

## Medical Journals and Industry Torts, rorts, and...trusted reports?

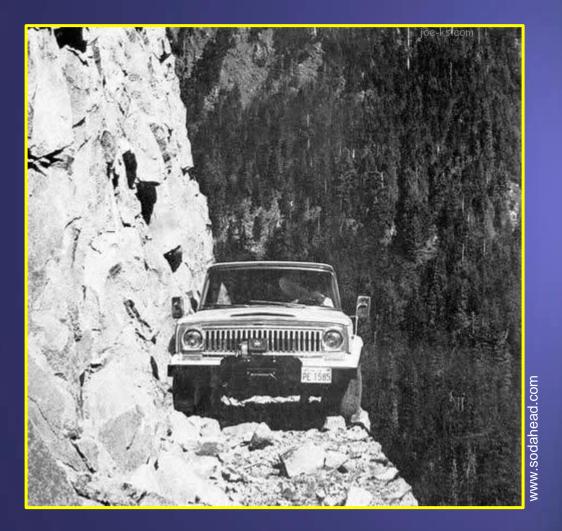
September 7 2011 ARCS, Sydney

Organised by the ARCS Medical Writing Interest Area Committee



| Chair         | <b>Professor Karen Woolley</b><br>ProScribe Medical Communications<br>Uni of Qld; Uni of the Sunshine Coast |
|---------------|---|
| Guest speaker | <b>Dr Larry Hirsch</b><br>Worldwide VP, Med Affairs Diabetes<br>BD (Becton Dickinson)                       |
| Panellists    | <b>Dr Annette Katelaris</b><br>Editor, The Medical Journal of Australia (MJA)                               |
|               | <b>Niamh Mullen</b><br>Journalist, Pharma In Focus  |
|               |   |

# Between a rock and a hard place



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# Publication planning - do you or don't you?

3. *Publication Activities*. To the extent that engages HCPs or HCIs to produce articles or other publications relating to Government Reimbursed Products (collectively "Dublication Activities") such LICDs on LICLs shall be referred to as Authors

## bioethics

Bioethics ISSN 0269-9702 (print); 1467-8519 (online) Volume 24 Number 6 2010 pp 273-283 doi:10.1111/j.1467-8519.2008.01702.x

### PUBLICATION ETHICS AND THE GHOST MANAGEMENT OF MEDICAL PUBLICATION

### SERGIO SISMONDO AND MATHIEU DOUCET

Publication Activities. Compliance personnel shall be involved in the review and approval of such annual Publications Plans, including any modification of an approved plan. The purpose of this review shall be to ensure that Publication Activities and related events are used for legitimate purposes in accordance with Policies and Procedures.



## People in glass houses shouldn't throw stones

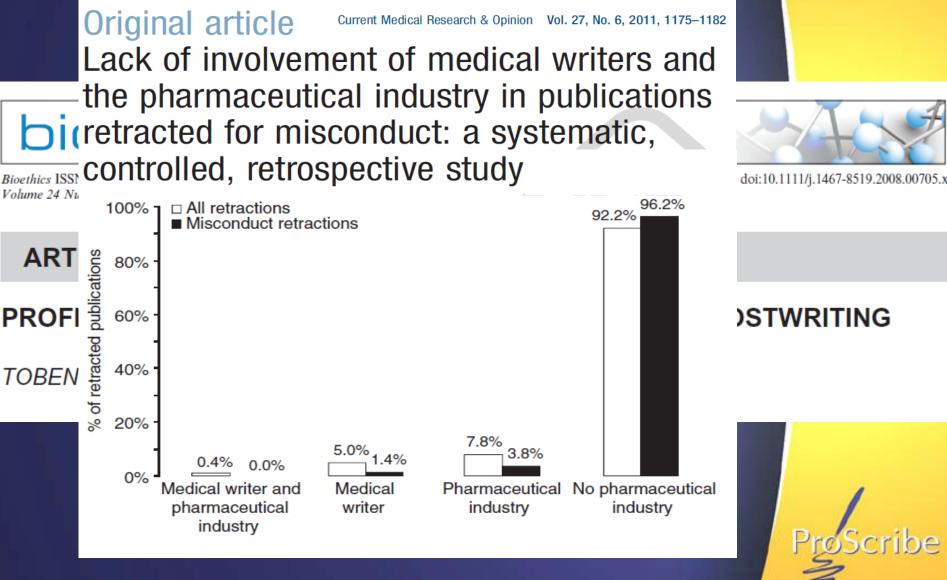


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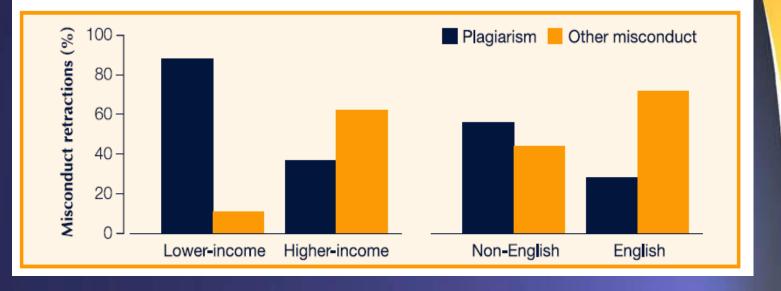


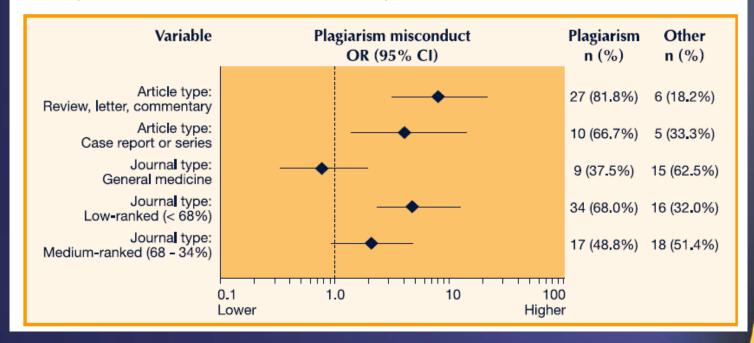
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# Who is the (most) guilty?



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## Stone the crows



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### Building trust with some editors?

"Anesthesia & Analgesia has experienced its share of fraud. Not a single case, including this one, has involved a study directly sponsored by a drug or device company. Sponsored studies are very closely audited, with each case report form checked against patient and laboratory data... Also, the process by which companies can perform highly detailed source-level audits offers an example for those engaged in investigator initiated trials of how to audit research in real time and maintain records of source-level documentation."

> Shafer SL. Shadow of Doubt . Anesthesia & Analgesia 2011; 112(3):498-500

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### Building trust with some editors?

"Auditing can not only uncover major problems, such as misconduct, but it can also expose minor ones, such as errors and irregularities in recording data. Auditing is a common practice in research sponsored by pharmaceutical or biotechnology companies, because the companies want to produce data that regulatory agencies will accept, and problems with the data can cost companies time and money."

> Zeng W. Resnik D. Research Integrity in China: Problems and Prospects. Developing World Bioethics 2010; 10(3):164-171



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### **Guest speaker**

Dr Larry Hirsch Worldwide VP, Med Affairs Diabetes BD (Becton Dickinson)