

Correction

Correction: Nelson et al. *UGT1A1* Guided Cancer Therapy: Review of the Evidence and Considerations for Clinical Implementation. *Cancers* 2021, 13, 1566

Ryan S. Nelson ^{1,2}, Nathan D. Seligson ^{3,4} , Sal Bottiglieri ⁵ , Estrella Carballido ^{6,7}, Alex Del Cueto ², Iman Imanirad ^{6,7}, Richard Levine ^{6,8}, Alexander S. Parker ⁹, Sandra M. Swain ¹⁰, Emma M. Tillman ¹¹  and J. Kevin Hicks ^{2,6,*} 

- ¹ Department of Consultative Services, ARUP Laboratories, Salt Lake City, UT 84108, USA; ryan.nelson@aruplab.com
- ² Department of Individualized Cancer Management, Moffitt Cancer Center, Tampa, FL 33612, USA; alex.delcueto@moffitt.org
- ³ Department of Pharmacotherapy and Translational Research, The University of Florida, Jacksonville, FL 32610, USA; nseligson@cop.ufl.edu
- ⁴ Department of Hematology and Oncology, Nemours Children’s Specialty Care, Jacksonville, FL 32207, USA
- ⁵ Department of Pharmacy, Moffitt Cancer Center, Tampa, FL 33612, USA; salvatore.bottiglieri@moffitt.org
- ⁶ Department of Oncological Sciences, University of South Florida, Tampa, FL 33612, USA; estrella.carballido@moffitt.org (E.C.); iman.imanirad@moffitt.org (I.I.); richard.levine@moffitt.org (R.L.)
- ⁷ Department of Gastrointestinal Oncology, Moffitt Cancer Center, Tampa, FL 33612, USA
- ⁸ Department of Satellite and Community Oncology, Moffitt Cancer Center, Tampa, FL 33612, USA
- ⁹ College of Medicine, University of Florida, Jacksonville, FL 32209, USA; alexander.parker@jax.ufl.edu
- ¹⁰ Georgetown University Medical Center, MedStar Health, Washington, DC 20007, USA; sandra.swain@georgetown.edu
- ¹¹ Indiana University School of Medicine, Indianapolis, IN 46202, USA; emtillma@iu.edu
- * Correspondence: james.hicks@moffitt.org; Tel.: +1-(813)-745-4668



Citation: Nelson, R.S.; Seligson, N.D.; Bottiglieri, S.; Carballido, E.; Cueto, A.D.; Imanirad, I.; Levine, R.; Parker, A.S.; Swain, S.M.; Tillman, E.M.; et al. Correction: Nelson et al. *UGT1A1* Guided Cancer Therapy: Review of the Evidence and Considerations for Clinical Implementation. *Cancers* 2021, 13, 1566. *Cancers* 2024, 16, 3595. <https://doi.org/10.3390/cancers16213595>

Received: 1 October 2024
Accepted: 9 October 2024
Published: 25 October 2024



Copyright: © 2024 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (<https://creativecommons.org/licenses/by/4.0/>).

Error in Table

In the original publication [1], there was a mistake in Table 1 as published. In certain instances, *UGT1A1* allele frequencies were inadvertently used to describe phenotype frequencies. The corrected Table 1 appears below. The authors apologize for any inconvenience caused and state that the scientific conclusions are unaffected. This correction was approved by the Academic Editor. The original publication has also been updated.

Table 1. Example *UGT1A1* alleles, predicted phenotype function, and frequencies among racial/ethnic groups.

Example <i>UGT1A1</i> Alleles and Predicted Function		
Star Nomenclature	Variant Type	Allele Function ^α
<i>UGT1A1</i> *36	(TA) ₅	Increased Function
<i>UGT1A1</i> *1	(TA) ₆	Normal Function
<i>UGT1A1</i> *6	(211G > A)	Decreased Function
<i>UGT1A1</i> *28	(TA) ₇	Decreased Function
<i>UGT1A1</i> *37	(TA) ₈	Decreased Function
Predicted <i>UGT1A1</i> Phenotypes Based on Commonly Observed Diplotypes		
Predicted <i>UGT1A1</i> Phenotype	Frequently Reported Diplotypes [Less Commonly Investigated Diplotypes] ^β	
	*1/*1 [*1/*36, *36/*36]	
Normal Metabolizer (NM)	*1/*28, *1/*6 [*1/*37, *6/*36, *28/*36, *36/*37]	
Intermediate Metabolizer (IM)		

Table 1. Cont.

Poor Metabolizer (PM)						
	*6/*6, *6/*28, *28/*28 [*6/*37, *28/*37, *37/*37]					
UGT1A1 Allele Frequencies Among Race/Ethnic Groups ^μ						
UGT1A1 Allele	African American/ Afro-Caribbean	Central/ South Asian	East Asian	European	Latino	Sub-Saharan African
*1	3%	54%	71%	36%	21%	49%
*6	<1%	4%	15%	<1%	1%	0%
*28	37%	41%	15%	32%	40%	40%
*36	8%	0%	0%	0%	0%	7%
*37	6%	0%	0%	<1%	0%	4%

α : UGT1A1 allele function per CPIC and prior investigations [6,11]. β : While allelic diversity continues to be recognized, reference laboratories may only test for certain polymorphisms such as *1, *6, and *28. μ : Table recreated from CPIC UGT1A1 Frequency Table [6,12].

Reference

- Nelson, R.S.; Seligson, N.D.; Bottiglieri, S.; Carballido, E.; Cueto, A.D.; Imanirad, I.; Levine, R.; Parker, A.S.; Swain, S.M.; Tillman, E.M.; et al. UGT1A1 Guided Cancer Therapy: Review of the Evidence and Considerations for Clinical Implementation. *Cancers* **2021**, *13*, 1566. [[CrossRef](#)] [[PubMed](#)]

Disclaimer/Publisher's Note: The statements, opinions and data contained in all publications are solely those of the individual author(s) and contributor(s) and not of MDPI and/or the editor(s). MDPI and/or the editor(s) disclaim responsibility for any injury to people or property resulting from any ideas, methods, instructions or products referred to in the content.