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Stably transfected MDCK cells expressing eGFP-kAE1 were seeded on Transwell filters and grown for 4 days. kAE1 expression was induced for 4, 6, 8, 10, 12, 16, 20, or 24 h. Cell surface proteins were isolated by biotinylation, then separated by SDS-PAGE followed by Western blot analysis of the same blot using anti-AE1 (Bric170) or anti-GP135 as described in 'Methods'. Steady state surface expression of eGFP-kAE1 (confirmed by densitometry) was observed at 12 and 16 h.