Surface disinfection device	Whitebox
EC Declaration of Conformity	**Rev. 03** **Date 2023-03-20**

The EC declaration of conformity is issued under the sole responsibility of the manufacturer Whitebox AB.

Whitebox with registered office at Dahlbergsgatan 8, 931 33 Skelleftea, Sweden and manufacturer for device defined below, declares the following:

Manufacturer	Whitebox AB: SE 559232-3371
SRN (single registration number)	SE-MF-000032348
Product name/article number	Whitebox Microbox article number: 016-501
Basic UDI number	7350136350018P
EU-MDR	https://ec.europa.eu/tools/eudamed/#/screen/search-device/ccadddf3-4dba-47bc-8005-80d2ebae8ffc
Classification rules (Annex VIII Regulations (EU) 2017/45	Medical device of risk Class 1
Following EU directives	EN 55014–1: 2017+A1:2020, EN/IEC 61003-3-2: 2019, EN 62233 (EMF) EN/IC 60 335-1-2-65

The device complies with the requirements of the European Medical Device Regulation MDR 2017/45. The device does not have any measuring function or used as a sterilization unit.

Skelleftea 2023-03-20

Peter Karlsten CEO Whitebox AB

CE

Surface disinfection device	Whitebox
EC Declaration of Conformity	**Rev. 03** **Date 2023-03-20**

The EC declaration of conformity is issued under the sole responsibility of the manufacturer Whitebox AB.

Whitebox with registered office at Dahlbergsgatan 8, 931 33 Skelleftea, Sweden and manufacturer for device defined below, declares the following:

Manufacturer	Whitebox AB: SE 559232-3371
SRN (single registration number)	SE-MF-000032348
Product name/article number	Whitebox L-Box article number: 009-501
Basic UDI number	7350136350038T
EU-MDR	https://ec.europa.eu/tools/eudamed/#/screen/search-device/ccadddf3-4dba-47bc-8005-80d2ebae8ffc
Classification rules (Annex VIII Regulations (EU) 2017/45	Medical device of risk Class 1
Following EU directives	EN 55014–1: 2017+A1:2020, EN/IEC 61003-3-2: 2019, EN 62233 (EMF) EN/IC 60 335-1-2-65

The device complies with the requirements of the European Medical Device Regulation MDR 2017/45. The device does not have any measuring function or used as a sterilization unit.

Skelleftea 2023-03-20

Peter Karlsten CEO Whitebox AB

CE

Surface disinfection device	Whitebox
EC Declaration of Conformity	**Rev. 03** **Date 2023-03-20**

The EC declaration of conformity is issued under the sole responsibility of the manufacturer Whitebox AB.

Whitebox with registered office at Dahlbergsgatan 8, 931 33 Skelleftea, Sweden and manufacturer for device defined below, declares the following:

Manufacturer	Whitebox AB: SE 559232-3371
SRN (single registration number)	SE-MF-000032348
Product name/article number	Whitebox Minibox article number: 015-501
Basic UDI number	7350136350028R
EU-MDR	https://ec.europa.eu/tools/eudamed/#/screen/search-device/ccadddf3-4dba-47bc-8005-80d2ebae8ffc
Classification rules (Annex VIII Regulations (EU) 2017/45	Medical device of risk Class 1
Following EU directives	EN 55014–1: 2017+A1:2020, EN/IEC 61003-3-2: 2019, EN 62233 (EMF) EN/IC 60 335-1-2-65

The device complies with the requirements of the European Medical Device Regulation MDR 2017/45. The device does not have any measuring function or used as a sterilization unit.

Skelleftea 2023-03-20

Peter Karlsten CEO Whitebox AB

CE

Air disinfection device	Whitebox
EC Declaration of Conformity	**Rev. 03** **Date 2023-03-20**

The EC declaration of conformity is issued under the sole responsibility of the manufacturer Whitebox AB.

Whitebox with registered office at Dahlbergsgatan 8, 931 33 Skelleftea, Sweden and manufacturer for device defined below, declares the following:

Manufacturer	Whitebox AB: SE 559232-3371
SRN (single registration number)	SE-MF-000032348
Product name/article number	Whitebox Air Cleaner with article number: 010-501
Basic UDI number	7350136350048V
EU-MDR	https://ec.europa.eu/tools/eudamed/#/screen/search-device/5da75ce7-af12-4788-9661-d3377519008d
Classification rules (Annex VIII Regulations (EU) 2017/45	Medical device of risk Class 1
Following EU directives	EN 55014–1: 2017+A1:2020, EN/IEC 61003-3-2: 2019, EN 62233 (EMF) EN/IC 60 335-1-2-65

The device complies with the requirements of the European Medical Device Regulation MDR 2017/45. The device does not have any measuring function or used as a sterilization unit.

Skelleftea 2023-03-20

Peter Karlsten CEO Whitebox AB

CE