

Safety Communication

رسالة سلامة

Risk of not pre-programmed insulin pump

Device/ Product Description:	MiniMed 600 and 700 series insulin pump	
Affected product:	MiniMed™ 640G	MMT-1711, MMT-1712, MMT-1751, MMT-1752
	MiniMed™ 670G	MMT-1780, MMT-1781, MMT-1782, MMT-1760, MMT-1761, MMT-1762, MMT-1741, MMT-1742
	MiniMed™ 720G	MMT-1809, MMT-1810, MMT-1859, MMT-1860
	MiniMed™ 740G	MMT-1811, MMT-1812, MMT-1861, MMT-1862
	MiniMed™ 770G	MMT-1881, MMT-1882, MMT-1892, MMT-1891
	MiniMed™ 780G	MMT-1885, MMT-1886, MMT-1895, MMT-1896
Manufacturer:	Medtronic SA	
Problem:	The new/replacement pump was NOT pre-programmed with the patient's basal rates or other verified settings (i.e., bolus wizard settings, sensor settings, etc.), which must be set up and saved on their pump prior to use.	

Recommendation /Actions:	<ul style="list-style-type: none">- Inform impacted users of the MiniMed™ 600 and 700 series insulin pump using the enclosed letter in the attached FSN. <p>For more information, please click here</p> <p>If you think you had a problem with your device or a device your patient uses, please report the problem to SFDA through: NCMDR Vigilance system (19999) unified call center</p>	
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