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人工颈椎间盘置换与颈前路减压植骨融合术治疗脊髓型颈椎病 近期疗效的回顾性研究*

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摘要 目的: 回顾性分析人工颈椎间盘置换(ADR)与颈前路减压植骨融合术(ACDF)治疗脊髓型颈椎病(CSM)的近期疗效。**方法:** 回顾性选取 2016 年 7 月~2018 年 12 月期间我院收治的 CSM 患者 120 例, 上述患者根据手术方式的不同分为 A 组(n=58, ACDF 治疗)和 B 组(n=62, ADR 治疗), 比较两组患者疗效、围术期指标、生活质量简表(SF-12)评分、日本矫形外科协会(JOA)评分、颈椎活动度及并发症发生情况。**结果:** B 组术后 12 个月的优良率为 74.19%(46/62), 高于 A 组的 53.45%(31/58)($P<0.05$)。两组患者术后 6 个月、术后 12 个月躯体健康评分、精神健康评分均较术前升高, 且 B 组高于 A 组($P<0.05$)。B 组术后 1 个月、术后 3 个月、术后 6 个月、术后 12 个月 JOA 评分和颈椎活动度均高于 A 组($P<0.05$)。B 组住院时间、术后颈托固定时间、术后恢复工作时间均短于 A 组($P<0.05$)。两组术后并发症发生率比较差异无统计学意义($P>0.05$)。**结论:** 与 ACDF 治疗相比, ADR 治疗 CSM 的近期疗效显著, 可有效改善患者脊椎功能及生活质量, 且安全性较好, 临床应用价值较高。

关键词: 人工颈椎间盘置换术; 颈前路减压植骨融合术; 脊髓型颈椎病; 近期; 疗效

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A Retrospective Study on the Short-term Effect of Artificial Cervical Disc Replacement and Anterior Cervical Decompression and Fusion in the Treatment of Cervical Spondylotic Myelopathy*

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ABSTRACT Objective: To retrospectively analyze the short-term effect of artificial cervical disc replacement (ADR) and anterior cervical decompression and fusion (ACDF) in the treatment of cervical spondylotic myelopathy (CSM). **Methods:** A total of 120 patients with CSM in our hospital from July 2016 to December 2018 were retrospectively selected. The above patients were divided into group A (n=58, ACDF treatment) and group B (n=62, ADR treatment) according to different surgical methods. The curative effect, perioperative index, quality of life (SF-12) score, Japan Orthopaedic Association (JOA) score, cervical vertebra activity and complications happened of the two groups were compared. **Results:** The excellent and good rate of group B was 74.19% (46/62), which was higher than 53.45% (31/58) of group A ($P<0.05$). The scores of physical health and mental health of the patients of the two groups at 6 months and 12 months after operation were higher than those before operation, and those of group B were higher than those of group A ($P<0.05$). The JOA score and cervical vertebra activity of group B at 1 month, 3 months, 6 months and 12 months after operation were higher than that of group A ($P<0.05$). The hospitalization time, fixation time of cervical bracket after operation and recovery time to work after operation of group B were shorter than those of group A ($P<0.05$). There was no significant difference in the incidence of postoperative complications between the two groups ($P>0.05$). **Conclusion:** Compared with ACDF, ADR can improve the spinal function and quality of life of patients with CSM in the short term, which has better safety, and higher clinical application value.

Key words: Artificial cervical disc replacement; Anterior cervical decompression and fusion; Cervical spondylotic myelopathy; Short-term; Curative effect

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前言

脊髓型颈椎病(CSM)是由于颈椎或其相邻组织发生退行

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性改变,进而引起脊髓神经运动、反射、感觉及排便等一系列的功能障碍的疾病^[1-3]。CSM发病隐匿且起病缓慢,常规的保守治疗效果一般,手术是治疗CSM的有效方法^[4-6]。既往临床通常采用颈前路减压植骨融合术(ACDF)治疗CSM,可在一定程度上缓解患者临床症状^[7,8]。但近年来的临床实践证实,ACDF术后可产生较多并发症,如植骨块脱落、塌陷、椎间高度丢失、假关节丢失、邻近节段应力增加导致的退变等,一定程度上影响治疗效果^[9,10]。人工颈椎间盘置换术(ADR)由Goffin等学者于2002年首次报道,在实现彻底减压的同时能维持置换节段的活动度和邻近椎间盘内的应力,为CSM的手术治疗提供了一种新的选择^[11,12]。鉴于此,本研究通过对比ADR、ACDF治疗CSM后的近期治疗效果,以期为临床治疗CSM的术式选择提供参考,整理如下。

1 资料与方法

1.1 一般资料

回顾性选取2016年7月~2018年12月期间我院收治的CSM患者120例,纳入标准:(1)经X线片、颈椎MRI及CT等影像学确诊为CSM;(2)患者临床资料完整;(3)术前经3个月以上的正规保守治疗无效者;(4)术后获得完整随访。排除标准:(1)合并心、肺、肝、肾功能疾病者;(2)既往有脊柱手术病史者;(3)合并脊柱结核、肿瘤者;(4)病变节段活动度明显丢失或椎间高度丢失>50%者;(5)合并凝血功能障碍者;(6)脊柱存在明显不稳者。上述患者根据手术方式的不同分为A组(n=58,ACDF治疗)和B组(n=62,ADR治疗),其中A组男33例,女25例,年龄34~65岁,平均(42.81±5.64)岁;病程3~12月,平均(7.25±0.94)月;单节段置换34例,双节段置换24例;体质量指数21.3~26.5 kg/m²,平均(23.18±0.84)kg/m²。B组男35例,女27例,年龄36~63岁,平均(42.06±4.97)岁;病程4~10月,平均(7.42±0.94)月;单节段置换36例,双节段置换26例;体质量指数20.9~26.8 kg/m²,平均(23.36±0.97)kg/m²。两组一般资料对比无统计学差异($P>0.05$)。

1.2 方法

A组给予ACDF治疗,全麻,取仰卧位,固定头部,经颈前

入路显露椎体前方,切开前纵韧带,将病变椎间盘及压迫组织切除后,于减压间隙处植入相应大小的自体髂骨,选取适宜长度的钛板行前方椎体间固定。B组给予ADR治疗,术前参考颈椎CT片,制作相应的Bryan假体(美国Sofamor公司),全麻,取仰卧位,固定头部,经颈前入路显露椎体前方,C臂机透视下确定病变椎间隙,处理椎体前方的骨赘及病变椎间盘,选择性切除后纵韧带显露硬脊膜减压。彻底清除椎体后缘的压迫组织和骨赘并彻底减压后,植入Bryan假体,位置满意后常规关闭切口。

1.3 观察指标

(1)记录两组住院时间、术后恢复工作时间、术后颈托固定时间。(2)两组术后采用门诊复查的形式随访12个月。评价两组患者术后12个月的疗效。按照Odom标准进行临床疗效评价^[13],具体如下:差:症状无改善或恶化;可:术前症状部分改善,日常活动受到限制;良:较少不适症状,但未明显影响工作;优:术前症状和体征全部缓解,可进行日常活动。优良率=优率+良率。(3)于术前、术后1个月、术后3个月、术后6个月、术后12个月采用日本矫形外科协会(JOA)^[14]评分评价患者椎椎功能,其中JOA总分17分,分数越高,椎椎功能越好。于术前、术后1个月、术后3个月、术后6个月、术后12个月行X线检测颈椎活动度。(4)于术前、术后6个月、术后12个月采用生活质量简表(SF-12)^[15]评分评价患者生活质量,SF-12包括躯体健康评分、精神健康评分,每个项目各100分,分数越高,生活质量越高。(5)记录两组并发症发生情况。

1.4 统计学方法

本研究数据的处理软件为SPSS21.0,计量资料以($\bar{x} \pm s$)表示,采用t检验,计数资料以[n(%)]表示,采用 χ^2 检验,检验水准设置为 $\alpha=0.05$ 。

2 结果

2.1 两组疗效比较

B组术后12个月的优良率为74.19%(46/62),高于A组的53.45%(31/58)($P<0.05$);详见表1。

表1 两组疗效比较例(%)

Table 1 Comparison of efficacy between two groups n(%)

Groups	Excellent	Good	Can	Bad	Excellent and good rate
Group A(n=58)	10(17.24)	21(36.21)	16(27.59)	11(18.97)	31(53.45)
Group B(n=62)	16(25.81)	30(48.39)	12(19.35)	4(6.45)	46(74.19)
χ^2					4.977
P					0.025

2.2 两组生活质量比较

两组术前躯体健康评分、精神健康评分比较无差异($P>0.05$);两组患者术后6个月、术后12个月躯体健康评分、精神健康评分均较术前升高,且B组高于A组($P<0.05$);详见表2。

2.3 两组JOA评分和颈椎活动度比较

两组术前JOA评分和颈椎活动度比较差异无统计学

意义($P>0.05$);两组术前、术后1个月、术后3个月、术后6个月、术后12个月JOA评分呈升高趋势,颈椎活动度呈先下降后升高趋势($P<0.05$);B组术后1个月、术后3个月、术后6个月、术后12个月JOA评分和颈椎活动度均高于A组($P<0.05$);详见表3。

表 2 两组生活质量比较($\bar{x} \pm s$, 分)

Table 2 Comparison of quality of life between the two groups($\bar{x} \pm s$, score)

Groups	Score of physical health			Score of mental health		
	Before operation	6 months after operation	12 months after operation	Before operation	6 months after operation	12 months after operation
Group A(n=58)	46.38± 5.87	58.10± 7.34 ^a	75.20± 6.83 ^{ab}	45.63± 6.82	62.60± 6.55 ^a	74.29± 6.24 ^{ab}
Group B(n=62)	46.52± 6.02	63.22± 6.12 ^a	89.18± 7.90 ^{ab}	44.78± 7.51	74.67± 5.71 ^a	88.30± 5.07 ^{ab}
t	0.129	4.160	10.338	0.648	10.778	13.537
P	0.898	0.000	0.000	0.518	0.000	0.000

Notes: compared with before operation, ^a $P<0.05$; compared with 6 months after operation, ^b $P<0.05$.

表 3 两组 JOA 评分和颈椎活动度比较($\bar{x} \pm s$)

Table 3 Comparison of JOA score and cervical vertebra activity between the two groups($\bar{x} \pm s$)

Groups	Time point	JOA score(score)	Cervical vertebra activity (°)
Group A(n=58)	Before operation	5.36± 0.84	49.75± 5.70
	1 month after operation	6.77± 0.76 ^a	25.42± 4.65 ^a
	3 months after operation	7.28± 0.62 ^{ab}	32.88± 5.22 ^{ab}
	6 months after operation	9.73± 0.65 ^{abc}	37.32± 6.15 ^{abc}
	12 months after operation	11.87± 0.64 ^{abcd}	41.71± 6.47 ^{abcd}
Group B(n=62)	Before operation	5.61± 0.69	49.12± 6.31
	1 month after operation	8.82± 0.65 ^{ac}	30.28± 6.51 ^{ac}
	3 months after operation	11.86± 0.69 ^{abc}	36.97± 5.45 ^{abc}
	6 months after operation	13.48± 0.50 ^{abce}	42.73± 6.20 ^{abce}
	12 months after operation	16.05± 0.38 ^{abcde}	48.20± 7.13 ^{bcde}

Notes: compared with before operation, ^a $P<0.05$; compared with 1 month after operation, ^b $P<0.05$; compared with 3 months after operation, ^c $P<0.05$; compared with 6 months after operation, ^d $P<0.05$; compared with group A, ^e $P<0.05$.

2.4 两组围术期指标比较

短于A组($P<0.05$);详见表4。

B组住院时间、术后颈托固定时间、术后恢复工作时间均

表 4 两组围术期指标比较($\bar{x} \pm s, d$)

Table 4 Comparison of perioperative indexes between the two groups($\bar{x} \pm s, d$)

Groups	Hospitalization time	Fixation time of cervical bracket after operation	Recovery time to work after operation
Group A(n=58)	11.88± 1.59	44.14± 2.30	49.07± 3.66
Group B(n=62)	6.34± 1.83	26.18± 1.54	28.41± 3.75
t	17.650	50.560	26.941
P	0.000	0.000	0.000

2.5 两组并发症发生率比较

A组术后出现植骨块脱落1例、植骨块塌陷2例、椎间高度丢失2例,并发症发生率为8.62%(5/58);B组术后出现椎间高度丢失1例,并发症发生率为1.61%(1/62);两组术后并发症发生率比较无差异($\chi^2=3.098, P=0.078$)。

3 讨论

CSM是中老年群体中常见的一种退行性病变,在各种类型的颈椎病中危害最为严重。该病的主要病变基础为椎间盘退变,并伴有脊髓和(或)支配脊髓的血管慢性、进行性受压致神经组织缺血受损^[16-18]。针对症状明显且神经功能进行性下降的

CSM患者通常需给予手术干预以改善脊髓血供,进而恢复脊髓功能^[19]。ACDF主要通过颈前路途径直接解除脊髓前方压迫,并通过融合稳定颈椎,起到即刻稳定及消除后方结构再退变的作用,是目前治疗CSM的常用方案^[20,21]。但随着时间的推移,融合手术使融合节段的生理活动范围丧失,导致局部生物力学改变,邻近节段的负荷增加,部分患者融合节段的相邻节段会发生退行性改变,甚至需要接受二次手术治疗,远期预后效果一般^[22,23]。因此,如何保留CSM患者术后病变节段的运动功能,减轻融合节段的相邻节段应力,已成为目前临床CSM手术治疗的研究热点。ADR是目前常用的非融合技术,通过置入人工颈椎间盘假体代替病变节段的结构和功能,发挥较好的对

邻近节段椎间盘的保护作用,近年来成为治疗CSM的重要手段方式^[24]。但有关两种术式的疗效优劣尚需通过在后续研究中扩大样本量进一步分析以证实。

本次研究结果显示,B组术后12个月的优良率、围术期指标改善优于A组,可见与ACDF治疗相比,ADR治疗CSM,可进一步提高患者预后。这主要是因为ADR可在切除突出的椎间盘组织的同时保留椎间高度和关节活动度,既保留了颈椎节段间的活动度,又可减轻邻近节段的应力负荷,进而促进患者恢复,改善患者预后^[25,26]。此外,ADR治疗者的JOA评分和颈椎活动度均优于ACDF治疗者,由于颈椎的稳定性重建需牺牲手术节段的运动功能,理论上会增加邻近节段退变发生的速度。由于ACDF多采用人工骨植骨,一定程度上降低植骨融合率^[27]。而ADR治疗过程中不刻意强调坚固的骨性愈合,在实现彻底减压的同时维持了手术节段的高度和活动度,相对来说更注重恢复脊柱的生理功能,从而有效维持颈椎活动度^[28,29]。而两组患者术后不同时间点生活质量均有所改善,且ADR治疗者的改善效果更佳。这主要是因为ADR治疗的患者术后恢复更快,脊柱的生理功能恢复的更为完整,可促使患者早日回归正常生活,改善其生活质量^[30]。另两组术后并发症发生率比较差异无统计学意义,可见ADR治疗安全性较好。值得注意的是,本次研究样本量偏少,未能观察ADR治疗是否能进一步减少并发症发生率,且由于人工假体不可避免会发生磨损,本次研究随访时间较短,有关假体磨损是否会影响ADR的远期预后尚待进一步研究。

综上所述,与ACDF治疗相比,ADR治疗CSM的近期疗效显著,可有效改善患者脊椎功能及生活质量,且安全性较好,临床应用价值较高。

参考文献(References)

- [1] Sun B, Shi C, Wu H, et al. Application of Zero-profile Spacer in the Treatment of Three-level Cervical Spondylotic Myelopathy: 5-year Follow-up Results[J]. Spine (Phila Pa 1976), 2020, 45(8): 504-511
- [2] Yuan H, Zhang X, Zhang LM, et al. Comparative study of curative effect of spinal endoscopic surgery and anterior cervical decompression for cervical spondylotic myelopathy [J]. J Spine Surg, 2020, 6(Suppl 1): S186-S196
- [3] McCormick JR, Sama AJ, Schiller NC, et al. Cervical Spondylotic Myelopathy: A Guide to Diagnosis and Management [J]. J Am Board Fam Med, 2020, 33(2): 303-313
- [4] Goyal DKC, Murphy HA, Hollern DA, et al. Is the Neck Disability Index an Appropriate Measure for Changes in Physical Function After Surgery for Cervical Spondylotic Myelopathy? [J]. Int J Spine Surg, 2020, 14(1): 53-58
- [5] Yeh KT, Chen IH, Lee RP, et al. Two surgical strategies for treating multilevel cervical spondylotic myelopathy combined with kyphotic deformity[J]. Medicine (Baltimore), 2020, 99(7): e19215
- [6] Xu N, Zhang Y, Zhou G, et al. The value of dynamic MRI in the treatment of cervical spondylotic myelopathy: a protocol for a prospective randomized clinical trial[J]. BMC Musculoskelet Disord, 2020, 21(1): 83
- [7] Lo YL, Zhu L, Soh RC, et al. Intraoperative Motor Evoked Potential Improvement in Cervical Spondylotic Myelopathy: Comparison of Cortical Stimulation Parameters [J]. J Clin Neurol, 2020, 16 (1): 102-107
- [8] Kong W, Xin Z, Du Q, et al. Anterior percutaneous full-endoscopic transcorporeal decompression of the spinal cord for single-segment cervical spondylotic myelopathy: The technical interpretation and 2 years of clinical follow-up[J]. J Orthop Surg Res, 2019, 14(1): 461
- [9] Wu A, Jin H, Dou H, et al. Anterior decompression through transoral axis slide and rotation osteotomy for salvage of failed posterior occipitocervical fusion: a novel technique note [J]. Ann Transl Med, 2020, 8(4): 129
- [10] Jun DS, Baik JM, Lee SK. A case report: white cord syndrome following anterior cervical discectomy and fusion: importance of prompt diagnosis and treatment[J]. BMC Musculoskelet Disord, 2020, 21(1): 157
- [11] Moon JH, Chung CK, Kim CH, et al. Longitudinal change of cervical artificial disc motion following replacement [J]. PLoS One, 2020, 15 (2): e0228628
- [12] Hou WX, Zhang HX, Wang X, et al. Application of a modified surgical position in anterior approach for total cervical artificial disc replacement[J]. World J Clin Cases, 2020, 8(1): 38-45
- [13] 陶海鹰, 杨博, 卫爱林, 等. Centerpiece 钛板系统治疗多节段脊髓型颈椎病的早期疗效分析 [J]. 临床外科杂志, 2019, 27(12): 1060-1062
- [14] 苟波, 苏权. 颈椎后路3种手术方式对多节段脊髓型颈椎病生理曲度的影响及远期疗效观察[J]. 空军医学杂志, 2018, 34(1): 28-31, 35
- [15] 周逸彬, 李西成. C2-C7活动度预测脊髓型颈椎病症状严重程度的临床价值[J]. 颈腰痛杂志, 2020, 41(1): 45-47
- [16] 贾斌, 梁冠楠, 陈宇飞, 等. 脊髓型颈椎病的发病机制及治疗的研究进展[J]. 现代生物医学进展, 2015, 15(19): 3766-3769, 3776
- [17] Fan XW, Wang ZW, Gao XD, et al. The change of cervical sagittal parameters plays an important role in clinical outcomes of cervical-spondylotic myelopathy after multi-level anterior cervical discectomy and fusion[J]. J Orthop Surg Res, 2019, 14(1): 429
- [18] Chen C, Yang C, Yang S, et al. Clinical and Radiographic Outcomes of Modified Unilateral Open-door Laminoplasty with Posterior Muscle-Ligament Complex Preservation for Cervical Spondylotic Myelopathy[J]. Spine (Phila Pa 1976), 2019, 44(24): 1697-1704
- [19] Kim J, Kim JY, Lee JM, et al. Progressive Cervical Spondylotic Myelopathy Caused by Tic Disorders in a Young Adult with Tourette Syndrome[J]. Korean J Neurotrauma, 2019, 15(2): 199-203
- [20] Camillo FX, Mitchell SM. Cervical Spine Decompression and Fusion Outcomes in Trauma Patients Actively Receiving Anticoagulation Treatment for Cerebrovascular Injury: A Retrospective Comparative Study[J]. Int J Spine Surg, 2020, 14(1): 66-71
- [21] Zhao XF, Zhao YB, Lu XD, et al. Development and Clinical Application of a New Open-Powered Nail Anterior Cervical Plate System[J]. Orthop Surg, 2020, 12(1): 248-253
- [22] 王建, 陈杰妮, 李德彬, 等. 颈椎前路减压植骨融合钢板内固定术后邻近节段退变的危险因素研究 [J]. 颈腰痛杂志, 2019, 40(6): 806-808
- [23] 孙永彪, 艾合买提·吾买尔, 赵岩, 等. 颈椎前路"杂交式"与颈椎后路治疗多节段脊髓型颈椎病的临床比较 [J]. 现代生物医学进展, 2017, 17(22): 4262-4267

- in High Risk Women for endometrial carcinoma [J]. *Cell Mol Biol (Noisy-le-grand)*, 2018, 64(11): 88-91
- [8] Cui J, Qu Z, Harata-Lee Y, et al. Cell Cycle, Energy Metabolism and DNA Repair Pathways in Cancer Cells are Suppressed by Compound Kushen Injection[J]. *BMC Cancer*, 2019, 19(1): e103
- [9] Kim H, Kim YM. Pan-cancer analysis of somatic mutations and transcriptomes reveals common functional gene clusters shared by multiple cancer types[J]. *Entific Reports*, 2018, 8(1): e6041
- [10] Bignottle E, Zanotti I, Calza S, et al. Trop-2 protein overexpression is an independent marker for predicting disease recurrence in endometrioid endometrial carcinoma[J]. *BMC Clinical Pathology*, 2012, 12(1): e22
- [11] 龙腾飞, 闫芸芳, 彭广兰, 等. 人滋养细胞表面抗原在宫颈病变中表达的研究[J]. *安徽医科大学学报*, 2013, 48(10): 1240-1242
- [12] BrandonLuke L Seagle, Amy L Alexander, Taliya Lantsman, et al. Prognosis and treatment of positive peritoneal cytology in early endometrial cancer: matched cohort analyses from the National Cancer Database[J]. *Ameri J Obstet Gynecol*, 2018, 218(3): 329.e1
- [13] A KH, A KF, A IS, et al. Immunohistochemical analysis using cell block technique leads accurate diagnosis of ovarian malignant lymphoma: A case report[J]. *Inter J Surgery Case Reports*, 2020, 69: 1-4
- [14] Ei Sewedy T, Fornaro M, Alberti S. Cloning of the murine TROP2 gene: conservation of a PIP2-binding sequence in the cytoplasmic domain of TROP-2[J]. *Int J Cancer*, 1998, 75(2): 324-330
- [15] Whitaker M, Larman MG. Calcium and mitosis [J]. *Semin Cell Dev Biol*, 2001, 12(1): 53-58
- [16] Varughese J, Cocco E, Bellone S, et al. Cervical carcinomas overexpress human trophoblast cell-surface marker (Trop-2) and are highly sensitive to immunotherapy with hRS7, a humanized monoclonal anti-Trop-2 antibody[J]. *Am J Obstet Gynecol*, 2011, 205(6): 567.e1-7
- [17] Zeybek B, Manzano A, Bianchi A. et al. Cervical carcinomas that overexpress human trophoblast cell-surface marker (Trop-2) are highly sensitive to the antibody-drug conjugate sacituzumab govitecan [J]. *Sci Rep*, 2020, 10: e973
- [18] Bignotti E, Todeschini P, Calza S, et al. Trop-2 overexpression as an independent marker for poor overall survival in ovarian carcinoma patients[J]. *Eur J Cancer*, 2020, 46(5): 944-953
- [19] Raji R, Guzzo F, Carrara L, et al. Uterine and ovarian carcinosarcomas overexpressing Trop-2 are sensitive to hRS7, a humanized anti-Trop-2 antibody[J]. *J Exp Clin Cancer Res*, 2011, 30: e106
- [20] Goldenbeg DM, Cardillo TM, Govindan SV, et al. Trop-2 is a novel target for solid cancer therapy with sacituzumabgovitecan (IMMU-132), an antibody-drug conjugate (ADC)[J]. *Oncotarget*, 2015, 6(26): 22496-22512
- [21] Lipinski M, Parks DR, Rouse RV, et al. Human trophoblast cell-surface antigens defined by monoclonal antibodies [J]. *Proc Natl Acad Sci USA*, 1981, 78(8): 5147-5150
- [22] 仇金荣, 唐奇, 林红, 等. 人滋养层细胞表面抗原 2 在人胰腺癌中的表达及其临床意义[J]. *中华医学杂志*, 2011, 91(2): 103-106
- [23] Saif, Zaman, Hassan, et al. Targeting Trop-2 in solid tumors: future prospects[J]. *Oncotargets Therapy*, 2019, 12: 1781-1789
- [24] Zhao W, Ding G, Wen J, et al. Correlation between Trop2 and amphiregulin coexpression and overall survival in gastric cancer[J]. *Cancer Medicine*, 2017, 6(5): 994-1001
- [25] Shunjin C, Xiaoying W, Jintian H, et al. Expression of Trop2 in nasopharyngeal carcinoma and its relationship with microvascular density[J]. *J Guangxi Medical University*, 2019, 36(6): 939-943
- [26] 张健, 马海, 杨柳, 等. Trop-2 基因在肝癌组织中的表达及对 HepG2 细胞增殖的影响[J]. *癌症进展*, 2019, 17(16): 1902-1906
- [27] Cubas R, Zhang S, Li M, et al. Trop2 expression contributes to tumor pathogenesis by activating the ERK MAPK pathway [J]. *Mol Cancer*, 2020, 9: 253-266
- [28] Stoyanova T, Goldstein AS, Cai H, et al. Regulated proteolysis of Trop2 drives epithelial hyperplasia and stem cell self-renewal via β -catenin signaling [J]. *Genes & Development*, 2012, 26 (20): 2271-2285
- [29] Heist RS, Guarino MJ, Masters G, et al. Therapy of Advanced Non-Small-Cell Lung Cancer with an SN-38-Anti-Trop-2 Drug Conjugate, Sacituzumab Govitecan [J]. *J Clin Oncol*, 2017, 35 (24): 2790-2797
- [30] Goldenberg DM, Sharkey RM. Antibody-drug conjugates targeting TROP-2 and incorporating SN-38: A case study of anti-TROP-2 sacituzumab govitecan[J]. *Mabs*, 2019, 11(6): 987-995
- [31] Liu JR, Yang DZ, Yin ZN, et al. A novel human monoclonal Trop2-IgG antibody inhibits ovarian cancer growth in vitro and in vivo[J]. *BiochemBioph Res Co*, 2019, 512: 276-282
- [32] 林宏坚. 靶向抑制人滋养层细胞表面抗原-2 基因表达诱导宫颈癌细胞凋亡的机制研究[J]. *中国妇幼保健*, 2017, 32(16): 3934-3937

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- [24] Huang K, Wu T, Wang B, et al. Biomechanical analysis of cervical range of motion and facet contact force after a novel artificialcervical disc replacement[J]. *Am J Transl Res*, 2019, 11(10): 6699-6700
- [25] Othman YA, Verma R, Qureshi SA. Artificial disc replacement in spine surgery[J]. *Ann Transl Med*, 2019, 7(Suppl 5): S170
- [26] Mehren C, Wuertz-Kozak K, Sauer D, et al. Implant Design and the Anchoring Mechanism Influence the Incidence of Heterotopic Ossification in Cervical Total Disc Replacement at 2-year Follow-up[J]. *Spine (Phila Pa 1976)*, 2019, 44(21): 1471-1480
- [27] Zhao X, Yuan W. Biomechanical analysis of cervical range of motion and facet contact force after a novel artificialcervical disc replacement [J]. *Am J Transl Res*, 2019, 11(5): 3109-3115
- [28] Choi H, Baisden JL, Yoganandan N. A Comparative in vivo Study of Semi-constrained and Unconstrained Cervical Artificial Disc Prostheses[J]. *Mil Med*, 2019, 184(Suppl 1): 637-643
- [29] MacDowall A, Canto Moreira N, Marques C, et al. Artificial disc replacement?versus fusion in patients with cervical degenerative disc disease and radiculopathy: a randomized controlled trial with 5-year outcomes[J]. *J Neurosurg Spine*, 2019, 30(3): 323-331
- [30] Zhu Y, Fang J, Xu G, et al. A hybrid technique for treating multilevel cervical myelopathy: Cervical artificial discreplacement combined with fusion[J]. *Oncol Lett*, 2019, 17(1): 360-364