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CORRECTION

# Correction to: An evaluation of the federal adverse events reporting system data on adverse effects of 5-alpha reductase inhibitors

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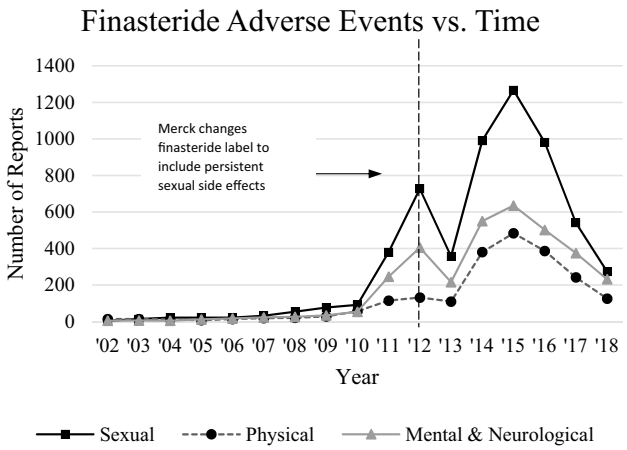
Figure 3a shows the number of adverse event reports for Finasteride monotherapy by year and 2010 as the year of label change by dashed lines. However, the year of label change was incorrectly labeled. The corrected Fig. 3a now reports the year of label change as 2012.




As a result, the Abstract should read as given below:

Case report submissions rose following FDA-mandated finasteride label change.

Accordingly, the Results section should read as below:

After isolating cases of monotherapy, an increase in frequency of case reports for finasteride was seen following the FDA mandated label change.

The Conclusion should read as below:

Reports to FAERS from non-healthcare personnel increased following the FDA-mandated finasteride label change.

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