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Papers

Pharmaceutical industry sponsorship and research outcome and quality: systematic review

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Abstract

Objective To investigate whether funding of drug studies by the pharmaceutical industry is associated with outcomes that are favourable to the funder and whether the methods of trials funded by pharmaceutical companies differ from the methods in trials with other sources of support.

Methods Medline (January 1966 to December 2002) and Embase (January 1980 to December 2002) searches were supplemented with material identified in the references and in the authors' personal files. Data were independently abstracted by three of the authors and disagreements were resolved by consensus.

Results 30 studies were included. Research funded by drug companies was less likely to be published than research funded by other sources. Studies sponsored by pharmaceutical companies were more likely to have outcomes favouring the sponsor than were studies with other sponsors (odds ratio 4.05; 95% confidence interval 2.98 to 5.51; 18 comparisons). None of the 13 studies that analysed methods reported that studies funded by industry was of poorer quality.

Conclusion Systematic bias favours products which are made by the company funding the research. Explanations include the selection of an inappropriate comparator to the product being investigated and publication bias.

Introduction

Clinical research sponsored by the pharmaceutical industry affects how doctors practise medicine.¹ An increasing number of clinical trials at all stages in a product's life cycle are funded by the pharmaceutical industry,^{2,3} probably reflecting the fact that the pharmaceutical industry now spends more on medical research than do the National Institutes of Health in the United States.⁴ Most pharmacoeconomic studies are either done in-house by the drug companies or externally by consultants who are paid for by the company.^{5,6}

Results that are unfavourable to the sponsor—that is, trials that find a drug is less clinically effective or cost effective or less safe than other drugs used to treat the same condition—can pose considerable financial risks to companies. Pressure to show that the drug causes a

favourable outcome may result in biases in design, outcome, and reporting of industry sponsored research.⁷

A recent systematic review of the impact of financial conflicts on biomedical research found that studies financed by industry, although as rigorous as other studies, always found outcomes favourable to the sponsoring company.⁸ However, this review looked for papers published only in English, excluded reports in letters and abstracts, and looked at studies funded by other industries. We reviewed the relation between the source of funding of the research and the reported outcomes and investigated whether quality of the methods in studies funded by pharmaceutical companies differs from that in other studies.

Methods

Study selection

We included only studies that specifically stated that they analysed research sponsored by a pharmaceutical company, compared methodological quality or outcomes with studies with other sources of funding, and reported the results in quantitative terms. Outcomes of interest were conclusions about differences in drug effectiveness, adverse effects, cost outcomes, or publication status between industry funded trials and other trials. Work published in any language was eligible for inclusion.

Some studies analysed both pharmacological and non-pharmacological trials and combined research funded by drug companies and other industries into one group. In these cases, if most were non-pharmaceutical trials and were funded by other industries they were excluded.

Search strategy

We searched Medline from January 1966 to December 2002 using a combination of terms as both MESH subject headings (exploded) and key words ("clinical trials," "conflict of interest," "drug industry," "financial support," "publication bias" (subject heading only), "research design," and "research support.") We searched Embase from January 1980 to December 2002 using a combination of terms as subject headings (exploded) and key words ("clinical trials" (subject heading only), "drug industry," "ethics," "financial management," "methodology," and "ethics." To find more studies, we scanned the reference lists from each of the articles and searched the Cochrane methodology register. We placed messages on two email drug

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discussion groups, contacted content experts, and searched our personal libraries. In cases where the reported results were incomplete, we contacted the lead author and asked for further details. A single author (JL) did the initial selection of studies and sent copies of each of these studies to the other three authors for validation of the inclusion criteria.

Data collection

From each study, we extracted the study design, type of research assessed in the study, design of research assessed in the study, search strategy used to locate research, time period covered, drug or drug class, disease, number of industry and non-industry funded articles analysed in each study, how industry funding was defined, criteria used to assess methodological quality of the research, results with respect to methodological quality or outcome of the research, and primary purpose of study.

We provide a critical description of each included study, but do not assess methodological quality (see table 1). Since our included studies had a variety of designs—that is, cohort collections of trials, meta-analyses, and economic studies—and since we included letters and abstracts with limited descriptions of methods, we had no valid and reliable quality assessment instrument available for assessing their methodological quality. We did not use a component approach to assess their quality since this approach applies to randomised controlled trials.^{9 10}

Three of us (LB, OC, JL), who were not blinded to study authors or results, independently abstracted information. We resolved disagreements by consensus.

On the basis of the rationale that funding does affect the direction of effect, we did a meta-analysis on the studies that reported the effects of funding on the outcome of either pharmacoeconomic analyses or clinical trials in cases where odds ratios could be computed. The homogeneity test showed that the effect size did not differ between the studies ($P=0.17$). Using a Mantel-Haenszel test, we constructed a pooled odds ratio.¹¹ We used the program StatsDirect and considered $P < 0.05$ significant.

Results

Search results

The combined searches and other data sources found 3351 potential titles. We scanned titles and abstracts (where available) for mention of the pharmaceutical industry in either the title or the abstract or any suggestion that the study would deal with industry funding. We read 103 articles in full (eight in languages other than English); we retained 30 articles for analysis. Reasons for exclusion are detailed in the QUOROM statement (fig 1).

The studies by Friedberg et al²⁵ and by Knox et al²⁹ analysed the same set of 44 trials but looked at different aspects of the trials: conclusions about the usefulness of products in one case,²⁵ and how the trials had been reported in the other.²⁹ Only one article was duplicated in the two studies reported by the group including Chard, Tallon, and Dieppe^{15 18} (J Chard, personal communication, 2002). Seven of the nine articles in Kemmeren's²⁷ meta-analysis of third generation oral contraceptives were also included in Vandenbroucke's

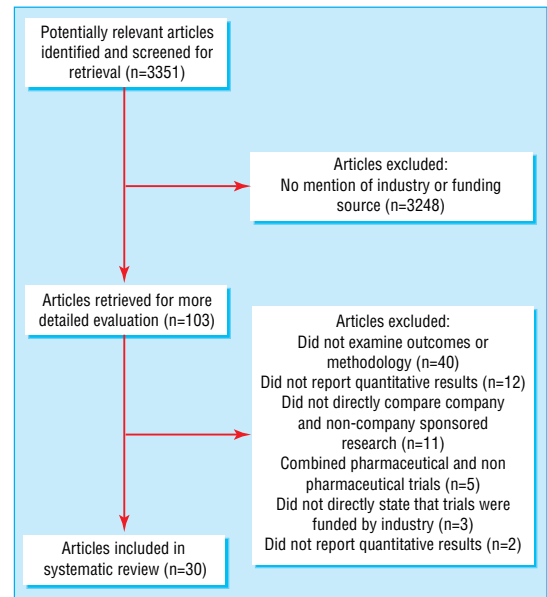


Fig 1 QUOROM statement

meta-analysis.³⁹ We found no other cases of double counting, but as some of the papers did not provide a full list of references we could not exclude the possibility of further overlap.

Characteristics of included studies

Table 1 gives the characteristics of the 30 studies included in this analysis.¹²⁻⁴¹ Six were reviews of pharmacoeconomic reports,^{12 23 26 29 34 36} two reviewed meta-analyses and systematic reviews,^{18 25} and the remaining 22 analysed groups of clinical trials.^{13-17 19-22 24 27 28 30-33 35 37-41} Eleven papers mentioned that some trials were funded by industry but offered no further definition of industry funding.^{17 18 22 24-26 28 34 36 39 40} In the other 15 papers the definition varied from a statement acknowledging industry funding in the article^{12 32} to a more comprehensive definition.³⁵

Relationship between source of funding and outcome

A total of 26 of the 30 studies reported results on the association of the outcome of the research and the source of funding: six examined the effects on publication,^{17 21 24 31 33 37} five looked at the outcome of pharmacoeconomic studies,^{12 23 26 34 36} and 16 analysed the outcome of clinical trials and meta-analyses of clinical trials (table 2).^{13-16 18-20 22 24 27 30 32 38-41}

Funding source and publication status

Research funded by drug companies was less likely to be published or presented than research funded by other sources (table 2).^{17 21} Three studies looked at time to publication,^{24 31 37} and two of these found that company sponsored research took longer to be published than research with other sources of funding.^{24 31} Research funded by drug companies was also more likely to be published in the proceedings of symposiums than non-industry sponsored research.³³

Funding source and economic outcomes

Pharmacoeconomic studies sponsored by the drug industry were more likely to report results favouring

Table 1 General characteristics of studies

Study (first author)	Type of publication	Research assessed	Design of research	Search strategy used to find research	Time covered	Number of separate trials funded by industry	Number of separate trials not funded by industry	How was industry funding defined
Azimi et al ¹²	Journal article, systematic review	Pharmacoeconomic studies	Cost effectiveness	Abridged Index Medicus	1 January 1990 to 30 June 1996	10	34	Explicit acknowledgment of support from private industry in article
Chard et al ¹³	Journal article, systematic review	Clinical trials	Prospective experimental, retrospective or prospective cohort studies, general disease reviews/specific intervention reviews, studies reporting pooled data	BIDS Institute for Scientific Information, Cochrane Library, Embase, Medline	1950 to March 1998	114 with pharmaceutical sponsorship out of 128 with commercial sponsorship	802	Source of funding declared in published article
Cho et al ¹⁴	Journal article, systematic review	Clinical trials	Experimental (randomised and non-randomised), observational	Computer generated list of random numbers from 625 symposiums, symposiums' articles matched to article from parent journal	Not stated	For quality score 50; for outcome 40	For quality score 77; for outcome: 112	Article acknowledged drug company providing funding or drugs or any authors employed by drug company
Clifford et al ¹⁵	Journal article, review	Clinical trials	Randomised controlled	Manual search for randomised trials in 5 general medical journals (convenience sample of 100 trials; 20/journal)	January 1999 to October 2000	44 by company only; 22 by company and at least 1 not for profit source	28	Acknowledged grant support; drug company employee listed as author; drug supplied by company; other types of company support
Davidson ¹⁶	Journal article, review	Clinical trials	Concurrent or crossover control group, before and after	Search of 5 major general medicine journals	1984	37	70	Acknowledgments stated support from company (excluding supply of drug or placebo) or any authors worked for company
Dickersin et al ¹⁷	Journal article, cohort study	Clinical trials	Observational, clinical trial, other	Manual search Institutional Review Board logs for studies receiving approval at one university	Studies approved in 1980 or before 1980 and still ongoing	42	472	Not stated
Dieppe et al ¹⁸	Journal letter, review	Meta-analyses	Meta-analysis	Search for meta-analyses in 8 high-impact journals	1993 to 1997	18	117	Not stated
Djulgovic et al ¹⁹	Abstract, systematic review	Clinical trials	Randomised controlled	From list of trials maintained by American Society of Hematology	1980 to 2000	23	7	Source of funding obtained from paper or investigator; association with manufacturer if drug supplied or grant support
Djulgovic et al ²⁰	Journal article	Clinical trials	Randomised controlled	Cochrane search strategy	1996 to 1998	35	101	Source of funding obtained from paper or investigator; association with manufacturer if drug supplied or grant support
Easterbrook et al ²¹	Journal article, retrospective cohort study	Clinical trials	Observational, experimental (clinical and non-clinical)	Manual search ethics committee records at 1 university	1 January 1984 to 31 December 1987	108	177	Direct contact with investigator or co-investigator
Freemantle et al ²²	Journal article, systematic review	Clinical trials	Randomised controlled	Embase, Medline, reviewed reference lists, contacted authors	1982 to 1997	Not stated	Not stated (total 105 trials)	Not stated
Freidberg et al ²³	Journal article, systematic review	Pharmacoeconomic studies	Cost minimisation, cost identification, cost effectiveness	HealthStar, Medline	1988 to 1998	20	24	Statement in article or by contacting investigator if no statement
Ioannidis ²⁴	Journal article, retrospective cohort study	Clinical trials	Randomised	Trials conducted under auspices of two AIDS trial groups	1986 to 1996	11	98	Not stated

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Table 1 General characteristics of studies—*continued from previous page*

Study (first author)	Type of publication	Research assessed	Design of research	Search strategy used to find research	Time covered	Number of separate trials funded by industry	Number of separate trials not funded by industry	How was industry funding defined
Jadad et al ²⁵	Journal article, systematic review	Systematic reviews, meta-analyses	Systematic review, meta-analysis	CINAHL, Cochrane Library, Embase, HealthStar, Medline, personal collections, reference lists	1988 to 1998	6	44	Not stated
Kamal-Bahl et al ²⁶	Journal article, systematic review	Pharmacoeconomic studies	Cost effectiveness, cost benefit, incremental cost analysis	HealthStar, Medline, contacted experts, reference lists, Cochrane Database of Systematic Reviews, National Health Service Economic Evaluation Database, National Institute for Clinical Excellence, Canadian Coordinating Office of Health Technology Assessment	1 January 1990 to 31 August 2001	4	8	Not stated
Kemmeren et al ²⁷	Journal article, meta-analysis	Clinical trials	Cohort, case-control	Medline, contacted experts, reference lists	October 1995 to December 2000	4	5	Industry funding explicitly stated in acknowledgement
Kjaergard et al ²⁸	Journal article, retrospective cohort study	Clinical trials	Randomised controlled and quasi-randomised controlled	Handsearch of <i>Hepatology</i> , Medline	January 1981 to August 1998	Not stated	Not stated (total 235)	Not stated
Knox et al ²⁹	Journal article, systematic review	Pharmacoeconomic studies	Cost minimisation, cost identification, cost effectiveness	HealthStar, Medline	1988 to 1998	20	24	Statement in article or by contacting investigator if no statement
Koeppe et al ³⁰	Journal letter, comment on meta-analysis	Clinical trials	Not stated	Studies were reported in previously published meta-analysis	Not stated	8	5	Authors employed by manufacturer, received grants, held stocks or options, in-house research of methodological or statistical analysis provided by manufacturer
Liebeskind et al ³¹	Abstract, systematic review	Clinical trials	Randomised controlled	Medline search, international trial registries	1957 to 1997	Not stated	Not stated (total 127)	Not stated but review discriminated between corporate sponsorship and corporate provision of drug
Mandelkern ³²	Journal letter, review	Clinical trials	Not stated	Manual review of one year of regular issues of <i>Journal of Clinical Psychiatry</i>	1997	16	16	Stated at front of article
Massie et al ³³	Journal letter, review	Clinical trials	Not stated	Articles published over 3 year period in 8 major journals	1980 to 1982	59	23	From article and by direct contact with authors
Neumann et al ³⁴	Journal article, systematic review	Pharmacoeconomic studies	Cost effectiveness	Medline, HealthStar, CancerLit	1975 to 1997	cost effectiveness 77; quality 32§	cost effectiveness 538; quality 118§	Not stated
Rochon et al ³⁵	Journal article, systematic review	Clinical trials	Randomised controlled	Medline search of 3 journals	January 1990 to November 1992	142 by company only; 25 both company and government/foundation	16 government/foundation; 59 no source	Acknowledged grant support; drug company employee listed as author; drug supplied by company; other types of company support
Sacristan et al ³⁶	Journal letter, review	Pharmacoeconomic studies	Cost effectiveness	Medline search in 6 general medical journals; hand search of <i>Pharmacoeconomics</i>	Medline: 1988 to 1994; <i>Pharmacoeconomics</i> 1993 to 1994	General medical journals 6; <i>Pharmacoeconomics</i> 18	General medical journals 63; <i>Pharmacoeconomics</i> 6	Not stated
Stern et al ³⁷	Journal article, retrospective cohort study	clinical trials	Randomised controlled, concurrent and historical controlled, uncontrolled	Search ethics committee submissions at 1 hospital	September 1979 to December 1988	Not stated	Not stated	Stated in submission to ethics committee

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Table 1 General characteristics of studies—*continued from previous page*

Study (first author)	Type of publication	Research assessed	Design of research	Search strategy used to find research	Time covered	Number of separate trials funded by industry	Number of separate trials not funded by industry	How was industry funding defined
Thomas et al ³⁸	Journal letter, systematic review	Clinical trials	Original research	Medline	1966 to 2001	48 (30 by company A; 18 by company B)	22	Funded by company; author was company employee; correspondence address used company address; statistical analysis done by company
Vandenbroucke et al ³⁹	Journal letter, review	Clinical trials	Not stated	Not stated	Not stated	4	9	Not stated
Wahlbeck et al ⁴⁰	Journal letter, comment on systematic review	Clinical trials	Not stated	Cochrane search strategy	Not stated	16	13	Not stated
Yaphe et al ⁴¹	Journal article, systematic review	Clinical trials	Randomised controlled	Hand search of 5 major medical journals	8 October 1992 to 1 October 1994	209	96	Funded by company; provision of study materials by company; author was company employee; statistical analysis done by company

*Additional data from author.

†Additional data from Freidberg et al.²³

‡Additional data from Massie et al.⁴²

§For cost effectiveness: number of ratios (studies contained more than one ratio; 70 ratios from pharmaceutical company funded studies and 7 from device company funded studies); for quality: number of studies.

the sponsor's product than studies with other sources of funding in all five articles that examined this question.^{12 23 26 34 36} In three cases, however, the bias in favour of industry funded research depended on the particular question being posed^{23 26} or on where the pharmacoeconomic analyses were published.³⁶

Funding source and outcomes of clinical trials and meta-analyses

Sixteen studies investigated the relationship between funding source and the outcomes of clinical trials and meta-analyses. Of these, 13 found that clinical trials and meta-analyses sponsored by drug companies favoured the product produced by the funder. Statistical significance for this finding was reported in eight of the 13 studies,^{13 14 16 19 20 38 40 41} and in another two there was a trend towards statistical significance.^{18 22} These studies covered a wide range of diseases, such as osteoarthritis of the knee,^{13 18} multiple myeloma,²⁰ various psychiatric problems,^{32 40} Alzheimer's disease,³⁰ and venous thromboembolism,³⁹ and a wide range of drugs, such as tacrine,³⁰ clozapine,⁴⁰ third generation oral contraceptives,³⁹ erythropoietin,¹⁹ antidepressants,²² and topical glucocorticosteroids.³⁸ One study that found no difference looked at the outcome of trials of treatment for HIV and associated complications and in this case the trials were monitored by the National Institutes of Health.²⁴ In one meta-analysis of third generation oral contraceptives,²⁷ the risk of venous thromboembolism for non-industry funded research was higher than that for industry sponsored trials, although the increased risk for thromboembolic disease was significant in both cases. Another study found no difference in outcomes in research published in five leading medical journals.¹⁵

Figure 2 shows the individual odds ratios and summary odds ratio for 18 different comparisons (15 studies) of the outcomes of industry funded and non-industry funded studies—seven from pharmacoeconomic analyses and 11 from clinical trials or meta-

analyses of clinical trials. The summary odds ratio was 4.05 (95% confidence interval 2.98 to 5.51).

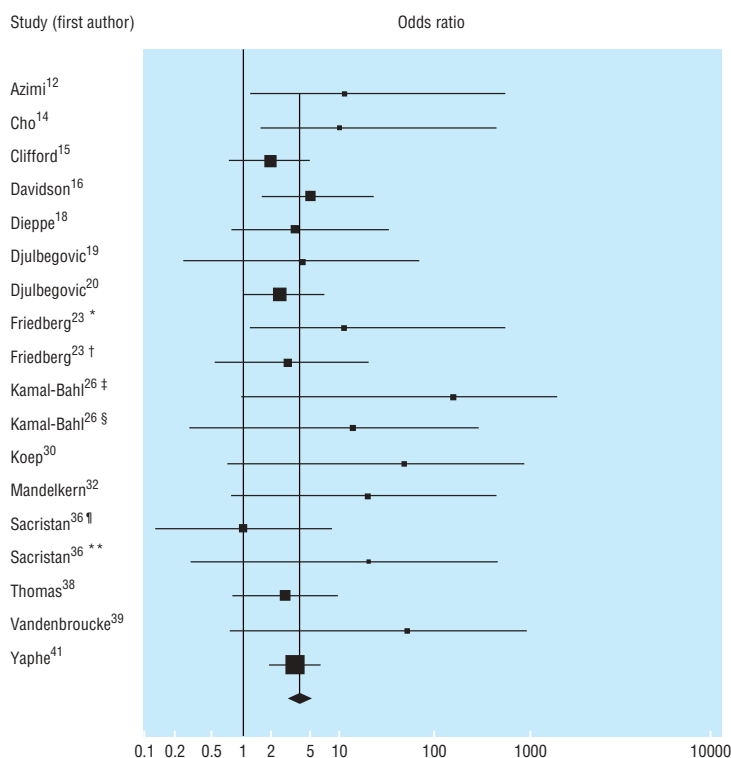


Fig 2 Source of funding and outcome in pharmacoeconomic analyses, clinical trials, and meta-analyses of clinical trials of drug treatments; for references see bmj.com (*Favourable qualitative results; †Overstatement of quantitative results; ‡Reporting possibility of cost effectiveness or cost savings of prophylaxis in entire high risk infant population either in point estimates or sensitivity analysis; §Reporting cost effectiveness or cost savings in either entire high risk populations or specific infant subgroups compared across studies; ¶Analyses reported in general medical journals; **Analyses reported in *Pharmacoeconomics*)

Table 2 Relation between source of funding and outcome of research

Author	Outcome question assessed by research	Results
Funding source and publication status		
Dickersin et al ¹⁷	Factors associated with publication of findings from clinical trials	Publication rate at one of two centres was considerably higher for studies funded by National Institutes of Health than for studies funded by drug industry (90.5% v 65.0%), but no indication that there was any difference in the tendency to publish significant results
Easterbrook et al ²¹	Are clinical trials with statistically significant results more likely to be published and what is the magnitude of this bias	Drug company sponsored trials of any type are significantly less likely to be published or presented compared to unfunded studies (OR=0.36; 95% CI 0.16 to 0.81); drug company sponsored clinical trials are significantly less likely to be published or presented compared to unfunded studies (0.17; 0.05 to 0.53)
Ioannidis ²⁴	Is time to completion and time to publication of clinical trials affected by statistical significance of results; description of natural history of trials	Trials where data managed by industry of shorter duration than those federally sponsored (P<0.01) but no difference in time to appearance in peer reviewed literature (P=0.33)†
Liesbeskind et al ³¹	Evidence of publication bias in randomised controlled trials	Median time between enrolment and publication of industry funded studies with positive results 3.5 years v 4.7 years for negative studies; time difference for studies with any type of funding 3.5 v 4.4 years for industry funded studies
Massie et al ³³	Whether source of funding of clinical trials on angina affected publication process	48% of drug company funded trials published in symposiums compared with 26% of studies with other types of funding or where source of funding not stated
Stern et al ³⁷	Extent to which publication is influenced by study outcome	Pharmaceutical industry funding not a statistically significant predictor of time to publication
Funding source and economic outcomes		
Azimi et al ¹²	How often do cost effectiveness analyses encourage a strategy requiring additional expenditures	Industry funded studies more likely to support a strategy requiring additional expenditures compared with those without such funding (9/10 v 15/34, P=0.01); among the 39 articles that supported additional expenditure the median cost effectiveness ratio of the 9 studies funded by industry was significantly higher than for the remainder (P=0.02)
Freidberg et al ²³	Relation between drug company sponsorship and economic assessment of oncology drugs	Drug company sponsored studies more likely to report favourable qualitative conclusions (19/20 v 15/24, P=0.04); overstatement of quantitative results not significantly different (6/20 v 3/24, P=0.26)
Kamal-Bahl et al ²⁶	Relation between drug company sponsorship and economic assessment of drugs for respiratory syncytial virus prophylaxis	Industry funded studies more likely to report possibility of cost effectiveness or cost savings of prophylaxis in entire high risk infant population either in point estimates or sensitivity analysis (4/4 v 0/8, P=0.002); when likelihood of reporting cost effectiveness or cost savings in either entire high risk populations or specific infant subgroups compared across studies, no difference between industry funded and non-industry funded studies (4/4 v 3/8, P=0.08)
Neumann et al ³⁴	Relation between drug company sponsorship and cost effectiveness ratios and per cent cost saving reported	For incremental cost effectiveness ratio: industry funded studies \$6000 v non-industry funded studies \$13,000 (P=0.003); for per cent cost saving: industry funded studies 21% v non-industry funded studies 9% (P=0.002)‡
Sacristan et al ³⁶	Relation between drug company sponsorship and results of cost effectiveness studies	General medical journals: 3/6 cost-effectiveness studies with industry funding had positive results v 31/63 with no funding or other source of funding; <i>Pharmacoeconomics</i> : 18/18 cost effectiveness studies with industry funding had positive results v 4/6 with no funding or other source of funding
Funding source and outcomes of clinical trials and meta-analyses		
Chard et al ¹³	Results of clinical trials for osteoarthritis of the knee	Projects commercially funded more likely to support intervention than non-commercially or non-specified funding source studies (P=0.024)*
Cho et al ¹⁴	Relation between drug company sponsorship of clinical trials and study outcome	Proportion of trials with favourable outcome higher in drug company sponsored research than in trials without company sponsorship (39/40 v 89/112, P<0.01)
Clifford et al ¹⁵	Relation between funding source and trial outcome	Studies favouring new product: industry only funding 30/44; mixed industry and non-industry funding 16/22; not for profit funding 15/28 (P=0.461)
Davidson ¹⁶	Relation between drug company sponsorship of clinical trials and study outcome	Studies supported by companies significantly more likely to support new therapies than trials with other sources of funding or where funding not stated (33/37 v 43/70, P=0.002)
Dieppe et al ¹⁸	Relation between drug company sponsorship and outcome of meta-analyses of treatment of osteoarthritis of the knee	Trend for meta-analyses sponsored by pharmaceutical industry to show a significant beneficial effect compared with meta-analyses with other source of funding or unknown sources of funding (16/18 v 81/117, P=0.084)
Djilbegovic et al ¹⁹	Relation between drug company sponsorship of clinical trials of erythropoietin and study outcome	Drug company funded trials showed positive results in 21/23 studies (P<0.0001) compared with 5/7 studies funded partially or completely with public resources (P=0.096)
Djilbegovic et al ²⁰	Relation between drug company sponsorship of clinical trials of the treatment of multiple myeloma and study outcome	Drug company funded trials showed positive results in 26/35 studies (P=0.004) compared with 54/101 studies funded by non-profit making organisations (P=0.608)
Freemantle et al ²²	Investigate potentially confounding factors which may affect clinical trials that assessed the relative efficacy of antidepressants	Most important structural predictor of outcome was trial sponsorship—trend towards increased efficacy of sponsor's drug (coefficient 0.097; 95% CI -0.03 to 0.23)
Ioannidis ²⁴	Is time to completion and time to publication of clinical trials affected by statistical significance of results; description of course of trials	No significant correlation between presence of statistical significance, study accrual, and whether data managed by industry or not (all correlation coefficients P<0.2)§
Kemmeren et al ²⁷	Relation between drug company sponsorship and outcome of studies comparing effects of second and third generation oral contraceptives on risk of venous thromboembolism	Odds ratio 1.3 (95% CI 1.0 to 1.7) for studies directly funded by pharmaceutical industry v 2.3 (1.7 to 3.2) in other studies

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Table 2 Relation between source of funding and outcome of research—*continued from previous page*

Author	Outcome question assessed by research	Results
Koeps et al ³⁰	Relation between drug company sponsorship of clinical trials on tacrine and study outcome	0/5 studies without corporate support found clinical benefit; 6/7 studies with corporate support found benefit (1 study could not be located)
Mandelkern ³²	Relation between drug company sponsorship of clinical trials in psychiatry and study outcome	16/16 drug company funded studies reported favourable outcome; 10/16 non-industry funded studies reported favourable outcome
Thomas et al ³⁸	Relation between drug company funding studies comparing topical glucocorticosteroids and whether study favours product made by sponsoring company¶	Product tested favoured in trial: 37/48 in company funded trials v 12/22 in trials with other sources of funding (P<0.001)
Vandenbroucke et al ³⁹	Relation between drug company sponsorship and outcome of studies comparing effects of second and third generation oral contraceptives on risk of venous thromboembolism	1/9 studies without industry funding found no higher risk of venous thromboembolism (relative risk 1.5-4.0, summary relative risk 2.4); 4/4 industry funded studies found no higher risk (relative risk 0.8-1.5, summary relative risk 1.1)
Wahlbeck et al ⁴⁰	Relation between drug company sponsorship of clinical trials on clozapine and study outcome	Odds of relapsing significantly in favour of clozapine in drug company sponsored trials (odds ratio 0.5; 95% CI 0.3 to 0.7) compared with non-sponsored trials (0.4; 0.1 to 1.4); both drug company sponsored and non-sponsored trials suggested clozapine mediates clinically important improvement compared with older drugs but industry sponsored trials were more positive (0.4; 0.2 to 0.7 v 0.3; 0.1 to 0.7); industry sponsored trials reported significantly fewer early dropouts compared with older drugs (0.5; 0.4 to 0.7) but non-sponsored trials did not (0.6; 0.3 to 1.2)
Yaphe et al ⁴¹	Relation between sources of support of research and published outcomes of randomised controlled drug trials	Positive outcomes: 181/209 trials with industry funding v 62/96 with non-industry funding (odds ratio 3.45, 95% CI 1.90 to 6.62)

*Results for all 128 commercially sponsored trials (114 funded by pharmaceutical industry).

†56 completed trials (of 98 where data was not managed by pharmaceutical industry) included in analysis.

‡70 trials funded by pharmaceutical companies and 7 funded by device companies.

§90 trials where full patient accrual completed (of 98 where data was not managed by pharmaceutical industry) included in analysis

¶Outcome of studies assessed by independent evaluators.

Relationship between source of funding and methodologic quality

A total of 13 studies examined the relationship between the source of funding and the methodological quality of the research (table 3).^{14-16 19 20 25 28 29 31-35} None of the 13 reported that industry funded studies had poorer methodological quality. Of the nine that provided statistical analyses, four found that drug company sponsored research had better quality scores.^{14 28 31 33}

Nine of the studies on clinical trials used well established methods of assessing quality.^{14 15 19 20 24 28 29 31 35} The single study that reported on the methods of pharmaco-economic analyses used commonly accepted criteria for assessing cost effectiveness.³⁴

One study evaluated the appropriateness of the comparators in clinical trials and found that a greater proportion of industry sponsored studies compared innovative treatment to either placebo or no therapy than did studies sponsored by public resources (60% v 21%; P<0.001).²⁰

Discussion

Research sponsored by the drug industry was more likely to produce results favouring the product made by the company sponsoring the research than studies funded by other sources. The results apply across a wide range of disease states, drugs, and drug classes, over at least two decades and regardless of the type of research being assessed—pharmaco-economic studies, clinical trials, or meta-analyses of clinical trials. The totality of the evidence reported in our meta-analysis of a subset of homogeneous studies suggests that there is some kind of systematic bias to the outcome of published research funded by the pharmaceutical industry.

Our results confirm and extend those reported by Bekelman et al.⁸ They identified only five studies that compared outcomes in research funded by pharmaceutical companies and other sources,^{14 16 20 23 41} and our study adds another 16

studies.^{12 13 15 18 19 22 24 26 27 30 32 34 36 38-40} Our results are also supported by Rochon and coworkers⁴³ (we excluded this paper because all of the trials were sponsored by drug companies and were, therefore, not comparable with trials lacking company funding.) They found that trials supported by manufacturers of non-steroidal anti-inflammatory agents almost always reported that the sponsor's drug was as or more effective and less toxic than the comparison drug.

Explanations

At least four possible explanations exist for favourable results seen in industry sponsored research. Firstly, pharmaceutical companies may selectively fund trials on drugs that they consider to be superior to the competition. Data collected so far, however, indicate that researchers cannot predict results of trials in advance.⁴⁴

Secondly, positive results could be the consequence of poor quality research conducted by industry. For example, low quality trials exaggerate the benefits of treatment by an average of 34%.^{45 46} We found that the research methods of trials sponsored by drug companies is at least as good as that of non-industry funded research and in many cases better. This does not guarantee the absence of bias in studies sponsored by the industry since outcome could be influenced by factors left out of quality scores, such as the question asked or the conduct or reporting of the study.^{7 47}

Thirdly, selecting an appropriate comparator is a key issue in planning a clinical trial.^{7 20 44} In the study by Rochon et al, in most cases in which the doses of the study and comparator drugs were not equivalent, the drug given at the higher dose was that of the supporting manufacturer.⁴³ As the authors saw, higher doses may bias the results in favour of effectiveness of the manufacturer's product. Safer also reports that in trials of psychiatric drugs the comparator drug is often given in doses outside the usual range or there is a rapid and substantial dose increase in the drug not manufactured by the sponsoring company.⁴⁸ In another instance, research funded by the company marketing flucona-

Table 3 Relation between source of funding and methodological quality of research

Study	Criteria used to assess methodological quality of research	Results
Cho et al ¹⁴	22 item validated scoring system	Study design in drug company sponsored clinical trials better than in research where no stated sponsorship (P=0.04)
Clifford et al ¹⁵	5 item validated scoring system (Jadad) plus component (individual items on Jadad scale and adequacy of concealment) approach	No difference by funding source for adequacy of allocation concealment (P=0.377); no difference by funding source for overall/composite score on Jadad scale (P=0.143)
Davidson ¹⁶	Sample size, blinding	For all trials higher rate of blinding for ones with industry sponsorship (67.5% v 41.8%, P=0.01); for trials investigating medications no difference in blinding (P=0.46); for sample size no difference between clinical trials supported by drug companies and those with other sources of funding or where funding not stated
Djulgovic et al ¹⁹	5 item validated scoring system (Jadad)	No difference in quality scores between randomised controlled trials funded solely by industry (mean 3.3 (SD 1.4); median: 3.5) and trials supported by public sources (mean 2 (SD 0.96); median: 2) (P=0.308)
Djulgovic et al ²⁰	5 item validated scoring system (Jadad)	Randomised controlled trials funded solely or partly by industry had trend towards higher quality scores (mean 2.94 (SD 1.3); median: 3) than trials supported by government or other non-profit organisations (mean 2.4 (SD 0.8); median: 2) (P=0.06)
Jadad et al ²⁵	7 point validated scoring system (Guyatt and Oxman)	6/6 industry funded systematic reviews and meta-analyses had serious flaws versus 34/44 non-industry funded reviews
Kjaergard et al ²⁸	5 point validated scale including: concealment of allocation, generation of allocation sequence, double blinding, dropouts/withdrawals, sample size	Clinical trials funded by either drug or device industry had higher quality than trials with no external funding (P<0.001); quality of publicly funded trials same as trials funded by drug or device industry (P=0.68)
Knox et al ²⁹	9 item scale developed for this study, including clinical design, generalisable data sources, statistical tests of significance performed on appropriate outcomes, statement regarding perspective, description of costs of the main included resources, description of time horizon, description of source of total costs differences, discussion of limitations, comparisons with other published studies	Drug company sponsored pharmacoeconomic analyses less likely to formally report on study generalisability, but were more likely to provide information on the key components of the methods section than were non-profit sponsored analyses
Liebeskind et al ³¹	100 point scale addressing 5 aspects of trial design and reporting: randomisation, outcome, inclusion/exclusion criteria, description of therapeutic regimen, statistical analysis	Clinical trials with corporate support had better quality than trials with non-profit support (mean 73.1 (95% CI 3.9) v 53.4 (9.8); P<0.0001)
Mandelkern ³²	Presence or absence of placebo control	5/16 industry funded clinical trials had placebo controls compared with 3/16 non-industry funded trials
Massie et al ³³	Not stated	Higher proportion of industry funded clinical trials were adequately controlled and designed than were trials with other sources of funding (71% v 33%, P<0.01)
Neumann et al ³⁴	Adherence to recommended protocols for cost effectiveness studies (adequate description of alternatives, study perspective clearly stated, discounted both costs and QALYs if needed, incremental analyses performed correctly) plus quality as judged by readers (scale of 1 to 7)	No difference between industry and non-industry funded studies on any measure: adequate description of alternatives P=0.30; study perspective clearly stated P=0.98; discounted both costs and QALYs P=0.65; incremental analyses performed correctly P=0.73; quality as judged by readers P=0.49
Rochon et al ³⁵	Modified version of Chalmers score including 14 items: control appearance and/or regimen, randomisation, blinding, patients blinded, observers blinded to treatment and results, previous estimate of numbers, testing compliance, results of randomisation on pretreatment variables and inclusion in analysis, major end points, post-beta estimate, confidence limits, statistical analyses, withdrawals after randomisation, side effects discussion	No difference in quality score between industry only funded clinical trials and those funded by government or foundations (mean 36.9% (SD 17.6%) v 37.1% (17.8%), P=0.271)

QALY=quality adjusted life year.

zole compared it with oral amphotericin B, a drug known to be poorly absorbed, thereby creating a bias in favour of fluconazole.⁴⁹ We did not consider who is finally responsible for the selection of the comparator—investigators, regulatory agencies, or sponsors.

Finally, our results suggest that publication bias may explain our finding of bias in favour of outcomes of research funded by industry. Although research sponsored by industry was less likely to be published than research with other sources of funding, the two studies with this finding did not specifically examine whether non-publication applied just to research with non-significant outcomes.^{17 21} In the past few years, manufacturers have attempted to prevent studies which are unfavourable to their products from being published in several high profile cases.^{50–52}

Massie and colleagues raise another possible source of publication bias.³³ They showed that research funded by industry appears more often in symposiums. Studies in symposiums are known to lack peer review and to favour the sponsor's product.^{14 53} Although the methods of industry funded trials are at

least equal to those in studies funded by other sources, the absence of peer review may result in an overly favourable interpretation of the results of a trial. Rochon and colleagues noted that claims of superiority for the sponsor's product were often not supported by the data.⁴³

Limitations

Our study has several limitations, primarily the difficulty in locating research examining the effects of company sponsorship. Our Medline and Embase searches found only 13 of the papers that we included,^{16–18 20 23 27–30 32 38 40 41} and the remaining 17 came from a search of the authors' personal files, suggestions by outside experts, or scanning of reference lists.^{12–15 19 21 22 24–26 31 33–37 39} The inability to critically evaluate the methodology in the abstracts and journal letters is another possible source of bias, and the conclusions of two of the letters that we included^{30 40} have been criticised.^{54 55}

Methods in studies sponsored by industry were at least as good as in studies with other sources of funding. Conclusions about overall quality can be

What is already known on this topic

When a pharmaceutical company funds research into drugs, studies are likely to produce results favourable to the sponsoring company's product

What this study adds

Research funded by drug companies was more likely to have outcomes that favour the sponsor's product than research funded by other sources

This cannot be explained by the reported quality of the methods in research sponsored by industry

The result may be due to inappropriate comparators or to publication bias

influenced by the instrument used,⁹ and some of the scales may have missed important criteria.

Research sponsored by the pharmaceutical industry is facing a number of challenges. Questions have been raised about the mismatch between the research agendas of the pharmaceutical industry and consumers of research.⁵⁶ Meta-analysts are confronted with the problems of duplicate publication of data from company funded trials and the withholding of data.^{49 57 58}

Leading medical journals recently decided to establish more rigorous criteria for the acceptance of research sponsored by industry; this is a step in the right direction towards increasing the credibility of studies paid for by drug companies.⁵⁸ The revised CONSORT statement should also help improve the quality of clinical research.^{59 60} In addition, authors and editors should consider including a statement concerning prior beliefs of the investigators about the uncertainty of the treatments that are reported. Finally, all clinical trials should be registered prospectively as the only way to prevent publication bias.⁶¹ The proposal to do so which was put forward in 1986⁶² has been periodically renewed,⁶³⁻⁶⁵ but to this date has not been implemented.

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