Caution: For Research Use. This product is intended for animal research only and not for use in humans. Not for human or animal therapeutic or diagnostic use.

# Staphylococcus aureus S. aureus 8325-4 (Xen8.1)

**Product No.: 119239** 

#### Material Provided: 1 Agar Plate Storage Conditions: -80°C

## In vitro Characteristics

## **Genetic Characteristics**

Staphylococcus aureus-Xen8.1 was derived from the parental strain *S. aureus* 8325-4. *S. aureus*-Xen8.1 was engineered through transposition of Tn4001 luxABCDE on plasmid pXen-5. Xen8.1 possesses a single stable copy of the modified P. luminescens lux operon that was inserted in the  $\delta$ toxin coding region in the RNAIII transcript downstream of the agr P3 promoter on the bacterial chromosome.

#### **Growth Characteristics**

*S. aureus*-Xen8.1 grows well in Luria Bertani (LB) medium at 37°C under ambient aeration. *S. aureus*-Xen8.1 may also be grown selectively on LB or BHI agar containing 200µg/mL kanamycin.

#### **Colonial Morphology**

On LB agar, *S. aureus*-Xen8.1 appears as small (~1.5mm), cream-colored, opaque, smooth, circular colonies.

## **Growth Curve**

*S.aureus*-Xen8.1 displays peak bioluminescence during early log-phase growth. Log-phase growth can be achieved after 1 to 1.5 hours of subculture in LB broth at 37°C, shaking at 150-200 rpm. For

these broth culture conditions, an absorbance measurement at 600nm (against a LB blank) of 0.6 is roughly equivalent to  $5 \times 10^8$  cfu/mL of *S. aureus*-Xen8.1 and the relative light intensity is 0.9 photons/sec/cell.



## **Virulence Factors**

**Hemolysis:**  $\beta$ -hemolysis on TSA + 5% sheep blood **Capsule:** literature cites that parental lacks a capsule (CP5 negative), cap5 mutant FEMS Microbiol Lett 1999 Jan 1;170(1):97-103.



**DNAse:** Negative **NaCl:** Tolerant via growth Mannitol Salts Agar **Coagulase:** Positive in 24hrs

## **Biochemical Profile**

A biochemical profile was obtained for *S. aureus*–Xen8.1 using the api 20 STAPH system available from bioMérieux.

Sugar Utilization		
D-Glucose	+	
D-Fructose	+	
D-Mannose	+	
Maltose	+	
Lactose	+	
Trehalose	+	
D-Mannitol	+	
Xylitol	-	
Raffinose	-	
Xylose	-	
D-Melibiose	+	
Sucrose	+	

Other Tests	
Nitrate Reduction	+
Alkaline Phosphatase	+
Voges Proskauer	-
$\alpha$ -methyl-D-glucoside	-
N-acetyl-glucoside	+
Arginine dihydrolase	+
Urease	+

# **MIC and MBC Data**

MIC and MBC were determined using the macrodilution methods specified in the NCCLS Approved Standard M7-A5.

NCCLS Macrodilution MIC/MBC			
Antibiotic	MIC (µg/mL)	MBC (µg/mL)	
Ceftriaxone	>32	NA	
Ciprofloxacin	>8.0	NA	
Erythromycin	1.0	1.0	
Gentamicin	2.0-4.0	4.0-8.0	
Kanamycin	>125	NA	
Penicillin G	0.013-0.063	0.063-0.125	
Tetracycline	0.25	0.5	
		(trailing endpoint)	

#### References

1. Photochem. Photobiol. Sci., 2004, 3, 451 – 458

2. Photochem. Photobiol. Sci., 2005, 4, 503 - 509

3. Biomaterials, Volume 27, Issue 22, August 2006, Pages 4157-4164

4. Wound repair and regeneration, Volume 16 Issue 3, Pages 425 - 431

#### **Product Information**

#### Warranty

#### **Antibiotic Susceptibility**

**Disk Diffusion Data** Disk diffusion tests were performed according to methods outlined in the NCCLS Approved Standard M2-A7.

Kirby-Bauer Disk Diffusion Test		
Sensitive to:	Resistant to:	
Carbenicillin 100	Kanamycin 30	
Gentamicin 20		
Penicillin G 10U		
Vancomycin 30		

PerkinElmer warrants that cells will be viable upon shipment from PerkinElmer for a period of thirty days, provided they have been properly stored and handled during this period.

#### **Disclaimers**

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