



The Equity of Virtual Ward Models of Care Across Sociodemographic Groups in the United Kingdom – A Systematic Review

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The Equity of Virtual Ward Models of Care Across Sociodemographic Groups in the United Kingdom – A Systematic Review

Journal of Integrated Care

Abstract

Purpose – Virtual Wards (VWs), or “Hospital-at-Home (HAH)” services, deliver hospital-level care in patients’ homes by integrating remote monitoring and treatment, facilitating early discharge and reducing inpatient admissions. However, the equity of these care models in the UK remains unclear. This paper aims to synthesise existing evidence on VW’s effectiveness, suitability, and acceptability across diverse sociodemographic groups using the PROGRESS-Plus equity framework.

Design/methodology/approach – Systematic literature search of PubMed, CINAHL-PLUS (EbscoHost), and MedRxiv databases was conducted, alongside grey literature from government and NHS sources. Eligible studies were appraised using the Mixed Methods Appraisal Tool, and findings were synthesized narratively.

Findings – A total of 8,011 records were identified, with 7 studies and 3 government reports included in the final analysis. Using the PROGRESS-Plus framework, the review found that race/ethnicity/culture/language, socioeconomic status and occupation, social capital, age, and health status and comorbidities shaped the effectiveness, suitability, and acceptability of VWs. Barriers included cultural and communication mismatches, inadequate housing, lack of caregiving support, digital exclusion among older adults, and reduced benefit or engagement for those with complex health needs. Targeted interventions are needed for equitable VW implementation across the UK.

Originality – This is the first UK-based systematic review to examine the equity implications of VW implementation using the PROGRESS-Plus framework. While VWs offer system-wide benefits, there is a need for inclusive design and targeted interventions to ensure equitable access and outcomes across all population groups.

Keywords Virtual Wards, Hospital-at-Home, Health Inequity, UK

Paper type Literature review

Background

Virtual wards (VWs) provide hospital-level care in patients' homes through integrated technology and multidisciplinary medical services (NHS England & NHS Improvement, 2022). As part of the broader "Hospital-at-Home (HAH)" spectrum, VWs offer acute care, monitoring, and treatment at home to prevent hospital admissions (the step-up model) or support early discharge (the step-down model) (Chappell *et al.*, 2024; NHS England & NHS Improvement, 2022).

VWs gained prominence during COVID-19 to ease hospital capacity (Vindrola-Padros *et al.*, 2021). Proponents suggest multiple potential benefits, including enhanced patient autonomy, personalised treatment, reduced hospital stress, and convenience for families (Chua *et al.*, 2021; NHS England, 2022). System-wide, VWs may alleviate hospital bed pressure, optimise resources, and reduce costs by potentially shortening hospital stays and leveraging home infrastructure (NHS England, 2022). Emerging evidence shows VWs may match or outperform traditional care, including possible reduced readmissions, shorter stays, and improved survival for chronic conditions like heart failure (Chauhan *et al.*, 2022; Shi *et al.*, 2024). They may also improve quality of life and reduce emergency department (ED) visits, particularly among frail patients (Low *et al.*, 2017; Westby *et al.*, 2024).

Therefore, the NHS has prioritised VW expansion, targeting clinical pathways such as frailty, acute respiratory infections, and heart failure, aiming for consistent utilisation rates above 80%, including children and young people (NHS England, 2024). Achieving this requires advanced remote monitoring, rapid access to diagnostics, and cross-service collaboration to enhance patient flow, as part of a broader strategy to modernise healthcare delivery and improve patient outcomes (NHS England, 2024).

However, there are challenges to further expanding use of VWs such as digital literacy gaps, housing suitability, and risks for isolated individuals (Chua *et al.*, 2021; Norman *et al.*, 2023). While research highlights clinical efficacy, evidence on effectiveness, suitability, and

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3 acceptability across populations remains limited (Chappell *et al.*, 2024). Digital inclusion is a key
4 barrier, with language, technology access, and IT skills disproportionately impacting vulnerable
5 groups. For example, a UK study found that 31.5% of patients referred to VWs could not use the
6 required app (Fox *et al.*, 2022). Similar global concerns include cultural, ethnic, and
7 socioeconomic factors further hindering virtual care access (Mistry *et al.*, 2022).
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14 Given well-documented health inequalities in UK healthcare driven by factors like deprivation,
15 gender, age, and ethnicity (Kulkarni *et al.*, 2022; Saunders *et al.*, 2021), it would be informative
16 to understand whether VWs are equitable, and whether they address or exacerbate existing
17 disparities. For example, while VWs might enhance healthcare access for underserved areas,
18 they risk excluding groups with limited digital access or poor living conditions, potentially
19 further disadvantaging vulnerable populations.
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27 This review synthesises existing evidence on the impact of VWs across diverse
28 sociodemographic groups, focusing on effectiveness, suitability, and acceptability to
29 understand their role in addressing or exacerbating health inequalities, and to offer insights for
30 healthcare providers and policymakers.
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36 **Box 1. Definitions**

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39 • **Effectiveness:** Examines whether VWs improve intended health outcomes and reduce
40 harm across different sociodemographic groups.
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42 • **Suitability:** Evaluates how well VWs align with the needs, contexts, and resources of
43 diverse populations of the UK.
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45 • **Acceptability:** Examines how VWs are perceived by patients, families, and healthcare
46 providers, including satisfaction, comfort, and willingness to engage.
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Methods

This systematic review adhered to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) 2020 guidelines and was carried out between July and November 2024.

Eligibility Criteria

This review followed the PICOS (Population, Intervention, Comparison, Outcome, Study design) framework (Amir-Behghadami and Janati, 2020). Publications were included if they focused on UK adults (19+); described VW or HAH hospital-level care at home; and involve a comparison to conventional inpatient care via step-up (prevent admission) or step-down (early discharge) models. Eligible outcomes included indicators of effectiveness (e.g., access to care, recovery, readmission, mortality), suitability (e.g., alignment with patient needs/resources), and acceptability (e.g., patient/family satisfaction, engagement). Acceptable study designs included empirical literature, descriptive or evaluative government reports, or theses employing randomized controlled trials (RCTs), cohort, case-control, cross-sectional, observational, evaluation, and mix methods. Only English-language publications from 2000 onward were considered.

Publications were excluded if they were abstracts, editorials, commentaries, letters to the editor, or opinion pieces; published before 2000; focused on populations outside the UK; or were non-human or non-English studies.

Information Sources

The review searched PubMed, CINAHL-PLUS (EbscoHost), and MedRxiv, and also included resources from NHS England's VW [evidence catalogue](#) (accessible through the FutureNHS learning platform), which contains both academic papers and grey literature like government and policy reports. Reference lists from included studies were manually reviewed for additional relevant literature.

Search Strategy

The search strategy focused on two main concepts: VW services and the UK, with specific search terms detailed in Table I.

[Insert Table I here]

Selection Process

Results were imported into EndNote (<https://endnote.com/>) for deduplication. Relevant papers were first identified through title and abstract screening by the first author (GP), then full texts of the retrieved articles were further assessed for eligibility by two authors independently (GP and DS). The first author thoroughly screened the NHS catalogue of VW resources, including articles, government reports, and NHS publications. Potentially relevant documents were then reviewed by the second author, with any uncertainties resolved jointly. The selection process is summarised using the PRISMA 2020 diagram.

Data Extraction

Key study details, including aims, outcomes, samples, and findings, were extracted into a summary table. Health inequality data were extracted using the PROGRESS-Plus framework to identify relevant factors, such as place of residence, race/ethnicity/culture/language, occupation, gender/sex, religion, education, socioeconomic status, and social capital (O'Neill *et al.*, 2013).

Quality Assessment of Studies

Study quality was evaluated using the Mixed Methods Appraisal Tool (MMAT) across five criteria: research relevance, design appropriateness, data reliability, analysis robustness, and findings validity, rated as “yes,” “no,” or “can’t tell” (Hong *et al.*, 2018). Following MMAT guidelines, overall quality scores were not calculated; instead, detailed ratings were provided to capture methodological strengths and weaknesses. Sensitivity analysis was omitted due to study design variability, and no studies were excluded to ensure comprehensive evidence.

Data Synthesis

Findings were synthesized using a narrative approach, as meta-analysis was unsuitable due to variability in study designs, outcome measures, and reporting formats, along with the inclusion of qualitative data that could not be quantitatively aggregated. Potential equity implications were inferred from reported differences across PROGRESS-Plus categories. These interpretations were indirect and hypothesis-generating rather than conclusive, given the absence of formal subgroup analyses and limited contextual information (e.g., population demographics) in most included studies.

Results

The literature search (July-November 2024) yielded 8,011 records, with 22 additional records identified through NHS England's evidence catalogue. After deduplication, 8,024 records were screened by title and abstract, with 73 selected for full-text review. Four were unavailable, and 59 were excluded based on study design (n = 4), intervention (n = 11), outcomes (n = 36), or geographical location (n = 8). Ultimately, 10 records (7 studies and 3 government reports) met the inclusion criteria. See Figure 1 for the PRISMA 2020 flow diagram.

[Insert Figure 1 here]

Study Characteristics

Table II summarises the characteristics of the included studies and government reports, which span from 2000 to 2024: mixed methods (n = 5), RCTs (n = 2), quantitative observational (n = 2), and qualitative (n = 1). Only Crellin *et al.* (2024) directly examined health disparities in virtual care, while Tsiachristas *et al.* (2019) addressed them indirectly through demographic comparisons. Other studies did not explicitly investigate outcome variations, but relevant findings were inferred indirectly.

[Insert Table II here]

How sociodemographic factors influence the effectiveness, suitability, and acceptability of VWs in the UK?

Thematic analysis using the PROGRESS-Plus framework identified how equity-related factors may moderate VW effectiveness, suitability, and acceptability.

Race, ethnicity, culture, and language

Ethnic and cultural background shaped VW acceptability, with individuals from minority ethnic groups facing language barriers, cultural mismatches, and limited service understanding, reducing trust and participation (Crellin et al., 2024). Underrepresentation of these groups in VW cohorts suggests systemic issues in access and inclusivity (The Wessex Academic Health Science Network, 2022).

Evidence on clinical effectiveness by ethnicity was limited; Nunan et al. (2020) identified potential differences in readmission rates, although statistical significance was unclear. Cultural considerations also influenced suitability, as services were inconsistently adapted to diverse needs, limiting their relevance and accessibility.

Socioeconomic status and occupation

Socioeconomic factors influenced participation through structural barriers and lifestyle constraints. Housing quality and stability were key to suitability, as low-income patients often lived in environments unsuitable for remote care (The Health Foundation, 2023). Overcrowded or insecure housing limited privacy, space, and technology needed for remote monitoring.

Effectiveness was impacted by environmental and financial barriers like overcrowded housing and unstable employment, limiting consistent patient engagement. Crellin et al. (2024) found financial hardship reduced sustained participation. Acceptability was similarly shaped by socioeconomic conditions, with some viewing VWs as impractical without additional support.

Social capital

Living arrangements and caregiving influenced patient engagement and suitability. In terms of effectiveness, Lewis et al. (2017) found no significant differences in unplanned hospital admissions or ED visits between those living alone and with others. Cotton et al. (2000) reported similar findings on readmission rates.

Caregiver presence was key to suitability. Crellin et al. (2024) and Health Foundation (2023) found patients with caregivers were better able to use monitoring technology and remain engaged in their care. In contrast, those without support, particularly individuals living alone, faced more difficulties with technologies. Acceptability was lower among some elderly living alone due to fear or lack of confidence managing alone (Health Innovation Network, 2021).

Age

Age appeared to shape digital engagement and perceptions of VWs. Contrary to assumptions, acceptability increased with age, with older individuals more willing to be treated remotely than younger patients, particularly when adequate support was available (Health Foundation, 2023).

However, suitability challenges related to digital literacy were common among older adults, who often struggled with devices and information. Crellin et al. (2024) and Health Innovation Network (2021) noted this sometimes led to refusal or disengagement.

Health status and comorbidities

Patients with complex needs, chronic conditions, or disabilities faced barriers to VW engagement. Tsiachristas et al. (2019) found higher mortality risks for those with multiple comorbidities in home care; Crellin et al. (2024) noted severe conditions required more support and lowered engagement.

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3 Suitability and acceptability concerns included struggles to access equivalent remote care
4 (Health Foundation, 2023) and emotional barriers. Cotton et al. (2000) found anxiety or
5 depression caused discomfort with remote care, with a preference for in-person visits.
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7 Isolation, fear of being alone, and reduced confidence commonly led to declining participation
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9 (Health Innovation Network, 2021; Dismore et al., 2019).
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13 14 Discussion

15 16 Summary of Findings

17 This systematic review evaluated how VWs impact health inequalities in the UK by examining
18 variations in effectiveness, suitability, and acceptability across sociodemographic groups. The
19 findings aim to inform more equitable implementation of virtual care models.
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25 Across the 10 included studies, only two directly examined health disparities in VW services,
26 while the rest provided indirect or subgroup-level insights. This underscores a significant gap in
27 the UK literature. While evidence on clinical efficacy is growing (Jeppesen *et al.*, 2012; Lee *et al.*,
28 2022), few studies examine the intersection of VWs with key equity dimensions. This limits our
29 ability to draw robust conclusions about whether VWs mitigate or exacerbate health
30 inequalities. More focused research is needed to understand how VWs influence diverse
31 populations and ensure they promote equity in healthcare delivery.
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40 Findings on effectiveness suggest patients with multiple comorbidities, limited social support,
41 or unstable housing may benefit less from VW services. Suitability challenges included
42 inadequate living environments, lack of caregiver support, and cultural or language mismatches
43 that hindered patient engagement. Acceptability was shaped by trust, confidence using digital
44 tools, and perceived relevance, particularly among individuals from ethnic minority groups,
45 older adults, and those facing digital exclusion.
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Comparisons with Previous Literature

This review echoes findings from an Australian systematic review, which identified cultural and ethnic inequities in virtual care (Mistry *et al.*, 2022). Similar to our findings, this review found older adults, people from lower socioeconomic backgrounds and minority ethnic groups were underrepresented in virtual services due to digital, structural, and cultural barriers. Both reviews note that older adults and low-income individuals face more barriers from limited digital access and confidence, while younger individuals were generally more engaged with virtual services.

Budhwani *et al.* (2022) highlight barriers like limited internet access, high costs, systemic racism, and mistrust in digital tools, disproportionately impacting socioeconomically disadvantaged and racialized groups. Our review supports this, showing these challenges often reduce VW suitability, which frequently lacks culturally sensitive design and tailored engagement for individuals from minority communities. Limited outreach and engagement strategies likely lowered engagement, though this was not systematically studied. These findings underscore the need for culturally responsive design and infrastructure investment to ensure equitable virtual care implementation. Recent work by Yahya *et al.* (2025) similarly cautions that without intentional equity planning, VWs may inadvertently exacerbate existing health disparities.

Methodological Considerations

A key strength of this review is its comprehensive inclusion of diverse studies and government reports, capturing broad perspectives on VW implementation, including real-world experiences from patients, the public, and NHS staff. Additionally, screening was conducted by two authors who reached consensus, and a study quality assessment was performed to ensure robust analysis. One limitation is that due to time constraints, this review was not prospectively registered on PROSPERO. Title and abstract screening was conducted by one author, which may introduce selection bias, though this was mitigated by dual screening at the full-text stage.

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3 Other limitations include the small number of studies directly addressing health outcome
4 variations across sociodemographic groups and reliance on indirect evidence.
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9 To support an equity-focused analysis, the PROGRESS-Plus framework was applied to explore
10 potential implications from observed differences across sociodemographic groups. As most
11 studies lacked formal equity analyses, these interpretations remain indirect and exploratory,
12 highlighting areas for future research rather than definitive conclusions on health inequalities.
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17 18 Conclusion

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20 This review demonstrates that while VWs have the potential to improve care delivery in the UK,
21 their effectiveness, suitability, and acceptability vary across sociodemographic groups. Barriers
22 like poor housing, limited digital access, cultural mismatches, and lack of caregiver support
23 disproportionately affect individuals from lower socioeconomic backgrounds, ethnic minorities,
24 and those with complex needs. Although some older adults are receptive, engagement remains
25 uneven.
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33 Equitable VW implementation in the UK requires culturally appropriate communication,
34 accessible technology, and targeted outreach. Monitoring and evaluation should incorporate
35 equity-focused indicators from PROGRESS-Plus domains. Future UK-based research should
36 prioritise disaggregated data and strategies to reduce structural barriers for underserved
37 groups.
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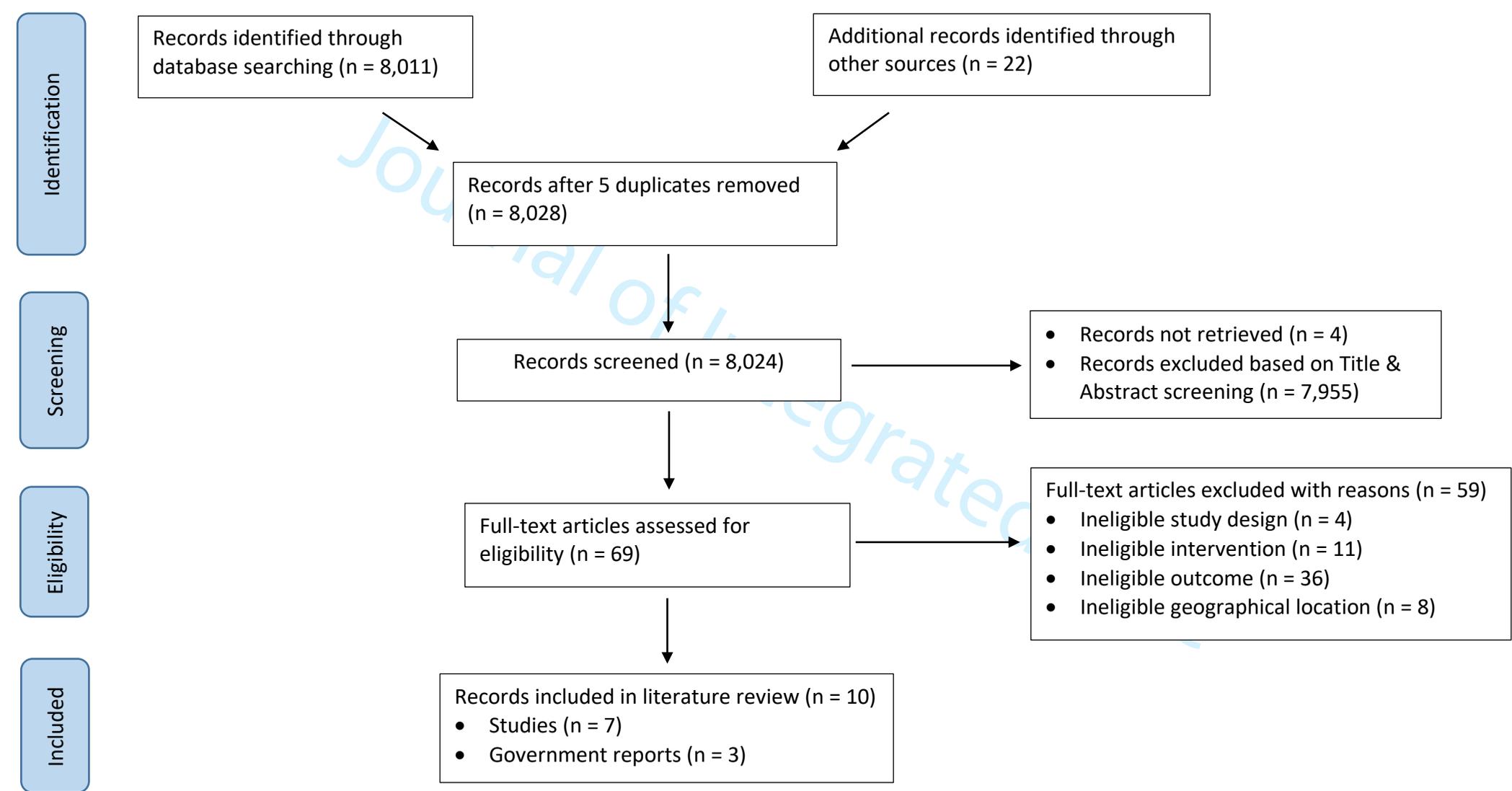
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Table I. Search Terms

Term between search terms	Concept 1: Virtual ward services	AND	Concept 2: United Kingdom
	Virtual ward		United Kingdom
OR	Virtual care model		UK
OR	Virtual hospital		England
OR	Virtual inpatient		Scotland
OR	Home hospitalization		Wales
OR	Hospital@home		Northern Ireland
OR	Home care		Great Britain
OR	Home care services		NHS
OR	Step down care		
OR	Step up care		
OR	Remote monitoring		
OR	Telehealth		
OR	Telemedicine		
OR	Telemonitoring		
OR	Telemanagement		
OR	Remote patient management		

Figure 1. PRISMA 2020 Flow Diagram



Source(s): Author's own creation/work

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Table II. Summary of methods and key results of included studies

Author & Year	Setting	Aim	Study Design	VW Model of Care	Sample Size	Outcomes Reported in the Study	Main Findings Relevant to Inequalities	PROGRESS-Plus Categories	MMAT Quality Assessment
Cotton <i>et al.</i> , 2000	Glasgow, Scotland	To evaluate early discharge with home respiratory nurse support versus standard hospital care for COPD exacerbations.	RCT	Step-down	81	<ul style="list-style-type: none"> Additional hospital days within 60 days of initial admission. Mortality during the 60-day period. Readmission rate. 	Living alone was not a risk factor for readmission, but social isolation, increased anxiety, and depression may be.	<ul style="list-style-type: none"> Health status and comorbidities (anxiety, depression) Social capital (living alone, social isolation, caregiver access) 	Good
Dar <i>et al.</i> , 2009	London, England	To assess home telemonitoring on heart failure patients discharged from three hospitals.	RCT	Step-down	182	<ul style="list-style-type: none"> Clinic visits. Days alive and out of hospital. Hospitalizations due to heart failure. Quality of life. 	Telemonitoring was acceptable and usable in this elderly, multi-ethnic sample, with good satisfaction reported, though outcome differences by sociodemographic subgroup were not examined.	<ul style="list-style-type: none"> Age (elderly participants) Race, ethnicity, culture, and language (multi-ethnic cohort) Social capital (living alone, caregiver support) 	Good
Lewis <i>et al.</i> , 2017	Dublin, Ireland	To assess whether a community VW reduces unplanned hospital visits and examine predictors of long-term care needs.	Quantitative Observational	Step-up	54	<ul style="list-style-type: none"> ED presentations. Number of bed days and hospital admissions. Hospital admission risk/activity, comparing individuals living alone versus those with a caregiver. 	Before community VW admission, those living alone had fewer hospital and ED visits. Post-VW, no differences were found in unplanned admissions or ED presentations between those living alone and others.	<ul style="list-style-type: none"> Social capital (family support, living alone) 	Good

1 2 3 4 5 6 7	Dismore <i>et al.</i> , 2019	North East England	To identify facilitators and barriers to implementing HAH.	Qualitative	Step-down	89	<ul style="list-style-type: none"> Barriers and facilitators to implementing HAH. 	Fear of being alone, especially among patients living alone or without informal support, was a common barrier to participation.	<ul style="list-style-type: none"> Social capital (fear of being alone, resistance linked to lack of informal support) 	Good
8 9 10 11 12 13 14 15 16 17 18	Tsiachristas <i>et al.</i> , 2019	Across Scotland	To compare populations admitted to HAH services versus hospitals and evaluate their impact on costs and mortality.	Retrospective observational cohort	Step-up	22,610	<ul style="list-style-type: none"> Healthcare costs Mortality rates 	Patients in HAH were older, more socioeconomically disadvantaged, and had frequent hospitalizations. The HAH cohort also included more women and individuals with multiple chronic conditions than those admitted to hospital. HAH admission was linked to higher mortality risk during follow-up.	<ul style="list-style-type: none"> Age Health status and comorbidities (multiple chronic conditions) Socioeconomic status and occupation (lower SES) 	Good
19 20 21 22 23 24 25 26 27 28 29	Nunan <i>et al.</i> , 2020	Reading, England	To evaluate the feasibility of a triage and follow-up pathway for mild COVID-19 pneumonia.	Mixed Methods	Step-down	273	<ul style="list-style-type: none"> Patient satisfaction Perceived safety and reassurance from daily contact and home monitoring National Early Warning Score (NEWS-2). Oxygen saturation at presentation. 	No significant differences in re-attendance rates, but possible associations with ethnicity, baseline oxygen saturation levels, post-walk saturation changes, and total NEWS scores.	<ul style="list-style-type: none"> Race, ethnicity, culture, and language (ethnicity-related differences noted) 	Good
30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47	Health Innovation Network, 2021	London, England	To examine the impact of a VW in Croydon, including demographics, effectiveness, acceptability, experiences, healthcare utilization, cost savings, and outcomes.	Mixed Methods Evaluation	Step-up, step-down	250	<ul style="list-style-type: none"> General improvement in clinical outcomes Economic impact and cost analysis Health service utilization Health status Patient and staff feedback 	VW patients, primarily older adults, found remote monitoring kits user-friendly, though 4% declined—often due to low confidence, especially if living alone. Satisfaction and quality of life were high, but a minimum digital literacy was needed.	<ul style="list-style-type: none"> Age Health status and comorbidities (some support needs noted in older adults) Race, ethnicity, culture, and language (implied through user preference and design suitability) 	Good

						<ul style="list-style-type: none"> • Representativeness of VW users 		<ul style="list-style-type: none"> • Social capital (living alone, caregiver access, confidence managing care alone) 	
Wessex Academic Health Science Network, 2022	Hampshire, England	To evaluate the HHFT VW program's demographics, referrals, outcomes, bed usage, workforce redesign, technology, and patient experiences, focusing on step-up and step-down processes.	Mixed Methods Evaluation	Step-up, step-down	1,223	<ul style="list-style-type: none"> • Representativeness of VW users • Patient outcomes (e.g., health conditions, discharge outcomes) • Patient experience • Healthcare utilization • Cost savings • Service expansion 	Most patients in the HHFT VW program were White British (92%), reporting high satisfaction and valuing regular consultations, staff expertise, and telephone communication.	<ul style="list-style-type: none"> • Race, ethnicity, culture, and language (underrepresentation of ethnic minorities) 	Good
The Health Foundation, 2023	Across UK	To explore public and NHS staff perceptions of VWs and identify key factors for effective implementation.	Mixed Methods	Step-up, step-down	8,351	<ul style="list-style-type: none"> • Demographic variations in support • Digital exclusion and access • Key factors for successful VW implementation • Perceived barriers to VWs • Public and staff support for VWs 	Disabled individuals and those with caregivers showed higher support, while people from lower socioeconomic backgrounds were less enthusiastic, citing unsuitable homes. Confidence in VW technologies grew with familiarity.	<ul style="list-style-type: none"> • Health status and comorbidities (disability) • Race, ethnicity, culture, and language (cultural relevance, communication preferences) • Social capital (caregiver presence, confidence in care) • Socioeconomic status and occupation (housing insecurity, financial hardship) 	Good
Crellin <i>et al.</i> , 2024	Across England	To identify disparities in access, engagement, and	Mixed-Methods Evaluation	Step-down	1,361	<ul style="list-style-type: none"> • Barriers to engagement • Health disparities 	Patient engagement varied by age, health status, ethnicity, gender, and employment. Older	<ul style="list-style-type: none"> • Age 	Good

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		experiences with COVID-19 remote monitoring, focusing on inclusivity adaptations and patient factors.				<ul style="list-style-type: none"> • Patient engagement and experiences • Services effectiveness • Staff and carer perspectives 	patients found the service less helpful, minority ethnic groups faced greater comprehension challenges, and illness severity impacted support needs. Living situation and social connections influenced feelings of reassurance.	<ul style="list-style-type: none"> • Health status and comorbidities (illness severity) • Race, ethnicity, culture, and language (minority groups experienced communication barriers) • Social capital (caregiver presence, social connectedness) • Socioeconomic status and occupation (employment status affected engagement) 	
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Appendices

Appendix A – The PRISMA 2020 Checklist

Section and Topic	Item #	Checklist item	Location where item is reported
TITLE			
Title	1	Identify the report as a systematic review.	Page 1
ABSTRACT			
Abstract	2	See the PRISMA 2020 for Abstracts checklist.	Pages 2
INTRODUCTION			
Rationale	3	Describe the rationale for the review in the context of existing knowledge.	Pages 3-4
Objectives	4	Provide an explicit statement of the objective(s) or question(s) the review addresses.	Page 4
METHODS			
Eligibility criteria	5	Specify the inclusion and exclusion criteria for the review and how studies were grouped for the syntheses.	Pages 5-7
Information sources	6	Specify all databases, registers, websites, organisations, reference lists and other sources searched or consulted to identify studies. Specify the date when each source was last searched or consulted.	Pages 5
Search strategy	7	Present the full search strategies for all databases, registers and websites, including any filters and limits used.	Page 6, Appendix B
Selection process	8	Specify the methods used to decide whether a study met the inclusion criteria of the review, including how many reviewers screened each record and each report retrieved, whether they worked independently, and if applicable, details of automation tools used in the process.	Page 6
Data collection process	9	Specify the methods used to collect data from reports, including how many reviewers collected data from each report, whether they worked independently, any processes for obtaining or confirming data from study investigators, and if applicable, details of automation tools used in the process.	Page 6
Data items	10a	List and define all outcomes for which data were sought. Specify whether all results that were compatible with each outcome domain in each study were sought (e.g. for all measures, time points, analyses), and if not, the methods used to decide which results to collect.	Pages 6-7
	10b	List and define all other variables for which data were sought (e.g. participant and intervention characteristics, funding sources). Describe any assumptions made about any missing or unclear information.	Pages 6-7
Study risk of bias assessment	11	Specify the methods used to assess risk of bias in the included studies, including details of the tool(s) used, how many reviewers assessed each study and whether they worked independently, and if applicable, details of automation tools used in the process.	N/A
Effect measures	12	Specify for each outcome the effect measure(s) (e.g. risk ratio, mean difference) used in the synthesis or presentation of results.	N/A
Synthesis methods	13a	Describe the processes used to decide which studies were eligible for each synthesis (e.g. tabulating the study intervention characteristics and comparing against the planned groups for each synthesis (item #5)).	Page 6, Table II
	13b	Describe any methods required to prepare the data for presentation or synthesis, such as handling of missing summary statistics, or data conversions.	N/A
	13c	Describe any methods used to tabulate or visually display results of individual studies and syntheses.	Page 6

Section and Topic	Item #	Checklist item	Location where item is reported
	13d	Describe any methods used to synthesize results and provide a rationale for the choice(s). If meta-analysis was performed, describe the model(s), method(s) to identify the presence and extent of statistical heterogeneity, and software package(s) used.	Page 6
	13e	Describe any methods used to explore possible causes of heterogeneity among study results (e.g. subgroup analysis, meta-regression).	N/A
	13f	Describe any sensitivity analyses conducted to assess robustness of the synthesized results.	N/A
Reporting bias assessment	14	Describe any methods used to assess risk of bias due to missing results in a synthesis (arising from reporting biases).	N/A
Certainty assessment	15	Describe any methods used to assess certainty (or confidence) in the body of evidence for an outcome.	Pages 6
RESULTS			
Study selection	16a	Describe the results of the search and selection process, from the number of records identified in the search to the number of studies included in the review, ideally using a flow diagram.	Pages 7, Figure 1
	16b	Cite studies that might appear to meet the inclusion criteria, but which were excluded, and explain why they were excluded.	Pages 7, Figure 1
Study characteristics	17	Cite each included study and present its characteristics.	Page 7, Table II
Risk of bias in studies	18	Present assessments of risk of bias for each included study.	N/A
Results of individual studies	19	For all outcomes, present, for each study: (a) summary statistics for each group (where appropriate) and (b) an effect estimate and its precision (e.g. confidence/credible interval), ideally using structured tables or plots.	Table II
Results of syntheses	20a	For each synthesis, briefly summarise the characteristics and risk of bias among contributing studies.	Pages 8-10, Table II
	20b	Present results of all statistical syntheses conducted. If meta-analysis was done, present for each the summary estimate and its precision (e.g. confidence/credible interval) and measures of statistical heterogeneity. If comparing groups, describe the direction of the effect.	N/A
	20c	Present results of all investigations of possible causes of heterogeneity among study results.	N/A
	20d	Present results of all sensitivity analyses conducted to assess the robustness of the synthesized results.	N/A
Reporting biases	21	Present assessments of risk of bias due to missing results (arising from reporting biases) for each synthesis assessed.	N/A
Certainty of evidence	22	Present assessments of certainty (or confidence) in the body of evidence for each outcome assessed.	Table II, Appendix C
DISCUSSION			
Discussion	23a	Provide a general interpretation of the results in the context of other evidence.	Page 10
	23b	Discuss any limitations of the evidence included in the review.	Pages 11-12
	23c	Discuss any limitations of the review processes used.	Pages 11-12

Section and Topic	Item #	Checklist item	Location where item is reported
	23d	Discuss implications of the results for practice, policy, and future research.	Pages 10-12
OTHER INFORMATION			
Registration and protocol	24a	Provide registration information for the review, including register name and registration number, or state that the review was not registered.	N/A
	24b	Indicate where the review protocol can be accessed, or state that a protocol was not prepared.	N/A
	24c	Describe and explain any amendments to information provided at registration or in the protocol.	N/A
Support	25	Describe sources of financial or non-financial support for the review, and the role of the funders or sponsors in the review.	N/A
Competing interests	26	Declare any competing interests of review authors.	N/A
Availability of data, code and other materials	27	Report which of the following are publicly available and where they can be found: template data collection forms; data extracted from included studies; data used for all analyses; analytic code; any other materials used in the review.	N/A

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Appendix B – Full Search Strategy

PubMed	
("virtual ward" OR "virtual care model" OR "virtual hospital" OR "virtual inpatient" OR "home hospitalization" OR "hospital@home" OR "home care" OR "home care services" OR "step down care" OR "step up care" OR "remote monitoring" OR "telehealth" OR "telemedicine" OR "telemonitoring" OR "telemangement" OR "remote patient management")	
AND	
("United Kingdom" OR UK OR England OR Scotland OR Wales OR "Northern Ireland" OR "Great Britain" OR NHS)	
Filters - Free full text, Full text, Clinical Trial, Comparative Study, Controlled Clinical Trial, Evaluation Study, Government Publication, Observational Study, Randomized Controlled Trial, English, Humans, Adult: 19+ years, 2000 to present	
CINAHL-PLUS (EbscoHost)	
Search ID	Search Terms
S1	TI "virtual ward" OR AB "virtual ward"
S2	TI "virtual care model" OR AB "virtual care model"
S3	TI "virtual hospital" OR AB "virtual hospital"
S4	TI "virtual inpatient" OR AB "virtual inpatient"
S5	TI "home hospitalization" OR AB "home hospitalization"
S6	TI "hospital at home" OR AB "hospital at home"
S7	TI "home care" OR AB "home care"
S8	TI "home care services" OR AB "home care services"
S9	TI "step down care" OR AB "step down care"
S10	TI "step up care" OR AB "step up care"
S11	TI "remote monitoring" OR AB "remote monitoring"
S12	TI "telehealth" OR AB "telehealth"
S13	TI "telemedicine" OR AB "telemedicine"
S14	TI "telemonitoring" OR AB "telemonitoring"
S15	TI "telemangement" OR AB "telemangement"
S16	TI "remote patient management" OR AB "remote patient management"
S17	S1 OR S2 OR S3 OR S4 OR S5 OR S6 OR S7 OR S8 OR S9 OR S10 OR S11 OR S12 OR S13 OR S14 OR S15 OR S16
S18	TI "united kingdom" OR AB "united kingdom"
S19	TI "uk" OR AB "uk"
S20	TI "england" OR AB "england"
S21	TI "scotland" OR AB "scotland"
S22	TI "wales" OR AB "wales"
S23	TI "northern ireland" OR AB "northern ireland"
S24	TI "great britain" OR AB "great britain"
S25	TI "nhs" OR AB "nhs"
S26	S18 OR S19 OR S20 OR S21 OR S22 OR S23 OR S24 OR S25
S27	S17 AND S26
Limiters - Full Text; References Available; Abstract Available; Publication Year: 2000-2024; Publication Date: 20000101-20241130; Peer Reviewed; English Language; Research Article; Human; Geographic Subset: UK & Ireland; Age Groups: Adult, 19-44 years, Middle Age, 45-64 years, Aged, 65+ years, Aged, 80 and over; Language: English	
Expanders - Apply related words; Apply equivalent subjects	
Search modes - Find any of my search terms	
MedRxiv	
("virtual ward" OR "virtual care model" OR "virtual hospital" OR "virtual inpatient" OR "home hospitalization" OR "hospital@home" OR "home care" OR "home care services" OR "step down care" OR "step up care" OR "remote monitoring" OR "telehealth" OR "telemedicine" OR "telemonitoring" OR "telemangement" OR "remote patient management")	
AND	
("United Kingdom" OR UK OR England OR Scotland OR Wales OR "Northern Ireland" OR "Great Britain" OR NHS)	

Source: Author's own work

Appendix C – Quality Assessment of Included Studies Using the MMAT Checklist

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Author & Date	Screening Questions		Qualitative Studies					
	Are there clear research questions?	Do the collected data allow to address the research questions?	Is the qualitative approach appropriate to answer the research question?	Are the qualitative data collection methods adequate to address the research question?	Are the findings adequately derived from the data?	Is the interpretation of results sufficiently substantiated by data?	Is there coherence between qualitative data sources, collection, analysis and interpretation?	Comments
Dismore et al., 2019	Yes	Yes	Yes	Yes	Yes	Yes	Yes	N/A
			Quantitative RCTs					
			Is randomization appropriately performed?	Are the groups comparable at baseline?	Are there complete outcome data?	Are outcome assessors blinded to the intervention provided?	Did the participants adhere to the assigned intervention?	Comments
Cotton et al., 2000	Yes	Yes	Yes	Yes	Yes	No	Yes	Outcome assessors were not blinded; the nature of the interventions (home-based care vs. traditional hospital care) made blinding impractical, and the studies did not report blinding procedures.
Dar et al., 2009	Yes	Yes	Yes	Yes	Yes	No	Yes	

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Quantitative Non-Randomized Studies								
			Are the participants representative of the target population?	Are measurements appropriate regarding both the outcome and intervention (or exposure)?	Are there complete outcome data?	Are the confounders accounted for in the design and analysis?	During the study period, is the intervention administered (or exposure occurred) as intended?	Comments
Bewis et al., 2017	Yes	Yes	Yes	Yes	Yes	Unclear	Yes	The study did not clearly report whether potential confounders were controlled for in the design or analysis.
Mixed Methods Studies								
			Is there an adequate rationale for using a mixed methods design to address the research question?	Are the different components of the study effectively integrated to answer the research question?	Are the outputs of the integration of qualitative and quantitative components adequately interpreted?	Are divergences and inconsistencies between quantitative and qualitative results adequately addressed?	Do the different components of the study adhere to the quality criteria of each tradition of the methods involved?	Comments
Nunan et al., 2020	Yes	Yes	Yes	Yes	Yes	Unclear	Yes	Limited information was provided in the included records regarding how the authors addressed divergences and inconsistencies
The Health Foundation, 2023	Yes	Yes	Yes	Yes	Yes	Unclear	Yes	
Wessex Academic	Yes	Yes	Yes	Yes	Yes	Unclear	Yes	

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Health Science Network, 2022								between the quantitative and qualitative data.
Health Innovation Network South London, 2021	Yes	Yes	Yes	Yes	Yes	Unclear	Yes	

Source: Author's own work

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