

# H.R. 3580, Food and Drug Administration Act of 2007

Introduced by John D. Dingell September 19, 2007

This legislation improves current drug safety provisions, encourages greater transparency and timeliness in the drug review process, and reauthorizes the Food and Drug Administration's essential user fee programs. This bill includes compromise language based on H.R. 2900/S. 10 negotiations.

Summary of H.R. 3580

Text of H.R. 3580

Explanation of Differences Between H.R. 2900 and the Food and Drug Administration Act of 2007

"This legislation would significantly improve our postmarket safety programs, thereby preventing many of the drug and device injuries and deaths that occur today." -- Read Chairman Dingell's Floor Statement on H.R. 3580

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