

# Effects of Uni- and Bi-Directional Interaction During Dyadic Ankle and Wrist Tracking

Matthew R. Short<sup>1</sup>, Daniel Ludvig<sup>1</sup>, Francesco Di Tommaso<sup>1</sup>, Lorenzo Vianello<sup>1</sup>,  
Emek Bariş Küçütabak<sup>1</sup>, *Student Member, IEEE*, Eric J. Perreault<sup>1</sup>, *Member, IEEE*,  
Levi Hargrove, *Member, IEEE*, Kevin Lynch<sup>1</sup>, *Fellow, IEEE*, Etienne Burdet<sup>2</sup>,  
and Jose L. Pons<sup>1</sup>, *Senior Member, IEEE*

**Abstract**—Haptic human-robot-human interaction allows users to feel and respond to one another's forces while interfacing with separate robotic devices, providing customizable infrastructure for studying physical interaction during motor tasks (e.g., physical rehabilitation). For upper- and lower-limb tracking tasks, previous work has shown that virtual interactions with

a partner can improve motor performance depending on the skill level of each partner. However, whether the mechanism explaining these improvements is identical in the upper and lower limbs is an open question. In this work, we investigate the effects of haptic interaction between healthy individuals during a trajectory tracking task involving single-joint movements at the wrist and ankle. We compare tracking performance and muscle activation during haptic conditions where pairs of participants were uni- and bidirectionally connected to investigate the contribution of real-time responses from a partner during the interaction. Findings showed similar improvements in tracking performance during bidirectional interaction for both the wrist and ankle. This was observed despite distinct strategies in muscle co-contraction between joints, as co-contraction was dependent on partner ability for the wrist but not the ankle. For each joint, bidirectional and unidirectional interaction resulted in similar improvements for the worse partner in the dyad. For the better partner, bidirectional interaction resulted in greater improvements than unidirectional interaction. While these results suggest that unidirectional interaction is sufficient for error correction of less skilled individuals during simple motor tasks, they also highlight the mutual benefits of bidirectional interaction which are consistent across the upper and lower limbs.

**Index Terms**—Human-human physical interaction, tracking performance, upper and lower limb.

## I. INTRODUCTION

IN DAILY life, humans interact through physical touch to assist and learn from one another. During physical rehabilitation, for instance, a therapist may interact with a patient by applying corrective forces during dynamic movements to stimulate (re)learning of motor behaviors. However, systematically characterizing these interactions is difficult due to the challenge of measuring contact forces between two individuals. To this end, robotic systems can be leveraged to study various aspects of human-human physical interaction. This is typically accomplished by rendering virtual haptic connections (e.g., spring-damper) between two robotic interfaces [1], [2], [3], [4], [5], [6], [7], [8], [9], allowing users to feel and respond to one another's forces during motor tasks. Such systems have been used to quantify the effects of collaborative training while haptically interacting with a partner, in terms of task

Received 10 December 2024; revised 28 April 2025; accepted 23 May 2025. Date of publication 27 May 2025; date of current version 2 June 2025. This work was supported in part by the National Science Foundation/National Robotics Initiative under Grant 2024488 and in part by Northwestern University. (Corresponding author: Matthew R. Short.)

This work involved human subjects or animals in its research. Approval of all ethical and experimental procedures and protocols was granted by Northwestern University under Application No. STU00212684.

Matthew R. Short and Daniel Ludvig are with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, and also with the Department of Biomedical Engineering, Northwestern University, Evanston, IL 60208 USA (e-mail: mshort@u.northwestern.edu).

Francesco Di Tommaso is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, also with the Department of Physical Medicine and Rehabilitation, Northwestern University, Chicago, IL 60611 USA, and also with the Departmental Faculty of Engineering, Università Campus Bio-Medico di Roma, 00128 Rome, Italy.

Lorenzo Vianello is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, and also with the Department of Physical Medicine and Rehabilitation, Northwestern University, Chicago, IL 60611 USA.

Emek Bariş Küçütabak is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, also with the Department of Mechanical Engineering, Northwestern University, Evanston, IL 60208 USA, and also with the Center for Robotics and Biosystems, Evanston, IL 60208 USA.

Eric J. Perreault is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, also with the Department of Biomedical Engineering, Northwestern University, Evanston, IL 60208 USA, also with the Center for Robotics and Biosystems, Evanston, IL 60208 USA, and also with the Department of Physical Medicine and Rehabilitation, Northwestern University, Chicago, IL 60611 USA.

Levi Hargrove is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, also with the Center for Robotics and Biosystems, Evanston, IL 60208 USA, and also with the Department of Physical Medicine and Rehabilitation, Northwestern University, Chicago, IL 60611 USA.

Kevin Lynch is with the Department of Mechanical Engineering, Northwestern University, Evanston, IL 60208 USA, and also with the Center for Robotics and Biosystems, Evanston, IL 60208 USA.

Etienne Burdet is with the Department of Bioengineering, Imperial College of Science, Technology and Medicine, SW7 2AZ London, U.K.

Jose L. Pons is with the Shirley Ryan AbilityLab, Chicago, IL 60611 USA, also with the Department of Biomedical Engineering and the Department of Mechanical Engineering, Northwestern University, Evanston, IL 60208 USA, also with the Center for Robotics and Biosystems, Evanston, IL 60208 USA, and also with the Department of Physical Medicine and Rehabilitation, Northwestern University, Chicago, IL 60611 USA.

Digital Object Identifier 10.1109/TNSRE.2025.3573956

performance and learning as well as the muscle activation strategy of each partner.

Previous studies, involving both upper- [1], [3], [5], [6] and lower-limb [8], [9] systems, have shown that pairs of healthy individuals (i.e., dyads) perform dynamic tracking tasks better while connected compared to tracking alone. These improvements depend on the ability (i.e., relative skill level) of each partner, as well as the stiffness of the virtual connection [3], [9]. Furthermore, better partners in each dyad exert greater effort, measured as an increase in muscle activation or co-contraction, to compensate for the inferior ability of the worse partner during the interaction [5]. However, this can be addressed by implementing an asymmetric connection, with a higher stiffness for the worse partner, to improve tracking performance without additional effort exerted by either partner [6]. Particularly relevant to physical rehabilitation, a few upper-limb studies have found that individuals can learn tracking tasks more effectively after haptic interaction with a partner than after training alone [1], [7] or with conventional robotic guidance toward a reference trajectory [4].

Though these benefits of human-human physical interaction have been confirmed with various robotic interfaces, the mechanism of these improvements is disputed. Previous work on upper-limb dyadic behaviors has proposed that individuals mutually adapt their movement strategy in response to the haptic feedback received from their partner [2], [3], [10]. When tracking a randomly moving target, individuals are suggested to improve their tracking performance as they infer the target's movement from the information perceived while interacting with a partner, which they combine with their own information in a stochastically optimal way. However, our work in the lower limb suggests a mechanism for these improvements based solely on the interaction mechanics [8], [9]. In these works, we simulated dyad trials by modeling the haptic interaction as three springs in series, where each participant's simulated angle was influenced by the physiological stiffness of their own ankle, the stiffness of the virtual spring, and the physiological stiffness of their partner's ankle. Using trajectories recorded during unconnected trials to represent each partner's planned trajectory, simulated trajectories are effectively a weighted average of two partially correlated signals (i.e., two users following a common trajectory). A key difference in these proposed models is the consideration of the planned trajectory. In the mutual planning model [2], [3], [10], the planned movements of each partner are estimated considering the haptic information received during the interaction. In the mechanics-based model [8], [9], planned movements are assumed to be the same with or without the interaction.

Due to differences in robotic infrastructure and task constraints across the aforementioned dyadic studies, it is unclear whether a single mechanism can explain these improvements, or whether the two proposed mechanisms, mutual planning [3] and interaction mechanics [9], are specific to the upper and lower limbs, respectively. If the basis for these mechanisms is mechanical and unrelated to mutual adaptations in planning, then changes in the movement strategy of each

partner should have a limited effect on the resulting behavior. In this case, one would expect similar tracking improvements whether two individuals are connected bidirectionally (i.e., two-way spring-damper) or unidirectionally (i.e., one-way spring-damper). Unidirectional interaction has been studied less extensively in dyads, but is similar to common approaches in robotic training where guidance is provided towards a reference trajectory [1], [4], [11]; in a dyadic case, partners would be unable to share information of their motor plan as only one individual receives haptic feedback. Conversely, if mutual changes in planning contribute significantly to performance improvements, we would expect individuals to perform worse during unidirectional interaction, compared to bidirectional.

In the context of physical rehabilitation, comparing these approaches is important as it can inform the design of partnered training for individuals with sensorimotor impairments (e.g., stroke) in future studies. Trajectory tracking tasks can be useful visuomotor exercises to assess and train individuals post-stroke [12], [13], using robotic devices to isolate the impaired joint(s) of interest. Given the learning benefits observed during tracking tasks in healthy individuals, it may be similarly advantageous to haptically couple an individual with sensorimotor impairments to a healthy individual (e.g., a physical therapist) during training. Waters et al. presented a preliminary investigation on task performance and motor learning during a 1-DoF upper-limb tracking task where an individual post-stroke was haptically paired with a healthy individual, facilitated via bidirectional interaction between two robotic manipulators [14]. While the results from this work are promising, it still remains an open question whether real-time interaction with a partner (i.e., bidirectional) is necessary for improvements reported in previous dyadic studies. Observing identical changes in task performance between uni- and bidirectional interaction could simplify setups in future studies with patient populations, for both upper- and lower-limb training paradigms. For instance, a recorded database of human trajectories could be used as the "reference" for assistance, obviating the need for a second user.

To address these questions in the upper and lower limb, we present results comparing dyadic behaviors during a 1-DoF trajectory tracking task at the wrist (i.e., flexion and extension) and ankle (i.e., plantarflexion and dorsiflexion) in healthy individuals. Despite differences in the functional roles of each joint, previous works have shown that the wrist and ankle exhibit similar velocity profiles when performing single-joint "pointing" tasks [15], [16], suggesting an invariant strategy employed by the central nervous system to minimize end-effector error across these joints. Therefore, we expect connected partners to improve similarly during wrist and ankle tasks according to their partner's ability, with the better partner increasing their muscle activation to compensate for the worse partner's performance. In addition, we expect that tracking improvements at the wrist and ankle can be explained by the mechanics of the interaction. This means that partners should improve similarly whether they are connected uni- or bidirectionally, based on our previous simulation work during 1-DoF tracking [8], [9] and the limited effects of haptic assistance

during spatial tasks with reduced complexity [13]. Our analysis focuses on changes in tracking error to assess improvements in performance as well as muscle co-contraction to suggest changes in human mechanics as a result of the connection. For the haptic conditions, we compare these measures between bidirectional interactions where individuals are connected to their partner in real-time and unidirectional interactions where individuals are connected to a recording of their partner's trajectory.

## II. METHODS

### A. Participants

We recruited 26 healthy individuals (12 females and 14 males;  $26.5 \pm 3.7$  years) to participate in this study and paired them into sex- and age-matched dyads, separated by a maximum of 5 years. Participants gave informed consent for their participation. The study protocol (registered on clinicaltrials.gov as NCT04578665) was conducted in accordance with the Declaration of Helsinki and approved by the institutional review board of Northwestern University (STU00212684). All participants were determined to be right-handed, assessed by the preferred writing hand. All but two participants were determined to be right-footed, assessed by the preferred foot when kicking a ball.

### B. Experimental Design

Pairs of participants used either their dominant wrist or ankle to perform 1-DoF movements while strapped into two commercially available robots (M1, Fourier Intelligence, Singapore; Fig. 1A). These robots are designed for 1-DoF exercises (e.g., flexion and extension) and are equipped with a sensor to measure the interaction torque between the user and robot. In our previous work [17], a custom interaction torque controller was developed to allow transparent motion (i.e., near zero interaction torque) for each robot and to render virtual haptic interactions between users interfacing with different robots. The desired interaction torque between users  $i$  and  $j$  was calculated as

$$\tau_i^{\text{int}}(t) = K_v[\theta_j(t) - \theta_i(t)] + C_v[\dot{\theta}_j(t) - \dot{\theta}_i(t)] = -\tau_j^{\text{int}}(t), \quad (1)$$

where  $\tau^{\text{int}}(t)$  is the interaction torque applied to either user,  $\theta(t)$  is the angular position measured by each robot,  $\dot{\theta}(t)$  is the angular velocity,  $K_v$  is the virtual stiffness that is applied between the angles of the two users, and  $C_v$  is the virtual damping that is applied between the velocities. Motor torque commands and sensor measurements including joint angle, velocity, and interaction torque data were updated at 333 Hz.

To allow practical comparisons between the wrist and ankle, we made a few modifications to the robot hardware and controllers for each joint. Specifically, each robot was equipped with either a pedal for interfacing with the foot, or a manufactured handle for interfacing with the hand with the fingers splayed. The position of each participant's foot/hand was adjusted such that their ankle/wrist joint was aligned with the robot's center of rotation for flexion and extension movements. The interaction torque controller for the wrist

and ankle configured robots was identical; manual tuning of parameters in the feedback control loop [17] was performed to achieve similar transparent tracking behaviors across joints. These parameters were identified prior to study recruitment, therefore the same set of robot parameters was used across all participants during experimentation.

The experiment was divided into two phases, one for the wrist and the other for the ankle. Each phase began with an electromyography (EMG) calibration procedure followed by sets of tracking trials. The order of the wrist and ankle phases was randomized for each dyad. For the ankle experimentation, participants were seated with their knee slightly flexed and restricted to using dorsi- and plantarflexion; EMG sensors (Bagnoli, Delsys Inc., USA) were placed on the tibialis anterior (TA), gastrocnemius medialis (GM), gastrocnemius lateralis (GL) and soleus (SOL). For the wrist, participants were seated with their forearm supported and restricted to flexion and extension with the wrist in  $90^\circ$  of pronation; EMG sensors were placed on the extensor carpi radialis longus (ECRL) and flexor carpi radialis (FCR). A ground electrode, for both the wrist and ankle measurements, was placed on the non-dominant elbow of each participant. EMG data were collected at 333 Hz and synchronized with data from the robots using a data acquisition board (USB-6218, National Instruments, USA) and a custom Python script. EMG data from the isometric calibration and tracking trials were high-pass filtered at 20 Hz (Butterworth, second order), rectified and low-pass filtered at 5 Hz (Butterworth, second order) to obtain the linear envelope of each muscle's activation.

### C. Trajectory Tracking Trials

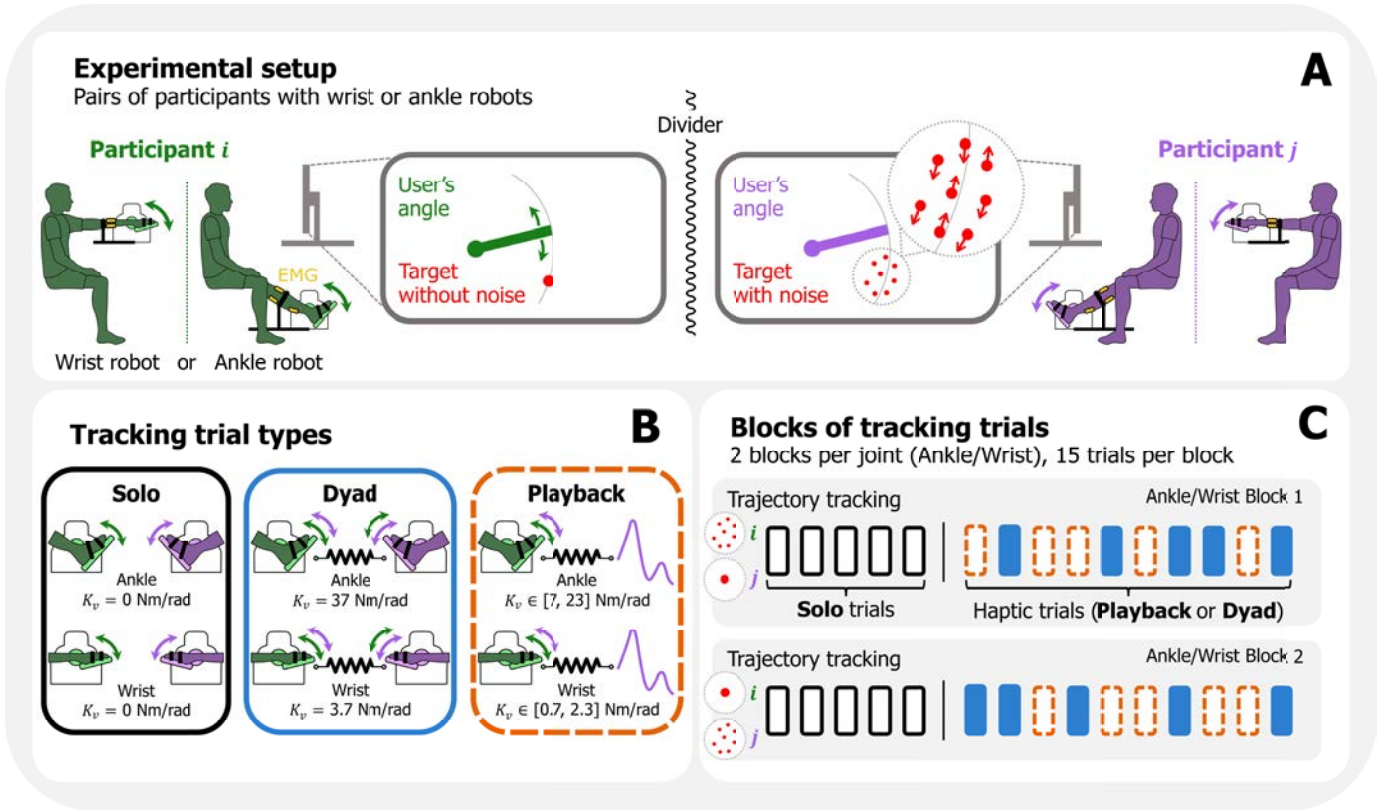
To study the effects of physical interaction during voluntary movement, we designed a continuous, dynamic tracking task involving 1-DoF. During tracking trials, dyads tried to match either their wrist or ankle angle,  $\theta(t)$ , to a visually-displayed, sinusoidal trajectory while their robots were commanded with the interaction torque controller described previously.  $\theta(t)$  was offset such that  $0^\circ$  corresponded to the center of each participant's active range of motion. The target appeared on the display for a duration of 20 seconds and varied according to a multi-sine function:

$$\theta^{\text{des}}(t) = 6.6^\circ [\sin(\pi t^*) + \sin(0.6 \pi t^*) + \sin(0.32 \pi t^*)], \quad (2)$$

where  $\theta^{\text{des}}(t)$  is the instantaneous target angle presented to both participants in the dyad at a given time point ( $t$ ). A time shift,  $t^* = t + t_r$ , with  $t_r$  randomly selected from a uniform distribution in the interval  $[0, 20]$  s was used to change the starting point of the trajectory for each trial and minimize memorization of the target's movement.

To evaluate how tracking performance during physical interaction is affected by the ability or "skill" of each partner, we added visual noise [3] to the target of one participant in each dyad to increase their tracking error (Fig. 1A); adding visual noise in this way increases the relative differences in tracking performance between two partners, resulting in "better" and "worse" partners during the interaction. Without visual noise, the target was presented as a single 10 mm





**Fig. 1.** Experimental setup and block design. (A) Using either their wrists or ankles, each partner tracked multi-sine targets with different visual feedback conditions (i.e., without or with visual noise). (B) During tracking trials, three types of interaction between partners were tested: no interaction (**Solo**), bidirectional (**Dyad**) and unidirectional (**Playback**). In Dyad trials, partners interacted in real-time through compliant spring-damper elements. In Playback trials, each participant was connected to a recording of their partner's Solo trajectory with a virtual stiffness dependent on the ability of their partner ( $1e_2^p$ ), resulting in a range of stiffnesses used across participants during this condition. (C) Participants completed four blocks of tracking trials, 20 seconds in duration, for each combination of visual noise and joint (i.e., ankle or wrist); each block consisted of 5 Solo trials, followed by 10 haptic trials (Dyad or Playback in a randomized order).

diameter disk on the display. With visual noise, the target was presented as a cloud of eight 5 mm diameter disks. Each disk was characterized by the following randomly selected parameters: the orthogonal distance to the target  $\eta \in N(0, 10 \text{ mm})$ , the angular distance to the target  $\eta_\theta \in N(0, 4.58^\circ)$ , and the angular velocity  $\eta_{\dot{\theta}} \in N(0, 4.58^\circ/\text{s})$ . The eight disks were sequentially replaced and updated with parameters every 250 ms.

In total, the experiment consisted of four blocks, for each combination of visual noise (i.e.,  $i$  with noise and  $j$  without,  $j$  with noise and  $i$  without) and joint (i.e., wrist, ankle) (Fig. 1C). At the start of each block, participants were given time to familiarize themselves with the robot's transparent control for the given visual condition; this involved following a multi-sine trajectory,  $\theta^{\text{des}}(t) = 6.6^\circ[\sin(0.8\pi t) + \sin(0.4\pi t)]$ , for 60 seconds followed by one tracking trial defined in Eq. (2). Familiarization was provided in order to minimize the confound of learning during subsequent trials, as our primary analysis focuses on changes in tracking behaviors during haptic interaction. For the remaining trials, participants were informed that they might experience some forces from their robot, but they were blinded to the nature of these forces (i.e., the virtual connection). Within each block, dyads performed 5 **Solo** trials where the robots allowed transparent motion and 10 haptic trials (**Dyad** or **Playback**) where the robots rendered a spring-damper between the joint angles of each partner

according to Eq. (1). The order of Dyad or Playback trials was randomized in the set of 10 haptic trials to mitigate the influence of trial number on tracking error for our primary comparison in this study. The same time shifts ( $t_r$ ) selected for the Solo trials were used in the Dyad and Playback trials, such that partners were presented the same 5 trajectories across the three trial types within a block.

**1) Dyad Trials:** In the Dyad trials, the two partners received and applied haptic feedback to one another via bidirectional interaction. This interaction was achieved using the real-time joint angles of each partner and a virtual stiffness ( $K_v^{\text{wrist}} = 3.7 \text{ Nm/rad}$ ;  $K_v^{\text{ankle}} = 37 \text{ Nm/rad}$ ) and damping ( $C_v^{\text{wrist}} = 0.86 \text{ Nms/rad}$ ;  $C_v^{\text{ankle}} = 2.72 \text{ Nms/rad}$ ). These virtual stiffnesses were selected using a similar approach to the haptic tracking experiment in [3]. In pilot experiments, participants were virtually connected, using two different sets of virtual stiffnesses for the wrist and ankle, to a target multi-sine trajectory; participants were asked to track the target's movement by minimizing the interaction torques experienced without the aid of visual feedback. Tracking errors, relative to the virtual stiffnesses, were characterized by an exponential decay function, as errors asymptotically decreased as the virtual stiffness increased. The virtual stiffness values which produced similar tracking errors for the ankle and wrist were selected for each joint, respectively, for implementation in the Dyad

trials. The set of virtual stiffnesses tested for each joint during piloting was chosen based on active stiffness values reported in previous works for the wrist [18] and ankle [19]. Damping parameters were chosen such that the damping ratio ( $\zeta \propto C_v/\sqrt{K_v} = 1.4$ ) of each connection was the same; this ratio was decided empirically to avoid oscillation and maintain the responsiveness of the interaction.

**2) Playback Trials:** In the Playback trials, partners received haptic feedback via unidirectional interaction. Participants were connected to a recording of their partner's solo trajectory (featuring the same initial time shift  $t_r$ ) with a virtual stiffness that was variable across participants ( $K_v^{\text{wrist}} \in [0.7, 2.3]$  Nm/rad;  $K_v^{\text{ankle}} \in [7, 23]$  Nm/rad) and damping consistent with the Dyad trials. Illustrated in Fig. 2, a range of virtual stiffnesses in this condition was implemented to account for the spring-series connection between the joints (i.e., human stiffness elements) of two users [9]. This spring-series model attempts to relate each partner's planned movement to their actual movement during connected trials as a result of the interaction mechanics. When two partners interact in real-time, the influence of one partner on the other depends on each partner's joint stiffness, relative to the stiffness of the virtual connection. For instance, if partner A is much stiffer than partner B, partner A will be less influenced by the planned movement of partner B; in turn, partner B will be more influenced by the planned movement of partner A. This results in a smaller displacement between partner A's planned and actual movements, and a larger displacement between partner B's planned and actual movements. This effectively means that the virtual connection stiffness perceived by partner A is smaller than that of partner B. In the spring-series model, we can compute the sum of two stiffnesses in series, the virtual connection stiffness and partner B's joint stiffness, to calculate the virtual stiffness perceived by partner A, and vice versa. With this in mind, we rescaled the virtual stiffness in the Playback condition in an attempt to reproduce the virtual stiffness perceived by each partner during the Dyad trials.

Partner stiffness values ( $K$ ) were computed as a function of each partner's ability ( $\Delta e_s^p$ , defined in section II-E), assuming that worse partners were more relaxed and better partners stiffer [5], [9]. Based on the literature values of active joint stiffness referenced previously [18], [19], we empirically generated sigmoid functions to relate differences in Solo trial performance across pairs of participants to their expected joint stiffness during connected trials,

$$K = a_0 + \frac{a_1 - a_0}{1 + \exp(-0.02 \Delta e_s^p)}, \quad (3)$$

where  $a_0, a_1$  are the sigmoid parameters defined separately for the wrist ( $a_0 = 0.1$  Nm/rad,  $a_1 = 8.0$  Nm/rad) and ankle ( $a_0 = 1.0$  Nm/rad,  $a_1 = 80$  Nm/rad). Sigmoid functions, displayed in Fig. 2 as the partner stiffness, were used to avoid negative stiffness values of  $K$ . The Playback virtual stiffness was calculated by taking the series equivalent of the virtual connection stiffness in the Dyad trials and the computed partner stiffness:  $(1/K_v + 1/K)^{-1}$ . Rescaling the virtual stiffness to account for the expected interaction mechanics results in a lower magnitude of the virtual stiffness implemented during

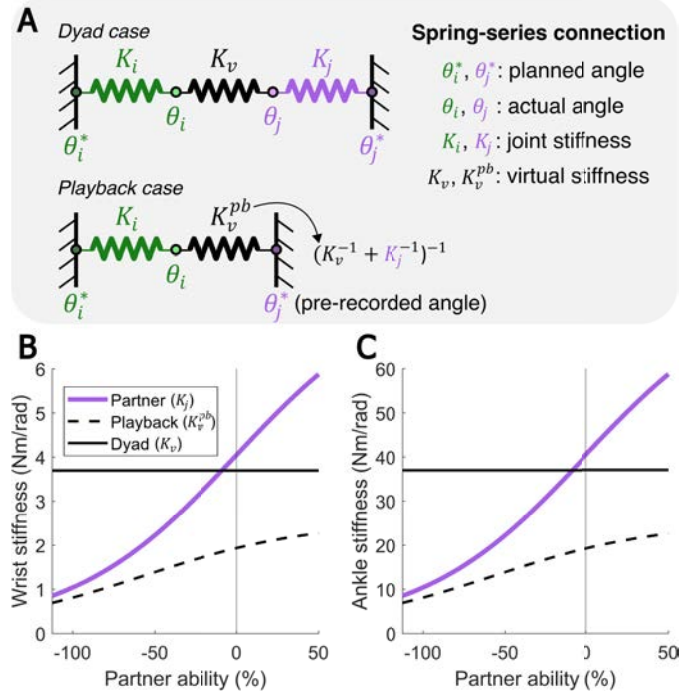


Fig. 2. (A) Spring-series representation of the mechanical interaction between two users, presented for the Dyad case (i.e., real-time interaction) and Playback case (i.e., interaction with pre-recorded trajectory). Implemented values of partner joint stiffness ( $K_i$ ) and virtual connection stiffness ( $K_v$ ) are plot with respect to partner ability and displayed for the (B) wrist and (C) ankle. Dyad virtual stiffness was defined as a constant value ( $K_v^{\text{wrist}} = 3.7$  Nm/rad;  $K_v^{\text{ankle}} = 37$  Nm/rad) for all participants. Playback virtual stiffness was defined separately for each participant, by taking the series equivalent of the Dyad virtual stiffness and partner stiffness  $(1/K_v + 1/K_i)^{-1}$ , resulting in a range of stiffness values during these trials ( $K_v^{\text{wrist}} = [0.7, 2.3]$  Nm/rad;  $K_v^{\text{ankle}} = [7, 23]$  Nm/rad).

Playback compared to Dyad trials, particularly for the better partner (Fig. 2B).

#### D. EMG Calibration

EMG data from isometric torque matching trials were used to estimate torques exerted during the trajectory tracking trials. Participants performed isometric trials while fixed in the center of their active range of motion, matching a visual display of their applied torque to a series of target torque values for a period of 7 seconds (wrist:  $[-3, -2, -1, 1, 2, 3]$  Nm; ankle:  $[-6, -4, -2, 2, 4, 6]$  Nm). These torque magnitudes represent the range of torques participants were expected to exert during the dynamic tracking trials, determined through pilot testing. Trials were repeated 2 to 3 times for each target torque with 10 seconds of rest between trials. The envelopes of EMG activity obtained during these isometric matching trials and at rest were regressed with the applied torque to generate models relating muscle activation to torque.

Two models characterized the flexor (plantarflexor) and extensor (dorsiflexor) torques ( $\tau_f(t)$ ,  $\tau_e(t)$ ) for each joint:

$$\begin{aligned} \tau_f(t) &= \alpha_f u_f(t), & \alpha_f &> 0 \\ \tau_e(t) &= \alpha_e u_e(t), & \alpha_e &> 0 \end{aligned} \quad (4)$$

where  $u_f(t), u_e(t)$  are the EMG envelopes for the flexor and extensor muscles, respectively. For the wrist, the ECRL

represents the extensor muscle and the FCR represents the flexor in these equations. For the ankle, TA represents the extensor muscle and one of MG, LG or SOL was selected to represent the flexor. This was decided based on the highest variance explained when using each muscle as an input for the flexor model in Eq. (4). Across all participants, this procedure resulted in appropriate linear fits of the isometric data (Wrist:  $R^2 = 0.89 \pm 0.09$ , Ankle:  $R^2 = 0.85 \pm 0.08$ ), with respect to similar EMG-torque calibration models used in related studies [5], [20]. We used predictions of torque from EMG signals as a method to normalize the muscle activation of the flexors and extensors on a subject-specific basis during the dynamic tracking trials. Predicting torque in this way also takes into account the relative torque production of the flexors and extensors, respectively, which is necessary for our analysis of muscle co-contraction detailed in the following section.

### E. Outcome Measures

Our primary outcome measures during the trajectory tracking trials were the changes in tracking error and muscle co-contraction during the haptic conditions. All measures defined below were averaged across trials of the same condition, with respect to each joint (i.e., wrist or ankle), visual condition (i.e., with or without noise) and tracking trial type (i.e., Solo, Playback or Dyad). Each participant's tracking error ( $e$ ) was quantified for each trial as the root-mean-square error ( $^\circ$ ) between the target and joint trajectories,

$$e = \left( \frac{1}{T - t_0} \int_{t_0}^T [\theta^{\text{des}}(t) - \theta(t)]^2 dt \right)^{\frac{1}{2}}, \quad t_0 = 2 \text{ s}, \quad T = 20 \text{ s}, \quad (5)$$

computed in the range of 2 to 20 seconds, excluding the first 2 seconds of each trial to account for initiation of the task. For each joint and visual noise condition, tracking improvements were measured by taking the normalized difference between mean Solo ( $e_s$ ) and Dyad ( $e_d$ ) or Playback ( $e_{pb}$ ) tracking errors. Positive values of the tracking improvements indicate better performance for Dyad ( $\Delta e_d = 100(1 - e_d/e_s)$ ) or Playback ( $\Delta e_{pb} = 100(1 - e_{pb}/e_s)$ ) compared to Solo trials. Partner ability was measured by taking the normalized difference between the mean Solo tracking errors of each partner in a dyad ( $\Delta e_s^p = 100(1 - e_s^p/e_s)$ ), with respect to each block. Positive values of partner ability indicate the worse partner in the pair, evaluated when performing the task alone with a given joint and visual noise condition.

Subject-specific models relating EMG to torque were used to quantify the mechanical contribution of agonist-antagonist co-activation [5], [21], termed co-contraction ( $c$ ):

$$c = \frac{1}{T - t_0} \int_{t_0}^T \min\{|\tau_f(t)|, \tau_e(t)\} dt, \quad t_0 = 2 \text{ s}, \quad T = 20 \text{ s}, \quad (6)$$

and computed in the range of 2 to 20 seconds. For each participant, co-contraction values were normalized based on the maximum ( $c_{\max}$ ) and minimum ( $c_{\min}$ ) values observed across all trials for each joint:

$$\Delta c = 100 \left( \frac{c - c_{\min}}{c_{\max} - c_{\min}} \right), \quad (7)$$

reported as mean percentages for the Dyad ( $\Delta c_d$ ) and Playback ( $\Delta c_{pb}$ ) conditions. We used this specific measure of co-contraction to be consistent with previous dyadic studies [5], [20], allowing us to interpret results in the context of these works. Normalizing co-contraction based on the minimum and maximum values observed within each participant gives an indication of how each participant changes their muscle activation strategy during the Playback and Dyad conditions. Additionally, because each participant exhibited different tracking errors when presented targets with or without visual noise, leading to distinct partner ability values ( $\Delta e_s^p$ ) for each visual condition, this normalization method enables more effective within-subject comparisons of changes in co-contraction when a participant performed “better” or “worse” than their partner.

### F. Statistical Analysis

The goal of this study was to assess how task performance improvements and muscle activation strategies change for the wrist and ankle during uni- and bidirectional dyadic interaction. We tested two primary hypotheses for our experimental results: (1) for both the wrist and ankle, task performance improvements are different between the unidirectional and bidirectional tracking conditions, (2) improvements are different during the bidirectional condition between the wrist and ankle. As a secondary analysis, we assessed whether distinct muscle activation strategies were utilized between the wrist and ankle during haptic trials, with the expectation that better partners will exhibit higher levels of muscle co-contraction compared to worse partners for both joints.

To test our hypotheses related to changes in task performance, we used a mixed effects model with the tracking improvement in the haptic trials as a dependent variable, haptic trial type (Dyad or Playback) and joint (ankle or wrist) as categorical variables, and partner ability with linear and quadratic forms as continuous variables, as well as the interaction between each of these predictors. We used a mixed effects model with similar structure to evaluate differences in muscle activation, with normalized co-contraction in the haptic trials as a dependent variable, haptic trial type and joint as categorical variables, and partner ability as a continuous variable. The DoF of the mixed effects models were estimated using a Satterthwaite approximation [22]. Significance was set to 0.05 for all hypotheses related to tracking improvements and co-contraction changes. Results are presented as mean  $\pm$  standard error unless otherwise specified.

## III. RESULTS

Errors during Solo trials were similar between the ankle and wrist when tracking without visual noise (ankle:  $4.1 \pm 0.7^\circ$ , wrist:  $4.0 \pm 0.8^\circ$ ;  $t_{25} = 0.6$ ,  $p = 0.6$ ) and with visual noise (ankle:  $5.0 \pm 0.7^\circ$ , wrist:  $5.3 \pm 1.0^\circ$ ;  $t_{25} = -1.7$ ,  $p = 0.1$ ). Shown in Fig. 3, these errors were positively correlated between the ankle and wrist for both visual conditions (without visual noise:  $r = 0.87$ ,  $p < 0.001$ ; with visual noise:  $r = 0.74$ ,  $p < 0.001$ ). Despite these similarities, the range of partner abilities was larger for the wrist ( $[-110, 50]\%$ ) compared



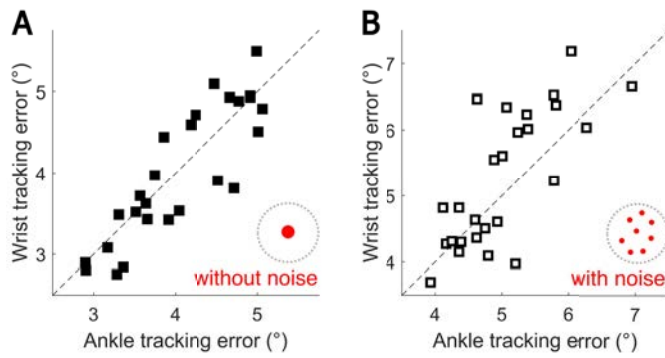


Fig. 3. Solo tracking errors were similar for the wrist and ankle, for both visual conditions. Mean tracking error (°) of each participant during Solo trials, plot for the wrist with respect to the ankle, when tracking targets (A) without visual noise and (B) with visual noise. Dashed lines indicate unity between Solo errors for the wrist and ankle.

to the ankle ( $[-65, 40]\%$ ), meaning that there was a wider distribution of partner performances during wrist tracking. Shown in Fig. 4A,B, improvements during the haptic trials, normalized with respect to Solo performance, increased with partner ability ( $\Delta e_s^p$ ). This means that worse partners, compared to better partners, demonstrated greater improvements when haptically connected in both the Dyad and Playback trials. This relationship between partner ability and tracking improvements was characterized by a second order polynomial, including data for both the wrist and ankle during Dyad and Playback trials ( $R^2 = 0.79$ ).

#### A. Bidirectional Improvements Are Similar Between the Ankle and Wrist

The improvements in tracking performance observed during the Dyad trials were similar for the wrist and ankle. Mixed effects model fits comparing ankle and wrist tracking improvements with respect to partner ability were not significantly different ( $F_{3,186} = 0.5$ ,  $p = 0.7$ ). Assessed over the range of partner abilities common to both joints ( $[-65, 40]\%$ ), there were no significant differences ( $p > 0.3$ ) in the improvements seen during Dyad trials between the wrist and ankle (Fig. 4C). To confirm that these findings were not dependent on the polynomial fit of the mixed effects model, we also compared wrist and ankle improvements separately for better ( $\Delta e_s^p < 0\%$ ) and worse ( $\Delta e_s^p > 0\%$ ) partners. Again, we did not observe significant differences in wrist and ankle improvements for the better ( $\delta = 1.1 \pm 1.8\%$ ,  $t_{200} = 0.4$ ,  $p = 0.7$ ) or worse partners ( $\delta = -0.2 \pm 1.8\%$ ,  $t_{200} = -0.1$ ,  $p = 0.9$ ).

#### B. Better Partners Improve Less During Unidirectional Interaction

Better partners tracked the trajectory more accurately during the bidirectional trials (Dyad) compared to the unidirectional trials (Playback); though worse partners showed no difference in tracking performance between the two trial types. Shown in Fig. 4D,E, we observed that the tracking performance in the Playback and Dyad conditions significantly differed for both the wrist ( $F_{3,184} = 6.4$ ,  $p < 0.001$ ) and ankle ( $F_{3,184} = 5.6$ ,  $p < 0.01$ ). With the wrist, participants had significantly

lower tracking improvements during Playback trials only when partner ability was below  $0\%$  ( $p < 0.05$ ). With the ankle, participants had significantly lower tracking improvements during Playback trials when partner ability was below  $-16\%$  ( $p < 0.05$ ). Comparing improvements during Playback and Dyad trials separately for the better and worse partners, the performance of the better partners was significantly deteriorated for the wrist ( $\delta = -7.2 \pm 1.8\%$ ,  $t_{200} = -2.9$ ,  $p < 0.01$ ) and ankle ( $\delta = -5.9 \pm 1.8\%$ ,  $t_{200} = -2.4$ ,  $p < 0.05$ ) during Playback trials; the performance of the worse partners during Playback and Dyad trials was not significantly different for the wrist ( $\delta = 0.04 \pm 1.8\%$ ,  $t_{200} = 0.02$ ,  $p = 1.0$ ) or ankle ( $\delta = -0.3 \pm 1.8\%$ ,  $t_{200} = -0.1$ ,  $p = 0.9$ ).

#### C. Muscle Co-Contraction Is Modulated at the Wrist, but Not at the Ankle, When Interacting With a Partner

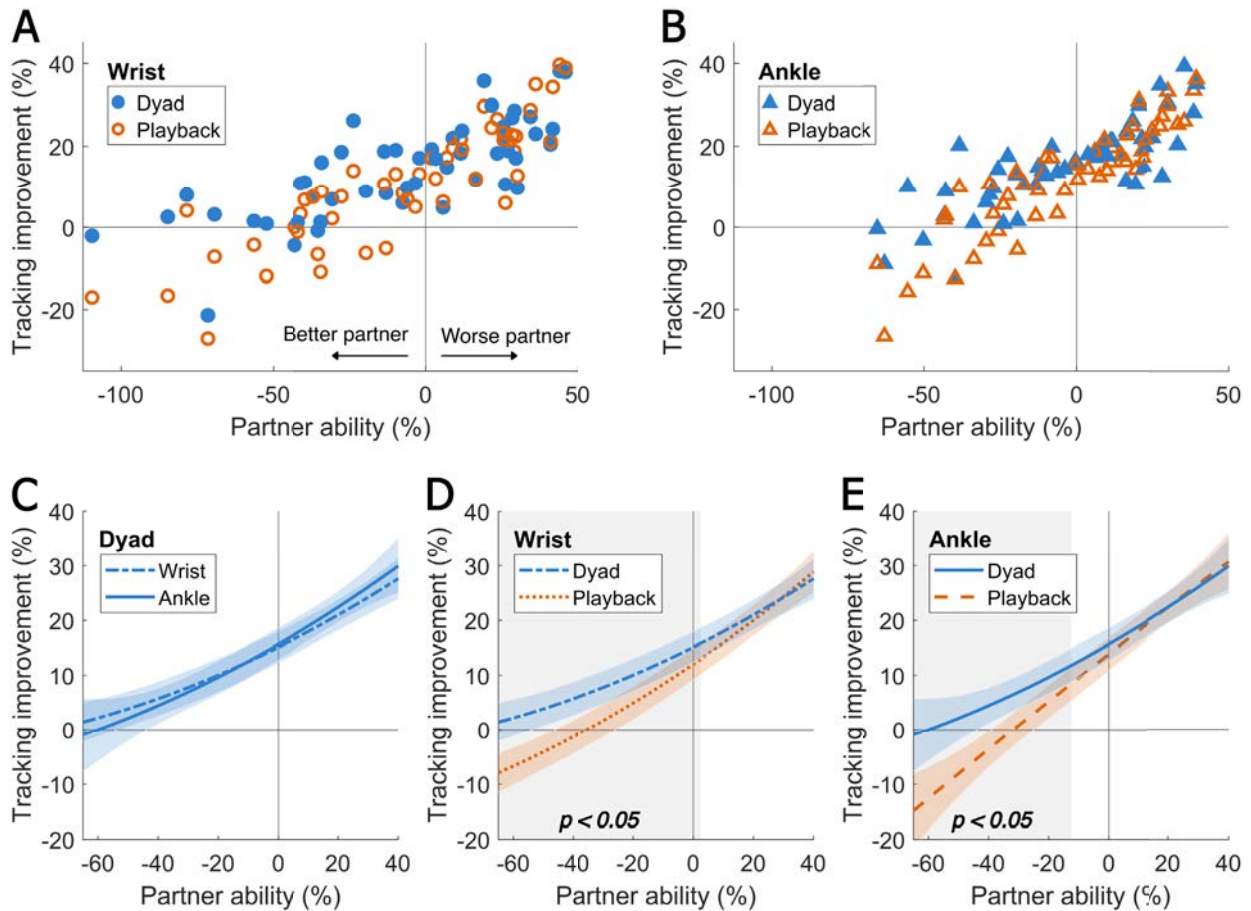
Only at the wrist did we observe changes in muscle co-contraction dependent on partner ability during the haptic trials (Fig. 5A). Mixed effects model fits comparing ankle and wrist co-contraction during bidirectional (Dyad) trials were significantly different ( $F_{2,200} = 5.6$ ,  $p < 0.01$ ). At the wrist, better partners co-contracted more than worse partners during Dyad trials, indicated by a significant change in co-contraction relative to partner ability ( $-33.1 \pm 6.4\%/%$ ,  $t_{200} = -5.2$ ,  $p < 0.001$ ). This trend was also observed during Playback trials. Though we found no significant difference in the change in co-contraction relative to partner ability ( $\delta = -4.4 \pm 6.4\%/%$ ,  $t_{200} = -0.5$ ,  $p = 0.6$ ), there was a significant increase in co-contraction ( $\delta = 7.1 \pm 2.5\%$ ,  $t_{200} = 2.0$ ,  $p < 0.05$ ) during Playback compared to Dyad trials at the wrist.

At the ankle, participants exhibited similar magnitudes of co-contraction independent of their partner's ability (Fig. 5B). There was no such modulation of co-contraction in the Dyad trials, as we observed a change in co-contraction relative to partner ability that was not significantly different from zero in our mixed effects model analysis ( $-3.1 \pm 8.7\%/%$ ,  $t_{200} = -0.4$ ,  $p = 0.7$ ). This was consistent for the Playback trials as well, with no significant difference in changes in co-contraction ( $\delta = -3.8 \pm 8.7\%/%$ ,  $t_{200} = -0.3$ ,  $p = 0.8$ ) between the two haptic conditions.

While normalized co-contraction was significantly modulated at the wrist according to partner ability, the magnitude of these changes was relatively small. To provide context, we report the group averaged, maximum and minimum co-contraction values observed at the wrist ( $c_{\min} = 0.28 \pm 0.17$  Nm;  $c_{\max} = 0.42 \pm 0.22$  Nm) and ankle ( $c_{\min} = 0.87 \pm 0.35$  Nm;  $c_{\max} = 0.97 \pm 0.39$  Nm). Based on the trend observed in normalized co-contraction for the wrist, our model predicts that a much better partner ( $\Delta e_s^p = -110\%$ ) exhibits a co-contraction value that is approximately 0.1 Nm greater than a much worse partner ( $\Delta e_s^p = 50\%$ ).

## IV. DISCUSSION

In this study, we investigated the effect of uni- and bidirectional physical interaction between healthy individuals on performance during a dynamic tracking task, in addition to a comparison of wrist and ankle behaviors during these haptic



**Fig. 4.** Dyadic tracking improvements were similar for the wrist and ankle, relative to each partner's ability; better partners perform worse when connected to a recording of their partner's trajectory (i.e., Playback), compared to a real-time connection (i.e., Dyad). Mean tracking improvements during Dyad and Playback trials plot with respect to the difference in each partner's Solo performances are shown for the (A) wrist and (B) ankle. Linear mixed effects model fits are compared between (C) the wrist and ankle during Dyad trials, (D) Dyad and Playback trials for the wrist and (E) Dyad and Playback trials for the ankle. Vertical shaded areas indicate significant differences in the hypothesis tests that individuals improved differently as a function of partner ability for each combination of conditions.

conditions. To the best of our knowledge, this was the first study to compare such behaviors across the upper and lower limbs using the same robotic infrastructure for the joints tested (i.e., wrist and ankle).

#### A. Similar Tracking Improvements for the Wrist and Ankle

Comparing the wrist and ankle, we observed very similar trends in tracking improvements during real-time, bidirectional interaction with a partner. At both joints, individuals performed the 1-DoF tracking task better while connected to their partner in real-time versus alone; the magnitude of these tracking improvements were dependent on each partner's ability (i.e., worse partners improved more than better partners). This finding is well-established and consistent with previous upper- [1], [3] and lower-limb [8], [9] studies involving compliant physical interaction between healthy individuals. However, generalization of findings across the upper and lower limbs was previously limited as these studies utilized different robots and limb postures in their experimental setups. Our work fills this gap, showing that dyadic tracking improvements can be similarly leveraged at the wrist and ankle, as long as the virtual connection stiffness is selected appropriately for the joints of interest.

A secondary, but noteworthy finding in this work was the similarities observed for the ankle and wrist during trials without interaction between partners (i.e., tracking alone). Whether trajectories were presented without or with visual noise, we found no difference in participants' tracking errors across joints. Solo tracking errors were highly correlated between the ankle and wrist (without noise:  $r = 0.86$ ; with noise:  $r = 0.74$ ), meaning that the skill level of each participant generalized well across joints. Daily tasks involving these human systems are quite different, as the upper limb is typically involved in discrete, goal-directed activities like reaching and manipulation while the lower limb performs rhythmic behaviors like stepping. In contrast to the cortical-driven control of the upper limb, it is known that spinal networks are essential for locomotion in the lower limb [23], and that their function tends to be less mutable over short-term periods than cortical networks [24], [25]. Despite these distinctions, our findings are in line with previous work involving goal-directed "pointing" tasks for the ankle and wrist [15], [16], suggesting that the central nervous system uses a similar feedforward strategy to control the position of the foot or hand during ballistic movements. Our results suggest that the wrist and ankle are capable of similar performances during continuous dynamic



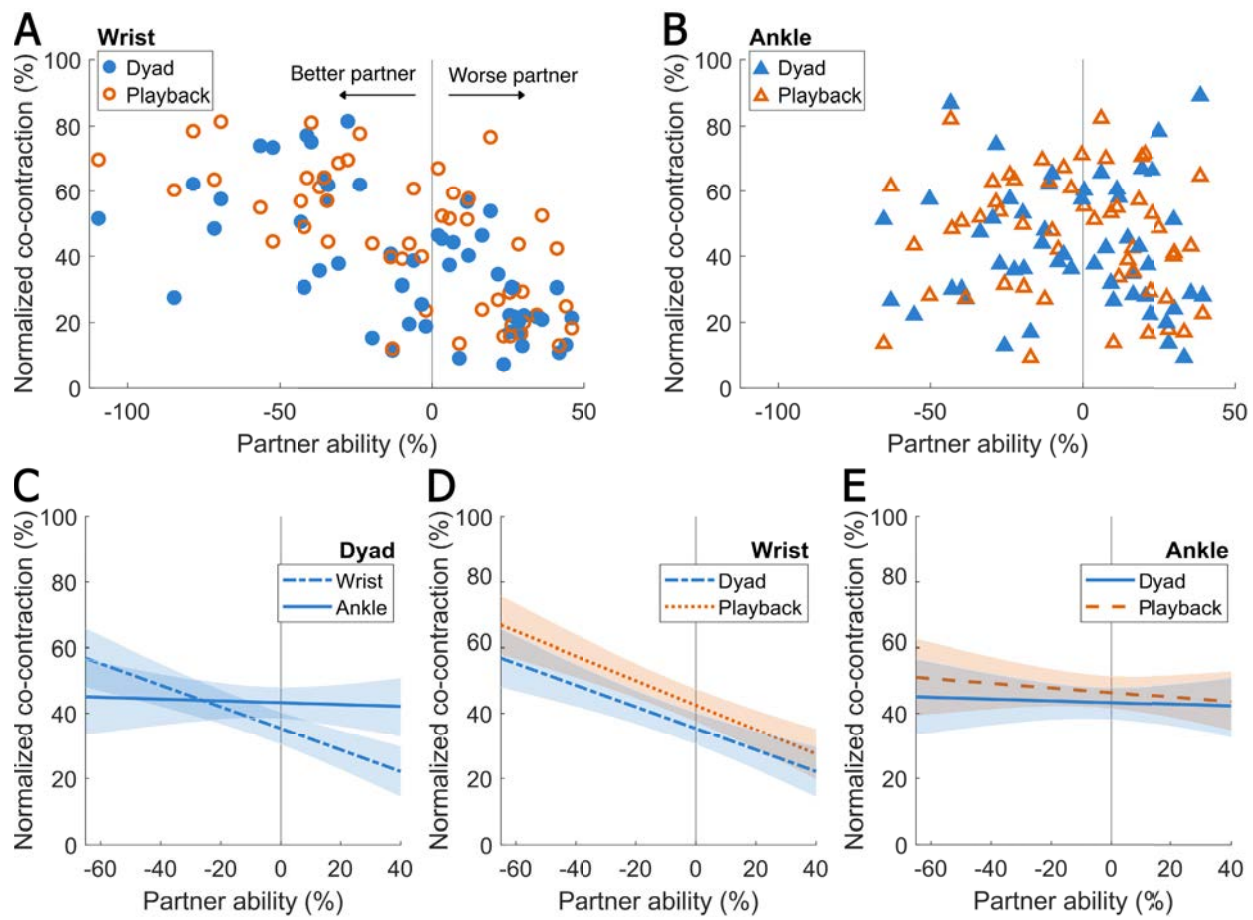


Fig. 5. Changes in co-contraction relative to partner ability were only observed for the wrist, as better partners co-contracted more than worse partners; for the ankle, participants maintained similar levels of co-contraction regardless of their partner's ability. Normalized co-contraction during Dyad and Playback trials plot with respect to the difference in each partner's solo performances are shown for the (A) wrist and (B) ankle. Linear mixed effects model fits are compared between (C) the wrist and ankle during Dyad trials, (D) Dyad and Playback trials for the wrist and (E) Dyad and Playback trials for the ankle.

tracking at relatively slow speeds ( $|\dot{\theta}| < 40^\circ/\text{s}$ ) as well, though kinematic modeling of these behaviors could be tested further in future work.

### B. Unique Muscle Co-Contraction Strategies Across Joints

Though we observed similar tracking improvements for the wrist and ankle during bidirectional interaction, the two joints differed in terms of muscle activation during the task. Specifically, at the wrist, individuals modulated co-activation of their antagonist-agonist muscles according to the ability of their partner, perceived via the virtual connection; this was demonstrated by larger normalized co-contraction values exerted by the better partner in the dyad compared to the worse partner. However, it is important to note that, though significant, the magnitude of these co-contraction changes was relatively small in magnitude. For the ankle, co-contraction was not dependent on partner ability and remained relatively constant for each individual across the tracking conditions. Despite these observed differences, our results for the ankle and wrist indicate that improvements in tracking performance can be achieved without or with changes in co-contraction of the interacting partners for these two joints, respectively.

In general, co-contraction changes in a predictable manner during the learning of novel tasks, as individuals decrease co-contraction to minimize metabolic cost and increase co-contraction to improve accuracy in response to the presented environment or task [26]. Co-contraction of the upper limb has been well-studied in the context of dynamic tasks like reaching [27], drumming [28], and trajectory tracking [5]. In the lower limb, co-contraction has primarily been studied during whole-body, loaded conditions such as walking [29] and balancing [30], and less extensively studied during isolated, dynamic tracking. Our previous work [9] showed modest changes in ankle co-contraction and overall muscle activation for better partners during dyadic tracking; however, these changes were primarily observed in a condition where individuals were rigidly connected with a very stiff spring. In the context of the compliant interaction studied in this work, it is possible that it is not advantageous or natural to modulate co-contraction with the ankle as observed at the wrist. This is supported by evidence that ankle co-contraction alone is not enough to ensure stability in unstable environments; instead, volitional descending control, associated with longer delays, is necessary to stabilize the system [31], [32], [33]. Therefore, our observed differences could indicate key differences in

stiffness control strategies between the upper and lower limbs during dynamic tracking tasks featuring external forces.

### C. Benefits of Bidirectional Interaction Depend on Partner Ability

Comparing the task performance effects of uni- and bidirectional interaction, we found that only the bidirectional interaction allowed *both* partners to improve their tracking performance, compared to tracking without any interaction, across the full range of partner abilities. This finding was observed for both the ankle and the wrist, as better partners experienced greater tracking improvements when connected to their partner in real-time compared to a recording of their partner's trajectory. It is important to note, however, that the worse partner in the dyad improved similarly whether or not they were connected to their partner in real-time. This differs from the results of Ganesh et al. [1], which reported that both better and worse partners tracked targets less accurately when connected to a pre-recorded trajectory during a 2-DoF upper-limb task. As this previous work involved 2-DoF reaching movements under an imposed visuomotor rotation, it is possible that this discrepancy in findings is related to task complexity during dyadic interaction; real-time partner responses may be more important during less constrained tasks involving adaptations to visual transformations, compared to simple 1-DoF tracking under visual noise. Our findings could also be attributed, in part, to our rescaling of the virtual stiffness during the unidirectional connection, emphasizing the importance of considering the human stiffness elements involved in the interaction which resulted in a lower virtual stiffness implemented during the unidirectional trials compared to bidirectional.

Similarities between the worse partners' uni- and bidirectional tracking improvements suggest that, for less skilled individuals, simply being connected to a reference trajectory which is closer to the target trajectory (i.e., the better partner's trajectory) is sufficient for improving task performance during training; this strategy of providing assistance towards a predefined reference is often implemented in rehabilitation robotics to demonstrate desired movements to impaired individuals [34], but its effectiveness in enhancing motor learning can be limited depending on the task constraints and the skill level of the individual [35]. As our study focused on task performance effects rather than changes in individual learning, it is an open question whether interaction with a reactive partner, interaction with a human partner's trajectory, or interaction with a "perfect" reference trajectory is the most appropriate strategy for training in healthy individuals or populations with sensorimotor impairments.

### D. Modeling Dyadic Improvements

To explain the mechanism of improvements during bidirectional interaction with a partner, two distinct models have been proposed for the upper and lower limbs, respectively. Previous work on upper-limb dyadic behaviors has emphasized the role of mutual adaptations in movement planning by each partner during the interaction [2], [3], [10]; when tracking a randomly

moving target, individuals are suggested to improve their tracking performance as they infer the target's movement from the haptic information received from their partner. Instead, our work in the lower limb suggests a mechanism for these improvements based solely on the interaction mechanics [8], [9]; individuals are assumed to track targets independently and any observed improvements are effectively a weighted average of two partially correlated trajectories (i.e., two users following a common trajectory), considering the spring-series connection between two human joints which are virtually connected. In this mechanics-based model, changes in movement planning are not accounted for, therefore the real-time responses of each partner are considered negligible.

As we observed a deterioration in performance during unidirectional trials compared to bidirectional, this would suggest that real-time responses between partners contribute significantly to the tracking improvements observed during dyadic interaction at the wrist and ankle. Interaction mechanics may also contribute to these improvements, but it is likely that partners additionally co-adapt their movement strategies to maximally improve their tracking performance during the interaction. Future work could develop a model which considers both the interaction mechanics and changes in movement planning to characterize the relative contribution of each component during dyadic interaction. Based on our observed differences in co-contraction modulation across joints, changes in mechanics due to volitional muscle activation may contribute less to improvements during bidirectional interaction at the ankle compared to the wrist. It is also possible that movement-dependent changes in joint stiffness, namely passive and reflex stiffness components [36], may contribute differentially to mechanics-related improvements in the wrist and ankle during the interaction. However, both of these interpretations cannot be concluded with the current experimentation and require further investigation.

### E. Limitations

One limitation of this study is the selection of partner stiffness values used in the rescaling of the virtual stiffness during the unidirectional condition. Though we assumed that partner stiffness would change as a function of partner ability, this was only suggested by the trends in wrist co-contraction, but not for the ankle. Co-contraction can give an indication of how the active component of joint stiffness may change during the task, however, it is not a direct measure of joint stiffness. Furthermore, the range of partner stiffness values used in our experimentation was based on isometric data published previously for the wrist and ankle [18], [19] and it is known that stiffness of human joints can decrease considerably from postural tasks to dynamic movements [36]. Because we defined the partner stiffness values based on our hypothesis that stiffness would change as a function of partner ability, rather than attempting to estimate the stiffness during tracking, this may have affected the perceived virtual environment when participants experienced the unidirectional condition compared to the bidirectional interaction.

Despite this limitation related to the estimation of joint stiffness, we believe our presented findings are still relevant and

can be leveraged in future applications of this unidirectional configuration. Using our empirically generated functions to rescale the stiffness of the virtual connection during Playback trials resulted in a larger range of virtual stiffness values implemented for the better partner ( $-110\% < \Delta e_s^p < 0\%$ ;  $K_v^{\text{wrist}} \in [0.7, 1.9]$  Nm/rad,  $K_v^{\text{ankle}} \in [7, 19]$  Nm/rad) but a much smaller range for the worse partner ( $0\% < \Delta e_s^p < 50\%$ ;  $K_v^{\text{wrist}} \in [1.9, 2.3]$  Nm/rad,  $K_v^{\text{ankle}} \in [19, 23]$  Nm/rad). As we observed similar tracking improvements during Dyad and Playback trials, only for the worse partners in each pair, this suggests that the worse partner's improvements can be maintained by using a smaller virtual stiffness in the unidirectional application compared to the bidirectional application. The range of virtual stiffness values we implemented for the worse partner gives an idea of how much this virtual stiffness should be reduced (approximately 50%) with respect to the bidirectional implementation.

To address this limitation and appropriately quantify the joint stiffness of interacting users, wrist and ankle stiffness should be estimated using kinematic responses to torque perturbations during the tracking task [37]. This estimation method would also provide a continuous measure of joint stiffness, which would allow analysis of changes in mechanics within the trajectory tracking trials during different haptic conditions. However, this approach is not straightforward, as a large quantity of repeatable movements is necessary to produce accurate estimates of joint impedance [38]. In the context of our dynamic tracking task, this presents a challenge as the target trajectories are designed to appear "random" to prevent memorization of the target's movement, resulting in a few overlapping segments across multiple trials of the same haptic conditions. In addition, external torque perturbations, which are used to estimate stiffness, must be designed in a way that they do not interfere with the torque feedback transmitted between partners during the interaction.

One other limitation relates to the reported magnitudes of tracking improvements during the Haptic trials (i.e., Dyad, Playback). Tracking improvements were computed with respect to baseline errors during tracking trials without interaction (i.e., Solo), which always preceded the Haptic trials. It is possible that individual learning following the Solo trials contributed, in part, to the tracking improvements reported. However, it is important to note that learning (i.e., an improvement in Solo trial performance) would influence the magnitude of tracking improvements equally for Dyad and Playback trials: our primary comparison in this work. In addition, the range of improvements we observed in our Haptic trials was quite similar to those obtained in previous dyadic studies involving 1-DoF tracking at the wrist [3] and ankle [9], suggesting that the effect of learning was small if present.

## V. CONCLUSION

During a 1-DoF trajectory tracking task, we found that bidirectional physical interaction between healthy individuals coupled at the wrist and ankle, respectively, results in similar trends of tracking improvements across joints. This was observed despite distinct muscle activation strategies across

joints, as participants connected at the wrist co-contracted more when paired with a worse partner while participants connected at the ankle did not modulate co-contraction. In addition, we found that worse partners improved similarly during uni- and bidirectional interaction, while better partners were negatively affected by the unidirectional connection to the worse partner. Together, these results suggest that partners leverage some combination of the virtual interaction mechanics and adaptations in movement planning to maximally benefit while interacting in real-time; however, additional modeling work is necessary to characterize the relative contributions of mechanics and adaptations in planning for the wrist and ankle, respectively. Similarities in the response to uni- and bidirectional connections across the upper and lower limbs demonstrate that the benefits of human-human physical interaction generalize across these human systems for simple 1-DoF tracking tasks. As this work was tested in healthy individuals, future work should explore these behaviors in populations with sensorimotor impairments to determine the functional benefits of interacting with a partner during rehabilitation compared to conventional approaches in robotic assistance.

## REFERENCES

- [1] G. Ganesh, A. Takagi, R. Osu, T. Yoshioka, M. Kawato, and E. Burdet, "Two is better than one: Physical interactions improve motor performance in humans," *Sci. Rep.*, vol. 4, no. 1, p. 3824, Jan. 2014.
- [2] A. Takagi, G. Ganesh, T. Yoshioka, M. Kawato, and E. Burdet, "Physically interacting individuals estimate the partner's goal to enhance their movements," *Nature Hum. Behav.*, vol. 1, no. 3, p. 0054, Mar. 2017.
- [3] A. Takagi, F. Usai, G. Ganesh, V. Sanguineti, and E. Burdet, "Haptic communication between humans is tuned by the hard or soft mechanics of interaction," *PLOS Comput. Biol.*, vol. 14, no. 3, Mar. 2018, Art. no. e1005971.
- [4] E. Ivanova, J. Eden, G. Carboni, J. Krüger, and E. Burdet, "Interaction with a reactive partner improves learning in contrast to passive guidance," *Sci. Rep.*, vol. 12, no. 1, p. 15821, Sep. 2022.
- [5] H. Börner et al., "Physically interacting humans regulate muscle coactivation to improve visuo-haptic perception," *J. Neurophysiol.*, vol. 129, no. 2, pp. 494–499, Feb. 2023.
- [6] A. Noccaro et al., "Robot-mediated asymmetric connection between humans can improve performance without increasing effort," *IEEE Trans. Biomed. Eng.*, early access, Mar. 7, 2025, doi: 10.1109/TBME.2025.3548884.
- [7] J. P. Batson et al., "Haptic coupling in dyads improves motor learning in a simple force field," in *Proc. 42nd Annu. Int. Conf. IEEE Eng. Med. Biol. Soc. (EMBC)*, Jul. 2020, pp. 4795–4798.
- [8] S. J. Kim et al., "Effect of dyadic haptic collaboration on ankle motor learning and task performance," *IEEE Trans. Neural Syst. Rehabil. Eng.*, vol. 31, pp. 416–425, 2023.
- [9] M. R. Short et al., "Haptic human-human interaction during an ankle tracking task: Effects of virtual connection stiffness," *IEEE Trans. Neural Syst. Rehabil. Eng.*, vol. 31, pp. 3864–3873, 2023.
- [10] A. Takagi, M. Hirashima, D. Nozaki, and E. Burdet, "Individuals physically interacting in a group rapidly coordinate their movement by estimating the collective goal," *eLife*, vol. 8, p. 41328, Feb. 2019.
- [11] J. Cao, S. Q. Xie, R. Das, and G. L. Zhu, "Control strategies for effective robot assisted gait rehabilitation: The state of art and future prospects," *Med. Eng. Phys.*, vol. 36, no. 12, pp. 1555–1566, 2014.
- [12] N. Lodha, P. Patel, A. Casamento-Moran, E. Hays, S. N. Poisson, and E. A. Christou, "Strength or motor control: What matters in high-functioning stroke?" *Frontiers Neurol.*, vol. 9, p. 1160, Jan. 2019.
- [13] E. Basalp, P. Wolf, and L. Marchal-Crespo, "Haptic training: Which types facilitate (re)learning of which motor task and for whom? Answers by a review," *IEEE Trans. Haptics*, vol. 14, no. 4, pp. 722–739, Oct. 2021.



- [14] E. L. Waters, R. J. Mendonca, P. Z. Cacchione, and M. J. Johnson, "TheraDyad: Feasibility of an affordable robot for multi-user stroke rehabilitation," in *Proc. 10th IEEE RAS/EMBS Int. Conf. Biomed. Robot. Biomechatronics (BioRob)*, Sep. 2024, pp. 1498–1503.
- [15] L. Vaisman, L. Dipietro, and H. I. Krebs, "A comparative analysis of speed profile models for wrist pointing movements," *IEEE Trans. Neural Syst. Rehabil. Eng.*, vol. 21, no. 5, pp. 756–766, Sep. 2013.
- [16] K. P. Michmizos, L. Vaisman, and H. I. Krebs, "A comparative analysis of speed profile models for ankle pointing movements: Evidence that lower and upper extremity discrete movements are controlled by a single invariant strategy," *Frontiers Human Neurosci.*, vol. 8, p. 962, Nov. 2014.
- [17] S. J. Kim et al., "A framework for dyadic physical interaction studies during ankle motor tasks," *IEEE Robot. Autom. Lett.*, vol. 6, no. 4, pp. 6876–6883, Oct. 2021.
- [18] A. Leger and T. E. Milner, "Passive and active wrist joint stiffness following eccentric exercise," *Eur. J. Appl. Physiol.*, vol. 82, nos. 5–6, pp. 472–479, Aug. 2000.
- [19] P. L. Weiss, I. W. Hunter, and R. E. Kearney, "Human ankle joint stiffness over the full range of muscle activation levels," *J. Biomechanics*, vol. 21, no. 7, pp. 539–544, Jan. 1988.
- [20] A. Takagi, C. Bagnato, A. Melendez-Calderon, N. Jarrassé, G. Ganesh, and E. Burdet, "Competition increases the effort put into a physical interaction task," *IEEE Trans. Haptics*, vol. 16, no. 4, pp. 719–725, Oct. 2023.
- [21] E. Burdet, D. W. Franklin, and T. E. Milner, *Human Robotics: Neuro-mechanics and Motor Control*. Cambridge, MA, USA: MIT Press, 2013.
- [22] S. G. Luke, "Evaluating significance in linear mixed-effects models in R," *Behav. Res. Methods*, vol. 49, no. 4, pp. 1494–1502, Aug. 2017.
- [23] H. Hultborn and J. B. Nielsen, "Spinal control of locomotion—from cat to man," *Acta Physiologica*, vol. 189, no. 2, pp. 111–121, Feb. 2007.
- [24] J. Shemmell, M. A. Krutky, and E. J. Perreault, "Stretch sensitive reflexes as an adaptive mechanism for maintaining limb stability," *Clin. Neurophysiol.*, vol. 121, no. 10, pp. 1680–1689, Oct. 2010.
- [25] J. R. Wolpaw and A. M. Tennissen, "Activity-dependent spinal cord plasticity in health and disease," *Annu. Rev. Neurosci.*, vol. 24, no. 1, pp. 807–843, Mar. 2001.
- [26] K. A. Thoroughman and R. Shadmehr, "Electromyographic correlates of learning an internal model of reaching movements," *J. Neurosci.*, vol. 19, no. 19, pp. 8573–8588, Oct. 1999.
- [27] D. W. Franklin, R. Osu, E. Burdet, M. Kawato, and T. E. Milner, "Adaptation to stable and unstable dynamics achieved by combined impedance control and inverse dynamics model," *J. Neurophysiology*, vol. 90, no. 5, pp. 3270–3282, Nov. 2003.
- [28] S. Beveridge, S. A. Herff, B. Buck, G. B. Madden, and H.-C. Jabusch, "Expertise-related differences in wrist muscle co-contraction in drummers," *Frontiers Psychol.*, vol. 11, p. 1360, Jul. 2020.
- [29] F. Di Nardo, A. Mengarelli, E. Maranesi, L. Burattini, and S. Fioretti, "Assessment of the ankle muscle co-contraction during normal gait: A surface electromyography study," *J. Electromyogr. Kinesiol.*, vol. 25, no. 2, pp. 347–354, Apr. 2015.
- [30] D. Kim and J.-M. Hwang, "The center of pressure and ankle muscle co-contraction in response to anterior-posterior perturbations," *PLoS ONE*, vol. 13, no. 11, Nov. 2018, Art. no. e0207667.
- [31] P. Morasso, M. Casadio, D. De Santis, T. Nomura, F. Rea, and J. Zenzeri, "Stabilization strategies for unstable dynamics," *J. Electromyogr. Kinesiol.*, vol. 24, no. 6, pp. 803–814, 2014.
- [32] M. Casadio, P. G. Morasso, and V. Sanguineti, "Direct measurement of ankle stiffness during quiet standing: Implications for control modelling and clinical application," *Gait Posture*, vol. 21, no. 4, pp. 410–424, Jun. 2005.
- [33] J. M. Finley, Y. Y. Dhaher, and E. J. Perreault, "Contributions of feed-forward and feedback strategies at the human ankle during control of unstable loads," *Exp. Brain Res.*, vol. 217, no. 1, pp. 53–66, Mar. 2012.
- [34] A. Roy, L. W. Forrester, and R. F. Macko, "Short-term ankle motor performance with ankle robotics training in chronic hemiparetic stroke," *J. Rehabil. Res. Develop.*, vol. 48, no. 4, p. 417, 2011.
- [35] L. Marchal-Crespo, N. Rappo, and R. Riener, "The effectiveness of robotic training depends on motor task characteristics," *Exp. Brain Res.*, vol. 235, no. 12, pp. 3799–3816, Dec. 2017.
- [36] D. Ludvig, M. Plocharski, P. Plocharski, and E. J. Perreault, "Mechanisms contributing to reduced knee stiffness during movement," *Exp. Brain Res.*, vol. 235, no. 10, pp. 2959–2970, Oct. 2017.
- [37] D. Ludvig, M. W. Whitmore, and E. J. Perreault, "Leveraging joint mechanics simplifies the neural control of movement," *Frontiers Integrative Neurosci.*, vol. 16, Mar. 2021, Art. no. 802608.
- [38] D. Ludvig and E. J. Perreault, "System identification of physiological systems using short data segments," *IEEE Trans. Biomed. Eng.*, vol. 59, no. 12, pp. 3541–3549, Dec. 2012.