

Whose Signals Are Being Amplified? Toward a More Equitable Clinical Psychophysiology

Clinical Psychological Science
1–16


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DOI: 10.1177/21677026221112117

www.psychologicalscience.org/CPS

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Abstract

Research using psychophysiological methods holds great promise for refining clinical assessment, identifying risk factors, and informing treatment. Unfortunately, unique methodological features of existing approaches limit inclusive research participation and, consequently, generalizability. In this brief overview and commentary, we provide a snapshot of the current state of representation in clinical psychophysiology with a focus on the forms and consequences of ongoing exclusion of Black participants. We illustrate issues of inequity and exclusion that are unique to clinical psychophysiology and consider intersections among social constructions of Blackness and biased design of current technology used to measure electroencephalography, skin conductance, and other signals. We then highlight work by groups dedicated to quantifying and addressing these limitations. We discuss the need for reflection and input from a wider variety of affected individuals to develop and refine new technologies given the risk of further widening disparities. Finally, we provide broad recommendations for clinical-psychophysiology research.

Keywords

electrophysiology, minority groups, prejudice, psychophysiology, racial and ethnic attitudes and relations

Received 1/4/22; Revision accepted 5/30/22

Clinical-psychological science, like other fields of psychology, is beginning to reckon with historical and ongoing inequities, especially those related to race and racism. Race is socially constructed on the basis of perceptions of physical traits, such as skin color, hair, and facial features, and has been imbued with oppressive power (racism) through the creation and maintenance of a sociopolitical hierarchy (American Psychological Association [APA], 2021a, 2021b). Racism and other intersecting systems of oppression (e.g., sexism, homophobia, ableism, classism, xenophobia) are intertwined with clinical-psychological theory and practice (APA, 2021a; for a review, see Roberts et al., 2020).

In this overview and commentary, we aim to expand on recent work examining racial inequities in a subfield of clinical-psychological science that rests at the interface between body, brain, and mind: clinical psychophysiology. Psychophysiological methods hold great promise for refining assessment, identifying risk factors, and informing treatment (Hajcak et al., 2019). Clinical-psychophysiological research can bridge neurological

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and behavioral units of analysis, provide a medium for cross-species translation, and parse relevant mechanisms in clinical trials. This approach is a core priority of mental-health-research funding agencies, as exemplified by the National Institute of Mental Health Research Domain Criteria (Hajcak & Patrick, 2015; Insel, 2015; Patrick & Hajcak, 2016). Research questions catalyzed by the continued “neuroscientification of psychology” have increased demand for psychophysiological approaches, particularly in clinical psychology (Haslam et al., 2022). Furthermore, clinical psychophysiology continues to gain ground as the proliferation of more affordable and easy-to-use psychophysiological equipment makes these approaches more accessible to researchers (Kaye et al., 2016; Wilson, 2010).

Clinical psychophysiology is subject to the same issues of racial inequity that plague other subfields of psychology. Systematic reviews have shown mistrust in researchers to be a key barrier to participation of minoritized communities (George et al., 2014; Schmotzer, 2012). Although historical harm by the research and medical community has caused distrust and hesitance to participate in research among individuals with marginalized identities (Kennedy et al., 2007), some analyses suggest they are often as equally willing to participate in research when equitably recruited (Fisher & Kalbaugh, 2011; Wendler et al., 2005). Still, both historically and currently, psychopathology-research participant samples have been dominated by individuals with privileged identities, particularly White, college-educated, cisgender people (Fisher & Kalbaugh, 2011; Gatzke-Kopp, 2016; Roberts et al., 2020; Wendler et al., 2005). Because of long-standing systems of racial oppression, biased representation has been made invisible, and findings are generalized as if they were drawn from representative samples (Helms, 1993; Helms et al., 2006). As one example, a classic and well-replicated finding in clinical psychophysiology is that prepulse inhibition—the dampened neural response to a strong sensory stimulus when it is preceded by a weak sensory stimulus—is reduced in individuals with or at high risk for schizophrenia (Li et al., 2021; Swerdlow et al., 2014). This finding, considered a key biomarker of schizophrenia (Donati et al., 2020), is derived from studies of predominantly White participants, but the results are presumed to apply universally—even though schizophrenia is diagnosed up to 2 to 4 times more often in Black individuals than in White individuals (Jones & Gray, 1986; Olbert et al., 2018; Schwartz & Blankenship, 2014). Given complex issues surrounding racially biased diagnostic practices that may contribute to these figures, it remains to be seen whether this biomarker applies equally well across less homogeneous groups (Martinez, 2021).

However, psychophysiology broadly and clinical psychophysiology specifically also introduce unique forms of systemic bias that prove especially threatening to sample representativeness and, as a result, generalizability of findings. For example, electroencephalography (EEG) and electrodermal analysis (EDA) depend on technology originally developed for and refined with light skin and thin, straight hair (see Webb et al., 2022). As we discuss, individuals with darker skin and/or curly, tightly coiled, dense, or voluminous hair are systematically disqualified at disproportionate rates from clinical-psychophysiology research using these methods. These phenotypic characteristics, subject to discrimination because of technical limitations in clinical-psychophysiology research (i.e., “phenotypic bias”; Webb et al., 2022), are often observed in those racialized as Black, who are additionally subject to other forms of racial discrimination in clinical psychology more broadly. This intersection between forms of oppression in research—one that may be particularly glaring in clinical psychophysiology—results in unrepresentative samples and biased generalizations. As we discuss, recent advances in psychophysiological technology risk increasing this inequity in clinical psychophysiology as well as other related subfields of psychology (e.g., cognitive neuroscience).

Clinical and other psychophysiology researchers have a duty to publish robust, generalizable results. This duty is especially important given that studies including psychophysiological measures may be perceived by the public as particularly believable and trustworthy (Gatzke-Kopp, 2016; Weisberg et al., 2008). Because results using such measures are seen as more “objective,” they may be used to justify the development and implementation of nonrepresentative assessment tools and intervention approaches that are mistakenly applied universally. Thus, there is a critical and immediate need to address inequities in clinical psychophysiology.

Goals and Scope

We represent a range of career stages, cultural identities, and disciplines, including psychology, communication, neuroscience, engineering, private practice, and the hair-care industry. Our collective goal is to increase awareness of problems and potential solutions in addressing diversity, equity, and inclusion in all psychophysiology research using examples from clinical psychophysiology. To that end, in this commentary, we focus on issues of racial representation, particularly the systemic exclusion of Black research participants, in our subfield’s study samples, recognizing that this is

only one piece of a larger conversation with implications for multiple groups. First, we highlight recent efforts to quantify racial and ethnic representativeness in clinical-psychophysiology samples broadly and present results from a new literature review of three prominent academic journals. We then explore how phenotypic discrimination against dark skin and curly, tightly coiled hair in clinical-psychophysiology research intersects with anti-Black racism to contribute to sustained inequities and exclusion. In doing so, we emphasize that the problem of underrepresentation in psychophysiological research lies in the limitations of current equipment rather than the individuals whose phenotypic characteristics are not yet equitably served by psychophysiological technologies. Perhaps most importantly, we call attention to recent and burgeoning work on this topic, including snapshots of new data from groups led by researchers of color who are taking steps to increase inclusivity in this field. To conclude, we provide broad recommendations to help clinical psychophysiologicalists contribute to a more equitable clinical-psychological science.

The State of Racial/Ethnic Representativeness in Clinical Psychophysiology: A Preliminary Literature Review

In pursuing reform in clinical psychophysiology, a foundational step is to quantify existing issues with study-sample composition, including the degree to which participants of color—particularly Black individuals—are excluded from this research. However, most researchers using psychophysiology measures simply do not report such demographics for their final sample. In the introduction to a special issue on diversity and representation in *Psychophysiology*, the flagship journal of the Society for Psychophysiological Research, Gatzke-Kopp (2016) noted that greater than 85% of *Psychophysiology* articles in the preceding 3 years were missing a description of racial demographics. To our knowledge, this single statistic is the only existing quantification of the broad state of representativeness in psychophysiology research.

In an effort to develop a more nuanced picture over the last quarter century, we conducted a preliminary literature review of articles from *Psychophysiology*, which exclusively publishes research using such methods, and two prestigious clinical psychology journals that often publish articles using psychophysiological methods: *Journal of Psychopathology and Clinical Science* (*JPCS*; formerly *Journal of Abnormal Psychology*) and *Clinical Psychological Science* (*CPS*). To balance coverage and

efficiency, we used sparse sampling by reviewing and coding the main text and supplemental materials of every article published in *Psychophysiology* during 1997 to 2000, 2007 to 2010, and 2017 to 2020. We also identified articles from the same three time periods in *JPCS* and the latest time period in *CPS* (because *CPS* was established in 2013) that contained psychophysiology measures by searching in APAPsycNet and SAGE Journals and cross-referencing with PubMed for any of several psychophysiology-related terms derived from the *Psychophysiology* articles found (e.g., “skin conductance,” “electrophysiology”; for search terms, see the Supplemental Material available online). All articles were coded for article type (e.g., empirical report, literature review), region of sample recruitment (United States vs. non-United States), reported sample demographics (including sex, gender, race, ethnicity), and psychophysiological methods used. Here, we describe the proportion of articles that reported racial and/or ethnic sample composition, and among the articles that did so and used U.S.-based samples, we summarize the racial and/or ethnic demographics provided (see Figs. 1–3).

Out of 1,480 articles in *Psychophysiology* across all three time periods, we screened out “ineligible” articles that did not include empirical samples (e.g., commentaries, literature reviews), leaving 1,315 articles. Of these, only 182 (13.8%) included a description of the sample’s race and/or ethnicity. Reporting of race and/or ethnicity increased in 2017 to 2020 (17.7%) compared with 2007 to 2010 (12.1%) and 1997 to 2000 (8.0%). Although reporting of race and/or ethnicity has clearly improved following *Psychophysiology*’s special issue that strongly encouraged and emphasized the importance of doing so (Gatzke-Kopp, 2016), it remains inadequate, likely because it is still not explicitly required by the journal.

Out of 978 articles from the targeted time frame in *JPCS*, 161 included at least one psychophysiology term, of which 38 were deemed ineligible, mirroring the criteria for *Psychophysiology* (e.g., lacking an empirical sample or missing psychophysiological measurements), resulting in 123 articles for analysis. Among these, 62 (50.4%) included a description of sample race and/or ethnicity. Reporting of race and/or ethnicity improved across years (1997–2000: 29.0%; 2007–2010: 58.5%; 2017–2020: 56.9%) yet, like *Psychophysiology*, needs continued improvement. As an APA journal, *JPCS* officially follows the Journal Articles Reporting Standards (Kofalt, 2018), which include clear guidelines for demographics reporting across all article types. However, this reporting does not appear to be enforced by the journal.

Out of 155 articles from the 2017 to 2020 time frame reviewed in *CPS*, 87 included at least one psychophysiology term, of which 36 were deemed ineligible, resulting in 51 articles for analysis. Among these, 38 (74.5%)

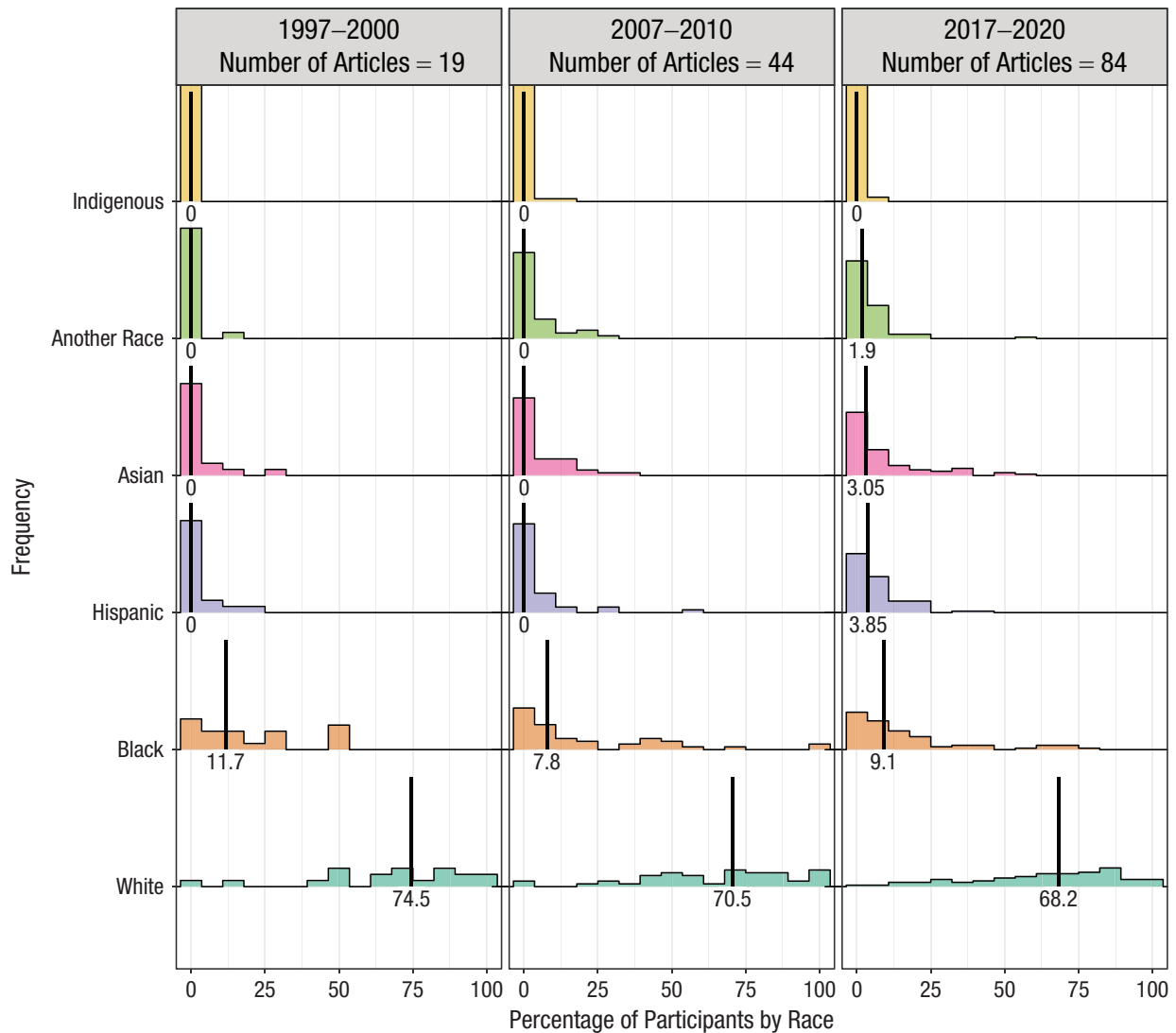


Fig. 1. Participant demographics among the 13.8% of *Psychophysiology* articles that reported sample race and ethnicity. Vertical black lines refer to the median percentage of each race/ethnicity across included articles. Note that medians are presented because of the clear overdispersion of percentage of races and ethnicities; medians will often not sum to 100%. “Another Race” includes all races not included in the listed categories, including biracial/multiracial individuals.

included a description of sample race and/or ethnicity. Reporting of race and/or ethnicity in *CPS* appears relatively frequent, and there is a high likelihood of continuous improvement with the implementation of new authorship guidelines. Under new editorship in 2021, *CPS* began explicitly requiring reporting of participant demographics including race and ethnicity.

When viewed across psychophysiology measures, these early results suggest that the prevalence of racial/ethnic demographic reporting in clinical-psychophysiology articles is beginning to increase but varies across journals and remains low in general. Given substantial variability in the number of psychophysiology articles per journal, differences in reporting rates between journals should be interpreted very cautiously. We speculate,

however, that articles in a methods-focused journal such as *Psychophysiology* may more often focus on assumed “basic” or “universal” mechanisms and processes than articles in clinical-psychology journals (i.e., *JPCS*, *CPS*) that may more often focus on clinically relevant mechanisms or applied processes that tend to be studied with greater consideration of the relevance of demographics such as racial or ethnic differences.

For those articles reporting racial and ethnic demographic data, we undertook informal analyses of the representativeness of their samples. Given our focus in this article on race and racism in the unique context of the United States (Ledgerwood et al., 2022), we restricted these analyses to U.S. samples. Such samples were identified according to the location of the

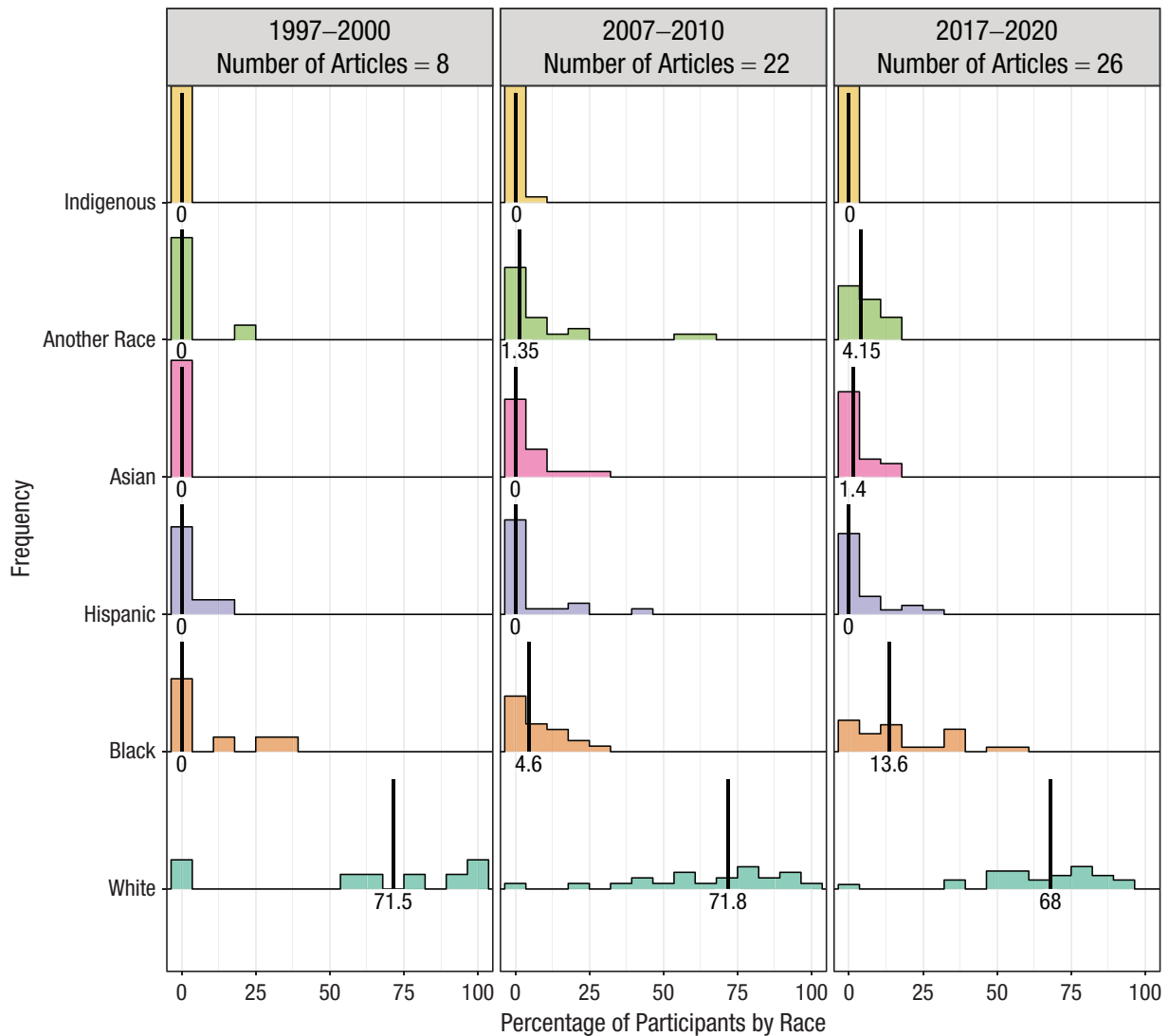


Fig. 2. Participant demographics among the 50.4% of *Journal of Psychopathology and Clinical Science* (*JPCS*; formerly *Journal of Abnormal Psychology*) articles that reported sample race and ethnicity. Vertical black lines refer to the median percentage of each race/ethnicity across included articles. Note that medians are presented because of the clear overdispersion of percentage of races and ethnicities; medians will often not sum to 100%. “Another Race” includes all races not included in the listed categories, including biracial/multiracial individuals.

institutional review board of record, explicitly stated region of recruitment in the Method section, and location of author affiliations. These inclusion criteria resulted in inclusion sample sizes of 147 articles from *Psychophysiology*, 56 from *JPCS*, and 33 from *CPS*. The demographics of these studies are depicted in Figures 1 through 3. In general, when demographics were reported, minoritized racial identities tended to be underrepresented compared with White participants. Note that no article in any of the three journals reported phenotypic descriptions such as skin color or hair texture, leading to a reliance on socially constructed categories of race.

The true representation of minoritized participants in research samples is likely to be even lower than these data suggest. There was a small proportion of articles reporting race and ethnicity; thus, these likely had a greater proportion of non-White populations than the broader literature (which often does not report racial demographics). It is possible that authors who report descriptions of race and ethnicity are more likely to recruit more diverse samples and in many cases are conducting research specifically focused on demographic differences, thus necessarily increasing the diversity of their samples. The potential reasons for exclusion of minoritized participants may also vary by

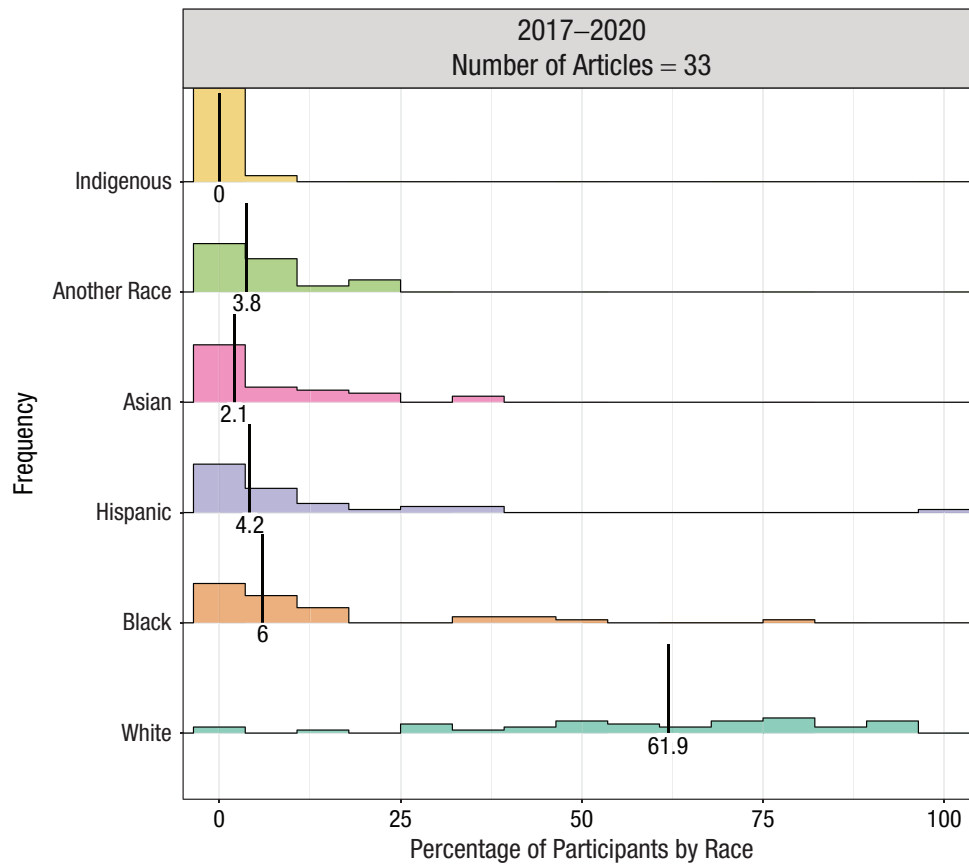


Fig. 3. Participant demographics among the 74.5% of *Clinical Psychological Science* (CPS) articles that reported sample race and ethnicity. Vertical black lines refer to the median percentage of each race/ethnicity across included articles. Note that medians are presented because of the clear overdispersion of percentage of races and ethnicities; medians will often not sum to 100%. “Another Race” includes all races not included in the listed categories, including biracial/multiracial individuals.

specific psychophysiology measure (for visualization of data for specific commonly used measures, see Figs. S1–S3 in the Supplemental Material; Literature review data and analysis code is available on Open Science Framework at <https://osf.io/hj3mq/>) and are discussed in greater depth below.

EEG: Recent Progress and Areas for Further Study

Of all clinical-psychophysiology measures, issues of equity in EEG appear to have gained the most attention and, consequently, steps toward progress. For researchers collecting EEG data, the problem is clear: Current EEG equipment is not designed for participants with curly, tightly coiled, dense, or otherwise voluminous hair.

EEG measures neuronally generated electrical signals on the scalp. As currently designed, this technology depends on contact between the electrode and scalp, either direct or through a conductive medium (e.g., saline gel); any impedance introduces noise into the signal and

degrades data quality (Luck, 2014; see Choy et al., 2022). Protocols for applying EEG caps prescribe simply moving strands of hair aside (Farrens et al., 2020), but this step presumes the participant has thin, straight hair that can be easily moved away from the intended scalp location. Moreover, in clinical-psychophysiology research, electrodes are typically embedded in a cloth cap, which imposes a further assumption that participants’ hair, when moved, will not push the cap so far away from the scalp as to impede scalp-electrode contact. Systems that rely on a chemical gel needed to bridge the scalp to the electrode require more gel for participants with curly, tightly coiled, dense, or otherwise voluminous hair. Larger amounts of gel can be detrimental for higher-density electrode montages (i.e., those with 64 channels and higher) because gel bridging between electrodes commonly produces signal artifacts. These technical limitations, based on manufacturer assumptions about participant hair, make scalp preparation for EEG more time-consuming, burdensome, and disruptive for individuals with curly, tightly coiled, dense, or voluminous hair.

For participants with these hair types, scalp preparation is sometimes terminated prematurely, but even when it is successfully completed, typical EEG equipment often produces noisy data for these individuals that is then excluded in postprocessing (Choy et al., 2022; Etienne et al., 2020). When data are included from participants with curly, tightly coiled hair, signals are often attenuated because of these technical limitations, risking the illusion that their neural responses are blunted when compared with participants with thinner, straighter hair (Etienne et al., 2020). Furthermore, some EEG researchers screen out potential participants with, for example, weaves, dreadlocks, or braids—all of which are typical hairstyles in the Black community—because of their perception that it will be difficult or impossible to secure the electrodes closely enough to the scalp to achieve an adequate level of contact (Choy et al., 2022). Note that curly, tightly coiled, dense, and voluminous hair is more prevalent among individuals of, for example, African, African American, and Caribbean ancestry (Loussouarn et al., 2007), who are more likely to be socially coded as Black. In these ways, such individuals are subject to not only phenotypic discrimination through the shortcomings of EEG equipment but also the myriad ways racism influences psychological research in general (e.g., the pervasive provision of cultural explanations for behavior of non-White individuals in contrast to psychological explanations for White behavior; Causadias et al., 2018).

Despite these issues, which are obvious to many in the field, quantification of the magnitude of the problem with representation in EEG is sparse. Choy and colleagues (2022) conducted a “proof of concept” survey of EEG research articles across 2 months in 2019, finding that only five of 81 articles (6.2%) reported that their sample included Black participants, although it was not stated how many of these articles reported any demographic information at all. Recognizing the need for more data, the Biomechanics, Rehabilitation, and Interdisciplinary Neuroscience (BRaIN) Lab at the University of Central Florida designed a detailed assessment of current practices in the field as part of its EEG Hair Project. This ongoing survey, available for participation at <https://hellobrainlab.com/research/eeg-hair-project/>, asks EEG researchers and participants about their own identities, EEG research experience, and perceptions about the influence of hair and other factors on participation in EEG studies.

As of this writing, more than 200 researchers have responded. Half ($\approx 49\%$) reported having recorded EEG data from fewer than five participants who identified as Black or African American, and the next highest percentage ($\approx 13\%$) reported having recorded data from only six to 10 such participants. A majority ($\approx 71\%$)

reported having excluded fewer than five participants with curly, tightly coiled, dense, or voluminous hair, but approximately 3% acknowledged excluding more than 50 such participants. These results, although highly preliminary, suggest that most researchers lack experience working with research participants with these hair types and that at least a small proportion of labs enroll but exclude them from analyses with regularity. These preliminary data suggest that the underrepresentation of Black participants across clinical EEG studies may be more attributable to initial underrecruitment than postenrollment exclusion for data-quality issues, at least in most labs.

The survey also asked about approaches to recording EEG from individuals with curly, tightly coiled, dense, or voluminous hair. Some researchers mentioned insightful solutions, such as flexible scheduling so data collection could occur just before salon appointments. Other responses highlighted the continuing demand for accessible solutions, with researchers noting that they “never found a solution” or simply “hope for the best” during data collection. Several respondents voiced an interest in specific training on collecting EEG from participants with often-excluded hair types.

Final results from the EEG Hair Project will deepen the understanding of the true magnitude of the problem as well as details about where in the pipeline (e.g., recruitment, data retention) the exclusion occurs. In the meantime, some researchers have initiated efforts to address these issues. For example, in the first project of its kind, Etienne et al. (2020) developed electrode casings that preferentially work on curly, tightly coiled, and dense hair. Their “Sèvo” product resembles a hair barrette containing a traditional electrode; these clips are then secured between cornrows that are braided in alignment with traditional EEG electrode layouts. The clips specifically harness the quantity and thickness of curly, tightly coiled hair to stabilize the electrode against the scalp at the specified site, thus celebrating these textures and empowering participants. Early trials of these electrodes suggest that they increase electrode-to-scalp contact and improve data quality for individuals with curly, tightly coiled, dense, or otherwise voluminous hair; future research is needed to determine whether such improvements extend to all hair types (Etienne et al., 2020; Kwasa et al., 2021).

Until such technology is refined and adopted by EEG hardware manufacturers, other practices can be implemented using existing equipment to obtain higher-quality data from participants with this hair type. For example, although presented in the context of hardware improvements, the braiding technique highlighted by Etienne et al. (2020) is a valuable approach to increase standard electrode contact. This technique

does not interfere with the standardized placement of electrodes, but it adds approximately 15 to 20 min of preparation time, depending on the size of the head and the coarseness of the hair. Given that hair braiding is a culturally familiar process and that scalp preparation is already often paired with other research tasks such as questionnaire completion, the relative costs of this extra time burden for some participants may be outweighed by the relative benefits of circumventing standard approaches that could prevent them from participating. Still, any approach that may systemically add time or other burden to a specific group of participants should be carefully considered from equity and research-integrity perspectives. More detailed guidance for hair preparation based on specific hair types—with multidisciplinary input, including a hair stylist with expertise in curly, tightly coiled hair—can be found in *A Guide to Hair Preparation for EEG Studies* (Richardson et al., 2021).

New organizations such as Black in Neuro (blackin neuro.com) and SPARK Society (sparksociety.org) are doing important work to raise awareness about systemic racism in brain sciences and empower researchers of color, including compiling educational and career resources online. Black in Neuro also hosts an annual #BlackinNeuro Week that includes panels, webinars, and other events discussing the existing biases in neuroscience, celebrating and encouraging researchers of color, and providing a call to action for others in the field. These organizations form the foundation of a necessary and overdue paradigm shift in awareness of inequity in clinical psychophysiology and related fields.

EDA: Initial Stirrings and Room for Growth

As inequity in EEG research gains greater attention, thanks to the efforts of advocacy groups, bias in EDA measurements was largely ignored until very recently. EDA has historically been considered a “gold-standard” psychophysiology measure of arousal and is often used to assess conditioned biological responses to a previously neutral stimulus that has been paired with an aversive stimulus (e.g., a mild electric shock; Fullana et al., 2020; Harnett et al., 2019; Kredlow et al., 2017; Rauch et al., 2012). These “fear learning” paradigms and associated fear-extinction processes serve as laboratory models of fear and anxiety. This work establishes an empirical basis for clinical treatments such as prolonged exposure therapy for posttraumatic stress disorder.

EDA involves administering small amounts of electrical current, usually to a participant’s hand, and measuring subtle changes in electrical impedance caused by

variations in the sweat response that accompanies arousal. Detecting these impedance changes is difficult for multiple reasons, and participant data are often discarded because of what is deemed too small of a response (i.e., “nonresponsiveness”). In addition, participants can be labeled “nonlearners” when differences are hard to detect between their response to a conditioned stimulus and a neutral stimulus, also leading to discarded data.

We propose that instead of labeling these individuals as nonresponders or nonlearners, researchers interrogate the inherent biases in EDA measurement. In particular, individuals who identify as African American or Black can appear to have lower skin-conductance levels and smaller conditioned responses than non-African American/Black individuals (Davis & Cowles, 1989; Janes et al., 1976; Johnson & Landon, 1965). As a result, Black participants are more likely than White participants to have their data discarded—something shown clearly in a review of fear-conditioning samples (Kredlow et al., 2017). Unlike EEG, whether EDA signals for Black participants are smaller solely because of technological limitations or additional sources of bias is unclear. Early research identified phenotypic factors that could affect EDA measurement fidelity, including number of active sweat glands (Boucsein, 1992; Kawahata & Adams, 1961; cf. Thomson, 1954; Wesley & Maibach, 2003), thickness of the upper layer of the skin (Berardesca & Maibach, 2003; Johnson & Corah, 1963; Weigand et al., 1974), electrolyte content of sweat (Johnson & Landon, 1965), skin resistance (Johnson & Corah, 1963; Juniper & Dykman, 1967), and skin temperature (Thomson, 1954). The majority of these studies on phenotypic factors were conducted in an era when research commonly relied on arbitrary social categories of race rather than thoughtful experimental design accounting for phenotypic differences between individuals. Unwarranted assumptions about biological differences between races (i.e., racial essentialism) risk exacerbating harmful disparities, akin to the harms imposed by societal belief in phrenology and eugenics. For example, Black individuals receive less treatment for pain because of erroneous assumptions about a shared racial biology that confers higher pain tolerance than White individuals possess (Hoffman et al., 2016).

Thus far, phenotypic factors have not been clearly demonstrated to account for racial differences in EDA. If they do, EDA technology should be made more robust to such phenotypic variation. Until then, skin-conductance data must be interpreted thoughtfully, accounting for all possible contributors to differences (e.g., Carter et al., 2021; Harnett et al., 2019). Although the harmful implications of discarding individuals’ EDA data have been touched on previously (Lonsdorf et al.,

2019), direct and clear attention to this equity issue has come only very recently (Webb et al., 2022).

Other Measures and New Technology: A Caution About the Future of Clinical Psychophysiology

Although EEG and EDA provide compelling and accessible examples of how psychophysiological approaches can lack inclusivity, especially for Black individuals, such problems extend beyond these measures and demographics. For example, the eye-blink startle response measured with electromyography is a translational measure widely used in studies touting direct clinical implications (Braff, 2010; Grillon & Baas, 2003), particularly in testing psychopharmacological therapeutics (Grillon & Ernst, 2020; Kaye et al., 2017). As with EDA, it is common practice to exclude participants with “small” general-startle reactivity at baseline, often using arbitrary criteria, because smaller or harder-to-detect signals are associated with noisier data (Blumenthal et al., 2005; Bradford et al., 2014, 2015). Given that recent studies suggest that individuals identifying as Hispanic, Asian, or Black exhibit decreased general-startle reactivity compared with White participants (Correa et al., 2021; Hasenkamp et al., 2008; Nelson et al., 2014), future work must ensure that individuals with marginalized social identities and/or phenotypic characteristics are not disproportionately excluded.

As psychophysiological technology advances, the field is at risk of magnifying bias embedded in these systems. Particularly worrisome is the rapidly advancing field of biomedical optics, which leverages the scattering and absorption properties of light in human tissue to infer underlying anatomy and function. Measurements by biomedical optical devices are influenced by the concentration of melanin (Mustafa et al., 2017; Sardar et al., 2001), the light-absorbing chromophore that gives skin its pigment. Thus far, optical studies do not typically report skin color and often use small samples of exclusively light-skinned participants. This technical limitation may skew interpretation of data from participants with darker skin, and the resulting issues of equity have not yet been discussed.

In one example of the use of optical technologies in clinical-psychophysiological research, increasing interest in measuring psychophysiology outside of the lab has led to the adoption of consumer-facing mobile devices for research purposes. Commonly used heart rate sensors, such as “fitness trackers” and “smart watches,” rely on light-based technology that is not currently optimized for darker pigmented (melanated) skin. These embedded biases have been highlighted multiple times in the popular press (Krisch & Schwartz, 2015; Rabin, 2020;

Tayag, 2020) but are only just beginning to be taken seriously by researchers (e.g., Shcherbina et al., 2017) and industry.

Just as mobile heart rate monitoring brings potential for increased bias, so do other optical technologies designed to be more affordable alternatives to their research-grade counterparts. For example, functional near-infrared spectroscopy (fNIRS) has been touted as a portable and affordable alternative to functional MRI. However, it has been known by NIRS experts for years that the technology does not work as well on darker skin (Wassenaar & Van den Brand, 2005). In addition, fNIRS requires good scalp contact, similar to EEG, but the suitability of current designs for diverse hair types has not been adequately discussed. It is an open secret in the field that fNIRS works best on fair skin and thin, blond (unpigmented) hair. In response, Meta Reality Labs recently funded six research groups to develop inclusive optical and other psychophysiology technology (Meta Research, 2021).

Interpreting Demographic Differences in Clinical-Psychophysiological Research

Thus far, we have largely focused on how psychophysiological technology is not currently optimized for phenotypic variations that map onto minoritized racial and ethnic groups. These technical limitations often lead to discriminatory exclusion of data from individuals with already marginalized identities. In the relatively rare cases when samples contain adequate representation of phenotypic and identity-based diversity, some differences have been demonstrated between racial groups. For example, as noted above, a blunted startle response has been observed for Black participants relative to White participants in some studies (e.g., Hasenkamp et al., 2008). Findings of this kind could be misinterpreted as reflecting true biological differences between racial groups, thus (erroneously) reifying the notion that race is physical. This is especially likely because of the privileged nature of psychophysiological findings, which are commonly viewed as more objective than other psychological research (Gatzke-Kopp, 2016; Weisberg et al., 2008). However, race is a social construction and must be differentiated from the phenotypic variations that only partially overlap with perceived racial categories (APA, 2021b). We have discussed how observed psychophysiological differences among racial groups (e.g., Black vs. White) could be attributable, at least in part, to limitations of psychophysiological technologies when working with phenotypic characteristics that are often observed in a given racial/ethnic group (e.g., dense, voluminous, curly hair vs. thin, straight hair). However, it is crucial not to dismiss all observed differences as spurious

consequences of technical limitations. Other factors, such as lived experience of racial discrimination, are likely to contribute to real differences in psychophysiology across identity groups, in tandem with these phenotypic influences.

Lessons for how to conceptualize these issues in clinical psychophysiology can be learned from cultural neuroscience. This field emerged in response to the presumed universality of the relationships among brain and behavior, regardless of cultural identity, in most neuroscience research (Chiao & Ambady, 2007). Note that cultural neuroscience is concerned with the interactional and multi-level influences of nuanced sociocultural and phenotypic factors. Cultural neuroscience was founded with ideas reminiscent of intersectionality (Crenshaw, 1989), a framework that “approaches conceptualizing the human experience through understanding multiple social identities and how they function in contextualized systems of inequality” (Grzanka, 2020, p. 249).

Cultural neuroscience provides a medium for interpreting racial and ethnic differences in psychophysiology. The body is intricately intertwined with other facets of human psychology such that lived experience influences physiological functions. For example, minority stress has downstream impacts on stress reactivity (Forrester et al., 2019; Huebner et al., 2021; Ruiz et al., 2018). Language differences are also associated with brain structure and function (see Kim & Sasaki, 2014). Furthermore, these social factors can interact with genetic differences—which contribute to phenotypic characteristics rather than explicitly to social categories of race—to produce observed physiological variability (Sasaki & Kim, 2017). Disentangling the influence of factors such as lived experience, language, and culture from the limitations of current psychophysiological measures is immensely challenging and will depend, in large part, on improvements in psychological science broadly.

In the meantime, one step may be to conduct post hoc analyses to test whether demographics account for variance in psychophysiological responses (Webb et al., 2022), but making such inferences is impossible without adequate representation of each demographic in such samples. Even with adequate representation, psychophysiology studies must continue to increase their typical sample sizes, which are often statistically underpowered for the detection of even medium-sized focal effects—much less the post hoc assessment of potentially small demographic differences (Correa et al., 2021). Another potential approach is to exclude as little data as possible by including data from participants with even the smallest signals. Researchers could then include participants’ general or baseline response as a covariate in statistical analysis to account for increased noise caused by shortcomings of the equipment (Bradford et al., 2014, 2015).

However, researchers must be careful when attempting to “control for” differences they believe are caused by bias in psychophysiological equipment because doing so may inadvertently parse out meaningful demographic variance of interest if it correlates with nuisance variance caused by equipment bias (Miller & Chapman, 2001). Finally, another approach to disentangling demographic differences attributable to technical issues with diverse phenotypes versus cultural differences of interest is to include multiple measures of social identity and lived experience. These decisions should be made in accordance with theory (see e.g., Mereish & Miranda, 2021). For example, if differences in EEG between Black and White participants are theorized to reflect minority stress, the study should explicitly measure minority stress and key covariates, such as self-reported hair texture, rather than relying on a single racial categorization as a proxy for both these potential mechanisms of physiological difference. Clearly, addressing these issues will require continued methodological and statistical work from a variety of affected individuals prioritizing social justice appropriately.

Broad Recommendations for Researchers Using Psychophysiology

Above, we suggested approaches to reduce inequity in EEG and EDA research. Below is a list of broad recommendations that cut across measures. We recognize that inequity in clinical psychophysiology is systemic, and addressing it will require a “top-down” approach with buy-in and action from powerful institutions (e.g., government agencies, funders, university departments). We wholeheartedly support the many thoughtful appeals previously made to individuals with the most power, including calls to address racial disparities in grant funding and earmark funding for equity-focused initiatives (Harnett & Ressler, 2021; Settles et al., 2020). At the same time, many readers of this article will not possess the institutional power to implement such suggestions. Accordingly, we provide actionable recommendations for researchers currently conducting or starting clinical-psychophysiological research. These recommendations have the potential to improve inclusiveness by increasing the fidelity of research with not only Black participants but also a range of people who differ from White individuals in terms of skin color and hair texture (e.g., Latine, South Asian).

Review your lab’s current practices

Recent public and scientific movements have provided a “conversation starter” to identify and address inequities in science. Practically speaking, clinical-psychophysiological data collection is slow-moving or even

paused because of the ongoing COVID-19 pandemic. These circumstances provide an opportunity for reflection and planning. Recommendations and resources highlighted above and below can provide a starting point. We acknowledge that adjusting recruitment and data-collection procedures to increase inclusivity requires substantial time and effort; this undertaking can and should be reported to evaluators as a valuable project during data-collection downtime. As funding agencies and academic institutions continue to publicly reinforce their stated commitments to diversity, equity, and inclusion, researchers must remind them to acknowledge (i.e., “count” and potentially reward) this valuable work. Once lab-level policies have been reformed, they should be revisited periodically as best practices evolve.

Be transparent in reporting of sample demographics and exclusions

As attention to diversity in clinical-psychophysiology samples grows, researchers may feel hesitant to highlight the lack of diversity in their studies and even more so to report that they disproportionately excluded individuals on the basis of phenotypic characteristics. As outlined above, significant barriers prevent researchers from collecting adequately representative samples in clinical-psychophysiology studies, and unfortunately, lack of diversity is still the norm (Rad et al., 2018). The demographics of final samples and excluded individuals must be reported transparently to aid efforts to quantify the magnitude of these problems and further justify (re)direction of resources toward their solutions. These efforts may be best aided if researchers collect and report not only self-reported race and ethnicity but also phenotypic characteristics, such as skin color and hair type, which may be more directly tied to the shortcomings of current psychophysiological approaches (e.g., Monk, 2015). Reforms such as fully online journals and exclusion of method sections from word counts further facilitate the thorough description of sample characteristics. For example, *CPS* now requires detailed reporting of participant demographics or acknowledgment of a lack of such detail as a weakness, consistent with the journal’s designation of diversity and representation as a priority area (Drew, 2020).

Bring typically excluded affected individuals to the table

Consistent with well-established but chronically underfunded approaches such as community-based participatory research (Chen et al., 2012; Ciccarella et al., 2018; Sandoval et al., 2012; Wallerstein & Duran, 2008),

researchers can engage affected individuals with diverse demographics characteristics and various lived experiences, allowing cultural translation through the expertise of native speakers and community partners. This will help ensure equity in all aspects of a research project. For example, researchers can organize focus groups with Black women to identify potential barriers to participation in EEG research and codevelop solutions. From information gathered during these focus groups, researchers might modify, for example, recruitment and study materials to use more inclusive and equitable content and language that address concerns about how EEG may affect hair or create an inconvenience, especially for individuals with hairstyles that require significant time and financial investment. For instance, recruitment materials might state that the hair gel is water-soluble or, if using dry electrodes, specify that study participation does not involve the use of hair gel. Researchers might also modify their recruitment procedures, such as explicitly offering flexible scheduling to accommodate participants’ individual hair-care routines or wash cycles.

Data collection for measures such as EEG can be aided by partnerships with local hair stylists who have expertise in specific hair types. Such partnerships can be facilitated by recruiting trainees with existing connections to their local communities. For example, the BRaIN Lab recruited a local Black hair expert only after successfully recruiting a Black graduate student who inquired with the student’s local network of natural-haired individuals. For researchers privileged enough to be in such a position, start-up or other funds could be earmarked for building these partnerships. Social media (e.g., Twitter, open-membership academic Slack groups) also facilitates connections among typically siloed groups and disciplines. In addition to directly improving representativeness and sociodemographic nuance in ongoing research, these practices would help to address the lack of diversity in authorship of published papers in psychology (Nunes, 2021).

Leverage the unique consumer power held by psychophysiology researchers to encourage inclusive change in hardware

Psychophysiology equipment is highly specialized, and manufacturers are exceptionally responsive to the requests of their small user bases. As a result, researchers have unique leverage to change problematic technical norms in clinical psychophysiology. We have personal experience with our mentors or colleagues requesting modifications to psychophysiology hardware to suit their lab’s specific needs (e.g., the addition of a new cable connection), with these changes implemented

in the product's next standard iteration. Likewise, clinical-psychophysiology researchers can advocate for the design and production of new equipment that addresses the pitfalls of current technology outlined above. With the continuing development of methods that ignore diversity in skin color and hair texture, phenotypic discrimination has the potential to become even more severe going forward if these limitations go unaddressed. Researchers must assert abandoning “color-blind” approaches in engineering psychophysiological equipment and hold manufacturers accountable as new technologies emerge.

Support existing efforts to improve clinical psychophysiology

As highlighted above, organizations such as Black in Neuro and SPARK Society are making great strides in increasing awareness of representation issues in clinical psychophysiology for both the general public and scientific communities. Interested researchers can contribute to such societies by joining and/or donating money and resources. Researchers can also participate in efforts such as the EEG Hair Project survey to aid in quantification and understanding of recruitment and exclusion practices. Aside from providing their own insight, one low-cost—but potentially high-impact—way for researchers to support these efforts is to share them in their circles through social media and laboratory or department websites.

Support can also occur at a more local level. Although much has been written about the need to recruit scholars with marginalized identities (McCormick-Huhn et al., 2019), there has been less discussion about providing the continued support and systemic restructuring needed to retain those individuals in science (De Los Reyes & Uddin, 2021; Settles et al., 2020). Researchers can work to support these individuals—even when they do not know them personally—by centering and elevating their voices in research (Sukhera & Palaniyappan, 2021). It is also important to adopt and encourage intersectional thinking while combating systemic biases stacked against marginalized researchers and their disciplines (Settles et al., 2020). Although these suggestions are not specific to clinical psychophysiology, the relative specialization of this discipline facilitates awareness of new research by junior scholars with marginalized identities as well as the ability to network at relevant conferences. Finally, even if researchers do not independently study issues of marginalization as a core content area, they can offer expertise in a given psychophysiological approach, quantitative modeling, or other specialized areas to such projects, thus contributing to the evolution of the field

in a more equitable direction (Uddin & De Los Reyes, 2021).

Concluding Remarks

Clinical-psychophysiological research holds great promise for refining assessment, identifying risk factors, and informing treatment. Despite continually increasing adoption of and enthusiasm for this approach, critical issues of representation and equity have been neglected. Consequently, the potential for this research to inform psychological interventions that decrease disparities and increase equity in mental health is lost. As anti-racism campaigns for systemic change slowly gain traction, researchers can take steps to increase inclusivity both immediately and over the long term and, in doing so, help ensure a more equitable future for clinical psychophysiology. Stated succinctly, clinical-psychological science is at an inflection point: “The time to reimagine our discipline is now” (Ledgerwood et al., 2022).

Transparency

Action Editor: Craig Rodriguez-Seijas

Editor: Jennifer L. Tackett

Author Contributions

D. E. Bradford and A. DeFalco contributed equally to this article. D. E. Bradford, A. DeFalco, E. R. Perkins, and K. J. Joyner conceptualized the manuscript. D. E. Bradford, A. DeFalco, E. R. Perkins, J. Kwasa, I. Carbajal, and F. R. Goodman drafted the manuscript, and all authors provided critical additions and revisions. A. DeFalco collected the literature review data under K. J. Joyner and D. E. Bradford's supervision. K. J. Joyner analyzed the literature review data and created the figures. L. N. S. Richardson, N. Woodley, L. Neuberger, J. A. Sandoval, and H. J. Huang designed, collected, and analyzed the survey data. All of the authors approved the final manuscript for submission.

Declaration of Conflicting Interests


The author(s) declared that there were no conflicts of interest with respect to the authorship or the publication of this article.

Funding


This project was partially supported by National Institutes of Health Grants 5R01AG054621-05 to H. J. Huang, F31MH122096 to E. R. Perkins, and F99 NS115331 to J. Kwasa.

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Acknowledgments

We acknowledge Justin Thomas, Harley Beach, and Danielle Jones at Florida State University and Jackson Cutler, Lindsay Beaman, Nico Doran, and Mia Tognoli at Oregon State University for their help collecting literature review data. We acknowledge E. K. Webb for early conceptual input and helpful comments on an initial draft.

Supplemental Material

Additional supporting information can be found at <http://journals.sagepub.com/doi/suppl/10.1177/21677026221112117>

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