

Manufacturer name:	Baxter Healthcare								
About the Recall:	Baxter Healthcare has issued a medical device correction notice for the Life2000 Ventilation system due to the ventilatory may fail to initiate the low gas pressure alarm if the pressure gas source (compressor, oxygen cylinder etc.) is not supplied to the ventilator before initiating therapy. Baxter is currently working on a software update to address the issue.								
Products(s) numbers/models that are affected:	<table border="1" data-bbox="389 819 1510 924"> <thead> <tr> <th data-bbox="393 823 581 861">Product Code</th> <th data-bbox="584 823 896 861">Product Description</th> <th data-bbox="899 823 1221 861">Impacted Devices</th> <th data-bbox="1224 823 1507 861">Device Identifier Number</th> </tr> </thead> <tbody> <tr> <td data-bbox="393 865 581 903">MS-01-0118</td> <td data-bbox="584 865 896 903">Life2000 Ventilator</td> <td data-bbox="899 865 1221 903">All Life2000 Ventilators with software version 06.08.00.00</td> <td data-bbox="1224 865 1507 903">00887761978089 or 00815410020537</td> </tr> </tbody> </table>	Product Code	Product Description	Impacted Devices	Device Identifier Number	MS-01-0118	Life2000 Ventilator	All Life2000 Ventilators with software version 06.08.00.00	00887761978089 or 00815410020537
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URL Link(s) to Manufacturer Notice/Web Site:	www.baxter.com								
Actions to be taken by customer/Patient:	Prior to starting therapy, ensure that the Life2000 ventilator is connected to a pressure gas source first (Life200 compressor, oxygen cylinder, or wall source) and then turn on. Contact your home medical equipment provider if need further instruction.								