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**Edited and reviewed by:**

Pietro Genovese,  
Boston Children's Hospital and  
Harvard Medical School,  
United States

**\*Correspondence:**

Mario Amendola  
mamendola@genethon.fr

**†Present address:**

Giulia Pavani,  
The Children's Hospital of  
Philadelphia, Raymond G. Perelman  
Center for Cellular and Molecular  
Therapeutics, Philadelphia, PA,  
United States

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# Corrigendum: Targeted Gene Delivery: Where to Land

**Giulia Pavani<sup>†</sup> and Mario Amendola\***

INTEGRARE, UMR\_S951, Genethon, Inserm, Univ Evry, Univ Paris-Saclay, Evry, France

**Keywords:** genome editing, gene therapy, nuclease, CRISPR, targeted integration (TI), knock-in, safe harbor, homologous recombination (HR)

## A Corrigendum on

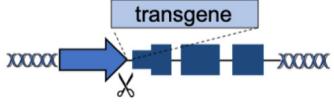
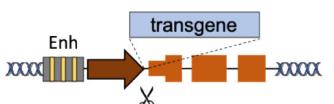
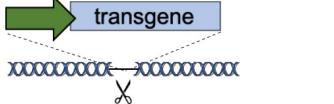
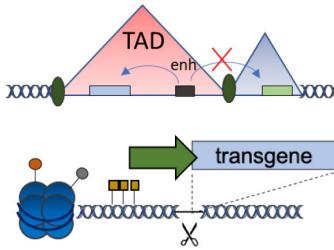
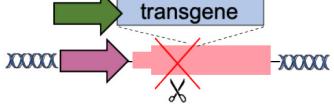
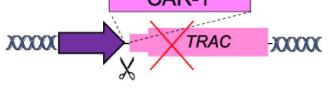
### Targeted Gene Delivery: Where to Land

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In the original article, there was a mistake in **Table 1** as published. The references indicated in row **B** are wrong. The corrected **Table 1** appears in the attached below.

The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

**TABLE 1 | (A–F)** The advantages and disadvantages of different integration strategies.

	Integration strategies	Advantages	Disadvantages	References	
A	Endogenous locus		Physiological transgene expression Corrects multiple mutations	Gene-specific strategy Limited to gene body mutations	Urnov et al., 2005; Lombardo et al., 2007; Li et al., 2011; Genovese et al., 2014; Voit et al., 2014; Dever et al., 2016; Hubbard et al., 2016; Schioli et al., 2017; Sweeney et al., 2017; Kuo et al., 2018; Wang et al., 2019; Rai et al., 2020; Wang L. et al., 2020
B	Superactive promoters (ALB, HBA)		Accommodates different transgenes Supraphysiological expression Few integrations required	Partial gene disruption Limited to non-cell autonomous disorders Extensive validation required	Barzel et al., 2015; Sharma et al., 2015; Davidoff and Nathwani, 2016; Laoharawee et al., 2018; Chen et al., 2019; Conway et al., 2019; De Caneva et al., 2019; Ou et al., 2019, 2020; Zhang et al., 2019; Wang Q. et al., 2020
C	Tolerant to integration (AAVS1, CCR5, Rosa26)		Accommodates different transgenes	Artificial promoters required Variable expression	De Ravin et al., 2016; Diez et al., 2017; Stephens et al., 2018, 2019; Gomez-Ospina et al., 2019; Scharenberg et al., 2020
D	Chromatin domains (NAD)		Fine gene regulation Far from oncogenic genes	No proof-of-principle in clinically relevant models	Schenkwein et al., 2020
E	Disease-modifier genes (CCR5, HBA)		Improve therapeutic effect Lower therapeutic threshold	Extensive validation required Limited to well-known diseases	Voit et al., 2013; Wiebking et al., 2018
F	Specificity Exchange (TCR, BCR)		Improved CAR expression and potency	Off-targets Translocations risk (for multiple edits)	Eyquem et al., 2017; MacLeod et al., 2017; Greiner et al., 2019; Hartweger et al., 2019; Moffett et al., 2019; Voss et al., 2019

Scissors: nuclease; Solid arrows: promoters; Enh, enhancers; TAD, topologically associating; d, domain; Solid ovals: histone modifications; Solid squares: DNA modifications.

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