Good reads:

- https://www.nejm.org/doi/10.1056/NEJMp2116224?url_ver=Z39.88-2003&rfr_id=ori:rid :crossref.org&rfr_dat=cr_pub%20%200pubmed
- How Should Educators and Publishers Eliminate Racial Essentialism? | Journal of Ethics | American Medical Association (ama-assn.org)

Preclinical lectures

- Pathologizing race in preclinical lectures
- https://www.nejm.org/doi/full/10.1056/NEJMms2025768

Issues in board prep/study questions

- The use of race in these questions can promote biases
- https://www.tandfonline.com/doi/abs/10.1080/10401334.2016.1268056
- How Should Educators and Publishers Eliminate Racial Essentialism? | Journal of Ethics | American Medical Association (ama-assn.org)

Use of race in patient presentation

- Race in one liners can pathologize race
- https://journalofethics.ama-assn.org/article/mention-patients-race-clinical-presentations/2 014-06
- Mitigating Racism and Implicit Bias in Psychiatric Notes: a Quality Improvement Project Addressing How Race and Ethnicity Are Documented | SpringerLink
- <u>First Impressions</u> <u>Should We Include Race or Ethnicity at the Beginning of Clinical Case Presentations?</u> | NEJM
- <u>Differential Documentation of Race in the First Line of the History of Present Illness</u> PMC (nih.gov)

Clinical signs in Black and Brown Skin

- Images used to depict skin conditions or other physical findings are underrepresented in textbooks and preclinical curriculum materials
- https://www.blackandbrownskin.co.uk/mindthegap
- https://www.sciencedirect.com/science/article/pii/S0277953618300790
- https://www.ingentaconnect.com/content/wk/acm/2016/00000091/00000007/art00031

Race in GFR calculations

- In 2021 a task force from the National Kidney Foundation and American Society of Nephrology recommended immediate implementation of the creatinine equation refit without the race variable in all laboratories
- https://jamanetwork.com/journals/jama/fullarticle/2735726
- https://www.ncbi.nlm.nih.gov/pmc/articles/PMC8638402/
- https://www.thelancet.com/journals/lancet/article/PIIS0140-6736(20)32716-1/fulltext

Race in clinical algorithms

- Race adjusted algorithms have the potential to direct more attention or resources to white patients than others
- https://www.nejm.org/doi/full/10.1056/NEJMms2004740

Issues in textbooks

- Textbooks often describe unique disease profiles between races that aren't supported by the literature
- https://journals.lww.com/academicmedicine/Fulltext/2022/10000/Race_in_the_Reading__
 A Study of Problematic Uses.30.aspx
- https://journals.lww.com/academicmedicine/fulltext/2011/10000/Unsupported_Labeling_ of Race as a Risk Factor for 33.aspx
- How Should Educators and Publishers Eliminate Racial Essentialism? | Journal of Ethics | American Medical Association (ama-assn.org)

Racialized Text Non-Racialized Text

- The allele responsible for sickle cell anemia is particularly common among people of African descent; about 9% of African Americans are heterozygous for this allele. About 0.2% are homozygous and therefore have the symptoms of sickle cell anemia. In some groups of people in Africa, up to 45% of all individuals are heterozygous for this allele, and 6% are homozygous. Why is sickle cell anemia so common in Africa? It turns out that carriers of sickle cell anemia are more resistant to malaria, a common and serious disease in central Africa." (Raven & Johnson, 2002, p. 260)
- http://www.mhhe.com/biosci/genbio/raven6b/ information/olc/samplechapter.mhtml. Use this link and then view page 260 of Raven & Johnson (2002) to see the figures used in the experimental text that depicted the distribution of malaria and SCA only in Africa.
- 3 "Perhaps the best example is cystic fibrosis (CF), the most common fatal genetic disorder among Caucasians" (Raven & Johnson, 2002, p. 261).
- 4 Frequency among human births: cystic fibrosis: 1/2,500 Caucasians sickle cell anemia: 1/625 African -Americans

- "About 2 million Americans (0.6%) are carriers of the allele responsible for sickle cell anemia. Around 72,000 people have the symptoms of the disease because they are homozygous. However, in some groups of people in the world, up to 45% of all individuals are heterozygous for this allele, and 6% are homozygous and therefore have the symptoms of sickle cell anemia. Why is sickle cell anemia so common in some groups of people? It turns out that carriers of sickle cell anemia are more resistant to malaria, a common and serious disease in many parts of the world."
- http://www.understandingrace.org/humvar/ sickle_01.html Use this link to view the figures used in the control condition that depicted the distribution of malaria and SCA in all world populations.
- "Perhaps the best example is cystic fibrosis (CF)" (Raven & Johnson, 2002, p. 261).
- Frequency among human births: cystic fibrosis: 1/3500 sickle cell anemia: 1/5,000
- o For example cystic fibrosis should not only be thought of as a disease of 'white people' https://jamanetwork.com/journals/jama/article-abstract/2780564
 - Further clarification should be provided saying that when doing carrier screening, the residual risk after a negative test is calculated based on ethnic background (due to the different prevalence of specific variants in various populations), genetic counselors are specifically trained to discuss this with patients and ask about self-identified ethnicity

How to report race and ethnicity in medical and science journals

• https://jamanetwork.com/journals/jama/article-abstract/2783090

Tool and Clinical Utility	Input Variables	Use of Race	Equity Concern
Cardiology			
Guidelines-Heart Failure® (https://www .mdcalc.com/gwtg-heart-failure-risk-score)	Systolic blood pressure Blood urea nitrogen Sodium	is identified as nonblack. This addition increases the estimated probability of	The original study envisioned using this score to "increase the use of recommended medical therapy in high-risk patients and
Predicts in-hospital mortality in patients with acute heart failure. Clinicians are advised to use	Age Heart rate History of COPD Race: black or nonblack	death (higher scores predict higher mortality).	reduce resource utilization in those at low risk." The race correction regards black patients as lower risk and may raise the threshold for using clinical resources for black patients.
Cardiac surgery			
Risk Calculator ¹⁰ (http://riskcalc.sts.org/ stswebriskcalc/calculate) Calculates a pairent's risks of complications and death with the most common cardiac surgeries. Considers >60 variables, some of which are listed	Operation type Age and sex Race: black/African American, Asian, American Indian/Alaskan Native, Native Hawaiian/Pacific Islander, or "Hispanic, Latino or Spanish ethnic- ity"; white race is the default setting. BMI	The risk score for operative mortality and major complications increases (in some cases, by 20%) if a patient is identified as black. Identification as another non-white race or ethnicity does not increase the risk score for death, but it does change the risk score for major complications such as renal failure, stroke, and prolonged ventilation.	When used preoperatively to assess a patient risk, these calculations could steer minori patients, deemed higher risk, away from these procedures.
Nephrology		prolonged ventuation.	
Estimated glomerular filtration rate (eGFR)	Serum creatinine Age and sex Race: black vs. white or other	The MDRD equation reports a higher eGFR (by a factor of 1.210) if the patient is identified as black. This adjustment is similar in magnitude to the correction for sex (0.742 if female). The CKD-EPI equation (which included a larger number of black patients in the study population), proposes a more modest race correction (by a factor	Both equations report higher eGFR values (given the same creatinine measurement) for patients identified as black, suggesting better kidney function. These higher eGFR values may delay referral to specialist care or listing for kidney transplantation.
		of 1.159) if the patient is identified as black. This correction is larger than the correction for sex (1.018 if female).	
optn.transplant.hrsa.gov/resources/allocation -calculators/kdpi-calculator/) Estimates predicted risk of donor kidney graft fail- ure, which is used to predict viability of potential kidney donor.†	Age Hypertension, diabetes Serum creatinine level Cause of death (e.g., cerebrovascular accident) Donation after cardiac death Hepatitis C Height and weight HLA matching Cold is chemia	Increases the predicted risk of kidney graft failure if the potential donor is identified as African American (coefficient, 0.179), a risk adjustment intermediate between those for hypertension (0.126) and diabetes (0.130) and that for elevated creatinine (0.209–0.220),	Use of this tool may reduce the pool of Africa American kidney donors in the United States, Since African-American patients a more likely to receive kidneys from African American donors, by reducing the pool of available kidneys, the KDRI could exacer- bate this racial inequity in access to kidney for transplantation.
	En bloc transplantation Double kidney transplantation Race: African American		
Obstetrics			
.edu/PublicBSC/MFMU/VGBirthCalc/vagbirth .html)	Age BMI Prior vaginal delivery Prior YBAC Recurring indication for cesarean sec-	tion factors subtract from the estimated success rate for any person identified as black or Hispanic. The decrement for black (0.671) or Hispanic (0.680) is	The VBAC score predicts a lower chance of st cess if the person is identified as black or Hispanic. These lower estimates may dis- suade clinicians from offering trials of lab to people of color.
	tion African-American race Hispanic ethnicity	almost as large as the benefit from prior vaginal delivery (0.888) or prior VBAC (1.003).	
Urology STONE Score ^{15,16}	Sex	Produces a score on a 13-point scale, with	By systematically reporting lower risk for blac
Predicts the risk of a ureteral stone in patients who present with flank pain	Acute onset of pain Race: black or nonblack Nausea or vomiting Hematuria	a higher score indicating a higher risk of a ureteral stone; 3 points are added for nonblack race. This adjustment is the same magnitude as for hematuria.	patients than for all nonblack patients, the calculator may steer clinicians away from aggressive evaluations of black patients.
uticalc.pitt.edu/) Estimates the risk of UTI in children 2–23 mo	Age <12 months Maximum temperature >39°C Race: Describes self as black (fully or partially)	Assigns a lower likelihood of UTI if the child is black (i.e., reports a roughly 2.5-times increased risk in patients who do not describe themselves as black).	By systematically reporting lower risk for blac children than for all nonblack children, thi calculator may deter clinicians from pursu- ing definitive diagnostic testing for black
	Female or uncircumcised male Other fever source		children presenting with symptoms of UT
Oncology			
www3.mdanderson.org/app/medcalc/index .cfm?pagename=rectumcancer) Estimates conditional survival 1–5 yr after diag-	Age and sex Race: white, black, other Grade Stage Surgical history	White patients are assigned a regression coefficient of 1, with higher coefficients (depending on stage) assigned to black patients (1.18–1.72).	The calculator predicts that black patients wil have shorter cancer-specific survival from rectal cancer than white patients. Clinicia might be more or less likely to offer inter- ventions to patients with lower predicted
nosis with rectal cancer	Current age, age at menarche, and age	The calculator returns lower risk estimates	survival rates. Though the model is intended to help concept
Assessment Tool (https://bcrisktool.cancer	at first live birth First-degree relatives with breast cancer Prior benign biopsies, atypical biopsies Race/ethnicity; white, African American, Hispanic/Latina, Asian American, American Indian/Alaska Native, unknown	for women who are African American, Hispanic/Latina, or Asian American (e.g., Chinese).	tualize risk and guide screening decisions it may inappropriately discourage more a gressive screening among some groups o nonwhite women.
Breast Cancer Surveillance Consortium Risk Calculator ¹⁹ (https://tools.bcsc-scc.org/ BCSyearRisk/calculator.htm)	Age Race/ethnicity: white, black, Asian, Native American, other/multiple races, unknown	The coefficients rank the race/ethnicity categories in the following descending order of risk: white, American Indian, black, Hispanic, Asian.	Returns lower risk estimates for all nonwhite race/ethnicity categories, potentially redu ing the likelihood of close surveillance in these patients.
cancer in women with no previous diagnosis of	BIRADS breast density score First-degree relative with breast cancer Pathology results from prior biopsies		
Endocrinology			
Osteoporosis Risk Estimation) ²⁰ (https://www .mdapp.co/osteoporosis-risk-score-calculator -316/)	Rheumatoid arthritis History of fracture Age Estrogen use Weight	Assigns 5 additional points (maximum score of 50, indicating highest risk) if the patient is identified as nonblack	By systematically lowering the estimated risk of osteoporosis in black patients, SCORE may discourage clinicians from pursuing further evaluation (e.g., DXA scan) in blac patients, potentially delaying diagnosis ar
ate, or high risk for low bone density in order to guide decisions about screening with DXA scan	Race: black or not black		intervention.
www.sheffield.ac.uk/FRAX/tool.aspx)	Age and sex Weight and height Previous fracture	The U.S. calculator returns a lower fracture risk if a female patient is identified as black (by a factor of 0.43), Asian (0.50),	The calculator reports 10-yr risk of major os- teoporotic fracture for black women as le- than half that for white women with ident
jor osteoporotic fracture on the basis of patient demographics and risk-factor profile. Calculators are country-specific.;\$	Parent who had a hip fracture Current smoking Glucocorticoid use Rheumatoid arthritis Secondary osteoporosis Alcohol use, 23 drinks per day Femoral neck bone mineral density	or Hispanic (0.53). Estimates are not provided for Native American patients or for multiracial patients.	cal risk factors. For Asian and Hispanic women, risk is estimated at about half the for white women. This lower risk reported for nonwhite women may delay interventi with osteoporosis therapy.
	Height	In the U.S., spirometers use correction factors for persons labeled as black	Inaccurate estimates of lung function may result in the misclassification of disease
Uses spirometry to measure lung volume and the rate of flow through airways in order to diagnose and monitor pulmonary disease	Race/ethnicity	(10-15%) or Asian (4-6%).	severity and impairment for racial/ethnic minorities (e.g., in asthma and COPD). ²³