

**FDLI'S** FOOD  
*and* DRUG  
POLICY FORUM

In the Aftermath of *Parens Patriae*:  
Can Private Copycats Still Sue?

**JAMES M. BECK**  
Counsel  
Reed Smith LLP

VOLUME 3, ISSUE 8 // APRIL 24, 2013

THE FOOD AND DRUG LAW INSTITUTE  
1155 15TH STREET NW, SUITE 800 // WASHINGTON, DC 20005  
[www.fdi.org](http://www.fdi.org)



## INFORMATION FOR SUBSCRIBERS AND PURCHASERS

### License Agreement (the “Agreement”) and Terms of Use for End Users of FDLI Digital Publication Product Services (the “Services”)

THIS IS AN AGREEMENT BETWEEN YOU, (THE “END USER”), AND THE FOOD AND DRUG LAW INSTITUTE (“FDLI”). FDLI IS THE PROVIDER OF THE SERVICES THAT PERMIT END USERS, (LIMITED TO FDLI MEMBERS OR NONMEMBER SUBSCRIBERS OR PURCHASERS OR OTHERS AS DETERMINED BY FDLI) TO LICENSE DIGITAL PUBLICATION PRODUCTS (THE “DIGITAL PUBLICATION PRODUCTS”) FOR END USER USE ONLY UNDER THE TERMS AND CONDITIONS SET FORTH IN THIS AGREEMENT. PLEASE READ THIS LICENSE AGREEMENT AND TERMS OF USE, AND ALL RULES AND POLICIES FOR THE SERVICES (INCLUDING, BUT NOT LIMITED TO, ANY RULES OR USAGE PROVISIONS SPECIFIED ON THE FDLI WEBSITE) BEFORE USING THE PRODUCTS. BY USING THE PRODUCTS, YOU AGREE TO BE BOUND BY THE TERMS OF THIS AGREEMENT.

### Digital Publication Products

**FDLI website:** The FDLI website enables the End User to download this Digital Publication Product to a personal computer or personal handheld device solely for personal use.

**Use of Digital Publication Products:** Upon your payment of the applicable fees, FDLI grants you the non-exclusive right to retain a permanent copy of the applicable Digital Publication Product and to view, print and use such Digital Publication Product an unlimited number of times, solely for your personal, non-commercial use.

**Restrictions:** The End User agrees that Digital Publication Products contain proprietary material that is owned by FDLI, and is protected by United States copyright laws. For reprint permissions or distribution inquiries, contact FDLI at (202) 371-1420.

For subscription or purchasing information, visit [www.fdpi.org](http://www.fdpi.org).

### Disclaimer

The Food and Drug Law Institute, founded in 1949, is a non-profit organization that provides a marketplace for discussing food and drug law issues through conferences, publications and member interaction.

The views, opinions and statements expressed in this article are those of the author(s). The Food and Drug Law Institute neither contributes to nor endorses Forum articles. *As a not-for-profit 501(c)(3) organization, FDLI does not engage in advocacy activities.*

©2013 FDLI

All rights reserved. ISSN pending.

Authorization to photocopy items for internal or personal use of specific clients is granted by the Food and Drug Law Institute, provided that the base fee of US \$.75 per page is paid directly to the Copyright Clearance Center (CCC), 222 Rosewood Drive, Danvers, MA 01923, USA. For those organizations that have been granted a photocopy license by CCC, a separate system of payment has been arranged. The fee code for users of the Transactional Reporting Service is: ISSN pending 02.75.

To order additional copies of this publication, please visit our website at [www.fdpi.org](http://www.fdpi.org).



1155 15th Street NW, Ste. 800, Washington, D.C. 20005

Tel: (202) 371-1420; Fax: (202) 371-0649

email: [comments@fdpi.org](mailto:comments@fdpi.org)

website: [www.fdpi.org](http://www.fdpi.org)

[www.fdpi.org](http://www.fdpi.org)

## FDLI'S FOOD AND DRUG POLICY FORUM

**Michael D. Levin-Epstein, J.D., M.Ed.**  
Editor-in-Chief

**Davina Rosen Marano, Esq.**  
Editor

---

## FDLI'S FOOD AND DRUG POLICY FORUM

**Joseph L. Fink III (Chair)**  
University of Kentucky

**Barbara A. Binzak Blumenfeld (Board Liaison)**  
Buchanan Ingersoll & Rooney, PC

**Sheila D. Walcoff (Vice Chair)**  
Goldbug Strategies, LLC

Gary C. Messplay  
Hunton & Williams, LLP

Christina Anderson Mooney  
Medtronic, Inc.

Peter Pitts  
Center for Medicine in the Public Interest

Peggy Armstrong  
International Dairy Foods Association

Mark Pollack  
Personal Care Products Council

Brendan Benner  
Medical Device Manufacturers Association

Lori M. Reilly  
PhRMA

Sandra B. Eskin  
The Pew Charitable Trusts

Robert Rosado  
Food Marketing Institute

Eric Feldman  
University of Pennsylvania

Timothy W. Schmidt  
Johnson Controls

Paul A. Franz  
The Procter & Gamble Company

David C. Spangler  
Consumer Healthcare Products Association

Robert L. Guenther  
United Fresh Produce Association

William Vodra  
Arnold & Porter, LLP

Mary Clare Kimber  
Plasma Protein Therapeutics Association

Pamela Wilger  
Cargill, Inc.

Patricia A. Maloney  
Quest Diagnostics

Lisa Ann Zoks  
Drug Information Association

## TABLE OF CONTENTS

I.	Introduction & Background.....	1
II.	Research & Response .....	1
	A. Defining <i>Parens Patriae</i> .....	1
	B. Application to <i>Curtis v. Altria Group, Inc.</i> .....	2
	C. Other Cases Binding Subsequent Private Plaintiffs .....	3
III.	Policy Implications .....	3
IV.	Conclusion.....	4
	About The Authors.....	6
	About the Food and Drug Policy Forum .....	6
	About FDLI .....	6

# In the Aftermath of *Parens Patriae*: Can Private Copycats Still Sue?

## I. INTRODUCTION & BACKGROUND

Pharmaceutical manufacturers and other state contractors are increasingly subject to suits brought by state attorneys general (AG). Such suits may claim a variety of alleged misdeeds, but frequently concern scientific statements about the effectiveness of off-label uses, regardless of the scientific accuracy of the information.<sup>1</sup> Because state attorneys general act on behalf of the population of an entire state, their litigation inherently carries with it the coercive potential of large damages. Thus, the pressure to settle such actions can be extreme.

A recent decision, however, demonstrates that there can be a silver lining, at least potentially, around at least some of those big, dark clouds of state attorney general litigation. Companies considering the settlement of such actions should be cognizant of the recent Minnesota Supreme Court decision, *Curtis v. Altria Group, Inc.*<sup>2</sup> The legal principles underlying the *Curtis* decision, under analogous facts, should be applicable to all attorney general actions, with respect to both the state litigation and subsequent private copycat actions, no matter the product or other allegations upon which such suits might be based.<sup>3</sup>

### POLICY RECOMMENDATIONS:

Since state attorneys general act on behalf of the public in the capacity of "*parens patriae*," their monetary settlements using general release language should also bind individual copycat plaintiffs, thus precluding relitigation of those claims.

1

## II. RESEARCH & RESPONSE

### A. Defining *Parens Patriae*

The relevant legal rule recognized in *Curtis* is that, since a state attorney general acts on behalf of the citizens of that state—that is to say in the capacity of "*parens patriae*"—any resolution (which usually means a settlement of some sort) of a state attorney general action will preclude private copycat plaintiffs from later suing the same defendant over the same alleged wrongdoing. *Parens patriae* actions are one of several methods of aggregating litigation.<sup>4</sup> In this type of action, a "[g]overnmental actor" has "authority to speak for citizens on matters of public concern,"<sup>5</sup> and in many ways resembles class actions—including their preclusive effect on subsequent actions by individual citizens. As discussed in the ALI's principles:

Where the interest to be protected is one held by members of the public at large, an action by a public official in behalf of that interest may be held

preemptive of private remedies and preclusive effects accordingly given to a judgment in an action involving the official.... The existence of such an interest is clearest when a government or public official sues *parens patriae*.

\*\*\*\*

The right to sue *parens patriae* being established..., it remained to be considered whether a judgment in a *parens patriae* action precludes a subsequent civil suit by a citizen brought to vindicate the same public interest. The U.S. Supreme Court answered affirmatively.... In principle, *parens patriae* actions can preclude large numbers of individuals from suing. In this respect, they resemble class actions.<sup>6</sup>

## B. Application to *Curtis v. Altria Group, Inc.*

The above is precisely what happened in *Curtis*.<sup>7</sup> In 1994, the Minnesota state AG brought a consumer fraud action against the defendants over alleged false advertising pertaining to “light” cigarettes, seeking among other things, restitution and damages.<sup>8</sup> This litigation action settled a few years later for over \$100 million.<sup>9</sup> Several years later, copycat Minnesota private plaintiffs brought duplicative suits asserting violations of the same consumer protection statutes, alleging the same purportedly violative conduct. They also sought the same recovery—restitution and damages.<sup>10</sup>

The trial court granted partial summary judgment on the ground that the copycat action was barred by the release that settled the prior attorney general action.<sup>11</sup> The copycat plaintiffs appealed, and the Minnesota intermediate appellate court reversed, finding the release to be inapplicable, notwithstanding the repetitive nature of the litigation.<sup>12</sup>

The Minnesota Supreme Court took the case and reversed, reinstating summary judgment. The basis for reversal, well-stated in the Supreme Court’s syllabus, was the *res judicata* effect of the state attorney general’s litigation and ultimate settlement of prior litigation brought in a *parens patriae* capacity:

Under [the Minnesota act], the Minnesota Attorney General (State AG) has the authority to bring a lawsuit...and to seek not only the relief available to the State..., but also the relief available to a private litigant.... It logically follows that **the State AG has the authority to settle and release a private litigant’s claims.**

The 1998 Settlement Agreement entered into by the State AG and [defendant] **expressly released** and barred [copycat plaintiffs’] consumer protection claims...and **is binding on [those plaintiffs].**<sup>13</sup>

2

### C. Other Cases Binding Subsequent Private Plaintiffs

Curtis is only the latest example of the *res judicata* effect of *parens patrie* actions binding subsequent private plaintiffs. The Supreme Court ruled, in a dispute over riparian rights, that a final judgment against the state “was effective, not only against [it], but also against its citizens...for they, in their common public rights as citizens...were represented by the State in those proceedings, and, like it, were bound.”<sup>14</sup> The *res judicata* effect of *parens patriae* actions arises particularly frequently in tobacco actions, since cigarettes were the first product to be targeted in attorney general actions. Thus, in Georgia, an action to recover punitive damages was barred by the *res judicata* effect of a consent judgment with the state’s attorney general.

Because punitive damages serve a public interest and are intended to protect the general public, as opposed to benefitting or rewarding particular private parties, we find the State, in seeking punitive damages in the suit against [defendant], did so as *parens patriae* and in this capacity represented the interests of all [of the state’s] citizens, including plaintiffs here. Accordingly, we conclude that the State and plaintiffs were privies in that action.<sup>15</sup>

### III. POLICY IMPLICATIONS

Attorneys general have now focused their *parens patriae* powers squarely on manufacturers of prescription drugs and medical devices. Thus, the precedents from the tobacco and environmental fields now are equally applicable to drugs and medical devices. To the extent that such manufacturers have the misfortune to be sued by one or more state attorneys general, and they elect to settle such actions, those settlements should likewise preclude copycat private litigation involving the same allegations. Particularly, since state attorneys generals can assert the public’s consumer protection rights, the court in Curtis concluded that private consumer protection litigation was merely “part of the broader authority of the State AG to bring a lawsuit...to enforce all remedies available to it,” including those remedies also provided to private litigants.<sup>16</sup> “[I]t logically follows” that since the attorney general can pursue private as well as public remedies, that office “has authority to settle and release” those claims, including those of subsequent private litigants.<sup>17</sup>

That “release,” in Curtis, moreover, was a broad, general one – including “all claims that the State of Minnesota made, or could have made.”<sup>18</sup> The release had the usual “broad and comprehensive” provisions that are ordinarily included in general releases: “any and all manner;” “known or unknown, suspected or unsuspected, accrued or unaccrued, whether legal, equitable or statutory;” “relating to the subject matter;” “directly or indirectly based on, arising out of or in any way related to.”<sup>19</sup> The court in Curtis accorded the general release language its ordinary broad scope. The copycat plaintiffs’ claims easily satisfied the “related to” test since they “assert[ed] violation of the same consumer protection statutes arising from the same fraudulent and deceptive misrepresentations.”<sup>20</sup>

The copycat litigants’ last stand in Curtis was a provision in the settlement providing that “no portion...shall bind any non-party.”<sup>21</sup> Because the state attorney general was acting on behalf of the public, members of the public could not be heard to claim they were “nonparties”:

We read the words “representatively” and “derivatively” to encompass [private copycats’] right as private litigants to bring [monetary] claims against [defendant]. Because the State AG brought and released those claims, including [monetary] consumer protection claims that could have been brought on behalf of private litigants, the release expressly determined [private copycats’] right to bring a [monetary] consumer protection claim.<sup>22</sup>

#### IV. CONCLUSION

Since prescription drug and medical device manufacturers are increasingly finding themselves on the receiving end of *parens patriae* litigation—both state attorney general actions and follow-up copycat litigation—Curtis provides reason to believe that the same silver lining to the dark cloud of such litigation should be equally available against copycat drug/device plaintiffs. The relevant state law would have to allow the state attorney to recover the same consumer protection damages as private litigants, but once that happens, the rest should flow. Since attorneys generals are acting on behalf of the public, their monetary settlements using general release language should also bind individual copycat plaintiffs and thus preclude relitigation of those claims.

#### ENDNOTES

1. *E.g., Caldwell v. Janssen Pharmaceuticals, Inc.*, 100 So. 3d 865 (La. App. 2012); *In re Zyprexa Products Liability Litigation*, 671 F. Supp.2d 397 (E.D.N.Y. 2009).
2. 813 N.W.2d 891 (Minn. 2012).
3. *Id.*
4. *See* Principles of the Law of Aggregate Litigation §1.02 & Reporters notes to comment b(1)(B) (ALI 2010).
5. *Id.*
6. *Id.* Reporters’ Notes at pp. 20-21.
7. *See supra*, note 1.
8. *See id.* at 896-97.
9. *See id.* at 897.
10. *See id.*
11. *Id.*
12. *Id.* at 898.



13. *Curtis*, 813 N.W.2d at 894, syllabus points 1, 3 (emphasis added) (syllabus point 2 dealt with the reverse situation, and thus was not relevant to the *res judicata* issue).
14. *City of Tacoma v. Taxpayers of Tacoma*, 357 U.S. 320, 341 (1958).
15. *Brown & Williamson Tobacco Corp. v. Gault*, 627 S.E.2d 549, 552 (Ga. 2006); accord *Fabiano v. Philip Morris Inc.*, 862 N.Y.S.2d 487, 490 (N.Y.A.D. 2008) (*parens patriae* action *res judicata* as to punitive damages because “punitive damages claims are quintessentially and exclusively public in their ultimate orientation and purpose”). See also *Alaska Sport Fishing Ass’n v. Exxon Corp.*, 34 F.3d 769, 773 (9th Cir. 1994) (“governments may act in their *parens patriae* capacity as representatives for all their citizens in a suit to recover damages for injury to a sovereign interest”); *Satsky v. Paramount Communications, Inc.*, 7 F.3d 1464, 1470 (10th Cir. 1993) (“[w]hen a state litigates common public rights, the citizens of that state are represented in such litigation by the state and are bound by the judgment”); *Badgley v. City of New York*, 606 F.2d 358, 364 (2d Cir.1979) (private citizens’ riparian rights were “conclusively determined by the terms of the [state consent d]ecree” because the rights of private “citizens cannot exceed those of [the state] itself”); *EPA v. City of Green Forest, Arkansas*, 921 F.2d 1394, 1401 (11th Cir. 1990) (pollution plaintiffs bound *parens patriae* settlement even though the governmental action was subsequently filed).
16. See *supra*, note 1 at 899.
17. *Id.* at 900.
18. *Id.* at 902.
19. *Id.*
20. *Id.* at 903 (emphasis original).
21. *Id.* at 904.
22. *Id.*

## ABOUT THE AUTHORS

**James M. Beck** is counsel to Reed Smith in Philadelphia in the Life Sciences Health Industries Group, handling primarily multi-district product liability litigation and other complex mass torts. Mr. Beck founded the Drug and Device Law Blog, a five-time ABA Top 100 Blog. For a decade he has edited the ABA newsletter “Mass Torts.” Mr. Beck sits on the national case selection committee for the Product Liability Advisory Committee (PLAC) and recently won PLAC’s John P. Raleigh Award, its highest honor. Mr. Beck’s 1997 Food & Drug Law Journal off-label use article was cited twice by the United States Supreme Court.

## ABOUT THE FOOD AND DRUG POLICY FORUM

FDLI’s Food and Drug Policy Forum provides a marketplace for the exchange of policy ideas regarding food and drug law issues. The Forum welcomes articles on cutting-edge state, national and international policy issues related to food and drug law.

FDLI’s Food and Drug Policy Forum is designed to provide a venue for the presentation of information, analysis and policy recommendations in these areas food, drugs, animal drugs, biologics, cosmetics, diagnostics, dietary supplements, medical devices and tobacco.

Each issue of the Forum presents an important policy topic in the form of a question, provides background information and detailed discussion of the issues involved in the policy question, relevant research, pertinent sources and policy recommendations. This publication is digital-only, peer-reviewed and smartphone enabled.

The Forum is published biweekly (24 times a year) and is provided as a complimentary benefit to FDLI members, and by subscription to members of associations on the Forum Editorial Advisory Board and non-members. Individual issues of the Forum are also available for separate purchase.

The 24-member Food and Drug Policy Forum Editorial Advisory Board, comprised of eight representatives of leading associations interested in food and drug law issues and 16 food and drug and healthcare professionals, provides peer review and guidance on articles considered for publication in the Forum.

## ABOUT FDLI

The Food and Drug Law Institute, founded in 1949, is a non-profit organization that provides a marketplace for discussing food and drug law issues through conferences, publications and member interaction. FDLI’s scope includes food, drugs, animal drugs, biologics, cosmetics, diagnostics, dietary supplements, medical devices and tobacco. As a not-for-profit 501(c)(3) organization, FDLI does not engage in advocacy activities.

6

FDLI's Mission is to provide education, training, and publications on food and drug law; act as a liaison to promote networking as a means to develop professional relationships and idea generation; and ensure an open, balanced marketplace of ideas to inform innovative public policy, law, and regulation.

In addition to the Forum, FDLI publishes the quarterly, peer-reviewed Food and Drug Law Journal presenting in-depth scholarly analysis of food and drug law developments; Update magazine, which provides members with concise analytical articles on cutting-edge food and drug issues; the FDLI Monograph Series, an annual six-publication set of practical guides on contemporary food and drug law topics, and numerous comprehensive new books each year.