California Device Recall Information Sheet

Food and Drug Branch – Device Recalls

Medtronic MiniMed, Inc. InPen smart insulin pen

Recall Date	Product Description	Recalling Firm	Recall Reason
04/21/2025	InPen smart insulin pen REF: MMT-105NNBLNA, MMT-105NNGYNA, MMT-105NNPKNA, MMT-105ELBLNA, MMT-105ELGYNA, MMT105ELPKNA	Medtronic MiniMed, Inc.	Insulin pens may have been incorrectly assembled therefore users could experience the following: either the insulin cartridge will not fit into the cartridge holder; or the cartridge holder may be difficult to remove from the insulin pen. If the issue is present, and if the user does not have backup insulin they may experience temporary hyperglycemia.

Recall Class	Product Identification	Distribution	Affected Dates
II	REF: MMT-105NNBLNA UDI-DI: 763000827182 Lot: D0020, D0034, D0036, D0038, D0045, D0046, D0050, D0057	259 units in California	March and prior

	REF: MMT-105NNGYNA UDI-DI: 763000827199 Lot: D0026, D0037, D0039, D0041, D0042, D0043, D0049 REF: MMT-105NNPKNA UDI-DI: 763000827205 Lot: D0029, D0030, D0040, D0051, D0062 REF: MMT-105ELBLNA UDI-DI: 763000827151 Lot: D0021, D0033, D0035, D0044, D0059, D0060, D0061 REF: MMT-105ELGYNA UDI-DI: 763000827168 Lot: D0031, D0032, D0047, D0068 REF: MMT105ELPKNA UDI-DI: 763000827175 Lot: D0027, D0028, D0048		

For additional information, please visit the FDA Website.