

Medical Devices; Cardiovascular Devices; Classification of the Hyperoxia Monitoring Device Adjunct to Pulse Oximetry

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The Food and Drug Administration (FDA) is classifying the hyperoxia monitoring device adjunct to pulse oximetry into class II (special controls). The special controls that apply to the device type are identified in this order and will be part of the codified language for classification of the hyperoxia monitoring device adjunct to pulse oximetry. We are taking this action because we have determined that classifying the device into class II will provide a reasonable assurance of the safety and effectiveness of the device. We believe this action will also enhance patients' access to beneficial innovative devices, in part by reducing regulatory burdens.

Agency: Health and Human Services Department, Food and Drug Administration.

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