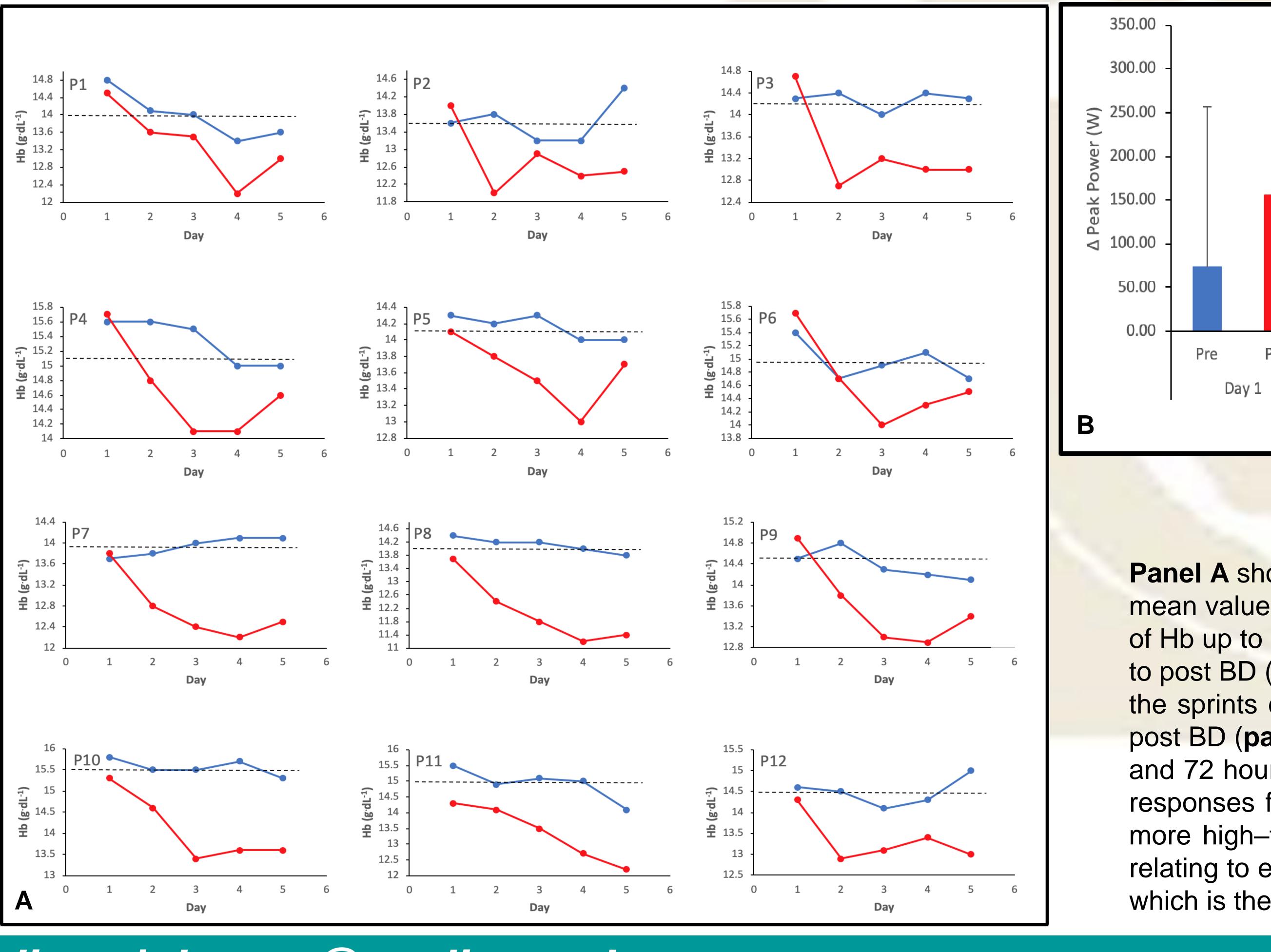


Anglia Ruskin University

PURPOSE: Previous studies investigating the effects of BD (~ 470ml) over a period of up to 96 hours on power output during four repeated 15s sprints. METHOD: Following local institutional ethical approval 12 participants, 9 males (age 31.2 ± 12.6 years, mass 84.3 ± 11.1 Kg, height 176.9 ± 6.9 cm) and 3 females (age 34.7 ± 10.3 years, mass 67.0 ± 8.5 Kg, height 167.1 ± 4.2 cm) volunteered to participate. Testing was conducted in the morning over a two week block with a rest week between testing phases. Visit one on week 1 (W1) established haematological levels and VO_{2max}. Subsequent visits during a Lode Excalibur Sport cycle ergometer. Week 2 (W2) was identical except the participant donated blood on visit 1 post haematological testing vo₂, cardiac output, stroke volume and heart rate using thoracic impedance cardiography was computed, while Near Infrared Spectroscopy measured O₂ delivery at the muscle. **RESULTS:** Hb on W2 day 1 (Hb=14.58 ± 0.69g·dL⁻¹, Hct=42.83 ± 2.04%) compared to day 2 post BD (Hb=13.52 ± 0.95g dL⁻¹, Hct=39.75 ± 2.93%) (P = 0.001), a percentage increase (1.5%) (P = 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 ± 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 (73.8 \pm 0.001) occurred between days 4-5. Δ PPO (W) pre-BD on day 1 182.5), 2 (46.6 ± 143.8), 3 (25.8 ± 75.3) and 4 (35.7 ± 95.6) compared to post BD on day 2 (P = 0.01). Mean PPO values reduced after each sprint on each testing day, with post BD values being a values being a values being a values being a values reduced after each sprint on each testing day, with post BD values being a values bevelocity being a values being a values bevelocity b generally higher than pre-BD values, however significance was only detected pre to post BD on day 2 (P = 0.02) and 3 (P = 0.02) in the first sprints. CONCLUSIONS: The data suggests that BD has significant effects on Hb and Hct up to 96 hours post BD. PPO appears most effected 48 and 72 hours following BD, possible explained by the decrement in Hb being greater 48-96-h post BD, this produced an effect of recruiting more high-threshold motor units due to the lack of circulating O₂. These data have possible ramifications for NHS guidelines relating to exercise post BD.

INTRODUCTION

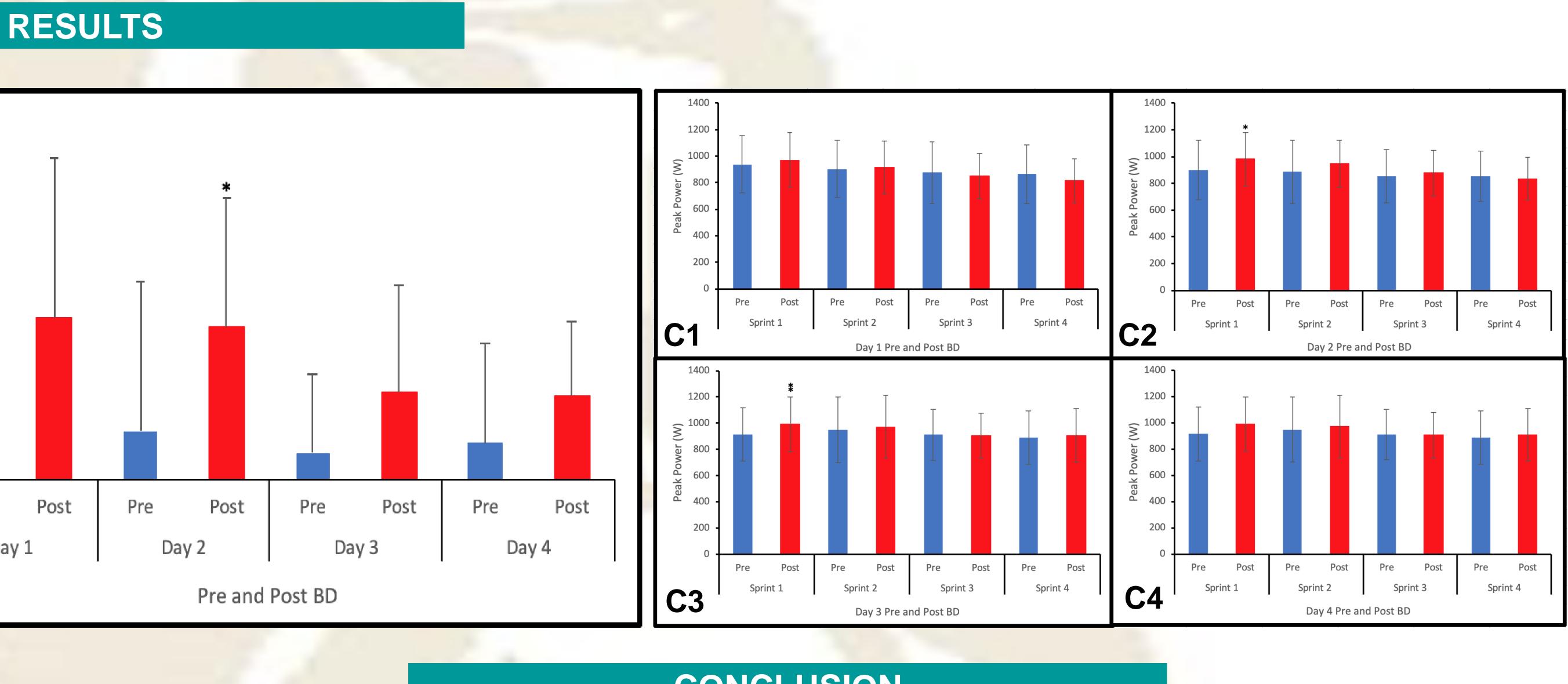
Haemoglobin (Hb) decreases following blood donation (BD) and post BD advice is to avoid strenuous Following local institutional ethical approval 12 participants, 9 males, 3 females volunteered to participate. exercise particularly in the first 12-24 hours. However, Hb levels appear to continue to decrease Testing was conducted in the morning over a two week period with a rest week between testing phases. beyond the first 24 hours post BD, while high intensity interval type classes appear ever popular Week one (W1) was pre-BD and week two (W2) was post BD (through NHS BD services). Visit one on W1 amongst the exercising population research is justified. Thus, this study determines the effects of BD established haematological levels and VO_{2max}. Subsequent visits during W1 were for Sprint Interval Testing over a period of up to 96 hours on Peak Power Output (PPO) during 4 repeated 15s sprints in a (SIT) (4 x 15s with 90s of unloaded pedalling prior to each effort), with a resistance of 0.07w kg⁻¹ of body moderately trained population. mass. W2 was identical except the participant donated blood on visit 1 post haematological testing only.



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Influence of whole blood donation (~ 470ml) on peak power outputs over a 96h period Diane Johnson¹, Justin Roberts¹, DanGordon¹, Viviane Merzbach¹, Dirk Dugdale¹, Henry Hodgkins¹, Everonika Gordon¹, Madie Rowland¹, Ellen Lockwood¹, Antonio Femminile¹, Flora Veres¹ ¹Cambridge Centre for Sport and Exercise Sciences, Anglia Ruskin University, Sport and Exercise Sciences Research Group, Cambridge, UK

ABSTRACT



CONCLUSION

Panel A shows the heterogeneous Hb pre (blue) and post (red) BD for all participants, with the dotted line indicating the mean value for week 1. The mean Hb pre to post BD significantly (P = 0.01) reduced, impacting the O₂ carrying capacity of Hb up to 96 hours post BD. Panel B signifies Δ PPO (from sprint 1 to 4) was significantly influenced up to 48 hours preto post BD (P = 0.01). The mean PPO values for sprints (panels C1 to C4) show reduced PPO throughout the course of the sprints conceivably indicating that with each sprint PCr restoration time is insufficient. However, 24 and 96 hours post BD (panel C1 and C4) an impairment of O₂ availability was not affecting PPO (P > 0.05), while PPO for sprint-1, 48 and 72 hours post BD was significantly (P = 0.001 and P = 0.02 respectively) increased. A possible explanation to the responses for PPO is that as the decrement in Hb was greater 48-96-h post BD, this produced an effect of recruiting more high-threshold motor units due to the lack of circulating O₂. These data have ramifications for NHS guidelines relating to exercise post BD because it appears that the continual fall in Hb is affecting physiology longer than 24 hours, which is the period suggested to be approached with caution regarding exercise.



METHOD