RECOMMENDATION D:

Madam Speaker, your Reference Committee recommends that the <u>title be changed</u> of Resolution 905 to read as follows:

RESEARCH AND TRANSPARENCY OF INGREDIENTS

IN MENSTRUAL HYGIENE PRODUCTS

HOD ACTION: Resolution 905 be <u>adopted as</u> <u>amended</u> with a <u>title change</u>.

RESOLVED, that our American Medical Association support more comprehensive research on contaminants in menstrual hygiene products (MHP), including but not limited to tampons, other MHPs, and vaginal wipes, and the absorption of toxins into systemic circulation in an effort to better understand their effects on health (New HOD Policy); and be it further

RESOLVED, that our AMA support regulations and legislation that mandate transparency, disclosure, and accurate labeling of contaminants in menstrual hygiene products. (New HOD Policy)

Your Reference Committee heard testimony that it is important to more fully understand menstrual hygiene product ingredients and their risks. However, there was some concern about preemptively supporting regulation and legislation without peer-reviewed scientific evidence of harms of ingredients to support this work. Therefore, Madam Speaker, your Reference Committee recommends that Resolution 905 be adopted as amended and the title be changed to reflect the policy therein.

(17) RESOLUTION 912 - ASSURING REPRESENTATION OF OLDER AGE ADULTS IN CLINICAL TRIALS

RECOMMENDATION A:

Madam Speaker, your Reference Committee recommends that the first Resolve of Resolution 912 be amended by addition and deletion to read as follows:

RESOLVED, that our American Medical Association specifically advocate for inclusion of older patients (both men and women) by amending H-460.911 as follows:

 H-460.911 Increasing Minority, Female, and other Underrepresented Group Participation in Clinical Research of People Identifying with Minoritized and Marginalized Groups

1. Our American Medical Association advocates that:

a. The Food and Drug Administration (FDA) and National Institutes of Health (NIH) conduct annual surveillance of clinical trials by gender, race, age and ethnicity, including consideration of pediatric and elderly populations, and disability status to determine if proportionate representation of people identifying with minoritized and marginalized groups, including by sex, gender, race, ethnicity, and age, women and minorities including older adults and children if appropriate and disability status is maintained in terms of enrollment and retention. This surveillance effort should be modeled after National Institute of Health guidelines on the inclusion of women and minority populations.

b. The FDA have a page on its web site that details the prevalence of people identifying with minoritized and marginalized groups, including sex, gender, race, ethnicity, and age, minorities and women and older adults including those over age 75 and disability status in its clinical trials and its efforts to increase their enrollment and participation in this research.

c. Resources be provided to community level agencies that work with people identifying with minoritized and marginalized groups, including by sex, gender, race, ethnicity, and age, those minorities, females, older adults including those over age 75 and disability status and other underrepresented groups who are not proportionately represented in clinical trials to address issues of lack of access, distrust, and lack of patient awareness of the benefits of trials in healthcare. These ethnic groups may minorities include Individuals/African Americans, Hispanics or Latino. Asians<mark>ł, Pacific Islanders/Native Hawaiians, <u>Middle</u></mark> Eastern or Northern African, and American Indian or Alaskan Natives Native Americans.

RECOMMENDATION B:

Madam Speaker, your Reference Committee recommends that the second Resolve of Resolution 912 be deleted:

RESOLVED, that our AMA monitor the effectiveness of H-460.911 on an annual basis (Directive to Take Action); and be it further

RECOMMENDATION C:

 Madam Speaker, your Reference Committee recommends that the third Resolve of Resolution 912 be amended by addition and deletion to read as follows:

RESOLVED, that our AMA collaborate with AHRQ, FDA, NIH and other relevant stakeholders interested parties to increase public and physician awareness and education on the topic of inclusivity in clinical trial participation (Directive to Take Action).

RECOMMENDATION D:

Madam Speaker, your Reference Committee recommends that <u>the fourth Resolve</u> of Resolution 912 be deleted:

RESOLVED, that our AMA specifically submit comments to the FDA on current proposed industry guidelines for inclusion of underrepresented populations in clinical trials¹ by September 2025.

RECOMMENDATION E:

Madam Speaker, your Reference Committee recommends that Resolution 912 be <u>adopted as amended</u>.

HOD ACTION: Resolution 912 be <u>adopted as amended</u>.

RESOLVED, that our American Medical Association specifically advocate for inclusion of older patients (both men and women) by amending H-460.911 as follows:

1. Our American Medical Association advocates that:

a. The Food and Drug Administration (FDA) and National Institutes of Health (NIH) conduct annual surveillance of clinical trials by gender, race, <u>age</u> and ethnicity, including consideration of pediatric and elderly populations, to determine if proportionate representation of women and minorities <u>including older adults and children if appropriate</u> is maintained in terms of enrollment and retention. This surveillance effort should be modeled after National Institute of Health guidelines on the inclusion of women and minority populations.

b. The FDA have a page on its web site that details the prevalence of minorities and women <u>and older adults including those over age 75</u> in its clinical trials and its efforts to increase their enrollment and participation in this research.

 c. Resources be provided to community level agencies that work with those minorities, females, <u>older adults including those over age 75</u> and other underrepresented groups who are not proportionately represented in clinical trials to address issues of lack of

access, distrust, and lack of patient awareness of the benefits of trials in healthcare. These minorities include Black Individuals/African Americans, Hispanics, Asians/Pacific Islanders/Native Hawaiians, and Native Americans (Directive to Take Action); and be it further

RESOLVED, that our AMA monitor the effectiveness of H-460.911 on an annual basis (Directive to Take Action); and be it further

RESOLVED, that our AMA collaborate with AHRQ, FDA, NIH and other relevant stakeholders to increase public awareness and education on the topic of inclusivity in clinical trial participation (Directive to Take Action); and be it further

 RESOLVED, that our AMA specifically submit comments to the FDA on current proposed industry guidelines for inclusion of underrepresented populations in clinical trials¹ by September 2025. (Directive to Take Action)

Your Reference Committee heard supportive testimony on this item, with some discussion on amendments to refine implementation. Per AMA policies, your Reference Committee however proposes amendments to update policy towards person-first language, and to make ethnicity categories consistent with recommendations from the Office of Management and Budget. One amendment was proposed to strike an annual report on this issue, as enrollment by age group is disclosed by the National Institutes of Health (here, hyperlink available in online report). Additionally, an amendment was proposed to strike reference to submitting comment on an FDA rule, as that docket has already been closed as of October 2024, however our AMA did submit comment that can be found online (hyperlink available in online report). Therefore, Madam Speaker, your Reference Committee recommends that Resolution 912 be adopted as amended.

(18) RESOLUTION 913 - SEXUALLY TRANSMITTED INFECTIONS ARE ON THE RISE IN THE SENIOR POPULATION

RECOMMENDATION A:

Madam Speaker, your Reference Committee recommends that the first Resolve of Resolution 913 be amended by addition and deletion to read as follows:

RESOLVED, that our American Medical Association advocate and promote the U.S. Preventive Services Task Force (USPSTF) recommendations for STI screening through interested senior older adult advocates such as AARP, specifically targeting chlamydia, gonorrhea, human immunodeficiency virus (HIV), HPV and syphilis, for the senior older adult population who are not regularly screened (Directive to Take Action); and be it further