

Standardized Alarm Resolution B. Braun Large Volume Infusion Pumps

Information presented Nov 23-Dec12, 2023



Targets for Manitoba Air in Line (AIL) & Downstream Occlusions (DSO)

AIL: 0.25 alarms per infusion Delivery DSO: 0.7 alarms per infusion Delivery

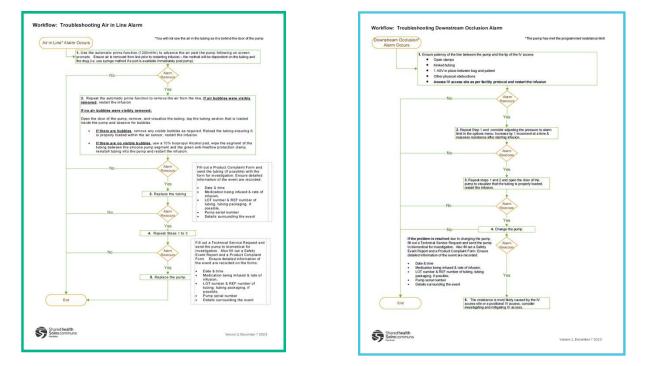
By the end of January, 2024



Resources to Mitigate Air in Line and Downstream Occlusion Alarms

Information on Shared Health MB:

B. Braun Large Volume Pumps & Administration Sets





Air in Line Alarms

If an air in line alarm occurs, we must treat the alarm as if there is air in the line.



Reducing Incidence of Air in line Alarms

To <u>Reduce</u> Air in line alarms we must decrease the amount of air that can be in the infusion line

- Bring cold fluids to room temperature prior to administration to avoid outgassing
- Ensure drip chambers are filled 1/2 to 2/3
- Ensure tubing is properly loaded in the air sensor of the pump
- Prime the IV lines on the pump
- Invert and tap injection sites during priming
- Ensure the ASV is in place at the distal end of the tubing



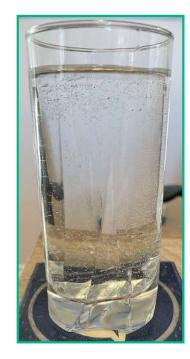




Reducing Incidence of Air in line Alarms

Bring cold fluids to room temperature prior to administration to avoid outgassing



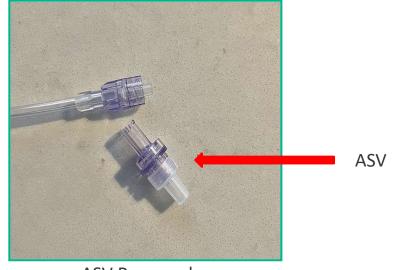


Reducing Incidence of Air in line Alarms

Ensure the ASV is in place at the distal end of the tubing



ASV on Tubing



ASV Removed

Troubleshooting Air in Line (AIL) Alarm



When an air in line alarm occurs You will not see the air in the tubing as it is behind the door of the pump

Use the automatic prime function (1200ml/hr) to remove the air from the line following on screen prompts, then restart the infusion



Step 2 *If the alarm reoccurs*

Repeat the automatic prime function to remove the air from the line



If air bubbles were visibly removed, restart the infusion

If no air bubbles were visibly removed,

Open the door of the pump, remove, and visualize the tubing, tap the tubing section that is loaded inside the pump and observe for bubbles.

- If there are bubbles, remove any visible bubbles as required. Reload the tubing ensuring it is properly loaded within the air sensor, restart the infusion.
- If there are no visible bubbles, use a 70% Isopropyl Alcohol pad, wipe the segment of the tubing between the silicone pump segment and the green anti-freeflow protection clamp, reinstall tubing into the pump and restart the infusion.





Step 3 *If the alarm reoccurs*

Replace the tubing, fill a product complaint form and send the tubing (if possible) with the product complaint form for investigation.

Ensure detailed information of the event are recorded:

- Date & time
- Medication being infused & rate of infusion,
- LOT number & REF number of tubing, tubing packaging, if possible,
- Pump serial number
- Details surrounding the event

Step 4 *If the alarm reoccurs*

Using new tubing after changing it Repeat Steps 1-3









Step 5 *If the alarm reoccurs*

Replace the pump, fill a **product complaint form**, and **send the pump to biomedical for investigation**.

Ensure detailed information of the event are recorded:

- Date & time
- Medication being infused & rate of infusion,
- LOT number & REF number of tubing, tubing packaging, if possible,
- Pump serial number
- Details surrounding the event

Downstream Occlusion Alarms

If the pump indicates there is a downstream occlusion, most likely because the pump has met the resistance threshold to create the alarm.

Causes

Resistance between the pump and the tip of the IV access Viscous fluids Higher rates of infusion

ASSESS > TROUBLESHOOT > RESTART PUMP

Without assessing & troubleshooting, the pump will most likely cause another alarm



Reducing Incidence of Downstream Occlusions

To <u>Reduce</u> downstream occlusion alarms we must decrease the amount of resistance in the infusion line

- Avoid placing IV access at places of flexion (INS Best Practice guidelines)
- If the only option is to insert IV into a flexion area, consider mitigation strategies to avoid flexion at the IV site
- Ensure the fluid never flows through more than 1 ASV
- Ensure all clamps in the fluid path are open
- Ensure patency of the IV access site



Troubleshooting Downstream Occlusion(DSO) Alarms



When a Downstream Occlusion occurs The pump has met the programmed resistance limit

Ensure patency of the line between the pump and the tip of the IV access

Open clamps Unkink any kinked tubing Ensure 1 ASV in place between bag and patient

Other physical obstructions

Assess IV access site as per facility protocol and restart the infusion



If the alarm reoccurs

Repeats Step 1

Consider adjusting pressure to alarm limit in options menu.

Increase by 1 increment at a time & reassess resistance after starting infusion





If the alarm reoccurs

Repeat Steps 1-2

Open door of pump and visualize that the tubing is properly loaded.

Restart the infusion





Step 4 *If the alarm reoccurs*

Change the pump

If the problem is resolved due to changing the pump, fill a **product complaint form** per facility protocol, and send the pump to biomedical for investigation.

Ensure detailed information of the event are recorded

- Date & time
- Medication being infused & rate of infusion •
- Pump serial number
- Details surrounding the event •
- LOT number & REF number of tubing (if possible)



If the alarm reoccurs with a new pump

The resistance is most likely caused by the IV access site or a positional IV access, consider investigating and mitigating IV access.

