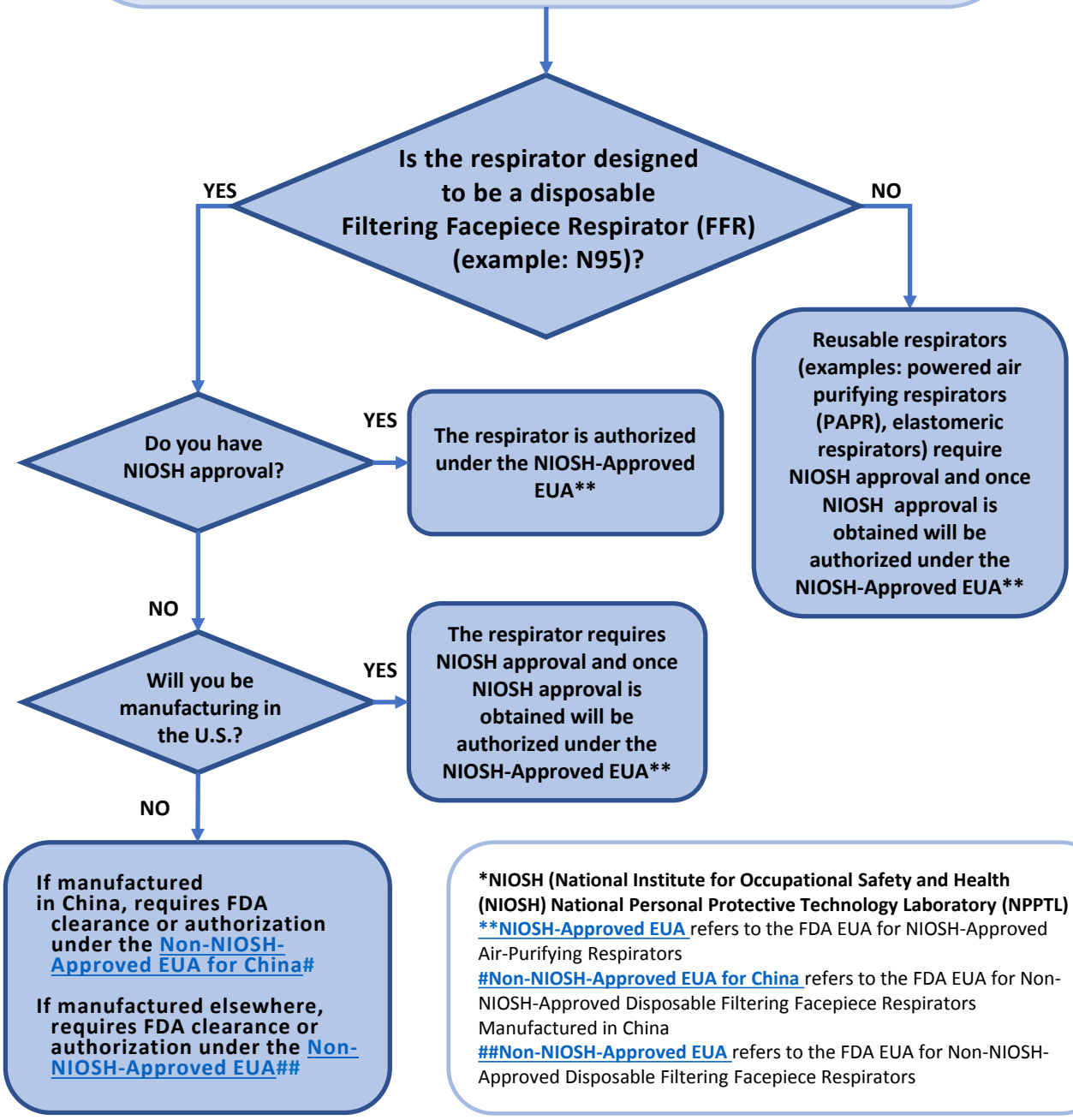
Manufacturing and Distributing Respirators for Health Care Use in the United States Under an Existing Emergency Use Authorization (EUA) During the COVID-19 Pandemic

- To distribute respirators for health care use in the United States, the respirators must be FDA cleared or be authorized under an FDA EUA.
- For information about the NIOSH Respirator Approval Program, consider the information provided by NIOSH NPPTL* at: https://www.cdc.gov/niosh/npptl/RespApprovalInfo.html
- If you are manufacturing outside the United States, see Information for Filing Personal Protective Equipment and Medical Devices During COVID-19



***NIOSH (National Institute for Occupational Safety and Health (NIOSH) National Personal Protective Technology Laboratory (NPPTL)**

**Non-NIOSH-Approved EUA refers to the FDA EUA for NIOSH-Approved Air-Purifying Respirators

#Non-NIOSH-Approved EUA for China refers to the FDA EUA for Non-NIOSH-Approved Disposable Filtering Facepiece Respirators Manufactured in China

##Non-NIOSH-Approved EUA refers to the FDA EUA for Non-NIOSH-Approved Disposable Filtering Facepiece Respirators