.

The *BMJ* study has a major impact on the meta-analysis, as pointed out in our 2003 *BMJ* letter [29] and our 2006 meta-analysis [39]. Note that inclusion of *BMJ* results yields a relative risk (RR) of CHD death in the U.S. of 1.05 (0.99–1.11), based on a comparison of current to never exposure to ETS. This is much less than the summary RR (exposed/unexposed) of 1.27 (1.19–1.36) contained in Figure 8.1. The Surgeon General's Report should have pointed out that the ETS and CHD relationship is much larger outside of the U.S. than it is within the U.S. We estimated that the RR outside the U.S. is approximately 1.5 [39] and the 1999 Thun meta-analysis found the RR was

1.41 (1.21–1.65) [32]. This large difference between the RRs within the U.S. and those outside of the U.S. is worthy of further discussion and investigation, in order to determine if it is a real difference or an anomaly due to methodological issues.

The Introduction of the Surgeon General's Report makes the statement that "about 50,000 excess deaths result annually from exposure to secondhand smoke (Cal/EPA 2005). Estimated annual excess deaths for the total U.S. population are about 3,400 (a range of 3,423 to 8,866) from lung cancer, 46,000 (a range of 22,700 to 69,600) from cardiac-related illnesses, and 430 from SIDS." [79]. Given the fact that the two largest epidemiologic studies on ETS and tobacco-related mortality [1,40] have been omitted from the Surgeon General's Report and the fact that these two U.S. studies suggest a substantially weaker ETS and mortality relationship in the US, the above estimate of excess deaths appears to be an intentional exaggeration of what the entire body of scientific evidence shows. A complete evaluation of all the peer-reviewed U.S. epidemiologic evidence suggests that ETS exposure is associated with a much smaller number of lung cancer and CHD deaths in U.S. never smokers. Furthermore, there is not a "causal" relationship by traditional epidemiologic standards.

An August 23, 2006 "research news and perspective" report in IAMA questioned various aspects of the Surgeon General's Report, particularly findings regarding the acute effects of small amounts of ETS exposure and the claim by the Surgeon General that "There is no safe level of exposure to secondhand smoke" [83]. This JAMA report is particularly noteworthy because it quotes two experts who have extensive experience regarding the ETS issue. Michael Siegel, MD, MPH, a professor of social and behavioral sciences at Boston University School of Public Health and a prominent tobacco control researcher, told JAMA "We're really risking our credibility [as public health professionals or officials] by putting out rather absurd claims that you can be exposed briefly to secondhand smoke and you are going to come down with heart disease or cancer. People are going to look at that and say that's ridiculous." Siegel's own paper expanding on this point is published alongside the present article [84]. Furthermore, since March 2005, Siegel has posted many detailed and insightful analyses regarding ETS and tobacco control on his personal website, "The Rest of the Story: Tobacco News Analysis and Commentary" [85]. Each post includes "Comments" from readers who provide additional insights. For instance, on June 28, 2006, he posted "Surgeon General's Communications Misrepresent Findings of Report; Tobacco Control Practitioners Appear Unable to Accurately Portray the Science" [86].

John C. Bailar III, MD, PhD, a prominent epidemiologist and biostatistician, who is Professor Emeritus at the University of Chicago, told JAMA "It doesn't make sense for the cardiovascular risk of secondhand smoke to be as high as one third of the risk from direct smoking. . . . That's a far bigger ratio than risk for lung cancer and it's hard for me to believe that it's real" [83]. These comments are similar to those in his March 25, 1999 NEJM editorial on ETS and coronary heart disease, in which he stated "I regretfully conclude that we still do not know, with accuracy, how much or even whether exposure to environmental tobacco smoke increases the risk of coronary heart disease" [87]. On June 7, 2006, just 20 days before the release of the Surgeon General's Report, the Select Committee on Economic Affairs of the House of Lords in London issued an important report on the management of risk, which suggests that passive smoking in England may be a relatively minor health risk [88]. The committee obtained testimony from Professor Sir Richard Peto of the University of Oxford on February 14, 2006 [89]. Sir Richard's testimony clearly states the substantial doubt that he has about the quantitative health risks of passive smoking [90,91]. The very fact that two major reports published in the same month, June 2006, come to substantially different conclusions about the health risks of ETS indicates that these risks are still uncertain and difficult to measure accurately.

Further evidence of the uncertainty regarding the health risks of ETS is contained in the June 28, 2007 *Nature* news article on ETS. Various claims made by Glantz about the acute and chronic health effects of ETS are questioned by Peto, Bailar, and Siegel, who restated their concerns that the dangers of ETS have been exaggerated [92]. For instance, Peto stated "Passive smoking must kill some people, but the big question is how many." This statement clearly underscores the existing uncertainty and directly contradicts the June 27, 2006 statement by U.S. Surgeon General Richard H. Carmona that "The debate is over" regarding the health effects of secondhand smoke [93].

Jonathan M. Samet, M.D., and United States of America v. Philip Morris USA, et al

One particularly pernicious aspect of the attack described above is the fact that my *BMJ* paper is now part of the largest (\$280 billion) Racketeer Influenced and Corrupt Organizations Act (RICO) lawsuit ever filed, United States of America v. Philip Morris USA, et al. [Civil Action No. 99-CV-02496(GK)] [94,95]. My research and I are described in a defamatory way on pages 821–830 within the section "Defendants Used Their Jointly Controlled Organizations to Promote Their Agenda Through Symposia, Publications and a Roster of Long-time Paid Scientists" of the 2543-page pretrial "UNITED STATES' FINAL PROPOSED FINDINGS OF FACT (July 2004)" prepared

by the U.S. Department of Justice (USDOJ) [96]. The trial took place in front of U.S. District Court Judge Gladys Kessler from September 2004 though June 2005 [94]. Additionally, my research and I are described in a defamatory way in several places in the 2454-page post-trial document "UNITED STATES' FINAL PROPOSED FINDINGS OF FACT (Incorporating Errata of August 16, 2005)" prepared by the USDOJ [97]. Specifically, my BMJ paper is listed on page vii of the Table of Contents under the category "Cooking the Books: The Manufacture of False Science to Support the Industry Position on ETS." On page 493 it is included among "examples of scientific fraud" and on page 589 it is described as "at best a contamination of the scientific literature and at worst a scientific fraud." It is discussed in detail on pages 609-615, where there are numerous false statements and distortions, such as, "the Enstrom/Kabat study is yet another self-serving, unreliable, and scientifically questionable product of the industry's unabated effort to attack the scientific consensus on passive smoking." Although no actual evidence was presented of errors in my study or of scientific misconduct on my part, the lawsuit makes it appear that I have engaged in scientific fraud.

The available evidence indicates that insertion of the BMJ paper was a collaborative effort of Glantz and Sharon Y. Eubanks (D.C. Bar No. 420147), Director of the USDOJ Tobacco Litigation Team from 1999 until December 2005, when she resigned from the USDOI [98]. The following brief in Civil No. 99-CV-02496 (GK), "REPLY IN SUPPORT OF THE UNITED STATES' THIRD MOTION TO COMPEL PRODUCTION OF DOCUMENTS WITHHELD BY BROWN & WILLIAMSON BASED ON ASSERTIONS OF PRIVILEGE OR PROTECTION," was prepared by Eubanks and signed on December 5, 2003. This brief is posted on the same listsery that Glantz has used to post other defamatory information about me [99]. Pages 8, 9, and 14 of this brief contain a misleading and distorted presentation of my alleged "ties" with the tobacco industry going back "nearly 30 years." This presentation later appeared in the July 2004 and August 2005 Findings of Fact of the USDOJ lawsuit. This 2003 brief does not present any evidence challenging my honesty as a scientist or the validity of the findings in my BMJ paper. It is simply an attempt to smear my reputation with inappropriately constructed "ties" to the tobacco industry, based on the fact that I had correspondence with the tobacco industry regarding my epidemiologic research.

On August 17, 2006 District Court Judge Gladys Kessler issued a 1,653 page Final Opinion concluding that the tobacco industry had engaged in racketeering [100,101]. Eleven key pages from her decision, including pages discussing my study, were assembled by Glantz and posted on a UCSF website [102]. The Kessler decision includes a

section entitled "The 2003 Enstrom/Kabat Study" on pages 1380–1383, as well as other references to my study. The Judge repeated in her opinion a number of the misleading and inaccurate statements about my study that are contained in the 2004 and 2005 Findings of Fact. However, the Judge identified no specific errors in the study and identified no scientific misconduct by me. At no time was I ever given an opportunity to challenge or refute the statements made about me and my research in the USDOJ Findings of Fact, in the trial itself, or in the Kessler opinion. I am now in the process of clearing my name in connection with this lawsuit and this paper represents a major step in that process. Furthermore, on October 31, 2006 the U.S. Court of Appeals of the District of Columbia Circuit granted the tobacco industry's emergency motion to stay Judge Kessler's final judgment and remedial order pending appeal [103]. On May 22, 2007 the U.S. Court of Appeals issued an order setting the briefing schedule for the appeal [104].

In formulating her comments about my study, Judge Kessler relied heavily on the testimony of Samet. On page 765 of her decision she states "Dr. Jonathan Samet, a Government expert with extraordinary qualifications, is a physician and epidemiologist with extensive experience treating patients with lung cancer and COPD." On page 1232 she states: "Dr. Samet is professor and chair of the Department of Epidemiology at the Johns Hopkins Bloomberg School of Public Health. He is also a licensed physician who is board certified in pulmonary and internal medicine. Dr. Samet is a member of the National Academy of Sciences' Institute of Medicine, the Board of Scientific Counselors of the National Cancer Institute, and EPA's Clean Air Scientific Advisory Committee. He is a recipient of the Surgeon General's Medallion and has participated as an author and/or editor of nine Surgeon General's Reports, including as Consulting Scientific Editor and author for the 1986 Report. He has participated in four NCI monographs in its series on smoking and health. He chaired the 2002 review of active and passive smoking and health for the International Agency for Research on Cancer of the World Health Organization. . . . after considering Dr. Samet's superb academic credentials, his vast experience working on Surgeon General Reports and NCI monographs, his continuing practice of medicine, as well as his demeanor and responsiveness to cross-examination, the Court fully credits his testimony." On page 1234 she states: "The Court accepts and credits Dr. Samet's conclusions, based on his expertise, as well as the other factual findings herein, that exposure to secondhand smoke causes lung cancer and coronary heart disease in adults and a number of respiratory diseases in children."

It is worth repeating the allegations in the Kessler decision, first to point out that they are the same false and mis-

leading claims about the Enstrom/Kabat study by the ACS, Samet, Glantz, and others that are described above, and second to show how obviously incorrect they are. The Enstrom/Kabat study was not "CIAR-funded and managed" and was not "funded and managed by the tobacco industry through CIAR and Philip Morris." Although the study was partially funded by CIAR, it was not managed by either CIAR or Philip Morris. Indeed, CIAR assigned its entire award for the study to UCLA in 1999 just before CIAR was dissolved as a condition of the Master Settlement Agreement [105]. CIAR did not even exist when my study was being completed. The study was conducted and published without any influence from the tobacco industry. The claim that the "American Cancer Society had repeatedly warned Enstrom that using its CPS-I data in the manner he was using it would lead to unreliable results" is utterly false and the ACS has produced no documentation to support this claim. The claim "Enstrom and Kabat's conclusions are not supported by the weak evidence that they offer" made by Samet and others is utterly false because our conclusions are fully supported by the evidence in our BMJ paper, as stated earlier.

In addition, Samet made an inaccurate and incomplete statement in his Written Direct testimony of September 20, 2004 (page 184, lines 8-9): "When the 2002 metaanalysis carried out by IARC was redone in 2004 to include this [Enstrom and Kabat] study, the positive findings were unchanged." [106]. This statement is inaccurate because the August 30, 2003 BMJ letter signed by Samet correctly states: "Adding the result from Enstrom and Kabat to the IARC analysis reduces the pooled estimate to 1.23." [73]. In addition, this statement is incomplete because Samet failed to state that the Enstrom and Kabat results reduced the pooled risk ratio estimates for U.S. studies to about 1.10 for lung cancer and to about 1.05 for coronary heart disease [39]. The Enstrom/Kabat summary risk ratios are far below the widely stated summary risk ratios of about 1.25 and are not consistent with the estimate that "about 50,000 excess deaths results annually from exposure to secondhand smoke" in the US, as stated on page 8 of the Surgeon General's Report [79].

Samet made a false statement in this September 20, 2004 testimony when he claimed (page 192, lines 21–23): "Except for the analyses of CPS I and CPS II presented by LeVois and Layard in 1995, all other studies have demonstrated at least a modest increase in risk for fatal and nonfatal CHD due to secondhand smoke exposure." [106]. Our *BMJ* study showed no increase in risk for fatal CHD, other than the insignificant statistical fluctuation that was also present in the LeVois and Layard paper, and reference to our study should have been included in Samet's testimony.

Since no errors had been found in our paper, and since Kabat and I had clearly declared there was no tobacco industry influence on our results (and no one has found any evidence to the contrary), our research did not warrant inclusion in the USDOJ lawsuit. The citation of our study in the Kessler decision appears to be primarily due to the false and misleading statements about our research made by Samet. All of this casts doubt on the ability of Samet to be objective regarding the subject of ETS.

Further evidence of Samet's campaign against me appeared in the May 4, 2007 Chronicle of Higher Education as a two-page, 15-inch by 22-inch advertisement "Why do the University of California Regents still cash checks from tobacco racketeers?" [107]. This advertisement by "Campaign to Defend Academic Integrity" [108] is an appeal to UC Regents to implement a tobacco funding ban and it makes direct reference to me and my tobacco industry funding. Statements throughout the advertisement falsely characterize me and my research: "To make vivid how Big Tobacco co-opted world-class research institutions for its disinformation and legal defense strategies, the Court cited the misuse of American Cancer Society data by a non-faculty researcher at UCLA. . . Big Tobacco's investment in UCLA bought it the chance to argue falsely, using UCLA's name, that the science on secondhand smoke was inconclusive, to battle public health measures. Whatever the tobacco industry gains from the University, the University loses. The public loses, too." This compounding of the defamation in the court papers through paid advertising was signed by 21 prominent individuals who identify themselves as "among those who support action by the University of California Regents to refuse all future tobacco industry funding." The signatories include both Samet and Eubanks, who obviously have been directly involved in lobbying the UC Regents, a position that compromises their objectivity with regard to my inclusion in the USDOJ lawsuit. Given the obsessive focus on my tobacco industry funding, it is noteworthy that there is no indication of the funding and competing interests of those associated with this advertisement. The Chronicle of Higher Education website states that a "tabloid-page spread" advertisement like this one costs \$22,630 [109], a sum unlikely to have been paid by the signatories themselves.

Based on the record presented above, Eubanks has obviously dealt extensively with both Glantz and Samet regarding the issue of my *BMJ* paper and the USDOJ lawsuit. She injected herself directly into the UC tobacco industry funding ban issue with a lecture before the Regents on July 18, 2007, when she described the USDOJ lawsuit and its connection to UC [110]. She claimed that Judge Kessler was "a neutral fact finder, a federal judge, who made her findings of conspiratorial conduct objectively" based on "a full and fair record." However, she

knows that the record is not objective and that I was never given any opportunity to defend myself and my *BMJ* paper during the trial. In an eloquent defense of academic freedom at UC, the 2006–2007 UC Academic Senate Chair John B. Oakley challenged Eubank's linkage of the USDOJ lawsuit to UC and raised the issue of whether Judge Kessler's opinion would ultimately be upheld upon appeal [11188d]. A clearer understanding of this entire issue can be gained by carefully listening to the Eubanks and Oakley audio files [110,111].

Jonathan M. Samet, M.D., and Conflict of Interest

Samet has not revealed his competing interests on the subject of ETS as they relate to the BMJ rr [20], the BMJ letter [73], the IARC Report [50], the JNCI article [52], the Surgeon General's Report [79], his USDOJ lawsuit testimony [106], or the Chronicle of Higher Education advertisement [107]. Given that Samet has criticized persons who disagree with his views on ETS because of their competing interests, it is fair and reasonable to ask why he has failed to report his own substantial competing interests. A careful examination of the Surgeon General's Report reveals that it contains no conflict of interest disclosures for Senior Scientific Editor Samet or for any of the other editors or reviewers. In addition, an examination of the other items above reveals the Samet has not disclosed a financial conflict of interest which could have compromised his objectivity on ETS. This imbalance further suggests that the attacks on my research have nothing to do with a principled concern about conflicts of interest, but are purely a matter of not liking the results.

The article, "smoke out!", in the Spring 2003 issue of *Johns Hopkins Public Health*, "The Magazine of the Johns Hopkins Bloomberg School of Public Health" [112] reveals that, "After three years of preparation, Samet testified in the landmark 1998 Minnesota tobacco trial that smoking causes certain diseases like lung cancer" and that Samet was "working on the federal government's \$289 billion lawsuit that accuses tobacco companies of 50 years of deceptive marketing," which is the USDOJ lawsuit discussed above. Later, the article stated "In March, the Flight Attendant Medical Research Institute honored Samet with the '...Dr. William Cahan Distinguished Professor' Award and \$600,000 over 3 years to combat tobacco-related disease."

According to the Flight Attendant Medical Research Institute (FAMRI) website, the 'Dr. William Cahan Distinguished Professor' award to Samet during 2003–2006 was "made in recognition of the recipients' ongoing work in combating the diseases caused by exposure to second hand tobacco smoke" [113]. In addition, Samet has a prominent role in the current multi-million dollar Johns Hopkins FAMRI Center of Excellence [114]. This Center

was established in 2005 and currently has 30 FAMRIfunded research projects on "diseases and medical conditions caused from exposure to tobacco smoke," including one by Samet on "Reducing the Risks of Secondhand Tobacco Smoke Globally" [113].

FAMRI is a foundation established as a result of an October 1991 Class Action suit filed in Miami's Dade County Circuit Court in Florida, known as Broin v. Philip Morris [116]. This suit was filed against the tobacco industry on behalf of flight attendants who sought damages for diseases and deaths allegedly caused by their exposure to second hand tobacco smoke in airline cabins [117]. A settlement was reached in October 1997 between the plaintiffs and four tobacco companies. The Settlement Agreement included the establishment of a not-for-profit medical research foundation with funding by the tobacco industry of \$300 million. The Foundation was to have no tobacco company involvement, other than funding. The purpose of the foundation was "to sponsor scientific research with respect to the early detection and cure of diseases associated with cigarette smoking" [118]. FAMRI, as it was actually established, has a distinctly different mission, which is "to sponsor scientific and medical research for the early detection, prevention, treatment and cure of diseases and medical conditions caused from exposure to tobacco smoke." [117]. Since FAMRI's mission statement assumes that diseases like lung cancer and CHD are caused by "exposure to tobacco smoke," this funding source may have influenced Samet's decisions about which epidemiologic studies he chooses to believe and which ones he chooses to ignore, and thus should have been disclosed. As noted in an August 23, 2006 JAMA editorial, in published articles it is important "that readers are aware of the authors' financial relationships and potential conflicts of interest so that these readers can interpret the article in light of that information" [119].

Jonathan M. Samet, M.D., and the 1992 EPA Report

One might wonder how omissions, distortions, and exaggerations like those pointed out above could occur in a document as important as a Surgeon General's Report on ETS. To better understand this phenomena one must realize that Samet has dealt with the ETS issue in this manner for many years. In particular, he played a major role in the epidemiologic analysis for the December 1992 report on Health Effects of Passive Smoking: Lung Cancer and Other Disorders: The Report of the United States Environmental Protection Agency [120]. This EPA report classified ETS as a Group A human carcinogen, which causes about 3,000 lung cancer deaths per year in the U.S. The findings from this report were used in the Broin v. Philip Morris litigation described above.

The epidemiologic methodology and conclusions of the EPA report have been severely criticized. One of the harshest critiques is the 92-page Decision issued by Federal Judge William L. Osteen on July 17, 1998, which overturned the report in the U.S. District Court [121]. For instance, in his conclusion Judge Osteen wrote: "In conducting the Assessment, EPA deemed it biologically plausible that ETS was a carcinogen. EPA's theory was premised on the similarities between MS [mainstream smoke], SS [sidestream smoke], and ETS. In other chapters, the Agency used MS and ETS dissimilarities to justify methodology. Recognizing problems, EPA attempted to confirm the theory with epidemiologic studies. After choosing a portion of the studies, EPA did not find a statistically significant association. EPA then claimed the bioplausibility theory, renominated the a priori hypothesis, justified a more lenient methodology. With a new methodology, EPA demonstrated from the 88 selected studies a very low relative risk for lung cancer based on ETS exposure. Based on its original theory and the weak evidence of association, EPA concluded the evidence showed a causal relationship between cancer and ETS. The administrative record contains glaring deficiencies. . . . "

In order to more fully understand the EPA report and its inherent flaws, one must read the complete Osteen decision [121], as well as the books *Passive Smoke: The EPA's Betrayal of Science and Policy* by Drs. Gio B. Gori and John C. Luik [122], *Ashes to Ashes: America's Hundred-Year Cigarette War, the Public Health, and the Unabashed Triumph of Philip Morris* by Richard Kluger [123], *For Your Own Good: The Anti-Smoking Crusade and the Tyranny of Public Health* by Jacob Sullum [124], and the *Brill's Content* magazine article "Warning: Secondhand Smoke May NOT Kill You" by Nicholas Varchaver [125]. Finally, one must read the January 28, 1993 *Investors' Business Daily* article "Is EPA Blowing Its Own Smoke? How Much Science Is Behind Its Tobacco Finding?" by Michael Fumento, who stimulated my own interest in the ETS issue [126].

2006 Congress of Epidemiology and Trofim Denisovich Lysenko Analogy

In order to explain the phenomenon that has made this defense of my epidemiologic research necessary, Geoffrey Kabat, Sheldon Ungar, and I presented a symposium entitled "Reassessment of the Long-term Mortality Risks of Active and Passive Smoking" at the 2nd North American Congress of Epidemiology in Seattle, Washington on June 24, 2006 [127]. We described major misrepresentations that are currently occurring with regard to the epidemiology of both active and passive smoking, as well as the silencing of science associated with this area of epidemiology. I presented the rationale for the symposium based on the fact that important epidemiologic findings have been ignored or mischaracterized in prior assessments. Then I

presented evidence that the adverse effects of active smoking on mortality are less reversible by cessation than generally believed, based on randomized controlled trials involving smoking cessation and "natural experiments" involving the CA CPS I cohort and several other cohorts [31,128,129]. Kabat presented evidence that the relationship between passive smoking and mortality is weaker than generally believed, particularly within the United States, based on our two recent ETS papers [1,39]. Ungar described the "silencing of science" phenomenon with regard to our May 17, 2003 *BMJ* paper that he documented and described in his 2005 paper [27].

In this symposium we addressed several important issues: 1) the implications of our reassessment for the relative dangers of active and passive smoking; 2) the way in which ideological and political agendas have influenced the interpretation of epidemiologic evidence; and 3) the importance of separating non-scientific agendas from objective assessment of evidence. We made the case that: 1) all epidemiologic findings must be evaluated in a fair and consistent manner in order to obtain an accurate assessment of the mortality risks of active and passive smoking; 2) epidemiologic findings must be judged on their merits and not on extraneous factors; and 3) additional epidemiologic research in this area needs to be conducted free of partisanship. Our complete presentations are available on the Scientific Integrity Institute website [130], and they include our PowerPoint slides and the audio files for our lectures.

It is quite informative to compare our Symposium with the June 23, 2006 lecture "Using Epidemiologic Evidence to Advance Health: Dealing with Critics and Criticisms" given by Samet at the same Congress of Epidemiology [131]. Samet discussed the use of epidemiologic evidence in public health policy making with regard to the environmental epidemiology issues in which he has been involved. In particular, he discussed the epidemiologic evidence on the relationship between passive smoking and lung cancer just four days before the June 27, 2006 release of the Surgeon General's Report on involuntary smoking for which he was Senior Scientific Editor [79]. He talked about the criticism of weak epidemiologic relationships, such as those described in major documents like the 2006 Surgeon General's Report. But he failed to mention that much of this criticism is due to the fact that he has attempted to turn weak and inconsistent observational epidemiologic evidence into an undisputed causal relationship. He talked about how critics raise epidemiologic issues like confounding and bias, but he failed to acknowledge his own biased presentation of the evidence, including omitting my BMJ paper from the report and failing to acknowledge that the U.S. evidence is weaker than the evidence outside of the U.S.

Also, it is quite telling how Samet dismissed critics of the causal relationship between passive smoking and lung cancer by classifying them as "stakeholders" linked with the "tobacco industry." He implied that it is not necessary to address the merits of their criticisms simply because they are stakeholders in decisions related to passive smoking. However, he failed to disclose his own financial interests that surely put him in the stakeholder category. He certainly never mentions that his FAMRI money originates from the tobacco industry, making it remarkably similar to my CIAR funding. Samet's lecture provides insight into his thought processes and the ways in which he manipulates evidence to fit his vision of an epidemiologic relationship with public policy implications. The transcript of a key portion of his lecture is available [132], as is the audio file [133].

We concluded our Symposium by drawing an analogy between the current situation involving ETS epidemiology in the United States and the historical situation involving agronomist Trofim Denisovich Lysenko and plant genetics in the Soviet Union during the period of 1927-1962 [2]. While it is common to invoke George Orwell or Joseph McCarthy in discussions like this, I believe the lessons from the admittedly more extreme Lysenko case are more analogous and informative. Although ETS epidemiologic evidence has never been conclusive, several major reports have been issued with definitive conclusions about a "causal relationship" between ETS and mortality. All major U.S. government and private health agencies have declared that a causal relationship exists and these organizations have created "a regime of truth that cannot be intelligibly questioned." These organizations then use any means necessary to enforce this "regime of truth." Since the publication of the influential null findings in my BMJ paper, which contradict the "regime of truth," I have been subjected to a massive ad hominem attack, my career has been threatened, and my paper has been dismissed because of its politically incorrect findings. In addition, I was inserted into a massive lawsuit by my own government in a manner that makes it appear that I have committed "scientific fraud" and have been engaged in racketeering with the tobacco industry. There also has been the attempt to force the University of California to ban the tobacco industry funding that I have used and to restrict future research in the areas of tobacco-related diseases that I have been investigating.

Lysenko used his influence and backing by the Soviet government to create a "regime of truth" and to stop others' research in order to promote scientifically invalid "vernalization" and Lamarckian plant genetics. He was also successful in attacking and destroying his critics, like Nicolai Vavilov, who espoused proper Mendelian plant genetics. Because Lysenko prevailed for such a long period of time,

crop yields were low, Soviet agriculture regressed, and Soviet citizens suffered greatly and many faced starvation. During this same period, proper plant genetics were developed and implemented in the U.S. and this resulted in the greatly increased crop yields that have made U.S. food production so incredibly successful. The entire saga of "Lysenko pseudoscience" has been extensively described in websites about Lysenko [134], journal articles [2,135], and books [136-138].

Prominent U.S. epidemiologists and activists are wielding governmental influence to distort the epidemiology of both active and passive smoking in the U.S. and are contributing to a Lysenko-like research environment where it is virtually impossible to conduct research that produces politically incorrect findings, such as, those in my *BMJ* paper. Much additional research is needed because the primary tobacco-related disease, lung cancer, still causes 160,000 deaths per year in the U.S. and will not go away any time soon. This Lysenko-like research environment needs to end and epidemiologists must be free to conduct additional research on tobacco-related diseases with a variety of funding sources without fear of the kind of attacks that I have experienced.

A Challenge to ACS and Michael J. Thun, M.D.

Some of the controversy about the relation of ETS and tobacco-related mortality in the largest U.S. observational epidemiologic studies could be settled if Thun fully, fairly, and transparently analyses the CPS I and CPS II cohort data that the ACS currently possesses. Because of their size and length of mortality follow-up, these two cohorts contain the vast majority of the potentially available U.S. evidence on ETS, and are already the basis for important U.S. evidence on active smoking. Given the epidemiologic expertise of Thun and the availability of the appropriate CPS I and CPS II data, such an analysis could be conducted in a matter of weeks. In the interest of better understanding cancer etiology, the ACS should fully analyze these important data. I have provided sample Tables 3, 4, 5, 6 and 7 so that Thun can present results that are directly comparable to those presented in my BMJ paper [1].

In addition, Thun should analyze the CPS II cohort as a "natural experiment" of smoking cessation and mortality trends in a manner similar to what I have done. Such an analysis would test my hypothesis, based on analysis of the CA CPS I and three other U.S. cohorts, that the long-term adverse mortality effects of active smoking are more dangerous than generally believed because they are less reversible by cessation than generally believed [31,128,129]. The ACS owes it to the over two million Americans who are subjects in the CPS I and CPS II cohorts, as well as to those Americans who support the ACS, to produce epidemiologic findings that accurately

and completely describe the mortality risks of active and passive smoking in their data.

In order to determine if a full analysis of ETS and mortality in the CPS II cohort supports the analysis of ETS and mortality in the CA CPS I cohort presented in my *BMJ* paper, I sent Thun a June 21, 2007 email request that he complete Tables 5, 6 and 7. Thun replied with a June 26, 2007 letter in which he gave several reasons why he would not complete Tables 5, 6 and 7. He stated "In summary, I do not believe that the analyses you request in CPS-II would produce scientifically meaningful results" [139]. He indicated no willingness to do further CPS II analyses of any kind, even analyses of the relationship of ETS to mortality during the past fifteen years. This is the latest evidence supporting the extensive "silencing of science" phenomena that currently exists with regard to ETS epidemiology in the U.S.

To illustrate the existing bias in the release of ACS results, it is quite informative to note the response by Thun to the September 26, 1994 letter that he received from Glantz [140], regarding the CPS II analyses that LeVois and Layard conducted in 1994 and published in 1995 [40]. Thun sent Glantz a detailed November 4, 1994 letter which included preliminary CPS II analyses and criticisms and described plans to do further CPS II analyses [141]. Responses to Thun's CPS II analyses and criticisms were then made by LeVois [142] and Layard [143]. All of this correspondence and commentary reinforces the continuing need for a full and objective analysis of the CPS I and CPS II data possessed by ACS.

Conclusion

It is very disturbing that a major health organization like the ACS has made false and misleading statements about me and my May 17, 2003 *BMJ* paper for over four years. It is further disturbing that prominent individuals like Thun, Samet, and Glantz have continued to attack the findings in the *BMJ* paper, even though I have presented extensive evidence that supports the validity of these findings. In addition, it is reprehensible that the *BMJ* paper was inserted in the USDOJ RICO lawsuit and omitted from the 2006 Surgeon General's Report. These actions must be kept in mind when evaluating the honesty, integrity, and objectivity of those responsible.

These criticisms may sound personally defensive, and indeed when one is so personally attacked, some personal defense is necessary. But this is also a defense against epidemiology becoming "Lysenko pseudoscience," where the validity of methods and studies is based merely on those results that are preferred by influential advocates and researchers and contrary results are discredited using the tactics of Lysenko. Epidemiologic science is not inher-

Table 3: Level of spousal smoking related to deaths from lung cancer and coronary heart disease among never smokers in California CPS I cohort as of 1959. Relative risk (RR with 95% CI) comparing persons with each level of exposure to those without exposure. Proportional hazards linear model (PHREG) is used and RR is adjusted for age at entry. Results shown are from Tables 7 and 8 of 2003 BMJ paper [1]. See note below.*

			Lung cancer (ICD7 =	162–3, ICE	08&9 = 162)		Coronary heart disease(IC	D7 = 420, I	CD8&9 = 410-4)
			I Jan 60–30 Sep 72		I Jan 60–31 Dec 98	-	I Jan 60–30 Dec 72		I Jan 60–31 Dec 98
Spousal smoking (ETS index level) as of 1959	Subjects	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)
	nn,nnn	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)
<u>Males</u>									
Never (I)	7,458		1.00	65	1.00			1,860	1.00
Former (2)	624			5	0.92 (0.37–2.30)			126	0.94 (0.78–1.12)
Current									
I-9 cpd (3)	392							81	0.97 (0.78-1.21)
10–19' cpd(4)	513							99	0.86 (0.70-1.05)
20 cpd (5)	458							81	0.92 (0.74–1.15)
21-39 cpd (6)	129							27	1.16 (0.79–1.69)
40+ cpd (7)	45							13	1.29 (0.75–2.22)
Current – total	1,537			9	0.69 (0.34–1.39)			301	0.94 (0.83-1.07)
Ever	2,161			14	0.75 (0.42–1.35)			427	0.94 (0.85–1.05)
<u>Females</u>									
Never (I)	7,399		1.00	51	1.00		1.00	1,053	1.00
Former (2)	6,858			51	1.08 (0.73-1.60)			1,059	1.02 (0.93–1.11)
Current									
Pipe/cigar (3)	2,691							389	0.99 (0.88-1.11)
I-9 cpd (4)	1,102							183	1.13 (0.97–1.33)
10–19 cpd (5)	2,117							310	1.03 (0.91–1.17)
20 cpd (6)	3,288							412	1.04 (0.92–1.16)
21-39 cpd (7)	1,646							167	0.95 (0.80-1.12)
40+ cpd (8)	841							72	0.83 (0.65-1.06)
Current – total	11,685			75	0.93 (0.65-1.33)			1,533	1.01 (0.93–1.09)
Ever	18,543			126	0.99 (0.72-1.37)			2,592	1.01 (0.94-1.08)

^{*}Tables C-G should be completed by Michael J. Thun, M.D., of the ACS to fully present results on ETS and lung cancer and coronary heart disease mortality in the CPS I cohort during 1960–1972 and in the CPS II cohort during 1982–1998 in a format that is the same as that used for the 1960–1998 CA CPS I results in the 2003 BMJ paper (1), some of which are shown in Table C.

Table 4: Level of spousal smoking related to deaths from lung cancer and coronary heart disease among never smokers in 25-state CPS I cohort as of 1959. Relative risk (RR with 95% CI) comparing persons with each level of exposure to those without exposure. Proportional hazards linear model (PHREG) is used and RR is adjusted for age at entry.

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			Lung cancer (ICD7 =	= 162–3, 10	CD8 = 162)		Coronary heart disease (IC	CD7 = 420), ICD8 = 410–4).
			I Jan 60–31 Dec 65		I Jan 60–30 Sep 72		I Jan 60–31 Dec 65		I Jan 60–30 Sep 72
Spousal smoking (ETS index level) as of 1959	Subjects	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI
	nn,nnn	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)
<u>Males</u>									
Never (I)			1.00		1.00		1.00		1.00
Former (2)									
Current									
I-9 cpd (3)									
10-19 cpd(4)									
20 cpd (5)									
21–39 cpd (6)									
40+ cpd (7)									
Current – total									
Ever									
<u>Females</u>									
Never (I)			1.00		1.00		1.00		1.00
Former (2)									
Current									
Pipe/cigar (3)									
I-9 cpd (4)									
10–19 cpd (5)									
20 cpd (6)									
21–39 cpd (7)									
40+ cpd (8)									
Current – total									
Ever									

Table 5: Level of spousal smoking related to deaths from lung cancer and coronary heart disease among never smokers in CPS II cohort as of 1982. Relative risk (RR with 95% CI) comparing persons with each level of exposure to those without exposure. Proportional hazards linear model (PHREG) is used and RR is adjusted for age at entry.

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			Lung cancer	(ICD9 = I	62)		Coronary heart disc	ease (ICD	9 = 410–4)
			I Sep 82–31 Dec 89		I Sep 82–31 Dec 98		I Sep 82–31 Dec 89		I Sep 82–31 Dec 98
Spousal smoking (ETS index level) as of 1982	Subjects	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)
	nn,nnn	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)
<u>Males</u>									
Never (I)			1.00		1.00		1.00		1.00
Former (2)									
Current									
I-9 cpd (3)									
10-19 cpd(4)									
20 cpd (5)									
21-39 cpd (6)									
40+ cpd (7)									
Current – total									
Ever									
<u>Females</u>									
Never (I)			1.00		1.00		1.00		1.00
Former (2)									
Current									
Pipe/cigar (3)									
I–9 cpd (4)									
10-19 cpd (5)									
20 cpd (6)									
21–39 cpd (7)									
40+ cpd (8)									
Current – total									
Ever									

			Lung cance	er (ICD9 = 1	52)		Coronary heart d	lisease (ICD9	= 410–4)
			I Sep 82–31 Dec 89		I Sep 82–31 Dec 98	_	I Sep 82–31 Dec 89		I Sep 82–31 Dec 98
Total daily hours of ETS exposure as of 1982	Subjects	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)	Deaths	Age-adjusted RR (95% CI)
	nn,nnn	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)	nnn	x.xx (x.xx-x.xx)
<u>Males</u>									
0 hours			1.00		1.00		1.00		1.00
I									
2									
3									
4									
5									
6									
7									
8+									
Current total (I+)									
<u>Females</u>									
0 hours			1.00		1.00		1.00		1.00
I									
2									
3									
4									
5									
6									
7									
8+									
Current total (I+)									

Table 7: 1982 level of spousal smoking related to total self-reported ETS exposure among never smokers in 1982 CPS II cohort and 1992 CPS II Nutrition cohort.

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		Percent	distribution	of 1982 tota	ll daily hours of E	TS exposure		Perce	nt distributio	n of 1992 total v	veekly hours of E	ΓS exposure
Spousal smoking as of 1982	1982 subjects	0	ı	2	3–7	8+	1992 subjects	0	I–7	8–14	15–49	50+
<u>Males</u>												
Never (I)												
Former (2)												
Current												
I-9 cpd (3)												
10-19 cpd(4)												
20 cpd (5)												
21-39 cpd (6)												
40+ cpd (7)												
Current – total												
ver												
<u>emales</u>												
Never (I)												
Former (2)												
Current												
Pipe/cigar (3)												
I-9 cpd (4)												
10-19 cpd (5)												
20 cpd (6)												
21-39 cpd (7)												
40+ cpd (8)												
Current – total												

ently pseudoscience, of course, but the process that has led to many current claims about ETS is.

Hopefully, epidemiology can continue as a field in which all legitimate research findings can be published and objectively evaluated, including those findings considered to be controversial. However, this will happen only if advocacy organizations like the ACS and activists like Glantz refrain from unethically smearing honest scientists and putting out false and misleading statements. In addition, epidemiologists like Thun must honestly analyze all the epidemiologic evidence that they possess and fully report their results, and epidemiologists like Samet must not omit important and accurate research findings from a major document such as the Surgeon General's Report. Such omissions and actions have seriously distorted the evidence on the health effects of ETS exposure, particularly within the US.

Hopefully, this entire episode will help prevent similar episodes in the future. Furthermore, this episode will be particularly valuable if it eventually leads to a full and objective analysis of the important epidemiologic evidence that the ACS possesses on both active and passive smoking. In the meantime, epidemiologists and others interested in a full assessment of the available epidemiologic evidence on the health effects of ETS should carefully read and study this document and all the references and tables that are included in it.

Competing interests

Funding of this paper is the same as that of reference 39. The content of this paper is based on the knowledge I have acquired during my entire epidemiologic career, during which I have had many funding sources. My competing interests are fully discussed in the text of this paper and in reference 1 and are known worldwide thanks largely to the efforts of Glantz, Thun, and Samet. My personal stake in the matters discussed here should be self-evident. In order to address concerns about my competing interests, this paper is entirely transparent and its contents can be verified with the references cited.

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RESEARCH PAPER

The limits of competing interest disclosures

L A Bero, S Glantz, M-K Hong

Tobacco Control 2005;14:118-126

Objective: To assess the effectiveness of conflict of interest disclosure policies by comparing a competing interests disclosure statement that met the requirements established by the journal in a 2003 article on health effects of secondhand smoke based on the American Cancer Society CPS-I dataset with internal tobacco industry documents describing financial ties between the tobacco industry and authors of the study.

Design: Descriptive analysis of internal tobacco industry documents retrieved from the Legacy Tobacco Documents Library, University of California, San Francisco.

Results: Meeting the requirements for financial disclosure established by the journal did not provide the reader with a full picture of the tobacco industry's involvement with the study authors. The tobacco industry documents reveal that the authors had long standing financial and other working relationships with the tobacco industry.

Conclusion: These findings are another example of how simply requiring authors to disclose financial ties with the tobacco industry may not be adequate to give readers (and reviewers) a full picture of the author's relationship with the tobacco industry. The documents also reveal that the industry funds research to enhance its credibility and endeavours to work with respected scientists to advance its goals. These findings question the adequacy of current journal policies regarding competing interest disclosures and the acceptability of tobacco industry funding for academic research.

See end of article for authors' affiliations

Correspondence to: Lisa A Bero, PhD, University of California, 3333 California Street, Suite 420, Box 0613, San Francisco, CA 94143-0613 (94118 for express mail only), USA; bero@ medicine.ucsf.edu

n May 2003, the *British Medical Journal (BMJ)* published "Environmental tobacco smoke and tobacco related mortality in a prospective study of Californians, 1960–1998". In this paper, Enstrom and Kabat used a longitudinal cohort collected by the American Cancer Society beginning in 1959 (the CPS-I dataset, Cancer Prevention Study) to conclude that secondhand smoke exposure does not increase the risk of lung cancer and heart disease. This conclusion conflicts with comprehensive reviews which find that secondhand smoke increases these risks by 20–30%. The paper was quickly and widely cited in the lay press^{3 4} and by tobacco industry supported journalists who criticise government sponsored "junk science". 5

Subsequent criticisms of the paper focused on the methodology of the study and the authors' disclosed financial ties. The main methodological issue was that there was no real "unexposed" group in the CPS-I dataset that Enstrom and Kabat used.⁶ ⁷

The article included a lengthy statement of funding sources and competing interests, listing as one of its sponsors the tobacco industry's Center for Indoor Air Research (CIAR). Peer reviewers tend to be more critical of articles with industry sponsorship than those without such sponsorship. One rapid response to the Enstrom and Kabat article suggested that it might be useful for editors to "require a statement about the role of the funding source in the design, conduct, analysis and reporting of the data". Earlier tobacco industry funded studies of the health effects of secondhand smoke have failed to fully disclose the sponsor's role in the research. In In contrast to these earlier situations, the Enstrom and Kabat paper was consistent with the requirements for such statements established by the *BMJ*.

Financial disclosure policies for authors of journal articles are becoming the norm.^{12 13} However, experts on scientific

Editor's note

This article replaces a previous version that was posted on the *Tobacco Control* website on 9 December 2004 and taken down on 11 February 2005.

journal disclosure policies have questioned the adequacy of these policies.14 Studies comparing disclosures of financial ties in research articles with information on the financial ties of authors obtained from independent sources have shown that only a small percentage of research articles (less than 20%) fully disclose the financial ties of authors. 14 15 A more recent study investigated the lead and last authors of research articles who declared that they had no conflicts to disclose in 163 original articles that appeared in four journals (New England Journal of Medicine, Journal of the American Medical Association, Environmental Health Perspectives, Toxicology and Applied Pharmacology). By comparing publicly available databases and the journals' conflict of interest policies, the study found 13 articles (8%) where relevant conflicts of interest were not disclosed to readers.15 In most cases, the authors failed to disclose the relevant information to the journal.¹⁵

The inadequacy of journal article disclosures when compared to information on financial ties of authors obtained from other sources has led experts on disclosure to suggest more comprehensive policies and stronger enforcement of existing policies. ^{14 16} For example, experts have suggested that journals require disclosure of exact amounts and duration of financial ties, and that journals clarify how they define the "relevance" of a financial tie to the research. ¹⁵ A number of scholars have argued that corporate sponsorship of research or financial ties of authors should be prohibited. ^{14 17 18} These proposed bans eliminate the need for disclosure to "manage" the conflict of interest and protect against even the appearance of conflict.

The Enstrom and Kabat paper provides an interesting case study to test this suggestion precisely because they provided an extensive disclosure that met the current *BMJ* standards

Abbreviations: ACS, American Cancer Society; *BMJ, British Medical Journal*; CIAR, Center for Indoor Air Research; CPS, Cancer Prevention Study; CTR, Council for Tobacco Research; JAMA, *Journal of the American Medical Association*; SRRC, Philip Morris scientific research review committee; TRDRP, California Tobacco-Related Disease Research Program

for such disclosure. Using methods similar to that of Krimsky, we compared the disclosure in a scientific journal article with information on financial ties of the authors obtained from an independent source. ¹⁹ ²⁰ We compare the disclosure Enstrom and Kabat provided in their article with internal tobacco industry documents describing their relationship with the tobacco industry in order to obtain details of the history and nature of their association, with the goal of understanding whether a disclosure statement that met *BMJ's* requirements gave the reader a clear description of the role of the tobacco industry in the study and the historical relationship of the authors with the industry.

METHODS

We retrieved documents from the Legacy Tobacco Documents Library (www.legacy.library.ucsf.edu) using the terms "Enstrom", "Kabat", "CIAR", "CPSI", "CPSII", "Womble Carlyle", and "Shook, Hardy and Bacon" and names of key individuals, followed by "snowball searches". We collected 156 documents; 81 discussed the development of the CPS-I data analysis or other research by Enstrom or Kabat.

RESULTS

Table 1 summarises financial ties between the authors and the tobacco industry. The disclosure statement was long, 305 words, compared to the average of 127 words in the 20 other research articles published in *BMJ* that month. Of these 305 words, 98 related to the tobacco industry (table 2).

Acknowledgement of American Cancer Society and California Tobacco-Related Disease Research Program

The authors disclosed that the American Cancer Society (ACS) developed the CPS-I dataset.²¹ The disclosure notes

that some work was funded by the California Tobacco-Related Disease Research Program (TRDRP), then states that continued funding was "denied". The renewal for the project was not funded because it had inadequate scientific merit as determined by a peer review panel in a year that TRDRP had an 85% budget reduction.²²

Early interactions with the tobacco industry

The disclosure states that the research was supported by the tobacco industry after continuing support from TRDRP was denied and that "[i]n recent years" Enstrom received research funding from the tobacco industry because he was unable to obtain equivalent funds from other sources. Although these statements are accurate and comply with Enstrom's reporting obligations under BMJ policy, the tobacco industry documents also show that Enstrom had a long history of association with the industry. In 1975, he approached the Council for Tobacco Research (CTR), a tobacco industry research organisation, to fund a study of cancer among US Mormons.23 24 He argued that the study "should be helpful in assessing the possible role which other factors besides smoking play in the etiology of lung cancer".24 The proposal was deferred for discussion by the Executive Committee, Scientific Advisory Board of CTR in June 1975.²⁵ Although Enstrom requested a letter of support from the Tobacco Institute, the industry's trade organisation in the USA,²⁶ and modified his experimental design in April 1976,²⁷ there is no record that the proposal was funded. In 1978, Enstrom submitted another request to CTR to study "Smoking cessation and mortality trends among California physicians".28 There is no record that this proposal was funded.

In 1979, Enstrom published a review article that provoked concern and criticism from the tobacco industry.^{29–31} The

Year	Enstrom	Kabat
1975	First record of Enstrom approaching the tobacco industry for funding. Enstrom asks the Council for Tobacco Research (CTR) to fund a study of cancer among Mormons living in the USA	
1978	Enstrom submits another request for funding to CTR for a study entitled "Smoking cessation and mortality trends among California physicians"	
1979	Enstrom publishes a review article that provokes criticism from the tobacco industry	
1981	'	Kabat begins collaborations with Ernst Wynder, whose American Health Foundation (AHF) was tobacco industry funded. Tobacco industry support was not acknowledged in subsequent publications
1990	The tobacco industry initiates contact with Enstrom to critique research on secondhand tobacco smoke. Enstrom declines to comment	ŭ , ,
1990	Enstrom requests funding from Philip Morris for a study to support his work on lung cancer mortality trends among non-smokers. Enstrom is advised to seek funding from the Center for Indoor Air Research (CIAR)	
1991	Enstrom submits a pre-proposal to the Council for Tobacco Research (CTR) for a study entitled "Mortality Trends Among Smokers and Nonsmokers Study"	
1991–7	CTR funds the "Mortality Trends Among Smokers and Nonsmokers Study"	
1996	Enstrom submits to the <i>Journal of the American Medical Association (JAMA)</i> a reanalysis of an article by Elizabeth Fontham, <i>et al.</i> This reanalysis was financially supported by the tobacco industry.	
1997	Enstrom submits a proposal entitled "Relationship of low levels of active smoking to mortality" for funding to Philip Morris. The proposal is funded	Kabat co-authors a paper with tobacco industry consultants that criticises the evidence linking secondhand smoke with lung cancer
1996	Enstrom submits pre-proposals to the tobacco industry's CIAR to explore the possibility of funding research related to secondhand smoke using the California CPS-I cohort	Ü
1997 May	CIAR board of directors is informed that discussions had taken place with Enstrom and Kabat about "the possibility of their collaboration"	
1997	Max Eisenberg, director of CIAR, recommends that Enstrom's revised proposal,	
November	"Proposed research on passive smoking", be considered under the Directed Studies programme	
1997	The CIAR board of directors votes to fund Enstrom's proposal with modifications	

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Table 2 Comparison of competing interest disclosure with information from internal tobacco industry documents and other sources

Disclosure published in BMJ that met BMJ's standards

"The American Cancer Society (ACS) initiated CPS-I in 1959, conducted follow up until 1972, and has maintained the original database"

"Extended follow up until 1997 was conducted at the University of California at Los Angeles with initial support from the Tobacco-Related Disease Research Program, a University of California research organisation funded by the Proposition 99 cigarette surtax. After continuing support from the Tobacco-Related Disease Research program was denied,

follow up through 1999 and data analysis were conducted at University of California at Los Angeles with support from the Center for Indoor Air Research, a 1988–99 research organisation that receive funding primarily from US tobacco companies"

"In recent years JEE has received funds originating from the tobacco industry for his tobacco related epidemiological research because it has been impossible for him to obtain equivalent funds from other sources"

"GCK never received funds originating from the tobacco industry until last year, when he conducted an epidemiological review for a law firm which has several tobacco companies as clients. He has served as a consultant to the University of California at Los Angeles for this paper"

"JEE and GCK have no other competing interests. They are both lifelong non-smokers whose primary interest is an accurate determination of the health effects of tobacco."

Findings from tobacco industry documents and other sources that go beyond what was required by BMJ

ACS epidemiologists repeatedly cautioned Enstrom before he began the study that the CPS-I dataset was not appropriate to investigate the effects of environmental tobacco smoke (ETS)²¹

Dr Enstrom's application for continued funding from the California Tobacco-Related Disease Research Program was not funded because it had inadequate scientific merit in a year when the program's budget was drastically reduced.²²

The Center for Indoor Air Research funded grants that were peer reviewed by scientists and "special projects" that were reviewed by tobacco industry lawyers and executives. 38 The CPS-I analysis was funded through the same mechanism as the special projects JEE sought research funding from the tobacco industry beginning in 1975 and received his first funding in 1992. He has also received funding for serving as an expert witness, reviewing dissertation and arrent proposals

GCK has had an ongoing indirect relationship with the tobacco industry since at least 1981 though his collaborations with Ernst Wynder whose American Health Foundation was funded by Philip Morris.

The analysis of the CPS-I dataset was also funded by Philip Morris tobacco company and this was not disclosed

article, which was consistent with journal policy at the time, disclosed no sources of funding, used data from the Mormon cohort, US veteran cohort, CPS-I, and National Mortality Survey to conclude that lung cancer rates had been rising among non-smokers between 1914 and 1968 and that this rise was related to factors other than "personal cigarette smoking" including "environmental pollution including environmental tobacco smoke".³²

Developing a relationship with the tobacco industry

The industry documents reveal little interaction between the tobacco industry and Enstrom from 1979 until 1990, when the tobacco industry began to recruit Enstrom as a potential critic of research on secondhand smoke. In 1990, Thomas Borelli, Manager Scientific Issues, Philip Morris, asked Enstrom, among others, to comment on a PhD dissertation by Luis Varela.33 34 Borelli was interested in the dissertation because it contained findings suggesting that secondhand smoke exposure was not associated with disease. The industry frequently cited the dissertation as evidence that should be included in risk assessments of secondhand smoke.35 In contrast to the preliminary results in the dissertation, however, the final publication in the New England Journal of Medicine³⁶ concluded that childhood exposure to secondhand smoke was associated with adult lung cancer.

Enstrom declined to review the dissertation, stating: "My epidemiologic research does not deal directly with the issue of environmental tobacco smoke". 33 However, he used the invitation to inquire about funding from Philip Morris to update his work on lung cancer mortality trends among non-smokers. 33 Borelli responded that "Philip Morris does not usually fund research projects" and referred Enstrom to the industry's CIAR. 37 The CIAR was supported solely by the tobacco industry and funded research projects using two different mechanisms. Some projects were peer reviewed by scientists and were more likely to examine health effects of indoor air contaminants other than secondhand smoke. 38 Other projects, which were more likely to focus on secondhand smoke, were reviewed by tobacco industry executives and lawyers. 38

Enstrom sent Borelli a number of references that suggested a need for further research on lung cancer trends among non-smokers, noting that "Attention could be given to the issues of personal smoking and environmental tobacco smoke".³⁹ Myron Johnston, Research and Development, Philip Morris, was asked by Bob Pages, Director, Science and Technology, Philip Morris, to comment on the correspondence between Borelli and Enstrom and concluded that:

[Enstrom] wrote a section of the 1983 Surgeon General's Report but now seems to have changed his mind on the smoking/mortality relationship. I say seems to have changed his mind because his writing is a little obtuse...⁴⁰

Enstrom's section of the 1983 Surgeon General's Report concluded that smoking cessation decreases the risk of heart attack and coronary heart disease. However, the conclusion went on to suggest that other risk factors for heart disease, such as blood pressure and cholesterol levels, needed further study.⁴¹

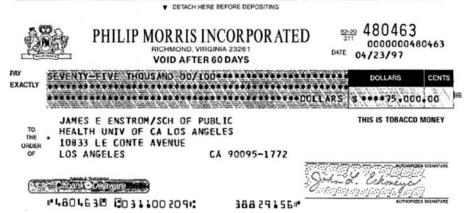
Enstrom submitted a pre-proposal to CTR for a study entitled "Mortality Trends Among Smokers and Non-smokers". In September 1991, two of the three CTR Scientific Advisory Board executive committee reviewers recommended that Enstrom be encouraged to submit a formal application. 42

In November 1991, a formal application was submitted to CTR⁴³ encouraged by the executive committee.⁴⁴ The project sought to investigate mortality trends from lung cancer, other smoking related causes, and all causes among smokers and non-smokers between 1966 and 1987 using data from the 1966–68 National Mortality Survey, 1986 National Mortality Survey, 1971–87 NHANES I Follow-up Study, and the American Cancer Society "follow-up of the 118,000 California residents in the 1959 Cancer Prevention Study (CPS-I)".⁴³ CTR funded the study for \$34 500 for the first year of a three year project in 1992 and funding was renewed in 1993⁴⁵ and 1997⁴⁶ ⁴⁷ with a \$25 000 supplement in 1997.⁴⁸

Part of the data from the CPS-I portion of the analysis was published in *Epidemiology*, but CTR funding was not acknowledged in the published paper. ⁴⁹ *Epidemiology* required authors to disclose their funding sources in a cover letter to the editor and the editor decides if these funding sources

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Figure 1 One of two cheques from Philip Morris funding the project "Relationship of low levels of active smoking to mortality" which included an analysis of the CPS-I dataset. Bates No. 2063610868, April 23, 1997. Legacy title: Check No. 0000000480463. Organisation authors: PM, Philip Morris. Person author: Eckmeyer JL. 75



should be published (www.epidem.com/pt/re/epidemiology/authorinfo.htm). Enstrom's cover letter is not available among the tobacco industry documents. Some of the data also are reported in the recent *BMJ* article by Enstrom and Kabat without acknowledging CTR funding (table 2). The analysis from the NHANES data was published with an acknowledgement of CTR funding.⁵⁰

Reanalysis of Fontham article

In April 1996, Enstrom submitted to the *Journal of the American Medical Association (JAMA)*⁵¹ a reanalysis of a major cohort study linking secondhand smoke and lung cancer published by Fontham and colleagues.⁵² ⁵³ The paper disclosed that the reanalysis was "supported in part by a special grant form the R.J. Reynolds Tobacco Company and Philip Morris".⁵⁴ Enstrom's analysis was conducted for Womble and Carlyle, one of the industry's law firms.⁶⁴ Before Enstrom submitted the manuscript to *JAMA*, he circulated it for comment among tobacco industry law firms, executives, and INBIFO (a Philip Morris scientific laboratory in Germany).⁵⁶

JAMA rejected the paper.⁶⁵ Although Enstrom's response to the peer reviewers' comments was circulated within the law firm Womble and Carlyle,⁶⁶ it is unclear whether the paper was resubmitted to *JAMA*. It was not published.

Strengthening the relationship with the tobacco industry

Relationship of low levels of active smoking to mortality

In January 1997, Enstrom submitted a research proposal to the Philip Morris Research Center,⁶⁷ where it was reviewed by the Scientific Research Review Committee (SRRC), a committee whose purpose was to "ensure that all scientific research, related to tobacco or smoking, conducted or funded by Philip Morris, ...serves relevant business needs". 68 The proposal, "Relationship of low levels of active smoking to mortality", sought to analyse data from four epidemiological cohorts: 1980–94 US Veterans study, 1971–92 NHANES I, 1976–92 NHANES II, and 1960–94 CPS-I in California. The CPS-I analysis was an expansion of the analysis funded by CTR. In his cover letter to Richard Carchman, Director of Scientific Affairs, Philip Morris, Enstrom stated:

These data are highly relevant to the ETS issue... A level of trust must be developed based on my past research on passive smoking and epidemiology in general in order to work out the best way for me to conduct this research. A substantial research commitment on your part is necessary in order for me to effectively compete against the large mountain of epidemiologic data and opinions that already exist regarding the health effects of ETS and active smoking.⁶⁹

Despite his status as a faculty member, Enstrom noted that a relationship of "mutual trust" with the tobacco industry in the context of its funding of his proposed research would minimise university involvement in the project. The proposal stated: "an unrestricted gift to James E. Enstrom / UCLA with mutual understanding/trust would minimize university restrictions and eliminate overhead costs." During January and February 1997, the proposal was reviewed by high level Philip Morris executives, lawyers and scientists who comprised the SRRC. To Tale At this time, the SRRC consisted of R Carchman, R Cox, C Ellis, A Kassman, J Nelson, GM Nixon, H Reif, W Reininghaus, and R Walk. Personnel at INBIFO

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commented that the amount of money requested seemed high when considering the amount of new scientific information that would likely be produced by the research, but stated that Enstrom "seems to have good connections/resources which might be useful in the future for other issues".⁷³

Philip Morris funded the project in April 1997 for \$150 000 to be paid in two instalments.^{74–79} (fig 1)

Research on secondhand smoke

During the 1990s, Enstrom also began to seek funding from the CIAR. In May and June 1996, Enstrom submitted a preproposal to CIAR for research on secondhand smoke, ^{80 81} including the re-analysis of the Fontham study (subsequently funded by RJ Reynolds and Philip Morris, as described above), reanalysis of the Varela dissertation data, a new casecontrol study, and follow up of several cohort studies.⁸¹ One of the proposed cohort studies was:

Conduct analysis of California CPS I cohort (51,000 males and 67,000 females) from 1960–94 and conduct follow-up of CPS I from selected states where individualized computerized death records are available (up to 200,000 and 300,000 females). Analysis will examine spousal smoking and death from lung cancer, coronary heart disease, and all causes.⁸¹

A formal proposal was submitted to Max Eisenberg, director CIAR, on 15 July 1996. In his cover letter, Enstrom noted that the results of his past research were favourable to the industry:

For the past three years I have done consulting and research on passive smoking for Jeffrey L. Furr of Womble Carlyle [law firm] on behalf of RJ Reynolds and Philip Morris. This research has found a number of results that raise serious questions about several published findings on the relationship of passive smoking to lung cancer and other diseases⁶⁴

Enstrom's proposal was discussed at the meeting on 14 August 1996 of CIAR's board of directors (consisting of tobacco industry executives and lawyers), which agreed that Eisenberg would meet with Enstrom to discuss the proposal in more detail.^{82 83}

Two sets of review comments on the 1996 Enstrom proposal were in the industry documents. One set, from INBIFO,⁸⁴ was critical of the proposal for its lack of a hypothesis and vague description of methods. The other review, probably from the CIAR board of directors,⁸⁵ was critical for similar reasons, as well as the fact that Enstrom once had acknowledged that secondhand smoke might be harmful. It noted:

Engstrom [sic] says that "[1]t is not possible to rule out a very weak relationship between passive smoking and mortality, especially for individual causes of death." How he reached a relationship for individuals when the statistics are not significant cannot be seen.⁸⁵

The reviewer went on to recommend an entirely different study for Enstrom:

In order to achieve something really new, CIAR could head for a pre-project which would check the protocols used for the above mentioned studies and put together the "ideal protocol" with all unthinkable adjustments for confounders and necessary investigations. This would give CIAR a "model protocol" which could then be used in order to create a "gold standard" against which other study protocols could be evaluated.⁸⁵

We did not find any documents showing that this "model protocol" was developed.

At its meeting on 25–26 November 1996, the CIAR board of directors decided that Max Eisenberg (Director, CIAR), Richard Carchman (Vice President, Scientific Affairs, Philip Morris), and Charles Green (CIAR, Chairman of Board and Principal Scientist, Research and Development, RJ Reynolds) would visit Enstrom at the University of California, Los Angeles. Enstrom's proposal was not funded at that time.

One result of the UCLA meeting was apparently to encourage Enstrom to collaborate with Geoffrey Kabat.⁸⁷ 88 Enstrom and Kabat had no prior record of collaboration. A search of PubMed on 25 July 2003 revealed that their only joint publication was the article in *BMJ* in 2003.¹

Kabat's ties with the tobacco industry

The documents provided no evidence that Kabat received any direct funding from the tobacco industry. Kabat had an ongoing indirect relationship with the tobacco industry since at least 1981, through Ernst Wynder, whose American Health Foundation had been funded by Philip Morris. Two of the 21 papers Wynder and Kabat co-authored between 1981 and 1995 were related to passive smoking, concluding that there was no association of secondhand smoke with lung cancer. He and Wynder published a review of secondhand smoke and lung cancer in a symposium proceeding which concluded that further evaluation of secondhand smoke and lung cancer risk was necessary. Although the symposium had been organised by individuals who were affiliated with the industry, tobacco industry funding was not expressly acknowledged in the symposium publication.

In 1997 Kabat co-authored a paper with several members of the tobacco industry's secret International ETS Consultants Project⁹⁴ critical of the evidence linking second-hand smoke with lung cancer. The project was run by industry lawyers to recruit, train, and pay scientists who would advocate the industry's position.¹¹ ⁹⁵

Receiving funding from CIAR

In September 1997, Carchman and Eisenberg had additional face-to-face meetings with Enstrom to discuss possible CIAR funding.96-103 On 3 November 1997, Eisenberg recommended that Enstrom's revised proposal, "Proposed research on passive smoking", be considered under the Directed Studies Program which is controlled by high level industry executives.104 The revised proposal only included the analysis of the CPS-I dataset.105 The proposal stressed Enstrom's past relationship with the ACS, noting that he "made use of this large and rich data base [CPS-I] with the cooperation of ACS". 105 Although Enstrom listed former ACS Vice Presidents for Epidemiology Clark Heath and Lawrence Garfinkel as unpaid consultants in the proposal (and in the BMJ disclosure, table 2), the proposal in the industry documents did not include letters of support from them, as is usual with consultants. In addition, ACS had advised against the use of the CPS-I dataset for Enstrom's analysis.6

In a memo commenting on the Enstrom proposal in November 1997, Chris Coggins of Lorillard Tobacco recommended funding the proposal. 106 An (unsigned) internal industry critique of the final proposal raised many of the same concerns about the Enstrom study that were subsequently raised by ACS6 and in the BMJ rapid responses

(http://bmj.bmjjournals.com/cgi/eletters/326/7398/1057). For example,

The proposal fails to distinguish between "ETS exposure" and "living with a spouse who smokes," makes light of substantial loss-to-follow-up expected in this cohort [CPS-I], freely assumes adequate adjustment can be made for the non-representativeness of the initial cohort, underplays the import of smoking cessation on the analysis, and is uncritical of positions established by anti-smoking community. ¹⁰⁷

Despite these concerns, the critic was "impressed" with the listing of Heath and Garfinkel from ACS as consultants on the proposal, as well as Kabat.¹⁰⁷ The reviewer suggested that their participation "should add credibility to the interpretation of the results".¹⁰⁷

On 19 November 1997 the CIAR board of directors voted to fund the Enstrom CPS-I proposal with modifications for \$525 000 from 1 June 1998 through 31 May 31 2001. 109

Additional financial ties

Enstrom had numerous other financial ties with the industry. He was paid to prepare analyses of scientific documents for tobacco industry law firms and served as a peer reviewer for Philip Morris' external research programme. 10 In 1998, Bill Rickert, chair of the consulting firm Labstat, Inc, was asked by the Canadian government to convene an expert committee to examine and make recommendations concerning cigarette toxicity. 11 Carchman, of Philip Morris, recommended Enstrom and long time tobacco industry consultant Peter Lee to serve on this committee. 12 Enstrom was a presenter at a scientific meeting organised by the tobacco industry, a June 2000 epidemiology conference organised by Philip Morris at which he discussed CPS-I results. 113 114

DISCUSSION

The published disclosure in the Enstrom and Kabat paper,1 which meets the requirements established by BMJ, does not reveal the full extent of the relationship the authors had with the tobacco industry. Tobacco industry funding of research is associated with favourable outcomes for the industry.115-117 Reasons for this observed association are complex and include sponsor involvement in the research questions asked, design, conduct, and publication of the study. 118 The Enstrom and Kabat paper is another example of how the content of even an extensive funding disclosure that meets the journal's requirements may not allow readers to understand fully the nature of the relationship between the authors and the research funder.10 11 119 The history of Enstrom's contacts and collaborations with the tobacco industry illustrates several reasons why the industry funds scientific research. First, although industry insiders were critical of Enstrom's methods, they nonetheless funded the research to enhance the industry's credibility, particularly by touting an association with the American Cancer Society. Second, the industry funded Enstrom because it perceived that his connections might be useful in the future. Enstrom had contributed to a Surgeon General's report on smoking and some of his early work suggested that secondhand smoke might be associated with cancer. Funding Enstrom allowed the industry to work with a scientist to advance its goals, as it has with other scientists.89 Third, funding Enstrom provided access for industry executives to seek to influence his research protocols by suggesting modifications. Lastly, funding Enstrom allowed the industry to connect him with other tobacco industry affiliated researchers, such as Kabat.

What this paper adds

Earlier studies of the health effects of secondhand smoke funded by the tobacco industry have failed to fully disclose the sponsor's role in the research. In addition to incomplete disclosures, industry sponsorship of research is often not entirely disclosed. Disclosure of financial ties can make reviewers more critical of manuscripts.

Comparing internal tobacco industry documents with a disclosure in a peer reviewed publication reveals that even an extensive financial disclosure statement that meets the journal's requirements can still provide an incomplete understanding of the tobacco industry's relationship with a project. The documents show how the tobacco industry funds research for multiple reasons, including gaining credibility or developing relationships with scientists that might be useful to the industry in the future. In addition to requiring financial disclosure, journals should require a statement in the Methods section of papers that clearly delineates the sponsor's role in designing, conducting, and reporting the results of a study.

The Enstrom and Kabat BMJ disclosure statement, when compared to the full extent of funding for the study as revealed in the internal industry documents, suggests that competing interest disclosures can be an inadequate mechanism for obtaining a complete understanding of the role of the sponsor in research. BMJ's competing interest disclosure policy specifically asks authors to disclose only competing financial interests related to their article; the disclosure of any other competing interests is left to the judgment of the author. Although the BMJ, like many other journals, asks for disclosure of relevant financial ties within the last five years, a requirement with which BMJ found that Enstrom and Kabat had complied, it might be more informative for readers to know that an author has a long, steady history of funding from a particular sponsor. Thus, journals could ask for disclosures of financial ties over a longer period of time.

Our findings raise the question of what additional steps journals can take to obtain the most meaningful disclosures from authors. An elaborate policing operation is not feasible or necessarily desirable, but a simple search could provide additional information. At least if the paper in question relates to tobacco, BMJ and other journals could conduct a quick search of the tobacco industry documents for the names of authors of papers on tobacco. The documents are freely available on the internet at legacy.library.ucsf.edu and bat.library.ucsf.edu. The industry documents could identify more details of the relationship between researchers and the tobacco industry than required in a simple funding disclosure. Beyond the possibility of identifying undisclosed funding, such searches could provide a description of the tobacco industry's role in the design, conduct or dissemination of the research. As an alternative to conducting tobacco industry document searchers themselves, journals could seek a peer reviewer with tobacco industry document research experience. Journals should discuss how the tobacco industry documents can be used most effectively to improve and inform the peer review process.

Journals should be alert to disclosures of funding from tobacco industry supported research organisations such as CIAR or the more recently formed Philip Morris External Research Program. Journals could keep on file peer reviewed articles that have investigated these tobacco industry programmes. These articles would describe the involvement of tobacco industry lawyers and executives in selecting projects for funding.³⁸ 120

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Readers of the Enstrom and Kabat paper are likely to critique it more rigorously if they are aware of the authors' long standing relationship with the tobacco industry and lack of support from the ACS. A randomised experiment to test the effects of financial disclosures on readers' evaluations of scientific journal articles found that readers rated articles with financial tie disclosures as less interesting, important, relevant, valid, and believable than those without financial disclosures.8 Requiring the disclosure of financial ties in all academic publications and presentations is frequently used as a way to manage financial conflicts of interest of researchers.14 16 121 An increasing number of scientific and medical journals are instituting requirements of disclosure of financial ties by authors of articles in the journals. Although a number of journals were initially opposed to financial disclosure policies,122 journal editors have begun to acknowledge that scientists might be influenced by financial interests and that disclosure is becoming the norm.12 13

In addition to reporting financial ties, it is also important to know what, if any, involvement the sponsor had in the design, conduct, and presentation of the research. One rapid response to the Enstrom and Kabat article suggested that it might be useful for editors to "require a statement about the role of the funding source in the design, conduct, analysis and reporting of the data".9 We believe that journals should require such a description as part of the Methods section in the paper, as required by Lancet (http://www.thelancet.com/ authorinfo).

The case of the Enstrom and Kabat paper also raises the issue of whether academic researchers should accept funding from the tobacco industry. A number of universities have adopted policies refusing tobacco industry funding for research.123 These institutions have decided to stop the tobacco industry's long standing efforts to manipulate research while hiding behind the respect and credibility of the academic institutions.

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Authors' affiliations

L A Bero, Department of Clinical Pharmacy, Institute for Health Policy Studies, University of California, San Francisco, San Francisco,

S Glantz, Center for Tobacco Control Research and Education, University of California, San Francisco

M-K Hong, Public Administration Analyst, Department of Clinical Pharmacy & Center for Tobacco Control Research and Education, University of California, San Francisco

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