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INTRODUCED BY Aubrughn Mon

A JOINT RESOLUTION OF THE SENATE AND THE HOUSE OF REPRESENTATIVES OF THE STATE OF
MONTANA URGING CONGRESS TO ENACT LEGISLATION TO REVISE THE PROCESS BY WHICH NEW
DRUGS, BIOLOGICAL PRODUCTS, AND MEDICAL DEVICES ARE APPROVED BY THE UNITED STATES
FOOD AND DRUG ADMINISTRATION.

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9 WHEREAS, improving patient access to quality health care is a paramount national goal; and
 10 WHEREAS, the key to improved health care, especially for persons with serious unmet medical
 11 needs, is the rapid approval of safe and effective new drugs, biological products, and medical devices; and
 12 WHEREAS, minimizing the delay between discovery and eventual approval of a new drug, biological
 13 product, or medical device derived from research conducted by innovative pharmaceutical and
 14 biotechnology companies could improve the lives of millions of Americans; and

WHEREAS, current limitations on the dissemination of information about pharmaceutical products reduce the availability of information to physicians, other health care professionals, and patients and unfairly restrict the right of free speech guaranteed by the first amendment to the United States Constitution; and WHEREAS, the current rules and practices governing the review of new drugs, biological products, and medical devices by the United States Food and Drug Administration can delay approvals and are usually unnecessarily expensive.

21

NOW, THEREFORE, BE IT RESOLVED BY THE SENATE AND THE HOUSE OF REPRESENTATIVES OF THE
 STATE OF MONTANA:

That the Legislature respectfully urge the Congress of the United States to address this important issue by enacting comprehensive legislation to facilitate the rapid review and approval of innovative new drugs, biological products, and medical devices, without compromising patient safety or product effectiveness.

BE IT FURTHER RESOLVED, that copies of this resolution be sent by the Secretary of State to the President of the United States, the Speaker of the United States House of Representatives, the President of the United States Senate, and to each member of the United States Senate and the House of



## 1 Representatives.

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SECOND READING

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House JOINT RESOLUTION NO. 18 aubrugen menn

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House JOINT RESOLUTION NO. 18 authurphin mon INTRODUCED BY

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THIRD READING

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## 1 Representatives.

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LC1158.01 APPROVED BY COM ON PUBLIC HEALTH, WELFARE & SAFETY

House JOINT RESOLUTION NO. 18 Automyster mon INTRODUCED BY

A JOINT RESOLUTION OF THE SENATE AND THE HOUSE OF REPRESENTATIVES OF THE STATE OF
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1 Representatives.

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Legislative Services Division

1 HOUSE JOINT RESOLUTION NO. 18 2 INTRODUCED BY DEBRUYCKER, MESAROS 3 4 A JOINT RESOLUTION OF THE SENATE AND THE HOUSE OF REPRESENTATIVES OF THE STATE OF MONTANA URGING CONGRESS TO ENACT LEGISLATION TO REVISE THE PROCESS BY WHICH NEW 5 6 DRUGS, BIOLOGICAL PRODUCTS, AND MEDICAL DEVICES ARE APPROVED BY THE UNITED STATES 7 FOOD AND DRUG ADMINISTRATION. 8 9 WHEREAS, improving patient access to quality health care is a paramount national goal; and 10 WHEREAS, the key to improved health care, especially for persons with serious unmet medical 11 needs, is the rapid approval of safe and effective new drugs, biological products, and medical devices; and 12 WHEREAS, minimizing the delay between discovery and eventual approval of a new drug, biological 13 product, or medical device derived from research conducted by innovative pharmaceutical and 14 biotechnology companies could improve the lives of millions of Americans; and 15 WHEREAS, current limitations on the dissemination of information about pharmaceutical products reduce the availability of information to physicians, other health care professionals, and patients and unfairly 16 17 restrict the right of free speech guaranteed by the first amendment to the United States Constitution; and 18 WHEREAS, the current rules and practices governing the review of new drugs, biological products, 19 and medical devices by the United States Food and Drug Administration can delay approvals and are usually 20 unnecessarily expensive. 21 22 NOW. THEREFORE, BE IT RESOLVED BY THE SENATE AND THE HOUSE OF REPRESENTATIVES OF THE

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