

29 March 2017

Our reference: FSCA 3388

Dear Customer

IMPORTANT: URGENT FIELD SAFETY NOTICE ETEST® OX 256 (OXACILLIN) Foam Packaging (Ref. 520518)

Our records indicate that your laboratory has received the following products. This letter is intended for all ETEST® OX 256 (OXACILLIN) Foam packaging (Ref. 520518) users (product reference and lot numbers included below).

Table 1- Product References impacted:

Reference	Description	Lot No.	Expiry date
520518	ETEST®OXACILLIN OX 256 WW F100 520518	1003055340	01-Apr-2017
		1003315740	28-Jul-2017
		1004070580	08-Jun-2018
		1004319590	22-Sep-2018
		1004818850	21-Apr-2019
		1004890270	24-May-2019
		1005366110	12-Dec-2019

Description of the issue

Based on QC failures (MIC out of range high) for *S.aureus* ATCC 29213 strain on ETEST® OX 256 (OXACILLIN) Foam packaging (Ref. 520518) reported from the field, bioMérieux initiated a complaint investigation to confirm product issue and determine root cause.

The following have been identified:

⇒The investigation confirmed a potential performance issue for *S. aureus* ATCC 29213 QC strain and clinical *Staphylococcus* strains on ETEST® OX 256 (OXACILLIN) Foam packaging (Ref. 520518) whatever the media used, when compared to the Agar Dilution reference method, that could lead to False Resistant results.





⇒The investigation states that ETEST® OX 256 (OXACILLIN) SPB configuration (Ref. 412431, 412432) performs within the expected specifications.

Impact to Patient/User:

As a result of the referenced issue, there is a potential performance issue on strain categorization for *Staphylococcus* strains that could lead to Major errors when compared to the AD (Agar Dilution) reference method (Resistant result instead of Susceptible result) when using 2017 CLSI or 2017 EUCAST standards.

Required actions:

- Please distribute this information to all appropriate personnel in your laboratory, retain a copy in your files, and forward this information to all parties that may use this product, including others to whom you may have transferred our product.
- Recommendations for the users of 2017 CLSI and 2017 EUCAST clinical guidelines,:

Laboratories can continue to use ETEST® OX256 FOAM (Ref.520518) listed in table 1 and can directly report the results for *Staphylococcus* spp. only when applying the following recommendations:

- 1- Perform systematically the Quality Control with *S. aureus* ATCC 29213 to detect performance issues. The expected range is $0.125-0.5 \,\mu g/mL$. An out of range QC result for the strain S. aureus ATCC 29213 would **invalidate the analysis** and **the patient result should not be reported**.
- 2 Oxacillin result can be directly reported when the following cases occur:
 - ETEST® OX256 is Susceptible i.e ≤2 μg/mL for *S. aureus*, *S. lugdunensis* or ≤0.25 μg/mL for all other species of Staphylococci –in accordance with CLSI 2017 breakpoints
 - ETEST® OX256 is Susceptible i.e ≤2 mg/L for S. aureus, S. lugdunensis, S. saprophyticus or ≤0.25 mg/L for coagulase-negative staphylococci except S. lugdunensis and S. saprophyticus in accordance to EUCAST 2017 breakpoints
- 3 Oxacillin result should be confirmed by an alternative method when the following cases occur:
 - ETEST® OX256 is Resistant i.e ≥4 µg/mL for S. aureus, S. lugdunensis or ≥0.5 µg/mL for all others Staphylococci according to CLSI 2017 breakpoints



- ETEST® OX256 is Resistant i.e >2 mg/L for *S. aureus*, *S. lugdunensis* and *S. saprophyticus* or >0.25 mg/L for the coagulase-negative staphylococci except *S. lugdunensis* and *S. saprophyticus* according to EUCAST 2017 breakpoints.
- Among tests previously performed, we are asking you to identify any possible false Resistant results, analyse the related risks and determine appropriate actions if relevant.
- Complete and return the Acknowledgement Form in Attachment A for the attention of UKVIGILANCE to Fax number 0044 (0) 1256 816863 or scan and email to <u>UKVIGILANCE@biomerieux.com</u> to confirm receipt of this notice.

bioMérieux is committed to providing our customers with the highest quality product possible. We sincerely apologise for any inconvenience that this may have caused you. If you require additional assistance or have any questions, please contact our Technical Service Department on 0044 (0) 1256 480711 or email uktechnical@biomerieux.com

Yours sincerely

UKVIGILANCE





Attachment A: Acknowledgement Form – FSCA 3388

It is a requirement of the Competent Authority that you complete and return this acknowledgement form

Please return for the attention of UK Vigilance Department to fax number 0044 (0) 1256 816863 or scan and email to UKVIGILANCE@biomerieux.com

Contact Name
Company
Address
Customer number
☐ I acknowledge the receipt of bioMérieux Urgent Field Safety Notice informing this laboratory on the ETEST® OX256 FOAM packaging (Ref.520518) product issue.
☐ I have followed the instructions and implemented the actions as indicated in the Urgent Field Safety Notice.
Have you received reports of illness or injury related to the identified issue? ☐ Yes or ☐ No
Comments:
Contact Name (Block Capitals)
Signature
Date

