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## CORRIGENDUM

### Integrins in cancer: biological implications and therapeutic opportunities

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On page 17 of this article, in the section Targeting  $\alpha v\beta 3$  and  $\alpha v\beta 5$  the sentence at the start of the second paragraph that reads "Cilengitide is an inhibitor of both  $\alpha v\beta 3$  and  $\alpha v\beta 5$  integrins, and it was selected in our laboratory by screening a library of cyclic RGD peptides in a cell-free receptor assay for their capacity to inhibit integrins  $\alpha v\beta 3$  and  $\alpha v\beta 5$  but not  $\alpha IIb\beta 3$  (REF. 130)." was incorrectly phrased. The corrected sentence with additional references is given below.

"Cilengitide is an inhibitor of both  $\alpha v\beta 3$  and  $\alpha v\beta 5$  integrins. We had shown that  $\alpha v\beta 3$  and  $\alpha v\beta 5$  integrins were important regulators of angiogenesis and tumour growth<sup>193,195,196</sup> and developed a cell-free receptor assay to select for antagonists of integrins  $\alpha v\beta 3$  and  $\alpha v\beta 5$  that did not effect integrin  $\alpha IIb\beta 3$  (REF. 130). This assay was used to screen a library of integrin binding cyclic RGD peptides designed and synthesized by H. Kessler and colleagues for  $\alpha v\beta 3$  activity and selectivity<sup>193–195</sup> from which cilengitide was developed<sup>196</sup>.

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