

Matrix-assisted laser desorption/ionization-time of flight mass-spectrometry (MALDI-TOF) based typing of an ESBL *Escherichia coli* outbreak in a hospital setting

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Purpose

- Rapid and specific identification to subspecies level and determination of strain identity are integral components of outbreak investigation and control.
- Conventional methods such as pulsed-field gel electrophoresis (PFGE) are labour intensive and require up to one week from sample collection to result.
- MALDI-TOF is able to identify bacterial species and subspecies rapidly.
- We aimed to assess the validity of MALDI-TOF for detection of strain identity as compared to PFGE.

MALDI-TOF technology

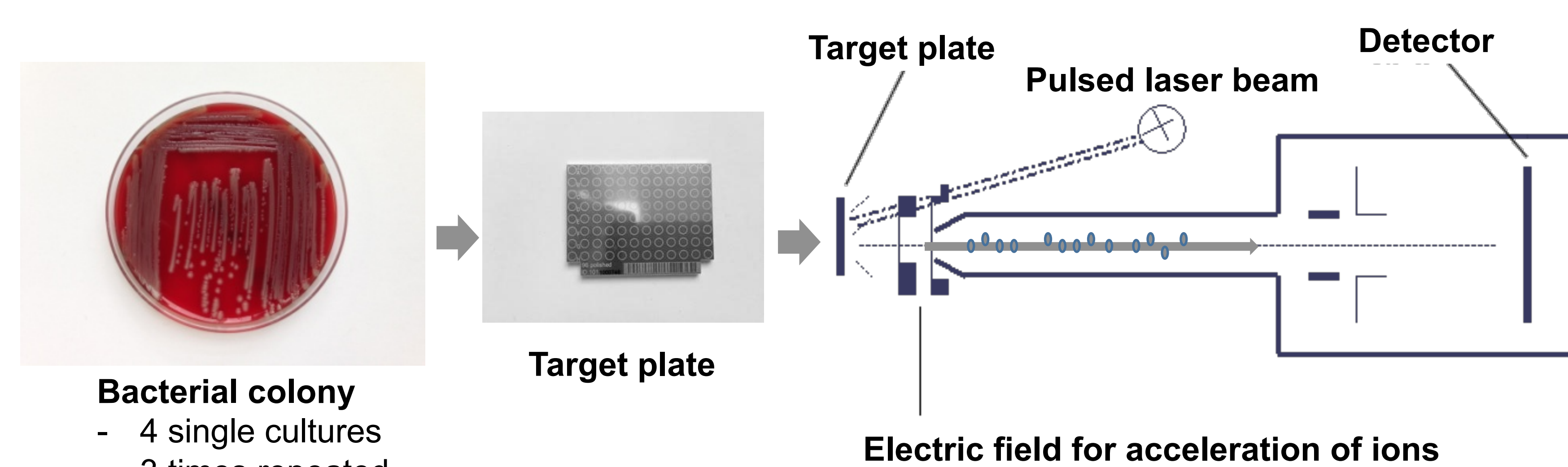


Figure 1. Work-flow for MALDI-TOF typing. The protein signature is highly specific for bacterial species

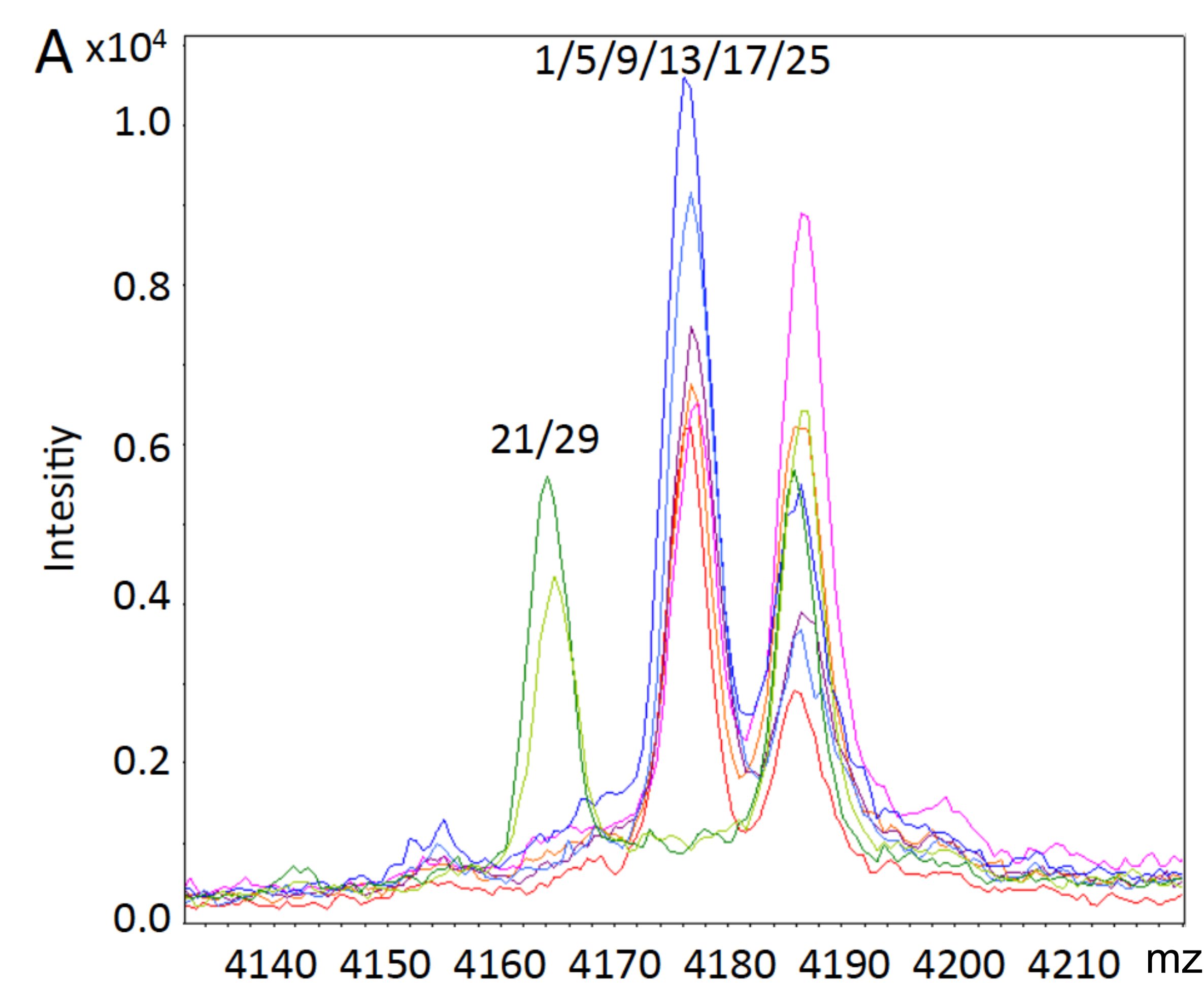


Figure 2. Frameshift analysis. Spectra were magnified and strain specific protein peaks were identified.

Methods

- ESBL-producing *E. coli* isolates from a previously investigated and published report from our institution were analysed using MALDI-TOF after a full protein extraction protocol.
- The spectra profiling was done in quadruplicates and repeated three times and analyzed using principal component analysis (PCA) to generate a dendrogram.
- Analysis was performed with Biotyper 3.0 software (Bruker Daltonic, Leipzig, Germany).

Results

- Outbreak on neonatal intensive care unit (Tschudin-Sutter S et al. Emerging Infect Dis 2010)

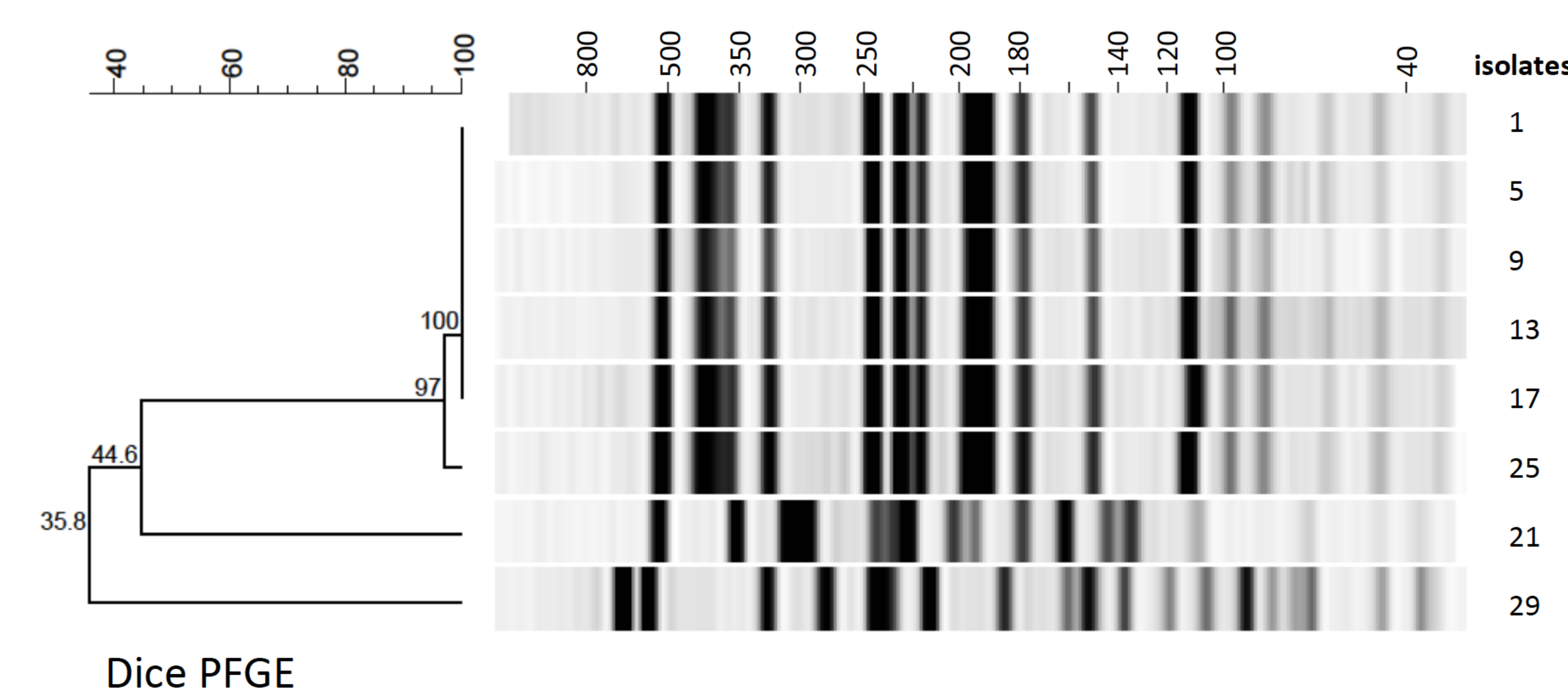


Figure 3. PFGE of ESBL-producing *E. coli* outbreak

- Eight isolates collected during an outbreak investigation in the neonatal care unit, identifying transmission of ESBL-producing *E. coli* from a mother to her newborn twins and subsequent spread to two other neonates and one healthcare worker, were analyzed by PFGE and MALDI-TOF.
- In comparison to PFGE, PCA-based typing with MALDI-TOF spectra reproduced a very similar dendrogram with identical clustering of six outbreak isolates. Two isolates of ESBL-producing *E. coli* not associated with the outbreak were correctly separated.

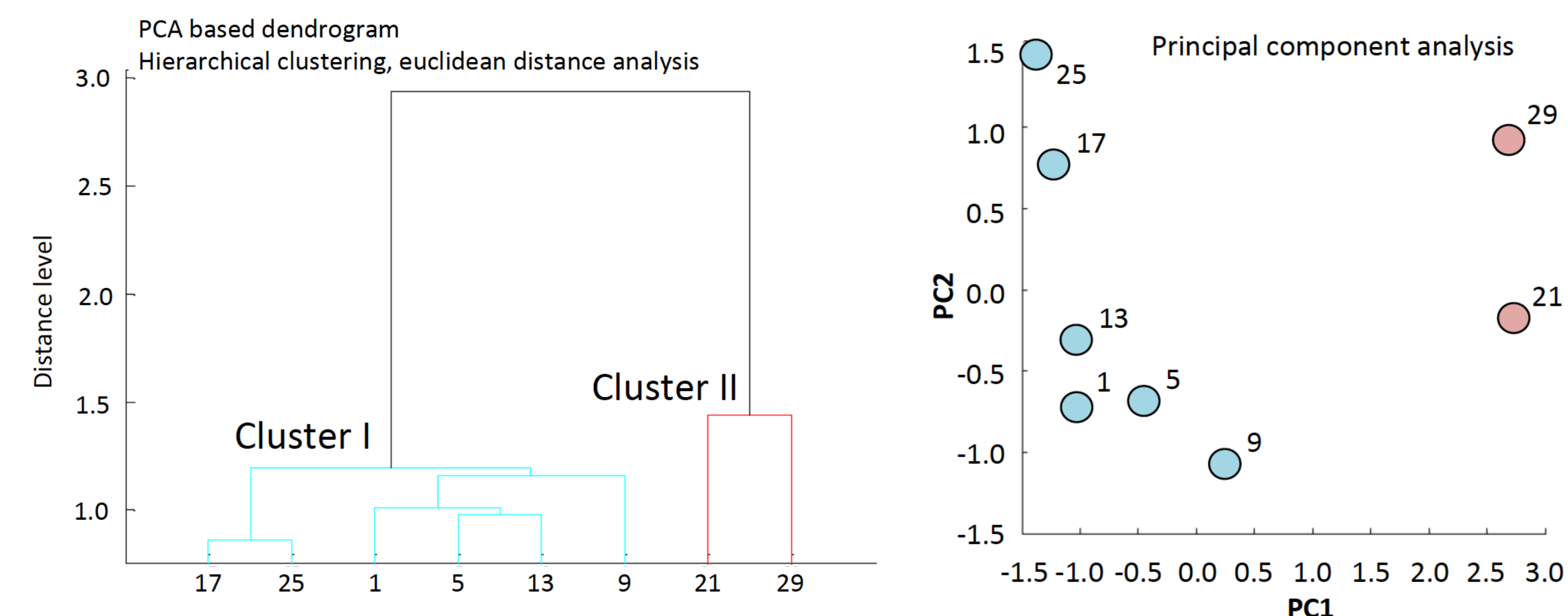


Figure 4. Principal component analysis of ESBL-producing *E. coli* isolates.

- PCA-based dendrogram and frameshift analysis of quadruplicates at three independent experiments showed a high reproducibility of the results.

Summary and Conclusions

- These results demonstrate that MALDI-TOF can provide a rapid means of typing of ESBL-producing *E. coli* isolates reliably separating outbreak-related from unrelated isolates.
- Our findings have important implications for outbreak investigations, which are hampered by the slow turnaround time of conventional typing techniques.

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