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Base-catalyzed hydrolysis and speciation-dependent photolysis of two cephalosporin antibiotics, Ceftiofur and Cefapirin

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12 Abstract

Lately, special attention has been given to veterinary cephalosporin antibiotics due to their broad 13 activity spectrum and significant consumption. Indeed, the determination of hydrolytic and 14 photolytic kinetics provides a better comprehension of the undesired persistence of cephalosporins 15 in aqueous matrices. In this work, the two widely used veterinary antibiotics ceftiofur (CEF) and 16 cefapirin (CEPA) showed high instability under alkaline conditions, degrading in few minutes at pH 17 >11. In buffered solutions at neutral pH and natural temperature (T = $22 \pm 1^{\circ}$ C), both drugs 18 presented moderate stability ($t^{1/2} = 3$ d, CEPA and 1.4 d, CEF). Our study also demonstrated that 19 CEPA and CEF speciation did not significantly influence the direct photolysis rates. Using a 20 simulated water disinfection set-up ($\lambda = 254$ nm), all ionic species of CEF and CEPA presented fast 21 22 and similar pseudo-first order degradation rates, kapp 0.0095 ± 0.0004 and 0.0092 ± 0.001 mJ cm-2, respectively. Furthermore, using surface water in hydrolysis experiments, CEF demonstrated 23 significant matrix-dependent stability with a half-life ($t^{1/2} = 14.7$ d) tenfold higher than in buffered 24 solutions. In contrast, CEPA presented a very similar hydrolysis rate in river water ($t\frac{1}{2} = 4.2$ d) and 25 26 a subtle faster photo-degradation rate in this same matrix (kapp 0.0128 ± 0.001 mJ cm-2),

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