ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

Effectiveness Of Speech Therapy Interventions In Parkinsonism: Mechanisms And Rehabilitation: A Systematic Review

Dr Arthi G K ¹, Dr Lavanya Sekhar ², Dr Anu Krishnan Kumar ³, Dr Muthu Prathibha K ⁴, Dr Meennakshi BS ⁵, Dr PriscillaJohnson ¹

¹.MD Postgraduate, Department of Physiology, SRMC & RI, SRIHER (DU) India;

Email Id - arthigk3499@gmail.com

¹ Associate Dean (Preclinical), Professor, Department of Physiology, SRMC & RI, SRIHER (DU) India; (Corresponding Author)

Email Id - priscillajohnson@sriramachandra.edu.in

² Assistant Professor, Dept. of Physiology, Panimalar Medical College Hospital and Research Institute, Chennai -123 India;

Email Id - drlavanyas1981@gmail.com

³Senior lecturer Department of Public Health Dentistry, Rajas Dental College & Hospital, Kavalkinaru Junction, Tirunelveli, India;

Email Id - anukrishnakumar1996@gmail.com

⁴Professor & HOD, Department of Physiology, SRMC & RI, SRIHER (DU) India

Email Id - prathibhakm@sriramachandra.edu.in

⁵Assistant Professor, Department of Physiology, SRMC & RI, SRIHER (DU) India;

Email Id - medico.meenaa@gmail.com

ABSTRACT

Background: Speech impairments, especially hypokinetic dysarthria, affect up to 89% of Parkinson's disease patients, impairing communication, and general lifestyle. Speech-language therapy (SLT) is a key non-drug approach, offering greater benefit than medications for speech issues.

Objective: This systematic review evaluates the efficiency of speech and language therapy in Parkinsonism, investigates underlying therapeutic mechanisms, and examines the impact of delivery modes and personalized intervention strategies.

Methods: A complete literature search was conducted following the PRISMA 2020 guidelines. Studies included adults with Parkinson's disease receiving speech and language therapy (SLT) interventions, such as Lee Silverman Voice Treatment (LSVT) LOUD, LSVT ARTICulation, clear speech strategies, music therapy, and telehealth-based programs. 13 studies met inclusion criteria: randomized controlled trials, experimental, and quasi-experimental designs.

Results: LSVT LOUD significantly improved vocal loudness, speech intelligibility, and quality of life. Telehealth-based SLT showed comparable efficacy to in-person delivery, improving accessibility. Multi-modal and customized approaches yielded positive outcomes in communication and functional independence. However, heterogeneity in protocols and outcome measures limited cross-study comparability.

Conclusion: SLT interventions, particularly LSVT LOUD, are effective in improving speech outcomes in PD. Emerging modalities such as telehealth and individualized, multi-disciplinary rehabilitation enhance therapy accessibility and impact. Future research should prioritize long-term, patient-centered studies and standardized outcome metrics to strengthen evidence-based clinical practice.

Keywords: Parkinson's Disease, Speech Therapy, Hypokinetic Dysarthria, Telehealth, Systematic Review, Communication Disorders

INTRODUCTION

Parkinsonism, including Parkinson's disease (PD) and related disorders, notably affects quality of life through motor symptoms like bradykinesia, tremor, rigidity, and speech difficulties [5]. Hypokinetic dysarthria, affecting up to 89% of PD patients, manifests as altered speech rhythm, monotonous pitch, decreased vocal volume, and impaired articulation [3,4]. These communication challenges contribute to social withdrawal and psychological distress, emphasizing the need for effective therapeutic strategies [1].

Dopamine depletion is the primary cause of speech impairment in PD, and while pharmacological treatments like Levodopa may alleviate some motor symptoms, their effectiveness on speech-related issues is inconsistent [4,6]. This has heightened the emphasis on speech and language therapy (SLT) as a crucial non-pharmacological treatment. SLT has shown significant improvements in speech parameters, though outcomes depend on dysarthria severity,

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

therapy duration, and adherence [1,3]. Intensive techniques like Lee Silverman Voice Treatment (LSVT LOUD) focus on vocal loudness and show promising outcomes [8,9].

Emerging modalities, including telehealth-based SLT, offer improved accessibility and hold potential for enhancing treatment engagement [2]. Research continues on how tailored SLT programs can optimize speech outcomes by addressing individual patient needs, while multidisciplinary approaches combining speech therapy with physical and occupational therapies have shown effectiveness in improving overall functional independence. Despite advances, there remains a need for standardized protocols, clearer outcome measures, and exploration of alternative interventions.

This systematic review aims to evaluate the efficiency of various SLT interventions in improving speech function in individuals with PD, explore their underlying mechanisms, and assess their role in fostering long-term communication abilities. By synthesizing recent clinical trials and meta-analyses, it provides evidence-based insights to guide clinicians and policymakers, improving patient care and quality of life (QoL).

METHODOLOGY

The review is guided by specific research questions and structured using the PICOS framework to ensure a comprehensive and focused analysis.

Research Questions:

- RQ.1 How effective are different speech therapy interventions in enhancing speech function in individuals with Parkinson's disease?
- RQ.2 Can customized treatment programs maximize outcomes?
- RQ.3 Evaluate the differences in intervention protocols, therapy durations, and outcome assessments?

PICOS Framework:

- Population: adults diagnosed with PD presenting with dysarthria or communication impairments;
- **Intervention**: speech therapy interventions, including LSVT LOUD, LSVT ARTIC, clear speech strategies, multidisciplinary rehabilitation, music therapy, and tele-rehabilitation approaches;
- Comparator: standard care, no treatment, sham therapy, or alternative speech therapy interventions;
- Outcomes: primary outcomes such as voice intensity (SPL), intelligibility, articulation, phonation measures, and secondary outcomes including QoL indices, VHI, and caregiver burden;
- **Study design**: randomized controlled trials (RCTs), parallel-group, and experimental studies. Eligible studies were screened and selected according to predefined inclusion criteria.

Search Strategy: A systematic literature search was conducted in accordance with PRISMA 2020 (Preferred Reporting Items for Systematic Reviews and Meta-Analyses) guidelines. The databases searched included PubMed, Embase, Scopus, Cochrane Library, Web of Science, and CINAHL. The search strategy utilized a combination of Medical Subject Headings (MeSH) terms and free-text keywords, with Boolean operators (AND, OR) to refine results. Key terms included "Parkinson's disease," "parkinsonism," "speech therapy," "speech and language therapy," "hypokinetic dysarthria," "LSVT LOUD," "telerehabilitation," "multi-modal therapy," "communication disorders," and "voice treatment." Manual searches of reference lists were also conducted to ensure comprehensive coverage. Selection Criteria: Among 638 articles screened, 13 were included in the review (Figure 1). These studies involved individuals diagnosed with Parkinson's disease or parkinsonism, focusing on RCTs, cohort studies, and systematic reviews/meta-analyses. Only studies evaluating SLT interventions and reporting quantitative speech-related outcomes, such as vocal intensity, articulation, and intelligibility, were included. Peer-reviewed full-text articles published in English were considered. Exclusion criteria included case reports, conference abstracts, editorials, and studies without measurable speech-related outcomes. Non-English publications without available translations were also excluded. This approach ensured the inclusion of high-quality evidence while minimizing bias.

Data Extraction and Synthesis: This was independently conducted by two reviewers (Dr. Arthi and Dr. Lavanya) using a standardized form. Extracted data included various parameters, as shown in Tables 1 and 2. Discrepancies were resolved through discussion and consensus. A qualitative synthesis of study characteristics and outcomes was performed, with quantitative comparisons made, considering variability in interventions, delivery modes, and outcome measures.

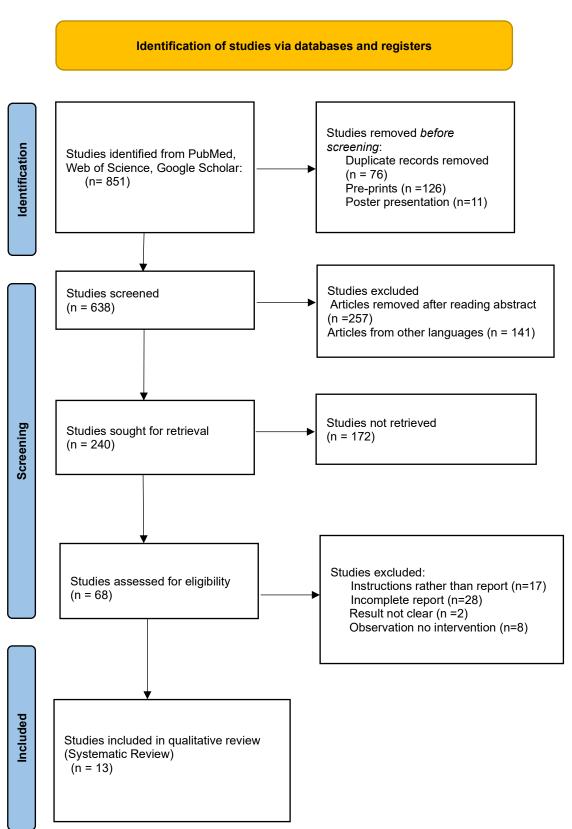
Risk of Bias Assessment: The methodological quality of RCTs was assessed using the Cochrane Risk of Bias 2.0 (RoB 2.0) [19] tool and the Newcastle-Ottawa Scale (NOS) [20]. RoB 2.0 evaluated 5 domains, with judgments

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

classified as low risk or some concerns. NOS assessed Selection, Comparability, and Exposure, assigning stars for quality.

Figure 1: PRISMA 2020 Flowchart for the review



ISSN: **2229-7359** Vol. 11 No. 7, 2025

Table 1: Summary of Key Studies on Speech Therapy Interventions in PD

Author (Year)	Design	n	Populatio n (Age/Gen der)	PD Details	Compari son	Setting	Outcomes & Conclusion
Theodo ros et al. (2016) [9]	Single- blind RCT	52	71.0 ± 8.8; 36M/16F	H&Y 1- 5; mild- mod dysarthri a	F2F (16), Online (15), Non- metro Online (21)	In- person/Teleh ealth	Online = F2F; ↑ SPL, intelligibility, QoL
Lam et al. (2016) [17]	Acoustic study	28	PD: 68.3 ± 6.7; C: 67.8 ± 6.5; 9M/5F PD	Idiopathi c PD; mild dysarthri a; some LSVT	Healthy controls	In-person	Overenuncia tion ↑ articulation; hearing-impaired ↑ prosody
Ramig et al. (2018)	Unblind ed RCT	64 PD; 20 HC	PD ~67 yrs, 68.8% M; HC ~64 yrs	H&Y I- IV; 5 yrs disease	LSVT LOUD, ARTIC, untreated	In-person	LSVT LOUD ↑ SPL, CETI- M at 1 & 7 mo
Ferrazzo li et al. (2018) [10]	Single- blind RCT	234	66.5 (exp), 66.9 (ctrl); 57% M	H&Y 2- 4; stable meds	No rehab	In-person	MIRT ↑ QoL, motor & functional outcomes
Sackley et al. (2020) [21]	Multicen tre RCT	546	Not reported	Idiopathi c PD	No Rx, NHS SLT, LSVT	In-person	SLT ↑ speech vs. no Rx
Levy et al. (2020) [7]	RCT	57	66.5 ± 8.5; 72% M	H&Y I- IV; ~4.9 yrs dx	LSVT LOUD, ARTIC, no Rx	In-person	LSVT LOUD ↑ intelligibility
Brabene c et al. (2021) [22]	Sham- controlle d RCT	33	71.7 (real), 71.5 (sham); 14M/6F	Mild- mod dysarthri a	Sham rTMS	In-person	rTMS ↑ phonetics, brain activation (8 wk effect)
Maas et al. (2022) [12]	RCT (Protoco l)	215	<46 to >66; stratified genders	All PD stages; ↓ intelligibi lity	Waitlist (8 wk)	Telehealth	Home voice app; evaluating QoL, speech
Mohsen i et al. (2023) [23]	RCT	33 (25M/ 8F)	Mean 58.9 yrs	H&Y 1- 2; dysphagia	ST, Music Tx	Telehealth (WhatsApp)	Combo ↑ swallowing; telerehab feasible

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

Maas et	RCT	214	68.1 yrs;	All	Waitlist	Telehealth	1
al.			27% F	stages;	(105)		communicati
(2024)				dysarthri			on QoL; no
[11]				a; mean			QoL change
				dx 7.8 yrs			
Sackley	Unblind	388	~70 yrs;	H&Y ≤3;	LSVT,	In-	LSVT ↓
et al.	ed RCT		74% M	dysarthri	NHS	person/home;	dysarthria
(2024)				a	SLT, no	LSVT remote	impact; no
[8]					SLT		AE
Sackley	Phase III	388	~70 yrs;	H&Y ≤2	LSVT,	In-	LSVT ↑
et al.	RCT		74% M	in 61%;	NHS	person/home	speech; NHS
(2024)				5-6 yrs	SLT, no		SLT =
[18]				PD	SLT		control
Steurer	RCT	95	≥60 yrs;	H&Y 2-	HiComm	In-	HiComm ↑
et al.	(EXPAN		MoCA	3; mild-	vs.	person/home	post-voice;
(2024)	d)		≥21	mod PD	HiBalanc		effect not
[24]					e		retained

Footnote: F2F: Face-to-Face; SPL: Sound Pressure Level; QoL: Quality of Life; H&Y: Hoehn and Yahr; LSVT: Lee Silverman Voice Treatment; SLT: Speech-Language Therapy; PD: Parkinson's Disease; VHI: Voice Handicap Index.

Table 2: Summary of Key Findings, Outcome Measures, and Efficacy of Speech Therapy Interventions in Parkinsonism

Auth or / Year	Interventi on Type	Primar y Outco mes	Secondary Outcomes	Main Findings	Statistica l Significa nce (p- value)	Effect Size	Mean Outcome Scores (Pre/Post/F ollow-up)	Durati on of Interve ntion / Follow -up
Theo doros DG et al. (2016) [9]	LSVT LOUD (face-to- face & telerehabil itation)	SPL in monolo gue (dB)	SPL in sustained phonation /reading; Max F0; intelligibili ty, loudness, pitch variability; Dysarthria Impact Profile; PDQ-39	Telereha b noninfer ior to face-to- face. Significa nt improve ments in loudness , intelligib ility; no differenc e between delivery modes.	p < .001 (time effect); no group × time interacti on (p > .05)	Not report ed	SPL (Mono): 69.6/77.1 (FTF), 70.2/76.4 (Online); Intelligibility: 107.3/127.1 (FTF), 120.5/125.9 (Online)	16 session s (4 weeks); post- treatm ent follow- up

Vol. 11 No. 7, 2025

Lam J et al. (2016) [17]	Clear Speech Variants (Habitual, Clear, Overenun ciate, Hearing Impaired)	Vowel Space Area (VSA), vowel duratio n, fricative duratio n	F0, SPL, articulatio n rate	Overenu nciate improve d VSA, vowel duration , articulati on rate. Hearing impaired increase d SPL and F0. Feasible interven tions for PD.	p < 0.05 for multiple measures	$\eta p^2 = 0.02 - 0.70$	Pre: 69.7–71.5; Post: 70.3–72.5; Follow-up: 73.2–76.6	Single- session tasks; no follow- up reporte d
Rami g L et al. (2018) [6]	LSVT LOUD vs. LSVT ARTIC (both intensive, PD- specific)	SPL in reading & spontan eous speech	CETI-M	LSVT LOUD improve d SPL significa ntly at 1 & 7 months. CETI-M improve d at 1 month for both groups but maintai ned at 7 months only in LOUD group.	SPL p < 0.05; CETI-M p = 0.02 (1 mo), p = 0.08 (7 mo)	SPL: ES = 1.53 (LOU D vs ARTI C), 2.19 (LOU D vs UNT XPD)	LSVT LOUD: Pre 70.5 / Post 76.8 / Follow-up 75.2; LSVT ARTIC: Pre 71.9 / Post 73.2 / Follow-up 72.5	month (16 session s); 7- month follow- up
Ferraz zoli D et al. (2018) [10]	Multidisci plinary Intensive Rehabilita tion Treatment (MIRT) incl. speech therapy	PDQ- 39 Global Index	UPDRS, PDDS, TUG, BBS, Levodopa dosage, neuropsyc hological tests	Significa nt QoL improve ment (PDQ-39 ↓ by 8.3 at 10 wks, maintai ned at 18 wks);	PDQ-39, UPDRS, PDDS, TUG, BBS: p < 0.0001	Not report ed	PDQ-39: 43.6 / 35.3 / 38.8; UPDRS: 39.6 / 27.2 / -; BBS: 46.8 / 52.5 / -	weeks; follow- up at 10 and 18 weeks

Vol. 11 No. 7, 2025

Sackl ey CM et al. (2020) [21]	NHS SLT, LSVT LOUD	VHI total score	PDQ-39, QASD, EQ-5D-5L, ICECAP- O	UPDRS, PDDS, TUG, BBS improve d. Levodop a dosage reduced. Both interven tions improve d outcome s over control.	p < 0.01	0.38	Not reported	3 month s/12 month s
Levy ES et al. (2020) [7]	LSVT LOUD vs LSVT ARTIC vs No Treatment	Transcr iption accurac y (TA)	None reported	LSVT LOUD significa ntly improve d intelligib ility (TA +31.5%, p < 0.0001); ARTIC had non- significa nt improve ment; No Treatme nt group declined	Voice vs ARTIC: p = 0.04; Voice vs No Tx: p = 0.0002	Voice vs ARTI C: ES = 1.0; Voice vs No Tx: ES = 1.8	Voice: 53.6 / 85.1; ARTIC: 44.8 / 51.6; No Tx: 64.4 / 52.5	4 weeks; post- treatm ent follow- up (6- month data collect ed but not reporte d)
Brabe nec L et al. (2021) [22]	rTMS over right superior temporal gyrus	Phonetics score (articulation, prosody, intelligibility)	Brain activation (fMRI), connectivit y changes	rTMS improve d phonetic s scores vs sham; effects up to 8 weeks; enhance d	Time × group p = 0.040	Cohe n's d = 1.391 (rTM S)	Phonetics score: Pre ~21 / Post ~23.5 / Follow-up ~24.5	10 session s (2 weeks); follow- up at 2, 6, 10 weeks

Vol. 11 No. 7, 2025

		1		. •				
				connecti vity in				
				motor				
				speech				
				regions.				
Maas	Personaliz	PDQ-	Voice	Ongoing	Not	Not	Not	8
JJL et	ed speech	39	quality	study.	reported	report	reported	weeks;
al.	therapy +	summar	(Radboud	Hypothe	- op or or or	ed		follow-
(2022	Voice	y index	Dysarthria	sized				up at 8
) [12]	Trainer		Assessmen	improve				& 32
	app via		t), VHI,	ments in				weeks
	Telehealth		intelligibili	speech				
			ty,	intelligib				
			caregiver	ility and				
			burden,	QoL,				
			mood/anx	maintai				
			iety,	ned at 6-				
			swallowing	month follow-				
Mohs	Combinati	Swallow	DHI	up. Combin	p < 0.05	Not	SDQ (CT):	12
eni Z	on therapy	ing	subscales	ation	p (0.03	report	16.27→4.64	session
et al.	(speech +	Disturb	(functional	therapy		ed	→4.73; DHI	s (4
(2023	music),	ance	, physical,	was			(CT):	weeks);
) [23]	speech	Questio	emotional)	most			29.18→11.6	3-
	therapy,	nnaire		effective.			4→11.91;	month
	music	(SDQ),		Music			ST & MT	follow-
	therapy	Dyspha		therapy			scores	up
		gia		improve			similar	
		Handic		d .			trends.	
		ap		emotion				
		Index		al				
		(DHI)		aspects				
				more than				
				physical.				
Maas	Personaliz	PDQ-	PDQ-39	Significa	PDQ-39	Not	PDQ-39:	8
JJ et	ed remote	39	communic	nt	summary	report	28.5 / 24.7	weeks;
al.	speech	summar	ation,	improve	p=0.056	ed	/ 26.6;	follow-
(2024	therapy via	y index	VHI,	ment in	(NS);		ROMP	up at
)[11]	Telehealth		ROMP,	commun	Commu		Speech: 17.8	32
			MLL, DIT	ication	nication		/ 15.55 /	weeks
				subdom	p=0.011;		16.8; VHI:	
				ain and	VHI		35.9 / 30.9	
				VHI; no	p=0.002		/ 32.0	
				overall				
				PDQ-39				
				improve				
				ment. VHI				
	<u> </u>	1		A 1 1 1		l		

Vol. 11 No. 7, 2025

Sackl ey CM et al. (2024) [8]	LSVT LOUD vs NHS SLT vs no SLT	VHI total score (3 months)	VHI subscales, QASD, PDQ-39, ICECAP- O, EQ-5D, PDQ- Carer	effect maintai ned at 32 weeks. LSVT LOUD improve d VHI at 3 months vs no SLT and NHS SLT. NHS SLT. showed no benefit. LSVT LOUD continue d benefit at 12	LSVT LOUD vs no SLT: p<0.001; NHS SLT vs no SLT: p=0.43	VHI ES ~0.38	VHI: 44.6 / 35.0 / 38.2; PDQ-39: 27.9 / 27.6 / 28.5; QASD: 29.9 / 23.5 / 25.6	LSVT LOUD: 4 weeks; NHS SLT: 6-8 weeks; 12-month follow-up
Sackl ey CM et al. (2024) [18]	LSVT LOUD vs NHS SLT	VHI total score (3 months)	PDQ-39, QASD, EQ-5D-5L, ICECAP- O, PDQ- Carer	months. LSVT LOUD reduced VHI significa ntly at 3 months vs control and NHS SLT; sustaine d effects at 12 months. NHS SLT showed no benefit. Significa	LSVT LOUD vs Control: p=0.000 1; LSVT LOUD vs NHS SLT: p<0.000 1; NHS SLT vs Control: p=0.4	VHI differ ence LSVT LOU D vs Contr ol: - 8.0 (99% CI: - 13.3 to - 2.6)	EQ-5D-5L: LSVT 0.637/0.605 /0.590; VHI: LSVT 44.6/36.7/3 8.2; PDQ- 39: LSVT 27.9/27.6/2 8.5	LSVT LOUD: 4 weeks; NHS SLT: 6 weeks; 12- month follow- up
er H et al.	HiComm unication (group-	voice sound level	AVQI, HNR	Significa nt improve	p = 0.003 (voice	Not report ed	Voice Sound Level (Text	Group session s;

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

(2024	based	(text	ments in	sound	Reading):	follow-
) [24]	speech &	reading)	voice	level), p	70.8 / 73.0	up at 6
	communic		sound	= 0.016	/ 70.4;	month
	ation		level and	(AVQI),	Voice	S
	training)		voice	p =	Sound Level	
			quality	0.014	(Monologue	
			post-	(HNR)): 69.0	
			interven		(truncated	
			tion; no		data)	
			sustaine			
			d effect			
			at 6			
			months.			

Footnote: dB SPL: decibel sound pressure level; SP: speech pathologist

RESULTS

Table 1 summarizes 13 studies evaluating speech therapy interventions in Parkinsonism, including RCTs, experimental, and acoustic studies. A total of 2,377 patients were assessed, with sample sizes ranging from 28 to 546 participants. Patient ages varied from 58.9 to 71.7 years, with a male predominance of up to 74%. Most studies included individuals with idiopathic PD, Hoehn and Yahr stages 1 to 5, and dysarthria severity ranging from mild to severe. Interventions examined were LSVT LOUD, LSVT ARTIC, repetitive transcranial magnetic stimulation (rTMS), music therapy, and telehealth-based programs. Primary outcomes measured speech intelligibility, sound pressure level (SPL), and quality of life (QoL). LSVT LOUD consistently improved loudness and QoL versus standard or no treatment [6,7,8,18,21]. Telehealth delivery showed non-inferiority to in-person therapy [9,11,12,23,24].

Table 2 summarizes the key findings, outcome measures, and efficacy of speech therapy interventions in PD. Interventions included LSVT LOUD, LSVT ARTIC, clear speech strategies, rTMS, music therapy, multidisciplinary rehabilitation, and telehealth-based approaches. The primary outcomes were speech loudness, intelligibility, articulation, and quality of life (QoL). LSVT LOUD consistently improved speech sound pressure level (SPL), increasing monologue SPL from 69.6 dB to 77.1 dB (p < 0.001) [9,12]. Clear speech variants enhanced vowel space area and articulation rate, with effect sizes up to ηp^2 = 0.70 [17]. Multidisciplinary programs reduced PDQ-39 scores by 8.3 points (p < 0.0001) [10]. Telehealth-delivered LSVT LOUD demonstrated non-inferiority to face-to-face interventions [9,11,12]. Voice Handicap Index (VHI) scores improved significantly post-LSVT LOUD, reducing by 8–9.6 points (p < 0.001) [7,8,18,21].

Table 3: Quality assessment done for the randomised control studies done using the Newcastle Ottawa Scale (NOS)

S.	Study	Sele	S2	S3	S4	Comparabil	Exposure	E	Е	Total
No		ctio				ity	E1	2	3	Score
		n								(Max
		S1								9
										Stars)
1	Theodoros DG et al. (2016) [9]	*	*	*	*	*	*	,	*	7

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

2.	Ramig L et al. (2018) [6]	*	*	*	*	**	*	*	*	9
3.	Ferrazzoli D et al. (2018) [10]	*	*	*	*	*	*	*	*	8
4.	Sackley CM et al. (2020) [21]	*	-	*	*	*	*	*	*	7
5.	Levy ES et al. (2020) [7]	*	*	-	*	**	*	*	*	8
6.	Brabenec L et al. (2021) [22]	*	*	*	*	*	*	*	*	8
7.	Maas JJL et al. (2022) [12]	*	*	*	*	**	*	*	*	9
8.	Mohseni Z et al. (2023) [23]	*	*	,	*	**	*	*	*	8
9.	Maas JJ et al. (2024) [11]	*	*	*	*	**	*	*	*	9
10	Sackley CM et al. (2024) [8]	*	*	*	*	**	*	*	*	9
1	Sackley CM et al. (2024) [18]	*	*	*	*	**	-	*	*	8
12	Steurer H et al. (2024) [24]	*	*	*	*	**	*	*	*	9

Table 3 presents the quality assessment of 12 RCTs using the NOS, with total scores ranging from 7 to 9 stars. Most studies demonstrated high methodological quality. 9 studies achieved 8 or more stars, including Ramig et al. (2018), Maas et al. (2022), and Steurer et al. (2024), each scoring the maximum 9 stars, indicating excellent selection, comparability, and outcome measures [6,11,12,24]. Theodoros et al. (2016) and Sackley et al. (2020) scored 7 stars due to limitations in comparability or exposure domains [9,21]. Consistency in selection criteria and exposure ascertainment was evident, ensuring the robustness of findings across the RCTs [7,8,10,18,22,23].

Table 4 details the quality appraisal of one quasi-experimental study, assessed using the JBI checklist. Lam et al. (2016) met 8 out of 9 quality criteria, indicating high methodological rigor [17]. The study adequately addressed participant selection, intervention clarity, outcome measurement, and statistical analysis. However, blinding was not reported, and one item was not applicable.

Table 4: Quality assessment done for quasi experimental study using the JBI tool for assessment

S. No	Study	Question Q1	Q2	Q3	Q4	Q5	Q6	Q7	Q8	Q9	Total Score (Max 9 Stars)
	Lam J et al. (2016) [17]	Yes	Yes	Yes	Yes	No	Not applicable	Yes	Yes	Yes	8 (high quality)

Figure 2 illustrates the individual risk of bias assessment for twelve RCTs using the ROB2 tool. The assessment covers five domains where most studies showed a low risk of bias in D2, D4, and D5, as indicated by green markers. However, concerns were identified in D1 and D3 in several studies, marked by yellow indicators. Specifically, Theodoros DG et al. (2016), Brabenec L et al. (2021), and Sackley CM

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

et al. (2024) had concerns related to randomization (D1) and missing data (D3). Overall, most studies demonstrated a low risk of bias, with concerns in specific domains.

Figure 2: Individual studies risk of bias assessment for RCT studies done using the ROB2 tool

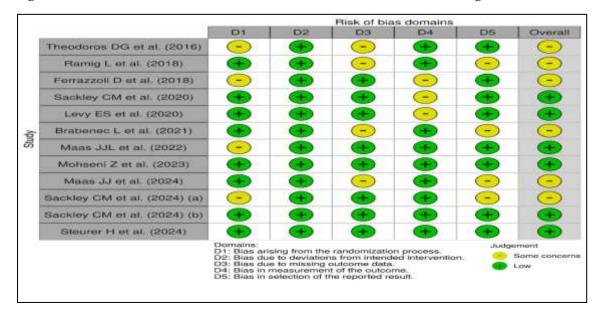


Figure 3 illustrates the overall risk of bias assessment for the included RCT studies based on the ROB2 tool. The bar chart summarizes the proportion of studies classified as low risk and those with some concerns in each domain. Approximately 50% of the studies show low risk in D1 (randomization), while the remaining studies have some concerns. In contrast, all studies show low risk in D2 (deviations from intended interventions). Domains D3 (missing outcome data), D4 (measurement of outcome), and D5 (selection of reported results) show higher proportions of studies with concerns, especially D3 and D5. Overall, more than half of the RCTs were assessed with some concerns, while the rest were classified as low risk.

Figure 3: Overall risk of bias assessment for RCT studies using the ROB2 tool

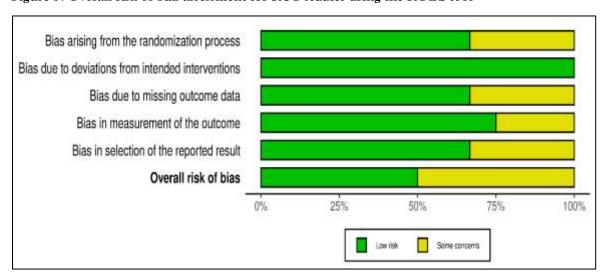


Figure 4 presents the individual risk of bias assessment for the quasi-experimental study by Lam J et al. (2016) [17], evaluated using the ROBINS-I tool across seven domains: D1 (confounding), D2 (selection of participants), D3 (classification of interventions), D4 (deviations from intended interventions), D5 (missing data), D6 (measurement of outcomes), and D7 (selection of reported results). The study shows

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

low risk in D1, D3, D4, and D5. However, D6 is rated as moderate risk, indicating concerns with outcome measurement, and D2 and D7 have no information, reflecting uncertainty in participant selection and reported results.

Figure 4: Individual studies risk of bias assessment for Quasi study using the ROBINS I tool

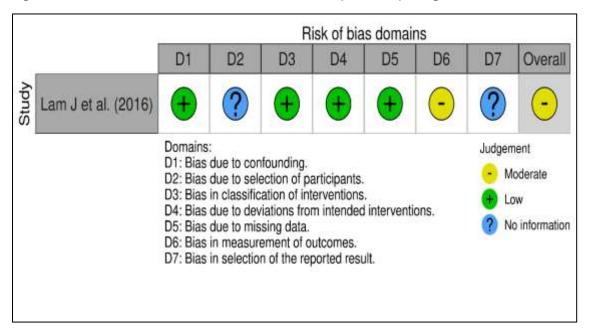


Figure 5: Overall risk of bias assessment for Quasi studies using the ROBINS tool

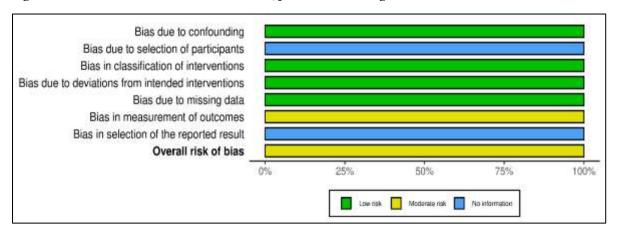


Figure 5 summarizes these findings, showing that while most domains indicate low risk, the study demonstrates a moderate overall risk of bias due to limitations in D6, D2, and D7.

DISCUSSION

PD is a progressive neurodegenerative disorder that impairs motor and non-motor functions, with hypokinetic dysarthria affecting up to 89% of individuals, reducing vocal loudness, pitch, and articulation [1,2,3,4]. This systematic review analysed 13 studies, including RCTs, prospective, and quasi-experimental designs, evaluating the effectiveness of various SLT interventions such as LSVT LOUD, LSVT ARTIC, and Muscle Intensive Respiratory Therapy (MIRT). The findings consistently support the efficacy of LSVT LOUD in improving speech outcomes, particularly vocal loudness and intelligibility [8,6,11,12,18].

Effectiveness of interventions:(RQ.1)

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

LSVT LOUD emerged as the most consistently effective intervention, with multiple RCTs showing significant gains in sound pressure levels (SPL) articulation, and speech intelligibility. [6,7,8,]. Improvements in vowel space area and articulation rate have also been reported, reflecting enhanced speech intelligibility [17,25]. Multiple RCTs and meta-analyses report significant gains in sound pressure levels (SPL), from 69.6 dB to 77.1 dB with face-to-face delivery [9]. LSVT LOUD outperforms LSVT ARTIC and NHS SLT, with an SPL effect size of 2.19 [6,18]. The underlying mechanisms of LSVT LOUD involve compensatory activation of neural pathways that address deficits in motor control caused by dopamine depletion, which contributes to hypokinetic dysarthria [3,5].

Randomized controlled trials confirm that remotely delivered interventions are non-inferior to face-to-face therapies in enhancing speech intelligibility, vocal loudness, and quality of life [9,11,29]. Besides, Tele-rehabilitation delivery of LSVT LOUD demonstrated non-inferior outcomes to in-person therapy, offering a feasible, effective option for patients with mobility limitations, geographic barriers and limited access to in-person services [11,12,26,27,28]. The PERSPECTIVE study supports personalized remote interventions aimed at improving conversational abilities [12]. Constantinescu et al. [23] validated telehealth delivery through a non-inferiority trial, showing sustained speech improvements. However, as novel interventions and technologies continue to emerge, there is a need for further research directly comparing the efficacy, long-term outcomes, and patient adherence of tele rehabilitation of LSVT LOUD with these newer approaches [7,24]. Such comparative studies would help determine the most effective and sustainable speech therapy strategies for individuals with Parkinsonism. While technological advancements offer real-time feedback and better adherence [9,29], heterogeneity in protocols and outcomes underscores the need for standardized methodologies in future studies [2,30].

Multi-modal rehabilitation strategies combining speech therapy with physical, occupational, and cognitive interventions show significant potential in improving communication and functional independence in individuals with PD [5,16,30]. Ferrazzoli et al. [10] reported notable QoL improvements, with PDQ-39 scores reduced by -8.3 ± 18.0 (p<0.0001). These outcomes support previous studies emphasizing the importance of addressing both motor and non-motor symptoms in PD management [30,31]. Comprehensive programs incorporating rhythmic cueing, respiratory exercises, and cognitive training further enhance speech production and communicative effectiveness [5,16]. The addition of music therapy to speech-language therapy (SLT) has been associated with improved speech clarity and vocal loudness [32]. Neuroimaging studies suggest interventions like LSVT LOUD and HiCommunication promote neuroplasticity and strengthen neural connectivity [7,19]. However, heterogeneity in protocols and outcomes complicates comparisons, highlighting the need for standardized methodologies and long-term follow-up research.

Although LSVT LOUD consistently demonstrates superior improvements in vocal loudness and speech intelligibility in individuals with PD, alternative interventions like LSVT ARTIC, MIRT (Multidisciplinary Intensive Rehabilitation Therapy), and HiCommunication offer notable benefits in specific speech domains. LSVT ARTIC, emphasizing articulatory precision rather than vocal loudness, has been shown to improve speech clarity, although its effects on loudness and overall intelligibility are less pronounced compared to LSVT LOUD [6,33]. Similarly, Lam and Tjaden (2016) found that clear speech strategies, particularly the "overenunciate" condition, enhanced vowel space area and reduced articulation rate, but did not significantly improve vocal loudness in conversational speech [17]. HiCommunication, combining speech and cognitive training, demonstrated efficacy in maintaining communication abilities and cognitive performance over time in PD patients [22]. These findings highlight the importance of comprehensive, multi-modal interventions addressing PD's complex speech impairments [5,10].

A key finding of this review is the sustained long-term benefits of LSVT LOUD. Several randomized controlled trials (RCTs) and longitudinal studies reported that improvements in vocal loudness, speech intelligibility, and communicative participation persisted for up to 12 months post-treatment [6,8,18].

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

These outcomes align with earlier findings by Ramig et al. (2008), who documented durable improvements in vocal loudness and intonation following LSVT LOUD [1]. Recent trials also demonstrated that telehealth-based delivery of LSVT LOUD yields comparable long-term outcomes to inperson therapy, making it a viable alternative when accessibility and adherence are concerns [11,9]. Beyond objective speech outcomes, studies have highlighted positive effects on quality of life. LSVT LOUD significantly reduced Voice Handicap Index (VHI) scores by 8 to 9.6 points compared to standard NHS speech-language therapy [8,18]. Additionally, multidisciplinary programs demonstrated improvements in PDQ-39 scores, supporting integrated care approaches [5,10,22,31].

Personalization of Therapy (RQ.2)

Personalized therapy is increasingly recognized as essential in PD care. Customizing treatment based on individual symptom profiles and disease progression has been shown to maximize treatment efficacy [4,6,9] Xu et al. [4], pointed out that by attending to the demands of each patient, customized treatment programs can maximize outcomes. Multi-modal approaches combining SLT with physical and occupational therapy offer a more holistic strategy to address the diverse impairments seen in PD [6] Research is still being done on the neurophysiological alterations brought about by therapies like LSVT LOUD, especially in respect to their association with long-lasting functional gains [7]. Tailored treatment strategies that consider each patient's unique demands and symptoms are crucial for optimizing results due to the diversity of Parkinson's disease and its unpredictable course [9]. The HiCommunication program, which integrates cognitive and speech training, reflects a promising direction in this domain [19].

Differences in Protocols and Outcomes (RQ3)

Significant heterogeneity was observed in intervention protocols, therapy durations, and outcome measures across the studies. While LSVT LOUD typically followed a high-intensity, 4-week protocol, other interventions varied in frequency, duration, and delivery mode. Despite this, robust RCTs confirmed the efficacy of structured interventions over no treatment or sham therapy [10].

Techniques for remote rehabilitation, such SLT based on telehealth, have been investigated as substitutes for in-person therapy. The growing use of telehealth platforms enhances accessibility, particularly for rural or mobility-limited patients. Studies such as Maas et al. (2022) and the PERSPECTIVE trial demonstrate the effectiveness of remote interventions, though more research is needed to standardize protocols and assess long-term outcome. [11,12,14].

Strengths and Limitations: Most included RCTs were of high methodological quality, with 9 out of 12 achieving 8–9 stars on the Newcastle-Ottawa Scale. Robust study designs, including proper randomization and control groups, enhanced the reliability of findings. However, some concerns regarding randomization concealment and missing data were identified using the ROB2 tool. The quasi-experimental study by Lam and Tjaden (2016) had moderate risk of bias due to issues in participant selection and outcome measurement [17]. While the evidence strongly supports LSVT LOUD as a gold-standard intervention, the lack of standardized outcome measures remains a challenge for comparability. Furthermore, most studies focus on short-to-medium term outcomes, with few assessing real-world functional communication over longer durations. These findings align with earlier systematic reviews identifying LSVT LOUD as the most effective intervention for hypokinetic dysarthria in PD. This review further extends evidence by incorporating recent trials on telehealth and hybrid models, addressing accessibility gaps. Standardized outcome measures remain necessary to enhance comparability.

Future Directions: This review highlights key limitations in SLT research for hypokinetic dysarthria in PD. Variability in intervention protocols, therapy durations, and outcome measures complicates comparisons and underscores the need for standardization. Although LSVT LOUD shows sustained

ISSN: **2229-7359** Vol. 11 No. 7, 2025

https://www.theaspd.com/ijes.php

benefits, further studies are necessary to confirm its long-term effects on functional communication in daily life. Moreover, emerging digital technologies and personalized, patient-centered interventions offer promising opportunities to improve adherence and outcomes, highlighting the need for comprehensive longitudinal and cost-effectiveness research.

CONCLUSION

Innovative approaches such as telerehabilitation, multimodal interventions, and technology-enhanced therapies present valuable opportunities to improve the accessibility and personalization of SLT for individuals with Parkinson's disease (PD). Despite these advancements, the current lack of standardization in intervention protocols and outcome measures limits the comparability of findings and underscores the need for consistent, long-term evaluations. Future research should emphasize patient-centered and individualized treatment strategies that address both the motor and non-motor components of communication impairment. Integrating pharmacological therapies with SLT may further optimize functional outcomes. Additionally, expanding the use of telehealth and digital platforms can help reduce access barriers, especially for underserved populations. Ultimately, clinicians should adopt comprehensive, tailored approaches to enhance adherence, therapeutic effectiveness, and overall health of individuals with PD.

REFERENCES

- 1. Ramig LO, Fox C, Sapir S. Speech treatment for Parkinson's disease. Expert review of neurotherapeutics. 2008 Feb 1;8(2):297-309.
- Muñoz-Vigueras N, Prados-Román E, Valenza MC, Granados-Santiago M, Cabrera-Martos I, Rodríguez-Torres J, Torres-Sánchez I. Speech and language therapy treatment on hypokinetic dysarthria in Parkinson disease: Systematic review and meta-analysis. Clin Rehabil. 2021 May;35(5):639-655. doi: 10.1177/0269215520976267. Epub 2020 Nov 24. PMID: 33233932.
- 3. Xu H, Bao Z, Liang D, Li M, Wei M, Ge X, Liu J, Li J. Speech and Language Therapy for Voice Problems in Parkinson's Disease: A Meta-Analysis. J Neuropsychiatry Clin Neurosci. 2020 Fall;32(4):344-351. doi: 10.1176/appi.neuropsych.19020044. Epub 2020 May 6. PMID: 32374650.
- 4. Campbell P, Rooney S, Nicoll A, Brady MC, Smith CH, Deane KHO, Herd CP, Tomlinson CL, Clarke CE, Sackley CM. Speech and language therapy interventions for speech problems in Parkinson's disease. Cochrane Database Syst Rev. 2022 Jun 6;2022(6):CD015009. doi: 10.1002/14651858.CD015009. PMCID: PMC9169534.
- 5. Saluja A, Goyal V, Dhamija RK. Multi-modal rehabilitation therapy in Parkinson's disease and related disorders. Annals of Indian Academy of Neurology. 2023 Jan 1;26(Suppl 1):S15-25.
- 6. Ramig L, Halpern A, Spielman J, Fox C, Freeman K. Speech treatment in Parkinson's disease: Randomized controlled trial (RCT). Movement Disorders. 2018 Nov;33(11):1777-91.
- 7. Levy ES, Moya-Galé G, Chang YH, Freeman K, Forrest K, Brin MF, Ramig LA. The effects of intensive speech treatment on intelligibility in Parkinson's disease: a randomised controlled trial. EClinicalMedicine. 2020 Jul 1;24.
- 8. Sackley CM, Rick C, Brady MC, Woolley R, Burton C, Patel S, Masterson-Algar P, Nicoll A, Smith CH, Jowett S, Ives N, Beaton G, Dickson S, Ottridge R, Sharp L, Nankervis H, Clarke CE; PD COMM collaborative group. Lee Silverman voice treatment versus NHS speech and language therapy versus control for dysarthria in people with Parkinson's disease (PD COMM): pragmatic, UK based, multicentre, three arm, parallel group, unblinded, randomised controlled trial. BMJ. 2024 Jul 10;386:e078341. doi: 10.1136/bmj-2023-078341. PMID: 38986549; PMCID: PMC11232530.
- 9. Theodoros DG, Hill AJ, Russell TG. Clinical and quality of life outcomes of speech treatment for Parkinson's disease delivered to the home via telerehabilitation: A noninferiority randomized controlled trial. American Journal of Speech-Language Pathology. 2016 May;25(2):214-32.
- 10. Ferrazzoli D, Ortelli P, Zivi I, Cian V, Urso E, Ghilardi MF, Maestri R, Frazzitta G. Efficacy of intensive multidisciplinary rehabilitation in Parkinson's disease: a randomised controlled study. Journal of Neurology, Neurosurgery & Psychiatry. 2018 Aug 1;89(8):828-35.
- 11. Maas JJ, de Vries NM, IntHout J, Bloem BR, Kalf JG. Effectiveness of remotely delivered speech therapy in persons with Parkinson's disease-a randomised controlled trial. EClinicalMedicine. 2024 Oct 1;76.

ISSN: **2229-7359** Vol. 11 No. 7, 2025

- 12. Maas JJL, De Vries NM, Bloem BR, Kalf JG. Design of the PERSPECTIVE study: PERsonalized SPEeCh Therapy for actIVE conversation in Parkinson's disease (randomized controlled trial). Trials. 2022 Apr 8;23(1):274. doi: 10.1186/s13063-022-06160-9. PMID: 35395953; PMCID: PMC8990485.
- 13. Trail M, Fox C, Ramig LO, Sapir S, Howard J, Lai EC. Speech treatment for Parkinson's disease. NeuroRehabilitation. 2005 Jan 1;20(3):205-21.
- 14. Herd CP, Tomlinson CL, Deane KH, Brady MC, Smith CH, Sackley CM, Clarke CE, Cochrane Movement Disorders Group. Speech and language therapy versus placebo or no intervention for speech problems in Parkinson's disease. Cochrane Database of Systematic Reviews. 1996 Sep 1;2012(8).
- 15. Behrman A, Cody J, Elandary S, Flom P, Chitnis S. The Effect of SPEAK OUT! and The LOUD Crowd on Dysarthria Due to Parkinson's Disease. Am J Speech Lang Pathol. 2020 Aug 4;29(3):1448-1465. doi: 10.1044/2020_AJSLP-19-00024. Epub 2020 May 18. PMID: 32421347; PMCID: PMC7893519.
- 16. París AP, Saleta HG, de la Cruz Crespo Maraver M, Silvestre E, Freixa MG, Torrellas CP, Pont SA, Nadal MF, Garcia SA, Bartolomé MV, Fernández VL. Blind randomized controlled study of the efficacy of cognitive training in Parkinson's disease. Movement Disorders. 2011 Jun;26(7):1251-8.
- 17. Lam J, Tjaden K. Clear Speech Variants: An Acoustic Study in Parkinson's Disease. J Speech Lang Hear Res. 2016 Aug 1;59(4):631-46. doi: 10.1044/2015_JSLHR-S-15-0216. PMID: 27355431; PMCID: PMC5280060.
- 18. Sackley CM, Rick C, Brady MC, Burton C, Jowett S, Patel S, Woolley R, Masterson-Algar P, Nicoll A, Smith CH, Abdali Z. The effect of two speech and language approaches on speech problems in people with Parkinson's disease: the PD COMM RCT. Health Technology Assessment (Winchester, England). 2024 Oct;28(58):1.
- 19. Sterne JA, Hernán MA, Reeves BC, Savović J, Berkman ND, Viswanathan M, Henry D, Altman DG, Ansari MT, Boutron I, Carpenter JR, Chan AW, Churchill R, Deeks JJ, Hróbjartsson A, Kirkham J, Jüni P, Loke YK, Pigott TD, Ramsay CR, Regidor D, Rothstein HR, Sandhu L, Santaguida PL, Schünemann HJ, Shea B, Shrier I, Tugwell P, Turner L, Valentine JC, Waddington H, Waters E, Wells GA, Whiting PF, Higgins JP. ROBINS-I: a tool for assessing risk of bias in non-randomised studies of interventions. BMJ. 2016 Oct 12;355:i4919. doi: 10.1136/bmj.i4919. PMID: 27733354; PMCID: PMC5062054.
- 20. Wells GA, Shea B, O'Connell D, Peterson J, Welch V, Losos M, Tugwell P. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses. University of Ottawa. 2001. Accessed 10/03/2025 https://www.ohri.ca/programs/clinical_epidemiology/oxford.asp
- 21. Sackley CM, Rick C, Au P, Brady MC, Beaton G, Burton C, Caulfield M, Dickson S, Dowling F, Hughes M, Ives N, Jowett S, Masterson-Algar P, Nicoll A, Patel S, Smith CH, Woolley R, Clarke CE; PD COMM Collaborative Group. A multicentre, randomised controlled trial to compare the clinical and cost-effectiveness of Lee Silverman Voice Treatment versus standard NHS Speech and Language Therapy versus control in Parkinson's disease: a study protocol for a randomised controlled trial. Trials. 2020 May 27;21(1):436. doi: 10.1186/s13063-020-04354-7. PMID: 32460885; PMCID: PMC7251680.
- 22. Brabenec L, Klobusiakova P, Simko P, Kostalova M, Mekyska J, Rektorova I. Non-invasive brain stimulation for speech in Parkinson's disease: A randomized controlled trial. Brain Stimul. 2021 May-Jun;14(3):571-578. doi: 10.1016/j.brs.2021.03.010. Epub 2021 Mar 26. PMID: 33781956.
- 23. Mohseni Z, Saffarian A, Mohamadi R, Abolghasemi J, Habibi SA. Effect of conventional speech therapy combined with music therapy on swallowing in patients with Parkinson's disease (Telerehabilitation): a randomized-controlled trial. Middle East J Rehabil Health Stud. 2023;10(1):e131572.
- 24. Steurer H, Albrecht F, Gustafsson JK, Razi A, Franzén E, Schalling E. Speech and neuroimaging effects following HiCommunication: a randomized controlled group intervention trial in Parkinson's disease. Brain Commun. 2024 Jul 12;6(4):fcae235. doi: 10.1093/braincomms/fcae235. PMID: 39051026; PMCID: PMC11267236.
- 25. Saffarian A, Amiri Shavaki Y, Shahidi GA, Hadavi S, Jafari Z. Lee Silverman voice treatment (LSVT) mitigates voice difficulties in mild Parkinson's disease. Med J Islam Repub Iran. 2019 Feb 13;33:5. doi: 10.34171/mjiri.33.5. PMID: 31086784; PMCID: PMC6504915.
- 26. Constantinescu G, Theodoros D, Russell T, Ward E, Wilson S, Wootton R. Treating disordered speech and voice in Parkinson's disease online: a randomized controlled non-inferiority trial. Int J Lang Commun Disord. 2011 Jan-Feb;46(1):1-16. doi: 10.3109/13682822.2010.484848. PMID: 21281410.
- 27. Kearney E, Shellikeri S, Martino R, Yunusova Y. Augmented visual feedback-aided interventions for motor rehabilitation in Parkinson's disease: a systematic review. Disabil Rehabil. 2019 May;41(9):995-1011. doi: 10.1080/09638288.2017.1419292. Epub 2018 Jan 9. PMID: 29316816.

ISSN: **2229-7359** Vol. 11 No. 7, 2025

- 28. Armstrong MJ, Okun MS. Diagnosis and Treatment of Parkinson Disease: A Review. JAMA. 2020 Feb 11;323(6):548-560. doi: 10.1001/jama.2019.22360. PMID: 32044947.
- 29. Mahler LA, Ramig LO, Fox C. Evidence-based treatment of voice and speech disorders in Parkinson disease. Curr Opin Otolaryngol Head Neck Surg. 2015 Jun;23(3):209-15. doi: 10.1097/MOO.000000000000151. PMID: 25943966.
- 30. Abbruzzese G, Marchese R, Avanzino L, Pelosin E. Rehabilitation for Parkinson's disease: Current outlook and future challenges. Parkinsonism Relat Disord. 2016 Jan;22 Suppl 1:S60-4. doi: 10.1016/j.parkreldis.2015.09.005. Epub 2015 Sep 3. PMID: 26360239.
- 31. Bloem BR, de Vries NM, Ebersbach G. Nonpharmacological treatments for patients with Parkinson's disease. Mov Disord. 2015 Sep 15;30(11):1504-20. doi: 10.1002/mds.26363. Epub 2015 Aug 14. PMID: 26274930.
- 32. Kamioka H, Tsutani K, Yamada M, Park H, Okuizumi H, Tsuruoka K, Honda T, Okada S, Park SJ, Kitayuguchi J, Abe T, Handa S, Oshio T, Mutoh Y. Effectiveness of music therapy: a summary of systematic reviews based on randomized controlled trials of music interventions. Patient Prefer Adherence. 2014 May 16;8:727-54. doi: 10.2147/PPA.S61340. PMID: 24876768; PMCID: PMC4036702.
- 33. Schulz G, Halpern A, Spielman J, Ramig L, Panzer I, Sharpley A, Freeman K. Single Word Intelligibility of Individuals with Parkinson's Disease in Noise: Pre-Specified Secondary Outcome Variables from a Randomized Control Trial (RCT) Comparing Two Intensive Speech Treatments (LSVT LOUD vs. LSVT ARTIC). Brain Sci. 2021 Jun 27;11(7):857. doi: 10.3390/brainsci11070857. PMID: 34199093; PMCID: PMC8301858.