

RESOLUTION 01.2020-31 (A-20)
COUNTRY OF MEDICATION MANUFACTURE DISCLOSURE

ISMS Board Action Taken on 01/25/2020

Adopted Resolution 01.2020-31 (A-20), as amended by the Board of Trustees:

~~RESOLVED, that the ISMS delegation to the American Medical Association (AMA) submit a resolution requesting that the AMA to introduce and/or support legislation requiring that containers in which each and every medication is dispensed to the public clearly state the country or countries of manufacture of such medication thereby allowing the public to know such locations prior to consuming the contents thereof; and be it further~~

RESOLVED, that the ISMS delegation to the American Medical Association (AMA) submit a resolution requesting that the AMA introduce and/or support legislation requiring the U.S. Food and Drug Administration (FDA) to resume safety testing for all drug manufacturing facilities on a frequent and rigorous basis, as done in the past; and be it further

~~RESOLVED, that the ISMS delegation to the AMA submit a resolution requesting that legislation introduced and/or supported by the AMA further require FDA testing of all medications, imported and produced domestically, for safety on a frequent and rigorous basis; and be it further~~

RESOLVED, that the ISMS delegation to the AMA submit a resolution requesting that the AMA call for the FDA to reaffirm the safety and adequacy of volume in the pipeline of the manufacture of drugs.

Council on Medical Service Recommendation to ISMS Board on 01/25/2020

The Council on Medical Service agreed that contamination of medications and lack of adequate testing by the FDA is an important problem that could be addressed by helping consumers to be informed.

The Council on Medical Services recommends that the ISMS Board of Trustees adopt Resolution 01.2020-31 (A-20), as amended:

38 RESOLVED, that the ISMS delegation to the American Medical Association
39 (AMA) submit a resolution requesting that the AMA to introduce and/or support
40 legislation requiring that containers in which each and every medication is dispensed to
41 the public clearly state the country or countries of manufacture of such medication
42 thereby allowing the public to know such locations prior to consuming the contents
43 thereof; and be it further
44

45 RESOLVED, that the ISMS delegation to the AMA submit a resolution
46 requesting that the AMA introduce and/or support legislation requiring the U.S. Food
47 and Drug Administration (FDA) to resume safety testing for all drug manufacturing
48 facilities on a frequent and rigorous basis, as done in the past; and be it further
49

50 RESOLVED, that the ISMS delegation to the AMA submit a resolution
51 requesting that legislation introduced and/or supported by the AMA further require FDA
52 testing of all medications, imported and produced domestically, for safety on a frequent
53 and rigorous basis; and be it further
54

55 RESOLVED, that the ISMS resolution to the AMA submit a resolution
56 requesting that the AMA call for the FDA to reaffirm the safety and adequacy of volume
57 in the pipeline of the manufacture of drugs.