

口腔外科专题

磁共振引导的咀嚼肌疼痛放松训练疗效评估

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[摘要] 目的·应用磁共振成像（magnetic resonance imaging, MRI）的 Dixon 技术评估非结构错乱的颞下颌关节紊乱病（temporomandibular joint disorder, TMD）的咀嚼肌疼痛患者的肌肉改变，并评价MRI引导的物理手法治疗咀嚼肌疼痛的临床疗效。方法·连续纳入自2021年6月—2022年9月就诊于上海交通大学医学院附属第九人民医院口腔外科、首次诊断为TMD肌筋膜疼痛（不伴有盘移位）的患者共29例。收集到增强MRI的Dixon序列并进行手法治疗的患者9例。治疗前，利用DICOM Viewer工作站获得9例患者的Dixon图像咀嚼肌疼痛区域值（Z1）、同侧非疼痛区域值（Z2）和对侧对应咀嚼肌区域值（Z3）。依据工作站获得的咀嚼肌异常区域，针对性地进行手法放松治疗。治疗后1~4周进行随访，对比治疗前后主动最大张口度（maximum mouth opening, MMO）和疼痛视觉评分（visual analogue scale, VAS）指标，并评价临床疗效。结果·Dixon序列Z2均值为 66.23 ± 32.90 , Z3均值为 66.27 ± 33.87 ; 而咀嚼肌疼痛区域Z1为 131.94 ± 83.99 , 明显高于Z2和Z3。手法治疗在改善MMO、VAS等方面有明显效果，有效率达88.89%。结论·Dixon序列的影像表现与临床主诉的疼痛点具有很大的相关性，MRI引导的手法放松治疗对改善非结构错乱的颞下颌关节紊乱病的张口度和疼痛有明显疗效。

[关键词] 颞下颌关节紊乱病；咀嚼肌；疼痛；磁共振；手法治疗

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Evaluation of clinical effect of manipulation on masticatory muscle pain guided by MRI

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[Abstract] Objective·To assess muscle changes of patients of temporomandibular joint disorder with nonstructural disorder by using the Dixon technique for MRI of masticatory muscle, and evaluate the clinical effect of manipulation on masticatory muscle pain guided by MRI. Methods·A total of 29 patients with TMD masticatory muscle pain (without disc displacement) who were diagnosed for the first time in the Department of Oral Surgery, Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine from June 2021 to September 2022 were included. Among them, 9 cases who were assessed with Dixon technique for MRI of masticatory muscle were collected and treated by manipulation. Before treatment, the DICOM Viewer workstation was used to compare the value of masticatory muscle pain area (Z1), the value of ipsilateral non-pain area (Z2) and the value of contralateral corresponding masticatory muscle area (Z3) in Dixon image. Manipulation therapy was performed according to the area of abnormal threshold. Follow-up was performed for 1-4 weeks after treatment, the active maximum mouth opening (MMO) and Visual Analogue Scale (VAS) before and after treatment were compared, and the value of masticatory muscle pain area in Dixon image after treatment was obtained again to evaluate the clinical efficacy. Results·The mean value of Z2 and Z3 in Dixon was (66.23 ± 32.90) and (66.27 ± 33.87), while Z1 in masticatory muscle pain region was (131.94 ± 83.99), which was significantly higher than Z2 and Z3. Manipulation therapy showed significant improvement in MMO and VAS, and the effective rate was 88.89%. Conclusion·There is a significant correlation between the imaging findings of Dixon technique for MRI and the pain points of the masticatory muscle reported by the clinical complaints. The manipulation therapy guided by Dixon technique for MRI has a significant effect on improving the degree of mouth opening and pain.

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[Key words] temporomandibular joint disorder (TMD); masticatory muscle; pain; magnetic resonance imaging; manipulation

颞下颌关节紊乱病 (temporomandibular joint disorder, TMD) 是指一系列涉及咀嚼肌组织、关节及其相关结构的疼痛性和非疼痛性疾病。关节杂音、疼痛以及张口受限等症状会显著影响患者的日常生活, 其中咀嚼肌疼痛是一类常见的TMD, 也是仅次于慢性下腰痛的第二大常见的肌肉骨骼疾病^[1]。针对咀嚼肌疼痛的保守治疗, 目前有药物治疗^[2-3]、咬合板治疗^[4]、手法治疗^[5-6]或联合治疗等, 但对咀嚼肌疼痛位点的判断多采用触压法, 缺乏特定的影像学评估, 从而使得手法放松治疗多以医师经验为指导, 导致临床疗效不稳定。

磁共振成像 (magnetic resonance image, MRI) 不仅是确定盘髁关系的金标准, 而且可以评估咀嚼肌异常^[7]。MRI Dixon技术又称水-脂分离技术^[8], 即利用脂肪质子和水质子共振频率的差异分离水和脂肪, 从而获得同相位、反相位、水像、脂像的图像。有学者将其应用于呼吸肌^[9]和大腿肌群^[10]的评估。Dixon技术在咀嚼肌方面的研究目前较少。本研究纳入正常盘髁关系的咀嚼肌疼痛患者, 通过 Dixon技术定量评估咀嚼肌疼痛患者的肌肉改变, 指导后续物理手法治疗, 并进行有效性评估。

1 对象与方法

本研究为前瞻性研究, 连续纳入2021年6月—2022年9月就诊于上海交通大学医学院附属第九人民医院口腔外科、首次诊断为TMD肌筋膜疼痛的患者。

纳入标准: ①年龄20~50岁。②根据2014扩展版 Diagnostic Criteria for Temporomandibular Disorders (30), 确诊为咀嚼肌疼痛患者。③颞下颌关节 (temporomandibular joint, TMJ) MRI显示盘髁关系正常。④就诊前2周内未经其他治疗。⑤身体质量指数 (body mass index, BMI) <26。排除标准: ①活动性特异性或非特异性关节炎。②急性创伤。③5年内恶性肿瘤史。④精神、神经性疾病史。

1.1 影像学检查

MRI检查采用德国Siemens Aera 1.5T MRI扫描仪, 增强剂采用国产马根维显 (有效成分为钆喷

酸葡胺)注射液15 mL, 扫描范围为面部。获取 Dixon图像, 根据主诉咀嚼肌的压痛点, 通过DICOM Viewer工作站获得患者的Dixon图像: 咀嚼肌疼痛区域值 (Z1)、同侧非疼痛区域值 (Z2)、对侧对应咀嚼肌区域值 (Z3)。工作站可将咀嚼肌的 Dixon图像在某区域的最大值、最小值以及均值直观显示 (图1)。为了减少误差, 取压痛的咀嚼肌的均值作为参考, 并多次测量 ($n=3$), 取平均值。

1.2 手法治疗

参考MRI显示异常区域, 医师进行放松手法治疗, 方法: ①咬肌: 施加压迫手法, 并要求患者咬紧牙齿或使用动态的纵向或横向按摩以进行放松 (图2)。②颞肌: 用2个手指从颅 (颤) 到尾 (下颌) 方向进行纵向或横向按摩, 进行放松。③翼内肌: 让患者张口, 用示指沿牙弓之间探入末端, 与翼内肌的中部接触, 并进行静力压迫以进行放松。④翼外肌: 将示指沿着口腔前庭, 与上颌骨牙槽突的上半部分平行的方向, 对上颌结节到翼突外侧板施以压迫手法以进行放松。每周治疗3次, 隔天1次, 共计8~10次。由于疼痛主观感受差异, 医师或治疗师手法治疗的力度以轻柔耐受为适度, 以不引发新的疼痛为限, 治疗的时间为每次15 min左右。

1.3 临床疗效评价指标

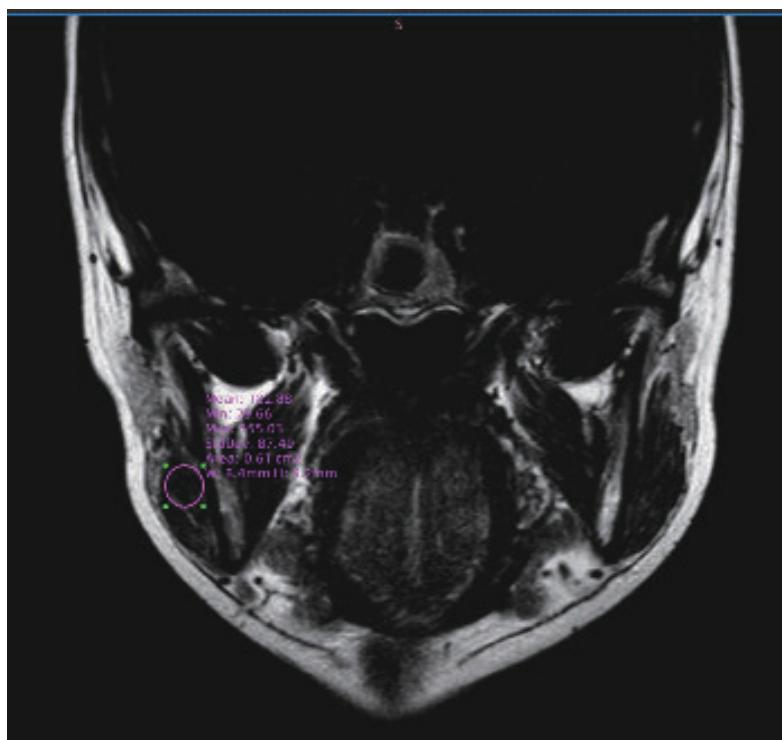
放松训练4周后评价患者切牙颌间主动最大张口度 (maximum mouth opening, MMO) 和疼痛视觉模拟评分 (visual analogue scale, VAS)。

1.3.1 MMO测量 嘱患者主动张口至最大, 用尺测量其上下切牙的距离。

1.3.2 VAS测量 在10 cm的横线上, 一端标为0, 表示无痛; 另一端标为10, 表示剧痛; 中间部分表示不同程度的疼痛。

1.3.3 疗效评估 根据临床症状对患者生活改善的程度分为3级: ①优, MMO≥35 mm, 下颌功能运动与压诊无痛。②良, 治疗后MMO改善≥5 mm, VAS较治疗前的基线水平下降≥2分。③差, MMO与VAS无改善、复发或加重。





Note: Getting the maximum, minimum and average values of the zone at the workstation of DICOM Viewer for 3 times, and then taking the mean values.

图1 咀嚼肌的Dixon图像测量

Fig 1 Measurement of masticatory muscle in the Dixon image

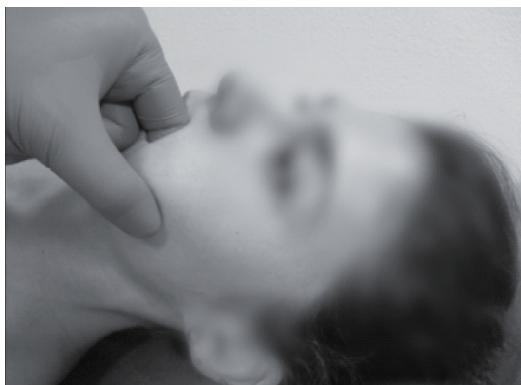


图2 咬肌的手法放松

Fig 2 Manipulation of masseter muscle

1.4 统计学方法

采用SPSS 20.0软件进行统计学分析。符合正态分布的定量资料以 $\bar{x} \pm s$ 表示。

表1 患者疼痛侧的区域值比较($n=9$)

Tab 1 Comparison of the masticatory muscle pain area of patients ($n=9$)

Case No.	Tender area of masticatory muscle	Z		
		Z1	Z2	Z3
1	Left coronoid process	124.00	82.70	83.16
2	Right coronoid process	323.39	112.79	115.87
3	Left maxillary tuberosity	80.63	34.08	34.62
4	Right superior segment of masseter	213.55	108.51	107.27

2 结果

本研究共纳入29例患者，排除患有空间幽闭恐惧症、对增强剂钆喷酸葡胺过敏、地域偏远未能规律治疗的患者，最终纳入9例，均为女性。入组患者平均年龄(39.78 ± 10.44)岁；平均病程(7.89 ± 10.84)个月(1~36个月)，MMO(24.56 ± 5.87)mm，触压诊疼痛位点在咬肌上段2例、咬肌中段1例、咬肌下段2例、颞区1例、冠突区2例和上颌结节区1例。

2.1 患者工作站检查结果

9例患者Z2均值为 66.23 ± 32.90 ，Z3均值为 66.27 ± 33.87 ；咀嚼肌疼痛区域Z1为 131.94 ± 83.99 ，明显高于Z2和Z3。DICOM Viewer工作站测得结果见表1。

Continued Tab

Case No.	Tender area of masticatory muscle	Z		
		Z1	Z2	Z3
5	Left middle segment of masseter	77.07	52.85	52.85
6	Right inferior segment of masseter	154.34	87.40	87.45
7	Right temporal region	115.10	72.45	74.36
8	Left inferior segment of masseter	41.26	25.55	24.49
9	Right superior segment of masseter	58.07	19.73	16.72
Mean value		131.94±83.99	66.23±32.90	66.27±33.87

2.2 患者手法治疗结果

9例患者均进行手法治疗，治疗前后的MMO和VAS结果见表2，疗效优为1例，良为7例，差为1例，有效率为88.89%。

表2 手法治疗前后的MMO和VAS比较

Table 2 Comparison of MMO and VAS before and after manipulation therapy

Case No.	MMO/mm		VAS/score		Efficacy evaluation
	Before	After	Before	After	
1	21	38	6	2	good
2	15	30	7	4	good
3	26	35	7	4	good
4	19	38	6	2	good
5	24	35	5	1	good
6	31	39	4	2	good
7	36	40	3	2	poor
8	25	35	5	2	good
9	24	37	0	0	excellent
Mean value	24.56±5.87	36.33±2.83	4.78±2.10	2.11±1.20	

3 讨论与结论

TMD咀嚼肌疼痛是由多种因素引起的，如咬合干扰、咀嚼肌活动过度、磨牙症、应激以及损伤、不良姿势以及退化因素等。疼痛通常与受影响肌肉的疲劳、张力、炎症或水肿有关。早期对轻微的压力（触诊）较敏感^[11-12]。并且这种疼痛存在性别差异，女性受影响的频率高于男性^[13]。本研究纳入的9例病例均为女性，有显著的性别倾向。

对咀嚼肌疼痛部位的判断多以患者自述和体检压诊决定，存在一定的主观性。MRI不仅是确定TMD盘髁关系的金标准，也经常用于评估肌肉内水分或脂肪含量的变化^[14-15]。特别是T2加权图像可显示脂肪浸润、水肿或炎症导致的脂肪和水分增加。YUTAKA等^[16]发现主诉有压痛的咬肌，其MRI的

T2值明显高于无疼痛侧咬肌，提示压痛患者咬肌的游离水含量较高。在此基础上的Dixon技术利用脂肪和水质子共振频率的差异分离水和脂肪，应用于眼部、头颈部等部位能够获得更高质量的图像。如今 Dixon在骨骼肌中的应用越发广泛，有研究^[17]利用 Dixon技术量化组织个体中脂肪和水的分布，计算肌肉的脂肪分数，以此评估呼吸肌受累程度^[9]。有研究^[10]用此评估控制性肢端肥大症患者每条大腿肌肉中脂肪分数的百分比，结果显示患者的肌间脂肪分数高于对照组。

本研究首次应用 Dixon技术获得咀嚼肌图像，通过工作站软件分析得到压痛侧咀嚼肌病灶区域的平均值，并对比其非病灶区和健侧咀嚼肌，发现其值明显高于后两者，而非病灶区和健侧咀嚼肌均值几乎相当。

针对咀嚼肌疼痛的治疗有药物治疗、咬合板治疗、手法治疗等。其中药物可以改善炎症，减轻疼痛。咬合板能缓解口颌肌肉功能亢进和高张力状态。上述2种方法的缺点是无法针对具体的疼痛肌肉病灶。而手法可以针对疼痛位点进行快速有效的治疗。有研究^[18]表明，手法治疗在治疗TMD方面显示出较好的临床效果。另有报道^[19]称手法治疗和运动干预在改善TMD患者的症状和功能方面有积极作用。针对TMD咀嚼肌疼痛部位，手法可以通过激活低阈值A β 纤维来调节疼痛^[20-21]。URBAŃSKI等^[22]将60名肌筋膜疼痛患者分为等长收缩后放松治疗组和肌筋膜松解治疗组，2组均进行了10次治疗，评估咀嚼肌疼痛等，结果显示咀嚼肌的疼痛显著下降。然而，以上研究由于治疗前缺乏影像学指导，只能依照患者主诉和临床触压诊获得主观疼痛程度和模糊范围。现针对咀嚼肌疼痛的 Dixon技术可显示异常肌肉区域，有利于疼痛区域主诉不明确或压诊无异常的患者，可辅助医师确定异常肌肉的位点，进行临床治疗。

咀嚼肌疼痛会影响日常生活但程度不一。本研究初始就医的29名患者中，仅9名因疼痛导致张口受限



的患者愿意选择手法治疗，其他无明显张口受限的患者更愿意选择家庭治疗，如自我热敷或其他治疗，因此无法对比2组的长期预后。希望将来可以开展更进一步的研究。

利益冲突声明/Conflict of Interests

所有作者声明不存在利益冲突

All authors disclose no relevant conflict of interests.

伦理批准和知情同意/Ethics Approval and Patient Consent

本研究涉及的所有实验均已通过上海交通大学医学院附属第九人民医院医学伦理委员会的审核批准（文件号SH9H-20190T316-2）。所有试验过程均遵照国家卫计委和国家药品监督管理总局有关法规进行。受试对象或其亲属已经签署知情同意书。

All experimental protocols in this study were reviewed and approved by Shanghai Ninth People's Hospital, Shanghai Jiao Tong University School of Medicine, (Approval Letter SH9H-20190T316-2, dated 15/

01/2020), and all experimental protocols were carried out by following the guidelines of National Health and Family Planning Commission of the People's Republic of China and National Medical Products Administration. Consent letters have been signed by the research participants or their relatives.

作者贡献/Authors' Contributions

杨海霞、陈敏洁参与了试验设计；杨海霞参与了论文的写作和修改；王博成参与了影像数据分析，徐丽丽参与了临床数据分析。所有作者均阅读并同意了最终稿件的提交。

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