Complicated sea urchin-induced wound infection caused by \textit{Vibrio alginolyticus} and \textit{Staphylococcus lugdunensis} in a 14-year-old boy

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\textbf{Introduction:} Wound infections with \textit{Vibrio alginolyticus}, a Gram-negative bacterium found in all temperate oceans, are rarely reported. However, a rising incidence of wound infections caused by \textit{V. alginolyticus} requires better knowledge about this infectious agent.

\textbf{Case presentation:} We report the case of a 14-year-old boy suffering from a wound infection caused by \textit{V. alginolyticus} and \textit{Staphylococcus lugdunensis} after stepping on a sea urchin. Despite wound debridement and antibiotic therapy with cefaclor, the lesion did not heal over several weeks. After identification of the pathogens and antibiotic-susceptibility testing, antibiotic therapy was switched to ciprofloxacin, followed by trimethoprim/sulfamethoxazole. Two months after the accident the wound was re-epithelialized. Follow up after 6 months revealed a painful scar.

\textbf{Conclusion:} Non-cholera vibrios like \textit{V. alginolyticus} should be considered as possible causative agents in seawater-contaminated wounds. \textit{S. lugdunensis} is a relevant pathogen in mixed wound infections. Early microbiological diagnosis and antibiotic-susceptibility testing is crucial to prevent therapeutic failure.

\textbf{Keywords:} Vibrio; \textit{Vibrio alginolyticus}; coagulase-negative staphylococci; \textit{Staphylococcus lugdunensis}; wound; sea urchin.

\textbf{Introduction}

\textit{Vibrio alginolyticus} is a Gram-negative halophilic bacterium found in temperate oceans all over the world. Most isolates show resistance to penicillins and second-generation cephalosporins (French et al., 1989; Li et al., 1999), and are capable of causing serious wound infections, even in association with only minor lesions (Gomez et al., 2003; Reilly et al., 2011).

Infections in humans are rarely reported, e.g. the incidence was only 0.048 per 100 000 population in the USA in 2011 (Conrad, 2013). Nevertheless, like other \textit{Vibrio}-mediated infections there has been an increase in incidence over many years, possibly induced by an elevation of marine temperature due to global warming (Conrad, 2013; Vezzulli et al., 2012).

Herein, we report the clinical course and follow up of a 14-year-old immunocompetent boy who was suffering from a mixed wound infection caused by \textit{V. alginolyticus} and \textit{Staphylococcus lugdunensis}. In the case of wounds acquired from contact with seawater or marine organisms, clinicians should be aware of \textit{Vibrio} infections. Nevertheless, these infections are unknown to many physicians even in high-incidence countries (Osaka et al., 2004) and require careful microbiological work-up.

\textbf{Case report}

The patient was swimming in the Red Sea while on vacation in Egypt, near Hurghada, when he stepped on a sea urchin. He was injured on the medial plantar margin of his left foot.
methoxazole was continued for 20 days. Susceptible. Antibiotic therapy with trimethoprim/sulfadiazine (160/800 mg twice daily) to which both isolates were resistant. After receiving antibiotic-susceptibility test results, therapy was switched to trimethoprim/sulfamethoxazole (160/800 mg twice daily) to which both isolates were susceptible. Antibiotic therapy with trimethoprim/sulfamethoxazole was continued for 20 days.

Two bacterial isolates were obtained from the wound swab and identified as *V. alginolyticus* and *S. lugdunensis*. Numerous colonies were present up to the second (*V. alginolyticus*) and third (*S. lugdunensis*) streak area of the initial streak plates. After receiving antibiotic-susceptibility test results, therapy was switched to trimethoprim/sulfamethoxazole (160/800 mg twice daily) to which both isolates were susceptible. Antibiotic therapy with trimethoprim/sulfamethoxazole was continued for 20 days.

**Investigations**

Aerobic and anaerobic cultures were performed using Columbia blood agar, chocolate agar, Columbia CNA agar, MacConkey agar and Schaedler agar with/without kanamycin, using standard microbiological procedures. A Gram-negative rod and coagulase-negative staphylococci were grown and identified to the species level using appropriate VITEK 2 ID cards (VITEK 2 GN and GP-cartridge; bio-Mérieux) as *V. alginolyticus* and *S. lugdunensis*. Antibiotic-susceptibility testing was performed using appropriate VITEK 2 cards, AST N263 and AST P619, respectively, and interpreted according to current Clinical and Laboratory Standards Institute (CLSI) guidelines (Table 1).

For confirmation of the *V. alginolyticus* identification, the strain was sub-cultured on *Vibrio* selective agar (thiosulfate-citrate-bile-sucrose agar; Becton-Dickinson) at 37°C. After 24 h, colonies grew on the plates turning the colour of the agar to yellow as expected for *V. alginolyticus*. Additionally, matrix-associated laser desorption ionization-time of flight MS identification using a MALDI Biotyper (Bruker Daltonics) using software version 3.1 was performed, revealing *V. alginolyticus* with a score of 1.94. The next most closely related species was *Vibrio mytili* with a score of 1.84.

As the MALDI Biotyper revealed only an identification at the probable genus level, further confirmation was sought using 16S rRNA gene sequencing applying the method of Harmsen et al. (2003). The resulting 0.5 kbp amplicons were sequenced with a 3500X Genetic Analyzer (Thermo Fisher Scientific). Using the curated database of EZbiocloud (Kim et al., 2012) and criteria for microbial identification using DNA target sequences (CLSI, 2008), similarities larger than 99% were found for numerous species of the genus *Vibrio*, including *V. alginolyticus*, without sufficient discrimination for identification at the species level. Similarly, the 16S rRNA gene sequence was analysed using *BLASTN* 2.2.26 and the DNA Database of Japan (DDBJ) due to its large number of well-documented *Vibrio* spp. genome sequences (http://ddbj.nig.ac.jp/blast/; Altschul et al., 1997). More than 200 strains of *Vibrio* spp. shared the best-reached similarity of 98% to our isolate, including 10 different species (*Vibrio fischeri*, *Vibrio para-haemolyticus*, *Vibrio harveyi*, *V. alginolyticus*, *Vibrio campbellii*, *Vibrio communis*, *Vibrio orientalis*, *Vibrio rotiferianus*, *Vibrio owensii* and *Vibrio antiquarius*).

In the next step, additional multiplex PCR for the conserved transcriptional regulator genes *VptoxR*, *VctoxR* and *VvtoxR* (Osorio & Kloese, 2000), according to Bauer & Rørvik (2007), was performed. There was a negative result for all *toxR* genes, leading to an exclusion of the species *V. para-haemolyticus*, *Vibrio cholerae* and *Vibrio vulnificus* from the identification.

Finally, 16S sequencing applying the method and primers of Tarr et al. (2007) delivered two sequences of the 16S rRNA gene (456 bp upstream, 528 bp downstream), which were analysed using DDBJ and *BLASTN* as described above.

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**Fig. 1.** Lesion on the medial margin of the sole of the left foot 28 days after the accident. Diameter approximately 2 cm.
(http://ddbj.nig.ac.jp/blast/; Altschul et al., 1997). Our isolate showed identity of 100% to more than 50 strains of *V. alginolyticus* (upstream) and 99% to more than 200 strains of *V. alginolyticus* (downstream). There was one 99% match with *V. harveyi* (upstream and downstream) and one with *V. parahaemolyticus* (upstream). The sequence data were analysed with Bionumerics (Applied Maths, version 7.1; Applied Maths) and compared to previously published sequences of the most common pathogenic *Vibrio* spp. in the National Center for Biotechnology Information (NCBI) database (http://www.ncbi.nlm.nih.gov/nuccore). To visualize the phylogenetic relationship, the unweighted pair group method with arithmetic mean based on multiple alignments between the *rpoB* sequences was used (Fig. 2).

Because of the former biochemical identification results and the large number of perfect homologies to strains identified in various taxonomic studies (Ki et al., 2009; Oberbeckmann et al., 2011), we accepted *V. alginolyticus* as the final identification. For coagulase-negative staphylococci, biochemical identification is widely used and commonly accepted (Becker et al., 2014). Therefore, we accepted the Vitek 2-based identification of *S. lugdunensis* described above.

**Diagnosis**

*V. alginolyticus* and *S. lugdunensis* co-infection of a sea urchin-induced wound.

**Outcome and follow-up**

In the following months, wound healing continued slowly until the wound was epithelialized about 2 months later. When examined for follow-up 6 months after the initial accident, it was noticed that there remained an induration of the former wound with tenderness on palpation.

**Discussion**

In this case, we identified three major reasons for the prolonged, complicated wound infection. First of all, insufficient first aid and the resulting burn necrosis led to an environment where *V. alginolyticus* and *S. lugdunensis* could survive repeated debridement and disinfection. Lack of protection because of burned skin enables secondary bacterial infections.

Second, *V. alginolyticus* is well known for its numerous chromosomal and plasmid-mediated antibiotic-resistance

<table>
<thead>
<tr>
<th>Antibiotic</th>
<th><em>V. alginolyticus</em></th>
<th>Interpretation</th>
<th><em>S. lugdunensis</em></th>
<th>Interpretation</th>
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<tbody>
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<td>MIC (mg l⁻¹)</td>
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<td>Interpretation</td>
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<td>≥0.5</td>
<td>R</td>
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<td>Ampicillin</td>
<td>≥32</td>
<td>R</td>
<td>NT*</td>
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<td>S</td>
<td>NT*</td>
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<td>≤20</td>
<td>S</td>
<td>≤10</td>
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</tbody>
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I, Intermediate susceptibility; NT, not tested; R, resistant; S, susceptible.

*Testing not recommended for this species.

Table 1. MIC and antimicrobial-susceptibility test. Interpretive categories were according to the CLSI guidelines for *Vibrio* spp. (CLSI, 2015a) and *Staphylococcus* spp. with special regard to *S. lugdunensis* (CLSI, 2015b).
determinants (French et al., 1989; Li et al., 1999). Many of the expressed β-lactamases lead to resistance to ampicillin and second-generation cephalosporins, as seen in our isolate (Li et al., 1999). Resistance to trimethoprim/sulfamethoxazole is commonly reported (Li et al., 1999). Some isolates also show resistance to third-generation cephalosporins and fluoroquinolones, as reported by Ye et al. (2016). According to this evolution of antibiotic resistance and because of the typical mixed flora in chronic wound infections (like *S. lugdunensis* in our case; Altoparlak et al., 2004), early antibiotic-susceptibility testing is important to prevent therapeutic failure.

Lastly, the presence of *S. lugdunensis* may have triggered the progression of the disease. Compared to many other coagulase-negative staphylococci, *S. lugdunensis* has higher pathogenic potential. It can cause serious infections, i.e. soft tissue and wound infections as well as infective endocarditis, and has to be considered as a relevant pathogen (Becker et al., 2014).

When treating the patient, the chosen therapy in the hospital with ciprofloxacin was an appropriate choice for the infection. However, fluoroquinolone use in children is still off label for many indications (except, for example, cystic fibrosis), especially if there is an alternative treatment (Bradley et al., 2011). Therefore, therapy was changed successfully to trimethoprim/sulfamethoxazole.

Altogether, this case and its course are an example of the need to consider *Vibrio*-mediated infections in similar circumstances. Even if it is a rare disease at present, a rising incidence has been observed, as indicated above. Warming of the oceans will probably make this a global trend as the first cases from northern shores suggest (Reilly et al., 2011; Schets et al., 2006). Identification of *V. alginolyticus* is less than straightforward and requires a combination of classical biochemical identification methods, as well as appropriate selective media and advanced molecular identification methodology.

References


