



Adverse Events Reporting Aggregate Reviews

An aggregate review is a type of root cause analysis. It is used by facilities to analyze root cause and develop corrective action plans when there are *multiple occurrences of the same type of event* in the same three month reporting period (quarter). Aggregate review is an effective way for facilities to analyze adverse events in a more comprehensive, system-focused method; to identify common root causes and to identify and implement a comprehensive prevention strategy throughout the facility.

To conduct an aggregate review:

- Continue the usual process to notify the Department of Health (DOH) within 48 hours of confirming an adverse event. Use the [Notification Form](#).
- Maintain a quarterly, chronological log of your confirmed adverse events. Include a short summary of the event, the root cause and casual statement(s).
- When two or more of the same type of adverse event occur within the same quarter, identify common causes/circumstances that contributed to the multiple occurrences.
- Develop a corrective action plan that addresses all components of your overall prevention strategy. This may include care coordination, equipment, documentation, clinical practice/knowledge, special clinical issues or organizational issues. You will need to:
 - Highlight the risk reduction plans for each component.
 - Identify who is responsible for making changes/corrections, a timeline for implementation and your measure of success.
 - Identify the members of your root cause analysis team and describe how you report this work to your leaders.
 - Examples of aggregate reviews:
 - [Unintended retention of surgical sponge](#) (PDF)
 - [Stage 3 or 4 pressure ulcer acquired after admission](#) (PDF)
- Prepare an aggregate review report. Include your chronological log and your corrective action plan. Send your aggregate review report to DOH according to this schedule:

Reporting Quarter	Due date
January 1 – March 31	April 15
April 1 – June 30	July 15
July 1 – September 30	October 15
October 1 – December 31	January 15

- Send your aggregate review report to:

DOH, Adverse Events
Attn: Nicole Fernandus
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