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### **Analysis of conflicts of interest in pharmaceutical payments made to Food and Drug Administration physician advisers after dermatologic drug approval**



*To the Editor:* Academic physicians have regularly served as voting members of advisory committees for the United States Food and Drug Administration (FDA).<sup>1-3</sup> These committee members serve as external experts to the FDA in determining whether a new medical therapy is fit for the United States market.<sup>2,3</sup> Considering that financial conflicts of interest have been shown to impact member voting habits, physician financial ties have been a topic of ongoing discussion.<sup>2-5</sup> A recent study found that many former FDA advisers receive large sums of money from pharmaceutical companies once their advisory role is complete.<sup>3</sup> Some fear these relationships may incentivize future committee

members to expect after-the-fact rewards for favorable voting habits.<sup>3</sup> The FDA has guidelines to minimize financial conflicts of interest but does not make stipulations about postadvisory role financial relationships.<sup>3</sup>

Investigations using publicly available Open Payment data from the Centers for Medicare and Medicaid Services database have analyzed post hoc advisory role payments to physician advisers in the approval of psychopharmacologic, rheumatologic, cardiac, and renal drugs.<sup>2</sup> To our knowledge, these analyses have not been reported for dermatologic drug advisers. The purpose of this study was to review Open Payment data for industry payments made to former dermatologic drug committee members.

Payments made to United States physicians who advised FDA committees during the approval of 8 dermatologic therapies were analyzed using the Open Payment database. Drugs were chosen from recently approved dermatology medications from [CenterWatch.com](http://CenterWatch.com) and included those used frequently and infrequently by dermatologists: brodalumab, dalbavancin, deoxycholic acid, dupilumab, peginterferon alfa-2b, secukinumab, tedizolid phosphate, and ustekinumab.<sup>3</sup> Data were collected from 2013 to 2017. This range was selected because no data are available before 2013, and 2017 correlated with the most recent data available.<sup>4</sup> Payments classified as general payments were recorded and included consulting fees, speaking fees, educational expenses, food and beverage expenses, and travel and lodging expenses.

Of the 61 physician advisers, 33 (54%) received at least 1 industry payment after dermatologic drug approval. Of the 33 receiving 1 payment, 9 (27%) accepted more than \$1000, 6 (18%) accepted more than \$10,000, 5 (15%) accepted more than \$50,000, and 3 (9%) accepted more than \$100,000. The 33 physician advisers received a mean  $\pm$  standard deviation of \$47,860.62  $\pm$  \$85,938. The standard deviation was larger than the mean due to the nonnormal distribution of the data. For the drugs examined, payments from competing pharmaceutical companies outnumbered payments from the drug manufacturer (Table 1).

Our research concludes former FDA committee advisers for dermatologic drugs frequently received payments from industry. Furthermore, a significant proportion (24%) of those who accepted payments received funds in excess of \$50,000. Critics of such industry-physician relationships argue these payments could incentivize FDA advisers to alter their voting habits.<sup>3</sup> Others conclude post hoc industry payments should not be discouraged

**Table I.** Payments made to voting members of Food and Drug Administration drug committees by pharmaceutical companies\*

Drug	Amount, \$
<b>Brodalumab</b>	
Competitor payments	85,249.89
Manufacturer (Valeant Pharmaceuticals) payments	0.00
Total payments	85,249.89
<b>Dalbavancin</b>	
Competitor payments	210,478.76
Manufacturer (Allergan, Inc) payments	2,262.77
Total payments	212,741.53
<b>Deoxycholic acid</b>	
Competitor payments	218,910.73
Manufacturer (Kythera Biopharma) payments	0.00
Total payments	218,910.73
<b>Dupilumab</b>	
Competitor payments	44,226.33
Manufacturer (Sanofi & Regeneron) payments	29,587.09
Total payments	73,813.42
<b>Peginterferon alpha</b>	
Competitor payments	792,090.19
Manufacturer (Merck) payments	71,907.98
Total payments	863,998.17
<b>Secukinumab</b>	
Competitor payments	101,392.93
Manufacturer (Novartis) payments	0.00
Total payments	101,392.93
<b>Tedizolid phosphate</b>	
Competitor payments	212,762.97
Manufacturer (Cubist Pharmaceuticals) payments	0.00
Total payments	212,762.97
<b>Ustekinumab</b>	
Competitor payments	120,788.88
Manufacturer (Janssen Biotech) payments	218.76
Total payments	121,007.64

\*Table excludes payments for research purposes.

because quality researchers are likely to be pursued by industry, and these mergers can lead to scientific advancements that benefit society.<sup>2,3</sup> Regardless of the ongoing debate, awareness of physician–industry financial ties is worthy of continued discussion. Future evaluations may help assess trends in post hoc advisory payments.

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#### Parental support for sun-protection policies in schools: A cross-sectional analysis



*To the Editor:* In 2014, the US Surgeon General issued a “Call to Action” for skin cancer awareness and prevention, recommending a community-wide effort that included providing shade, educating patients in a health care setting, and teaching children in school about sun safety.<sup>1</sup> Arizona was the first to mandate that a sun-safety curriculum be taught in all schools from kindergarten through eighth grade.<sup>1,2</sup> Since then, Washington, Oregon, Utah, Texas, and New York have enacted laws that support sun protection