

# 杜梨组培生根过程中多胺、内源激素及相关氧化酶活性的变化

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**摘要:**【目的】探究杜梨组培苗生根诱导过程中内源激素、多胺类物质及相关氧化酶活性变化的生理响应机制。【方法】以长势相近的杜梨继代苗为试材,筛选优化了生根培养基配方,分别在生根诱导0、3、7、10、15 d后,测定分析了基部茎段多胺类物质、内源激素含量及关联酶活性。【结果】(1)激素配比2.0 mg·L<sup>-1</sup>IAA+0.5 mg·L<sup>-1</sup>IBA的生根效果良好,IAA和IBA两种激素共同诱导杜梨生根效果要优于单一激素处理;(2)在生根诱导0~15 d内,精胺(Spm)和亚精胺(Spd)含量呈现下降→升高→下降→升高的变化趋势,腐胺(Put)含量则呈现出下降→上升的变化趋势;内源激素IAA、ABA含量均呈现先下降后升高的趋势,最低值出现在第7天,GA含量也呈现先下降后上升的变化趋势,最低值出现在第3天,ZR含量呈现出下降→升高→下降的变化趋势;两种关联酶IAAO和PPO酶活性均呈现增高→降低→增高的趋势;(3)Spd与IAA、ABA、IAA/ABA IAA/ZR及IAAO酶活性均存在极显著正相关关系( $p < 0.01$ ),IAA与Spd、GA、ABA及IAA/ABA、IAA/ZR、PPO酶活性在 $\alpha = 0.01$ 水平达到极显著正相关。【结论】杜梨较适宜的生根培养基为:1/2MS+2.0 mg·L<sup>-1</sup>IAA+0.5 mg·L<sup>-1</sup>IBA+7.5 g·L<sup>-1</sup>琼脂+25.0 g·L<sup>-1</sup>蔗糖;IAA/ABA、IAA/ZR比值增大更有利于杜梨不定根的发生;多胺类物质中Spd含量与杜梨不定根发生紧密相关;生根诱导后期,GA含量增加对杜梨不定根发生并无明显的抑制作用。

关键词:杜梨;组织培养;多胺;内源激素;关联酶;相关性

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## Dynamic changes in polyamines, endogenous hormones and oxidase activities during rooting of *in vitro* plantlets of *Pyrus betulifolia* Bunge

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**Abstract:**【Objective】*Pyrus betulifolia* Bunge is widely used as a pear rootstock in China. Propagation with seeds has large genetic variation and poor consistency. Difficulty in rooting limits propagation by traditional vegetative propagation methods such as layering and cutting. With the increasingly perfecting of *in vitro* propagation technology, virus-free seedlings can be produced by micropropagation method. Root induction is one of the important processes in *in vitro* propagation. The transplanting survival rate and seedling quality are affected by rooting efficiency. There are many internal and external factors affecting the rooting process, including endogenous hormones, polyamines, related oxidases and exogenous substances such as ethylene and salicylic acid. The study aimed at revealing the roles of en-

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dogenous hormones, polyamines and related oxidases in root induction of tissue-cultured seedlings.

**【Methods】** Subcultured plantlets of *Pyrus betulifolia* Bunge were cultured on rooting medium with 1/2MS, 7.5 g·L<sup>-1</sup> agar, 25.0 g·L<sup>-1</sup> sucrose and different ratios of hormones. A total of 24 treatments using five different concentrations (0, 0.2, 0.5, 1 and 2 mg·L<sup>-1</sup>) of IAA and IBA were assessed in terms of rooting percentages at 40 days after transfer to the media. Each treatment consisted of 6 bottles each comprising 5 subcultured plantlets. Uniform subcultured plantlets were selected to be cultured on the optimized rooting medium. Approximately 500 plantlets were used to monitor the dynamic changes in polyamines, endogenous hormones and related enzyme activities during root regeneration. All of the cultures were incubated at 25 °C under a 16 h light/8 h dark cycle and 2 000-3 000 lx light intensity. The basal stem cuttings were taken 0, 3, 7, 10 and 15 days after transfer to media. Polyamines, endogenous hormones and related enzyme activities were determined. **【Results】** The treatment of 2.0 mg·L<sup>-1</sup> IAA + 0.5 mg·L<sup>-1</sup> IBA increased rooting rate and the percentage of plantlets with ≥3 roots compared with the other treatments. The rooting rate was 86.7% and the percentage of plantlets with ≥3 roots was 53.3%. The combination of IAA and IBA was better than single hormone. Callus was observed between the 7th and 10th day, growing fast from the 10th to 15th day. The roots generated 20 days after transfer to media. The contents of three polyamines declined after 3 days in the root induction medium. The content of spermine (Spm) and spermidine (Spd) fluctuated during root induction period. The content of putrescine declined in the first three days and rose between the 3th day and the 15th day. The contents of IAA and IBA decreased firstly and then increased, and were the lowest at the 7th day. The content of GA was the lowest at the 3rd day and increased rapidly from the 3th to the 10th day, but then the increasing speed slowed down. The variation in ZR content followed a pattern of "decrease→increase→decrease", lowest at the 7th day. The ratio of IAA/ABA was relatively stable during the first seven days, and increased from the 7th day to the 15th day. The ratio of IAA/ZR decreased slightly within the first 3 days and then increased from the 3th day to the 15th day. The activity of IAAO remained stable from the 7th day to the 10th day, and increased rapidly from the 10th to the 15th day, indicating that decomposition of IAA was accelerated during this period. There was a significant negative correlation between Spm and ZR contents ( $p < 0.01$ ,  $r = -0.898$ ). Spd content had significant positive correlations ( $p < 0.01$ ) with the contents of IAA and ABA, the ratios of IAA/ABA and IAA/ZR and the activity of IAAO. Put content was positively correlated with GA ( $p < 0.01$ ,  $r = 0.646$ ), IAA and ABA ( $p < 0.05$ ). IAAO activity was significantly positively correlated with PPO activity ( $r = 0.843$ ), Spd content ( $r = 0.643$ ) and IAA/ZR ratio ( $r = 0.662$ ). PPO enzyme activity was significantly positively correlated with IAA and ABA contents and the ratio of IAA/ZR ( $p < 0.01$ ). The content of IAA had moderate correlations with the contents of Spd, GA and ABA, PPO activity and the ratios of IAA/ZR and IAA/ABA. The content of ABA had moderate correlations with the contents of IAA, Spd and GA and the ratios of IAA/ZR and IAA/ABA. The results showed that Spd, IAA and ABA played important regulation roles in rooting process.

**【Conclusion】** The suitable rooting medium for *Pyrus betulifolia* Bunge was 1/2MS + 2.0 mg·L<sup>-1</sup> IAA + 0.5 mg·L<sup>-1</sup> IBA + 7.5 g·L<sup>-1</sup> agar + 25.0 g·L<sup>-1</sup> sucrose. The increase in IAA/ABA and IAA/ZR ratios induced adventitious roots formation. Spd might be closely related to the adventitious root formation. In the late stage of root induction, the increase in GA content had no significant effect on the adventitious root formation.

**Key words:** *Pyrus betulifolia* Bunge; Tissue culture; Polyamine; Endogenous hormone; Associated enzyme; Correlation

植物不定根的发生受多种内外因素的影响。已有研究表明:植物不定根发生与内源激素水平紧密相关,其中生长素对不定根发生的促进作用最大<sup>[1-2]</sup>,其他外源物质也是通过影响激素的代谢或信号传导而影响不定根的发生,如外源水杨酸、乙烯等<sup>[3-4]</sup>;无论低浓度还是高浓度的赤霉素(GA)对不定根的发生都有抑制作用;低浓度的脱落酸(ABA)可以促进生根,高浓度的则抑制生根<sup>[5]</sup>。除内源激素影响外,多胺是普遍存在于植物体内的另一种植物生长发育调节物质,以精胺(spermine, Spm)、亚精胺(spermidine, Spd)、腐胺(putrescine, Put)3种最为常见。国内外许多学者发现,亚精胺对植物的生长发育有重要影响,可以促进根系生长,增加不定根数量<sup>[6-8]</sup>。辛蓓<sup>[9]</sup>研究也表明:多胺也可以促进苹果砧木M26根系形成,Faivre-Rampant等<sup>[10]</sup>发现在无生长素培养基中添加多胺能促进烟草不定根的发生。

吲哚乙酸氧化酶(IAAO)和多酚氧化酶(PPO)是植物不定根发生过程中重要的关联酶,两类酶活性高低与植物生根难易程度紧密相关<sup>[11]</sup>。相关研究发现,IAAO酶活性越高不定根的发生就越困难,而PPO酶活性越高越易生根,因为PPO酶可以催化酚类物质与IAA结合成一种“IAA-酚酸复合物”,此复合物对植物不定根的发生具有促进作用<sup>[12]</sup>。

杜梨作为梨砧木应用广泛,但通过压条或扦插等常规无性繁殖方式不易生根,而利用组培技术快速繁育脱毒苗木在实际生产中应用越来越广泛。生根诱导是组培快繁技术体系中的重要环节,生根质量的好坏直接影响苗木田间移栽成活率和苗木质量,因此探究如何调控组培苗快速生根,提高生根数量及质量就显得十分重要。目前,前人研究多集中在杜梨组培快繁体系建立方面,并且取得了较多的科研进展<sup>[13-15]</sup>,但对诱导组培杜梨生根过程中的多胺类物质、内源激素及相关氧化酶活性影响的相关报道较少。笔者通过探究生长素诱导杜梨生根过程中多胺类物质含量、内源激素含量及关联酶活性的变化,力图揭示杜梨茎段基部上述物质在不定根发生过程中的生理响应机制,以期为今后杜梨生根调控的相关研究提供理论依据。

## 1 材料和方法

### 1.1 材料

生长健壮、长势相近的杜梨继代苗:8~12枚叶

片,基茎粗度1.0~2.0 mm,株高2.5~3.0 cm。

### 1.2 试验设计

1.2.1 杜梨生根培养基激素浓度配比的筛选 选取健壮、长势相近的杜梨继代苗,接种到1/2MS+不同激素配比+7.5 g·L<sup>-1</sup>琼脂+25.0 g·L<sup>-1</sup>蔗糖的培养基中。激素浓度配比采用正交试验设计L<sub>25</sub>(5<sup>3</sup>),考虑2个因素5个水平:IAA(0、0.2、0.5、1.0、2.0)、IBA(0、0.2、0.5、1.0、2.0)(表1)。每种培养基接种6瓶,每瓶接种5株,在培养室进行培养,培养40 d后统计生根状况。培养条件:温度控制在25 °C左右,光培养时间16 h,暗培养时间8 h,光强2 000~3 000 lx。

表1 不同培养基激素浓度配比情况

Table 1 Combinations of different hormones  
in the culture media

处理 Treatment	$\rho(\text{IAA})/($ $\text{mg} \cdot \text{L}^{-1})$	$\rho(\text{IBA})/($ $\text{mg} \cdot \text{L}^{-1})$	处理 Treatment	$\rho(\text{IAA})/($ $\text{mg} \cdot \text{L}^{-1})$	$\rho(\text{IBA})/($ $\text{mg} \cdot \text{L}^{-1})$
A1	0.0	0.0	C4	1.0	0.5
A2	0.2	0.0	C5	2.0	0.5
A3	0.5	0.0	D1	0.0	1.0
A4	1.0	0.0	D2	0.2	1.0
A5	2.0	0.0	D3	0.5	1.0
B1	0.0	0.2	D4	1.0	1.0
B2	0.2	0.2	D5	2.0	1.0
B3	0.5	0.2	E1	0.0	2.0
B4	1.0	0.2	E2	0.2	2.0
B5	2.0	0.2	E3	0.5	2.0
C1	0.0	0.5	E4	1.0	2.0
C2	0.2	0.5	E5	2.0	2.0
C3	0.5	0.5	-	-	-

1.2.2 诱导生根过程中基部茎段生理生化指标的测定 选取健壮、长势相近的继代苗,接种到上述优化后的生根培养基上,共接种100瓶,每瓶5株,分别在处理后0,3,7,10,15 d的上午10:00进行取样,每次20瓶,取基部茎段0.5 cm左右,用去离子水清洗干净,用纱布擦干后用液氮冷冻研磨后保存于-80 °C的超低温冰箱中,用于多胺类物质、内源激素含量及关联酶活的测定。

### 1.3 测定方法

利用酶联免疫法<sup>[16]</sup>测定基部茎段4种内源激素含量(吲哚乙酸(IAA)、赤霉素(GA<sub>3</sub>)、玉米素核苷(ZR)和脱落酸(ABA))以及3种多胺类物质含量(精胺(Spm)、亚精胺(Spd)和腐胺(Put));IAAO酶和PPO酶的活性测定分别参照王小玲等<sup>[12]</sup>和卢绪娟等<sup>[17]</sup>的方法。

### 1.4 数据处理

所有试验数据分别选用Excel 2010和SPSS 20.0数据分析软件处理。

## 2 结果与分析

### 2.1 杜梨生根培养基配方的筛选与优化

由表2可知,不同浓度的IAA和IBA对杜梨生根状况影响较明显,C3处理的生根率最高,达到了88.9%,但生根数 $\geq 3$ 占比仅为33.3%;C5处理生根数 $\geq 3$ 占比为53.3%,在 $\alpha = 0.05$ 水平下显著高于其他处理,同时生根率达到86.7%,综合考虑生根率及生根数 $\geq 3$ 的占比可知,杜梨较适宜的生根培养基为:1/2MS+2.0 mg·L<sup>-1</sup>IAA+0.5 mg·L<sup>-1</sup>IBA+7.5 g·L<sup>-1</sup>琼脂+25.0 g·L<sup>-1</sup>蔗糖,诱导生根状况见图1。由结果可以看出,IAA和IBA两种激素共同诱导生根效果要优于单一激素处理。在0~2.0 mg·L<sup>-1</sup>浓度范围内,单一激素IBA对杜梨进行生根诱导,低浓度生根效果要优于高浓度,而IAA诱导生根的效果相反。另外,各处理愈伤组织出现在7~10 d开始出现,10~15 d迅速增大,之后增大趋势减缓,在20 d以后各处理相继长出根系。

### 2.2 诱导生根过程中多胺物质含量的变化

由图2可知,3种多胺类物质在诱导3 d后均出现下降趋势,分别比未经诱导的下降14.55%、4.58%和30.55%。Spm和Spd含量呈现下降→升高→下降→升高的“波浪型”变化趋势,均在诱导后0~10 d

表2 不同激素浓度配比杜梨生根状况  
Table 2 Ratio of different hormones for the rooting of *Pyrus betulifolia* Bunge plantlets

处理 Treatment	生根率 Rate of rooting/%	生根数 $\geq 3$ 占比 Rooting number $\geq 3$ /%
A1	-	-
A2	24.0±4.0 hi	8.0±2.0 gh
A3	15.0±4.3 i	5.0±1.3 hi
A4	36.0±6.5 g	4.0±1.0 hi
A5	36.0±4.0 g	8.0±1.7 gh
B1	36.0±3.6 g	8.0±3.0 gh
B2	56.0±6.0 f	16.0±1.7 ef
B3	76.0±10.6 cd	4.0±1.7 hi
B4	68.0±3.5 de	12.0±2.0 fg
B5	72.0±2.0 cd	12.0±3.0 fg
C1	60.0±5.6 ef	20.0±3.4 e
C2	87.5±7.0 ab	31.3±3.6 d
C3	88.9±8.8 a	33.3±4.7 cd
C4	68.8±8.0 de	18.8±4.1 e
C5	86.7±8.2 ab	53.3±3.5 a
D1	66.7±9.2 de	33.3±2.9 cd
D2	74.4±3.8 cd	31.1±3.5 d
D3	66.7±6.2 de	33.3±5.0 cd
D4	60.0±6.2 ef	40.0±9.1 bc
D5	77.8±3.5 bc	44.4±5.1 b
E1	16.7±6.0 i	0.0 i
E2	80.0±5.8 abc	30.0±5.6 d
E3	77.8±4.0 cd	44.4±5.6 b
E4	33.3±2.7 gh	33.3±4.6 cd
E5	55.6±3.2 f	22.2±5.6 e

注:每列数据后字母表示 $\alpha = 0.05$ 水平下差异显著性水平。

Note: The words indicate the significant difference at  $\alpha = 0.05$  level after each column of data.



A. 生根前生长状况;B. 初始生根状况;C. 最终生根状况。

A. Growth status before rooting; B. Initial rooting status; C. Ultimate rooting status.

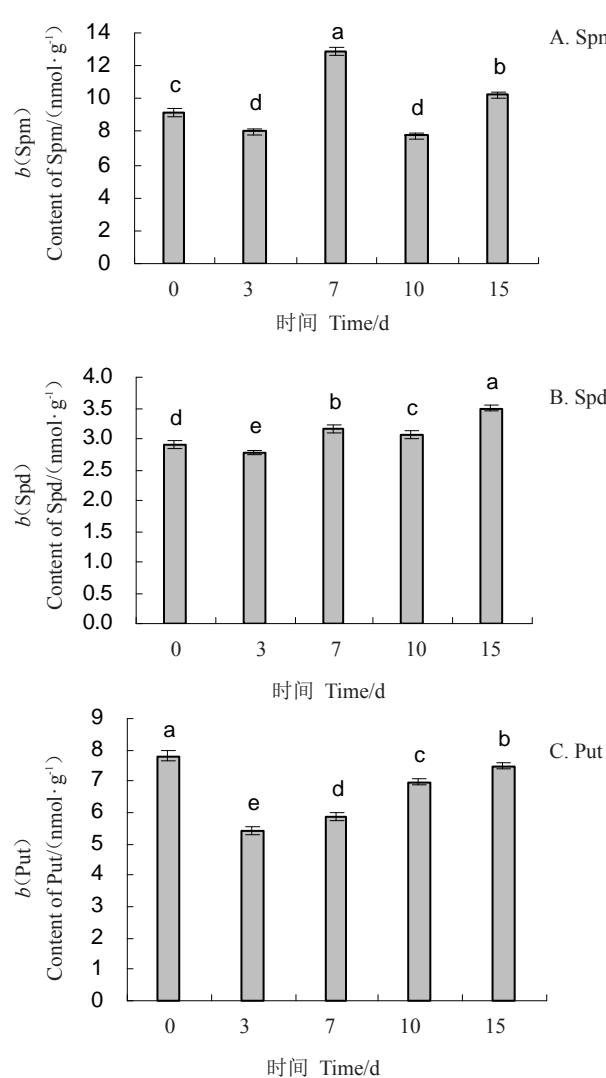
图1 杜梨生根状况  
Fig. 1 Rooting of *Pyrus betulifolia* Bunge

形成“单峰”,峰值分别为12.88 nmol·g<sup>-1</sup>和3.16 nmol·g<sup>-1</sup>,10~15 d二者含量逐步提高。Put含量则呈现出下降→上升的变化趋势,0~3 d含量下降,3~15 d含量增加,但后期增加速率减缓。

### 2.3 诱导生根过程中内源激素含量及比值的变化

由图3可知,内源激素IAA、ABA、GA含量均呈现下降→升高的变化趋势,ZR含量则呈现出下降→

升高→下降的趋势。IAA含量在7 d时最低,7~10 d含量上升迅速,10~15 d上升速度减缓,处理15 d时含量(w,后同)最高,为112.44 ng·g<sup>-1</sup>。ZR含量呈现下降→上升→下降的趋势,处理7 d后含量最低,为76.57 ng·g<sup>-1</sup>。GA含量呈现下降→上升的变化趋势,在3 d出现最低值,仅为3.49 ng·g<sup>-1</sup>,3~15 d逐步上升,3~10 d含量上升迅速,之后上升速度减缓。



同一图中不同小写字母表示  $\alpha = 0.05$  水平下差异的显著性。下同。

Different small letters represent the significant difference at  $\alpha = 0.05$  level in the same figure. The same below.

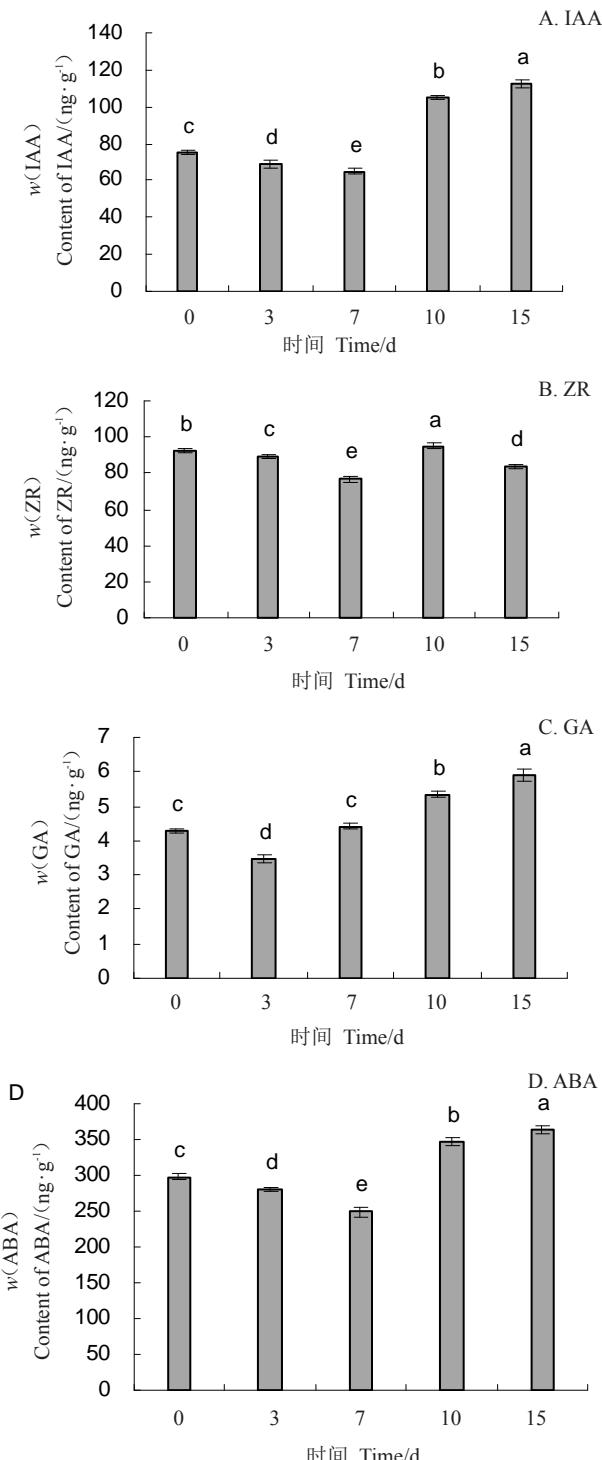
**图 2 不同处理时间 3 种多胺类物质含量**  
**Fig. 2 Contents of three polyamines in different periods**

ABA 含量呈现下降→上升的变化趋势, 0~7 d 逐步降低, 第 7 天时含量最低, 测定值为  $249.07 \text{ ng} \cdot \text{g}^{-1}$ , 7~10 d 呈现较快速率的增长, 10~15 d 增长平缓。

IAA/ABA、IAA/ZR 的比值与植物的生根调控紧密相关<sup>[18-19]</sup>。由图 4 可以看出, IAA/ABA 在 0~7 d 基本无变化, 7~15 d 开始呈现增长趋势; IAA/ZR 呈现下降→升高的趋势, 表明后期 IAA/ZR 值的升高更有利不定根的发生。

#### 2.4 诱导生根过程中 2 种关联酶活性的变化

由图 5 可以看出, 诱导杜梨生根过程中 IAAO 和 PPO 酶活性呈现增高→降低→增高的趋势, IAAO



**图 3 不同处理时间内源激素含量**

**Fig. 3 Content of endogenous hormones in different periods**

酶活最低值出现在第 10 天, PPO 酶活性则在第 7 天; 二者在 0~3 d 显著上升, 3~7 d 下降, 10~15 d 酶活性均出现上升趋势。

#### 2.5 诱导生根过程中多胺类物质、内源激素含量及关联酶酶活性的相关性分析

由表 3 可知, Spm 与内源激素 ZR 呈极显著性负

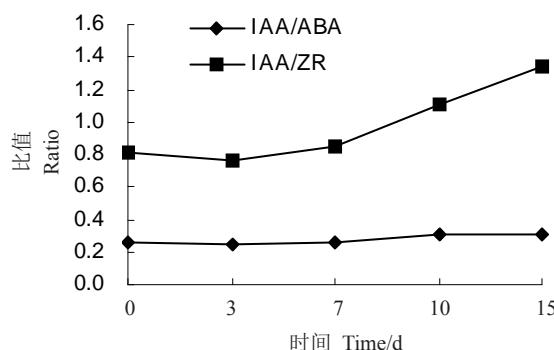


图 4 不同激素比值的变化

Fig. 4 Changes in different hormone ratios

相关( $p < 0.01$ )，相关系数  $r = -0.898$ ，Spd 与 IAA、ABA、IAA/ABA、IAA/ZR 及 IAAO 酶活性均存在极显著正相关关系( $p < 0.01$ )，表明 Spd 在诱导杜梨生根过程中发挥着重要的调控作用。Put 与 GA 呈极显著性正相关( $p < 0.01$ )，相关系数  $r = 0.646$ ，Put 与 IAA、ABA 呈显著正相关关系( $p < 0.05$ )。

PPO 酶活性与 IAA、ABA、IAA/ZR 之间存在极显著正相关关系( $p < 0.01$ )，IAAO 酶活性与 PPO 酶活性、Spd、IAA/ZR 极显著正相关，相关系数分别为 0.843、0.643、0.662。

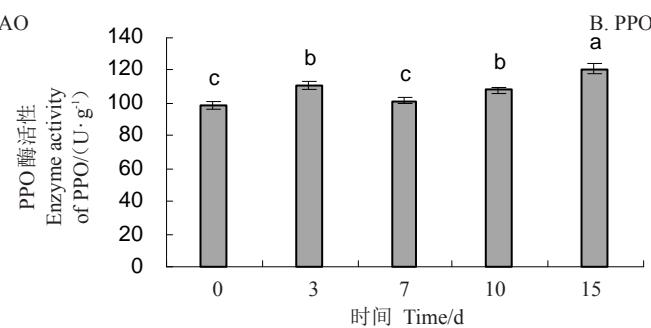
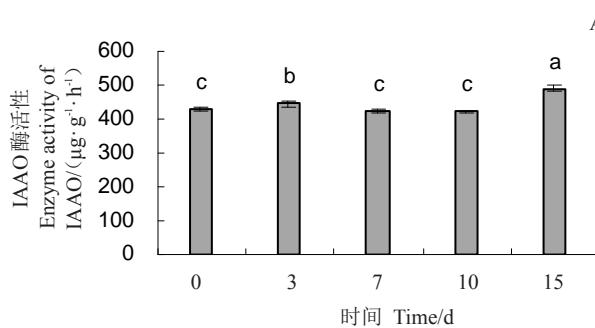


图 5 不同处理时间 2 种关联酶活性

Fig. 5 Changes in the activities of two related enzyme in different periods

表 3 不同生理生化指标及比值的相关性分析

Table 3 Correlation analysis of different physiological and biochemical parameters

	Spm	Spd	Put	IAA	ZR	GA	ABA	比值 1 Ratio 1	比值 2 Ratio 2	PPO	IAAO
Spm	1	0.481	-0.082	-0.290	-0.898**	-0.440	0.114	-0.052	-0.006	-0.206	0.029
Spd		1	0.396	0.658**	-0.463	0.527*	0.852**	0.749**	0.845**	0.555*	0.643**
Put			1	0.594*	0.348	0.646**	0.621*	0.512	0.514	0.057	0.236
IAA				1	0.292	0.967**	0.901**	0.932**	0.944**	0.677**	0.525*
ZR					1	0.450	-0.039	0.084	-0.035	-0.046	-0.230
GA						1	0.808**	0.813**	0.864**	0.674**	0.529*
ABA							1	0.942**	0.948**	0.532*	0.469
比值 1 Ratio 1								1	0.931**	0.531*	0.396
比值 2 Ratio 2									1	0.744**	0.662**
PPO										1	0.843**
IAAO											1

注：“表示在 0.01 水平上显著相关；\*表示在 0.05 水平上显著相关；比值 1 代表 IAA/ABA，比值 2 代表 IAA/ZR。

Note: \*\* Indicates significant correlation at  $\alpha = 0.01$  level; \* Indicates significant correlation at  $\alpha = 0.05$  level; Ratio 1 represents IAA/ABA, ratio 2 represents IAA/ZR.

由内源激素与多胺类物质、两种关联酶活相关分析可以看出，IAA 与 GA、ABA、Spd、IAA/ABA、IAA/ZR、PPO 酶活极显著正相关( $p < 0.01$ )，相关系数分别为 0.967、0.901、0.658、0.932、0.944、0.677。除 IAA 外，ABA 与 GA、Spd、IAA/ZR、IAA/ABA 也存在极显著正相关关系( $p < 0.01$ )，相关系数分别为 0.808、0.852、0.942、0.948，表明 IAA、ABA 在杜梨诱

导生根过程中发挥着重要的调控作用。

### 3 讨 论

#### 3.1 杜梨生根诱导与多胺类物质含量变化的关系

在高等植物中，多胺合成途径有 2 种：鸟氨酸脱羧途径(ODC 途径)和鲱精胺途径(ADC 途径)，Put 与脱羧 S-腺苷甲硫氨酸在亚精胺合成酶催化下生成

Spd, Spd与脱羧S-腺苷甲硫氨酸在精胺合成酶作用下生成Spm<sup>[20]</sup>。植物可以通过调节多胺物质的合成与代谢来调控多项生命活动,多胺含量的增加和降解可以有效激活或降低关联酶的活性从而提高植物体的抗逆性<sup>[21-22]</sup>,还可以影响植物的花芽分化<sup>[23-24]</sup>等。本试验研究结果中初始阶段多胺类物质保持较高水平,处理3 d后含量都有不同程度的下降,分析原因可能是由于茎段基部受伤,多胺类物质含量升高主要是调节植物体抵御外界胁迫,而随着培养时间延长,愈伤组织形成后多胺类物质则主要参与不定根根原基的诱导。有学者研究发现Put的合成与代谢在植物不定根的诱导发生过程中发挥着重要作用<sup>[25]</sup>,通过本试验诱导生根过程中不同生理生化指标间相关性分析可知,Spd与多种激素和2种关联酶活均有一定程度的相关性,而Put含量变化仅与GA存在显著相关性,表明Spd在杜梨不定根诱导和发生过程中起重要调节作用,这一结果与杨洪强<sup>[6]</sup>等在多胺与精胺对苹果实生根系影响的研究结果一致。

### 3.2 杜梨生根诱导与内源激素含量及相关酶活变化的关系

植物不定根形成的步骤大概为:薄壁细胞脱分化形成潜在的根起始点,后经细胞分裂、增大形成形成层,进一步形成根原基,在此过程中,植物激素起到重要调节作用,生长素尤为重要<sup>[26]</sup>。本试验中IAA含量在0~7 d出现下降趋势,可能是IAAO酶活性增强促使IAA降解,或者是PPO酶催化IAA和酚类物质结合形成“IAA-酚酸复合物”,此复合物是不定根发生和伸长的辅助因子。大多数试验证明GA会抑制不定根的发生,但有学者认为抑制与否与GA施用时期紧密相关,后期添加可以促进不定根生成<sup>[27-29]</sup>。本试验结果中GA含量呈现先下降后升高的趋势,表明在诱导初期高浓度GA不利于细胞脱分化,随着后期愈伤组织形成,根源基开始形成,其调节作用才逐步显现,且后期GA含量增加对杜梨不定根发生并无明显的抑制作用。高浓度ABA抑制植物不定根的发生,低浓度ABA对不定根的发生有促进作用<sup>[5]</sup>,本试验中在诱导生根初期ABA含量降低,分析与低浓度ABA有利于愈伤组织的形成及分化有关,后期ABA浓度升高,分析原因可能是在一定浓度范围内ABA升高可以促进杜梨不定根的发生,有学者认为ABA绝对含量的降低并不是促

进生根的主要原因<sup>[30]</sup>,具体原因有待于进一步研究。在诱导生根后期IAA/ABA比值呈略微升高趋势,IAA/ZR比值上升迅速,这与多人研究结果一致<sup>[7, 18-19, 31-33]</sup>,表明IAA/ABA和IAA/ZR比值增大更有利于杜梨不定根的发生和伸长。另外,试验发现无论是生长型还是抑制型激素含量在诱导初期均会降低,随着愈伤组织形成,二者含量也相应增加,可能是初期高浓度激素水平抑制了植株不定根的发生。

## 4 结 论

杜梨较适宜的生根培养基为:1/2MS+2.0 mg·L<sup>-1</sup>IAA+0.5 mg·L<sup>-1</sup>IBA+7.5 g·L<sup>-1</sup>琼脂+25.0 g·L<sup>-1</sup>蔗糖;IAA/ABA、IAA/ZR比值增大有利于杜梨不定根发生;多胺类物质中Spd含量与杜梨不定根发生紧密相关;生根诱导后期,GA含量增加对杜梨不定根发生并无明显的抑制作用。

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