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Título	Selection and validation of reference genes for gene expression studies in <i>Klebsiella pneumoniae</i> using Reverse Transcription Quantitative real-time PCR
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Resumo	For reliable results, Reverse Transcription Quantitative real-time Polymerase Chain Reaction (RT-qPCR) analyses depend on stably expressed reference genes for data normalization purposes. <i>Klebsiella pneumoniae</i> is an opportunistic Gram-negative bacterium that has become a serious threat worldwide. Unfortunately, there is no consensus for an ideal reference gene for RT-qPCR data normalization on <i>K. pneumoniae</i> . In this study, the expression profile of eleven candidate reference genes was assessed in <i>K. pneumoniae</i> cells submitted to various experimental conditions, and the expression stability of these candidate genes was evaluated using statistical algorithms BestKeeper, NormFinder, geNorm, Delta C _T and RefFinder. The statistical analyses ranked <i>recA</i> , <i>rho</i> , <i>proC</i> and <i>rpoD</i> as the most suitable reference genes for accurate RT-qPCR data normalization in <i>K. pneumoniae</i> . The reliability of the proposed reference genes was validated by normalizing the relative expression of iron-regulated genes in <i>K. pneumoniae</i> cells submitted to iron-replete and iron-limited conditions. This work emphasizes that the stable expression of any potential reference candidate gene must be validated in each physiological condition or experimental treatment under study.
Fomento	