Table 3: Properties of different categories of protein-based building blocks with potential applications in nanosci internacional)

drug delivery. Main pitfalls are stressed.

Category	<u>Origin</u>	<u>Procedure</u>	Cage/shell formation	Functional versatility	Architectonic versatility	Delivered drugs	Representative examples or reviews
Natural viruses	<u>Natural</u>	Cell culture	<u>Yes</u>	Limited (tropism can be re- directed; severe side effects)	None	Mainly DNA	(Giacca, M. et al. 2012)
VLPs	<u>Natural</u>	Recombinant	<u>Yes</u>	Limited (tropism can be re- directed)	Limited (disassembling and reassembling can be controlled in vitro)	Nucleic acids, proteins and chemicals	(Fang, C. Y. et al. 2012;Kaczmarc zyk, S. J. et al. 2011;Zhao, Q. et al. 2011)
BMCs	<u>Natural</u>	Recombinant	Yes	Limited (BMC proteins can be engineered)	Very limited	Proteins and chemicals	(Seebeck, F. P. et al. 2006;Sutter, M. et al. 2008)
Vaults	Natural	Recombinant	<u>Yes</u>	Limited (vault proteins can be engineered)	Very limited	Proteins, chemicals and gold probes	(Goldsmith, L. E. et al. 2009;Gopinath, S. C. et al. 2005;Kickhoefer , V. A. et al. 2005)
<u>IBs</u>	Bioinspired	Recombinant	<u>No</u>	High (essentially any protein can form IBs; tropism can be conferred)	Limited (Size and geometry are regulatable)	Proteins	(Liovic, M. et al. 2012; Talafova, K., Hrabarova, E., Chorvat, D. A., & Nahalka, J. 2013; Vazquez, E. et al. 2012)
Peptides	Usually bioinspired	Chemical synthesis	<u>No</u>	Moderate (essentially any short aa sequence can be produced)	High (self-assembling can be engineered)	Nucleic acids and chemicals	(Hosseinkhani, H., Hosseinkhani, M., Khademhosseini , A., Kobayashi, H., & Tabata, Y. 2006;Hsieh, P. C., Davis, M. E.,

							Gannon, J., MacGillivray, C., & Lee, R. T. 2006)
Modular proteins	Bioinspired	Recombinant	<u>Yes</u>	High (essentially any protein and peptide set can be combined)	Moderate (through protein engineering)	Nucleic acids and proteins	(Unzueta, U. et al. 2012;Unzueta, U. et al. 2012;Vazquez, E. et al. 2010)

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