

# The University of Bradford Institutional Repository

<http://bradscholars.brad.ac.uk>

This work is made available online in accordance with publisher policies. Please refer to the repository record for this item and our Policy Document available from the repository home page for further information.

To see the final version of this work please visit the publisher's website. Access to the published online version may require a subscription.

**Link to publisher's version:** <http://dx.doi.org/10.1021/jm5000547>

**Citation:** Sheldrake HM and Patterson LH (2014) Strategies to inhibit tumour associated integrin receptors: rationale for dual and multi-antagonists. *Journal of Medicinal Chemistry*. 57(15): 6301-6315.

**Copyright statement:** This document is the Accepted Manuscript version of a Published Work that appeared in final form in *Journal of Medicinal Chemistry*, copyright © 2014 American Chemical Society after peer review and technical editing by the publisher. To access the final edited and published work see <http://dx.doi.org/10.1021/jm5000547>

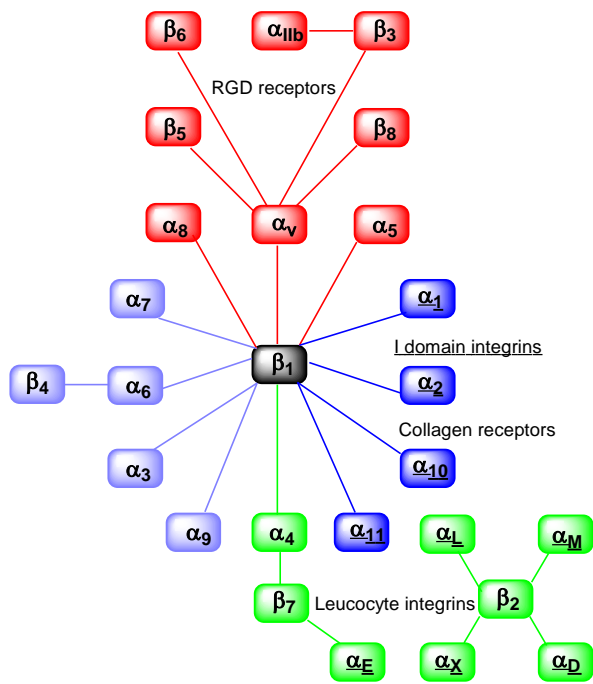
## **Strategies to inhibit tumor associated integrin receptors: rationale for dual and multi-antagonists**

Helen M. Sheldrake\* and Laurence H. Patterson

Institute of Cancer Therapeutics, University of Bradford, Bradford, UK.

**Abstract:** The integrins are a family of 24 heterodimeric transmembrane cell surface receptors. Involvement in cell attachment to the extracellular matrix, motility, and proliferation identifies integrins as therapeutic targets in cancer and associated conditions; thrombosis, angiogenesis and osteoporosis. The most reported strategy for drug development is synthesis of an agent that is highly selective for a single integrin receptor. However, the ability of cancer cells to change their integrin repertoire in response to drug treatment renders this approach vulnerable to the development of resistance and paradoxical promotion of tumor growth. Here, we review progress towards development of antagonists targeting two or more members of the RGD-binding integrins, notably  $\alpha_v\beta_3$ ,  $\alpha_v\beta_5$ ,  $\alpha_v\beta_6$ ,  $\alpha_v\beta_8$ ,  $\alpha_5\beta_1$ , and  $\alpha_{IIb}\beta_3$ , as anticancer therapeutics.

### **Integrin structure and implications for drug development**



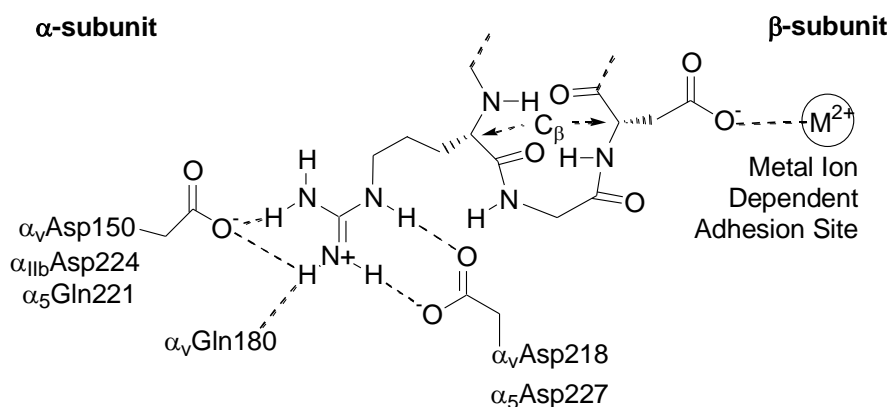
**Figure 1** Combinations of integrin subunits which form the 24 human receptors

Recognition sequence	Integrins	Major Ligands
RGD	$\alpha_v\beta_1$ , $\alpha_v\beta_3$ , $\alpha_v\beta_5$ , $\alpha_v\beta_6$ , $\alpha_v\beta_8$ , $\alpha_5\beta_1$ , $\alpha_8\beta_1$ , $\alpha_{11b}\beta_3$	Vitronectin, Fibronectin, Osteopontin, Fibrinogen
LDV and related sequences	$\alpha_4\beta_1$ , $\alpha_9\beta_1$ , $\alpha_4\beta_7$ , $\alpha_E\beta_7$ , $\alpha_L\beta_2$ , $\alpha_M\beta_2$ , $\alpha_X\beta_2$ , $\alpha_D\beta_2$	Fibronectin, Vascular Cell Adhesion Molecule 1, Mucosal Addressin Cell Adhesion Molecule-1, Intercellular Cell Adhesion Molecule-1
GFOGER	$\alpha_1\beta_1$ , $\alpha_2\beta_1$ , $\alpha_{10}\beta_1$ , $\alpha_{11}\beta_1$	Collagen, Laminin
Other	$\alpha_3\beta_1$ , $\alpha_6\beta_1$ , $\alpha_7\beta_1$ , $\alpha_6\beta_4$	Laminin

**Table 1** Integrin ligands and recognition sequences

The integrin family comprises 24 transmembrane receptors, each a heterodimeric combination of 1 of 18  $\alpha$  and 1 of 8  $\beta$  protein subunits. Their function is to integrate adhesion and interaction with the extracellular microenvironment with intracellular signalling and cytoskeletal rearrangement through transmitting signals across the cell membrane on ligand binding. They are usually classified into subfamilies based on the major protein recognised, type of cell expressing the receptor, or the presence of an I-domain in the  $\alpha$  subunit (Figure 1). The largest subfamily recognise the RGD tripeptide found in a range of extracellular matrix (ECM) ligands (Table 1), although these integrins also contain interaction sites for other proteins such as matrix metalloproteinases, and are able to bind other sequences. Integrins involved in immune functions recognise LDV and related sequences such as LDT and IDS. A subset of the collagen and laminin binding integrins recognise GFOGER, and the remaining laminin binding integrins recognise a range of peptide sequences with no common features currently reported.<sup>1</sup>

The binding site for RGD and related sequences is located at the junction of the  $\alpha$  and  $\beta$  subunits of RGD-binding integrins such as  $\alpha_v\beta_3$  and  $\alpha_5\beta_1$ . Key features of the RGD binding site are shown in Figure 2. Ligand binding has been explored through NMR studies, mutagenesis, X-ray structures of bound ligands and homology modelling.<sup>2,3</sup> Ligand binding to the  $\beta$  subunit is primarily through an electrostatic interaction between a carboxylate group on the ligand and a positively charged metal ion (usually  $Mg^{2+}$ ) associated with the integrin subunit. The  $\alpha$  subunit binds the basic arginine sidechain through interactions with several subunit specific acidic residues.



**Figure 2 Key features of the RGD binding site**

Differences in binding site size and the identity of the arginine-binding residues have allowed development of integrin specific ligand mimetics which act as competitive antagonists; however, the information amassed in determining ligand specificity should also be applicable to the design of multi-integrin antagonists. NMR studies on peptides have shown that molecules binding to  $\alpha_{IIIb}\beta_3$  typically have a distance of 7.5-8.5 Å between the  $\beta$  carbons of arginine and aspartate, and compounds binding  $\alpha_v\beta_3$  and  $\alpha_5\beta_1$  less than 6.7 Å.<sup>4</sup> For peptidomimetic ligands, it is easier to measure the distance between acidic and basic groups acting as sidechain mimetics; typically 13.5-15.5 Å is required for anti- $\alpha_{IIIb}\beta_3$  activity and 10.0-13.8 Å for anti- $\alpha_v\beta_3$ .<sup>5,6</sup>  $\alpha_5\beta_1$  also binds molecules spanning ~13 Å, although its binding site is larger than  $\alpha_v\beta_3$  with space for bulky residues adjacent to the RGD site and allowing more lipophilic interactions.<sup>7</sup>  $\alpha_5\beta_1$  binding to the arginine sidechain has similarities to both  $\alpha_{IIIb}$  and  $\alpha_v$ .<sup>3</sup>  $\alpha_v\beta_5$  is highly similar to  $\alpha_v\beta_3$  in the ligand binding site region although homology modelling suggests the  $\alpha_v\beta_5$  binding pocket is somewhat smaller and cannot accommodate large groups adjacent to the metal-ion dependent adhesion site.<sup>8</sup> The structures of the other RGD-binding integrins,  $\alpha_v\beta_6$ ,  $\alpha_v\beta_8$ ,  $\alpha_v\beta_1$  and  $\alpha_8\beta_1$ , have not yet been studied in detail, although two recently reported homology models for  $\alpha_v\beta_6$ <sup>9,10</sup> provide an important starting point to develop an understanding of antagonist selectivity. It has been shown experimentally that the common  $\alpha_v$  subunit and distinct  $\beta$  subunits allow the synthesis of small molecules with

varying degrees of selectivity within the subfamily.  $\alpha_8\beta_1$  has not yet been proven to be a target for therapeutic intervention. It has some functions in common with other members of the subfamily, suggesting there may be some value in multi-targeting, but further investigations are required to establish whether this will be safe, effective, and possible.

Integrins possess redundancy in ligand recognition, adhesion and signalling functions, and cells invariably express multiple integrins from several subfamilies. Crosstalk between integrins can affect cell functions, for example *via* trans-dominant inhibition where competition for intracellular signalling effectors means inhibition of the function of one integrin promotes activation, adhesion and signalling of another.<sup>11</sup>

Despite the intense interest in integrins as targets in a range of diseases,<sup>12,13</sup> the main strategy to date has been to develop a highly selective antagonist of a single integrin, whilst the deliberate development of dual antagonists has been little explored,<sup>14</sup> with the exception of the application of  $\alpha_4\beta_1/\alpha_4\beta_7$  antagonists in autoimmune disorders.<sup>15</sup> The drive towards selectivity for a single integrin may reduce the effectiveness or safety of the developed antagonists, leaving them vulnerable to the development of resistance, or even paradoxical effects, as targeting one receptor promotes the upregulation of a related receptor binding the same ligand to maintain adhesion and signalling.

This Perspective review summarises evidence of the importance, and interactions, of pairs and wider combinations of integrins in cancer progression and dissemination, and progress in the development of dual and multi-antagonists to efficiently target these processes.

## **Dual antagonism**

### **$\alpha_{IIb}\beta_3/\alpha_v\beta_3$**

$\alpha_{IIb}\beta_3$  is normally expressed only on platelets and megakaryocytes. On platelet activation by any agonist,  $\alpha_{IIb}\beta_3$  is activated and binds fibrinogen through its RGD and KQAGDV motifs

causing platelet crosslinking thus thrombus formation.<sup>16</sup> Antagonism of  $\alpha_{IIb}\beta_3$  prevents platelet adhesion and aggregation, therefore it has been a popular target in the development of broad spectrum anti-thrombotic therapy.<sup>17</sup> Three  $\alpha_{IIb}\beta_3$  antagonists are currently approved for clinical use in the treatment of acute coronary syndrome and during percutaneous coronary intervention; the cyclic peptide Eptifibatide, small molecule Tirofiban and antibody Abciximab. Unlike the former two, Abciximab is not selective [IC<sub>50</sub> (cell adhesion)  $\alpha_{IIb}\beta_3$  6.5 nM,  $\alpha_v\beta_3$  9.8 nM,  $\alpha_M\beta_2$  160 nM]<sup>18</sup> and its ability to antagonise both  $\beta_3$  integrins and potentially reduce inflammation has been suggested as the reason for its better performance at preventing restenosis compared to  $\alpha_{IIb}\beta_3$ -specific antagonists.<sup>19,20</sup>

The most important sites of expression of  $\alpha_v\beta_3$  are osteoclasts, where it controls attachment to the bone surface to form a sealing zone for bone resorption,<sup>21</sup> and on active endothelial cells where it mediates vascular angiogenesis.<sup>22</sup> The role of integrins in angiogenesis is complex; antagonism or knockout of  $\alpha_v\beta_3$  can promote as well as inhibit angiogenesis, but a recent study has confirmed that  $\alpha_v\beta_3$  is a valid target for preventing the early stages of angiogenesis.<sup>23</sup>

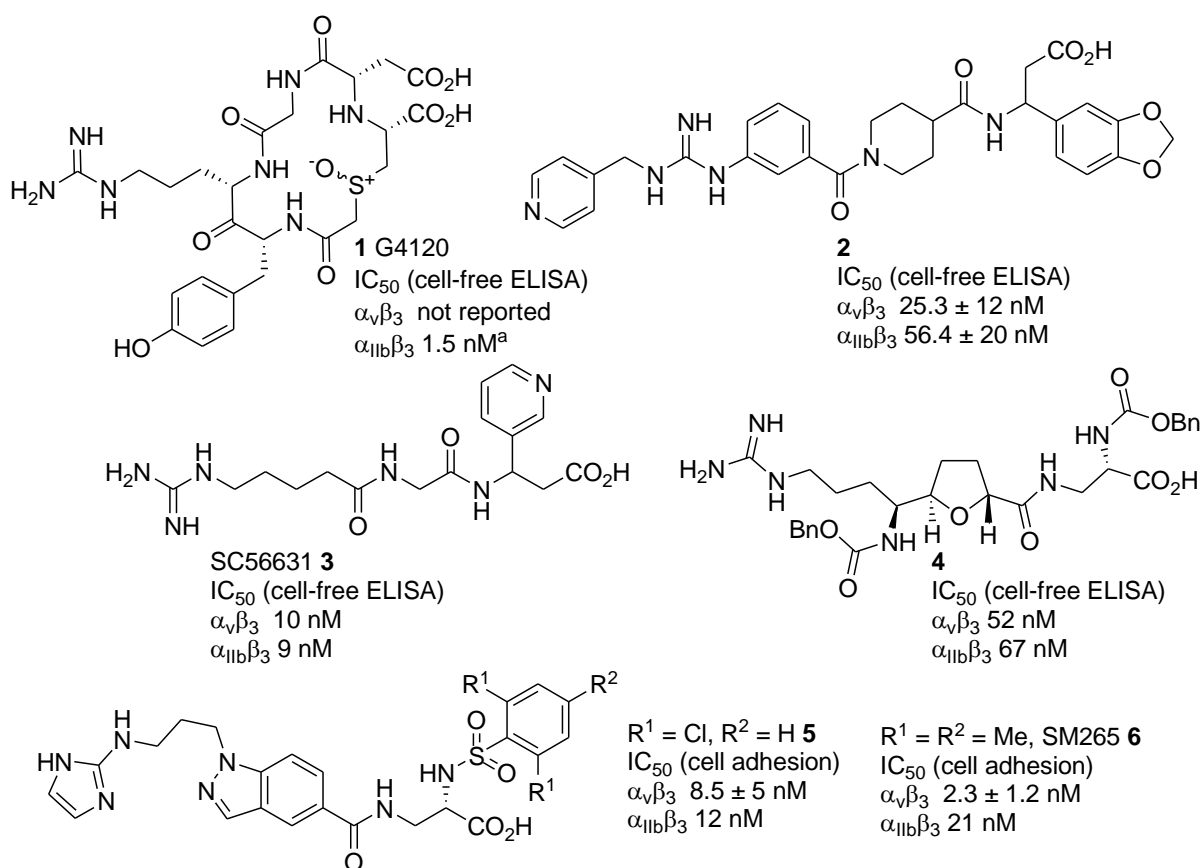
$\alpha_v\beta_3$  is expressed on a wide range of tumors and associated vasculature, where it is associated with invasion, metastasis and poor prognosis.<sup>24-27</sup>  $\alpha_v\beta_3$  promotes site-specific metastasis to the lungs and bone; adhesion to fibronectin, vitronectin, osteopontin or bone sialoprotein allows the metastatic deposit to become established<sup>28-32</sup> and tumoral  $\alpha_v\beta_3$  signalling is required for bone deposition in osteoblastic lesions, in contrast with the recognised role of osteoclast  $\alpha_v\beta_3$  in osteolytic metastases.<sup>30,33-35</sup>

$\alpha_{IIb}\beta_3$  is abnormally expressed in melanoma and prostate tumors and is associated with increased tumor growth, recurrence and metastasis.<sup>36-38</sup> Cells co-expressing  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_3$  show increased growth and angiogenesis *in vivo*, but reduced  $\alpha_v\beta_3$  function due to its displacement or trans-dominant inhibition by  $\alpha_{IIb}\beta_3$ , suggesting that selectively targeted anti- $\alpha_v\beta_3$  antagonists will prove clinically ineffective in dual  $\beta_3$  expressing tumors.<sup>36,39</sup>

The full potential of dual  $\alpha_v\beta_3/\alpha_{IIb}\beta_3$  integrin antagonism is evident when considering their roles in metastasis.  $\beta_3$  integrins promote lymphatic metastasis, and hematogenous metastasis through promoting extravasation from the primary tumor, cell adhesion, intravasation and tumor growth at the metastatic site.  $\beta_3$ -mediated interactions between tumor cells and platelets promote platelet aggregation and release of growth factors, and increase cell survival through formation of a tumor-platelet microthrombus which protects circulating tumor cells from immune targeting and facilitates arrest and adhesion in blood vessels.<sup>40-45</sup>

Both selective anti- $\alpha_{IIb}\beta_3$  antagonists, and  $\alpha_v\beta_3$  antagonists have shown a wide range of anticancer effects (for a previous review see<sup>26</sup>). Studies using Abciximab and related murine antibodies are particularly important in demonstrating the potential of dual  $\beta_3$  antagonism, although the problems of immunogenicity and bleeding associated with Abciximab suggest further development of small molecules is required to provide an attractive clinical candidate. Dual  $\beta_3$  antagonism with these antibodies was effective at blocking tumor growth and angiogenesis through targeting tumor cells' interaction with platelets and endothelial cells, in addition to direct effects on tumor tissue.<sup>46-48</sup> In a model of bone metastasis, daily treatment with m7E3 F(ab')<sub>2</sub> was effective at reducing growth of  $\beta_3$  negative tumors implanted in the tibia through acting on platelets and the bone microenvironment, indicating that this target will be applicable to a wide range of tumors.<sup>47</sup> m7E3 F(ab')<sub>2</sub> pretreatment was also shown to reduce the development of lung metastases after intravenous injection of tumor cells as a model of the early stages of hematogenous dissemination.<sup>49</sup> Combining two small molecule  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_3$  antagonists has also been shown to be more effective than either agent alone at preventing tumor cell adhesion to endothelial cell or ECM.<sup>50,51</sup>



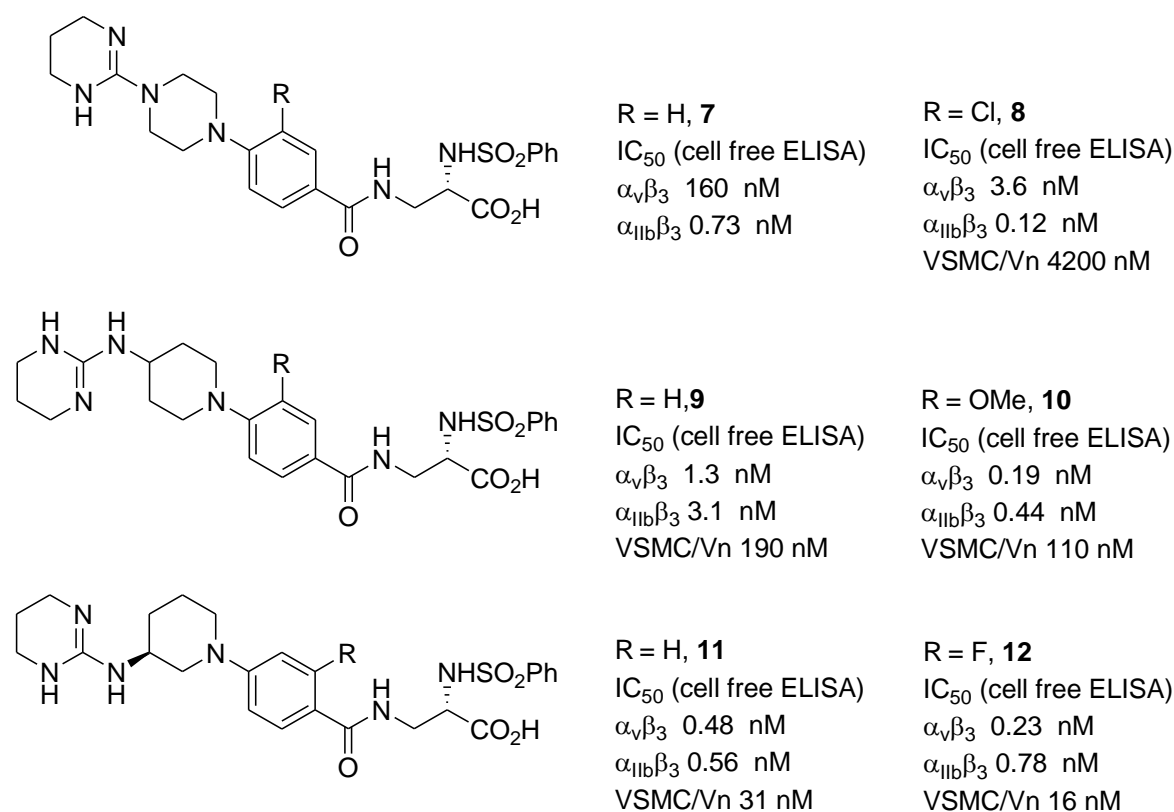


**Figure 3**  $\alpha_{IIb}\beta_3/\alpha_v\beta_3$  antagonists <sup>a</sup>The absence of standard deviation from a measurement indicates this data was not reported in the original reference.

Dual antagonists have been obtained incidentally during the development of highly active selective  $\alpha_{IIb}\beta_3$  or  $\alpha_v\beta_3$  antagonists eg **1-4** (Figure 3).<sup>52-54</sup>

The cyclic peptides G4120 **1** and G3580 (a related structure which unfortunately cannot be unambiguously identified)<sup>55</sup> were found to be non-specific<sup>56</sup> and have been used as tools in integrin biology and to demonstrate that inhibition of both  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_3$  was effective at reducing restenosis, although there was no advantage in using G3580 [ $IC_{50}$  (cell free ELISA)  $\alpha_{IIb}\beta_3$  1.5 nM;  $\alpha_v\beta_3$  8 nM] compared to a highly selective  $\alpha_{IIb}\beta_3$  or  $\alpha_v\beta_3$  antagonist alone.<sup>57</sup> SC56631 **3**, developed in an anti- $\alpha_v\beta_3$  antiosteoporotic programme, has been used as a tool compound to demonstrate that  $\alpha_v\beta_3$  antagonism is cytotoxic to cysteine-rich protein 61 overexpressing cancer cells which depend on integrin survival signalling; in this  $\alpha_{IIb}\beta_3$  negative *in vitro* model it was less effective when compared to antagonists with subnanomolar  $IC_{50}$ s on  $\alpha_v\beta_3$ .<sup>58</sup> SM265 **6** has demonstrated anticancer effects *in vitro* and *in*

*in vivo* through binding to  $\alpha_v\beta_3$  but possible contribution of  $\alpha_{IIb}\beta_3$  to its *in vivo* efficacy has not been investigated.<sup>59,60</sup>



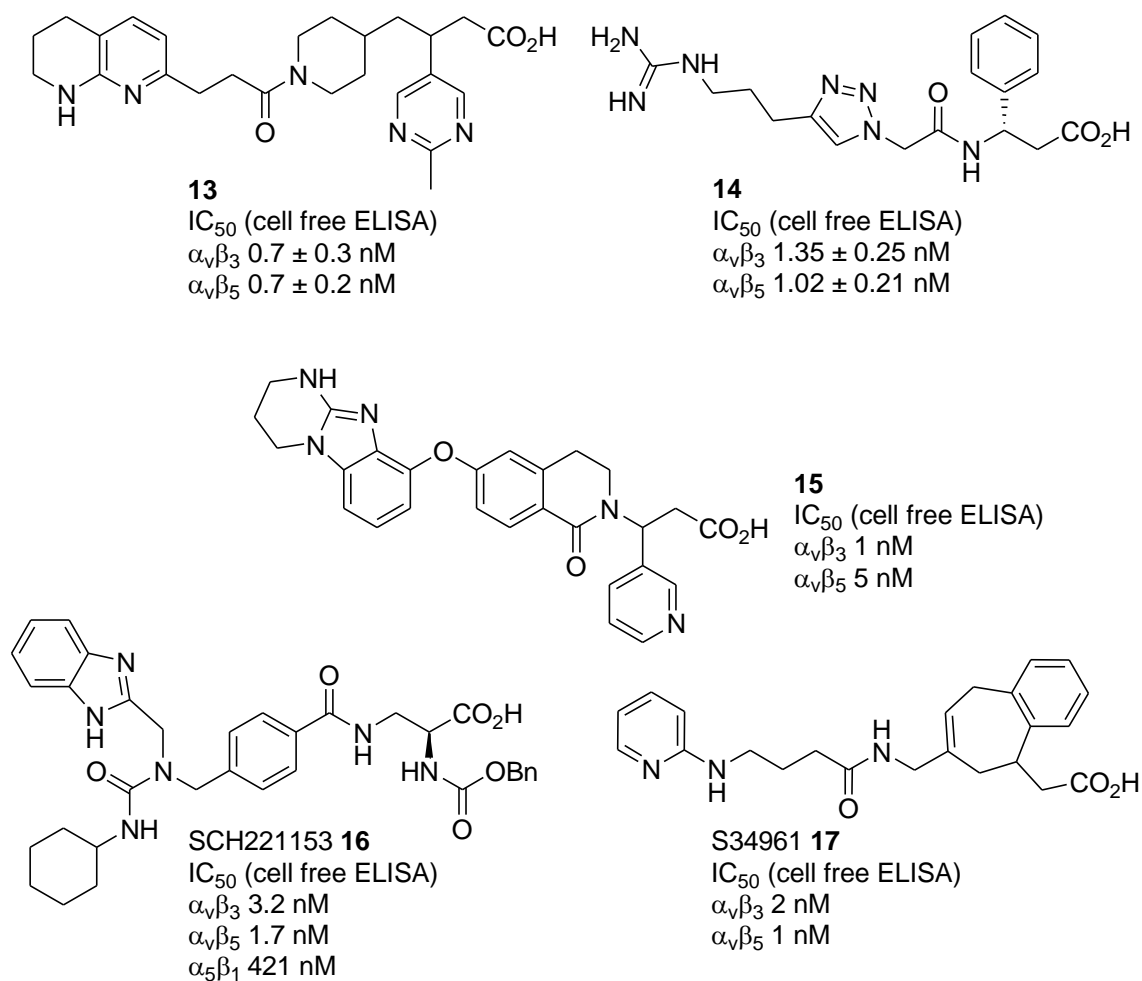
**Figure 4 MN447 and related antagonists**

MN447 **10** (Figure 4) is the only small molecule deliberately developed as a dual  $\beta_3$  antagonist, intended as a small molecule analogue of Abciximab for the treatment of acute thrombotic events. Starting from a selective  $\alpha_{IIb}\beta_3$  antagonist **7**, altering substituents on the central aromatic ring gave compound **8** with nanomolar anti- $\alpha_v\beta_3$  and anti- $\alpha_{IIb}\beta_3$  activity in cell free ELISA but relatively ineffective at inhibiting  $\alpha_v\beta_3$ -mediated cell adhesion to vitronectin.<sup>61</sup> *Para*-substitution about the central aromatic ring is required for  $\alpha_{IIb}\beta_3$  activity; meta-substitution shortens the molecule thus yielding selective  $\alpha_v\beta_3$  antagonists.<sup>62</sup> Changing to a 4-aminopiperidine linker **9** improved the anti-adhesion activity,<sup>61</sup> and adding either R = F, OH or OMe further increased potency and resulted in good water solubility.<sup>63</sup> 3-aminopiperidine also afforded a highly active scaffold eg **11**, **12**.<sup>64</sup> MN447 **9** (R = OMe) was effective in treating thromboembolus-induced dysfunction of the heart muscle *in vivo* and is

currently in preclinical development. Importantly, both MN447 **10** and SC56631 **3** have been demonstrated not to prolong bleeding time.<sup>63,65</sup>

### **$\alpha_v\beta_3/\alpha_v\beta_5$**

$\alpha_v\beta_5$  has a similar expression pattern and function to  $\alpha_v\beta_3$ ; both are highly expressed by activated endothelial cells and have similar roles in angiogenesis, promoting the angiogenic response to different growth factors.<sup>66</sup>  $\alpha_v\beta_5$  has been shown to be highly expressed on a wide range of tumor types in both cell lines and clinical material;<sup>67</sup> this ubiquitous expression suggests it may be a good target receptor for developing tumor-directed therapies to benefit the widest range of patients. However, a number of tumors and associated vasculature co-express  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$ ;<sup>68-70</sup> given that the two integrins engage the same ECM ligands and activate complementary cell signalling pathways<sup>71-73</sup> to promote tumor progression, dual targeting will be necessary to effectively treat such tumors.



**Figure 5**  $\alpha_v\beta_3/\alpha_v\beta_5$  antagonists

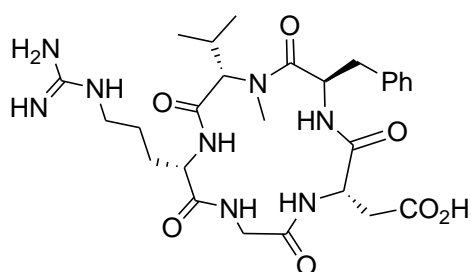
Screening of compounds which are highly active against  $\alpha_v\beta_3$  reveals they frequently also possess similar activity against  $\alpha_v\beta_5$  eg **13**, **14** (Figure 5).<sup>53</sup> Alternatively, dual antagonists have been deliberately prepared through screening to identify and optimise compounds with similar affinity for both integrins eg **15**.<sup>74</sup> **14** is antiangiogenic and is highly effective at preventing adhesion of  $\alpha_v\beta_3/\alpha_v\beta_5$  expressing melanoma cells to a range of RGD-containing ECM ligands, particularly osteopontin, indicating that it may be effective in reducing bone metastasis. It appears to be selective for inhibiting the active form of the integrin. Decreasing the length of the guanidine-bearing sidechain gave an  $\alpha_v\beta_5$ -selective agent.<sup>75</sup>

SCH221153 **16** was developed by screening a combinatorial RGD-mimetic library. The diaminopropionic acid aspartate mimetic was identified as essential for activity, and compounds optimised for high  $\alpha_v\beta_3$  activity, resulting in the selective dual  $\alpha_v\beta_3/\alpha_v\beta_5$

antagonist **16**.<sup>76</sup> Despite a short half-life (12 minutes), **16** significantly reduced the growth of  $\alpha_v$ -negative melanoma xenografts, indicating that continuous exposure to high blood levels of integrin antagonist is not required to effectively inhibit angiogenesis, and contrasting with a subsequent report that drops in blood levels of an  $\alpha_v\beta_3$  antagonist to subtherapeutic concentrations promote tumor growth.<sup>77</sup>

S34961 **17** was developed using the cycloheptene ring system to mimic the conformation of the RGD motif in natural integrin ligands. Other arginine sidechain mimetics such as tetrahydropyrimidine, tetrahydronaphthyridine, imidazoline and benzimidazoline also gave high dual antagonist activity.<sup>78</sup> *In vitro*, **17** caused cell detachment and sensitized quiescent colon cancer cells to mitogen-activated protein kinase inhibition by reducing integrin survival signaling.<sup>79</sup> The resolved *S* isomer of **17** (S 36578-2) was shown to cause anoikis of endothelial cells.<sup>80</sup>

### Cilengitide



Cilengitide **18**  
IC<sub>50</sub> (cell-free ELISA)  
 $\alpha_v\beta_3$  0.65 ± 0.07 nM  
 $\alpha_v\beta_5$  11.7 ± 1.5 nM  
 $\alpha_5\beta_1$  13.2 ± 0.6 nM

### Figure 6 Cilengitide

The cyclic pentapeptide Cilengitide **18** (Figure 6) is the first non-antibody integrin antagonist to progress to Phase III clinical trials for the treatment of cancer. The development of Cilengitide has been previously described by its inventors,<sup>81</sup> so will be briefly summarised here. Briefly, Cilengitide was derived from the cRGDfV pentapeptide by *N*-methylation;

methylation of valine was found to be the preferred position for increasing activity against  $\alpha_v\beta_3$  whilst maintaining selectivity over  $\alpha_{IIb}\beta_3$ .<sup>82,83</sup> Cilengitide is less selective against other RGD-binding integrins; it is usually described as an  $\alpha_v\beta_3/\alpha_v\beta_5$  antagonist but has similar activity (12 nM) against both  $\alpha_v\beta_5$  and  $\alpha_5\beta_1$ . Addition of a second *N*-methyl group to the glycine, aspartic acid or D-phenylalanine residue improved the  $\alpha_v\beta_3$  selectivity of the peptide.<sup>84</sup>

The progress of Cilengitide through clinical trials was most recently summarised by Scaringi *et al.*<sup>85</sup> It showed a good safety profile in all applications, and reached Phase III clinical trials against glioblastoma. Unfortunately, this first Phase III trial recently failed to reach its primary endpoint of increased progression-free survival, discouraging further development.<sup>86</sup> Preclinical and some clinical results indicate that Cilengitide should be used as a combination with radiotherapy or other chemotherapies.<sup>87</sup> Notably, combination with cytotoxic chemotherapy in head and neck squamous cell carcinoma achieved 100% disease control in the Phase I cohort.<sup>88</sup> Other preclinical studies indicate it may be effective in preventing or treating bone metastasis.<sup>89,90</sup>

### **$\alpha_v\beta_3/\alpha_5\beta_1$**

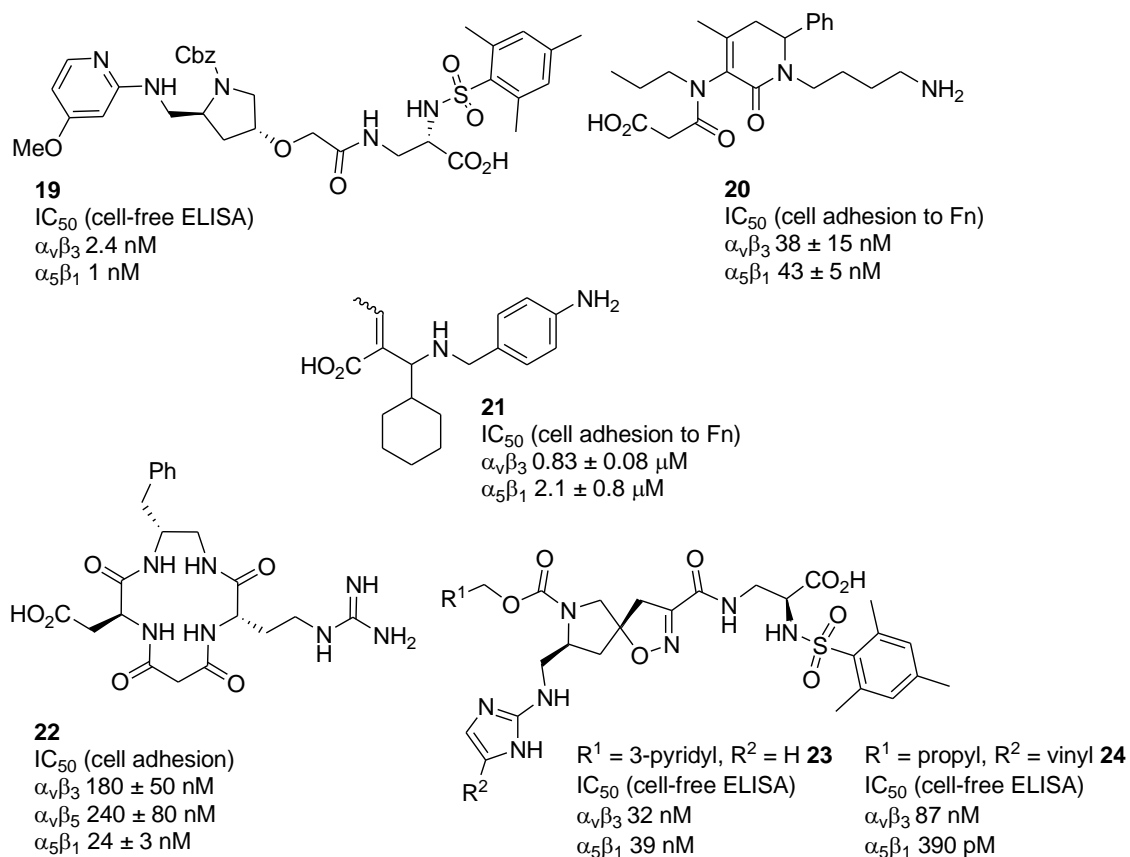
Tumors often overexpress  $\alpha_5\beta_1$  along with  $\alpha_v\beta_3$  and other  $\alpha_v$  integrins.<sup>91,92</sup> Both  $\alpha_v\beta_3$  and  $\alpha_5\beta_1$  allow interaction between cancer cells, endothelial cells and the ECM through common ligands, most notably fibronectin.<sup>93</sup>  $\alpha_v\beta_3$  has been shown to regulate the function of  $\alpha_5\beta_1$  and *vice versa*; selective inhibition of  $\alpha_v\beta_3$  allows  $\alpha_5\beta_1$  mediated adhesion but not migration, whereas inhibition of both integrins more effectively prevented cancer cell adhesion.<sup>94,95</sup>  $\alpha_5\beta_1$  binding fibronectin promoted endothelial cells binding to vitronectin, and angiogenesis.<sup>96</sup>

Inhibition of  $\beta_1$  integrins has been shown to promote tumor progression and metastasis by increasing the expression of  $\beta_3$  indicating that selective inhibition of  $\alpha_5\beta_1$  is likely to be ineffective as an anticancer therapy due to  $\beta_3$ -mediated resistance. However,  $\alpha_v\beta_3$  antagonism

or combination  $\beta_1/\beta_3$  antagonism is not vulnerable to this resistance.<sup>97</sup> In contrast,  $\alpha_v\beta_3$  antagonism can increase tumor cell invasiveness in the presence of high levels of fibronectin due to increased  $\alpha_5\beta_1$  recycling; the effect of dual  $\beta_1/\beta_3$  antagonism in these circumstances remains to be established.<sup>98</sup> Such changes in receptor expression and usage in response to the chemical environment of a tumor may explain the contradictory results obtained in a minority of pathological studies, which show decreased expression of  $\alpha_5\beta_1$  as disease progresses.

$\alpha_5\beta_1$  has a similar proangiogenic function to  $\alpha_v\beta_3$  and knock-out of both  $\alpha_5$  and  $\alpha_v$  is required to prevent vascular development since loss of one is generally compensated by the other.<sup>99</sup> Changes in the conformation of fibronectin favor binding by  $\alpha_v\beta_3$  rather than  $\alpha_5\beta_1$  in the tumor microenvironment and promote vascular endothelial growth factor (VEGF) secretion; selective  $\alpha_5\beta_1$  inhibition stimulated VEGF secretion, but  $\alpha_v\beta_3$  inhibition reduced it, suggesting that targeting  $\alpha_5\beta_1$  alone may have undesirable pro-angiogenic effects but targeting  $\alpha_5\beta_1/\alpha_v\beta_3$  should not.<sup>100</sup>

Selective  $\alpha_5\beta_1$  and  $\alpha_v\beta_3$  antibodies had little effect on angiogenesis in the stromal microenvironment as single agents, but combination of selective antibodies completely blocked endothelial tube formation.<sup>101</sup> Taken together, biological studies suggest that the safety and effectiveness of selective anti- $\alpha_5\beta_1$  therapy requires further evaluation, whereas targeting  $\alpha_5\beta_1$  in combination with  $\alpha_v\beta_3$  is likely to prove more efficient.



### Figure 7 $\alpha_v\beta_3/\alpha_5\beta_1$ antagonists

Pyrrolidine derivatives eg **19** (Figure 7) with sulfonamides on the diaminopropanoate moiety proved to be dual  $\alpha_v\beta_3/\alpha_5\beta_1$  antagonists with moderate to good selectivity over  $\alpha_v\beta_5$ . Increasing the lipophilicity of the sidechains increased anti-integrin activity non-selectively, whereas switching from sulfonamide to carbamate gave  $\alpha_5\beta_1$  selective antagonists which was rationalized by the greater flexibility of the sulfonamide group allowing it to adopt conformations to fit both integrins' binding pockets.<sup>102</sup>

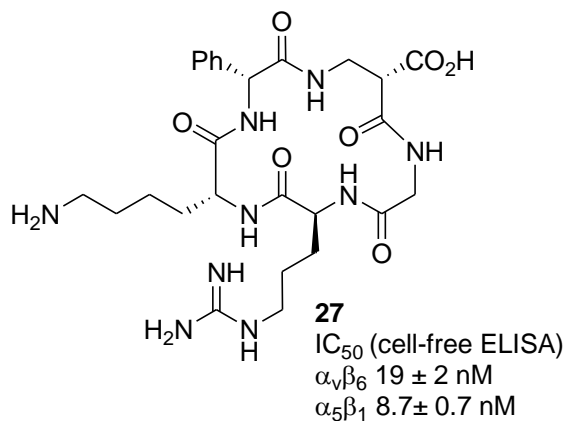
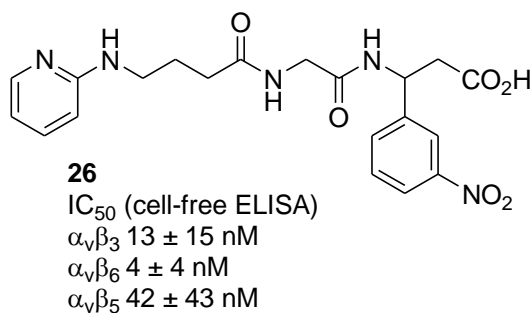
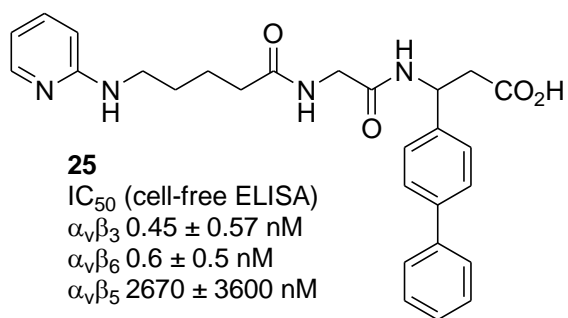
Dihydropyridinone antagonists were developed to provide more efficient antiangiogenic agents. These molecules lack the exosite binding motif seen in many RGD mimetic antagonists, and also contain very simple arginine mimetics. A primary amine was better than a guanidine group for activity on both receptors, and the most active compound **20** contained a simple butylamine chain.<sup>103</sup> Dehydro  $\beta$ -amino acid **21** with a similar primary amines also showed dual antagonist activity.<sup>104</sup>



Tetrapeptide mimetic **22** was also designed as an anti-angiogenic agent. A number of stereoisomers were effective in reducing the adhesion of cancer cells with similar levels of affinity for  $\alpha_v\beta_3$ ,  $\alpha_5\beta_1$  and  $\alpha_v\beta_5$ , however only those containing *S*-aspartic acid inhibited human vascular endothelial cell tube formation *in vitro*.<sup>105</sup>

Modifications of isoxazoline compounds to increase potency by introducing a spirocyclic linker gave selective or dual active  $\alpha_v\beta_3/\alpha_5\beta_1$  antagonists eg **23**. Selectivity could be changed to favor  $\alpha_5\beta_1$  by introducing a carbamate group to the portion of the molecule interacting with the  $\alpha$ -subunits.<sup>106</sup> A related molecule **24** was used as a dual targeting agent to selectively deliver nanoparticles to tumors. Dual targeting was more effective at reducing angiogenesis than nanoparticles targeting  $\alpha_v\beta_3$  alone which may be due to increased drug delivery when binding 2 receptors or to targeting a wider population of tumor and/or endothelial cells.<sup>107</sup>

**$\alpha_v\beta_6/\alpha_v\beta_3$  and  $\alpha_v\beta_6/\alpha_5\beta_1$**



### Figure 8 Dual $\alpha_v\beta_6$ antagonists

$\alpha_v\beta_6$  is upregulated during inflammation and cancer progression, particularly in colon, head and neck and pancreatic cancers.<sup>108</sup> It shares with  $\alpha_v\beta_3$  the ability to localize and activate matrix metalloproteinases (MMPs) and activate transforming growth factor- $\beta$ 1 (TGF- $\beta$ 1). There are few examples of small molecules targeting  $\alpha_v\beta_6$ ; the first selective antagonist was developed alongside dual and tri-integrin antagonists **25** and **26** (Figure 8). Sterically demanding aromatic sidechains on the aspartate sidechain mimetic reduce affinity for  $\alpha_v\beta_5$  while increasing it for  $\alpha_v\beta_6$ .<sup>109</sup> The cell-based adhesion assays used in this study provide a useful illustration of experimental design to selectively measure inhibition of a single integrin using a cell line expressing several RGD-binding integrins.

A cyclic peptide **27** with high affinity for  $\alpha_v\beta_6$  and  $\alpha_5\beta_1$  yet completely inactive against  $\alpha_v\beta_3$ ,  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_5$  has recently been developed.<sup>9</sup> Docking with a homology model of  $\alpha_v\beta_6$  suggested that the high affinity for  $\alpha_v\beta_6$  and selectivity over  $\alpha_v\beta_3$  is due to D-phenylglycine occupying a hydrophobic binding pocket in the  $\beta_6$  subunit which is larger than the corresponding pocket in  $\beta_3$  thus directing the lysine sidechain to form a salt bridge to the  $\beta_6$  subunit

### **Multi-integrin antagonism**

The majority of cell types express a range of integrins from several subfamilies on their surfaces. It is extremely unlikely that a single antagonist could affect the entire family of integrin receptors due to significant differences in structures of both  $\alpha$  and  $\beta$  subunits. The most obvious difference is the presence or absence of an I-domain in the  $\alpha$  subunit controlling the site and mode of ligand binding, but differences in structure of the cytoplasmic domains such as the long tail present in  $\beta_4$  also modulate integrin function and the binding of cell-penetrating molecules. Despite the differences in structure, it is possible to develop an antagonist with similar affinities for most or all members of a particular subfamily.

Recent development of rabbit monoclonal antibodies suitable for use in formalin-fixed paraffin-embedded tissue samples has paved the way for extensive studies of the expression of the  $\alpha_v$  integrin subfamily in primary tumors and metastases, with a particular focus on brain tumors.<sup>67</sup>  $\alpha_v\beta_3$ ,  $\alpha_v\beta_5$  and  $\alpha_v\beta_8$  are frequently coexpressed on gliomas, and increased gene expression is associated with decreased survival;  $\alpha_v\beta_5$  and  $\alpha_v\beta_8$  were more common in glioblastomas compared to less aggressive tumor types.<sup>110</sup> Evaluation of integrin expression in primary tumors and metastases to the brain revealed  $\alpha_v$  integrins were invariably present on tumors and upregulated in brain metastases with the specific heterodimers involved depending on the tumor type;  $\alpha_v\beta_5$  is present in most primary tumors,  $\alpha_v\beta_3$  and  $\alpha_v\beta_8$  are

upregulated in metastases compared to primary tumors. The relative levels of  $\alpha_v\beta_6$  in primary tumor and metastasis vary between studies.<sup>111,112</sup>

Based on these results, several suggestions have been made regarding appropriate combination of integrins to target;  $\alpha_v\beta_3/\alpha_v\beta_5$  antagonists will target blood vessels but not tumor cells and metastases which often express  $\alpha_v\beta_6$  and  $\alpha_v\beta_8$  as well as  $\alpha_v\beta_3$ .  $\alpha_v\beta_6/\alpha_v\beta_8$  antagonists may control primary tumor growth and metastasis without normal tissue toxicity, but  $\alpha_v\beta_8$  has not been as extensively studied as the other integrins thus its role in tumorigenesis requires further investigation. Overall, the range of integrins expressed on tumors suggests pan- $\alpha_v$  antagonism will be required for efficient prevention and treatment of metastases to the brain since it is the only approach that can target both tumors and the supporting stroma and endothelial cells.<sup>111,112</sup>

Pan- $\alpha_v$  antagonism should be broadly applicable in the treatment of metastasis.  $\alpha_v$  signalling has been shown to be highly significant in prostate tumor progression and dissemination. Knockdown of  $\alpha_v$  inhibited the growth of  $\alpha_v\beta_1/\alpha_v\beta_5$  expressing prostate cancer xenografts in bone through interfering with tumor-microenvironment interactions,<sup>113</sup> and reduced orthotopic tumor growth and bone metastasis initiated by intracardiac injection of prostate cancer stem cells.<sup>114</sup> Antibodies and small molecules targeting  $\alpha_v$  have demonstrated antimetastatic effects in a number of tumor types and are reviewed below.

The expression of  $\alpha_v\beta_1$  on tumors and metastases has not yet been investigated, and the studies described above do not consider the contribution of non- $\alpha_v$  RGD-binding integrins to tumor progression and metastasis.  $\alpha_5\beta_1$  and  $\alpha_{IIb}\beta_3$  respectively support angiogenesis and hematogenous metastasis, and can both promote tumor growth, thus should also be considered when designing multi-targeted antagonists.

### **Anti- $\alpha_v$ antibodies**

Antibodies binding all  $\alpha_v$  integrins have progressed to clinical trials. Intetumumab (CNTO95), an anti- $\alpha_v\beta_1/\alpha_v\beta_3/\alpha_v\beta_5/\alpha_v\beta_6$  antibody developed to target a wide range of cancer types, was effective at reducing or preventing tumor growth in preclinical models through both tumor specific effects on  $\alpha_v\beta_3/\alpha_v\beta_5$ , and more clinically relevant action on integrins on both the tumor and surrounding cells.<sup>115</sup> It also diminished hematogenous metastasis to the brain and lungs by reducing the ability of cells to migrate and invade from the vasculature to the metastatic site.<sup>116,117</sup> The anti-metastatic effect was independent of treatment timing, but treatment at the time of removing the primary tumor has been suggested to prevent the formation of metastases from tumor cells released by surgery.<sup>116</sup> Importantly for the development of further pan- $\alpha_v$  targeted drugs, intetumumab has been shown to be safe in both preclinical and clinical studies, despite binding to  $\alpha_v$  integrins in a wide range of normal tissues.<sup>118</sup>

In Phase I trials of intetumumab, partial responses occurred in patients with a number of cancer types including metastatic melanoma, angiosarcoma and ovarian cancer,<sup>119,120</sup> and a larger trial on melanoma showed anecdotal evidence of improved overall survival.<sup>121</sup> Despite encouraging results in a Phase I trial on metastatic prostate cancer, in Phase II, intetumumab did not improve progression free survival, but did reduce expression of biomarkers of bone turnover.<sup>122,123</sup> Although these results seem to refute the assertion that multi-integrin inhibition will be effective clinically they must be treated with caution since both low exposure to intetumumab resulting from unfavorable pharmacokinetics and varying levels of target expression in the patients enrolled have been noted during trials.<sup>121</sup> Use of a small molecule with better penetration and half-life in the body may provide more reliable evidence.

In preclinical studies, the humanised deimmunised anti- $\alpha_v$  antibody DI17E6 prevented the growth of melanoma xenografts in mice through acting on human  $\alpha_v\beta_3$  expressed on tumors.<sup>124</sup> DI17E6 was well-tolerated in Phase I trials in healthy subjects, and those with

progressing prostate cancer bone metastasis. Clinical trials in prostate cancer are ongoing after multiple patients on the Phase I trial showed stable disease or partial response.<sup>125,126</sup>

## **Disintegrins**

Snake venom disintegrins are a family of small proteins (typically 45–84 amino acids in length) that inhibit multiple integrins including the  $\alpha_v$  subfamily,  $\alpha_{IIb}\beta_3$ ,  $\alpha_5\beta_1$  and others in the  $\beta_1$  family. As a result of these wide and variable anti-integrin profiles, disintegrins have become useful as probe molecules (eg echistatin), and starting points for anticancer and antithrombotic drug development; the clinically used anti- $\alpha_{IIb}\beta_3$  cyclopeptide Eptifibatid was developed based on the KGD sequence of the  $\alpha_{IIb}\beta_3$  specific disintegrin barbourin.<sup>127,128</sup>

Some recent studies investigating the anticancer effects of disintegrins and other snake venom sourced molecules which bind multiple integrins are summarised in Table 2. The majority of these examples contain the RGD recognition sequence, however this is not a requirement for disintegrin activity. Other related sequences also confer high activity on the RGD-binding subfamily,<sup>129</sup> or promote binding to  $\alpha_4$  and  $\beta_1$  integrins. Not all disintegrins have been fully characterised for their integrin binding profile; many other examples<sup>130-132</sup> show interesting anti-platelet and anti-tumor effects.

Binding a wider range of integrins than many currently used selective antibodies and small molecules is likely to make disintegrins more efficient in controlling cancer growth and dissemination since they will be able to block multiple redundant adhesion mechanisms and signalling pathways. For example, acurhagin C inhibits growth of melanoma cell lines which are unaffected by the  $\alpha_v\beta_3/\alpha_v\beta_5$  antagonist cRGDFV.<sup>133</sup> Many disintegrins inhibit platelet aggregation, and thus tumor-platelet interactions involving  $\alpha_{IIb}\beta_3$ . They are notably effective at preventing the formation of lung metastases when coinjected intravenously with cancer cells.<sup>131,134,135</sup> Vicrostatin, an echistatin/contortrostatin chimera, is selective for activated

platelets and caused no bleeding side effects *in vivo*, suggesting anti- $\alpha_{IIb}\beta_3$  activity can be safely incorporated into an effective anti-metastatic agent.<sup>136</sup>

The clinical use of disintegrins presents some challenges. Naturally produced supply is limited, expensive and hazardous to collect, whereas recombinant disintegrins may have different properties to the naturally sourced material. The echistatin/contortrostatin chimera vicrostatin can be reliably produced recombinantly, avoiding difficulties in supplying contortrostatin in large quantities,<sup>136</sup> and contortrostatin itself has been produced recombinantly<sup>137</sup> and formulated as liposomes to improve half-life and facilitate intravenous administration.<sup>138</sup>

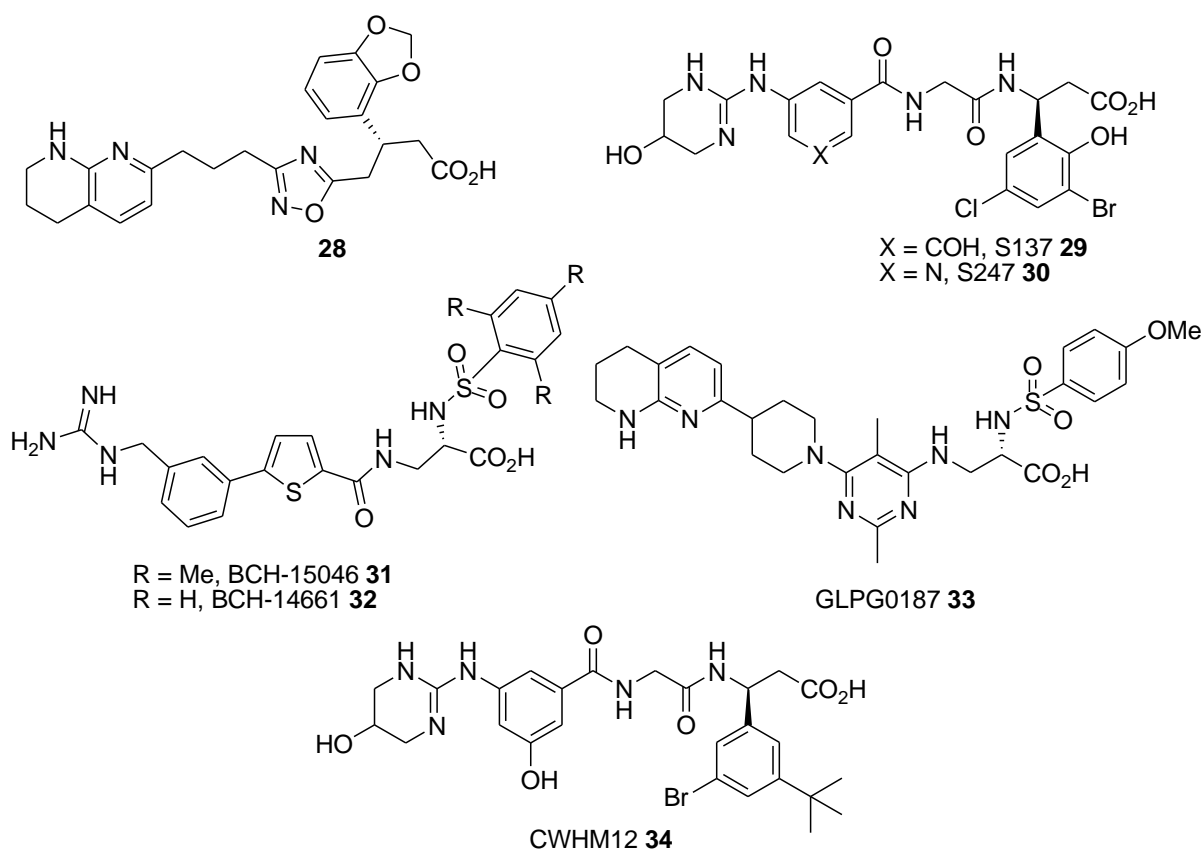
Preclinical experiments with disintegrins indicate that multi-antagonists of the RGD-binding integrins are effective, and generally safe, anticancer agents. Knowledge of disintegrin structures, specifically the conformation of the loop containing the recognition motif, and the nature of the amino acids flanking the RGD tripeptide will provide inspiration for small molecule development.

<b>Disintegrin</b>	<b>Integrins bound</b>	<b>Recognition sequence</b>	<b>Anticancer effects</b>	<b>Ref</b>
Salmosin	$\alpha_{IIb}\beta_3$ , $\alpha_v$ subfamily, $\beta_1$ subfamily	RGD	Reduces hematogenous metastasis. Blocks cell survival signalling.	<sup>134,139</sup>
Contortrostatin	$\alpha_{IIb}\beta_3$ , $\alpha_v\beta_3$ , $\alpha_v\beta_5$ , $\alpha_5\beta_1$	RGD	Anti-angiogenic. Reduces xenograft growth in bone. Reduces hematogenous metastasis. Synergistic with docetaxel	<sup>135,140</sup>
Vicrostatin	$\alpha_{IIb}\beta_3$ , $\alpha_v\beta_3$ , $\alpha_v\beta_5$ , $\alpha_5\beta_1$	RGD	Anti-angiogenic: disrupts endothelial cell cytoskeleton.	<sup>136</sup>

			Reduces tumor growth	
Acurhagin-C	$\alpha_v$ -subfamily (excluding $\alpha_v\beta_5$ ), $\alpha_5\beta_1$	ECD	Inhibits melanoma cell proliferation and migration. Synergistic with methotrexate	<sup>133</sup>
PIVL	$\alpha_v\beta_3$ , $\alpha_v\beta_5$ , $\alpha_v\beta_6$ , $\alpha_1\beta_1$ , $\alpha_5\beta_1$	Not determined	Inhibits glioblastoma cell invasion and migration	<sup>141</sup>
Eristostatin	$\alpha_{IIb}\beta_3$ , $\alpha_4\beta_1$ , $\alpha_v$ subfamily	RGD	Reduces hematogenous metastasis. Affects natural killer cell function and interaction with melanoma cells.	<sup>142</sup>
Viridistatin	$\alpha_{IIb}\beta_3$ , $\alpha_v\beta_3$ , other $\alpha_v$ subfamily	RGD	Inhibits tumor cell invasion and migration. Reduces hematogenous metastasis.	<sup>131</sup>

**Table 2 Recent examples of disintegrins with anti-cancer activity**





**Figure 9 Multi-integrin antagonists**

	$\alpha_v\beta_1$	$\alpha_v\beta_3$	$\alpha_v\beta_5$	$\alpha_v\beta_6$	$\alpha_v\beta_8$	$\alpha_5\beta_1$	$\alpha_2\beta_1$	$\alpha_4\beta_7$	$\alpha_{10}\beta_1$	$\alpha_{IIb}\beta_3$	Ref
<b>28</b>	2.6 <sup>a</sup>	0.65 <sup>a</sup>	1.7 <sup>a</sup>	2660 <sup>a</sup>	NT	NT	NT	NT	NT	27 <sup>b</sup>	143
S137 <b>29</b>	10.4 ± 7.3 <sup>b</sup>	1.6 ± 1 <sup>b</sup>	18.1 ± 14.6 <sup>b</sup>	11.7 ± 10.7 <sup>b</sup>	1.71 ± 0.05 <sup>b</sup>	316 ± 228 <sup>b</sup>	NT	NT	NT	257 ± 29 <sup>b</sup>	144
S247 <b>30</b>	1.59 ± 0.41 <sup>b</sup>	0.40 ± 0.24 <sup>b</sup>	1.50 ± 1.26 <sup>b</sup>	1.13 ± 0.55 <sup>b</sup>	0.65 ± 0.21 <sup>b</sup>	64 ± 5 <sup>b</sup>	NT	NT	NT	380 ± 92 <sup>b</sup>	144
BCH- 15046 <b>31</b>	NT	3.5 ± 0.9 <sup>a</sup>	60 ± 50 <sup>a</sup>	NT	NT	20 ± 4 <sup>a</sup>	NT	NT	NT	0.2 <sup>b</sup>	145

BCH- 14661 <b>32</b>	NT	0.033 ± 0.016 <sup>a</sup>	72 ± 37 <sup>a</sup>	NT	NT	640 ± 256 <sup>a</sup>	NT	NT	NT	0.3 <sup>b</sup>	145
SC56631 <b>3</b>	317 <sup>a</sup>	10 <sup>b</sup> 120 <sup>a</sup>	23 <sup>a</sup>	NT	NT	~70000 <sup>a</sup>	NT	NT	NT	9 <sup>b</sup>	65
GLPG0187 <b>33</b>	1.3 ± 0.1 <sup>b</sup>	3.7 ± 0.6 <sup>b</sup>	2.0 ± 0.6 <sup>b</sup>	1.4 ± 0.3 <sup>b</sup>	1.2 ± 0.3 <sup>b</sup>	7.7 ± 4.0 <sup>b</sup>	>10 <sup>4,b</sup>	>10 <sup>4,b</sup>	NT	>10 <sup>5,b</sup>	146
CWHM12 <b>34</b>	1.8 ± 0.9 <sup>b</sup>	0.8 ± 0.6 <sup>a</sup>	61 ± 17 <sup>a</sup>	1.5 ± 2.5 <sup>a</sup>	0.2 ± 0.1 <sup>b</sup>	NR <sup>a</sup>	>5000 <sup>b</sup>	NT	>5000 <sup>b</sup>	>5000 <sup>b</sup>	147

**Table 3.** IC<sub>50</sub> (nM) of multi-RGD antagonists in cell-based (<sup>a</sup>) and cell-free (<sup>b</sup>) ELISA. NT = not tested. NR = tested but not reported.

A smaller number of small molecule multi-RGD antagonists have been reported compared with  $\alpha_v\beta_3$  or  $\alpha_v\beta_3/\alpha_v\beta_5$  targeted agents, although complete testing for selectivity may reveal more; for example screening ‘selective’  $\alpha_v\beta_3$  antagonists eg. **28** and SC56631 **3** revealed activity on  $\alpha_v\beta_1$  as well as  $\alpha_v\beta_5$ . In this regard the orally bioavailable S137 **29** was described initially as an  $\alpha_v\beta_3$  antagonist, but proved to be a pan- $\alpha_v$  antagonist with highest activity against  $\alpha_v\beta_3/\alpha_v\beta_8$ . Switching to a pyridine ring system gave S247 **30** which has increased affinity for all  $\alpha_v$  integrins and measurable activity against  $\alpha_5\beta_1$ .<sup>144</sup> S247 **30** decreased liver metastasis from established colon cancer splenic xenografts, and continuous infusion reduced lung metastasis from orthotopic implantation of an aggressive breast cancer cell line although it did not reduce growth of the primary tumor in either case.<sup>144,148</sup> In some studies, S247 **30** appeared to exert most effect on the early stages of metastasis,<sup>149</sup> however S137 **29** reduced metastatic tumor burden when administered orally after the primary tumor was removed.<sup>144</sup> Since this mimics the situation of patients with advanced cancer following surgery, it suggests anti- $\alpha_v$  agents will be useful as adjuvant or palliative therapy.

The anti-angiogenic BCH-15046 **31** was shown to be more effective in a range of *in vitro* angiogenesis models than the related  $\alpha_v\beta_3$  selective compound BCH-14461 **32** as a result of its ability to prevent both MMP-dependent and collagen-dependent cell proliferation and survival which require  $\alpha_5\beta_1$ .<sup>145</sup> The high activity of both **31** and **32** against  $\alpha_{IIb}\beta_3$  suggests they would show further improved antitumor effects *in vivo*. GLPG0187 is a nanomolar antagonist of 6 integrins developed as a treatment for bone metastasis in breast and prostate cancers;<sup>146,150</sup> its structure has not been reported, but patent analysis suggests **33**. In preclinical studies, GLPG0187 has been shown to be a potent anti-angiogenic and anti-osteoporotic agent, effective in reducing both new and established bone metastases.<sup>151,152</sup> It was shown to be safe and effective at reducing biomarkers of bone turnover in a Phase I trial in healthy volunteers, and has now progressed into trials in patients with advanced cancers. An isopropyl malonate prodrug to increase oral bioavailability is also being investigated.<sup>153</sup> CWHM12 **34** is a nanomolar antagonist of all  $\alpha_v$  integrins structurally related to S137 **29**. CWHM12 **34** has not been investigated in cancer, but was effective in preventing and treating renal and pulmonary fibrosis.<sup>147</sup> Cilengitide was not effective here, suggesting that blockade of  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  alone is not sufficient to have therapeutic effect in fibrosis perhaps because it leaves other  $\alpha_v$  integrins able to activate TGF- $\beta$ 1. The effects of subtle differences in integrin affinity between GLPG0187 and CWHM12 **34**, most notably on  $\alpha_v\beta_5$  and  $\alpha_5\beta_1$ , remain to be investigated.

## Discussion

Despite the large number of integrin antagonists reported in the medicinal chemistry literature, a very small number of agents have progressed successfully into the clinic. Multiple hypotheses have been proposed to explain the lack of efficacy seen in clinical trials of agents such as Cilengitide, ranging from reversibly binding antagonists promoting changes in integrin receptor conformation and signalling at low concentrations to issues of selectivity between cell types. In particular, lack of selectivity between the members of an integrin

subfamily has been suggested as a safety issue complicating clinical applications of targeted therapy; to overcome this, the use of highly selective small molecules to provide a tightly aimed targeting of a single integrin expressed by a particular cancer has been proposed.<sup>86</sup>

The use of small molecules targeted to specific integrins as carriers may be useful in delivering cytotoxic agents to cancer cells. However, it does not directly target the integrin mediated processes which are key to tumor proliferation and dissemination, will require the development of a large number of individual targeting agents, and is still liable to failure as cancer cells alter their integrin expression in response to changes in the surroundings or to drug treatment. In contrast, a single small molecule multi-integrin antagonist will provide an anticancer agent applicable to a wide range of tumors which is not vulnerable to the development of resistance or paradoxical effects by changes in expression of particular members of an integrin subfamily. Studies using disintegrins and anti- $\alpha_v$  antibodies have shown that multi-integrin antagonism is likely to be both safe and effective; the ability to bind integrins on multiple cell types does not cause unacceptable normal tissue toxicities, and should result in improved anticancer effects through inhibiting a wide range of tumor and stromal cell interactions with the microenvironment. Further work is required to determine the optimum combination(s) of integrins that should be targeted by an efficient multi-antagonist. A number of the RGD-binding subfamily,  $\alpha_v\beta_3$ ,  $\alpha_v\beta_5$  and  $\alpha_v\beta_6$ , are already well-established anticancer targets. The biological functions of others, such as  $\alpha_v\beta_8$  and  $\alpha_8\beta_1$ , still largely remain to be defined. Both  $\alpha_v$  and  $\beta_1$  are widely expressed in cells, and the  $\alpha_v\beta_1$  heterodimer may become more important as other related receptors are inhibited. Antagonism of  $\alpha_{IIb}\beta_3$  is frequently a cause for concern due to its involvement in platelet aggregation and hence the possibility of vascular bleeding as a serious side effect. Early selective  $\alpha_{IIb}\beta_3$  antagonists did have a poor safety profile; however, it is noteworthy that the few small molecule dual  $\alpha_{IIb}\beta_3/\alpha_v\beta_3$  antagonists investigated to date do not cause prolongation of bleeding time or related side effects. That accepted, the potential benefit of  $\alpha_{IIb}\beta_3/\alpha_v\beta_3$

antagonism on preventing life-threatening tumor associated thrombus embolism should not be underestimated. Given the great potential of such compounds in combating tumor dissemination, it is important that development continues and  $\alpha_{IIb}\beta_3$  is considered for inclusion as part of the target profile for multi-integrin antagonists.

The medicinal chemistry literature contains a vast number of integrin antagonists. Most of these are selective and designed as antagonists of a single receptor. However, their selectivity has usually been determined against only 2 or 3 related integrins. Full screening of compounds against an entire subfamily is required to determine whether an antagonist is genuinely selective or an unrecognised dual or multi-antagonist. This knowledge will improve the interpretation of biological results obtained using integrin antagonists as probe molecules, and provide structure-activity data to assist with the design of further antagonists with defined selectivity. Combined with full analysis of the integrin expression profile in different diseases, and how it changes in response to drug treatments, such investigations will provide lead molecules not just for cancer, but also other diseases involving angiogenesis, atherosclerosis, fibrosis, and viral infections.

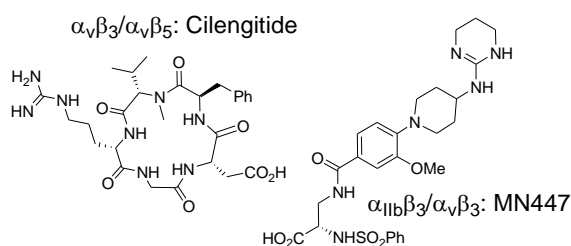
**Corresponding author information:** Email: [h.sheldrake@bradford.ac.uk](mailto:h.sheldrake@bradford.ac.uk). Telephone: 01274 236858

**Abbreviations:** ECM extracellular matrix; ELISA enzyme-linked immunosorbent assay; MMP matrix metalloproteinase; VEGF vascular endothelial growth factor.

**Author biography:** Helen M. Sheldrake read Natural Sciences at the University of Cambridge, U.K., and subsequently received a Ph.D. in Organic Chemistry with Dr Jonathan Burton. After postdoctoral work with Dr Tim Wallace at The University of Manchester, she took up a RCUK Academic Fellowship in Medicinal Chemistry at the Institute of Cancer Therapeutics, University of Bradford where she leads a team focussing on the development of novel dual/multi-integrin antagonists to prevent cancer dissemination.

Laurence H Patterson has over 30 years' experience in pharma and academia undertaking small molecule drug discovery and mechanisms of action. He has been employed at Fisons Pharma (now AZ), School of Pharmacy, DeMontfort University and The School of Pharmacy, University of London. He is a founding director of Biostatus Ltd and a founding shareholder of Incanthera Ltd. In his present role as Director of the Institute of Cancer Therapeutics he has instigated and oversees a portfolio of drug discovery projects focused on tumor selective chemotherapy and antimetastatic agents. He earned his BSc (hons) at Hatfield Polytechnic/Atomic Energy Research Establishment, Harwell, Oxon and PhD at Chelsea School of Pharmacy, King's College, London with John Gorrod.

Table of Contents graphic:



- (1) J. D. Humphries; A. Byron; M. J. Humphries. Integrin ligands at a glance. *J. Cell Science* **2006**, *119*, 3901-3903.
- (2) T. Xiao; J. Takagi; B. S. Coller; J.-H. Wang; T. A. Springer. Structural Basis for Allostery in Integrins and Binding to Fibrinogen-mimetic Therapeutics. *Nature* **2004**, *432*, 59-67.
- (3) M. Nagae; S. Re; E. Mihara; T. Nogi; Y. Sugita; J. Takagi. Crystal structure of  $\alpha_5\beta_1$  integrin ectodomain: Atomic details of the fibronectin receptor. *J. Cell Biol.* **2012**, *197*, 131-140.
- (4) M. Pfaff; K. Tangemann; M. Müller; M. Gurrath; G. Müller; H. Kessler; R. Timpl; J. Engel. Selective Recognition of Cyclic RGD Peptides of NMR Defined Conformation by  $\alpha_{IIb}\beta_3$ ,  $\alpha_v\beta_3$ , and  $\alpha_5\beta_1$  Integrins. *J. Biol. Chem.* **1994**, *269*, 20233-20238.
- (5) C. E. Peishoff; F. E. Ali; J. W. Bean; R. Calve; C. A. D'Ambroeo; D. S. Weston; Hwang, S. M.; T. P. Kline; P. F. Koeter; A. Nichols; D. Powers; T. Romoff; J. M. Samanen; J. Stade; J. A. Vasko; K. D. Kopple. Investigation of Conformational Specificity at GPIIb/IIIa: Evaluation of Conformationally Constrained RGD Peptidest. *J. Med. Chem.* **1992**, *35*, 3962-3969.
- (6) L. Marinelli; A. Lavecchia; K.-E. Gottschalk; E. Novellino; H. Kessler. Docking Studies on  $\alpha_v\beta_3$  Integrin Ligands: Pharmacophore Refinement and Implications for Drug Design. *J. Med. Chem.* **2003**, *46*, 4393-4404.

- (7) L. Marinelli; A. Meyer; D. Heckmann; A. Lavecchia; E. Novellino; H. Kessler. Ligand Binding Analysis for Human  $\alpha_5\beta_1$  Integrin: Strategies for Designing New  $\alpha_5\beta_1$  Integrin Antagonists. *J. Med. Chem.* **2005**, *48*, 4204-4207.
- (8) L. Marinelli; K. E. Gottschalk; A. Meyer; E. Novellino; H. Kessler. Human integrin  $\alpha_v\beta_5$ : homology modeling and ligand binding. *J. Med. Chem.* **2004**, *47*, 4166-4177.
- (9) A. Bochen; U. Kiran Marelli; E. Otto; D. Pallarola; C. Mas-Moruno; F. Saverio Di Leva; H. Boehm; J. P. Spatz; E. Novellino; H. Kessler; L. Marinell. Biselectivity of isoDGR Peptides for Fibronectin Binding Integrin Subtypes  $\alpha_5\beta_1$  and  $\alpha_v\beta_6$ : Conformational Control through Flanking Amino Acids. *J. Med. Chem.* **2013**, *56*, 1509-1519.
- (10) G. Sowmya; J. M. Khan; S. Anand; S. B. Ahn; M. S. Baker; S. Ranganathan. A site for direct integrin  $\alpha_v\beta_6$ •uPAR interaction from structural modelling and docking. *J. Struct. Biol.* **2014**, DOI:10.1016/j.jsb.2014.01.001.
- (11) D. A. Calderwood; V. Tai; G. Di Paolo; P. De Camilli; M. H. Ginsberg. Competition for Talin Results in Trans-dominant Inhibition of Integrin Activation. *J. Biol. Chem.* **2004**, *279*, 28889-28895.
- (12) D. Cox; M. Brennan; N. Moran. Integrins as therapeutic targets: lessons and opportunities. *Nature Rev. Drug Discovery* **2010**, *9*, 804-820.
- (13) S. L. Goodman; M. Picard. Integrins as therapeutic targets. *Trends Pharm. Sci.* **2012**, *33*, 405-412.
- (14) K. Nadrah; M. S. Dolenc. Dual antagonists of integrins. *Curr. Med. Chem.* **2005**, *12*, 1449-1466.
- (15) R. J. Davenport; J. R. Munday. Alpha4-integrin antagonism - an effective approach for the treatment of inflammatory diseases? *Drug Discovery Today* **2007**, *12*, 569-576.



- (16) J. S. Bennett. Structure and Function of the Platelet Integrin  $\alpha_{IIb}\beta_3$ . *J. Clin. Invest.* **2005**, *115*, 3363-3369.
- (17) R. M. Scarborough; D. D. Gretler. Platelet Glycoprotein IIb-IIIa Antagonists as Prototypical Integrin Blockers: Novel Parenteral and Potential Oral Antithrombotic Agents. *J. Med. Chem.* **2000**, *43*, 3453-3473.
- (18) S. A. Cohen; M. Trikha; M. A. Mascelli. Potential Future Clinical Applications for the GPIIb/IIIa Antagonist, Abciximab in Thrombosis, Vascular and Oncological Indications. *Pathol. Onc. Res.* **2000**, *6*, 163-174.
- (19) E. J. Topol; R. M. Califf; H. F. Weisman; S. G. Ellis; J. E. Tchong; S. Worley; R. Ivanhoe; B. S. George; D. Fintel; M. Weston; K. Sigmon; K. M. Anderson; K. L. Lee; J. T. Willerson. Randomised Trial of Coronary Intervention with Antibody Against Platelet IIb/IIIa Integrin for Reduction of Clinical Restenosis: Results at Six Months. *Lancet* **1994**, *343*, 881-886.
- (20) The IMPACT-II Investigators. Randomised Placebo-Controlled Trial of the Effect of Epifibatide on Complications of Percutaneous Coronary Intervention: IMPACT-II. *Lancet* **1997**, *349*, 1422-1428.
- (21) F. P. Ross; J. Chappel; J. I. Alvarez; D. Sander; W. T. Butler; M. C. Farach-Carson; K. A. Mintz; P. Gehron Robey; S. L. Teitelbaum; D. A. Cheresch. Interactions between the Bone Matrix Proteins Osteopontin and Bone Sialoprotein and the Osteoclast Integrin  $\alpha_v\beta_3$  Potentiate Bone Resorption. *J. Biol. Chem.* **1993**, *268*, 9901-9907.
- (22) P. C. Brooks; A. M. P. Montgomery; M. Rosenfeld; R. A. Reisfeld; T. Hu; G. Klier; D. A. Cheresch. Integrin  $\alpha_v\beta_3$  Antagonists Promote Tumor Regression by Inducing Apoptosis of Angiogenic Blood Vessels. *Cell* **1994**, *79*, 1157-1164.
- (23) V. Steri; T. S. Ellison; A. M. Gontarczyk; K. Weilbaecher; J. G. Schneider; D. Edwards; M. Fruttiger; K. M. Hodivala-Dilke; S. D. Robinson. Acute Depletion of

Endothelial  $\beta_3$ -Integrin Transiently Inhibits Tumor Growth and Angiogenesis in Mice. *Circ. Res.* **2014**, *114*, 79-91.

- (24) R. Max; R. R. C. M. Gerritsen; P. T. G. A. Nooijen; S. L. Goodman; A. Sutter; U. Keilholz; D. J. Ruiter; R. M. W. De Waal. Immunohistochemical Analysis of  $\alpha_v\beta_3$  Expression on Tumour Associated Vessels of Human Carcinomas. *Int. J. Cancer* **1997**, *71*, 320-324.
- (25) G. Gasparini; P. C. Brooks; E. Biganzoli; P. B. Vermeulen; E. Bonoldi; L. Y. Dirix; G. Ranieri; R. Miceli; D. A. Cheresh. Vascular Integrin  $\alpha_v\beta_3$ : A New Prognostic Indicator in Breast Cancer. *Clin. Cancer Res.* **1998**, *4*, 2625-2634.
- (26) H. M. Sheldrake; L. H. Patterson. Function and antagonism of  $\beta_3$  integrins in the development of cancer therapy. *Curr. Cancer Drug Targets* **2009**, *9*, 519-540.
- (27) S. M. Albelda; S. A. Mette; D. E. Elder; R. Stewart; L. Damjanovich; M. Herlyn; C. A. Buck. Integrin Distribution in Malignant Melanoma: Association of the  $\beta_3$  Subunit with Tumor Progression. *Cancer Res.* **1990**, *50*, 6757-6764.
- (28) G. Malik; L. M. Knowles; R. Dhir; S. Xu; S. Yang; E. Ruoslahti; J. Pilch. Plasma Fibronectin Promotes Lung Metastasis by Contributions to Fibrin Clots and Tumor Cell Invasion. *Cancer Res.* **2010**, *70*, 4327-4334.
- (29) X. P. Duan; S. F. Jia; Z. C. Zhou; R. R. Langley; M. F. Bolontrade; E. S. Kleinerman. Association of  $\alpha_v\beta_3$  Integrin Expression with the Metastatic Potential and Migratory and Chemotactic Ability of Human Osteosarcoma Cells. *Clin. Exp. Metastasis* **2004**, *21*, 747-753.
- (30) N. P. McCabe; S. De; A. VasANJI; J. Brainard; T. V. Byzova. Prostate Cancer Specific Integrin  $\alpha_v\beta_3$  Modulates Bone Metastatic Growth and Tissue Remodeling. *Oncogene* **2007**, *26*, 6238-6243.

- (31) L. M. Knowles; L. A. Gurski; C. Engel; J. R. Gnarra; J. K. Maranchie; J. Pilch. Integrin  $\alpha_v\beta_3$  and fibronectin upregulate Slug in cancer cells to promote clot invasion and metastasis. *Cancer Res.* **2013**, *73*, 6175-6184.
- (32) J. G. Schneider; S. H. Amend; K. N. Weilbaecher. Integrins and bone metastasis: Integrating tumor cell and stromal cell interactions. *Bone* **2011**, *48*, 54-65.
- (33) Y. Zhao; R. Bachelier; I. Treilleux; P. Pujuguet; O. Peyruchand; R. Baron; P. Clement-Lacroix; P. Clezardin. Tumor  $\alpha_v\beta_3$  Integrin Is a Therapeutic Target for Breast Cancer Bone Metastases. *Cancer Res.* **2007**, *67*, 5821-5830.
- (34) E. K. Sloan; N. Pouliot; K. L. Stanley; J. Chia; J. M. Moseley; D. K. Hards; R. L. Anderson. Tumor-specific expression of  $\alpha_v\beta_3$  integrin promotes spontaneous metastasis of breast cancer to bone. *Breast Cancer Res.* **2006**, *8*, R20.
- (35) I. Pecheur; O. Peyruchaud; C.-M. Serre; J. Guglielmi; C. Voland; F. Bourre; C. Margue; M. Cohen-Solal; A. Buffet; P. Clezardin. Integrin  $\alpha_v\beta_3$  Expression Confers on Tumor Cells a Greater Propensity to Metastasize to Bone. *FASEB J.* **2002**, *16*, 1266-1268.
- (36) M. Trikha; J. Timar; A. Zacharek; J. A. Nemeth; Y. Cai; B. Dome; B. Somlai; E. Raso; A. Ladanyi; K. V. Honn. Role for  $\beta_3$  Integrins in Human Melanoma Growth and Survival. *Int. J. Cancer* **2002**, *101*, 156-167.
- (37) J. Timar; M. Trikha; K. Szekeres; R. Bazaz; K. Honn. Expression and Function of the High Affinity  $\alpha_{IIb}\beta_3$  Integrin in Murine Melanoma Cells. *Clin. Exp. Metastasis* **1998**, *16*, 437-445.
- (38) J. Pontes-Júnior; S. T. Reis; L. C. Neves de Oliveira; A. C. Sant'Anna; Dall'Oglio, M. F.; A. A. Antunes; L. Ribeiro-Filho; P. A. Carvalho; J. Cury; M. Srougi; K. R. Moreira Leite. Association Between Integrin Expression and Prognosis in Localized Prostate Cancer. *Prostate* **2010**, *70*, 1189-1195.

- (39) B. Dome; E. Raso; J. Dobos; L. Meszaros; N. Varga; L. G. Puskas; L. Z. Feher; T. Lorincz; A. Ladanyi; M. Trikha; K. V. Honn; J. Timar. Parallel Expression of  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_3$  Integrins in Human Melanoma Cells Upregulates bFGF Expression and Promotes their Angiogenic Phenotype. *Int. J. Cancer* **2005**, *116*, 27-35.
- (40) L. Oleksowicz; Z. M. E. Schwartz; M. Khorshidit; J. P. Dutcher; E. Puszkin. Characterisation of Tumor-Induced Platelet Aggregation: The Role of Immunoregulated GPIIb and GPIIb/IIIa Expression by MCF-7 Breast Cancer Cells. *Thromb. Res.* **1995**, *79*, 261-274.
- (41) D. Buergy; F. Wenz; C. Groden; M. A. Brockmann. Tumor-platelet interaction in solid tumors. *Int. J. Cancer* **2012**, *130*, 2747–2760.
- (42) A. S. Lonsdorf; B. F. Krämer; M. Fahrleitner; T. Schönberger; S. Gnerlich; S. Ring; S. Gehring; S. W. Schneider; M. J. Kruhlak; S. G. Meuth; B. Nieswandt; M. Gawaz; A. H. Enk; H. F. Langer. Engagement of  $\alpha_{IIb}\beta_3$  (GPIIb/IIIa) with  $\alpha_v\beta_3$  mediates interaction of melanoma cells with platelets – a connection to hematogenous metastasis. *J. Biol. Chem.* **2011**, *287*, 2168-2178.
- (43) R. Dardik; Y. Kaufmann; N. Savion; N. Rosenberg; B. Shenkman; D. Varon. Platelets Mediate Tumor Cell Adhesion to the Subendothelium under Flow Conditions: Involvement of Platelet GPIIb/IIIa and Tumor Cell  $\alpha_v$  Integrins. *Int. J. Cancer* **1997**, *70*, 201-207.
- (44) J. S. Palumbo; K. E. Talmage; J. V. Massari; C. M. La Jeunesse; M. J. Flick; K. W. Kombrinck; M. Jirouskova; J. L. Degen. Platelets and Fibrin(ogen) Increase Metastatic Potential by impeding Natural Killer Cell-mediated Elimination of Tumor Cells. *Blood* **2005**, *105*, 178-185.
- (45) F. Zhao; L. Li; L. Guan; H. Yang; C. Wu; Y. Liu. Roles for GPIIb/IIIa and  $\alpha_v\beta_3$  integrins in MDA-MB-231 cell invasion and shear flow-induced cancer cell mechanotransduction. *Cancer Lett.* **2014**, *344*, 62-73.

- (46) M. Trikha; Z. Zhou; J. Timar; E. Raso; M. Kennel; E. Emmell; M. Nakada. Multiple Roles for Platelet GPIIb/IIIa and  $\alpha_v\beta_3$  Integrins in Tumor Growth, Angiogenesis, and Metastasis. *Cancer Res.* **2002**, *62*, 2824-2833.
- (47) O. Engebraaten; M. Trikha; S. Juell; S. Garman-Vik; O. Fodstad. Inhibition of In Vivo Tumour Growth by the Blocking of Host  $\alpha_v\beta_3$  and  $\alpha_{IIb}\beta_3$  Integrins. *Anticancer Res.* **2009**, *29*, 131-137.
- (48) W. Zhang; S. Dang; T. Hong; J. Tang; J. Fan; D. Bu; Y. Sun; Z. Wang; T. Wisniewski. A humanized single-chain antibody against  $\beta_3$  integrin inhibits pulmonary metastasis by preferentially fragmenting activated platelets in the tumor microenvironment. *Blood* **2012**, *120*, 2889-2898.
- (49) A. Amirkhosravi; M. Amaya; F. Siddiqui; J. P. Biggerstaff; T. V. Meyer; J. L. Francis. Blockade of GpIIb/IIIa Inhibits the Release of Vascular Endothelial Growth Factor (VEGF) from Tumor Cell-Activated Platelets and Experimental Metastasis. *Platelets* **1999**, *10*, 285-292.
- (50) N. Gomes; J. Vassy; C. Lebos; B. Arbeille; C. Legrand; F. Fauvel-Lefevre. Breast Adenocarcinoma Cell Adhesion to the Vascular Subendothelium in Whole Blood and under Flow Conditions: Effects of  $\alpha_v\beta_3$  and  $\alpha_{IIb}\beta_3$  Antagonists. *Clin. Exp. Metastasis* **2004**, *21*, 553-561.
- (51) Y. Liu; F. Zhao; W. Gu; H. Yang; Q. Meng; Y. Zhang; H. Yang; Q. Duan. The Roles of Platelet GPIIb/IIIa and  $\alpha_v\beta_3$  Integrins during HeLa Cells Adhesion, Migration, and Invasion to Monolayer Endothelium under Static and Dynamic Shear Flow. *J. Biomed. Biotech.* **2009**, *2009*, 829243.
- (52) D. G. Batt; J. J. Petraitis; G. C. Houghton; D. P. Modi; G. A. Cain; M. H. Corjay; S. A. Mousa; P. J. Bouchard; M. S. Forsythe; P. P. Harlow; F. A. Barbera; S. M. Spitz; R. R. Wexler; P. K. Jadhav. Disubstituted Indazoles as Potent Antagonists of the Integrin  $\alpha_v\beta_3$ . *J. Med. Chem.* **2000**, *43*, 41-58.

- (53) B. L. De Corte; W. A. Kinney; L. Liu; S. Ghosh; L. Brunner; W. J. Hoekstra; R. J. Santulli; R. W. Tuman; J. Baker; C. Burns; J. C. Proost; B. A. Tounge; B. P. Damiano; B. E. Maryanoff; D. L. Johnson; R. A. J. Galemmo. Piperidine-containing  $\beta$ -Arylpropionic Acids as Potent Antagonists of  $\alpha_v\beta_3/\alpha_v\beta_5$  Integrins. *Bioorg. Med. Chem. Lett.* **2004**, *14*, 5227-5232.
- (54) F. Osterkamp; B. Ziemer; U. Koert; M. Weisner; P. Raddatz; S. L. Goodman. Synthesis and Biological Evaluation of Integrin Antagonists Containing *trans*- and *cis*-2,5-Disubstituted THF Rings. *Chem. Eur. J.* **2000**, *6*, 666-683.
- (55) P. L. Barker; S. Bullens; S. Bunting; D. J. Burdick; K. S. Chan; T. Deisher; C. Eigenbrot; T. R. Gadek; R. Gantzog; M. T. Lipari; C. D. Muir; M. A. Napier; R. M. Pitti; A. Padua; C. Quan; M. Stanley; M. Struble; J. Y. K. Tom; J. P. Burnier. Cyclic RGD Peptide Analogues as Antiplatelet Antithrombotics. *J. Med. Chem.* **1992**, *35*, 2040-2048.
- (56) H. Matsuno; J. M. Stassen; J. Vermynen; H. Deckmyn. Inhibition of integrin function by a cyclic RGD-containing peptide prevents neointima formation. *Circulation* **1994**, *90*, 2203-2206.
- (57) T. J. A. Chico; J. Chamberlain; J. Gunn; N. Arnold; S. L. Bullens; T. R. Gadek; S. E. Francis; S. Bunting; M. Horton; L. Shepherd; M. T. Lipari; C. Quan; J. Knolle; H. U. Stiliz; A. Peyman; D. C. Crossman. Effect of Selective or Combined Inhibition of Integrins  $\alpha_{IIb}\beta_3$  and  $\alpha_v\beta_3$  on Thrombosis and Neointima After Oversized Porcine Coronary Angioplasty. *Circulation* **2001**, *103*, 1135-1141.
- (58) J. A. Menendez; L. Vellon; I. Mehmi; P. K. Teng; D. W. Griggs; R. Lupu. A Novel CYR61-triggered 'CYR61- $\alpha_v\beta_3$  Integrin Loop' Regulates Breast Cancer Cell Survival and Chemosensitivity through Activation of ERK1/ERK2 MAPK Signaling Pathway. *Oncogene* **2005**, *24*, 761-779.

- (59) J. S. Kerr; R. S. Wexler; S. A. Mousa; C. S. Robinson; E. J. Wexler; S. Mohamed; M. E. Voss; J. J. Devenny; P. M. Czerniak; A. Gudzelak; A. M. Slee. Novel Small Molecule  $\alpha_v$  Integrin Antagonists: Comparative Anti-cancer Efficacy with Known Angiogenesis Inhibitors. *Anticancer Res.* **1999**, *19*, 959-968.
- (60) F. B. Davis; H.-Y. Tang; A. Shih; T. Keating; L. Lansing; A. Hercbergs; R. A. Fenstermaker; A. Mousa; S. A. Mousa; P. J. Davis; H.-Y. Lin. Acting via a Cell Surface Receptor, Thyroid Hormone Is a Growth Factor for Glioma Cells. *Cancer Res.* **2006**, *66*, 7270-7275.
- (61) D. Kubota; M. Ishikawa; M. Yamamoto; S. Murakami; M. Hachisu; K. Katano; K. Ajito. Tricyclic pharmacophore-based molecules as novel integrin  $\alpha_v\beta_3$  antagonists. Part 1: Design and synthesis of a lead compound exhibiting  $\alpha_v\beta_3/\alpha_{IIb}\beta_3$  dual antagonistic activity. *Bioorg. Med. Chem.* **2006**, *14*, 2089-2108.
- (62) D. Kubota; M. Ishikawa; M. Ishikawa; N. Yahata; S. Murakami; K. Fujishima; M. Kitakaze; K. Ajito. Tricyclic pharmacophore-based molecules as novel integrin  $\alpha_v\beta_3$  antagonists. Part IV: Preliminary control of  $\alpha_v\beta_3$  selectivity by meta-oriented substitution. *Bioorg. Med. Chem.* **2006**, *14*, 4158-4181.
- (63) M. Ishikawa; D. Kubota; M. Yamamoto; C. Kuroda; M. Iguchi; A. Koyanagi; S. Murakami; K. Ajito. Tricyclic pharmacophore-based molecules as novel integrin  $\alpha_v\beta_3$  antagonists. Part 2: Synthesis of potent  $\alpha_v\beta_3/\alpha_{IIb}\beta_3$  dual antagonists. *Bioorg. Med. Chem.* **2006**, *14*, 2109-2130.
- (64) M. Ishikawa; Y. Hiraiwa; D. Kubota; M. Tsushima; T. Watanabe; S. Murakami; S. Ouchi; K. Ajito. Tricyclic pharmacophore-based molecules as novel integrin  $\alpha_v\beta_3$  antagonists. Part III: Synthesis of potent antagonists with  $\alpha_v\beta_3/\alpha_{IIb}\beta_3$  dual activity and improved water solubility. *Bioorg. Med. Chem.* **2006**, *14*, 2131-2150.
- (65) V. W. Engleman; G. A. Nickols; F. P. Ross; M. A. Hortin; D. W. Griggs; S. L. Settle; P. G. Ruminiski; S. L. Teitelbaum. A Peptidomimetic Antagonist of the  $\alpha_v\beta_3$  Integrin

- Inhibits Bone Resorption In Vitro and Prevents Osteoporosis In Vivo. *J. Clin. Inv.* **1997**, *99*, 2284-2292.
- (66) M. Friedlander; P. C. Brooks; R. W. Shaffer; C. M. Kincaid; J. A. Varner; D. A. Cheresh. Definition of two angiogenic pathways by distinct  $\alpha_v$  integrins. *Science* **1995**, *270*, 1500-1502.
- (67) S. L. Goodman; H. J. Grote; C. Wilm. Matched rabbit monoclonal antibodies against  $\alpha_v$ -series integrins reveal a novel  $\alpha_v\beta_3$ -LIBS epitope, and permit routine staining of archival paraffin samples of human tumors. *Biology Open* **2012**, *1*, 329-340.
- (68) C. Böger; H. Kalthoff; S. L. Goodman; H. M. Behrens; C. Röcken. Integrins and their ligands are expressed in non-small cell lung cancer but not correlated with parameters of disease progression. *Virchows Arch.* **2014**, *464*, 69-78.
- (69) L. Bello; M. Francolini; P. Marthyn; J. Zhang; R. S. Carroll; D. C. Nikas; J. F. Strasser; R. Villani; D. A. Cheresh; P. M. Black.  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  Integrin Expression in Glioma Periphery. *Neurosurgery* **2001**, *49*, 380–390.
- (70) A. Erdreich-Epstein; H. Shimada; S. Groshen; M. Liu; L. S. Metelitsa; K. S. Kim; M. F. Stins; R. C. Seeger; D. L. Durden. Integrins  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  are expressed by endothelium of high-risk neuroblastoma and their inhibition is associated with increased endogenous ceramide. **2000**, *60*, 712-721.
- (71) S. De; J. Chen; N. V. Narizhneva; W. Heston; J. Brainard; E. H. Sage; T. V. Byzova. Molecular Pathway for Cancer Metastasis to Bone. *J. Biol. Chem.* **2003**, *278*, 39044-39050.
- (72) A. Bianchi-Smiraglia; S. Paesante; A. V. Bakin. Integrin  $\beta_5$  contributes to the tumorigenic potential of breast cancer cells through the Src-FAK and MEK-ERK signaling pathways. *Oncogene* **2013**, *32*, 3049-3058.



- (73) V. Sung; J. T. I. Stubbs; L. Fisher; A. D. Aaron; E. W. Thompson. Bone sialoprotein supports breast cancer cell adhesion proliferation and migration through differential usage of the  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  integrins. *J Cell Physiol.* **1998**, *176*, 482-494.
- (74) J. J. Letourneau; J. Liu; M. H. J. Ohlmeyer; C. Riviello; Y. Rong; H. Li; K. C. Appell; S. Bansal; B. Jacob; A. Wong; M. L. Webb. Synthesis and initial evaluation of novel, non-peptidic antagonists of the  $\alpha_v$ -integrins  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$ . *Bioorg. Med. Chem. Lett.* **2009**, *19*, 352-355.
- (75) A. Trabocchi; G. Menchi; N. Cini; F. Bianchini; S. Raspanti; A. Bottoncetti; A. Pupi; L. Calorini; A. Guarna. Click-Chemistry-Derived Triazole Ligands of Arginine-Glycine-Aspartate (RGD) Integrins with a Broad Capacity To Inhibit Adhesion of Melanoma Cells and Both in Vitro and in Vivo Angiogenesis. *J. Med. Chem.* **2010**, *53*, 7119-7128.
- (76) C. C. Kumar; M. Malkowski; Z. Yin; E. Tanghetti; B. Yaremko; T. Nechuta; J. Varner; M. X. Liu; E. M. Smith; B. Neustadt; M. Presta; L. Armstrong. Inhibition of Angiogenesis and Tumor Growth by SCH221153, a Dual  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  Integrin Receptor Antagonist. *Cancer Res.* **2001**, *61*, 2232-2238.
- (77) A. R. Reynolds; I. R. Hart; A. R. Watson; J. C. Welti; R. G. Silva; S. D. Robinson; G. Violante; M. Gourlaouen; M. Salih; M. C. Jones; D. T. Jones; G. Saunders; V. Kostourou; F. Perron-Sierra; J. C. Norman; G. C. Tucker; K. M. Hodivala-Dilke. Stimulation of tumor growth and angiogenesis by low concentrations of RGD-mimetic integrin inhibitors. *Nature Medicine* **2009**, *15*, 392-400.
- (78) F. Perron-Sierra; D. Saint Dizier; M. Bertrand; A. Genton; G. C. Tucker; P. Casara. Substituted Benzocycloheptenes as Potent and Selective  $\alpha_v$  Integrin Antagonists. *Bioorg. Med. Chem. Lett.* **2002**, *12*, 3291-3296.
- (79) M. F. Burbridge; V. Venot; P. J. Casara; F. Perron-Sierra; J. A. Hickman; G. C. Tucker. Decrease in Survival Threshold of Quiescent Colon Carcinoma Cells in the

- Presence of a Small Molecule Integrin Antagonist. *Mol. Pharmacol.* **2003**, *63*, 1281-1288.
- (80) S. Maubant; D. Saint-Dizier; M. Boutillon; F. Perron-Sierra; P. J. Casara; J. A. Hickman; G. C. Tucker; E. Van Obberghen-Schilling. Blockade of  $\alpha_v\beta_3$  and  $\alpha_v\beta_5$  integrins by RGD mimetics induces anoikis and not integrin-mediated death in human endothelial cells. *Blood* **2006**, *108*, 3035-3044.
- (81) C. Mas-Moruno; F. Rechenmacher; H. Kessler. Cilengitide: The First Anti-Angiogenic Small Molecule Drug Candidate. Design, Synthesis and Clinical Evaluation. *Anti-cancer Agents Med. Chem.* **2010**, *10*, 753-768.
- (82) M. Aumailley; M. Gurrath; G. Müller; J. Calvete; R. Timpl; H. Kessler. Arg-Gly-Asp constrained within cyclic pentapeptides. Strong and selective inhibitors of cell adhesion to vitronectin and laminin fragments. *FEBS Lett.* **1991**, *291*, 50-54.
- (83) M. A. Dechantsreiter; E. Planker; B. Matha; E. Lohof; G. Holzemann; A. Jonczyk; S. L. Goodman; H. Kessler. *N*-Methylated Cyclic RGD Peptides as Highly Active and Selective  $\alpha_v\beta_3$  Integrin Antagonists. *J. Med. Chem.* **1999**, *42*, 3033-3040.
- (84) C. Mas-Moruno; J. G. Beck; L. Doedens; A. O. Frank; L. Marinelli; S. Cosconati; E. Novellino; H. Kessler. Increasing  $\alpha_v\beta_3$  Selectivity of the Anti-Angiogenic Drug Cilengitide by *N*-Methylation. *Angew. Chem. Int. Ed.* **2011**, *50*, 9496-9500.
- (85) C. Scaringi; G. Minniti; P. Caporello; R. M. Enrici. Integrin inhibitor cilengitide for the treatment of glioblastoma: a brief overview of current clinical results. *Anticancer Res.* **2012**, *32*, 4213-4223.
- (86) U. K. Marelli; F. Rechenmacher; T. R. A. Sobahi; C. Mas-Moruno; H. Kessler. Tumor targeting via integrin ligands. *Front. Oncol.* **2013**, *3*, 222.
- (87) L. B. Nabors; T. Mikkelsen; M. E. Hegi; X. Ye; T. Batchelor; G. Lesser; D. Peereboom; M. R. Rosenfeld; J. Olsen; S. Brem; J. D. Fisher; S. A. Grossman; New Approaches to Brain Tumor Therapy Central Nervous System Consortium. A safety

- run-in and randomized phase 2 study of cilengitide combined with chemoradiation for newly diagnosed glioblastoma (NABTT 0306). *Cancer* **2012**, *118*, 5601-5607.
- (88) J. B. Vermorken; J. Guigay; R. Mesia; J. M. Trigo; U. Keilholz; A. Kerber; U. Bethe; M. Picard; T. H. Brummendorf. Phase I/II trial of cilengitide with cetuximab, cisplatin and 5-fluorouracil in recurrent and/or metastatic squamous cell cancer of the head and neck: findings of the phase I part. *Br. J. Cancer* **2011**, *104*, 1691-1696.
- (89) M. Bretsch; C. Cheng; H. Witt; A. Dimitrakopoulou-Strauss; L. G. Strauss; W. Semmler; T. Bäuerle. Cilengitide affects tumor compartment, vascularization and microenvironment in experimental bone metastases as shown by longitudinal <sup>18</sup>F-FDG PET and gene expression analysis. *J. Cancer Res. Clin. Oncol.* **2013**, *139*, 573-583.
- (90) M. Bretsch; M. Merz; D. Komljenovic; M. R. Berger; W. Semmler; T. Bauerle. Cilengitide inhibits metastatic bone colonization in a nude rat model. *Oncol. Rep.* **2011**, *26*, 843-851.
- (91) S. Martin; H. Janouskova; M. Dontenwill. Integrins and p53 pathways in glioblastoma resistance to temozolomide. *Front. Oncol.* **2012**, *2*, 157.
- (92) F. Schaffner; R. A. M.; M. Dontenwill. Integrin  $\alpha_5\beta_1$ , the Fibronectin Receptor, as a Pertinent Therapeutic Target in Solid Tumors. *Cancers* **2013**, *5*, 27-47.
- (93) V. I. Romanov; M. S. Goligorsky. RGD-Recognizing Integrins Mediate Interactions of Human Prostate Carcinoma Cells With Endothelial Cells In Vitro. *Prostate* **1999**, *39*, 108-118.
- (94) A. Stachurska; J. Elbanowski; H. M. Kowalczyńska. Role of  $\alpha_5\beta_1$  and  $\alpha_v\beta_3$  integrins in relation to adhesion and spreading dynamics of prostate cancer cells interacting with fibronectin under in vitro conditions. *Cell Biol. Int.* **2012**, *36*, 883-892.

- (95) K. O. Simon; E. M. Nutt; D. G. Abraham; G. A. Rodan; L. T. Duong. The  $\alpha_v\beta_3$  Integrin Regulates  $\alpha_5\beta_1$ -mediated Cell Migration toward Fibronectin. *J. Biol. Chem.* **1997**, 272, 29380-29389.
- (96) S. Kim; M. Harris; J. A. Varner. Regulation of Integrin  $\alpha_v\beta_3$ -mediated Endothelial Cell Migration and Angiogenesis by Integrin  $\alpha_5\beta_1$  and Protein Kinase A. *J. Biol. Chem.* **2000**, 275, 33920-33928.
- (97) J. G. Parvani; A. J. Galliher-Beckley; B. J. Schiemann; W. P. Schiemann. Targeted inactivation of  $\beta_1$  integrin induces  $\beta_3$  integrin switching that drives breast cancer metastasis by TGF- $\beta$ . *Mol. Biol. Cell* **2013**, 24, 3449-3459.
- (98) C. Christoforides; E. Rainero; K. K. Brown; J. C. Norman; A. Toker. PKD Controls  $\alpha_v\beta_3$  Integrin Recycling and Tumor Cell Invasive Migration Through its Substrate Rabaptin-5. *Dev. Cell* **2012**, 23, 560-572.
- (99) A. van der Flier; K. Badu-Nkansah; C. A. Whittaker; D. Crowley; R. T. Bronson; A. Lacy-Hulbert; R. O. Hynes. Endothelial  $\alpha_5$  and  $\alpha_v$  integrins cooperate in remodeling of the vasculature during development. *Development* **2010**, 137, 2439-2449.
- (100) A. M. Wan; E. M. Chandler; M. Madhavan; D. W. Infanger; C. K. Ober; D. Gourdon; G. G. Malliaras; C. Fischbach. Fibronectin conformation regulates the proangiogenic capability of tumor-associated adipogenic stromal cells. *Biochim. Biophys. Acta* **2013**, 1830, 4314-4320.
- (101) N. Laurens; M. A. Engelse; C. Jungerius; C. W. Löwik; V. W. van Hinsbergh; P. Koolwijk. Single and combined effects of  $\alpha_v\beta_3$  and  $\alpha_5\beta_1$  integrins on capillary tube formation in a human fibrinous matrix. *Angiogenesis* **2009**, 12, 275-285.
- (102) R. Stragies; F. Osterkamp; G. Zischinsky; D. Vossmeier; H. Kalkhof; U. Reimer; G. Zahn. Design and Synthesis of a New Class of Selective Integrin  $\alpha_5\beta_1$  Antagonists. *J. Med. Chem.* **2007**, 50, 3786-3794.

- (103) F. Benfatti; G. Cardillo; S. Fabbroni; P. Galzerano; L. Gentilucci; R. Juris; A. Tolomelli; M. Baiula; A. Sparta; S. Spampinato. Synthesis and biological evaluation of non-peptide  $\alpha_v\beta_3/\alpha_5\beta_1$  integrin dual antagonists containing 5,6-dihydropyridin-2-one scaffolds. *Bioorg. Med. Chem.* **2007**, *15*, 7380-7390.
- (104) A. Tolomelli; M. Baiula; L. Belvisi; A. Viola; L. Gentilucci; S. Troisi; S. Deianira Dattoli; S. Spampinato; M. Civera; E. Juaristi; M. Escudero. Modulation of  $\alpha_v\beta_3$ - and  $\alpha_5\beta_1$ -integrin-mediated adhesion by dehydro- $\beta$ -amino acids containing peptidomimetics. *Eur. J. Med. Chem.* **2013**, *66*, 258-268.
- (105) L. Gentilucci; G. Cardillo; S. Spampinato; A. Tolomelli; F. Squassabia; R. De Marco; A. Bedini; M. Baiula; L. Belvisi; M. Civera. Antiangiogenic Effect of Dual/Selective  $\alpha_5\beta_1/\alpha_v\beta_3$  Integrin Antagonists Designed on Partially Modified Retro-Inverso Cyclotetrapeptide Mimetics. *J. Med. Chem.* **2010**, *53*, 106-118.
- (106) J. M. Smallheer; C. A. Weigelt; F. J. Woerner; J. S. Wells; W. F. Daneker; S. A. Mousa; R. R. Wexler; P. K. Jadhav. Synthesis and biological evaluation of nonpeptide integrin antagonists containing spirocyclic scaffolds. *Bioorg. Med. Chem. Lett.* **2004**, *14*, 383-387.
- (107) A. H. Schmieder; S. D. Caruthers; H. Zhang; T. A. Williams; J. D. Robertson; S. A. Wickline; G. M. Lanza. Three-dimensional MR mapping of angiogenesis with  $\alpha_5\beta_1$  ( $\alpha_v\beta_3$ )-targeted theranostic nanoparticles in the MDA-MB-435 xenograft mouse model. *FASEB J.* **2008**, *22*, 4179-4189.
- (108) A. Bandyopadhyay; S. Raghavan. Defining the Role of Integrin  $\alpha_v\beta_6$  in Cancer. *Curr. Drug Targets* **2009**, *10*, 645-652.
- (109) S. L. Goodman; G. Holzemann; G. A. G. Sulyok; H. Kessler. Nanomolar Small Molecule Inhibitors for  $\alpha_v\beta_6$ ,  $\alpha_v\beta_5$ , and  $\alpha_v\beta_3$  Integrins. *J. Med. Chem.* **2002**, *45*, 1045-1051.

- (110) J. Schittenhelm; E. I. Schwab; J. Sperveslage; M. Tatagiba; R. Meyermann; F. Fend; S. L. Goodman; B. Sipos. Longitudinal Expression Analysis of  $\alpha_v$  Integrins in Human Gliomas Reveals Upregulation of Integrin  $\alpha_v\beta_3$  as a Negative Prognostic Factor. *J. Neuropathol. Exp. Neurol.* **2013**, 72, 194-210.
- (111) A. Vogetseder; S. Thies; B. Ingold; P. Roth; M. Weller; P. Schraml; S. L. Goodman; H. Holger Moch.  $\alpha_v$ -Integrin isoform expression in primary human tumors and brain metastases. *Int. J. Cancer* **2013**, 133, 2362–2371.
- (112) J. Schittenhelm; A. Klein; M. S. Tatagiba; R. Meyermann; F. Fend; S. L. Goodman; B. Sipos. Comparing the expression of integrins  $\alpha_v\beta_3$ ,  $\alpha_v\beta_5$ ,  $\alpha_v\beta_6$ ,  $\alpha_v\beta_8$ , fibronectin and fibrinogen in human brain metastases and their corresponding primary tumors. *Int. J. Clin. Exp. Pathol.* **2013**, 6, 2719-2732.
- (113) K. Bisanz; J. Yu; M. Edlund; B. Spohn; M.-C. Hung; L. W. K. Chung; C.-L. Hsieh. Targeting ECM–Integrin Interaction with Liposome-Encapsulated Small Interfering RNAs Inhibits the Growth of Human Prostate Cancer in a Bone Xenograft Imaging Model. *Mol. Ther.* **2005**, 12, 634-643.
- (114) C. van den Hoogen; G. van der Horst; H. Cheung; J. T. Buijs; R. C. M. Pelger; G. van der Pluijm. Integrin  $\alpha_v$  Expression Is Required for the Acquisition of a Metastatic Stem/Progenitor Cell Phenotype in Human Prostate Cancer. *Am. J. Pathol.* **2011**, 179, 2559-2568.
- (115) M. Trikha; Z. Zhou; J. A. Nemeth; Q. Chen; C. Sharp; E. Emmell; J. Giles-Komar; M. T. Nakada. CNTO 95, A fully human monoclonal antibody that inhibits  $\alpha_v$  integrins, has antitumor and antiangiogenic activity *in vivo*. *Int. J. Cancer* **2004**, 110, 326-335.
- (116) Y. J. Wu; L. L. Muldoon; S. Gahramanov; D. F. Kraemer; D. J. Marshall; E. A. Neuwelt. Targeting  $\alpha_v$ -integrins decreased metastasis and increased survival in a nude rat breast cancer brain metastasis model. *J Neurooncol.* **2012**, 110, 27-36.

- (117) Q. Chen; C. D. Manning; H. Millar; F. L. McCabe; C. Ferrante; C. Sharp; L. Shahied-Arruda; P. Doshi; M. T. Nakada; G. M. Anderson. CNTO 95, a fully human anti- $\alpha_v$  integrin antibody, inhibits cell signaling, migration, invasion, and spontaneous metastasis of human breast cancer cells. *Clin. Exp. Metastasis* **2008**, *25*, 139-148.
- (118) P. L. Martin; Q. Jiao; J. Cornacoff; W. Hall; B. Saville; J. A. Nemeth; A. Schantz; M. Mata; H. Jang; A. A. Fasanmade; L. Anderson; M. A. Graham; H. M. Davis; G. Treacy. Absence of Adverse Effects in Cynomolgus Macaques Treated with CNTO 95, a Fully Human Anti- $\alpha_v$  Integrin Monoclonal Antibody, Despite Widespread Tissue Binding. *Clin. Cancer Res.* **2005**, *11*, 6959-6965.
- (119) S. A. Mullahitha; N. C. Ton; G. J. Parker; A. Jackson; P. J. Julyan; C. Roberts; G. A. Buonaccorsi; Y. Watson; K. Davies; S. Cheung; L. Hope; J. W. Valle; J. A. Radford; J. Lawrance; M. P. Saunders; M. C. Munteanu; M. T. Nakada; J. A. Nemeth; H. M. Davis; Q. Jiao; U. Prabhakar; Z. Lang; R. E. Corringham; R. A. Beckman; G. C. Jayson. Phase I evaluation of a fully human anti- $\alpha_v$  integrin monoclonal antibody (CNTO 95) in patients with advanced solid tumors. *Clin. Cancer Res.* **2007**, *13*, 2128-2135.
- (120) S. J. O'Day; A. C. Pavlick; M. R. Albertini; O. Hamid; H. Schalch; Z. Lang; J. Ling; M. Mata; M. Reddy; B. Foster. Clinical and pharmacologic evaluation of two dose levels of intetumumab (CNTO 95) in patients with melanoma or angiosarcoma. *Invest. New Drugs* **2012**, *30*, 1074-1081.
- (121) S. O'Day; A. Pavlick; C. Loquai; D. Lawson; R. Gutzmer; J. Richards; D. Schadendorf; J. A. Thompson; R. Gonzalez; U. Trefzer; P. Mohr; C. Ottensmeier; D. Chao; B. Zhong; C. J. de Boer; C. Uhlir; D. Marshall; M. E. Gore; Z. Lang; W. Hait; P. Ho; S. O'Day; S. Monica; J. Richards; P. Ridge; A. Pavlick; L. Feun; T. Malpass; M. Gordon; R. Gonzalez; G. Daniels; J. Hainsworth; D. Lawson; J. A. Thompson; T. Sato; M. Sherman; D. Schadendorf; T. Tueting; A. Hauschild; U. Trefzer; C.

- Sunderkötter; P. Mohr; R. Gutzmer; C. Loquai; J. Freise; U. R. Hengge; M. Kaatz; M. Gore; M. Harries; C. Ottensmeier; P. Lorigan; D. Chao; P. Corrie; S. Danson. A randomised, phase II study of intetumumab, an anti- $\alpha_v$ -integrin mAb, alone and with dacarbazine in stage IV melanoma. *Br. J. Cancer* **2011**, *105*, 346-352.
- (122) F. M. Chu; J. Picus; P. M. Fracasso; R. Dreicer; Z. Lang; B. Foster. A phase 1, multicenter, open-label study of the safety of two dose levels of a human monoclonal antibody to human  $\alpha(v)$  integrins, intetumumab, in combination with docetaxel and prednisone in patients with castrate-resistant metastatic prostate cancer. *Invest. New Drugs* **2011**, *29*, 674-679.
- (123) A. Heidenreich; S. K. Rawal; K. Szkarlat; N. Bogdanova; L. Dirix; A. Stenzl; M. Welslau; G. Wang; F. Dawkins; C. J. de Boer; D. Schrijvers. A randomized, double-blind, multicenter, phase 2 study of a human monoclonal antibody to human  $\alpha_v$  integrins (intetumumab) in combination with docetaxel and prednisone for the first-line treatment of patients with metastatic castration-resistant prostate cancer. *Ann. Oncol.* **2013**, *24*, 329-336.
- (124) F. Mitjans; T. Meyer; C. Fittschen; S. Goodman; A. Jonczyk; J. F. Marshall; G. Reyes; J. Piulats. In Vivo Therapy of Malignant Melanoma by means of Antagonists of  $\alpha_v$  Integrins. *Int. J. Cancer* **2000**, *87*, 716-723.
- (125) W. Uhl; M. Zühlsdorf; T. Koernicke; U. Forssmann; A. Kovar. Safety, tolerability, and pharmacokinetics of the novel  $\alpha_v$ -integrin antibody EMD 525797 (DI17E6) in healthy subjects after ascending single intravenous doses. *Invest New Drugs*. **2013**, DOI:10.1007/s10637-013-038-5.
- (126) M. Wirth; A. Heidenreich; J. E. Gschwend; T. Gil; S. Zastrow; M. Laniado; J. Gerloff; M. Zühlsdorf; G. Mordenti; W. Uhl; H. Lannert. A Multicenter Phase 1 Study of EMD 525797 (DI17E6), a Novel Humanized Monoclonal Antibody



- Targeting  $\alpha_v$  Integrins, in Progressive Castration-resistant Prostate Cancer with Bone Metastases After Chemotherapy. *Eur. Urol.* **2013**, doi:10.1016/j.eururo.2013.05.051.
- (127) R. M. Scarborough. Development of Eptifibatide. *Am. Heart J.* **1999**, *138*, 1093-1104.
- (128) J. J. Calvete. The continuing saga of snake venom disintegrins. *Toxicon* **2013**, *62*, 40-49.
- (129) J. J. Calvete; J. W. Fox; A. Agelan; S. Niewiarowski; C. Marcinkiewicz. The Presence of the WGD Motif in CC8 Heterodimeric Disintegrin Increases Its Inhibitory Effect on  $\alpha_{IIb}\beta_3$ ,  $\alpha_v\beta_3$ , and  $\alpha_5\beta_1$  Integrins. *Biochemistry* **2002**, *41*, 2014-2021.
- (130) E. E. Sánchez; A. Rodríguez-Acosta; R. Palomar; S. E. Lucena; S. Bashir; J. G. Soto; J. C. Pérez. Colombistatin: a disintegrin isolated from the venom of the South American snake (*Bothrops colombiensis*) that effectively inhibits platelet aggregation and SK-Mel-28 cell adhesion. *Arch. Toxicol.* **2009**, *83*, 271-279.
- (131) S. E. Lucena; Y. Jia; J. G. Soto; J. Parral; E. Cantu; J. Brannon; K. Lardner; C. J. Ramos; A. I. Seoane; E. E. Sánchez. Anti-invasive and anti-adhesive activities of a recombinant disintegrin, r-viridistatin 2, derived from the Prairie rattlesnake (*Crotalus viridis viridis*). *Toxicon* **2012**, *60*, 31-39.
- (132) C. M. Carey; R. Bueno; D. A. Gutierrez; C. Petro; S. E. Lucena; E. E. Sanchez; J. G. Soto. Recombinant rubistatin (r-Rub), an MVD disintegrin, inhibits cell migration and proliferation, and is a strong apoptotic inducer of the human melanoma cell line SK-Mel-28. *Toxicon* **2012**, *59*, 241-248.
- (133) C.-H. Shih; T.-B. Chiang; W.-J. Wang. Inhibition of integrins  $\alpha_v/\alpha_5$ -dependent functions in melanoma cells by an ECD-disintegrin acurhagin-C. *Matrix Biol.* **2013**, *32*, 152-159.

- (134) I.-C. Kang; D.-S. Kim; Y. Jang; K.-H. Chung. Suppressive Mechanism of Salmosin, a Novel Disintegrin in B16 Melanoma Cell Metastasis. *Biochem. Biophys. Res. Comm.* **2000**, 275, 169-173.
- (135) M. Trikha; Y. A. De Clerck; F. S. Markland. Contortrostatin, a Snake Venom Disintegrin, Inhibits  $\beta_1$  Integrin-mediated Human Metastatic Melanoma Cell Adhesion and Blocks Experimental Metastasis. *Cancer Res.* **1994**, 54, 4993-4998.
- (136) R. O. Minea; C. M. Helchowski; S. J. Zidovetzki; F. K. Costa; S. D. Swenson; F. S. J. Markland. Vicrostatin – An Anti-Invasive Multi-Integrin Targeting Chimeric Disintegrin with Tumor Anti-Angiogenic and Pro-Apoptotic Activities. *PLoS One* **2010**, 5, e10929.
- (137) R. Minea; S. Swenson; F. Costa; T. C. Chen; F. S. Markland. Development of a Novel Recombinant Disintegrin, Contortrostatin, as an Effective Anti-Tumor and Anti-Angiogenic Agent. *Pathophysiol. Haemost. Thromb.* **2005**, 34, 177-183.
- (138) S. Swenson; F. Costa; R. Minea; R. P. Sherwin; W. Ernst; G. Fujii; D. Yang; F. S. Markland Jr. Intravenous liposomal delivery of the snake venom disintegrin contortrostatin limits breast cancer progression. *Mol. Cancer Ther.* **2004**, 3, 499-511.
- (139) K.-H. Chung; S.-H. Kim; K.-y. Han; Y.-D. Sohn; S.-I. Chang; K.-H. Baek; Y. Jang; D.-S. Kim; I.-C. Kang. Inhibitory effect of salmosin, a Korean snake venom derived disintegrin, on the integrin  $\alpha_v$ -mediated proliferation of SK-Mel-2 human melanoma cells. *J. Pharm. Pharmacol.* **2003**, 55, 1577-1582.
- (140) E. Lin; Q. Wang; S. Swenson; H. Jadvar; S. Groshen; W. Ye; F. S. Markland; J. Pinski. The Disintegrin Contortrostatin in Combination With Docetaxel Is a Potent Inhibitor of Prostate Cancer In Vitro and In Vivo. *Prostate* **2010**, 70, 1359-1370.
- (141) M. Morjen; O. Kallech-ziri; A. Bazaa; H. Othman; K. Mabrouk; R. Zouari-kessentini; L. Sanz; J. J. Calvete; N. Srairi-Abid; El Ayeb; J. Luis; N. Marrakchi. PIVL, a new

- serine protease inhibitor from *Macrovipera lebetina transmediterranea* venom, impairs motility of human glioblastoma cells. *Matrix Biol.* **2013**, *32*, 52-62.
- (142) S. Hailey; E. Adams; R. Penn; A. Wong; M. A. McLane. Effect of the disintegrin Eristostatin on melanoma-natural killer cell interactions. *Toxicon* **2013**, *61*, 83-93.
- (143) M. L. Boys; L. A. Schretzman; N. S. Chandrakumar; M. B. Tollefson; S. B. Mohler; V. L. Downs; T. D. Penning; M. A. Russell; J. A. Wendt; B. B. Chen; H. G. Stenmark; H. Wu; D. P. Spangler; M. Clare; B. N. Desai; I. K. Khanna; M. N. Nguyen; T. Duffin; V. W. Engleman; M. B. Finn; S. K. Freeman; M. L. Hanneke; J. L. Keene; J. A. Klover; G. A. Nickols; M. A. Nickols; C. N. Steininger; M. Westlin; W. Westlin; Y. X. Yu; Y. Wang; C. R. Dalton; S. A. Norring. Convergent, parallel synthesis of a series of  $\beta$ -substituted 1,2,4-oxadiazole butanoic acids as potent and selective  $\alpha_v\beta_3$  receptor antagonists. *Bioorg. Med. Chem. Lett.* **2006**, *16*, 839-844.
- (144) K. E. Shannon; J. L. Keene; S. L. Settle; T. D. Duffin; M. A. Nickols; M. Westlin; S. Schroeter; P. G. Ruminski; D. W. Griggs. Anti-Metastatic Properties of RGD-Peptidomimetic Agents S137 and S247. *Clin. Exp. Metastasis* **2004**, *21*, 129-138.
- (145) K. Meerovitch; F. Bergeron; L. Leblond; B. Grouix; C. Poirier; M. Bubenik; L. Chan; H. Gourdeau; T. Bolwin; G. Attardo. A novel RGD antagonist that targets both  $\alpha_v\beta_3$  and  $\alpha_5\beta_1$  induces apoptosis of angiogenic endothelial cells on type I collagen. *Vasc. Pharmacol.* **2003**, *40*, 77-89.
- (146) H. P. Naber; E. Wiercinska; E. Pardali; T. van Laar; E. Nirmala; A. Sundqvist; H. van Dam; G. van der Horst; G. van der Pluijm; B. Heckmann; E. H. Danen; P. Ten Dijke. BMP-7 inhibits TGF- $\beta$ -induced invasion of breast cancer cells through inhibition of integrin  $\beta_3$  expression. *Cell. Oncol. (Dordr)*. **2012**, *35*, 19-28.
- (147) N. C. Henderson; T. D. Arnold; Y. Katamura; M. M. Giacomini; J. D. Rodriguez; J. H. McCarty; A. Pellicoro; E. Raschperger; C. Betsholtz; P. G. Ruminski; D. W. Griggs; M. J. Prinsen; J. J. Maher; J. P. Iredale; A. Lacy-Hulbert; R. H. Adams; D.

- Sheppard. Targeting of  $\alpha_v$  integrin identifies a core molecular pathway that regulates fibrosis in several organs. *Nature Medicine* **2013**, *19*, 1617-1624.
- (148) N. Reinmuth; W. Liu; S. A. Ahmad; F. Fan; O. Stoeltzing; A. A. Parikh; C. D. Bucana; G. E. Gallick; M. A. Mickols; W. F. Westlin; L. M. Ellis.  $\alpha_v\beta_3$  Integrin Antagonist S247 Decreases Colon Cancer Metastasis and Angiogenesis and Improves Survival in Mice. *Cancer Res.* **2003**, *63*, 2079-2087.
- (149) J. F. Harms; D. R. Welch; R. S. Samant; L. A. Shevde; M. E. Miele; G. R. Babu; S. F. Goldberg; V. R. Gilman; D. M. Sosnowski; D. A. Campo; C. V. Gay; L. R. Budgeon; R. Mercer; J. Jewell; A. M. Mastro; H. J. Donahue; N. Erin; M. T. Debies; W. J. Meehan; A. L. Jones; G. Mbalaviele; A. Nickols; N. D. Christensen; R. Melly; L. N. Beck; J. Kent; R. K. Rader; J. J. Kotyk; M. D. Pagel; W. F. Westlin; D. W. Griggs. A Small Molecule Antagonist of the  $\alpha_v\beta_3$  Integrin Suppresses MDA-MB-435 Skeletal Metastasis. *Clin. Exp. Metastasis* **2004**, *21*, 119-128.
- (150) B. Heckmann; J.-M. Lefrancois Pyrimidines derivatives as antagonists of the vitronectine receptors. European Patent EP2368891A1, 28 September 2011.
- (151) G. van der Horst; C. van den Hoogen; J. T. Buijs; H. Cheung; R. C. M. Pelger; J. Feyen; P. Pujuguet; R. Blanque; P. Clément-Lacroix; G. van der Pluijm. Targeting of  $\alpha_v$  integrins, a potential marker for tumor-initiating cells, in prostate cancer and bone stroma inhibits bone metastasis formation. *Bone* **2010**, *47*, S320.
- (152) G. van der Horst; C. van den Hoogen; J. T. Buijs; H. Cheung; H. Bloys; R. C. M. Pelger; G. Lorenzon; B. Heckmann; J. Feyen; P. Pujuguet; R. Blanque; P. Clément-Lacroix; G. van der Pluijm. Targeting of  $\alpha_v$ -Integrins in Stem/Progenitor Cells and Supportive Microenvironment Impairs Bone Metastasis in Human Prostate Cancer. *Neoplasia* **2011**, *13*, 516-525.
- (153) Clinicaltrials.gov: [www.clinicaltrials.gov](http://www.clinicaltrials.gov). Accessed 27/11/13.