

Pavé du Moulin 59260 Lille-Hellemmes Tel +33 3 20 67 67 67 Fax + 33 3 20 67 67 68 web <u>vigilance@anios.com</u> To Healthcare Organisation Name Address

URGENT FIELD SAFETY NOTICE

<u>Date</u> : 22/01/19
<u>Object</u> : ☑ Batch recall ☑ Information and/or recommendations

Affected products:

Device Commercial Name	Article Code	Packaging
WIP'ANIOS PREMIUM	2088655BR /2088655C4 / 2088655C5	6 X 100 WIPES
WIP'ANIOS EXCEL	2446424C2 /2446424ZA	12X50 WIPES
	2446492GX	12X60 WIPES
	2446655C4 / 2446655C5 / 2446655CO / 2446655Y6	6 X 100 WIPES

Madam, Sir,

A microbial contamination source has been identified and localized in an outsourced manufacturing process and has brought a potential bacterial contamination to some batches of manufactured wipes.

In case of contamination triggered by contaminated wipes, immunocompromised patients would be at greater risk of infection. Globally, the risk assessment for potential health hazard linked to the use of those wipes results in low risk for the patient and the user. It takes into consideration the products' indication, the absence of any adverse event report linked to the use of the wipes, the probability of infection occurrence and the results of additional investigations (limited survival time of the bacteria on surfaces and sensitivity of this bacteria to antibiotics: piperacilline (PTZ) and ceftazidime (CZD).

Corrective actions to eliminate the contamination source have been implemented and are being closely monitored.

For precautionary reasons, we ask you not to use any longer any remaining units you may have in stock bearing the batch numbers documented in the enclosed appendix, as they may contain some contaminated wipes.

Customer n°: FSN_ANIOS_EN



We would be grateful if you could acknowledge receipt of the present communication by returning at your earliest convenience - but no later than 28/02/2019 - the attached reply form duly completed and signed.

If applicable, the proof of product's destruction has to be provided to close the current action.

Your sales contact remains at your entire disposal for any question or assistance that you may need.

The undersigns confirm that this notice has been reported to the appropriate Regulatory Agency

Please accept our apologizes for the inconvenience it may have caused.

Yours faithfully

Isabelle Prévost Quality Manager	Dr Monique Manche Materiovigilance responsible	Bertrand Letartre CEO
Awien!		

This notice needs to be passed on all those who need to be aware within your organisation or to any organisation where the potentially affected devices have been transferred.

This means, if you are a distributor, that this information has to be forwarded to any customers which was delivered with one of the affected batches.

Please maintain awareness on this notice and resulting action for an appropriate period to ensure effectiveness of the corrective action.

Customer n°: FSN_ANIOS_EN



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APPENDIX I CUSTOMER REPLY FORM

1. Field Safety Notice (FSN)

FSN Reference number: FSN_ANIOS_EN

FSN Date: January 22, 2019

Affected products: Please refer to appendix 2

2. Customer Details

Customer Number	
Healthcare Organisation Name*	
Organisation Address*	
Department/Unit	
Shipping address if different to above	
Contact Name*	
Title or Function	
Telephone number*	
Email*	
Print Name*	
Signature*	
Date*	

Mandatory fields are marked with *



3. Customer action undertaken on behalf of Healthcare Organisation

□ I confi	rm receipt of the Field Safety No	otice (FSN) and th	nat I read and un	derstood its content.	
□ I perfo	rmed all actions requested by the	ne FSN.			
☐ The in	formation and required actions	have been brougl	nt to the attention	of all relevant users	and
execu	ited, including end customers in	case of distributi	on of those produ	ucts	
□ I have	destroyed affected devices - n	umber of devices	destroyed is doo	cumented in the table	below
(proof	f of destruction have to be provi	ded to close the c	urrent action)		
	Device Commercial Name	Article Code	Batch N°	Packages Quantity (units)	
□ No aff	ected devices are available for	destruction	•		
□ Other	Action (Define):				

4. Return acknowledgement to sender

Email	vigilance@anios.com
Postal Address	Laboratoires ANIOS
	Service qualité
	Pavé du Moulin
	59260 Lille – Hellemmes - France
Fax	+33 3 20 67 67 68
Deadline for returning the customer reply form	28/02/2019

It is important that your organisation takes the actions detailed in the FSN and confirms that you have received the FSN.

Your organisation's reply is the evidence we need to monitor the progress of the corrective actions

APPENDIX 2

AFFECTED PRODUCTS

Medical device name	Model	Reference number	Lot/batch number
WIP'ANIOS EXCEL	12 packs of 50 wipes	2446424C2	A00324S
WIF ANIOS EXCEE	12 packs of 50 wipes	244042402	A003243 A04420S
			A06805S
			A000033 A08122S
			A21502S
			A21502S A23602S
			A25015S
			A26503S
			A203033 A28107S
			W01716S
			W017103 W02615S
			W04204S
			W07004S
			W17911S
			W19303S
			W25811S
		044040474	W26321S
		2446424ZA	A00807S
	40	04404000	A23602S
	12 packs of 60 wipes	2446492GX	A12712S
			A20701S
			A26215S
	0 1 1100 1	0.4.400550.4	W33121S
	6 packs of 100 wipes	2446655C4	A02518S
			A02615S
			A03211S
			A04305S
			A06805S
			A07305S
			A14911S
			A16210S
			A17404S
			A18712S
			A19311S
			A20108S
			A20409S
			A23320S
			A23602S
			A24126S
			A28107S
			W00513S
			W03015S
			W04204S
			W06116S
			W06823S
			W07917S
			W08310S
			W09318S

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AFFECTED PRODUCTS

	No1-1	Defenses assembles	1 -4/14-1
Medical device name	Model	Reference number	Lot/batch number
			W13516S
			W14205S
			W15010S
			W15111S
			W16619S
			W17823S
			W20720S
			W21215S
			W23603S
			W23804S
			W24717S
			W25104S
			W25811S
		2446655C5	A00807S
			A02615S
			A06805S
			A16210S
			A19311S
			A28107S
			W03015S
			W04722S
			W12423S
			W25603S
			W29906S
			W34720S
			W35612S
		2446655CO	W03015S
		2446655Y6	A24126S
			A24712S
			A24829S
WIP'ANIOS PREMIUM	6 packs of 100 wipes	2088655BR	W01618S
			W03821S
			W18418S
			W26111S
		2088655C4	A00219S
			A18104S
			A19123S
			U08405S
			U11821S
			U17703S
			U21512S
			U25702S
			U32001S
			W01618S
			W03821S
			W10912S
			W18418S
			W20304S

APPENDIX 2

AFFECTED PRODUCTS

Medical device name	Model	Reference number	Lot/batch number
incalcal acvice flame	MOGGI	- Actoronee maniber	W24313S
			W26111S
			W29209S
			W35214S
		2088655C5	A00219S
			A05504S
			A06105S
			A18104S
			A19123S
			A19725S
			U06705S
			U08405S
			U08412S
			U17605S
			U21404S
			U21512S
			U25602S
			U25702S
			U32001S
			U33419S
			U36401S
			W01618S
			W04410S
			W10104S
			W10912S
			W12905S
			W14705S
			W18418S
			W20304S
			W24313S
			W26111S
			W29209S
			W35214S