

Certificate of Irradiation

Product	Nunc Dish Nunclon 35 x 10	Date 2025-06-17
		Page 1
Item Number	153066	
Lot Number	185432	
Specified Dose Range	15.8 - 44.3 kGy	
SAL	10E-3	
Quantity	480 boxes	

Thermo Fisher Scientific confirms that the product supplied has been inspected and tested in accordance to relevant specifications.

Irradiation lot no.	PalletID	Date of irradiation	Dose kGy
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300427A	DK00397568	2025-06-11	18.9
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300802B	DK00397569	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300816B	DK00397570	2025-06-11	18.5
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300788B	DK00397571	2025-06-11	18.4
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300900A	DK00397572	2025-06-11	18.6
EC__1300760B	DK00397573	2025-06-11	18.6
EC__1300760B	DK00397573	2025-06-11	18.6
EC__1300760B	DK00397573	2025-06-11	18.6

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EC__1300760B	DK00397573	2025-06-11	18.6

Sterility is obtained through irradiation according to ISO 11137 "Sterilization of health care products - Radiation".

Quality assurance of the product is performed in accordance with the requirements of the Quality Management System ISO 9001:2015 "Quality Management System - Requirements" and ISO 13485:2016 "Medical Devices - Quality Management Systems - System requirements for regulatory purposes".

Roskilde Site



Per Bernack
Senior Quality Assurance Manager

Please note that this is an electronically generated signature