NOTICE OF VIRTUAL PUBLIC HEARING

As part of the mandate of the Food and Drug Administration (FDA) to protect public health and safety through regulation of health products including medical devices, the FDA through the Center for Device Regulation, Radiation Health and Research (CDRRHR) is in the process of developing guidelines on good storage and distribution practices for medical devices. Relative to this, the CDRRHR has finalized the draft FDA Circular entitled "Good Storage and Distribution Practices for Medical Devices".

In the interest of service and to seek the comments of the stakeholders that will be affected by the proposed policy, the FDA is inviting representatives from the medical device industry to attend the **Virtual Public Hearing** on **31 May 2022** via Google Meet. Eighty (80) slots have been allotted for those interested to attend the said activity. One (1) representative per company shall be accommodated.

To register and confirm your attendance, kindly accomplish the registration form through this link: https://forms.gle/pvBLkHJbzYrPcnTSA.

Thank you.