

By: Senator(s) McDaniel, Sojourner

To: Rules

SENATE CONCURRENT RESOLUTION NO. 539

1 A CONCURRENT RESOLUTION URGING THE UNITED STATES FOOD AND
2 DRUG ADMINISTRATION TO PROTECT AMERICAN CONSUMERS WITH RESPONSIBLE
3 KRATOM REGULATION.

4 WHEREAS, it is the policy of the State of Mississippi and its
5 local governments to ensure the safety of consumers using products
6 to protect and maintain their well-being; and

7 WHEREAS, the United States Department of Health and Human
8 Services (HHS) has formally rescinded the recommendation to
9 classify kratom as a Schedule I substance under the federal
10 Controlled Substances Act because the Food and Drug Administration
11 (FDA) failed to provide the evidence to meet the required criteria
12 for scheduling under the act; and

13 WHEREAS, HHS has found there is a significant risk of
14 immediate adverse health consequences for potentially millions of
15 Americans, including tens of thousands of Mississippi citizens, if
16 kratom were included in Schedule I; and

17 WHEREAS, HHS specifically identified that any ban on kratom
18 would force kratom consumers to switch to highly lethal opioids,



19 including potent and deadly prescription opioids, heroin, and/or
20 fentanyl, risking thousands of deaths from overdoses and
21 infectious diseases associated with IV drug use; and

22 WHEREAS, Mississippi has lost thousands of its citizens to
23 the scourge of the opioid overdose epidemic; and

24 WHEREAS, peer-reviewed published surveys show that among
25 kratom consumers in the United States, one-third use kratom for an
26 energy boost and increased focus, one-third use kratom for its
27 relaxing effects and reduction of anxiety, and one-third use it to
28 manage acute and chronic pain, including as a replacement for
29 highly addictive and potentially deadly opioids; and

30 WHEREAS, peer-reviewed published literature, including from
31 researchers from Johns Hopkins University, have concluded that
32 kratom can relieve opioid withdrawal symptoms among those using
33 kratom to treat opioid dependence, and 35% were free from opioids
34 within a year; and

35 WHEREAS, the National Institute on Drug Abuse (NIDA), HHS,
36 the FDA, and the United States Congress all concur that
37 unregulated adulterated kratom products laced with heroin,
38 morphine, or fentanyl put consumers at significant risk, including
39 death:

40 NOW, THEREFORE, BE IT RESOLVED BY THE SENATE OF THE STATE OF
41 MISSISSIPPI, THE HOUSE OF REPRESENTATIVES CONCURRING THEREIN, That
42 we urge the United States Food and Drug Administration to
43 immediately publish good manufacturing guidelines for kratom



44 manufacturers that will restrict adulteration, ban synthetic
45 enhancement of kratom's natural alkaloid contents to alter the
46 overall alkaloid fraction present in the natural plant, and
47 require appropriate labeling of kratom products to protect
48 consumers.

49 BE IT FURTHER RESOLVED, That this resolution be transmitted
50 to the members of Mississippi's congressional delegation, the
51 Secretary of the Department of Health and Human Services, the
52 Commissioner of the Food and Drug Administration, the President of
53 the United States, and made available to the Capitol Press Corps.

