

CERTIFICATE OF COMPLIANCE

Part # 58794 Lot # A8811

Product Description Secure 1 Leg Gravity Fill Set with Female Screw Connector

Date of Manufacture 2025-06-12

Irradiation Date 08-04-25 Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy Maximum Dose Result (kGy) 30.0 kGy

The above product has been manufactured according to *21 CFR 820 - FDA Quality System Regulations and ISO 13485 Medical Devices - Quality Management System* and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (*ISO 14644 Cleanrooms and associated controlled environments*).

The fluid pathway materials of this product have been tested and qualified to conform with *ISO 10993-1 Biological evaluation of medical devices* for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with *ISO 11137 Sterilization of health care products - Radiation* according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by *ISO 11137 Sterilization of health care products - Radiation*.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with *ANSI/AAMI ST72 "Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing"*. The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record.

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Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 64045 Lot # 64045-A8833

Product Description McKesson 250 mL Empty EVA Container with 2 Ports

Date of Manufacture 2025-06-13 Use by Date 2028-06-13

Irradiation Date 08-04-25 Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy Maximum Dose Result (kGy) 30.0 kGy

The above product, supplied to McKesson Medical-Surgical Inc., has been manufactured according to 21 CFR 820 - FDA Quality System Regulations and ISO 13485 Medical Devices - Quality Management System and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (ISO 14644 Cleanrooms and associated controlled environments).

The fluid pathway materials of this product have been tested and qualified to conform with ISO 10993-1 Biological evaluation of medical devices for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with ISO 11137 Sterilization of health care products - Radiation according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by ISO 11137 Sterilization of health care products - Radiation.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with ANSI/AAMI ST72 Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing. The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record. Materials are traceable and have not changed without prior notice to McKesson Medical-Surgical Inc.


Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 64050

Lot # 64050-A8834

Product Description McKesson 500 mL Empty EVA Container with 2 Ports

Date of Manufacture 2025-06-16

Use by Date 2028-06-16

Irradiation Date 08-04-25

Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy

Maximum Dose Result (kGy) 30.0 kGy

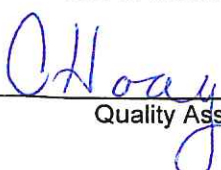
The above product, supplied to McKesson Medical-Surgical Inc., has been manufactured according to 21 CFR 820 - FDA Quality System Regulations and ISO 13485 Medical Devices - Quality Management System and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (ISO 14644 Cleanrooms and associated controlled environments).

The fluid pathway materials of this product have been tested and qualified to conform with ISO 10993-1 Biological evaluation of medical devices for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with ISO 11137 Sterilization of health care products - Radiation according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by ISO 11137 Sterilization of health care products - Radiation.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with ANSI/AAMI ST72 Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing. The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record. Materials are traceable and have not changed without prior notice to McKesson Medical-Surgical Inc.


Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 66041

Lot # 66041-A8807

Product Description Secure 100 mL Empty EVA Container with 2 Ports (Bulk Packed)

Date of Manufacture 2025-06-24

Use by Date 2028-06-24

Irradiation Date 08-04-25

Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy

Maximum Dose Result (kGy) 30.0 kGy

The above product has been manufactured according to 21 CFR 820 - FDA Quality System Regulations, Canadian Medical Devices Regulations Part 1- SOR 98/282, and ISO 13485 Medical Devices - Quality Management System and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (ISO 14644 Cleanrooms and associated controlled environments).

The fluid pathway materials of this product have been tested and qualified to conform with ISO 10993-1 Biological evaluation of medical devices for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with ISO 11137 Sterilization of health care products - Radiation according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by ISO 11137 Sterilization of health care products - Radiation.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with ANSI/AAMI ST72 Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing. The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record.


Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 66045

Lot # 66045-A8809

Product Description Secure 250 mL Empty EVA Container with 2 Ports

Date of Manufacture 2025-06-17

Use by Date 2028-06-17

Irradiation Date 08-04-25

Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy

Maximum Dose Result (kGy) 30.0 kGy

The above product has been manufactured according to 21 CFR 820 - FDA Quality System Regulations, Canadian Medical Devices Regulations Part 1- SOR 98/282, and ISO 13485 Medical Devices - Quality Management System and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (ISO 14644 Cleanrooms and associated controlled environments).

The fluid pathway materials of this product have been tested and qualified to conform with ISO 10993-1 Biological evaluation of medical devices for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with ISO 11137 Sterilization of health care products - Radiation according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by ISO 11137 Sterilization of health care products - Radiation.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with ANSI/AAMI ST72 "Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing". The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

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Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 66100 Lot # 66100-A8808

Product Description Secure 1000 mL Empty EVA Container with Detachable 2 Leg Transfer Set

Date of Manufacture 2025-06-24 Use by Date 2028-06-24

Irradiation Date 08-04-25 Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy Maximum Dose Result (kGy) 30.0 kGy

The above product has been manufactured according to *21 CFR 820 - FDA Quality System Regulations, Canadian Medical Devices Regulations Part 1- SOR 98/282, and ISO 13485 Medical Devices - Quality Management System* and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (*ISO 14644 Cleanrooms and associated controlled environments*).

The fluid pathway materials of this product have been tested and qualified to conform with *ISO 10993-1 Biological evaluation of medical devices* for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

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The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with *ANSI/AAMI ST72 "Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing"*. The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record.


Quality Assurance

8-11-25
Date

CERTIFICATE OF COMPLIANCE

Part # 66120 Lot # 66120-A8835

Product Description Secure 1000 mL Empty EVA Container with Detachable 3 Leg Transfer Set

Date of Manufacture 2025-06-18 Use by Date 2028-06-18

Irradiation Date 08-04-25 Irradiation Work Order No. 3795691

Minimum Dose Result (kGy) 22.8 kGy Maximum Dose Result (kGy) 30.0 kGy

The above product has been manufactured according to 21 CFR 820 - FDA Quality System Regulations, Canadian Medical Devices Regulations Part 1- SOR 98/282, and ISO 13485 Medical Devices - Quality Management System and complies with the relevant requirements. The product has been manufactured and assembled in accordance with internal specifications, with component traceability and lot number control in an ISO Class 8 cleanroom (ISO 14644 Cleanrooms and associated controlled environments).

The fluid pathway materials of this product have been tested and qualified to conform with ISO 10993-1 Biological evaluation of medical devices for biocompatibility test requirements. Incoming materials are inspected upon arrival to confirm compliance to specification and are assigned control numbers for traceability.

The product has been processed with Gamma Radiation to achieve a 10^{-6} sterility assurance level (SAL) in accordance with ISO 11137 Sterilization of health care products - Radiation according to a validated gamma irradiation procedure to yield a sterile fluid path. The certificate of processing is on file and has been reviewed to verify the minimum and maximum doses meet the validated dose range requirements. Sterilization dose audit and a determination of bioburden is completed quarterly on representative product to monitor the sterilization dose effectiveness as required by ISO 11137 Sterilization of health care products - Radiation.

The pyrogenicity of this product's fluid pathway has been tested for bacterial endotoxin content and meets the endotoxin criteria in accordance with established manufacturing operating procedures. The test method is in compliance with ANSI/AAMI ST72 "Bacterial endotoxins - Test methods, routine monitoring, and alternatives to batch testing". The laboratory test report is on file and has been reviewed to verify the test result is less than or equal to 20 EU/device (USP <161>).

Product testing and inspections have been completed to ensure conformity to the specifications and records are maintained on file as part of the device history record.


Quality Assurance

8-11-25
Date