

HOUSE OF REPRESENTATIVES

KENTUCKY GENERAL ASSEMBLY AMENDMENT FORM  
2026 REGULAR SESSION  
**Unofficial Document**

Amend printed copy of **HB 335**

On page 1, lines 7 to 8, delete "registered as a Class II medical device with the United States Food and Drug Administration" and insert in lieu thereof "authorized by the United States Food and Drug Administration for marketing and distribution"; and

On page 1, line 16, after "device", insert ", according to its labeled instructions for use,".

Amendment No. HFA 1

Rep. Rep. Matthew Lehman

Committee Amendment

Signed:

Floor Amendment

LRC Drafter: \_\_\_\_\_

Adopted: \_\_\_\_\_

Date: \_\_\_\_\_

Rejected: \_\_\_\_\_

Doc. ID: XXXX

**Not for Filing**