

**SDC 3 - Table - Characteristics of included studies**

First author, year, country.	Participants N (N women) Age Phase: acute or persistent	Intervention, Comparison, frequency, duration	Outcome Measures	Reported results (from the study authors and from the Meta-analysis)
Maerlender et al. 2015 (35) USA	<p><b>Total:</b> N= 28 (20 women)</p> <p><b>Intervention group:</b> N= 13 (8 women)</p> <p><b>Control group:</b> N= 15 (12 women)</p> <p><b>Age:</b> College athletes (no specific age group reported)</p> <p><b>Phase:</b> Acute SRC</p>	<p><b>Intervention:</b> Schwinn Airdyne stationary Bicycle at a perceived exertion level of mild to moderate (0 to 6 on 10-point RPE scale) for 20 minutes or until symptoms became uncomfortable</p> <p><b>Comparison:</b> Rest (no physical or mental exertion)</p> <p><b>Frequency:</b> Daily</p> <p><b>Duration:</b> Daily follow-up until symptoms-free. The maximum duration is unknown.</p>	<p><b>Primary:</b> Time to recovery (days)</p> <p><b>Secondary:</b> Time to recovery (subgroup with 2 weeks cutoff) ImPACT Actigraph Symptoms change after bike ride.</p>	<p><b>Time to recovery [median number of days (range)]:</b> Intervention: [15 (5 – 61)] Control: [13 (6 – 56)]</p> <p>No statistically significant difference between groups</p> <p><b>ImPACT:</b> No statistically significant difference between groups for all measures of the impact test</p> <p><b>Other secondary outcome measures:</b> NR</p>

		<b>Timing:</b> Exercise program began after randomization.		
Thomas et al. 2015 (34) USA	<p><b>Total:</b> N= 99 (16 women)</p> <p><b>Intervention group:</b> N= 50 (18 women)</p> <p><b>Control group:</b> N= 50 (18 women)</p> <p><b>Age:</b> 11-22 years old</p> <p><b>Phase:</b> Acute SRC</p>	<p><b>Intervention:</b> 1-2 days of rest followed by stepwise return to physical and mental activities</p> <p><b>Comparison:</b> Strict rest for 5 days followed by stepwise return to physical and mental activities</p> <p><b>Frequency:</b> Daily (starting after 2 days in the intervention group and 5 days in the Control group)</p> <p><b>Duration:</b> A total of 10 days</p> <p><b>Timing:</b> Initiation of stepwise physical activation was allowed 24h-48h post-injury.</p>	<p><b>Primary:</b> PCSS (19 symptoms version); ImPACT.</p> <p><b>Secondary:</b> BESS; Subjects experiencing symptom resolution</p>	<p><b>PCSS at the end of study mean (SD):</b> Intervention: 8 (13.00) Control: 13 (14.99)</p> <p>No statistically significant difference between groups</p> <p><b>Total PCSS scores over the 10 days:</b> Control: 187.9 Intervention: 131.9</p> <p>Statistically significant difference favoring Intervention group (P&lt;0.03)</p> <p><b>Number of symptoms over the 10 days:</b> Control: 70.4 Intervention: 50.2</p> <p>Statistically significant difference favoring Intervention group (P&lt;0.03)</p> <p><b>BESS (Mean and 95%CI):</b></p>

				<p>Intervention: 19 (15.9 – 22.1)</p> <p>Control: 21 (18.4 – 23.6)</p> <p>No statistically significant difference between groups</p> <p><b>ImPACT:</b></p> <p>No statistically significant difference between groups for all measures of the impact test</p>
<p>Kurowski et al. 2017 (38) USA</p>	<p><b>Total:</b> N= 30 (17 women)</p> <p><b>Intervention group:</b> N= 15 (10 women)</p> <p><b>Control group:</b> N= 15 (7 women)</p> <p><b>Age:</b> 12-17 years old</p> <p><b>Phase:</b> Persistent symptoms after SRC (4 to 6 weeks)</p>	<p><b>Intervention:</b> Sub-symptom exacerbation aerobic training (80% of time) on stationary bike during follow-up visits (mean 4.42) with an intensity target of 11-16 on Borg scale (weekly assessment) for a maximum of 30 minutes.</p> <p><b>Comparison:</b> full body stretching (program rotated every 2 weeks with a follow-up every week).</p> <p><b>Frequency:</b> 5-6 days/week at home</p> <p><b>Duration:</b> A total of 6 weeks (2</p>	<p><b>Primary:</b> PCSI (self-rating)</p> <p><b>Secondary:</b> PCSI (parent-rated) Adherence</p>	<p><b>PCSI at the end of study mean (SD):</b> Intervention: 4.17 (7.36) Control: 15.93 (20.18)</p> <p>Statistically significant difference favoring Intervention group (P&lt;0.044). Largest difference was at 4 weeks (ES: 0.81 (large))</p> <p><b>Number of subjects having symptoms</b> Intervention: 6/15 Control: 13/15</p> <p>Statistically significant difference favoring Intervention group</p>

		additional weeks if symptoms persist). <b>Timing:</b> Exercise program began one week after randomization.		
Chan et al. 2018 (39) Canada	<b>Total:</b> N= 19 (14 women) <b>Intervention group:</b> N= 9 (8 women) <b>Control group:</b> N= 10 (6 women) <b>Age:</b> 12-18 years old <b>Phase:</b> Persistent symptoms after SRC (more than 4 weeks)	<b>Intervention:</b> active rehab program consisting of: 1.- Sub-Symptoms threshold aerobic training for up to 15 minutes 2.- Coordination ex's 3.- Visualisation/imaging with physio 4.- home ex's program 5.- Symptoms-management and RTP advice, RTS facilitation, physiatry consultation if needed <b>Comparison:</b> Symptoms-management and RTP advice, RTS facilitation, physiatry consultation <b>Frequency:</b> 2-5 visits (mean 3,4)	<b>Primary:</b> PCSS (8 times over 6 weeks) <b>Secondary:</b> Recorded adverse events and exacerbation of Symptoms during aerobic training BESS ImPACT BDI PROMIS	<b>PCSS at the end of study [mean (SD)] :</b> Intervention: 51.5 (27.8) Control: 40.3 (29.4) Statistically significant difference favoring Intervention group (P<0.047) <b>BESS [mean (SD)] :</b> Intervention: 10.3 (3.2) Control: 11.4 (7.4) No statistically significant difference between groups <b>ImPACT:</b> No statistically significant difference between groups for all measures of the impact test <b>BDI:</b> No statistically significant difference between groups <b>PROMIS:</b>

		<p><b>Duration:</b> A total of 6 weeks</p> <p><b>Timing:</b> Exercise program began within one week after randomization.</p>		No statistically significant difference between groups for all measures of the PROMIS test
Micay et al. 2018 (40) Canada	<p><b>Total:</b> N= 16 (1 woman)</p> <p><b>Intervention group:</b> N= 8 (0 women)</p> <p><b>Control group:</b> N= 8 (1 woman)</p> <p><b>Age:</b> 14-18 years old</p> <p><b>Phase:</b> Acute SRC (6 days)</p>	<p><b>Intervention:</b> Stationary cycle ergometer at 50% to 70% of age-predicted maximal heart rate (Increasing gradually through the study period); first session duration 10 minutes, all other sessions 20 minutes.</p> <p><b>Comparison:</b> gradual return to activity using the six-stage progression of Berlin 2016 Guidelines</p> <p><b>Frequency:</b> 1 session per day for 2 days in a row followed by 1 day of rest</p> <p><b>Duration:</b> A total of 11 days</p> <p><b>Timing:</b> Exercise program began on day 6 following the injury.</p>	<p><b>Primary:</b> PCSS Time to medical clearance (in days)</p>	<p><b>PCSS at the end of study [mean (SD)] :</b> Intervention: 4.3 (4.1) Control: 10 (6.1)</p> <p><b>Time to medical clearance [mean (SD)] :</b> Intervention: 36.1 (18.5) Control: 29.6 (15.8)</p> <p>No statistically significant difference between groups</p>

<p>Bailey et al. 2019 (37) USA</p>	<p><b>Total:</b> N= 16 (7 woman)</p> <p><b>Intervention group:</b> N= 7 (N women NR)</p> <p><b>Control group:</b> N= 9 (1 woman)</p> <p><b>Age:</b> 14-18 years old</p> <p><b>Phase:</b> Persistent symptoms after SRC (mean 56 days)</p>	<p><b>Intervention:</b> intensive PT treatment consisting of a 20 minutes subthreshold ex's program (HR 80% of Symptoms exacerbation threshold) combined with education session</p> <p><b>Comparison:</b> light exercises (1<sup>st</sup> 3 weeks:5 daily stretching activity and last 3 weeks: 20 min of daily walking)</p> <p><b>Frequency:</b> 3X20min/week + home based daily</p> <p><b>Duration:</b> A total of 6 weeks</p> <p><b>Timing:</b> Exercise program began immediately after randomization.</p>	<p><b>Primary:</b> PCS-R</p> <p><b>Secondary:</b> BDI BESS ImPACT</p>	<p><b>PCS-R at the end of study [mean (SD)] :</b> Intervention: 18.17 (16.45) Control: 16.38 (19.23)</p> <p>No statistically significant difference between groups</p> <p><b>PCS-R at the end of study [Mean and (SD)] when removing influence of depression*:</b> Intervention: 63.3 (17.4) Control: 56.8 (27.8)</p> <p>Statistically significant difference favoring Intervention group (P&lt;0.05), large effect <math>n^2=0,32</math></p> <p>*Removed outlier who showed no improvement (higher depression Symptoms at baseline)</p> <p><b>Other secondary outcome measures:</b> NR</p>
<p>Leddy et al. 2019 (36) USA</p>	<p><b>Total:</b> N= 113 (48 woman)</p> <p><b>Intervention group:</b> N= 52 (24 women)</p>	<p><b>Intervention:</b> Sub-Symptoms threshold exercises (no stretching): bike or treadmill 80% of HR target. Max 20 min or until exacerbation of 2 points on</p>	<p><b>Primary:</b> Days to recovery since date of injury</p>	<p><b>PCSS at the end of study [mean (SD)] :</b> Intervention: 1.08 (3.91) Control: 5.47 (16.34)</p> <p>Significance not reported</p>

	<p><b>Control group:</b></p> <p>N= 51 (24 women)</p> <p><b>Age:</b> 13-18 years old</p> <p><b>Phase:</b> Acute SRC</p>	<p>symptoms score.</p> <p><b>Comparison:</b> rest plus whole body stretching 20 min/daily</p> <p><b>Frequency:</b> Daily</p> <p><b>Duration:</b> maximum of 30 days or until symptoms resolution (whichever came first).</p> <p><b>Timing:</b> Exercise program began 48h post-injury.</p>	<p><b>Secondary:</b></p> <p>PCSS</p> <p>BCTT</p>	<p><b>Time to recovery [median number of days (Interquartile range)]</b></p> <p>Intervention: 13 (10 - 18,5)</p> <p>Control: 17 (13 - 23)</p> <p>Statistically significant difference favoring Intervention group (P&lt;0.009)</p> <p><b>BCTT at the end of study:</b> NR</p>
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Abbreviations and outcomes: N= number of participants, SRC=Sport-related concussion, NR=Not reported, ImPACT: Immediate post-concussion assessment and cognitive testing, BDI: Beck depression inventory, Actigraph: wearable activity and sleep monitor, PCSS=Post-concussion symptoms scale, PSCI=Post-concussion symptoms inventory, PCS-R=Post-concussive scale-Revised, BESS=Balanced error scoring system, BCTT=Buffalo concussion treadmill test, PROMIS= Patient-Reported Outcomes Measurement Information System, pediatric short forms, SD=Standard deviation, CI=Confidence Interval.

