Appendix 1

1# "Fecal Incontinence" OR "Fecal Incontinence" OR "Incontinence, Fecal" OR "Bowel Incontinence" OR "Fecal Soiling" OR "Low anterior resection syndrome" OR "LARS" OR "Faecal incontinence" OR "Sphincter function" OR "Bowel function"

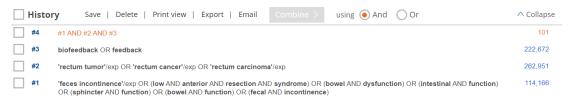
2# "Biofeedback" OR "BF"

3# "Colorectal Surgery" OR "Proctocolectomy" OR "Anal preservation" OR "Sphincter-sparing" OR "Sphincter-saving" OR "Sphincter sparing" OR "Sphincter saving" OR "Low anterior resection" OR "Anterior resection" OR "Intersphincteric resection" OR "Resection"

Pubmed

Conducted on 2020-6-18, number of hits: 130

Embase



Conducted on 2020-6-18, number of hits: 101

Cochrane Library



Conducted on 2020-6-18, number of hits: 14

ClinicalTrials.Gov

(rectal neoplasms OR rectal cancer) AND (biofeedback OR feedback)

Conducted on 2020-6-18, number of hits: 17

Basic information of literature

Review Title					
Date Re	eviewe	er		_	
Study Title					-
First author					
Year of publication					
Country of publication					
Publication type	Jo	urnal/Ab	ostract/O	ther	
Contact details					
Eligibility of included student	dies				
Study type					
Patients with intestinal	dysfu	nction	yes:	no:	
after rectal cancer surgery	y				
Biofeedback therapy			yes:	no:	
Note:					
Characteristics of particip	ants				
Characteristics of participa	Study				
Total number of participa	nts				
Number of groups in	n the				
intervention					
Number of groups in	n the				

control gro	up				
Participants	6	Age: MedianAverageRange			
		Sex:			
		Other:			
Intervention measures		*Record the specific contents, methods, operation specifications, participants, time, frequency and cycle of intervention, and whether the participants have received training*			
Control measures		*Record the specific contents, methods, operation specifications, participants, time, frequency and cycle of intervention, and whether the participants have received training*			
Follow up time					
Outcomes	Diagnostic criteria of	Evaluation of prognosis			
	disease				

Methods: Non randomized study quality assessment: (MINORS0)

Meth	Score				
1,	A clearly stated aim: the question addressed should be				
	precise and relevant in the light of available literature				
2、	Inclusion of consecutive patients: all patients potentially				
	fit for inclusion (satisfying the criteria for inclusion) have				
	been included in the study during the study period (no				
	exclusion or details about the reasons for exclusion)				
3、	Prospective collection of data:data were collected				
	according to a protocol established before the beginning				

of the study

- 4. Endpoints appropriate to the aim of the study:

 unambiguous explanation of the criteria used to evaluate

 the main outcome which should be in accordance with the

 question addressed by the study. Also, the endpoints

 should be assessed on an intention-to-treat basis.
- **5.** Unbiased assessment of the study endpoint: blind evaluation of objective endpoints and double-blind evaluation of subjective endpoints. Otherwise the reasons for not blinding should be stated
- 6. Follow-up period appropriate to the aim of the study:

 the follow-up should be sufficiently long to allow the

 assessment of the main endpoint and possible adverse

 events
- 7. Loss to follow up less than 5%: all patients should be included in the follow up. Otherwise, the proportion lost to follow up should not exceed the proportion experiencing the major endpoint
- 8. Prospective calculation of the study size: information of the size of detectable difference of interest with a calculation of 95% confidence interval, according to the expected incidence of the outcome event, and

information about the level for statistical significance and estimates of power when comparing the outcomes

- 9. An adequate control group: having a gold standard diagnostic test or therapeutic intervention recognized as the optimal intervention according to the available published data
- 10. Contemporary groups: control and studied group should be managed during the same time period (no historical comparison)
- 11. Baseline equivalence of groups: the groups should be similar regarding the criteria other than the studied endpoints. Absence of confounding factors that could bias the interpretation of the results
- 12. Adequate statistical analyses: whether the statistics were in accordance with the type of study with calculation of confidence intervals or relative risk

Note: 0 (Not reported) 1 (Insufficient) 2 (Sufficient)

Results

	Intervention group	Control group
Number		
Exclusion number		
Number of observations		

Number of lost follow-up	
Number of events	

Continuous	Intervention group		Control group			
variable (Before						
intervention)						
Outcomes	N	Mean	Standard	N	Mean	Standard
			deviation			deviation
			(SD)			(SD)
Continuous	Intervention group			Control group		
variable (After						
intervention)						
Outcomes	N	Mean	Standard	N	Mean	Standard
			deviation			deviation
			(SD)			(SD)
Continuous	Intervention group		Control group			

variable						
(Change)						
Outcomes	N	Mean	Standard	N	Mean	Standard
			deviation			deviation
			(SD)			(SD)

Appendix 3

- 1.Dubovyĭ,V.A. Biofeedback training in the treatment of anterior resection syndrome. Lik Sprava2006;(5-6): 55-60.
- 2.Fomenko O.Y,Kashnikov V. N,Alekseev M. V, et al. Rehabilitation program for patients with low anterior resection syndrome. Vopr Kurortol Fizioter Lech Fiz Kult.2020;97(5): 52-59.
- 3.Cohee W,Hurff A,Gazewood D,et al. Benign Anorectal Conditions: Evaluation and Management.Am Fam Physician.2020;101(1): 24-33.
- 4.Andromanakos N,Skandalakis P,Troupis T,et al.Constipation of anorectal outlet obstruction: pathophysiology, evaluation and management. J Gastroenterol Hepatol 2006;21(4): 638-646.
- 5.Andromanakos P,Filippou K,Pinis I,et al.Anorectal incontinence: A challenge in diagnostic and therapeutic approach. European Journal of Gastroenterology and

Hepatology 2013;25(11): 1247-1256.

6.Bartlett L,Sloots K,Nowak M,et al.Biofeedback for fecal incontinence: a randomized study comparing exercise regimens.Dis Colon Rectum 2011;54(7): 846-856.

7.Bartlett L,Sloots K,Nowak M,et al.Supplementary home biofeedback improves quality of life in younger patients with fecal incontinence. Journal of Clinical Gastroenterology 2015;49(5): 419-428.

8.Buhr J,Hoffmann W,Allemeyer E,et al.Clinical pathway for fecal incontinence-results in a 7-year follow-up.Colorectal disease 2019;21: 106.

9.Dalsgaard P,Emmertsen J,Juul T et al.Nurse-led personalized conservative treat ment in patients with Low Anterior Resection Syndrome.Colorectal disease 201 9;21: 112.

10.Dalsgaard P,Emmertsen J,Mekhael M,et al.Nurse-led standardized intervention for low anterior resection syndrome. A population-based pilot study.Colorectal Dis 2021;23(2): 434-443.

11.Ho H,Chiang M,Tan M,et al.Biofeedback therapy for excessive stool frequen cy and incontinence following anterior resection or total colectomy.Dis Colon R ectum 1996;39(11): 1289-1292.

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13.Kang B,T.G.Lee,Muscle regeneration: Research for the treatment of fecal inc ontinence.Journal of the Korean Society of Coloproctology 2010;26(1): 1-7.

14.Rimmer C,Stackhouse K,Cruickshank N,et al.A targeted biofeedback program me improves functional outcome following low anterior resection. Colorectal dise ase. 2012;14: 23.

15.Kim K,Jeon G,Song S,et al.Biofeedback Therapy Before Ileostomy Closure i n Patients Undergoing Sphincter-Saving Surgery for Rectal Cancer: A Pilot Study.Ann Coloproctol 2015;31(4): 138-143.

16.Arnaud A,Fretes R,Joly A,et al.Posterior approach to the rectum for treatme nt of selected benign lesions.Int J Colorectal Dis 1991;6(2): 100-102.

17.Stackhouse K,Clarson E,Smith, J,et al.A targeted biofeedback programme im proves functional outcome following low anterior resection. Colorectal disease 20 18;20: 43.