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Volume 58, no. 4, p 2030–2037, 2014. Page 2030, right column, line 6. Since the suspension of the European marketing authorization for telavancin has now been lifted, the following text in the introduction no longer applies: "At the time of submission, the telavancin European marketing authorization for the treatment of nosocomial pneumonia was suspended pending evidence of a new European Medicines Agency-approved supplier. Clinigen Healthcare Ltd., Theravance's commercialization partner for telavancin in Europe, is in the process of seeking approval of a new manufacturing source."

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