

硝普钠及其光解产物对日本晴水稻幼苗生长和 5 种激素标记基因表达的影响

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摘要:以粳稻品种日本晴水稻为材料,检测硝普钠及其光解产物 KNO_2 、 $\text{K}_4\text{Fe}(\text{CN})_6$ 对幼苗生长的影响,并采用 qRT-PCR 技术检测了 5 种植物激素标记基因在经上述处理后在幼苗根中的表达水平.结果表明 SNP 能够显著抑制水稻幼苗的根长和株高;SNP 对根生长的抑制主要是通过其光解产物 $\text{K}_4\text{Fe}(\text{CN})_6$ 实现的. SNP 和 $\text{K}_4\text{Fe}(\text{CN})_6$ 处理都能够抑制生长素、细胞分裂素、脱落酸和赤霉素 4 种激素标志基因在水稻根中的表达,但是 SNP 处理可以抑制一氧化氮标志基因 OsNOA1 的表达,而 $\text{K}_4\text{Fe}(\text{CN})_6$ 则上调该基因的表达.

关键词:硝普钠;水稻;幼苗;植物激素;基因表达

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在高等植物中,根是重要的营养器官之一,具有吸收、储藏和运输营养物质的功能.更重要的是,根部产生的植物激素影响着地上部分的生长和发育.以往的研究表明,多种植物激素参与调控根的生长发育,其中生长素、细胞分裂素及赤霉素均能促进根的生长^[1].近期的研究发现,气体信号分子一氧化氮(nitric oxide, NO)也参与植物根的发育调节过程^[2~3],其作用方式涉及植物激素信号传导网络,但是机制尚不清楚.因此,对植物根生长发育与 NO 等激素信号关系的研究有待深入.

水稻是重要的粮食作物和模式植物,已有的研究表明 NO 影响水稻根的形态建成,包括水稻侧根的形成^[4].硝普钠(sodium nitroprusside, SNP)作为研究常用的 NO 供体,经过光解后会产生 3 种产物即 NO, KNO_2 和 $\text{K}_4\text{Fe}(\text{CN})_6$,尽管已有 SNP 对水稻生长发育影响的检测^[5~7],但是 SNP 及其光解产物对水稻生长发育的系统研究尚未见报道.为此本文以粳稻品种日本晴为材料,以外源 NO 供体 SNP 以及光解产物 KNO_2 , $\text{K}_4\text{Fe}(\text{CN})_6$ 处理幼苗,并采用 qRT-PCR 检测在上述处理条件下水稻植物激素标记基因的表达水平,旨在检测其对水稻幼苗生长的影响,并明确 SNP 处理对生长素、细胞分裂素等植物激素水平的影响,为进一步鉴定 NO 在水稻生长发育中的功能奠定基础.

1 材料和方法

1.1 供试材料

水稻粳稻品种日本晴(*Oryza sativa L. japonica. cv. Nipponbare*)由本实验室保存.

1.2 供试试剂

硝普钠, $\text{K}_4\text{Fe}(\text{CN})_6$, KNO_2 购于北京鼎国生物有限公司;总 RNA 提取试剂 TRNzol, DNaseI 酶, Prime-Script RT Master Mix 购于大连宝生物有限公司;SYBR Green Master Mix 购于南京诺唯赞生物技术有限公司.

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1.3 方法

1.3.1 水稻幼苗的培养和处理

取水稻种子适量用75%的酒精消毒,无菌水冲洗,室温浸泡1 h后,置于人工气候箱28 ℃黑暗培养催芽,待种子发芽后转移至Yoshida完全营养液^[8]进行水培,培养温度28 ℃,光照强度为16 000 lx,光照时间为12 h/d.外源NO处理采用SNP添加到Yoshida完全营养液中,终浓度分别为0、0.02、0.04、0.06、0.08 mM,每4 d更换一次营养液,培养12 d取样,测量不同处理组水稻的根长和株高,每组样品重复3次.

1.3.2 总RNA的提取

取生长至12 d的水稻幼苗根,液氮研磨后,按照TRNzol说明书的要求提取总RNA,并采用超微量光度计检测RNA样品浓度及质量.

1.3.3 反转录

按照宝生物反转录试剂盒说明书的要求进行反转录,制备cDNA反应体系如下:5×PrimeScript RT Master Mix(2 μL)、Total RNA(≤500 ng)、RNase Free H₂O(补至10 μL).将上述反应体系短暂离心混匀后,在37 ℃保温15 min,然后85 ℃ 5 s终止反应.

1.3.4 实时定量PCR

用以上制备的cDNA为模板,内参基因选用*OsAct1*,其他检测基因引物见表1.按照SYBR Green Master Mix说明书进行配制20 μL体系:2×SYBR Green Master Mix(10 μL)、上游引物(0.4 μL)、下游引物(0.4 μL)、ROX Reference Dye II (50×)(0.4 μL)、cDNA(2 μL)和RNase Free H₂O(6.8 μL).采用两步法进行PCR扩增:95 ℃预变性,5 min;PCR反应95 ℃,10 s;然后60 ℃,30 s,并收集数据.设置未处理水稻为对照组,每个样品平行重复3次.

表1 实时定量PCR引物一览表

引物序号	引物序列 (5'-3')	基因名称	基因特征
P1	TCTTCCAGCCTTCCTTCA	<i>OsAct1</i>	肌动蛋白基因 ^[9]
P2	ATCCACGTCGCACTTCAT		
P3	ATAATCCTCGCCACAGGCTACA	<i>OsYUCCA1</i>	IAA 标记基因 ^[10]
P4	GCAGTGCAGACAGAAAGAAAA		
P5	TACGAGTGCTGCTTCCTCTGGG	<i>OsIPT</i>	CTK 标记基因 ^[11]
P6	AGATGCCCTGGAGTAGTCGGT		
P7	ATCGGCTGGAGATGAAGAGGG	<i>OsGA20ox1</i>	GA 标记基因 ^[12]
P8	GCGGCTCATCTCGTGCCAGT		
P9	ACTGTGTCGAGCCTGTG	<i>OsNCED2</i>	ABA 标记基因 ^[13]
P10	GGAAGGTCAGATTCGCATAG		
P11	TGAAGGATGGGTTGGTCTG	<i>OsNOA1</i>	NO 标记基因 ^[14]
P12	ACCCACCCGAGACCAGAAA		

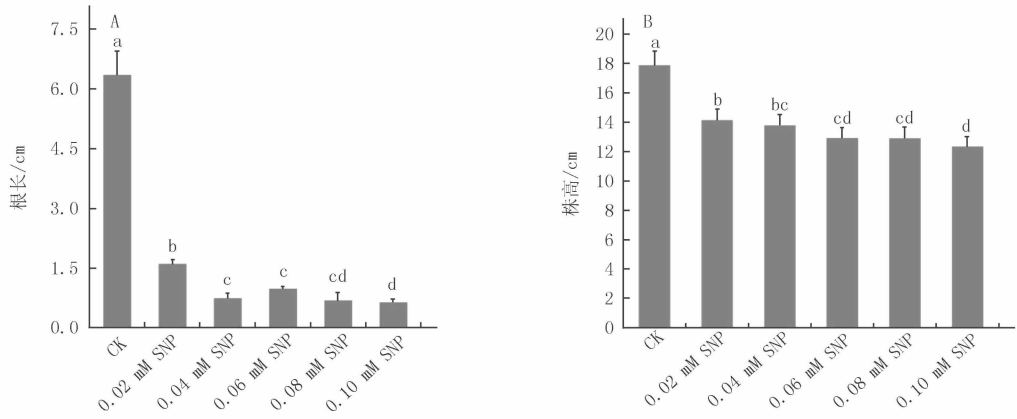
1.4 数据分析

采用DPS数据处理软件对水稻根长和株高数据进行方差分析,实时定量PCR的数据采用 $2^{-\Delta\Delta C_t}$ 法进行处理分析.数据分析结果用Origin 8.5软件绘制图表.

2 结果与分析

2.1 SNP对水稻幼苗生长的影响

采用不同浓度的SNP处理水稻幼苗,统计根长和株高,发现与未处理的水稻相比,不同浓度的SNP对幼苗的根和地上部分的生长均有抑制作用(图1).结果显示,随着SNP处理浓度的升高,水稻的根长和株高都明显的降低,统计学分析显示SNP对地上部分生长的抑制程度不如对根的抑制显著.与对照相比,随着SNP浓度的增加,根长降低幅度为88%~90%.



“a, b, c, d”代表 $P < 0.05$ 显著分析

图1 不同浓度SNP对水稻幼苗根长(A)和株高(B)的影响

2.2 SNP及其光解产物对水稻幼苗生长的抑制效应分析

为了解 SNP 抑制根生长的作用机制,进一步采用 SNP 及其光解产物 KNO_2 和 $\text{K}_4\text{Fe}(\text{CN})_6$ 分别处理幼苗,与对照根长进行比较,结果显示,SNP 和 $\text{K}_4\text{Fe}(\text{CN})_6$ 处理后,根的长度与对照相比显著降低,SNP 处理根长略微短于 $\text{K}_4\text{Fe}(\text{CN})_6$ 处理,而 KNO_2 处理后,水稻幼根长度与对照差别不大,甚至略有增加(图 2,图 3A).此外,对株高的统计也显示出类似的结果(图 3B),表明 SNP 对根生长的抑制效应中, $\text{K}_4\text{Fe}(\text{CN})_6$ 发挥主要作用.

2.3 SNP及其光解产物对激素标记基因在水稻幼苗根中表达的影响

分别选择 5 种植物激素生长素 (IAA)、细胞分裂素 (CTK)、赤霉素 (GA)、脱落酸 (ABA) 和 NO 的标记基因 OsYUCCA1 , OsIPT , OsGA20ox1 , OsNCED2 和 OsNOA1 , 采用 qRT-PCR 检测上述激素标记基因在经 SNP 及其光解产物处理后的水稻根中表达特征,结果显示,SNP 处理后,5 种基因表达都下调;而 $\text{K}_4\text{Fe}(\text{CN})_6$ 处理后,除 OsNOA1 基因表达上调外(图 4E),其余 4 种基因表达都下调;另外 KNO_2 处理后,除了 OsIPT (图 4B)和 OsNCED2 (图 4C)这两种基因表达与对照组没有显著差异外,其他 3 种标记基因表达水平都显著上调.

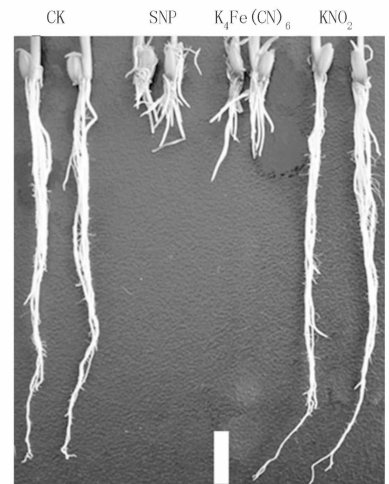


图2 SNP及其光解产物对水稻幼苗根长的影响
处理浓度为0.06 mM, Bar=1 cm

3 讨论

NO 已被证明是普遍存在于动植物中的气体信号分子.已有的报道表明,NO 参与植物多种生长发育过程的调节,例如植物根的形态建成^[15],植物细胞的程序性死亡^[16],气孔的关闭^[17]等.本研究用 SNP 处理水稻幼苗,结果显示水稻幼苗根系伸长生长被抑制,根的形态与处理前相比也发生明显的改变,侧根呈现短粗特征.SNP 处理后,水稻株高也明显的降低,可能是由于 SNP 对根生长的抑制,影响了根对营养的吸收,可能还引起植物激素水平降低,导致水稻株高的降低.

由于 SNP 光解时生成 KNO_2 , $\text{K}_4\text{Fe}(\text{CN})_6$ 和 NO 3 种物质,为此,本研究采用 SNP 及其光解产物分别处理水稻进一步研究 SNP 的作用机制.结果显示,SNP 与 $\text{K}_4\text{Fe}(\text{CN})_6$ 都能够抑制水稻根的生长,且两种处理水稻幼根的表型相似,提示在上述条件下 SNP 对根生长的抑制效应是 NO 和 $\text{K}_4\text{Fe}(\text{CN})_6$ 共同作用的结果,且以 $\text{K}_4\text{Fe}(\text{CN})_6$ 为主.

已有研究表明 NO 与多种植物激素信号途径相关,比如通过参与 IAA^[18-19],CTK^[20],ABA^[21],乙

烯^[22],水杨酸^[23]等植物激素之间的信号转导网络来调节植物生理过程. 因此在 SNP 及其光解产物处理的条件下,水稻幼苗生长受抑制的过程中,其他植物激素水平是否发生变化值得研究. 为此,本研究选取了5种水稻植物激素的标记基因,进行 qRT-PCR 检测,发现 SNP 和 $K_4Fe(CN)_6$ 处理都能下调 IAA,CTK 和 GA 标记基因在根中的表达水平,提示可能会因此导致上述激素含量的降低,说明外源 NO 和 $K_4Fe(CN)_6$ 可能通过降低水稻幼苗根中 IAA,CTK,GA 的水平,达到对根生长的抑制效应,在功能上与上述激素发生拮抗作用. SNP 处理的效应显著于 $K_4Fe(CN)_6$ 处理,也进一步证明 SNP 的复合效应. 另外,值得注意的是, KNO_2 和 $K_4Fe(CN)_6$ 处理都明显提高了 *OsNOA1* 基因的表达水平,推测由此导致水稻内源 NO 水平的提高,提示 SNP 不仅提供外源 NO,也影响水稻内源 NO 水平. 本研究为探究 NO 在抑制水稻根生长中的作用机制以及其与 IAA,CTK 等5种植物激素的关系提供了线索,为揭示水稻幼苗根发育过程中复杂的信号网络奠定理论基础.

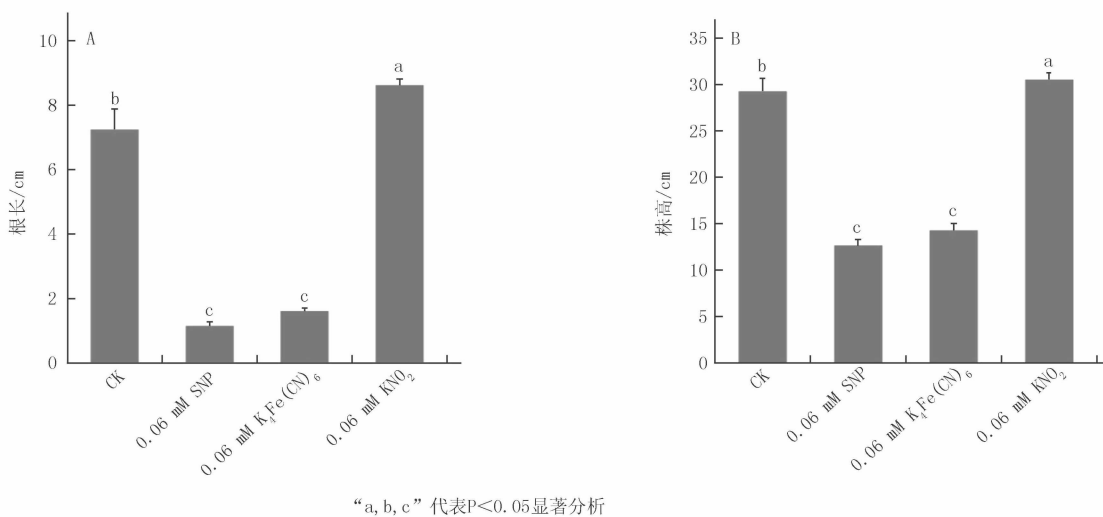
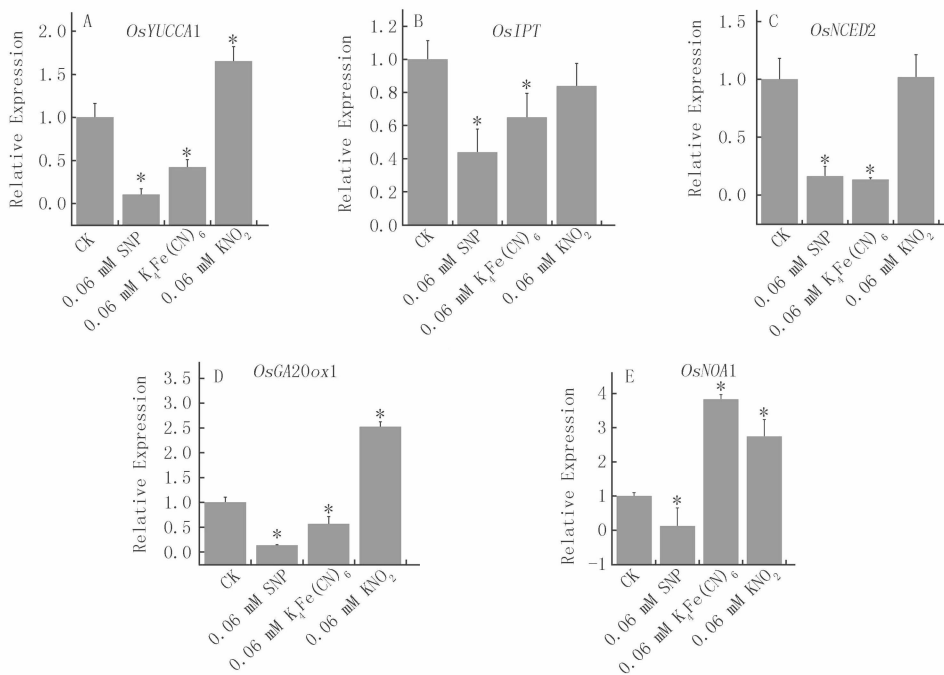


图3 SNP 及其光解产物对水稻幼苗根长(A)和株高(B)的统计学分析



*代表 $P < 0.05$ 显著分析

图4 SNP 及其光解产物对水稻5种植物激素标志基因在水稻根中表达的影响

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Extraction and Identification of Host-Plant Volatiles of *Acer mono* and EAG Responses of *Anoplophora nobilis* to the Primary Compounds of *A. mono* Volatiles

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Abstract: To explore the most appropriate hosts of nutrient supplement and oviposition for *Anoplophora nobilis* adults in Yili Prefecture, the volatiles composition of *Acer mono* were analyzed with GC-MS, and the electrophysiological activities of adults towards main volatiles were tested by EAG responses. The results showed that *A. nobilis* preferred fresh twigs of *A. mono* for diet supplement and carve twigs to oviposit in mixed plant area. Thirty-four volatiles were identified from shoots and leaves of *A. mono*, and terpenoids and esters were the major components, with the relative concentration up to 70.49% and 23.94%, respectively. (*Z*)- β -ocimene, sabinene, β -caryophyllene, α -phellandrene and cis-3-Hexenyl acetate represented a higher proportion in all components, with the relative concentration of 21.07%, 19.59%, 7.35%, 13.58% and 12.99%, respectively. With the concentration of 10 mg \cdot mL⁻¹, the EAG responses of female adults to β -caryophyllene was significantly higher than the other volatile compounds, and the male showed the most intensively EAG responses to ethyl acetate and α -pinene. The EAG responses of male adults to ethyl acetate and α -pinene were significantly stronger than female, while the female adults had more strongly EAG responses to β -caryophyllene and cis-3-Hexenyl-2-methyl butyrate compared to males. This study will contribute to further research of chemical fingerprinting relating to host location of *A. nobilis*, therefore have a certain practical significance to development plant source attractants.

Keywords: *Anoplophora nobilis*; host selection; plant volatiles; electroantennogram responses; olfactory communication

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Effects of SNP and Its Photolysis Products on the Seedlings Growth of Rice and Expression of Marker Genes for Five Plant Hormones

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Abstract: Taking japonica cultivar Nipponbare rice as materials, the effects of sodium nitroprusside and its photolysis products KNO₂, K₄Fe(CN)₆ on the rice seedlings growth were analyzed, and marker genes for five different plant hormones in rice seedlings roots under the above treatments were detected by qRT-PCR in the study. The results showed that SNP can obviously inhibit the root length and plant height of the two rice seedlings, which implement mainly through its photolysis products K₄Fe(CN)₆. Moreover, the expression of the marker genes of auxin, cytokinins, abscisic acid and gibberellic acid were inhibited after the treatment of SNP and K₄Fe(CN)₆ in rice roots, but the marker gene of nitric oxide was inhibited by SNP, induced by K₄Fe(CN)₆.

Keywords: sodium nitroprusside; rice; seedling; plant hormone; gene expression

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