Discrepancy between Results and Abstract Conclusions in Industry- vs Nonindustry-funded Studies Comparing Topical Prostaglandins

TARIQ ALASBALI, MICHAEL SMITH, NOA GEFFEN, GRAHAM E. TROPE, JOHN G. FLANAGAN, YAPING JIN, AND YVONNE M. BUYS

- PURPOSE: To investigate the relationship between industry- vs nonindustry-funded publications comparing the efficacy of topical prostaglandin analogs by evaluating the correspondence between the statistical significance of the publication's main outcome measure and its abstract conclusions.
- DESIGN: Retrospective, observational cohort study.
- METHODS: English publications comparing the ocular hypotensive efficacy between any or all of latanoprost, travoprost, and bimatoprost were searched from the MEDLINE database. Each article was reviewed by three independent observers and was evaluated for source of funding, study quality, statistically significant main outcome measure, correspondence between results of main outcome measure and abstract conclusion, number of intraocular pressure outcomes compared, and journal impact factor. Funding was determined by published disclosure or, in cases of no documented disclosure, the corresponding author was contacted directly to confirm industry funding. Discrepancies were resolved by consensus. The main outcome measure was correspondence between abstract conclusion and reported statistical significance of the publications' main outcome measure.
- RESULTS: Thirty-nine publications were included, of which 29 were industry funded and 10 were nonindustry funded. The published abstract conclusion was not consistent with the results of the main outcome measure in 18 (62%) of 29 of the industry-funded studies compared with zero (0%) of 10 of the nonindustry-funded studies (P = .0006). Twenty-six (90%) of the industry-funded studies had proindustry abstract conclusions.
- CONCLUSIONS: Twenty-four percent of the industry-funded publications had a statistically significant main outcome measure; however, 90% of the industry-funded

See accompanying Editorial on page 1. Accepted for publication Jul 1, 2008.

From the Department of Ophthalmology and Vision Sciences, University of Toronto, Toronto Western Hospital, Toronto, Ontario, Canada (T.A., M.S., N.G., G.E.T., J.G.F., Y.M.B.); the Department of Ophthalmology, King Faisal University, King Fahad Hospital of the University, Riyad, Saudi Arabia (T.A.); the School of Optometry, University of Waterloo, Waterloo, Ontario, Canada (J.G.F.); and the Department of Public Health Sciences, University of Toronto, Toronto, Ontario, Canada (Y.I.).

Inquiries to Yvonne M. Buys, Toronto Western Hospital, 399 Bathurst Street. EW6-405. Toronto. Ontario. Canada M5T 2S8: e-mail: v.buvs@ studies had proindustry abstract conclusions. Both readers and reviewers should scrutinize publications carefully to ensure that data support the authors' conclusions. (Am J Ophthalmol 2009;147:33–38. © 2009 by Elsevier Inc. All rights reserved.)

INANCIAL RELATIONSHIPS BETWEEN PHARMACEUTIcal companies and researchers and funding of medical research by drug companies has increased dramatically during the last two decades. 1-4 This can result in industry bias where the source of funding of clinical trials either affects the results in a systematic way or leads to selective presentation of the results. Industry funding often has been associated with proindustry results^{2,5–20} and publication bias, 21-23 which can affect the interpretation and presentation of outcomes resulting in conclusions that overstate results without statistical support. The purpose of this study was to investigate the relationship between industry- vs nonindustry-funded publications comparing ocular hypotensive efficacy of the topical prostaglandin analogs (PGA) latanoprost 0.005%, travoprost 0.004%, and bimatoprost 0.03% by evaluating the correspondence between the statistical significance of the publication's main outcome measure and its published abstract conclusions.

METHODS

A MEDLINE SEARCH FROM 1966 TO THE SECOND WEEK OF November 2007 using any combination of the keywords *latanoprost*, *travoprost*, and *bimatoprost* was conducted. The title and abstracts from the initial search were reviewed and those included were English language publications comparing the intraocular pressure (IOP)-lowering efficacy of any combination of latanoprost; travoprost; or bimatoprost. The complete articles were obtained and the references also were searched to identify relevant publications missed during the initial search.

Each publication was reviewed by three independent observers using a standardized data collection sheet evaluating: source of funding, industry author, study quality, main outcome measure, statistical significance (P < .05) of main outcome measure, abstract conclusion, correspon-



TABLE 1. Grading of Study Quality

Level	Criteria
1A: Meta-analysis (to assign this level, you must answer 'yes' to all questions)	Does the paper report a comprehensive search for evidence?
	Did the authors avoid bias in selecting articles for inclusion?
	Did the authors asses each article for validity?
	Does the paper report clear conclusions that are supported by the data and appropriate analysis?
1A: Large RCT (to assign this level, you must answer 'yes' to all questions)	Were patients randomly allocated to treatment groups?
	Was follow-up at least 80% complete?
	Were both the patients and the investigators blind to the treatment the patient received?
	Were the patients analyzed in the treatment groups to which they were assigned?
	Was the sample size large enough to detect the outcome of interest?
1B: NRCT	NRCT or cohort study with indisputable results
2: RCT	RCT or overview that did not meet level 1
3	NRCT or cohort study
4	Other (case series without controls, case report, expert opinion, etc.)

TABLE 2. Summary of Industry- vs Nonindustry-funded Studies Comparing Topical Prostaglandins

Outcome Studied	Industry-funded (n $= 29$)	Nonindustry-funded (n = 10)	P value
Noncorrespondence of main outcome and conclusions	18 (62%)	0 (0%)	.0006ª
Statistically significant (P < .05) main outcome	7 (24%)	2 (20%)	1.00 ^a
Number of IOP comparisons, mean \pm SD (median, range)	17.4 ± 11.6 (14, 1 to 45)	13.0 ± 11.4 (8, 1 to 30)	.31 ^b
Mean study quality	2.4 ± 1.1	2.0 ± 0.7	.27 ^b
Industry coauthor	18 (62%)	N/A	
Journal impact factor	2.14 ± 1.32	2.33 ± 1.51	.72 ^b

IOP = intraocular pressure; N/A = not applicable; SD = standard deviation.

outcome measure and abstract conclusion, total number of IOP outcomes compared, and journal impact factor. Any discrepancies between the three reviewers were resolved by consensus.

Funding was determined by published disclosure, or in cases of no documented disclosure, the corresponding author was contacted directly to confirm any direct funding of the study. In one case, the pharmaceutical company was contacted to verify funding. Study quality was assessed according to the criteria in Table 1.²⁴ Journal impact factors from 2006 were assigned to each publication.

The main outcome measure was the correspondence between the statistical significance of the publication's main outcome measure and its published abstract conclusion. Statistical analysis included the Fisher exact test for categorical data and the Student *t* test for

RESULTS

A TOTAL OF 180 ARTICLES WERE IDENTIFIED BY THE ORIGInal search. After reviewing the abstracts, 39 met the inclusion criteria and were included in the study. In reviewing the references of these publications, no additional publications were found. Of the 39 publications, 35 were studies that directly compared two or three of the PGAs and four were meta-analyses. Thirty-five of the publications included a disclosure statement, and four had no documented disclosure. The authors were contacted regarding these four publications; two publications^{25,26} confirmed industry funding of the study and two reported no funding. One of the publications for which the author denied industry funding subsequently was discovered to have received industry funding after direct communication



^aFisher exact test.

^bStudent t test.

industry funding. Twenty-nine (74%) of the publications were industry funded (18 by Allergan, Irvine, California, USA, ^{25–42} 10 by Alcon, Fort Worth, Texas, USA^{43–52} and one by Pfizer, New York, New York, USA⁵³) and 10 (26%) were nonindustry funded (nine had no funding ^{54–62} and one had government funding ⁶³). There was an industry coauthor in 18 (62%) of the industry-funded publications. The results are summarized in Table 2.

Statistically significant main outcome measures were reported in 7 (24%) industry-funded publications and in 2 (20%) nonindustry-funded publications (P = 1.00, Fisher exact test). Correspondence between the results of the main outcome measure and the abstract conclusions was found in 11 (38%) of the industry-funded publications vs 10 (100%) of the nonindustry-funded publications (P = .0006, Fisher exact test). Twenty-six (90%) of the industry-funded studies had proindustry conclusions.

The mean number of IOP comparisons reported were 17.4 ± 11.6 for industry-funded publications and 13.0 ± 11.4 for nonindustry-funded publications (P = .31, Student t test). The mean study quality was 2.4 ± 1.1 for industry-funded publications compared with 2.0 ± 0.7 for nonindustry-funded publications (P = .27, Student t test). The mean journal impact factor also was similar between industry-funded (2.14) and nonindustry-funded (2.33) publications (P = .72, Student t test).

DISCUSSION

WE FOUND THAT 62% OF THE INDUSTRY-FUNDED VS NONE of the nonindustry-funded studies' abstract conclusions did not correspond with the results of the main outcome measure (P = .0006, Fisher exact test). Although only 24% of the industry-funded publications had a statistically significant main outcome measure, 90% of the industry funded studies had a proindustry abstract conclusion.

The influence of industry on publications involving a wide range of diseases and drugs is well documented. 2,5-20 Kjaergard and Als-Nielsen reviewed 159 randomized controlled trials from 12 specialties and found that when financial interests were disclosed, the authors' conclusions significantly favored experimental intervention. 11 Lexchin and associates reviewed 30 pharmaceutical-sponsored studies and found "systematic bias to the outcome of published research funded by the pharmaceutical industry." 13 Als-Nielsen and associates evaluated 370 randomized controlled trials over a broad area of diseases and found that conclusions significantly favored experimental drugs in trials funded by for-profit organizations. 15 A review of 124 meta-analyses of antihypertensives found that industry support was not associated with more favorable results, but was associated with more favorable conclusions. 20 To our knowledge, ours is the first attempt to determine industry

Prostaglandin analogs currently are the first-line therapy for the treatment of glaucoma, representing 43.9% of glaucoma medications dispensed in Ontario, Canada, in 2007. Latanoprost 0.005% (Pfizer) was first available in Ontario in June 1997, followed by travoprost 0.004% (Alcon) in November 2001 and bimatoprost 0.03% (Allergan) in May 2002. These three medications belong to the same class and therefore are competing directly for the same market share. The use of PGAs is influenced by the number and quality of publications.

Of the 39 publications studied, 29 (74%) were industry funded. The high proportion of industry-funded studies is consistent with reports of increased funding of biomedical research by the biomedical industry. ^{1–4} Our definition of industry funding, however, may be considered conservative because we did not investigate the financial ties of each author and included only studies with direct industry funding. Evaluating financial disclosures of individual authors is difficult because many authors have support from several companies, although the amount of support per company may vary.

Similar to studies in other disciplines, we found no difference in significant main outcome measures, \$^{10,16,20,23}\$ study quality, \$^{2,7,10,11,13,15,16,23}\$ or journal impact factor \$^{8,15}\$ between industry- and nonindustry-funded studies comparing PGAs. Four of the seven industry-funded studies with significant main outcome measure were of the lowest level of quality (level 4). \$^{29,35,37,40}\$ There were no level four studies in the nonindustry-funded group. Six of the eight industry-funded publications with level four study quality were published in journals with a higher (> 2.2) impact factor. \$^{29,35-37,40,52}\$

The discrepancy between the results of the main outcome measure and abstract conclusions stems from the interpretation of surrogate outcomes or multiple comparisons assigning undue attention to significant results while minimizing nonsignificant results. This is commonly referred to as "spin." To evaluate for possible data dredging, we compared the total number of IOP outcome comparisons presented in the results and found no difference between industry- and nonindustry-funded studies. The mean number of IOP outcome measures was 17.4 ± 11.6 (range, one to 45) for industry-funded publications and 13.0 ± 11.4 (range, one to 30) for nonindustry-funded publications (P = .31).

This study raises concerns regarding undue industry influence in publications on PGAs. Less industry funding and increased funding by peer-reviewed governmental agencies or other organizations may remove this bias. Authors should provide transparency in the interpretation and conclusions of their study, and it is the role of journal editors and reviewers to ensure that data are not misrepresented. It is important that journals develop strong guidelines to limit potential bias by creating minimum standards in reporting disclosure and results. Full disclo-



the study is necessary. Three of the studies in our review were funded directly by industry; however, there was no published disclosure. The requirement of registration of all clinical trials at the time of design and before the collection of data and making available all data could minimize inappropriate data analysis and selective reporting of results. The inclusion in the abstract of a

heading specifying the main outcome measure and the statistical results of the main outcome measure may improve transparency of the study findings. Ultimately, however, it is the responsibility of the reader to scrutinize abstract conclusions carefully to ensure that they are supported by the data reported in the RESULTS SECTION of the article.

THE AUTHORS INDICATE NO FINANCIAL SUPPORT OR FINANCIAL CONFLICT OF INTEREST. DR BUYS IS AN ADVISORY BOARD member for Allergan Inc and Pfizer and received lecture fees from Alcon. Drs Buys and Trope receive grants from Allergan Inc, Pfizer, and Canadian Institutes of Health Research. Dr Smith received a travel grant from Alcon and Allergan Inc. Involved in design of study (Y.M.B.); literature search (Y.M.B., T.A.); reviewing publications (T.A., M.S., N.G., G.E.T., J.G.F., Y.M.B.); statistical analysis (Y.M.B., Y.J.); and preparation of manuscript (Y.M.B., T.A.). This study was a review of published literature and Institutional Review Board approval was not necessary.

REFERENCES

- Moses III H, Martin JB. Academic relationships with industry: a new model for biomedical research. JAMA 2001;285: 933–935.
- Bekelman JE, Li Y, Gross CP. Scope and impact of financial conflicts of interest in biomedical research: a systematic review. JAMA 2003;289:454–465.
- 3. Buchkowsky SS, Jewesson PJ. Industry sponsorship and authorship of clinical trials over 20 years. Ann Pharmacother 2004;38:579–585.
- 4. Patsopoulos NA, Ioannidis J, Analatos A. Origin and funding of the most frequently cited papers in medicine: database analysis. BMJ 2006;332:1061–1064.
- 5. Davidson RA. Source of funding and outcome of clinical trials. J Gen Intern Med 1986;1:155–158.
- Rochon PA, Gurwitz JH, Simms RW, et al. A study of manufacturer-supported trials of nonsteroidal anti-inflammatory drugs in the treatment of arthritis. Arch Intern Med 1994;154:157–163.
- 7. Cho MK, Bero LA. The quality of drug studies published in symposium proceedings. Ann Intern Med 1996;124:485–489.
- Friedberg M, Saffran B, Stinson TJ, Nelson W, Bennett CL. Evaluation of conflict of interest in economic analyses of new drugs used in oncology. JAMA 1999;282:1453–1457.
- Djulbegovic B, Lacevic M, Cantor A, et al. The uncertainty principle and industry-sponsored research. Lancet 2000;356: 635–638.
- 10. Clifford TJ, Barrowman NJ, Moher D. Funding source, trial outcome and reporting quality: are they related? Results of a pilot study. BMC Health Services Res 2002;2:18.
- Kjaergard LL, Als-Nielsen B. Association between competing interests and author's conclusions: epidemiological study of randomized clinical trials published in the BMJ. BMJ 2002;325:249–253.
- Baker CB, Johnsrud MT, Crismon ML, Rosenheck RA, Woods SW. Quantitative analysis of sponsorship bias in economic studies of antidepressants. Br J Psychiatry 2003; 183:498–506.
- 13. Lexchin J, Bero LA, Djulbegovic B, Clark O. Pharmaceutical industry sponsorship and research outcome and quality: systematic review. BMJ 2003;326:1167–1170.
- Leopold SS, Warme WJ, Fritz Braunlich E, Shott S. Association between funding source and study outcome in ortho-

- Als-Nielsen B, Chen W, Gluud C, Kjaergard LL. Association of funding and conclusions in randomized drug trials. A reflection of treatment effect or adverse events? JAMA 2003;290:921–928.
- Bhandari M, Busse JW, Jackowski D, et al. Association between industry funding and statistically significant proindustry findings in medical and surgical randomized trials. Can Med Assoc J 2004;170:481–483.
- 17. Finucane TE, Boult CE. Association of funding and findings of pharmaceutical research at a meeting of a medical professional society. Am J Med 2004;117:842–845.
- Shah RV, Albert TJ, Bruegel-Sanchez V, Vaccaro AR, Hilibrand AS, Grauer JN. Industry support and correlation to study outcome for papers published in Spine. Spine 2005;30: 1099–1104.
- 19. Perlis RH, Perlis CS, Wu Y, Hwang C, Joseph M, Nierenberg AA. Industry sponsorship and financial conflict of interest in the reporting of clinical trials in psychiatry. Am J Psychiatry 2005;162:1957–1960.
- Yank V, Rennie D, Bero LA. Financial ties and concordance between results and conclusions in meta-analysis: retrospective cohort study. BMJ 2007;335:1202–1205.
- 21. Easterbrook PJ, Berlin JA, Gopalan R, Matthews DR. Publication bias in clinical research. Lancet 1991;337:867–872.
- 22. Dickerson K, Min Y-I, Meinert CL. Factors influencing publication of research results. Follow-up of applications submitted to two institutional review boards. JAMA 1992; 267:374–378.
- Lynch JR, Cunningham MR, Warme WJ, Schaad DC, Wolf FM, Leopold SS. Commercially funded and United Statesbased research is more likely to be published; good-quality studies with negative outcomes are not. J Bone Joint Surg [Am] 2007;89:1010–1018.
- 24. Users' Guides to the Medical Literature: Essentials of Evidence-based Clinical Practice. Chicago, Illinois: American Medical Association, 2001.
- Gandolfi S, Simmons ST, Sturm R, Chen K, VanDenburgh AM. Three-month comparison of bimatoprost and latanoprost in patients with glaucoma and ocular hypertension. Adv Ther 2001;18:110–121.
- Dirks MS, Noecker RJ, Earl M, Roh S, Silverstein SM, Williams RD. A 3-month clinical trial comparing the IOPlowering efficacy of bimatoprost and latanoprost in patients



- 27. Simmons ST, Dirks MS, Noecker RJ. Bimatoprost versus latanoprost in lowering pressure in glaucoma and ocular hypertension: results from parallel-group comparison trials. Adv Ther 2004;21:247–262.
- DuBiner H, Cooke D, Dirks M, Stewart W, VanDenburgh A, Felix C. Efficacy and safety of bimatoprost in patients with elevated intraocular pressure: a 30-day comparison with latanoprost. Surv Ophthalmol 2001;45:S353–S360.
- 29. Choplin N, Bernestein P, Batoosingh A, Whitcup SM. A randomized, investigator-masked comparison of diurnal responder rates with bimatoprost and latanoprost in the lowering of intraocular pressure. Surv Ophthalmol 2003;49: S19–S25.
- Noecker R, Dirks M, Choplin N, et al. A six-month randomized clinical trial comparing the IOP-lowering efficacy of bimatoprost and latanoprost in patients with ocular hypertension or glaucoma. Am J Ophthalmol 2003;135:921– 927
- 31. Noecker RJ, Earl ML, Mundorf T, Peace J, Williams RD. Bimatoprost 0.03% versus travoprost 0.004% in black Americans with glaucoma or ocular hypertension. Adv Ther 2003;20:121–128.
- 32. Cantor LB, WuDunn D, Cortes A, Hoop J, Knotts S. Ocular hypotensive efficacy of bimatoprost 0.03% and travoprost 0.004% in patients with glaucoma or ocular hypertension. Surv Ophthalmol 2004;49:S12–S18.
- Walters TR, DuBiner HB, Carpenter SP, Khan B, VanDenburgh AM, Bimatoprost Circadian IOP Study Group. 24-Hour IOP control with once-daily bimatoprost, timolol gel-forming solution, or latanoprost: a 1-month, randomized, comparative clinical trial. Surv Ophthalmol 2004;49:S26–S35.
- Walt J, Lee J. A cost-effectiveness comparison of bimatoprost versus latanoprost in patients with glaucoma or ocular hypertension. Surv Ophthalmol 2004;49:36–44.
- Holmstrom S, Buchholz P, Walt J, Wickstrøm J, Aagren M. Analytic review of bimatoprost, latanoprost and travoprost in primary open-angle glaucoma. Curr Med Res Opin 2005; 21:1875–1883.
- Law SK, Song BJ, Fang E, Caprioli J. Feasibility and efficacy of a mass switch from latanoprost to bimatoprost in glaucoma patients in a prepaid health maintenance organization. Ophthalmology 2005;112:2123–2130.
- 37. Zeitz O, Matthiessen E, Reuss J, et al. Effects of glaucoma drugs on ocular hemodynamics in normal tension glaucoma: a randomized trial comparing bimatoprost and latanoprost with dorzolamide. BMC Ophthalmology 2005;5:6.
- Cantor LB, Hoop J, Morgan L, Wudunn D, Catoira Y, Bimatoprost-Travoprost Study Group. Intraocular pressurelowering efficacy of bimatoprost 0.03% and travoprost 0.004% in patients with glaucoma or ocular hypertension. Br J Ophthalmol 2006;90:1370–1373.
- Fiscella R, Walt J. Estimated comparative costs of achieving a 20% reduction in intraocular pressure with bimatoprost or latanoprost in patients with glaucoma or ocular hypertension. Drugs Aging 2006;23:39–47.
- 40. Noecker RJ, Earl ML, Mundorf TK, Silverstein SM, Phillips MP. Comparing bimatoprost and travoprost in black Americans. Curr Med Res Opin 2006;22:2175–2180.
- 41. Konstas A, Hollo G, Irkec M, et al. Diurnal IOP control with

- over observer-masked three-center study. Br J Ophthalmol 2007;91:757–760.
- Martinez A, Sanchez M. A comparison of the safety and intraocular pressure lowering of bimatoprost/timolol fixed combination versus latanoprost/timolol fixed combination in patients with open-angle glaucoma. Curr Med Res Opin 2007;23:1025–1032.
- Netland PA, Robertson SM, Sullivan EK, et al. Travoprost compared with latanoprost and timolol in patients with open-angle glaucoma or ocular hypertension. Am J Ophthalmol 2001;132:472–484.
- 44. Halpern MT, Covert DW, Robin AL. Projected impact of travoprost versus both timolol and latanoprost on visual field deficit progression and costs among black glaucoma subjects. Trans Am Ophthalmol Soc 2002;100:109–118.
- 45. Cardascia N, Vetrugno M, Trabucco T, Cantatore F, Sborgia C. Effect of travoprost eye drops on intraocular pressure and pulsatile ocular blood flow: a 180-day, randomized, double-masked comparison with latanoprost eye drops in patients with open-angle glaucoma. Curr Ther Res 2003;64:389–400.
- Nordmann JP, Lepen C, Lilliu H, Berdeaux G. Estimating the long-term visual field consequences of average daily intraocular pressure and variance: a clinical trial comparing timolol, latanoprost, and travoprost. Clin Drug Invest 2003; 23:431–438.
- DuBiner HB, Sircy MD, Landry T, et al. Comparison of the diurnal ocular hypotensive efficacy of travoprost and latanoprost over a 44-hour period in patients with elevated intraocular pressure. Clin Ther 2004;26:84–91.
- 48. Denis P, Launois R, Devaux M, Berdeaux G. Comparison of diurnal intraocular pressure control by latanoprost versus travoprost: results of an observational survey. Clin Drug Invest 2006;26:703–714.
- García-Feijoo J, Martínez-de-la-Casa JM, Castillo A, Méndez C, Fernández-Vidal A, García-Sánchez J. Circadian IOP-lowering efficacy of travoprost 0.004% ophthalmic solution compared to latanoprost 0.005%. Curr Med Res Opin 2006; 22:1689–1697.
- 50. Franks WA, Renard JP, Cunliffe IA, Rojanapongpun P. A 6-week, double-masked, parallel-group study of the efficacy and safety of travoprost 0.004% compared with latanoprost 0:005%/timolol 0.5% in patients with primary open-angle glaucoma or ocular hypertension. Clin Ther 2006;28:332–339.
- Schmier J, Halpern M, Covert D, Robin A. Travoprost versus latanoprost combinations in glaucoma: economic evaluation based on visual field deficit progression: Curr Med Res Opin 2006;22:1737–1743.
- Denis P, Lafuma A, Khoshnood B, Mimaud V, Berdeaux G. A meta-analysis of topical prostaglandin analogues intraocular pressure lowering in glaucoma therapy. Curr Med Res Opin 2007;23:601–608.
- Parrisk RK, Palmberg P, Sheu WP, XLT Study Group. A comparison of latanoprost, bimatoprost, and travoprost in patients with elevated intraocular pressure: a 12-week, randomized, masked-evaluator multicenter study. Am J Ophthalmol 2003;135:688–703.
- 54. Inan U, Ermis S, Orman A, et al. The comparative cardiovascular, pulmonary, ocular blood flow, and ocular



DOCKET

Explore Litigation Insights



Docket Alarm provides insights to develop a more informed litigation strategy and the peace of mind of knowing you're on top of things.

Real-Time Litigation Alerts



Keep your litigation team up-to-date with **real-time** alerts and advanced team management tools built for the enterprise, all while greatly reducing PACER spend.

Our comprehensive service means we can handle Federal, State, and Administrative courts across the country.

Advanced Docket Research



With over 230 million records, Docket Alarm's cloud-native docket research platform finds what other services can't. Coverage includes Federal, State, plus PTAB, TTAB, ITC and NLRB decisions, all in one place.

Identify arguments that have been successful in the past with full text, pinpoint searching. Link to case law cited within any court document via Fastcase.

Analytics At Your Fingertips



Learn what happened the last time a particular judge, opposing counsel or company faced cases similar to yours.

Advanced out-of-the-box PTAB and TTAB analytics are always at your fingertips.

API

Docket Alarm offers a powerful API (application programming interface) to developers that want to integrate case filings into their apps.

LAW FIRMS

Build custom dashboards for your attorneys and clients with live data direct from the court.

Automate many repetitive legal tasks like conflict checks, document management, and marketing.

FINANCIAL INSTITUTIONS

Litigation and bankruptcy checks for companies and debtors.

E-DISCOVERY AND LEGAL VENDORS

Sync your system to PACER to automate legal marketing.

