The Board of Pharmacy has received notice of the following product correction. The Board strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.

Medtronic has issued a Medical Device Correction notice regarding the items(s)/model number(s) below due to the pump not being pre-programmed with basal rates or other verified settings (i.e., bolus wizard settings, sensor settings, etc.), which must be set up and saved on the pump prior to use. This event is to the consumer level.

MiniMed™ 620G: MMT-1750

MiniMed™ 630G: MMT-1715, MMT-1755, MMT-1754

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-1740, 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-1880, MMT-1881, MMT-1882, MMT-1892, MMT-1891, MMT-1890

MiniMed™ 780G: MMT-1884, MMT-1885, MMT-1886, MMT-1895, MMT-1896