



HR1170

LRB095 20736 GRL 49206 r

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HOUSE RESOLUTION

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WHEREAS, The U.S. Food and Drug Administration (FDA) has  
3 the responsibility to determine if newly developed medical  
4 products are safe and effective; and

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WHEREAS, Any prescription medication, over the counter  
6 medication, medical device, or vaccination must be approved by  
7 the FDA prior to being marketed for general sale in the United  
8 States; and

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WHEREAS, Over the last few years, various widely-used  
prescription drugs have been removed from the market following  
documented deaths and serious harm to patients; and

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WHEREAS, These incidents call into question the standards  
currently used to review drugs and devices in response to the  
FDA Modernization Act of 1997; and

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WHEREAS, Many women have health conditions that require  
them to take medication throughout their lives, such as asthma,  
epilepsy, hypertension, and depression; at this time, while a  
rating system for drug safety does exist, statistics and  
research are lacking regarding the safety of taking these  
medications during pregnancy; and

1           WHEREAS, According to figures kept by birth defect research  
2 groups, the incidence of women using nonprescription medicines  
3 to treat headaches, colds, upset stomachs, and other ailments  
4 typical during pregnancy is on the rise; researchers have found  
5 that over 20% of pregnant women use four or more drugs during  
6 their pregnancy, with over the counter medicines being utilized  
7 most often; and

8           WHEREAS, The American College of Obstetricians and  
9 Gynecologists does not currently have guidelines for  
10 physicians prescribing medications during pregnancy; and

11           WHEREAS, Over 150,000 babies in the U.S. each year are born  
12 with serious birth defects; while some causes of birth defects  
13 are linked to genetic or chromosomal problems, the majority are  
14 caused by unknown reasons; therefore, be it

15           RESOLVED, BY THE HOUSE OF REPRESENTATIVES OF THE  
16 NINETY-FIFTH GENERAL ASSEMBLY OF THE STATE OF ILLINOIS, that we  
17 urge the U.S Food and Drug Administration to seek out better  
18 research and data collection pertaining to drug use during  
19 pregnancy; and be it further

20           RESOLVED, That we also urge the U.S. Food and Drug  
21 Administration, in cooperation with pharmacists and  
22 physicians, to do an informational campaign to provide

1 scientifically accurate information to both pregnant and  
2 lactating women regarding the hazards of taking prescription  
3 and over the counter medications, along with the potential  
4 impact on their child; and be it further

5       RESOLVED, That suitable copies of this resolution be  
6 delivered to the members of the Illinois congressional  
7 delegation, the Illinois State Medical Society, the Illinois  
8 Pharmacists Association, and the Illinois Department of Public  
9 Health.