

Associations Between Upper Extremity Function, Activities of Daily Living, and Functional Capacity in Patients with Heart Failure with Reduced Ejection Fraction

Düşük Ejeksiyon Fraksiyonlu Kalp Yetersizliği Hastalarında Üst Ekstremitte Fonksiyonu, Günlük Yaşam Aktiviteleri ve Fonksiyonel Kapasite Arasındaki İlişkiler

ABSTRACT

Objective: Limited information is available regarding the associations between upper extremity function, activities of daily living (ADLs), and functional capacity in patients with heart failure with reduced ejection fraction (HFrEF). This study aimed to investigate the associations between upper extremity function, ADLs, and functional capacity in patients with HFrEF.

Methods: This cross-sectional study included 31 patients with HFrEF. Demographic, anthropometric, and clinical data were recorded. Upper extremity function and ADLs were evaluated using the 6-Minute Pegboard and Ring Test (6PBRT) and the Glittre Activities of Daily Living Test (TGlittre), respectively. The 6-Minute Walk Test (6MWT) was administered to measure functional capacity. Heart rate (HR), systolic blood pressure (SBP), diastolic blood pressure (DBP), peripheral oxygen saturation (SpO₂), dyspnea, and fatigue were assessed at the beginning and end of each test.

Results: The 6PBRT was significantly correlated with TGlittre ($\rho = -0.718, P < 0.001$) and 6-minute walk distance (6MWD) ($r = 0.546, P = 0.001$). A significant correlation was also found between TGlittre and 6MWD ($\rho = -0.810, P < 0.001$). Changes in HR, SBP, and dyspnea were significantly different across the 6PBRT, TGlittre, and 6MWT ($P < 0.05$).

Conclusion: This study indicates that upper extremity function is associated with ADLs and functional capacity in patients with HFrEF. The 6PBRT requires lower cardiopulmonary demand than TGlittre and 6MWT in this patient population.

Keywords: Activities of daily living, functional capacity, heart failure, upper extremity function

ÖZET

Amaç: Düşük ejeksiyon fraksiyonlu kalp yetersizliği (DEFKY) olan hastalarda üst ekstremitte fonksiyonu, günlük yaşam aktiviteleri ve fonksiyonel kapasite arasındaki ilişkiler konusunda sınırlı bilgi vardır. Bu çalışmanın amacı DEFKY'li hastalarda üst ekstremitte fonksiyonu, günlük yaşam aktiviteleri (GYA) ve fonksiyonel kapasite arasındaki ilişkileri araştırmaktır.

Yöntem: Bu kesitsel çalışmaya toplam 31 DEFKY'li hasta katıldı. Demografik, antropometrik ve klinik veriler kaydedildi. Üst ekstremitte fonksiyonu ve GYA sırasıyla 6 Dakika Pegboard ve Ring Testi (6PBRT) ve Glittre ADL Testi (TGlittre) ile değerlendirildi. 6 Dakika Yürüme Testi (6DYT) fonksiyonel kapasiteyi ölçmek için kullanıldı. Her testin başında ve sonunda kalp hızı (KH), sistolik kan basıncı (SKB), diyastolik kan basıncı (DKB), periferik oksijen saturasyonu (SpO₂), dispne ve yorgunluk değerlendirildi.

Bulgular: 6PBRT, TGlittre ($\rho = -0,718, P < 0,001$) ve 6 dakika yürüme mesafesi (6DYM) ($r = 0,546, P = 0,001$) ile anlamlı düzeyde korele idi. TGlittre ile 6DYM arasında anlamlı bir korelasyon bulundu ($\rho = -0,810, P < 0,001$). KH, SKB ve dispnedeki değişiklikler 6PBRT, TGlittre ve 6DYT arasında anlamlı derecede farklıydı ($P < 0,05$).

Sonuç: Çalışmamız DEFKY'li hastalarda üst ekstremitte fonksiyonunun günlük yaşam aktiviteleri ve fonksiyonel kapasite ile ilişkili olduğunu göstermektedir. Bu hasta popülasyonunda 6PBRT, TGlittre ve 6DYT'ye göre daha düşük kardiyopulmoner talep gerektirir.

Anahtar Kelimeler: Günlük yaşam aktiviteleri, fonksiyonel kapasite, kalp yetersizliği, üst ekstremitte fonksiyonu

ORIGINAL ARTICLE KLİNİK ÇALIŞMA

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Heart failure (HF) is a significant public health problem, contributing to high mortality rates.¹ Globally, an estimated 64 million individuals are affected by HF, with its prevalence increasing in many middle- and low-income countries.^{1,2} Despite advancements in medical treatment, the number of years lived with disability due to HF has risen, especially in middle- and low-income countries.¹

HF can impair skeletal muscle metabolic activity in both the upper and lower extremities.³ Additionally, muscle mass in the upper and lower extremities has a prognostic impact on mortality in HF patients.⁴ However, upper extremity function has often been overlooked in this population. Upper extremity function can be assessed using different methods, one of which is the 6-Minute Pegboard and Ring Test (6PBRT), which reflects upper-extremity activities of daily living (ADLs).^{5,6} Upper extremity function as measured by the 6PBRT has been shown to correlate with functional capacity and ADLs in patients with pulmonary hypertension and chronic obstructive pulmonary disease (COPD).^{7,8}

It is likely that a strong association exists between low exercise tolerance and functional disability, as assessed by submaximal and functional tests, in patients with heart failure with reduced ejection fraction (HFrEF). However, limited information is available on the associations between upper extremity function, ADLs, and functional capacity in patients with HFrEF. Understanding these associations would enable healthcare professionals to develop more personalized and targeted disease management strategies. Therefore, this study aimed to investigate the associations between upper extremity function, ADLs, and functional capacity in patients with HFrEF. A secondary aim was to compare the physiological responses to the 6PBRT, Glitter Activities of Daily Living Test (TGlitter), and 6-Minute Walk Test (6MWT) to assess the extent of physiological capacity required to perform each test.

Materials and Methods

This prospective, cross-sectional study was conducted in the Department of Cardiology at Dokuz Eylül University Hospital from April 2023 to December 2023. Patients diagnosed with HFrEF according to international guidelines were enrolled in this study.^{9,10} Additional inclusion criteria were being over 18 years of age, clinically stable as classified by New York Heart Association (NYHA) functional class II-III, and willing to participate. Exclusion

criteria included a history of acute coronary syndrome within the past six months, any orthopedic, neurological, or cognitive condition that could limit test performance, or severe pulmonary disease.

This study received ethical approval from the Dokuz Eylül University Non-Invasive Research Ethical Committee (Approval Number: 2023/03-05, Date: 18.01.2023), and all participants provided written informed consent. The study was conducted in compliance with the Declaration of Helsinki.

Assessments

Functional Class

Functional class was assessed using the NYHA scale, a valid measure for patients with cardiac disease.¹¹ The NYHA functional classification system categorizes patients into classes 1 through 4 based on the impact of cardiac symptoms on daily activities. Higher NYHA functional classes indicate greater disease severity.

Dyspnea Perception

Dyspnea perception was assessed using the modified Medical Research Council (mMRC) scale. The mMRC scale consists of five items, where a score 0 indicates dyspnea only with strenuous exercise, and a score 4 indicates dyspnea even during dressing or undressing.

Pulmonary Function

Pulmonary function was assessed using a spirometer (Minispir® MIR s.r.l., Rome, Italy), following recommended guidelines.¹² Forced expiratory volume in one second (FEV₁), forced vital capacity (FVC), and the FEV₁/FVC ratio were recorded as percentages.

Functional Capacity

Functional capacity was measured using the 6-Minute Walk Test.¹³ The test was conducted following standard guidelines.¹⁴ Participants were instructed to walk a 20-meter straight hallway as quickly as possible over six minutes. After the test began, participants were informed of the remaining time each minute, and standardized encouragement was provided. The total distance covered during the test was recorded as the 6-minute walk distance (6MWD). The predicted percentage of the 6MWD (6MWD%_{predicted}) was calculated using reference equations.¹⁵

Activities of Daily Living (ADLs)

ADLs were assessed with the Glitter Activities of Daily Living Test (TGlitter), an applicable and reproducible tool for patients with HF.^{16,17} The test was conducted following the original protocol for TGlitter in patients with COPD.¹⁸ The TGlitter consists of a 10-meter circuit that includes a two-step staircase with each step measuring 27 cm deep and 17 cm high, along with two shelves positioned at shoulder and waist height for each participant. Participants wore a backpack weighing 5.0 kg for men and 2.5 kg for women throughout the TGlitter. The TGlitter began with participants rising from a seated position, followed by walking, ascending and descending the two steps, and walking again to reach the shelves. Three 1 kg weights placed on the upper shelf were moved one at a time to the lower shelf, then to the floor, back to the lower shelf, and finally returned to the upper shelf. Participants then turned around, walked back across the stairs to their chairs, sat down, and promptly began the next lap by

ABBREVIATIONS

6MWT	6-Minute Walk Test
6PBRT	6-Minute Pegboard and Ring Test
ADLs	Activities of daily living
COPD	Chronic obstructive pulmonary disease
DBP	Diastolic blood pressure
FEV ₁	Forced expiratory volume in one second
FVC	Forced vital capacity
HFrEF	Heart failure with reduced ejection fraction
HR	Heart rate
mMRC	Modified Medical Research Council
NYHA	New York Heart Association
SBP	Systolic blood pressure
SpO ₂	Peripheral oxygen saturation
TGlitter	Glitter Activities of Daily Living Test

standing up again. Participants were instructed to complete five rounds as quickly as possible, and the total duration of the TGlittre was recorded. The percentage of the predicted TGlittre duration ($TGlittre\%_{\text{predicted}}$) was calculated using the reference equation.¹⁹

Upper Extremity Function

Upper extremity function was assessed using the 6PBRT. The test was conducted following the protocol described in the initial study on the 6PBRT in patients with COPD.²⁰ A pegboard was positioned in front of each participant within arm's reach while they sat upright in a chair. The pegboard included two pegs at shoulder level and two additional pegs positioned 20 cm above shoulder level. Ten lightweight wooden rings were hung individually on each of the bottom two pegs. Participants were instructed to move one ring from each bottom peg to the top peg, using both hands simultaneously. Once all rings were transferred to the top pegs, participants moved the rings back to the bottom pegs. This task was repeated as quickly as possible for six minutes, with standardized verbal encouragement provided each minute. The total number of rings moved during the 6PBRT was recorded, and the percentage of the predicted 6PBRT value ($6PBRT\%_{\text{predicted}}$) was calculated using the reference equation.²¹

Procedure

The study procedure consisted of a single visit. First, demographic and anthropometric data were collected, and left ventricular ejection fraction and medication information were retrieved from the latest patient records. Next, functional class, dyspnea perception, and pulmonary function were assessed. Finally, the 6PBRT, TGlittre, and 6MWD were conducted in random order, determined by a computerized random number generator (www.random.org). Heart rate (HR), systolic blood pressure (SBP), diastolic blood pressure (DBP), peripheral oxygen saturation (SpO_2), dyspnea, and fatigue were measured at the beginning and end of each test. The next test was initiated only after these physiological parameters returned to their baseline levels evaluated at the start of the test.

Statistical Analysis

The study sample size was calculated using G*Power software (version 3.1.9.2, Düsseldorf, Germany), based on a previous study investigating the association between 6MWD and TGlittre in patients with HF.²² Considering an effect size of 0.484, a power of 0.80, and an alpha error probability of 0.05, the minimum required sample size was determined to be 28 participants.

Statistical analyses were conducted using IBM SPSS software (version 26, IBM, Armonk, NY, USA). The normality of the data distribution was assessed with the Shapiro-Wilk test and histograms. Descriptive and categorical data were expressed as mean \pm standard deviation (SD), median (interquartile range), or number (percentage), as appropriate. The correlation between the 6PBRT and 6MWD was analyzed using Pearson's product-moment correlation, as parametric conditions were met. Correlations among other variables were analyzed using Spearman's rank-order correlation, as nonparametric conditions were met. Correlation strength was classified as weak (0.20–0.39), moderate (0.40–0.59), strong (0.60–0.79), or very strong (0.80–1.0).²³ Comparisons of physiological values obtained before and after each test were analyzed using the Wilcoxon

test or paired sample t-test, as appropriate. Differences among physiological responses to the tests were analyzed using repeated measures Analysis of Variance (ANOVA) or the Friedman test, as appropriate. Post-hoc analyses were conducted with Bonferroni correction. Statistical significance was set at $P < 0.05$.

Results

A total of 31 patients with HFREF participated in this study. Table 1 presents the demographic, anthropometric, and clinical characteristics of the participants. The mean age was 62.13 ± 9.38 years. Most participants were men (87.1%), were classified as overweight (48.4%), and fell into NYHA functional class II

Table 1. Demographic and Clinical Features

Variables	Total Participants (n = 31)
Age, years	62.13 \pm 9.38
Gender, n (%)	
Women	4 (12.9)
Men	27 (87.1)
Body mass index, kg/m ²	27.97 (24.51–30.40)
LVEF, %	30.00 (20.00–35.00)
NYHA functional class, n (%)	
II	18 (58.1)
III	13 (41.9)
Medication, n (%)	
Beta-blockers	27 (87.1)
Ivabradine	6 (19.4)
ACEI/ARBs/ARNI	23 (74.2)
MRAs	15 (48.4)
Digoxin	4 (12.9)
Diuretics	22 (71.0)
mMRC dyspnea scale, n (%)	
I	11 (35.5)
II	14 (45.2)
III	5 (16.1)
IV	1 (3.2)
Pulmonary function	
FEV ₁ , %	85.00 (75.00–94.00)
FVC, %	86.00 (78.00–92.00)
FEV ₁ /FVC, %	79.40 (74.50–83.70)
FEF _{25–75} , %	70.58 \pm 27.37
PEF, %	75.74 \pm 28.62
Functional tests	
6PBRT, rings	133.48 \pm 33.51
6PBRT, % predicted	64.22 \pm 14.04
TGlittre, min	3.20 (2.53–4.15)
TGlittre, % predicted	108.39 (88.27–137.19)
6MWD, m	435.17 \pm 97.03
6MWD, % predicted	85.08 (69.37–92.92)

Data are expressed as mean \pm standard deviation (SD) or median (IQR) unless otherwise indicated. 6MWD, Six-Minute Walk Distance; 6PBRT, 6-Minute Pegboard and Ring Test; ACEI, Angiotensin-Converting Enzyme Inhibitor; ARBs, Angiotensin Receptor Blockers; ARNI, Angiotensin Receptor-Nephrilysin Inhibitor; FEF_{25–75}, Forced Expiratory Flow at 25–75%; FEV₁, Forced Expiratory Volume in One Second; FVC, Forced Vital Capacity; LVEF, Left Ventricular Ejection Fraction; mMRC, Modified Medical Research Council; MRAs, Mineralocorticoid Receptor Antagonists; NYHA, New York Heart Association; PEF, Peak Expiratory Flow; TGlittre, Glittre ADL test.

(58.1%). The majority of participants had a dyspnea perception score of mMRC II (45.2%). The mean total number of rings moved in the 6PBRT was 133.48 ± 33.51 , with a mean percentage of the predicted 6PBRT value at 64.22 ± 14.04 . The median duration for participants to complete the TGlitter was 3.20 (2.53-4.15), and the median percentage of the predicted TGlitter value was 108.39

(88.27-137.19). The mean 6MWD for participants was 435.17 ± 97.03 , with a median percentage of the predicted 6MWD value at 85.08 (69.37-92.92).

Figure 1 presents the correlations between the 6PBRT, TGlitter, and 6MWD. The 6PBRT was strongly correlated with TGlitter

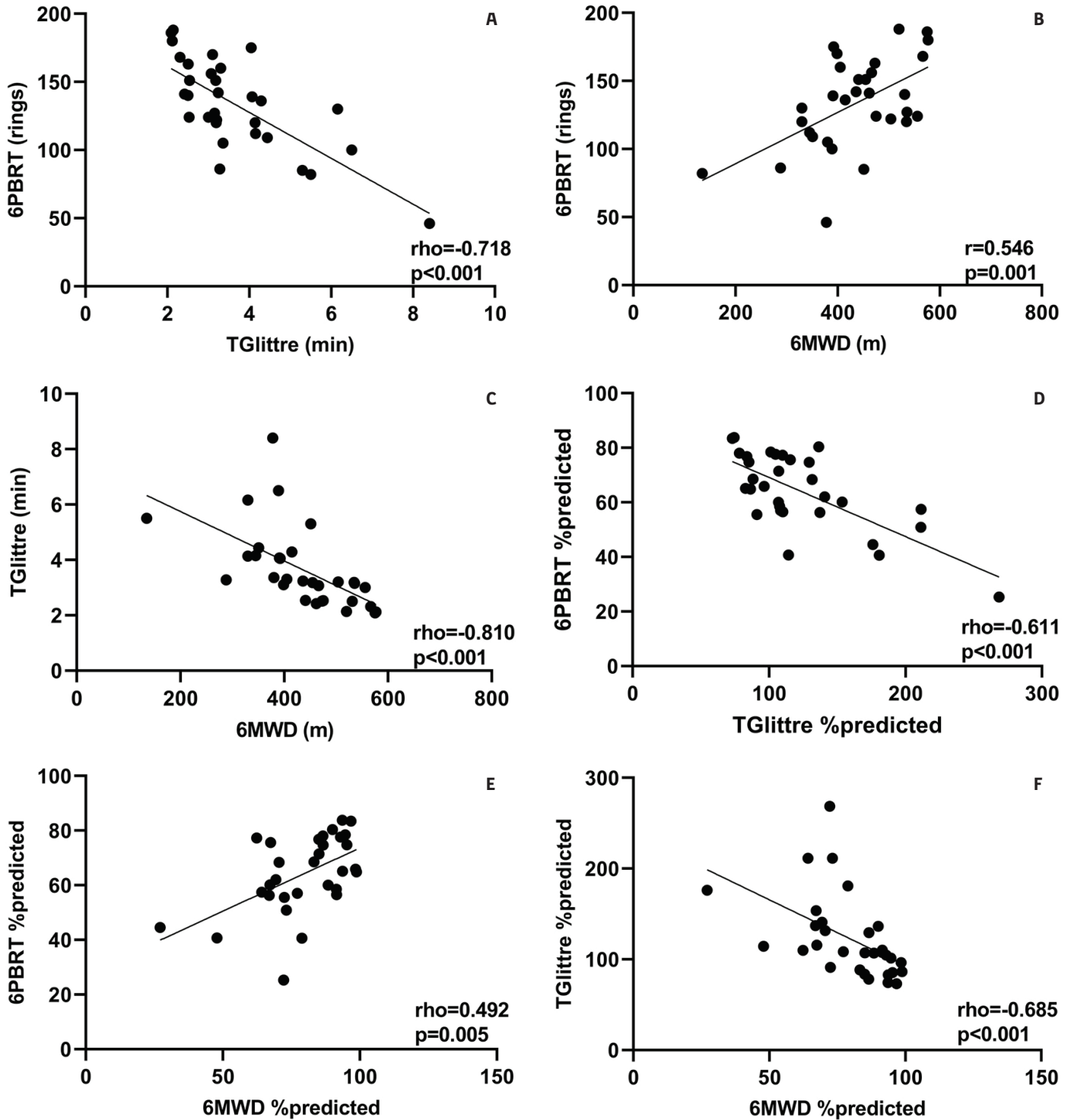


Figure 1. Correlations between: A) 6PBRT and TGlitter, B) 6PBRT and 6MWD, C) TGlitter and 6MWD, D) 6PBRT%_{predicted} and TGlitter%_{predicted}, E) 6PBRT%_{predicted} and 6MWD%_{predicted}, F) TGlitter%_{predicted} and 6MWD%_{predicted}. 6PBRT, 6-Minute Pegboard and Ring test; TGlitter, Glitter Activities of Daily Living Test; 6MWD, 6-Minute Walk Distance.

Table 2. Comparison of Physiological Values Obtained Pre- and Post-Test for 6PBRT, TGlitter, and 6MWT

	6PBRT (n = 31)			TGlitter (n = 31)			6MWT (n = 31)		
	Pre-test	Post-test	t/Z	Pre-test	Post-test	t/Z	Pre-test	Post-test	t/Z
HR (beats/min)	71.55 ± 10.99	74.71 ± 12.46	-3.408	70.90 ± 12.50	89.90 ± 14.29	-9.578	70.87 ± 11.56	86.32 ± 14.38	-8.464
SBP (mmHg)	120.13 ± 14.65	127.71 ± 17.14	-4.298	118.03 ± 15.80	136.87 ± 20.70	-8.316	119.52 ± 3.41	131.32 ± 3.86	-5.715
DBP (mmHg)	76.13 ± 8.66	78.71 ± 9.27	-2.515	75.94 ± 9.76	79.97 ± 10.97	-2.890	75.61 ± 10.53	78.48 ± 2.05	-2.048
SpO ₂ (%)	96.00 (96.00-97.00)	97.00 (96.00-98.00)	-1.762	97.00 (96.00-98.00)	98.00 (96.00-99.00)	-0.579	96.00 (95.00-97.00)	98.00 (96.00-98.00)	-2.895
Dyspnea (0-10)	0.00 (0.00-0.00)	0.00 (0.00-0.00)	-2.032	0.00 (0.00-0.00)	1.00 (0.00-4.00)	-3.838	0.00 (0.00-0.00)	1.00 (0.00-3.00)	-3.749
Arm Fatigue (0-10)	0.00 (0.00-0.00)	3.00 (2.00-5.00)	-4.640	0.00 (0.00-0.00)	0.00 (0.00-3.00)	-3.307	NA	NA	NA
Leg Fatigue (0-10)	NA	NA	NA	0.00 (0.00-0.00)	2.00 (0.00-3.00)	-4.219	0.00 (0.00-0.00)	0.00 (0.00-4.00)	-3.736
General Fatigue (0-10)	0.00 (0.00-0.00)	0.00 (0.00-3.00)	-3.428	0.00 (0.00-0.00)	2.00 (0.00-3.00)	-3.943	0.00 (0.00-0.00)	0.00 (0.00-4.00)	-3.528

Data are expressed as mean ± standard deviation (SD) or median (interquartile range). 6MWT, 6-Minute Walk Test; 6PBRT, 6-Minute Pegboard and Ring Test; DBP, Diastolic Blood Pressure; HR, Heart Rate; NA, Not Applicable; SBP, Systolic Blood Pressure; SpO₂, Peripheral Oxygen Saturation; TGlitter, Glitter ADL test.
* Paired sample t-test (t), ** Wilcoxon test (Z), P < 0.05.

(rho = -0.718, P < 0.001) and moderately correlated with 6MWD (r = 0.546, P = 0.001). A very strong correlation was observed between TGlitter and 6MWD (rho = -0.810, P < 0.001). Additionally, the 6PBRT%_{predicted} was strongly correlated with the TGlitter%_{predicted} (rho = -0.611, P < 0.001) and moderately correlated with 6MWD%_{predicted} (rho = 0.492, P = 0.005). A strong correlation was also found between TGlitter%_{predicted} and 6MWD%_{predicted} (rho = -0.685, P < 0.001).

Table 2 presents the comparisons of physiological values obtained before and after the 6PBRT, TGlitter, and 6MWT. Pre-test values of HR, SBP, DBP, SpO₂, dyspnea, and fatigue were similar across the 6PBRT, TGlitter, and 6MWT (P > 0.05). Significant differences were observed between pre- and post-test measurements for HR, SBP, DBP, dyspnea, and general fatigue across all three tests (P < 0.05). SpO₂ showed a significant difference between pre- and post-6MWT (P < 0.05) but remained similar between pre- and post-6PBRT and TGlitter (P > 0.05). Significant differences were found in arm fatigue between the pre- and post-6PBRT and TGlitter (P < 0.05) and in leg fatigue between the pre- and post-TGlitter and 6MWT (P < 0.05).

Table 3 presents the comparisons of physiological changes among the 6PBRT, TGlitter, and 6MWT. The percentage of maximal heart rate (HR_{max%}) differed significantly among the tests. HR_{max%} was lower in the 6PBRT than in both the TGlitter (P < 0.001) and 6MWT (P < 0.001). However, HR_{max%} was similar between the TGlitter and 6MWT (P = 0.347). Significant differences were observed in changes in HR, SBP, SpO₂, and dyspnea among the tests (P < 0.05). Post hoc analysis revealed that the change in HR was lower in the 6PBRT compared to both the TGlitter (P < 0.001) and the 6MWT (P < 0.001). HR changes were similar between the TGlitter and 6MWT (P = 0.344). The change in SBP was higher in the TGlitter than in the 6PBRT (P < 0.001) and 6MWT (P = 0.027), while SBP changes were similar between the 6PBRT and 6MWT (P = 0.264). The significant difference in SpO₂ changes across the tests disappeared after post hoc analysis. The change in dyspnea was lower in the 6PBRT than in the TGlitter (P = 0.001) and 6MWT (P = 0.047). The change in dyspnea was similar between the TGlitter and 6MWT (P = 0.612). No significant differences were found in changes in DBP and general fatigue among the 6PBRT, TGlitter, and 6MWT (P > 0.05).

Discussion

The main finding of our study is that there were statistically significant associations between upper extremity function, ADLs, and functional capacity in patients with HFREF. Additionally, the 6PBRT, TGlitter, and 6MWT elicited significant acute changes in physiological values. The percentage of maximum heart rate achieved was lowest in the 6PBRT. Changes in HR and dyspnea were also lowest in the 6PBRT, while the change in SBP was highest in the TGlitter. Changes in DBP and general fatigue were similar across the tests.

Takeda et al.⁶ found a correlation between upper extremity function and upper extremity ADLs assessed by activity counts in patients with COPD. Additionally, Calik-Kutukcu et al.²⁴ showed a correlation between upper extremity function and upper extremity ADLs assessed by the ADL simulation test in patients with moderate to severe COPD. This study similarly

Table 3. Comparison of Physiological Changes Between the 6PBRT, TGlitter, and 6MWT

	6PBRT (n = 31) Mean ± SD or Median (IQR)	TGlitter (n = 31) Mean ± SD or Median (IQR)	6MWT (n = 31) Mean ± SD or Median (IQR)	P
HR _{max%}	45.43 ± 7.42 ^a	54.63 ± 8.28	52.46 ± 8.31 ^c	<0.001*
ΔHR (beats/min)	3.16 ± 5.17 ^a	19.00 ± 11.05	15.45 ± 10.17 ^c	<0.001*
ΔSBP (mmHg)	7.58 ± 9.82 ^a	18.84 ± 12.61 ^b	11.81 ± 11.50	<0.001*
ΔDBP (mmHg)	2.58 ± 5.71	4.03 ± 7.77	2.87 ± 7.81	0.663*
ΔSpO ₂ (%)	0.00 (0.00–1.00)	0.00 (–1.00–1.00)	1.00 (0.00–2.00)	0.015**
ΔDyspnea (0–10)	0.00 (0.00–0.00) ^a	1.00 (0.00–4.00)	1.00 (0.00–3.00) ^c	<0.001**
ΔGeneral Fatigue (0–10)	0.00 (0.00–3.00)	2.00 (0.00–3.00)	1.00 (0.00–3.00)	0.446**

Data are expressed as mean ± standard deviation (SD) or median (interquartile range). 6MWT, 6-Minute Walk Test; 6PBRT, 6-Minute Pegboard and Ring Test; DBP, Diastolic Blood Pressure; HR, Heart Rate; SBP, Systolic Blood Pressure; SpO₂, Peripheral Oxygen Saturation; TGlitter, Glitter ADL test. D, change from pre-test to post-test values.
*Repeated measures Analysis of Variance (ANOVA), **Friedman test. ^aSignificant difference between 6PBRT and TGlitter; ^bSignificant difference between TGlitter and 6MWT; ^cSignificant difference between 6PBRT and 6MWT. P < 0.05.

demonstrated that upper extremity function was correlated with ADLs measured by the TGlitter in patients with HFrEF. Collectively, this evidence suggests that higher upper extremity function is associated with better ADLs in patients with HFrEF.

Ozsoy et al.⁷ reported that functional capacity assessed by the 6MWT was one of the determinants of upper extremity exercise capacity assessed by the 6PBRT in patients with COPD. This similarly study found that upper extremity function correlated with functional capacity assessed by the 6MWT, indicating that upper extremity function tends to impact functional capacity in patients with HFrEF.

Most patients with HF have report difficulty performing one or more ADLs, with these difficulties often worsening over time.²⁵ It has also been shown that patients with HF tend to reduce the intensity of their effort and increase the time required to complete ADLs when they can self-adjust exercise intensity.²⁶ Since patients with HF experience difficulties in ADLs that require both upper and lower extremity function—such as carrying objects, doing housework and heavy labor, and performing personal care—the functionality of the upper extremities is crucial for maintaining ADLs in this population.²⁷ Paneroni et al.²⁸ demonstrated that short-term cardiac rehabilitation improved the performance of routine ADLs, significantly decreasing the cumulative time required to complete these activities in elderly patients with HF. Additionally, Nyquist-Battie et al.²⁹ found that upper extremity exercise training increased cardiopulmonary exercise test duration in patients with HF. Considering these findings, the relationships between upper extremity function, ADLs, and functional capacity observed in patients with HFrEF in this study suggest that upper extremity training should be incorporated into cardiac rehabilitation programs.

HR, SBP, DBP, dyspnea, and fatigue significantly increased after the 6PBRT, the TGlitter, and the 6MWT, indicating that all three tests create physiological load. The greater systolic blood pressure response observed in the TGlitter compared to the 6PBRT and 6MWT may be partly due to the involvement of more muscle mass, as both arms and legs are used.³⁰ However, the lower physiological response observed in the 6PBRT compared

to the TGlitter and the 6MWT suggests that the 6PBRT can be used to assess physical function with lower physiological demand in patients with HFrEF. Further studies assessing physiological demand using a portable gas analyzer are needed to support these findings.

This study has several limitations. First, due to the cross-sectional design, causal relationships between these associations cannot be established. Second, although the post-hoc power analysis indicated adequate power and the tests were performed randomly, studies with larger sample sizes that control for confounders such as age, gender, body mass index, medication, and physical activity level would strengthen these results by reducing potential bias. Lastly, since this study included only patients with HFrEF, the findings may not be generalizable to all phenotypes of HF. Future studies with larger sample sizes are required to investigate the associations between upper extremity function, ADLs, and functional capacity in patients with HF with mildly reduced and preserved ejection fraction.

In a clinical context, the moderate to strong correlations between upper extremity function, ADLs, and functional capacity in patients with HFrEF suggest that changes in upper extremity function are closely linked to a patient's ability to perform daily tasks and maintain physical capacity. Identifying declines in upper extremity function may serve as an early indicator of reduced ADLs and functional capacity, potentially allowing for proactive measures to prevent further deterioration of physical function. Additionally, incorporating upper extremity exercises into cardiac rehabilitation programs could contribute to improvements in ADLs and functional capacity in patients with HFrEF. Furthermore, the fact that the 6PBRT requires lower physiological demand suggests that clinicians and researchers can use it safely to predict ADL capability and functional capacity in patients with HFrEF, particularly in those with more severe disease.

Conclusion

This study demonstrates that upper extremity function is associated with ADLs and functional capacity in patients with

HFREF. The 6PBRT requires lower cardiopulmonary demand than the TGlittre and the 6MWT in this patient population.

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Informed Consent: All participants provided written informed consent.

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