An Analysis on Medical Device Recalls and Cybersecurity
Implications on Patient Safety

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AN ANALYSIS OF MEDICAL DEVICE RECALLS

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INTRODUCTION

According to the Food and Drug Administration (FDA) a recall is the removal or correction of medical devices that do not abide by the laws governed by the FDA and threaten patient health and safety [1]. Even though medical devices have become more advanced and are able to connect to networks and other devices, cybersecurity has become an issue while ensuring proper patient care. A recent study examined cybersecurity attacks in medical devices, then characterized the vulnerabilities based on reports from the CVE and ICS-CERT databases [3]. This analysis of FDA reports [2] are the preliminary actions to categorize medical device recalls and eventually connect them to potential cybersecurity vulnerabilities.

METHODS AND DATA

Data was collected from the FDA Medical Device Recall Database from January 2010-March 2020 by entering “software” and “hardware” reasons into the search engine and yielded approximately 1300 results. The total number of software and hardware-related device recalls is shown by year in Figure 1.

The entries from the top ten manufacturers were categorized by utilizing a keyword-based method the manufacturer’s reason for recall. Keywords were manually grouped together into categories, shown in Table 1. The proportions of these categories are shown in Figure 3.

CONCLUSION/FUTURE WORK

● Our preliminary results indicate that there are many types of medical devices and reasons for recall. The manufacturer’s reason for recall is more specific than the FDA’s reason, so patients may not understand the errors in their devices and the impact these errors can have on their safety and privacy due to the report. Therefore, using the manufacturer’s reason may provide more transparency for the patient.

● Develop a Natural Processing Language (NPL) software to categorize current and future recalls into a more manageable database for ease of patient access, while also linking these reasons for recall to the CVE and ICS-CERT classifications of software and cybersecurity malfunctions.

REFERENCES

[1] Center for Devices and Radiological Health, FDA. “Recalls, Corrections and Removals (Devices).” U.S. Food and Drug Administration, FDA,


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