A Conceptual Framework for Addressing Digital Health Literacy and Access Gaps in U.S. Underrepresented Communities

Leesi Saturday Komi¹, Ernest Chinonso Chianumba², Adelaide Yeboah Forkuo³, Damilola Osamika⁴, Ashiata Yetunde Mustapha⁵

¹Independent Researcher, Chicago IL, USA, <u>komileesi@gmail.com</u>

² School of Computing / Department of Computer Science & Information Technology,

Montclair State University, <u>ernestdivine74@gmail.com</u>

³Independent Researcher, USA; <u>ayeboahforkuo@gmail.com</u>

⁴Independent Researcher, Ohio USA, <u>osamikadamilola@yahoo.com</u>

⁵Kwara State Ministry of Health, Nigeria, <u>mustaphaashiata@gmail.com</u>

Corresponding Author: <u>komileesi@gmail.com</u>

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Abstract

Digital health technologies are rapidly transforming healthcare delivery in the United States, yet significant disparities in digital health literacy and access persist, particularly among underrepresented communities. These gaps exacerbate existing health inequities, limit the effectiveness of digital health interventions, and hinder the achievement of national health goals. This paper proposes a conceptual framework to address digital health literacy and access gaps in underrepresented populations, including racial and ethnic minorities, lowincome groups, rural residents, and individuals with limited English proficiency or disabilities. The framework is built on four interconnected pillars: 1) Digital Access Infrastructure, focusing on equitable internet connectivity and device availability; 2) Digital Health Literacy Development, encompassing culturally appropriate training, language accessibility, and inclusive learning methods; 3) Community-Based Engagement, promoting trust-building, peerled education, and local partnerships to drive adoption; and 4) Policy and Systems Integration, ensuring that federal, state, and local health systems embed digital equity in their strategies and programs. The framework draws on insights from public health, education, information technology, and community development. It emphasizes a participatory approach, where community voices inform intervention design and implementation. Case examples from U.S. initiatives—such as mobile telehealth units in rural Appalachia, bilingual health portals in Hispanic communities, and digital skills training in urban public housing—illustrate how these pillars can be operationalized for measurable impact. Addressing digital health literacy and access is critical for advancing health equity, improving patient engagement, and ensuring that digital health innovations reach and benefit all populations. The framework provides actionable guidance for policymakers, healthcare providers, educators, and technology developers to collaboratively reduce barriers and enhance the usability, accessibility, and effectiveness of digital health solutions. This work advocates for embedding digital inclusion into national health strategies and calls for sustained investment in infrastructure, education, and inclusive design. By centering equity and community engagement, the proposed framework aims to build a digitally empowered population equipped to participate fully in the evolving landscape of U.S. healthcare.

Keywords: Digital Health Literacy, Health Equity, Underrepresented Communities, Digital Divide, Telehealth Access, Digital Inclusion, Health Disparities, U.S. Healthcare, Community Engagement, Health Technology.

1.0. Introduction

The digital transformation currently reshaping U.S. healthcare represents a pivotal shift towards patient-centered care, driven by technologies including telemedicine, electronic health records (EHRs), remote monitoring, and artificial intelligence (AI) systems. These innovations have significantly broadened access to healthcare services, improved operational efficiencies, and personalized health interventions, fostering a more tailored patient experience (Steinhauser, 2021: Bajwa et al., 2021). For instance, the COVID-19 pandemic has accelerated the adoption of digital technologies such as telemedicine, highlighting the readiness of stakeholders to engage with digital innovations when they recognize potential benefits (Steinhauser, 2021). The integration of AI further enhances patient care by supporting clinical decision-making and operational processes (Mohammed et al., 2022). However, despite these advancements, the realization of equal benefits from digital healthcare transformations remains elusive, particularly for marginalized communities—including racial and ethnic minorities, low-income groups, older adults, rural populations, and those with limited English proficiency (Ferreira et al., 2025: Novelli et al., 2021).

The persistent disparities in health outcomes and access to digital health resources can be traced to systemic inequities encompassing education, income, and social determinants of health. These inequities create barriers that hinder the effective use of digital health tools, exacerbating existing health disparities. For example, limited broadband access and digital literacy inhibit the ability of underrepresented groups to utilize healthcare technology, resulting in delayed care and poorer health outcomes (Novelli et al., 2021; Ferreira et al., 2025). The phenomenon of digital health literacy, defined as the ability to seek, understand, and utilize digital health information, is crucial in this context. There is a significant segment of the population lacking the necessary skills and resources for navigating digital platforms, which further impairs their ability to access preventive care and manage chronic conditions (Nazeha et al., 2020; Konopik, 2023). It is essential to address these barriers to ensure that the benefits of digital health are equitably distributed, avoiding the risk of reinforcing existing health inequities.

To tackle these issues, a conceptual framework is proposed, aimed at understanding and mitigating the gaps in digital health literacy and access in underrepresented communities. This framework draws insights from various disciplines, including public health, digital equity, behavioral science, and community engagement, to inform intervention design and policy development. It emphasizes the need for a holistic approach that considers individual capabilities, community infrastructures, healthcare system responsiveness, and sociocultural contexts (Olivera et al., 2022; Gleiß et al., 2021). By linking digital health literacy to health equity, the framework proposes actionable strategies that can expand access to digital health tools while ensuring they are accessible, trusted, and effective for all demographics. Ultimately, this approach seeks to create a more inclusive healthcare system where digital transformation serves to enhance, rather than impede, the well-being of historically underserved communities (Nazeha et al., 2020; Ferreira et al., 2025).

2.1. Methodology

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) method was employed to structure and streamline the methodology for developing a conceptual framework to address digital health literacy and access gaps in underrepresented U.S. communities. A comprehensive search and review of 252 published articles, policy documents, and technical papers was conducted across reputable databases, journals, and institutional repositories. This search aimed to identify patterns, strategies, interventions, and technological innovations related to digital health literacy, healthcare access equity, and the deployment of AI-driven tools in underserved regions.

Following the identification phase, duplicate records were excluded, resulting in 210 unique articles eligible for screening. Titles and abstracts were assessed for relevance to digital health interventions, community-based outreach models, and the socio-technological factors affecting marginalized populations. During this screening stage, emphasis was placed on publications that explicitly discussed health communication, public-private partnerships, CRM systems in healthcare, policy development, and innovative delivery mechanisms for digital health solutions.

In the eligibility phase, 85 full-text articles were critically reviewed based on predefined inclusion criteria, including geographical relevance (U.S. context), research quality, focus on underrepresented groups, and empirical or conceptual contributions to digital health literacy and access. Articles that lacked methodological rigor, context-specific relevance, or actionable frameworks were excluded.

The final selection included 34 studies that provided substantial qualitative insights and/or conceptual advancements. These studies, including those by Abass et al. (2024) and Alemede et al. (2024), were instrumental in the development of the proposed framework. The synthesis incorporated models on patient engagement through CRM systems, AI applications in public health, community-centered interventions, and integration of mobile health platforms to bridge access disparities.

The outcome of this methodology is a data-informed conceptual model integrating technology-enhanced learning, localized outreach, AI-driven diagnostics, and public policy alignment to strengthen digital health literacy and ensure equitable access to healthcare services. This approach supports evidence-based policymaking and enables targeted interventions tailored to the unique barriers faced by underserved U.S. populations.

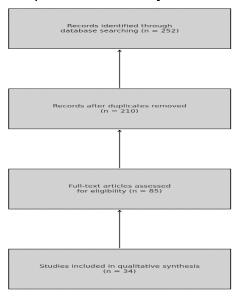


Figure 1: PRISMA Flow chart of the study methodology

2.2. Background and Context

Digital health literacy and digital access have become pivotal components of modern healthcare engagement in the United States. As healthcare systems increasingly rely on digital technologies for service delivery, patient education, monitoring, and communication, the ability to navigate and utilize these tools has become essential for patients to receive timely and appropriate care (Tomassoni, et al., 2012, Tomassoni, et al., 2013, Ugwu, et al., 2024, Zouo & Olamijuwon, 2024). Digital health literacy refers to an individual's capacity to seek, understand, evaluate, and apply health information from electronic sources to make informed

health decisions. It encompasses both technical proficiency with digital platforms and the cognitive skills required to interpret health-related data. Digital access, on the other hand, refers to the availability and usability of internet-connected devices, broadband connectivity, and supportive digital infrastructure that enable participation in online health services. The interplay between digital literacy and access determines the extent to which individuals and communities can benefit from innovations such as telehealth, mobile health applications, electronic health records, patient portals, and remote monitoring tools (Adelodun & Anyanwu, 2024, Chigboh, Zouo & Olamijuwon, 2024, Nwankwo, et al., 2024). Figure 2: Conceptual framework for use of digital health tools presented by Hernandez, et al., 2024.

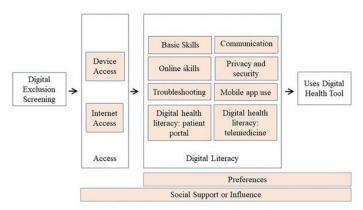


Figure 2: Conceptual framework for use of digital health tools (Hernandez, et al., 2024).

Underrepresented communities in the U.S. face significant challenges in both digital health literacy and access. These communities include racial and ethnic minorities, such as African Americans, Hispanic/Latino populations, Native Americans, and Pacific Islanders, who often experience structural inequities in education, income, and health. Low-income households, which may lack the financial resources to afford smartphones, tablets, computers, or broadband subscriptions, also fall disproportionately behind in digital participation (Ayo-Farai, et al., 2023, Chianumba, et al., 2023, Nnagha, et al., 2023). Rural residents face additional hurdles due to limited broadband infrastructure, geographic isolation, and lower density of healthcare providers offering telehealth services. People with disabilities encounter physical, sensory, or cognitive barriers that impede their ability to interact with digital interfaces, particularly when technologies are not designed with accessibility in mind. Non-English speakers, including immigrants and refugees, often encounter language-related barriers when navigating digital health content, which is predominantly available in English and often fails to reflect culturally relevant information or user-friendly design.

The persistence of the digital divide in the United States is reflected in national statistics. According to the Pew Research Center, as of 2021, approximately 25% of adults in households earning less than \$30,000 annually do not own a smartphone, and 40% lack a desktop or laptop computer. More than one-third of low-income households do not have broadband internet access (Akerele, et al., 2024, Edoh, et al., 2024, Ikese, et al., 2024, Olowe, et al., 2024). Among older adults, only about 60% report being confident in using digital technology, and digital adoption remains lower in communities of color. The Federal Communications Commission (FCC) has reported that rural communities continue to lag significantly behind urban areas in broadband coverage, with some areas lacking even basic internet connectivity. These disparities extend directly into the realm of health. A 2020 report from the National Digital Inclusion Alliance showed that counties with lower rates of broadband access also had higher rates of preventable hospitalizations, delayed diagnoses, and poor chronic disease management

outcomes. Conceptual framework for health literacy: The interaction of individual and systems domains presented by Gilder, et al., 2019, is shown in figure 3.

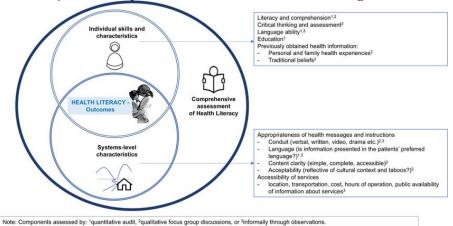


Figure 3: Conceptual framework for health literacy: The interaction of individual and systems domains (Gilder, et al., 2019).

Moreover, the COVID-19 pandemic further exposed and widened digital health disparities. As health systems pivoted to telehealth and digital communication tools to limit in-person contact, many patients in underrepresented communities were left behind. A study published in *JAMA Network Open* found that Black and Hispanic patients were significantly less likely to complete telemedicine visits than white patients, even after adjusting for socioeconomic factors. Similar disparities were observed among older adults, individuals with limited English proficiency, and those with Medicaid insurance (Nwankwo, Tomassoni & Tayebati, 2012, Olamijuwon, 2020, Tayebati, et al., 2010). These patterns highlight the compounding effects of digital exclusion on already vulnerable populations, amplifying existing health disparities and undermining public health efforts to deliver equitable care.

The barriers to digital health engagement are multifaceted and deeply rooted in broader structural inequalities. One of the most immediate barriers is limited access to technology and internet connectivity. For many households, the cost of broadband or cellular data plans is prohibitive, and public access points such as libraries or community centers may not be available or sufficiently equipped to support private health consultations (Abass, et al., 2024, Chianumba, et al., 2024, Matthew, et al., 2024). In rural areas, even when residents can afford internet services, infrastructure limitations may prevent reliable connectivity, limiting their ability to engage with telehealth or download essential applications. For individuals with disabilities, poorly designed interfaces, lack of compatibility with assistive technologies, and failure to follow accessibility standards often render digital health tools unusable or frustrating. Another major barrier is low digital literacy, particularly among older adults, individuals with lower educational attainment, and non-native English speakers. Many patients report difficulty using telemedicine platforms, patient portals, and mobile health apps due to confusing navigation, lack of guidance, or unfamiliar terminology. Without adequate support or training, these users are likely to abandon digital health tools or rely on intermediaries, such as family members, which may compromise privacy or autonomy. Even when digital literacy programs exist, they may not be tailored to the specific health contexts or cultural backgrounds of the users they aim to support (Alemede, et al., 2024, Chigboh, Zouo & Olamijuwon, 2024, Nwankwo, et al., 2024). Nkansah & Oldac, 2024, presented in figure 4, conceptual framework explaining the sources of digital literacy gap among students.

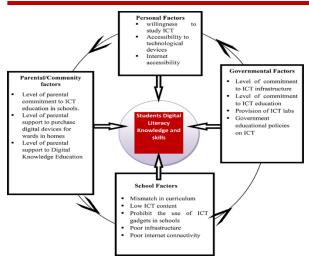


Figure 4: Conceptual framework explaining the sources of digital literacy gap among students (Nkansah & Oldac, 2024).

Trust also plays a crucial role in digital health engagement. Underrepresented communities, particularly Black and Native American populations, have a long history of experiencing systemic racism and exploitation in healthcare and research. This history contributes to skepticism toward new technologies, particularly those that collect personal data or appear to replace human interaction with automated systems. Data privacy concerns, fear of surveillance, and confusion about how digital information is used or shared further discourage participation. When digital health platforms fail to demonstrate transparency, security, and respect for user preferences, trust is eroded, and users disengage (Madu, et al., 2019, Matthew, et al., 2021, Nwankwo, et al., 2011, Tomassoni, et al., 2013).

Language and cultural barriers further inhibit meaningful digital health engagement. The dominance of English-language content and lack of cultural tailoring in digital health tools alienate users from diverse linguistic and ethnic backgrounds. For instance, a Hispanic patient navigating a telehealth platform may encounter difficulty if the system lacks Spanish translation or culturally relevant health information. Similarly, an immigrant from Southeast Asia may find it difficult to interpret digital diagnostic tools that do not consider traditional health beliefs or practices (Aderinwale, et al., 2025, Edwards, et al., 2025, Opia, et al., 2025). This lack of cultural competence in digital design and communication creates a disconnect between the tools and the people they are intended to serve.

Finally, healthcare system readiness and responsiveness also affect digital health engagement. In many cases, healthcare providers are not adequately trained or resourced to support patients in using digital tools. Clinics may lack staff to assist patients with portal registration, troubleshoot technology issues, or provide follow-up support for digital appointments. Some systems may not have integrated platforms that allow for seamless communication across digital and in-person care settings, leading to fragmentation and confusion (Balogun, et al., 2024, Edoh, et al., 2024, Ikese, et al., 2024, Olowe, et al., 2024). Providers may also unintentionally exclude patients with lower digital literacy by failing to offer alternative appointment types or assuming that all patients can engage with technology in the same way. In sum, the background and context for developing a conceptual framework to address digital health literacy and access gaps in U.S. underrepresented communities is characterized by both immense opportunity and significant inequity. While digital health innovations have the potential to revolutionize care delivery and improve outcomes, persistent digital divides threaten to leave behind those who are already most marginalized. The barriers to engagement are complex, spanning technological, educational, cultural, and systemic domains (Gabrielli,

et al., 2010, Imran, et al., 2019, Nwankwo, et al., 2012). A responsive and equity-focused framework must account for these intersecting challenges and guide the development of strategies that promote inclusion, build trust, and ensure that the digital transformation of healthcare truly benefits all individuals, regardless of their background or circumstances.

2.3. Review of Existing Literature and Models

The growing digitalization of healthcare in the United States has prompted the emergence of various frameworks and policies aimed at promoting digital health inclusion. These efforts recognize that equitable access to and use of digital health tools is essential for achieving health equity and improving outcomes for all populations. Numerous initiatives—both governmental and academic—have sought to define the principles, components, and implementation strategies necessary to enhance digital engagement across diverse communities (Adelodun & Anyanwu, 2025, Edwards, et al., 2025, Udegbe, et al., 2023). However, while progress has been made in laying foundational policies and highlighting the importance of digital health literacy, there remain significant gaps in the existing frameworks, especially in their ability to comprehensively address the unique challenges faced by underrepresented communities.

Several existing models provide a useful starting point for understanding digital health equity. For example, the *Office of the National Coordinator for Health Information Technology (ONC)* has developed a "Health IT Strategic Plan" which includes objectives to advance access, user-centered design, and digital literacy across the healthcare ecosystem. Similarly, the *National Digital Inclusion Alliance (NDIA)* advocates for equitable broadband access and digital literacy training through its digital inclusion frameworks (Edwards & Smallwood, 2023, Ekpechi, et al., 2023, Obianyo & Eremeeva, 2023). These efforts emphasize the importance of infrastructure, affordability, skills development, and device access as pillars of digital inclusion. The *Health Literacy Online* guide from the U.S. Department of Health and Human Services offers practical design strategies to make digital content more accessible and user-friendly, especially for people with limited health literacy.

In academic literature, the concept of eHealth literacy, as originally conceptualized by Norman and Skinner, has been widely used to define the multifaceted skills required for individuals to find, understand, evaluate, and apply health information from electronic sources. This model, known as the eHealth Literacy Lily Model, highlights six core literacies—traditional literacy, health literacy, information literacy, scientific literacy, media literacy, and computer literacy—as foundational to effective digital health engagement (Adegoke, et al., 2022, Chianumba, et al., 2022, Patel, et al., 2022). More recently, researchers have expanded the scope of eHealth literacy to incorporate social determinants of health and digital access barriers, particularly in underserved populations.

Furthermore, some community health models have begun to incorporate digital equity considerations into their interventions. The Digital Health Equity Framework (DHEF), proposed by Crawford and Serhal (2020), outlines how health system infrastructure, digital determinants of health, and individual digital health literacy interact to influence outcomes. The model emphasizes equity-oriented design and the role of systemic power dynamics in shaping access. Likewise, the Technology Acceptance Model (TAM) and its subsequent adaptations, such as TAM2 and the Unified Theory of Acceptance and Use of Technology (UTAUT), have been widely used to predict user adoption of technology, including in healthcare contexts (Kuo, et al., 2019, Matthew, et al., 2021, Nwankwo, et al., 2011, Tomassoni, et al., 2013). These models focus on perceived usefulness, ease of use, and user attitudes as predictors of digital engagement.

Despite the usefulness of these frameworks, several critical gaps persist in their applicability to underrepresented communities in the U.S. First, many existing models adopt a generalized, population-level approach to digital inclusion without sufficiently accounting for the

intersecting forms of disadvantage experienced by racial/ethnic minorities, low-income individuals, rural residents, non-English speakers, immigrants, and people with disabilities. These groups often face layered barriers that compound and interact—such as linguistic exclusion, cultural dissonance, systemic distrust, and historical marginalization—that are not adequately addressed by broad-stroke strategies (Ayo-Farai, et al., 2024, Edwards, et al., 2024, Nwankwo, et al., 2024).

Second, most frameworks focus primarily on the individual's digital competencies or on technological access without fully considering the sociocultural and systemic context in which digital health behaviors occur. For instance, while digital literacy training is vital, it is insufficient on its own if healthcare systems are not equipped to support patients with low literacy or provide culturally relevant, linguistically accessible, and trust-building communication (Akerele, et al., 2024, Edwards, et al., 2024, Ikhalea, et al., 2024, Zouo & Olamijuwon, 2024). Similarly, while improving broadband access is essential, this must be accompanied by community engagement and provider-side interventions to ensure digital tools are designed and delivered in a way that resonates with the lived experiences of underrepresented populations.

Another gap in current strategies is the limited integration of behavioral science and health equity principles into digital health frameworks. While the Technology Acceptance Model and similar approaches offer valuable insights into adoption behavior, they often lack an equity lens. They do not fully capture how social determinants—such as housing insecurity, unemployment, racial discrimination, or immigration status—shape an individual's readiness, motivation, or ability to engage with digital health tools (Babarinde, et al., 2023, Chianumba, et al., 2023, Ogundairo, et al., 2023). Moreover, current models rarely incorporate structural racism and historical exclusion as driving forces behind mistrust in digital platforms and health institutions. Without acknowledging these realities, digital health strategies risk reinforcing existing inequities and failing to reach those most in need.

Additionally, the current literature is limited in its guidance on how to operationalize digital health equity at the community level. While high-level policy goals are often articulated, there is little consensus or clarity on how to implement these strategies in diverse local contexts, particularly in under-resourced settings. Community health centers, local health departments, and grassroots organizations are often left without practical, adaptable models to guide their work in addressing digital health literacy and access disparities in their unique populations (Ariyibi, et al., 2024, Edwards, et al., 2024, Nwankwo, et al., 2024).

In light of these limitations, there is a strong rationale for developing a new conceptual framework tailored specifically to underrepresented communities in the U.S. Such a framework must move beyond a one-size-fits-all approach and instead embrace a multidimensional understanding of the barriers and facilitators to digital health engagement. It must be informed by equity principles, grounded in community realities, and designed to address the specific sociocultural, structural, and economic factors that influence health behaviors and access (Govender, et al., 2022, Matthew, Akinwale & Opia, 2022, Udegbe, et al., 2022).

A tailored framework should integrate key components such as structural determinants (e.g., broadband access, device affordability), healthcare system responsiveness (e.g., cultural competence, language accessibility, provider training), individual-level competencies (e.g., digital literacy, trust, user preferences), and community engagement strategies (e.g., co-design, local leadership, peer navigators). It must also recognize the importance of intersectoral collaboration between health, education, technology, and social service sectors to address the root causes of digital exclusion (Afolabi, Ajayi & Olulaja, 2024, Edwards, et al., 2024, Obianyo, Das & Adebile, 2024).

Moreover, the framework should be adaptive and iterative, allowing for continuous evaluation and refinement based on community feedback, emerging evidence, and technological changes.

It should include actionable steps for implementation, such as policy recommendations, programmatic interventions, workforce development initiatives, and evaluation metrics aligned with health equity outcomes. By grounding the framework in the realities of marginalized communities and centering their voices in its development, it can serve not only as a guide for intervention design but also as a tool for advocacy, funding allocation, and cross-sector collaboration (Nwankwo, Tomassoni & Tayebati, 2012, Tayebati, Nwankwo & Amenta, 2013, Tomassoni, et al., 2013).

In conclusion, while existing literature and models provide a valuable foundation for advancing digital health literacy and access, they fall short of fully addressing the needs of underrepresented communities in the U.S. A new, equity-driven conceptual framework is needed—one that reflects the complexities of digital exclusion, acknowledges historical and structural barriers, and provides clear, context-sensitive pathways for meaningful engagement. This framework must bridge theory and practice, centering inclusion, trust, and justice as core principles in the pursuit of digitally equitable healthcare for all (Adewuyi, et al., 2024, Edwards, Mallhi & Zhang, 2024, Ohalete, et al., 2024).

2.4. The Conceptual Framework

The conceptual framework for addressing digital health literacy and access gaps in U.S. underrepresented communities is grounded in a holistic and equity-centered approach that acknowledges the multifaceted barriers these populations face. It organizes the pathway to inclusive digital health transformation around four interdependent pillars: digital access infrastructure, digital health literacy development, community-based engagement, and policy and systems integration (Ayo-Farai, et al., 2023, Chianumba, et al., 2023, Katas, et al., 2023). Together, these pillars form a roadmap that not only addresses immediate technological deficits but also challenges systemic inequities, fosters trust, and builds sustainable, community-informed solutions that align with broader health equity goals.

The first pillar, digital access infrastructure, recognizes that without the foundational tools and connectivity to participate in digital healthcare, no amount of training or outreach can bridge the gap. Many underrepresented communities, particularly low-income households and rural residents, continue to face chronic deficits in internet access and device availability. Ensuring universal broadband coverage—both fixed and mobile—is essential. This includes expanding internet service to remote areas and subsidizing broadband costs for households that cannot afford monthly fees (Anyanwu, et al., 2024, Ekwebene, et al., 2024, Obianyo, et al., 2024). Device affordability and availability are equally critical. Smartphones, tablets, and computers must be accessible to every household, not just in terms of price, but also through programs that provide low-cost or free devices, particularly to seniors, persons with disabilities, and marginalized youth. Additionally, digital access must be designed inclusively. This means integrating accessibility tools for individuals with disabilities—such as screen readers, voice commands, alternative input devices, and adjustable visual settings—into all digital health platforms. Ensuring compliance with the Web Content Accessibility Guidelines (WCAG) is not optional but foundational to equitable digital health design.

The second pillar focuses on digital health literacy development, which addresses the knowledge and skills required to effectively use digital health technologies. Digital literacy is not merely about navigating a website or downloading an app—it is about understanding how to evaluate health information, manage personal health data, and engage confidently with telehealth, patient portals, or mobile apps (Ajayi, et al., 2024, Emeihe, et al., 2024, Johnson, et al., 2024, Olowe, et al., 2024). For underrepresented communities, effective literacy programs must be rooted in culturally and linguistically appropriate approaches. Educational content should be delivered in multiple languages, reflect diverse health beliefs and practices, and be created with sensitivity to the lived experiences of the target population. This includes using

trusted messengers and tailoring materials to community-specific norms. Content should also be presented using plain language, avoiding medical jargon or complex technical terminology. Visual aids, videos, infographics, and interactive learning tools can enhance understanding for users with limited reading proficiency or cognitive impairments. Public libraries, schools, health centers, and community hubs can serve as venues for delivering in-person and virtual training sessions focused on basic digital skills, privacy and security, and navigating specific digital health services. These efforts should be sustained and adaptive, recognizing that digital tools evolve, and continuous learning must be supported.

The third pillar of the framework, community-based engagement, emphasizes the role of trust, local leadership, and grassroots participation in closing digital health equity gaps. Many underrepresented communities harbor justified skepticism toward healthcare systems and government technologies, rooted in historical neglect, systemic discrimination, and previous exploitative practices. Building trust requires meeting people where they are, both literally and figuratively (Fuko, et al., 2025, Matthew, Nwaogelenya & Opia, 2025, Usuemerai, et al., 2024). One of the most effective strategies is the deployment of peer educators and digital navigators—community members trained to help others use digital health tools. These individuals not only offer technical assistance but also serve as cultural brokers who can translate unfamiliar systems into familiar contexts. Community health workers, promotores de salud, and members of faith-based organizations are especially effective in this role, as they often maintain close relationships with residents and have credibility that institutions lack. Faith-based institutions, local nonprofits, schools, and neighborhood associations can act as trusted intermediaries for digital health initiatives. These organizations can host digital health literacy workshops, facilitate access to devices, or serve as distribution points for educational materials. Importantly, the framework calls for participatory design approaches in developing digital health tools and programs (Adelodun & Anyanwu, 2024, Emeihe, et al., 2024, Majebi, Adelodun & Anyanwu, 2024). This means involving community members from the outsetduring needs assessments, program planning, and solution development—so that interventions reflect real-life challenges and preferences. Community feedback loops, user testing with target populations, and collaborative design sessions are essential to ensure that tools are intuitive, relevant, and empowering.

The fourth pillar is policy and systems integration, which focuses on institutionalizing digital equity within the healthcare system and broader policy environment. Embedding digital health equity into healthcare policies requires commitment at every level—from federal guidelines to local clinic protocols. Health systems must adopt policies that require digital health platforms to be inclusive, accessible, and available in multiple formats and languages (Akerele, et al., 2024, Emeihe, et al., 2024, Kelvin-Agwu, et al., 2024). Reimbursement models should incentivize healthcare providers to offer digital services equitably, including telehealth for Medicaid populations, services in multiple languages, and accommodations for patients with disabilities. Healthcare organizations should be required to collect and report digital access and usage data disaggregated by race, ethnicity, income, geography, and disability status to identify disparities and guide targeted interventions.

Incentivizing technology developers to design with equity in mind is also critical. Federal and state agencies can establish grants, procurement preferences, or certification programs that reward developers who meet accessibility and cultural responsiveness benchmarks. Health IT vendors should be encouraged to collaborate with communities during design and implementation, and to build platforms that are adaptable to the specific needs of safety-net providers and underserved populations (Abisoye & Olamijuwon, 2022, Chianumba, et al., 2022, Udegbe, et al., 2023). Moreover, public sector agencies—particularly the Department of Health and Human Services (HHS), Federal Communications Commission (FCC), and Centers for Medicare & Medicaid Services (CMS)—must coordinate on national digital equity

initiatives that address both health and infrastructure needs. These include expanding programs like the Affordable Connectivity Program, integrating digital literacy into Medicaid managed care contracts, and requiring digital health tools funded with public dollars to adhere to inclusive design principles.

State and local governments also play a crucial role in implementing digital health equity policies. They can lead statewide digital inclusion plans, fund community-based digital navigators, and integrate digital literacy support into public health campaigns and emergency preparedness plans. Health departments, especially those serving rural or underserved counties, can collaborate with schools, libraries, and local businesses to create digital health access points, similar to telehealth kiosks or digital health hubs, in accessible community spaces (Ayo-Farai, et al., 2024, Emeihe, et al., 2024, Kelvin-Agwu, et al., 2024). Importantly, all policies and systems-level interventions must be accompanied by accountability mechanisms. This includes periodic equity audits, community advisory councils, and public reporting of progress toward digital inclusion benchmarks.

Taken together, the four pillars of this conceptual framework offer a comprehensive and actionable model for addressing digital health literacy and access gaps in underrepresented U.S. communities. By ensuring foundational digital access, building user capabilities, engaging trusted local actors, and institutionalizing equity in policy and system design, this framework aims to close the digital divide not just technologically, but socially and structurally. The intersectionality of race, class, disability, geography, language, and history must inform every aspect of digital health planning and implementation (Adhikari, et al., 2024, Eze, et al., 2024, Johnson, et al., 2024). Only then can digital health tools truly serve as instruments of inclusion, rather than amplifiers of inequality. Through the coordinated action of community organizations, health systems, policymakers, and technology innovators, this framework envisions a future in which every person—regardless of background—can equitably access, understand, and benefit from the digital transformation of healthcare.

2.5. Operationalization of the Framework

Operationalizing a conceptual framework for addressing digital health literacy and access gaps in underrepresented communities in the United States requires translating abstract pillars into practical, context-sensitive actions. It entails embedding the framework into the daily operations of health systems, community organizations, and public institutions. This process demands a clear strategy for implementation, robust mechanisms for monitoring progress, and collaboration among a wide array of stakeholders (Elujide, et al., 2021, Khosrow Tayebati, et al., 2011, Nwankwo, et al., 2012). The success of operationalizing this framework hinges on real-world applications, meaningful metrics, and partnerships that bridge gaps between healthcare, technology, policy, and community leadership.

Across the country, a growing number of real-world initiatives exemplify how components of this framework are being effectively implemented. These initiatives provide concrete models for how digital access infrastructure, literacy development, community engagement, and policy alignment can come together to serve underrepresented populations. One such example is the use of rural telehealth vans in states like West Virginia, Arkansas, and Mississippi (Adelodun & Anyanwu, 2025, Ekpechi, et al., 2025, Usuemerai, et al., 2024). These mobile health units bring internet-enabled services directly to remote and medically underserved communities, eliminating travel barriers and providing both in-person and virtual care access. Outfitted with high-speed connectivity, diagnostic equipment, and trained staff, telehealth vans not only deliver immediate healthcare services but also serve as training hubs for digital health literacy. Patients are assisted in downloading and using health apps, setting up patient portal accounts, and learning how to navigate telehealth platforms. This direct, hands-on engagement in a

trusted environment helps reduce fear, increase confidence, and build digital familiarity among rural residents.

Another notable implementation occurs through digital health programs embedded in community centers and libraries. In cities such as Chicago, Detroit, and Baltimore, public libraries have transformed into digital equity hubs, offering free Wi-Fi, loanable devices, and personalized tech support. These facilities often host digital health literacy workshops cofacilitated by community health workers or digital navigators. Participants receive instruction on using telemedicine platforms, interpreting lab results online, accessing educational content, and safeguarding personal health data (Okoro, et al., 2024, Olamijuwon & Zouo, 2024, Olorunsogo, et al., 2024). Libraries serve as safe, trusted, and stigma-free spaces, which is crucial for reaching older adults, low-income individuals, and non-English speakers who may hesitate to engage with clinical institutions. Furthermore, some public libraries have partnered with local health departments to offer virtual provider consultations in private rooms, making digital health more accessible for people without home connectivity.

Public housing programs also offer valuable examples of operationalizing the framework. In cities like San Francisco and New York, housing authorities have implemented pilot programs that provide residents with subsidized internet, tablets, and ongoing tech support. Health-focused components of these programs include training residents to schedule virtual medical visits, refill prescriptions online, and monitor chronic conditions using mobile apps or Bluetooth-enabled devices (Maduka, et al., 2023, Majebi, et al., 2023, Ogundairo, et al., 2023). Importantly, these initiatives are co-designed with tenant associations, ensuring that the content and services are relevant, accessible, and culturally appropriate. This approach exemplifies participatory design and community-based engagement, two vital components of the framework. Residents not only receive services but also contribute to shaping how these services are delivered, increasing trust and uptake.

Measurement and evaluation are central to the operationalization process, providing evidence of impact, identifying areas for improvement, and justifying continued investment. Evaluation strategies must be both quantitative and qualitative, combining data-driven metrics with community-informed insights. Key performance indicators (KPIs) may include the number of individuals who gain internet access, complete digital literacy training, create patient portal accounts, or engage in telehealth visits (Alemede, et al., 2024, Eze, et al., 2024, Katas, et al., 2024, Obianyo, et al., 2024). Evaluators should disaggregate data by race, ethnicity, age, income level, disability status, and language to detect disparities and tailor interventions accordingly. For example, a telehealth program reporting high usage overall might still be failing to reach Spanish-speaking residents or older adults with cognitive impairments unless these details are surfaced through disaggregated analysis.

In addition to utilization metrics, evaluation should capture outcomes related to patient empowerment, care continuity, and health improvement. Surveys, focus groups, and interviews can be used to assess user satisfaction, confidence in using digital tools, perceived barriers, and trust in the healthcare system. Outcome-based indicators might include improvements in blood pressure control among patients using remote monitoring, reduced missed appointments due to increased telehealth adoption, or enhanced medication adherence following digital reminders (Abass, et al., 2024, Eze, et al., 2024, Johnson, et al., 2024, Olowe, et al., 2024). Longitudinal tracking is valuable for understanding the sustained impact of interventions and the extent to which digital health tools are integrated into patients' daily routines.

A rigorous feedback loop is essential. Insights from evaluation must be fed back into the design and implementation processes, allowing for continuous refinement. For example, if evaluations show low usage of a particular platform among individuals with disabilities, developers and program managers must work with accessibility advocates to redesign the interface, improve user instructions, or add compatibility with assistive technologies. In this way, evaluation

becomes not just a reporting tool, but a driver of innovation and inclusion (Chukwuma, et al., 2022, Gbadegesin, et al., 2022, Udegbe, et al., 2023).

Effective operationalization also depends heavily on coordinated stakeholder roles and cross-sector partnerships. Healthcare providers must be central actors, not only offering digital health services but also training their staff to identify and address digital literacy challenges among patients. Clinics should routinely assess patients' digital readiness and offer tailored support, just as they screen for social determinants of health. Providers can also partner with community organizations to offer outreach and support in non-clinical settings, expanding their reach and building trust with hard-to-engage populations (Kuo, et al., 2019, Madu, et al., 2020, Nwankwo, et al., 2012, Tayebati, et al., 2011).

Technology companies and developers play a crucial role in ensuring that digital health platforms are user-friendly, accessible, and inclusive. These companies must work closely with community representatives and health equity experts to incorporate universal design principles, language options, culturally relevant content, and privacy safeguards that reflect the concerns of marginalized users (Balogun, et al., 2023, Eyeghre, et al., 2023, Mgbecheta, et al., 2023). Moreover, funders—whether governmental, philanthropic, or corporate—should align their grant-making and investment strategies with the framework's pillars. They should prioritize multi-year funding that allows for program development, community input, evaluation, and scaling.

Government agencies at the federal, state, and local levels have a critical role in policy alignment and resource mobilization. Agencies like the Department of Health and Human Services (HHS), the Federal Communications Commission (FCC), and Centers for Medicare and Medicaid Services (CMS) must ensure that digital inclusion is embedded into health policy, broadband policy, and Medicaid and Medicare reimbursement schemes. This includes expanding support for telehealth in underserved areas, funding digital literacy initiatives, and mandating accessible design for federally funded digital health tools (Nwankwo, Tomassoni & Tayebati, 2012, Ogbonna, et al., 2012, Tayebati, et al., 2013). State health departments can support regional collaboratives, develop workforce training programs, and facilitate data sharing to monitor digital inclusion progress.

Lastly, community-based organizations, including faith-based institutions, advocacy groups, and neighborhood associations, must be recognized not as peripheral actors but as equal partners. Their intimate knowledge of local needs, histories, and networks enables culturally sensitive and contextually appropriate implementation. These organizations should be included in decision-making bodies, evaluation processes, and funding pipelines to ensure sustained community ownership (Adelodun & Anyanwu, 2024, Ezeamii, et al., 2024, Majebi, Adelodun & Anyanwu, 2024).

In sum, operationalizing the conceptual framework for addressing digital health literacy and access gaps requires a dynamic, multi-level approach grounded in equity, collaboration, and adaptability. It must bridge policy vision with ground-level action, marrying infrastructure expansion with human-centered design and community empowerment. Real-world initiatives offer promising blueprints, but scaling their success depends on rigorous evaluation, inclusive partnerships, and sustained political and financial commitment (Akerele, et al., 2024, Ezeamii, et al., 2024, Kelvin-Agwu, et al., 2024). With intentional effort and coordinated leadership, this framework can serve as a transformative guide for ensuring that digital health innovation benefits all communities—not just those already well-connected.

2.6. Challenges and Considerations

Implementing a conceptual framework for addressing digital health literacy and access gaps in underrepresented communities in the United States presents a promising vision for inclusive and equitable healthcare. However, translating this vision into practice is not without its

challenges. Despite compelling evidence supporting the potential of digital health innovations, several key considerations—namely sustainability and funding, privacy and trust concerns, and interoperability and technological adaptability—pose significant obstacles that must be addressed to ensure the effectiveness and longevity of these effort (Adaramola, et al., 2024, Ezeamii, et al., 2024, Ohalete, et al., 2024)s.

Sustainability remains one of the most persistent challenges facing digital health equity initiatives. Many programs targeting underserved populations are initially funded through time-limited grants, pilot projects, or philanthropic investments. While these resources often provide the necessary capital to launch digital literacy programs, purchase devices, or deploy telehealth platforms, they do not guarantee continuity. When external funding ends, programs risk scaling down or halting entirely, leaving communities with fragmented services and unfulfilled expectations (Adelodun & Anyanwu, 2025, Ekpechi, et al., 2025, Usuemerai, et al., 2024). This disruption not only wastes prior investments but can also erode community trust in future digital health efforts.

Ensuring long-term sustainability requires integrated funding strategies that move beyond temporary or project-based models. Federal and state governments must prioritize digital health equity in healthcare budgets, embedding these efforts into Medicaid and Medicare programs, public health initiatives, and broadband infrastructure investments. Additionally, reimbursement models must evolve to recognize and support digital inclusion activities (Okoro, et al., 2024, Olamijuwon, et al., 2024, Olorunsogo, et al., 2024). For instance, Medicaid could reimburse community health workers for digital navigation services, or Medicare Advantage plans could cover technology training for older adults. At the same time, public-private partnerships can play a vital role in sustaining programs, but these collaborations must be carefully designed to align with equity goals rather than commercial interests. The risk is that in the absence of strong regulatory guardrails, private sector involvement may prioritize profit over accessibility or contribute to widening the digital divide.

Another challenge is the complexity of evaluating return on investment (ROI) for digital inclusion programs. While clinical benefits such as improved chronic disease management or reduced hospitalizations may be evident, quantifying the economic and social returns of digital literacy training or broadband expansion can be difficult, especially in the short term. Policymakers and funders must adopt broader definitions of value—ones that incorporate empowerment, health autonomy, and community resilience as key outcomes worthy of sustained investment (Ayo-Farai, et al., 2024, Ezeamii, et al., 2024, Oboh, et al., 2024, Oshodi, et al., 2024).

In addition to funding concerns, privacy and trust present formidable barriers to engagement, particularly among historically marginalized populations. Underrepresented communities—especially Black, Latino, Indigenous, and immigrant populations—have experienced a long history of systemic discrimination, medical exploitation, and surveillance. These experiences contribute to deep-seated skepticism toward digital health tools, particularly those that collect sensitive health data. For example, undocumented immigrants may fear that their information could be shared with immigration enforcement agencies, while others may worry that digital platforms could be used to deny services or discriminate against them based on race, income, or health status (Adhikari, et al., 2024, Ezeamii, et al., 2024, Ogundairo, et al., 2024).

The challenge is compounded by a general lack of transparency in how health data is collected, stored, and shared. Many patients are unaware of what information is being tracked through patient portals, telehealth apps, or wearable devices. Even when privacy policies are provided, they are often written in technical or legal language that is difficult for laypersons to understand. This confusion undermines informed consent and reinforces mistrust, especially when coupled with reports of data breaches or unauthorized data sharing (Madu & Nwankwo, 2018, Nasuti, et al., 2008, Nwankwo, et al., 2011, Tayebati, et al., 2013).

To address these concerns, developers and healthcare organizations must prioritize transparency and user agency in all aspects of digital health design. Privacy notices should be written in plain language and offered in multiple languages. Platforms should allow users to opt in or out of specific types of data collection, and these preferences should be honored and easily revisited. Community health workers and digital navigators can also play a crucial role in building trust by explaining privacy protections, demonstrating how to use digital tools safely, and serving as intermediaries between patients and institutions (Babarinde, et al., 2023, Eyeghre, et al., 2023, Nwaonumah, et al., 2023).

Moreover, policies must be updated to reflect the unique risks and needs of underserved populations. Federal regulations such as HIPAA (Health Insurance Portability and Accountability Act) provide a baseline for data protection, but they do not cover all health-related apps or devices, especially those outside of traditional healthcare systems. Additional safeguards are needed to ensure that third-party vendors, technology companies, and insurers do not misuse health data. Policymakers must also invest in public education campaigns that inform people of their digital rights and how to protect their health information (Adelodun, et al., 2018, Chianumba, et al., 2021, Tayebati, et al., 2012, Tomassoni, et al., 2013).

Interoperability and technological adaptability present further complications in implementing a unified framework for digital health equity. Interoperability refers to the ability of different digital health systems, applications, and devices to exchange, interpret, and use data seamlessly. For underrepresented communities, this issue is critical because fragmented or siloed systems can lead to inconsistent care, data duplication, or complete exclusion from digital health services. For example, a patient who accesses telehealth through a community health center may not have their records integrated with hospital systems, leading to redundant tests or overlooked medical history (Akerele, et al., 2024, Fagbenro, et al., 2024, Kelvin-Agwu, et al., 2024).

The U.S. healthcare system is notoriously fragmented, with varying levels of digital maturity across providers, payers, and regions. While some systems boast integrated electronic health records and robust telehealth platforms, others struggle with outdated software, limited bandwidth, or incompatible databases. These disparities affect the ability to deliver cohesive care and make it difficult for public health agencies to monitor digital health equity outcomes across populations (Ajibola, et al., 2024, Folorunso, et al., 2024, Majebi, Adelodun & Anyanwu, 2024). Furthermore, non-standardized platforms may exclude community clinics or safety-net providers from accessing funding or participating in broader digital health networks. To address interoperability, national standards and regulatory mandates must be enforced, requiring all vendors and healthcare systems to use common protocols and data-sharing frameworks. The 21st Century Cures Act and related rules from the Office of the National Coordinator for Health Information Technology (ONC) have made strides in promoting open APIs and data portability, but full implementation remains uneven. Equity considerations must be central in interoperability planning to ensure that smaller, underfunded clinics serving marginalized populations are not left behind in the digital transformation (Madu & Nwankwo, 2018, Nwankwo, et al., 2012, Nwankwo, Tomassoni & Tayebati, 2012).

Technological adaptability is also a concern as digital health tools continue to evolve rapidly. New platforms, devices, and features emerge constantly, often requiring frequent updates, new training, or additional hardware. For populations with limited resources or digital literacy, keeping up with technological changes is a significant burden. Systems must therefore be designed with flexibility, backward compatibility, and low-bandwidth functionality in mind. Moreover, providers and patients alike must have ongoing access to support—not just during initial implementation but as technologies advance (Noah, et al., 2025, Opia & Matthew, 2025, Udegbe, et al., 2023, Usuemerai, et al., 2024).

The lack of culturally competent tech development also hinders adaptability. Many platforms are developed without user input from underrepresented communities, resulting in designs that are not intuitive, relevant, or accessible. This problem can be mitigated through participatory design processes, in which community members are directly involved in prototyping, testing, and evaluating tools intended for their use (Olowe, et al., 2024, Olulaja, Afolabi & Ajayi, 2024, Shittu, et al., 2024). By embedding adaptability into both technology and process, digital health interventions can remain relevant and useful over time, particularly for communities that have historically been excluded from the digital mainstream.

In conclusion, while the conceptual framework for addressing digital health literacy and access gaps provides a robust foundation for equitable digital healthcare, its implementation is fraught with complex challenges. Sustainability and funding remain precarious, requiring coordinated public investment and long-term planning. Privacy and trust concerns must be addressed through transparent communication, regulatory protections, and culturally sensitive engagement (Okon, Zouo & Sobowale, 2024, Olamijuwon, et al., 2024, Olorunsogo, et al., 2024). Interoperability and adaptability are technical and organizational hurdles that demand both standardization and inclusive innovation. Addressing these challenges will require not only technical solutions but also a firm commitment to equity, justice, and the lived experiences of the communities most affected by the digital divide. Only by doing so can we ensure that digital transformation enhances, rather than hinders, access to health for all.

2.7. Policy Recommendations

Addressing digital health literacy and access gaps in underrepresented communities in the United States requires a coordinated and sustained policy response that integrates public health, technology, education, and social equity. The conceptual framework for achieving digital inclusion in healthcare identifies the core pillars needed to advance equitable access, but for the framework to succeed, it must be operationalized through concrete policy measures across all levels of governance and within the private sector (Adigun, et al., 2024, Folorunso, et al., 2024, Kelvin-Agwu, et al., 2024). The development and implementation of inclusive digital health ecosystems must be a shared responsibility, and policy recommendations targeting government agencies, healthcare providers, and technology developers are essential to realizing this vision.

Government agencies at the federal, state, and local levels play a critical role in shaping the digital health landscape and have the authority to create enabling environments for equitable access. One of the most urgent policy actions is to formally embed digital health literacy and access objectives into national and state health agendas. This includes incorporating digital inclusion metrics into Healthy People 2030 objectives, as well as Medicaid and Medicare modernization plans (Uwumiro, et al., 2024, Wada, et al., 2025, Zouo & Olamijuwon, 2024). Federal agencies such as the Department of Health and Human Services (HHS), the Federal Communications Commission (FCC), and the Centers for Medicare and Medicaid Services (CMS) should align policies to ensure broadband access, affordable devices, and culturally appropriate digital health services are treated as foundational components of public health infrastructure.

Policy mandates must also ensure that digital health access is treated as a social determinant of health. This redefinition would enable the integration of digital inclusion assessments into community health needs assessments, electronic health records, and patient intake processes. It would also allow healthcare systems and insurers to use public funds to provide digital support services—such as training, navigation, and devices—to patients most in need (Balogun, et al., 2023, Ezeamii, et al., 2023, Katas, et al., 2023, Usuemerai, et al., 2024). Government funding mechanisms, including Section 1115 waivers, HRSA grants, and public health block grants, should be explicitly directed to support initiatives that close digital health

literacy gaps among Medicaid populations, residents of public housing, rural communities, and people with disabilities.

Healthcare providers and health systems must be incentivized through policy to proactively address digital inclusion. Reimbursement structures should recognize and support the role of healthcare workers, including community health workers and digital navigators, in educating patients about digital tools and supporting their use. Current reimbursement models for telehealth services often fail to account for the time and resources required to assist patients with low digital literacy or limited English proficiency in navigating digital platforms (Adelodun & Anyanwu, 2024, Ibikunle, et al., 2024, Ogugua, et al., 2024). Updating these models to cover digital literacy screening, technical support, and interpreter services for digital care encounters would reduce inequities and promote more meaningful digital engagement. In parallel, accreditation and licensing bodies should update provider education and continuing professional development requirements to include digital health equity competencies. Providers should be trained to assess digital literacy as part of routine care, recognize the signs of digital exclusion, and deliver information in formats that meet patients' literacy levels. Cultural humility, trauma-informed communication, and technology-demystifying practices should be incorporated into clinical training programs, ensuring providers can deliver patientcentered care in an increasingly digital environment (Ayo-Farai, et al., 2024, Ibikunle, et al., 2024, Oddie-Okeke, et al., 2024).

Technology developers, including electronic health record vendors, app creators, and telehealth platform providers, must be held accountable through policy and incentives to create inclusive, accessible, and user-friendly tools. Policymakers should establish clear design standards and certification requirements for digital health platforms that are funded through public dollars or used in federally qualified health centers and safety-net institutions (Anyanwu, et al., 2024, Idoko, et al., 2024, Kelvin-Agwu, et al., 2024). These standards must include multilingual support, accessibility for users with disabilities, offline functionality, low-bandwidth optimization, and compliance with plain language guidelines. The Office of the National Coordinator for Health Information Technology (ONC) could lead this effort by expanding its certification program to prioritize equity-driven benchmarks in addition to technical interoperability (Bello, et al., 2024, Igwama, et al., 2024, Katas, et al., 2024, Okobi, et al., 2024).

Incentivizing inclusive design and innovation will require both carrots and sticks. On the incentive side, the federal government and philanthropic foundations should increase investment in equity-centered technology development through challenge grants, innovation funds, and public-private partnerships. Programs such as the Small Business Innovation Research (SBIR) and Small Business Technology Transfer (STTR) should dedicate funding tracks for technologies specifically designed to close digital health disparities (Elujide, et al., 2021, Khosrow Tayebati, Ejike Nwankwo & Amenta, 2013), Tomassoni, et al., 2013). Tax credits and preferential contracting can be offered to companies that demonstrate a commitment to universal design principles and co-creation with underrepresented communities. Conversely, digital health tools that fail to meet accessibility standards or demonstrate disproportionate negative impacts on vulnerable populations should be ineligible for use in publicly funded healthcare settings (Gabrielli, et al., 2010, Khosrow Tayebati, et al., 2013, Nwankwo, et al., 2011).

Policies must also promote transparency and accountability. Federal and state health agencies should require health systems and vendors to collect and publicly report data on digital access, usage, and satisfaction disaggregated by race, ethnicity, income, disability status, and geography. This data will help identify gaps, drive targeted interventions, and ensure that digital transformation efforts are benefiting all communities equitably. Patient-reported outcome measures should be expanded to include digital engagement and confidence, and these metrics

should inform both quality improvement and value-based payment models (Abass, et al., 2024, Igwama, et al., 2024, Kelvin-Agwu, et al., 2024, Olowe, et al., 2024).

Integration into national and state agendas also requires interagency and cross-sector collaboration. Digital health equity must be coordinated with efforts in education, housing, workforce development, and broadband infrastructure. For instance, public housing authorities can partner with health systems to provide residents with connectivity and telehealth kiosks. Workforce development agencies can fund training programs for digital navigators and community health workers. School districts and libraries can serve as access points for digital literacy training and health education (Attah, et al., 2022, Chianumba, et al., 2022, Opia, Matthew & Matthew, 2022). A digital health equity task force, established at the federal level and mirrored by state-level counterparts, could help coordinate these efforts, monitor implementation, and elevate best practices.

Another critical policy recommendation is to develop and implement a national digital health equity strategy. This strategy should be shaped by community voices and guided by an equity framework that centers on historically marginalized populations. The strategy must articulate clear goals, timelines, funding commitments, and evaluation mechanisms. Importantly, it must include mechanisms for meaningful community participation at all levels of decision-making, from program design to evaluation. The strategy should be accompanied by a dedicated office or agency charged with implementation oversight, community engagement, and coordination across health and technology sectors (Adelodun & Anyanwu, 2024, Igwama, et al., 2024, Majebi, Adelodun & Anyanwu, 2024).

Finally, state and local governments must be empowered and resourced to adapt and implement these recommendations in a manner that is responsive to their unique demographic, geographic, and cultural contexts. States should be encouraged to develop their own digital health equity plans, aligned with national priorities but tailored to local realities. Local health departments, in particular, are well-positioned to implement community-based digital inclusion initiatives, assess local needs, and coordinate multisector partnerships (Ayo-Farai, et al., 2023, Ezeamii, et al., 2023, Katas, et al., 2023).

In conclusion, the successful implementation of the conceptual framework for addressing digital health literacy and access gaps depends on an ambitious and coordinated policy agenda. Government agencies must institutionalize digital health equity as a public health imperative, healthcare providers must be equipped and incentivized to support digital inclusion, and technology developers must be held to standards that prioritize accessibility and cultural relevance (Afolabi, Ajayi & Olulaja, 2024, Igwama, et al., 2024, Ohalete, et al., 2024). Integration into national and state health agendas ensures that digital equity is not an afterthought, but a fundamental component of 21st-century healthcare delivery. By aligning incentives with inclusive design and community empowerment, the digital divide can be closed—not only in terms of access, but in terms of trust, usability, and health outcomes for all (Al Hasan, Matthew & Toriola, 2024, Igwama, et al., 2024, Okhawere, et al., 2024).

2.8. Conclusion

The conceptual framework for addressing digital health literacy and access gaps in underrepresented U.S. communities presents a comprehensive, equity-centered approach to one of the most pressing challenges in modern healthcare. By articulating four interconnected pillars—digital access infrastructure, digital health literacy development, community-based engagement, and policy and systems integration—the framework provides a roadmap for transforming how healthcare is delivered and accessed in the digital age. It underscores the importance of not only providing internet and devices but also ensuring that individuals have the skills, support, and culturally relevant tools to engage meaningfully with digital health systems. It recognizes the indispensable role of communities, healthcare providers,

policymakers, and technology developers in creating inclusive solutions that reflect the lived experiences of those most often excluded.

The urgency of closing digital health equity gaps cannot be overstated. As healthcare becomes increasingly digitized, individuals without access to or fluency in digital tools are at risk of being further marginalized—exacerbating long-standing disparities in care, outcomes, and trust. The COVID-19 pandemic highlighted both the potential of digital health innovations and the cost of digital exclusion, particularly for low-income families, rural populations, older adults, people with disabilities, and communities of color. If left unaddressed, these gaps will deepen structural inequities and undermine national efforts toward health equity, population health improvement, and universal access to care. The moment demands decisive action, long-term investment, and a commitment to equity that extends beyond rhetoric.

This is a call to action for multi-sector collaboration. Government agencies must lead with bold policies that prioritize digital equity as a core public health goal. Healthcare systems must integrate digital inclusion into their models of care, treating it as essential to quality and patient-centered service. Technology companies must design with and for the people most at risk of exclusion, embedding accessibility and cultural relevance into every layer of innovation. And communities must be empowered—not merely engaged—to shape the solutions that affect their health and lives. Only through sustained, coordinated, and inclusive efforts can we ensure that digital health advancements reach and uplift every community, leaving no one behind in the digital transformation of care.

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