



中国科学引文数据库(CSCD)收录期刊
中文核心期刊要目总览收录期刊
中国科技论文统计源期刊

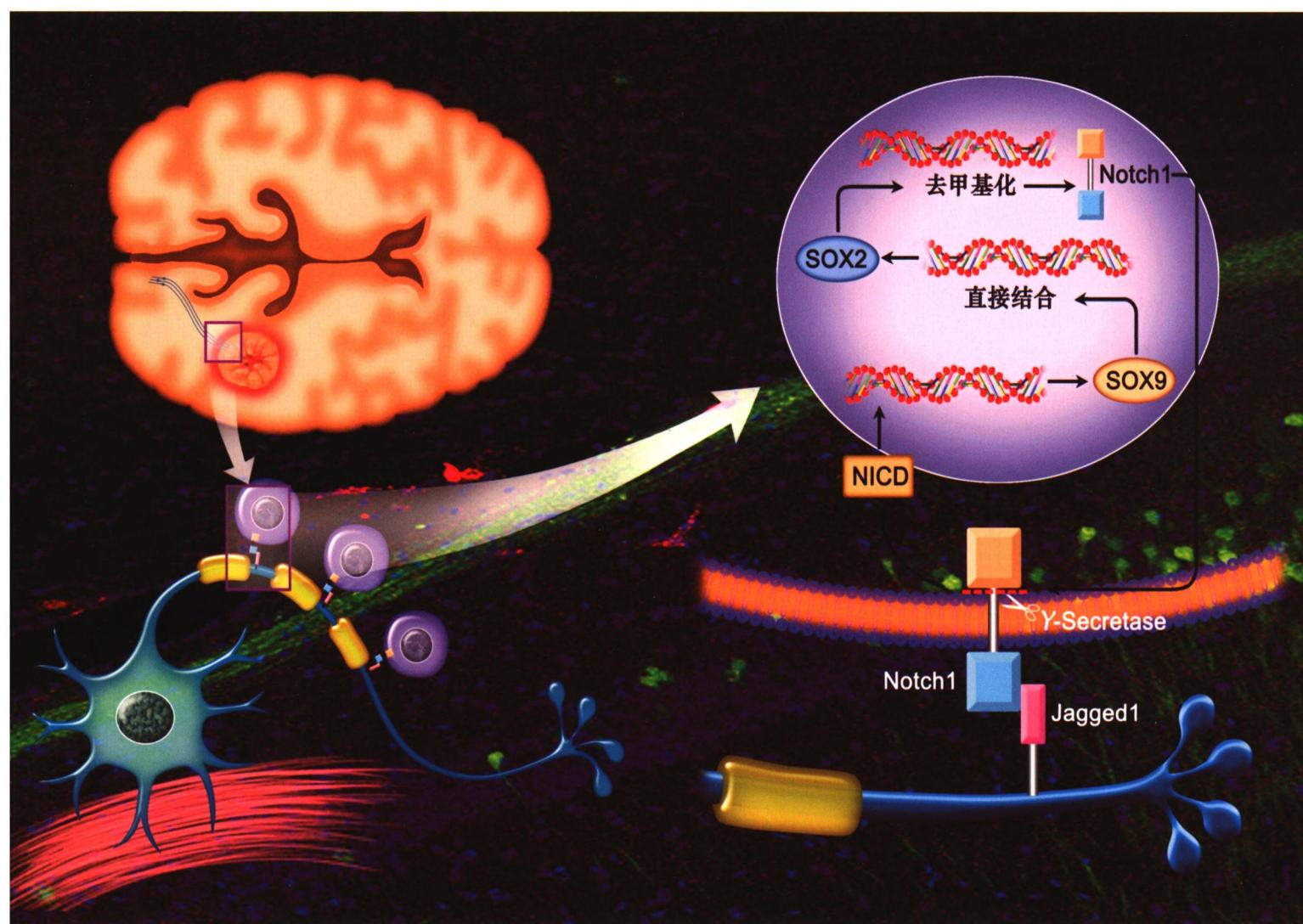


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I-1126/R
I DYXUE8

Q K 2 0 2 5 1 7 7

第三軍醫大學學報

Journal of Third Military Medical University



2019年度重庆市出版专项资金资助期刊

2020年6月15日 第42卷 第11期

Vol.42, No.11 Jun. 15 2020

ISSN 1000-5404



第三軍醫大學學報

半月刊 1979年1月创刊 第42卷 第11期 总第586期 2020年6月15日出版

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印 刷

普天印务(重庆)股份有限公司
401147, 重庆市渝北区黄泥磅工业园区1幢

发 行(公开)

国内:重庆市报刊发行局
国外:中国国际图书贸易总公司
(北京399信箱, 100044)
代号:SM6529

订 购

全国各地邮政局
邮发代号:78-91

邮 购

第三军医大学学报编辑部

定 价

国内:¥30.00/期, ¥720.00/年
国外:\$30.00/期, \$720.00/年

中国标准连续出版物号

ISSN 1000-5404
CN 50-1126/R

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封面故事:胶质瘤是中枢神经系统中最常见的原发性恶性肿瘤,治疗棘手。沿着白质神经纤维分布的早期侵袭性生长,是恶性胶质瘤突出的病理学特征之一,也是其治疗失败的主要原因。陆军军医大学第一附属医院全军临床病理学研究所、干细胞与再生医学科卞修武院士和余时沧教授团队研究发现,在脑胶质瘤侵袭前沿处,CD133+Notch1+胶质瘤干细胞(GSC)临近发生脱髓鞘的白质纤维分布,而这些神经纤维则表达Notch的配体Jagged1。Notch1通过转录因子Sox9促进SOX2的表达,而SOX2的上调则反过来通过降低NOTCH1启动子甲基化以增强其转录,揭示了NOTCH1-SOX2正反馈环路对GSC沿白质束侵袭性生长的调控作用。临床病理研究进一步证明,该正反馈环路的检测具有预测患者生存时间的意义。这些发现丰富了我们对恶性胶质瘤侵袭性生长和复发机制的认识,也为靶向治疗提供了新思路。研究成果被《Nat Neurosci》作为封面论文发表[WANG J, XU S L, DUAN J J, et al. Invasion of white matter tracts by glioma stem cells is regulated by a NOTCH1-SOX2 positive-feedback loop. Nat Neurosci, 2019, 22(1):91–105],并被《Cancer Stem Cell News》作为Top Story评介。论文发表6个月后成为ESI前1%高被引用论文。

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[期刊基本参数] CN 50-1126/R * 1979 * s * 16 * 96 * zh+en * P * ¥30.00 * 1500 * 14 * 2020-06

Journal of Third Military Medical University

DI-SAN JUNYI DAXUE XUEBAO

ACTA ACADEMIAE MEDICINAE MILITARIS TERTIAE

Semimonthly Established in January 1979 Vol. 42, No. 11(Gen. No. 586) June 15, 2020

Responsible Institution

Army Medical University
(Third Military Medical University)

Sponsor

Army Medical University
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http://aammt.tmmu.edu.cn

Printing

Chongqing Putian Printing Co., Ltd
No. 1 Building, Huangnibang Industrial Park,
Yubei District, Chongqing, 401147, China

Open Publishing

Domestic Distributor
Local Post Offices
Code No. 78-91

Overseas Distributor
China International
Book Trading Corporation
P. O. Box 399, Beijing, 100044, China
Code No. SM6529

Mail Order

Editorial Office of *Journal of Third Military Medical University*

Price

Domestic: ¥ 30.00/issue
Overseas: \$ 30.00/issue

CSSN

ISSN 1000-5404
CN 50-1126/R

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