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Sesquiterpene Synthase Gene and Protein

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Chappell et al.

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(45) **Date of Patent:** **Jun. 5, 2012**

(54) **SESQUITERPENE SYNTHASE GENE AND PROTEIN**

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(73) Assignee: **The University of Kentucky Research Foundation**, Lexington, KY (US)

(*) Notice: Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b) by 475 days.

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C12P 1/00 (2006.01)
C12P 7/00 (2006.01)
C12P 7/40 (2006.01)

(52) **U.S. Cl.** **435/41**; 435/71.1

(58) **Field of Classification Search** 435/41,
435/71.1

See application file for complete search history.

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(74) *Attorney, Agent, or Firm* — McKenna Long & Aldridge LLP; Stephanie Seidman

(57) **ABSTRACT**

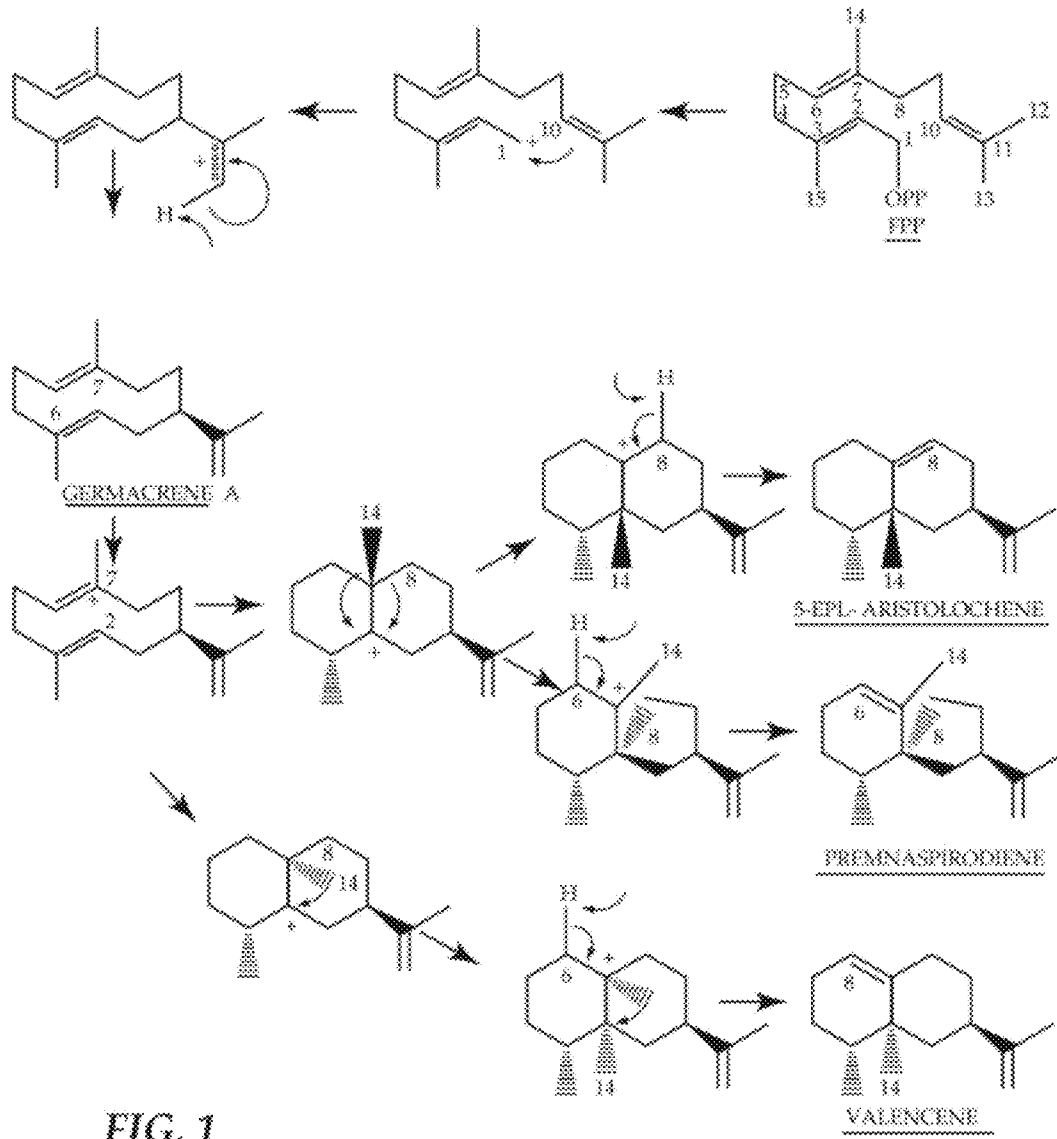
The invention relates to sesquiterpene synthases and methods for their production and use. Particularly, the invention provides nucleic acids comprising the nucleotide sequence of citrus valencene synthase (CVS) which codes for at least one CVS. The invention further provides nucleic acids comprising the nucleotide sequence coding for amino acid residues forming the tier 1 and tier 2 domains of CVS. The invention also provides for methods of making and using the nucleic acids and amino acids of the current invention.

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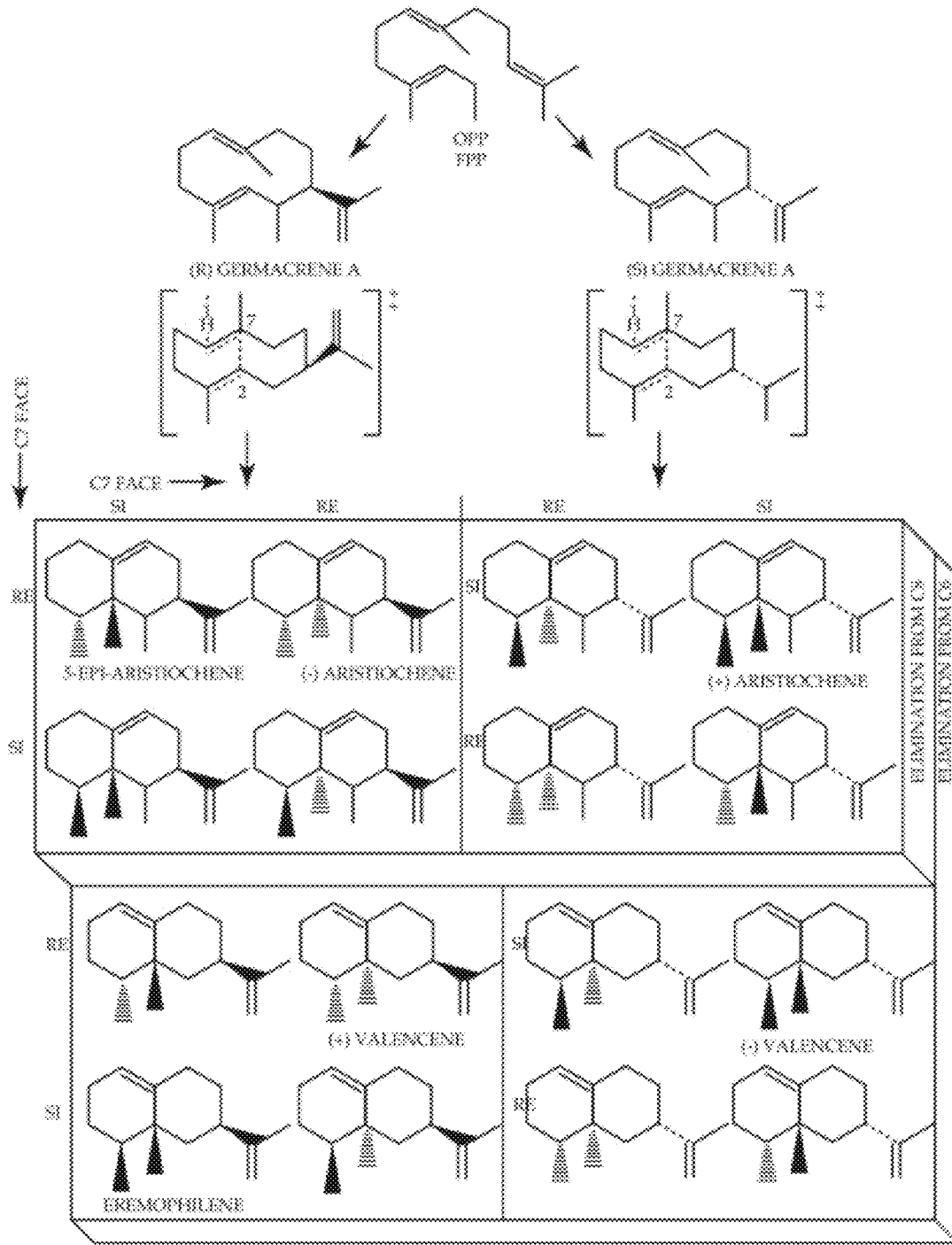


FIG. 2

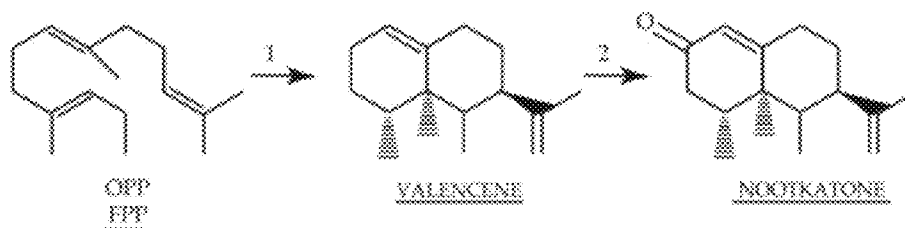


FIG. 3

SYNTHASE PRODUCT	FIRST TIER										SECOND TIER																															
	26	27	28	29	30	31	32	33	34	35	J/K LOOP										36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57
5-EPILABDSTEROLCHENE	R	W	T	Y	C	D	Y	D	Y	T	F	R	C	S	D	V	Y	T	T	L	S	R	D	T	L	I	V	T	H													
PREMNASINCHOLENE	R	W	T	Y	C	D	Y	D	Y	T	F	R	C	S	D	■	Y	T	■	L	■	R	D	T	L	L	■	T	H													
VALENCENE	R	W	■	Y	C	D	Y	D	Y	T	F	R	■	S	D	V	Y	■	■	■	R	D	■	L	■	■	T	H														

FIG. 4

NUCLEOTIDE (DNA) SEQUENCE FOR CITRUS VALENCENE SYNTHASE
GENE (CVS)
SEQ ID NO.: 1
LENGTH: 1710 NT

```
64  ATGTCGTCTGGAGAAACATTTTCGTCCTACTGCAGATTTCCATCCT
109 AGTTTATGGAGAAACCATTTCTCAAAGGTGCTTCTGATTTCAAG
154 ACAGTTGATCATACTGCAACTCAAGAACGACACGAGGCACTGAAA
199 GAAGAGGTAAGGAGAATGATAACAGATGCTGAAGATAAGCCTGTT
244 CAGAAGTTACGCTTGATTGATGAAGTACAACGCCTGGGGGTGGCT
289 TATCACTTTGAGAAAAGAAATAGAAGATGCAATACTAAAATTATGT
334 CCAATCTATATTGACACTAATAGAGCTGATCTCCACACCGTTTCC
379 CTTCAATTTTCGATTGATTAGGCAGCAAGGAATCAAGATTTTCAATG
424 GATGTGTTTGAGAAGTTCAAAGATGATGAGGGTAGATTCAAGTCA
469 TCGTTGATAAACGATGTTCAAGGGATGTTAAGTTTGTACGAGGCA
514 GCATACATGGCAGTTCGCGGAGAACATATATTAGATGAAGCCATT
559 GCTTTCACTACCACTCACCTGAAGTCATTGGTAGCTCAGGATCAT
604 GTAACCCCTAAGCTTGCAGAACAGATAAATCATGCTTTATACCGT
649 CCTCTTCGTA AAAACCCCTACCAAGATTAGAGGGCAGGATTTTATG
694 TCCATGATCAATTCAACAAGTGATCATTTATAACAATAAAACTCTG
739 CTGAATTTTGCAAAGTTAGATTTTAACATATTGCTAGAGCCGCAC
784 AAGGAGGAACTCAATGAATTAACAAAGTGCTGGAAAGATTAGAC
829 TTCACTACAAAACCTACCTTATGCAAGAGACAGATTAGTGGAGTTA
974 TAZTTTTGGGATTTAGGGACATACTTCGAGCCTCAATATGCATTT
919 GGGAGAAAGATAATGACCCAATTA AATTACATATTATCCATCATA
964 GATGATACTTATGATGCGTATGGTACACTTGAAGA AACTCAGCCTC
1009 TTTACTGAAGCAGTTCAAAGATGGAATATTGAGGCCCGTAGATATG
1054 CTTCCAGAATACATGAAATTGATTTACAGGACACTCTTAGATGCT
1099 TTTAATGAAATTGAGGAAGATATGGCCAAGCAAGGAAGATCACAC
1144 TGGGTACGTTATGCAAAAAGAGGAGAATCAAAAAGTAATTGGAGCA
1189 TACTCTGTTCAAGCCAAATGGTTCAAGTGAAGGTTACGTTCCAACA
1234 ATTGAGGAGTATATGCCTATTGCACTAACAAGTTGTGCTTACACA
1279 TTCGTCAFAACA AATTCCTTCCCTTGGCATGGGTTGATTTTGCAACT
1324 AAAGAGGTTTTTGAATGGATCTCCAATAACCCTAAGGTGTTAAAA
1369 GCAGCATCAGTTATCTGCAGACTCATGGATGACATGCAAGGTCAT
1414 GAGTTTGAGCAGAAGAGAGGACATGTTGCGTCAGCTATTGAATGT
1459 TACACGAAGCAGCATGGTGTOTCTAAGGAAGAGGCAATFAAAATG
1504 TTTGAAGAAGAAGTTGCA AATGCATGGAAAGATATTAACGAGGAG
1549 TTGATGATGAAGCCAACCGTTCGTTGCCCGACCACTGCTCGGGACG
1594 ATTC TTAATCTTTCGTCGTTGCAATTGATTTTATTTACAAAAGAGGAC
1639 GACGGCTATACGCATTCTTACCTAATTA AAGATCAA AATTGCTTCT
1684 G TGC TAGGAGACCACGT TCCATTTTGA 1710
```

FIG. 5

AMINO ACID SEQUENCE OF CITRUS VALENCENE SYNTHASE (CVS)															
SEQ ID NO.: 4															
LENGTH: 548 AA															
1	M	S	S	G	E	T	F	R	P	T	A	D	F	H	P
16	S	L	W	R	N	H	F	L	K	G	A	S	D	F	K
31	T	V	D	H	T	A	T	Q	E	R	H	E	A	L	K
46	E	E	V	R	R	M	I	T	D	A	E	D	K	P	V
61	Q	K	L	R	L	I	D	E	V	Q	R	L	G	V	A
76	Y	H	F	E	K	E	I	E	D	A	I	L	K	L	C
91	P	I	Y	I	D	S	N	R	A	D	L	H	T	V	S
106	L	H	F	R	L	L	R	Q	Q	G	I	K	I	S	C
121	D	V	F	E	K	F	K	D	D	E	G	R	F	K	S
136	S	L	I	N	D	V	Q	G	M	L	S	L	Y	E	A
151	A	Y	M	A	V	R	G	E	H	I	L	D	E	A	I
166	A	F	T	T	T	H	L	K	S	L	V	A	Q	D	H
181	V	T	P	K	L	A	E	Q	I	N	H	A	L	Y	R
196	P	L	R	K	T	L	P	R	L	E	A	R	Y	F	M
211	S	M	I	N	S	T	S	D	H	L	Y	N	K	T	L
226	L	N	F	A	K	L	D	F	N	I	L	L	E	P	H
241	K	E	E	L	N	E	L	T	K	W	W	D	D	L	D
256	F	T	T	K	L	P	Y	A	R	D	R	L	V	E	L
271	Y	F	W	D	L	G	T	Y	F	E	P	Q	Y	A	F
286	G	R	K	I	M	T	Q	L	N	Y	I	L	S	I	I
301	D	D	T	Y	D	A	Y	G	T	L	E	E	L	S	L
316	F	T	E	A	V	Q	R	W	N	I	E	A	V	D	M
331	L	P	E	Y	M	K	L	I	Y	R	T	L	L	D	A
346	F	N	E	I	E	E	D	M	A	K	Q	G	R	S	H
361	C	V	R	Y	A	K	E	E	N	Q	K	V	I	G	A
376	Y	S	V	Q	A	K	W	F	S	E	G	Y	V	P	T
391	I	E	E	Y	M	P	I	A	L	T	S	C	A	Y	T
406	F	V	I	T	N	S	F	L	G	M	G	D	F	A	T
421	K	E	V	F	E	W	I	S	N	N	P	K	V	V	K
436	A	A	S	V	I	C	R	L	M	D	D	M	Q	G	H
451	E	F	E	Q	K	R	G	H	V	A	S	A	I	E	C
466	Y	T	K	Q	H	G	V	S	K	E	E	A	I	K	M
481	F	E	E	E	V	A	N	A	W	K	D	I	N	E	E
496	L	M	M	K	P	T	V	V	A	R	P	L	L	G	T
511	I	L	N	L	A	R	A	I	D	F	I	Y	K	E	D
526	D	G	Y	T	H	S	Y	L	I	K	D	Q	I	A	S
541	V	L	G	D	H	V	P	F	*	548					

FIG. 6

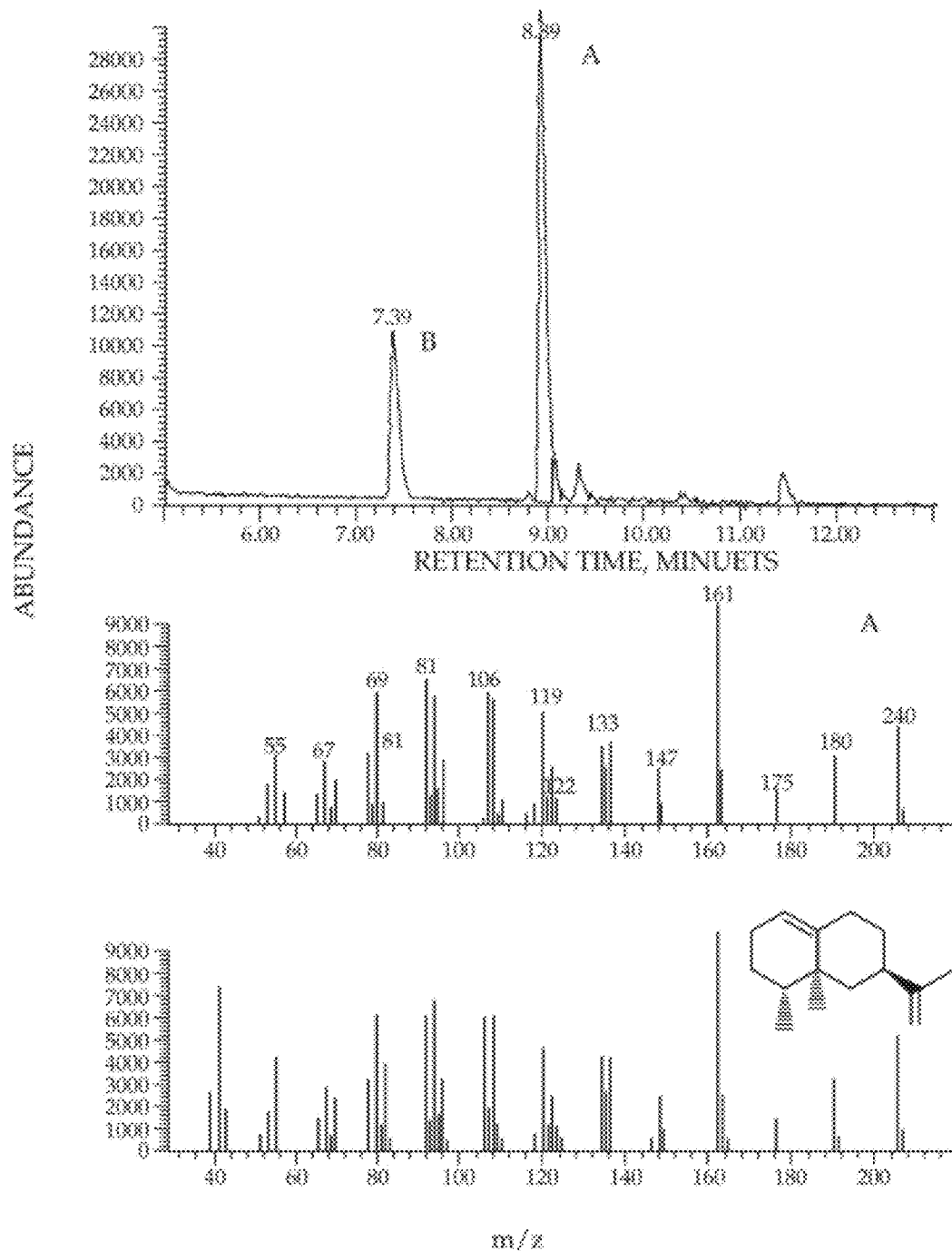


FIG. 7

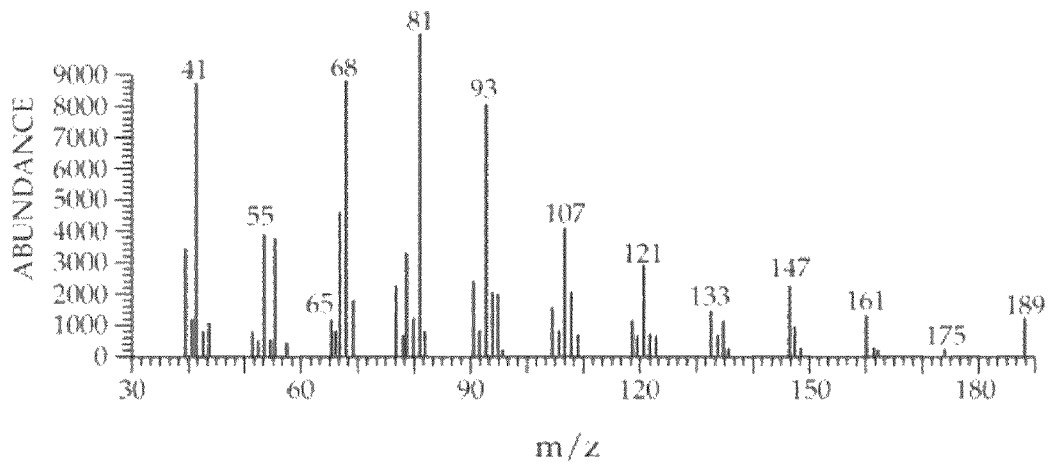
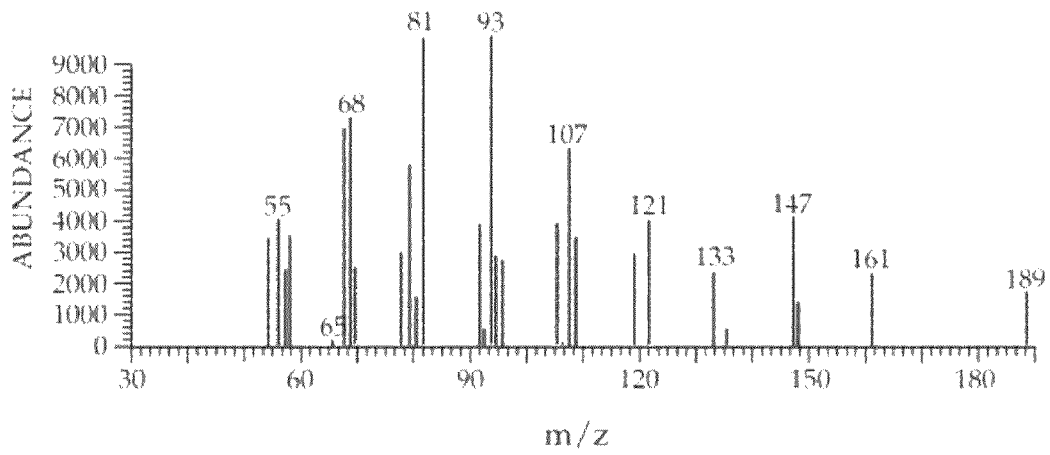


FIG. 8

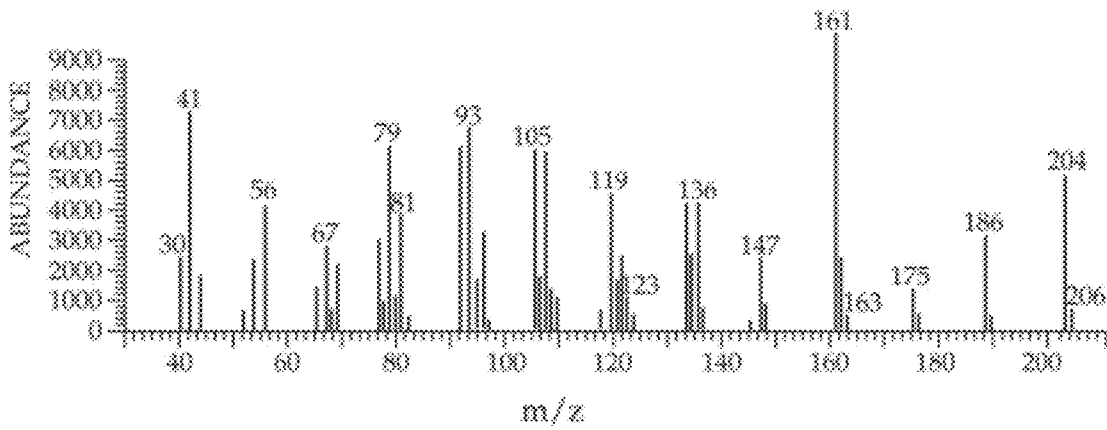
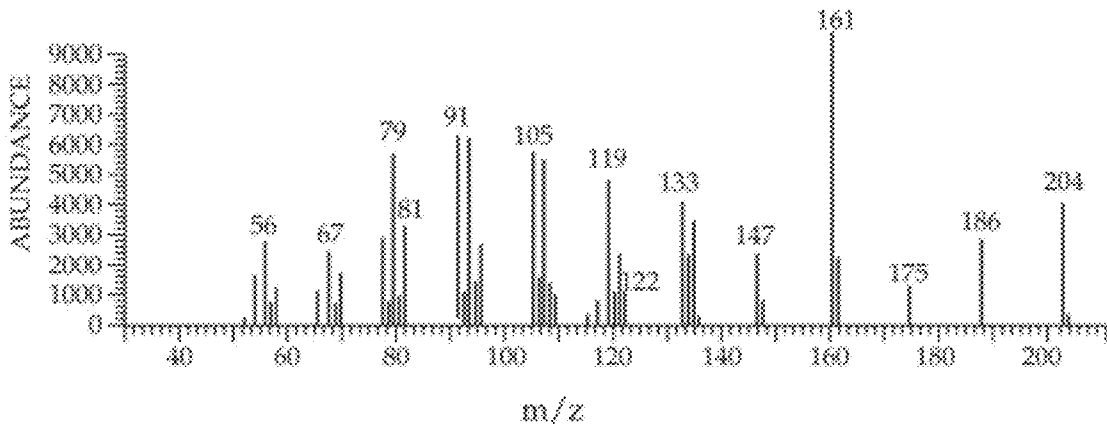


FIG. 9

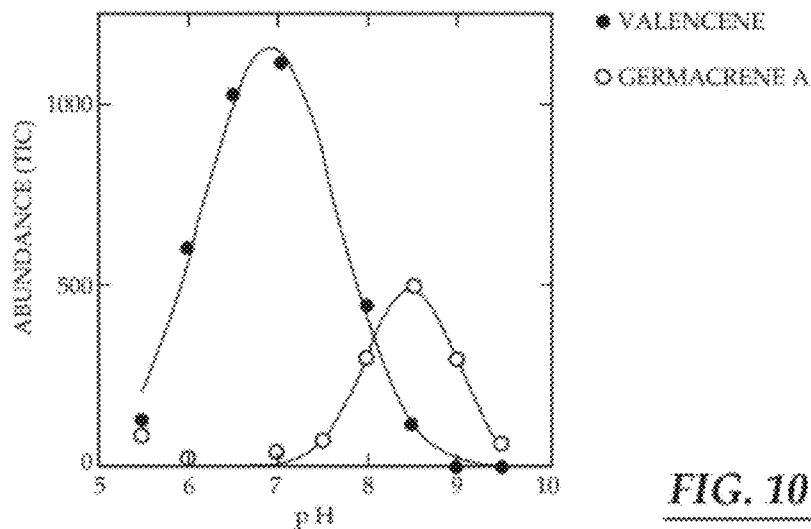


FIG. 10

FIG. 11

SYNTHASE PRODUCT	FIRST TIER									
	304	273	403	404	440	444	J/K LOOP			
5-EPI-ARISTOLOCHENE	R	W	T	Y	C	D	Y	D	Y	T
PREMNASPIRODIENE	R	W	T	Y	C	D	Y	D	Y	T
VALENCENE	R	W	■	Y	C	D	Y	D	Y	T

ABSOLUTE SEQ ID NO 4	COMPARATIVE SEQ ID NO 7	NUCLEOTIDE SEQ ID NO. 5
R264	R264	853aga
W273	W273	680ttg
A403	A403	1270gct
Y404	Y404	1273tac
0441	C440	1384tcg
C445	D444	1396gat
Y522	Y520	1627tac
D526	D525	1639gac
Y528	Y527	1645tat
T329	T328	1648acg

FIG. 12

ABSOLUTE SEQ ID NO 4	COMPARATIVE SEQ ID NO 8	NUCLEOTIDE SEQ ID NO. 6
F13	F20	100ttc
R266	R266	859aga
L270	L270	871tta
S298	S298	955tcc
D302	D302	967gat
V372	V372	1177gtb
Y376	Y376	1189tac
S401	S401	1264agt
C2402	C402	1267tgt
V407	V407	1282gtc
A407	A436	1372gca
B442	B441	1387aga
D446	D445	1399gac
G449	G448	1408ggf
L514	L512	1603ctt
A517	A515	1612gca
B518	B516	1615att
B521	B519	1624att
H530	H529	1651cat

FIG. 13

R264
W273
A403
Y404
C441
D445
Y522
D522b
Y528
T529

FIG. 14a

R264
W273
A403
Y404
C440
D444
Y520
D525
Y527
T526

FIG. 14b

F13
R266
L270
S298
D302
V372
Y376
S401
C402
V407
A437
R442
D446
G449
L514
A517
I518
I521
H530

FIG. 15a

F20
R266
L270
S298
D302
V372
Y376
S401
0402
C407
A436
R441
D445
G448
L512
A515
I516
I519
H529

FIG. 15b

TEAS protein	1	MASAAVANYEEEEIVRPVADFSPSLWGDQFLS--FSIKNQVAEKYAQEIEA	48
CVS - chappell	1	MSSG-----ETFRPTADFHPSLWRNHFLKGCASDFKTVDHTATQERHE	43
TEAS protein	49	LKEQTKNMLLATGMKLADTLNLIIDTIERLGISYHFEKEIDDILDQIYNQ-	97
CVS - chappell	44	LKEEVKRMITDAEDKFPVQKLRLLIDEVQRLCYAYHFEKEIEDAILELCPHY	93
TEAS protein	98	-NSCNDLCTSALQFKLLRQHGFNISPEIFSKFQDENGKFKESLASDVLG	146
CVS - chappell	94	IDSNEADLHTVSLHFRLRQQGKIKISCDFEKFKDDEGRFKSSLINDVQG	143
TEAS protein	147	LLNLYEASHVVRTHADDILEDALAPSTIHLES--AAPHLDSPLREQVTHAL	194
CVS - chappell	144	MLSLYEAAAYMAVRGEHILDEALAFTHLKSLVAQDHVTFKLAEQINHAL	193
TEAS protein	195	EQCLHKGVPVRETRFFISSIYDKE-QSKNNVLLRFKLDNFLLQMLHKQE	243
CVS - chappell	194	YRPLRETLPRLEARYFMSMINSTSDHLYNKTLLNFAKLDNFILLEPHKEE	243
TEAS protein	244	LAQVSRWVKDLDFFVTLLPYARLRKRVVECYHWLGVYFEPQYSQARVMLVKT	293
CVS - chappell	244	LNELTKWVKDLDFTTELPHYARDRLVELYHWDLGTYFEPQYAFGRKIMTQL	293
TEAS protein	294	ISISIVDDTRDAYGTVKELEAYTDATQRWDINEIDRLFDYMKISYKAIL	343
CVS - chappell	294	NYILSHDDTYDAYGTLLELSLFTAVQQRWNIEAVDMLPEYMKLIYRILL	343
TEAS protein	344	DLYKDYEEKELSSAGRSHIVCHAIERMKEYVRYNRYNVESTWPIEGYTPPVSE	393
CVS - chappell	344	DAFNEIEEDMAKQGRSHCVRYAKEENQKVICAYSVOAKWFESEGYVPTIEE	393
TEAS protein	394	YLSNALATITVYYLATTSSYLGMEK-ATEQDFEWLSEKPKILEASVHCRV	442
CVS - chappell	394	YMPALTSQAYFFVITNSFLGMGDFATKEYFEWISNPNKVVKAASVLCRL	443
TEAS protein	443	ITATYEVKSRGQIATGIECCMRDYCISTKEAMAKFQNMETAWDIN	492
CVS - chappell	444	MDPMQGHFEQKRGHVASAIECYTKQHGVSKEEAIKMFEEEVANAWKDIN	493
TEAS protein	493	EGLLRFTFPVSTEFELTPILNLARTVEVTHNMGVTIPEKVLKVLKPHINL	541
CVS - chappell	494	EELMMKPTVVARPLLGTHLNLARAIDHTEKEDGCTYH-SYLIKDQIASV	541
TEAS protein	542	LVDSIKIN 549	
CVS - chappell	542	LGDHVPP- 548	

FIG. 16

TEAS AMINO ACID NUMBER	CVS ABSOLUTE SEQ ID NO.: 4	COMPARATIVE SEQ ID NOS 7 AND 8
TIER 1		
R264	R264	R264
W273	W273	W273
T403	A403	A403
Y404	Y404	Y404
C440	C441	C440
D444	D445	D444
Y520	Y522	Y520
D525	D526	D525
Y527	Y528	Y527
T528	T529	T528
TIER 2		
F20	F13	F20
R266	R266	R266
C270	L270	L270
S298	S298	S298
D302	D302	D302
V372	V372	V372
Y376	Y376	Y376
T401	S401	S401
T402	C402	C402
L407	V407	V407
S436	A437	A436
R441	R442	R441
D445	D446	D445
T448	G449	O448
L512	LL514	L512
I515	A517	A515
V516	I518	I516
T519	I521	I519
H529	H530	H529

FIG. 17

SESQUITERPENE SYNTHASE GENE AND PROTEIN

RELATED APPLICATION

Benefit of priority under 35 U.S.C. 119(e) is claimed herein to U.S. Provisional Application No. 60/489,514, filed Jul. 24, 2003, and benefit of priority under 35 U.S.C. 120 is claimed herein to U.S. Utility application Ser. No. 10/899,356, filed on Jul. 26, 2006, now U.S. Pat. No. 7,442,758, issued Oct. 28, 2008. The disclosures of the above referenced applications are incorporated by reference in their entirety herein.

FIELD OF THE INVENTION

This invention relates generally to the field of production of valencene and nootkatone, and more particularly relates to the discovery of a valencene synthase gene and related protein, which provides a pathway for generating highly pure valencene which can be converted into the flavorant, nootkatone.

BACKGROUND

Terpenes are a diverse family of compounds with carbon skeletons composed of five-carbon isoprene units. Approximately 20,000 different terpenes and terpenoids (compounds of terpene origin whose carbon skeleton has been altered or rearranged) have been identified to date, representing only a small fraction of the estimated natural variation. Terpenes are commonly isolated from the essential oils of plants. Essential oils often have pleasant tastes or aromas, and they are widely used as flavorings, deodorants, and medicines.

Sesquiterpenes are terpenes with 15 carbon atoms (three isoprene units). The plant kingdom contains the highest diversity of sesquiterpenes. Often they play a role in defense of the plants against pathogens, insects and herbivores and for attraction of pollinating insects.

Valencene (1,2,3,5,6,7,8,8a-octahydro-7-isopropenyl-1,8a-dimethyl-naphthalene) and nootkatone (4,4a,5,6,7,8-hexahydro-6-isopropenyl-4,4-a-dimethyl-2(3H)-naphthalenone) are just two examples of sesquiterpenes that are derived from cyclization of the ubiquitous pyrophosphate intermediate farnesyl diphosphate. Nootkatone is formed by the oxidation of valencene.

Valencene and nootkatone are compounds of natural origin, and are natural constituents of citrus oils, such as orange and grapefruit. Because of its excellent organoleptic qualities and in particular its typical grapefruit taste, nootkatone is a widely used ingredient in perfumery and the flavor industry. Alternatively, nootkatone may be used as an insecticide. Valencene, the starting material for the generation of nootkatone (either biologically or chemically), is also used as a flavorant and fragrance.

Several methods to purify sesquiterpenes, such as valencene and nootkatone, from citrus fruits or to maintain high levels of these sesquiterpenes in citrus fruit extracts have been described in the prior art. These methods are described below.

Japikse et al., in U.S. Pat. No. 4,693,905, claimed a method of extracting concentrated orange flavor and aroma compositions from natural orange essence oil by using a dense solvent gas. Their procedure entailed (a) contacting natural orange essence oil with a solvent gas having a temperature between its critical temperature and 100° C., and having a reduced pressure between about 0.56 and about 1.31 (where reduced pressure is defined as the extraction pressure of the solvent gas divided by its critical pressure), to extract flavor

and aroma compounds; (b) separating the solvent gas and dissolved compounds from the remaining undissolved compounds; and (c) separating the dissolved compounds from the solvent gas.

5 Rich, in U.S. Pat. No. 4,973,485, discloses a method of producing aqueous orange stripper essences and orange stripper oils with high ratios of valencene to the less desirable orange flavor compounds. This procedure involves the following steps: (a) heating an orange fed juice stream to a temperature of 37.7° C. to 71° C.; (b) stripping the heated feed juice with steam at a steam:soluble solids ratio of 0.3 to 1.5, at a temperature of 37.7° C. to 71° C. and at a stripping column pressure of less than 9 inches of Hg, absolute; (c) condensing the stripped volatiles at a temperature of 40.6° C. to -196° C.; (d) centrifuging the condensate in a continuous stacked disk hermetic centrifuge to produce two clear phases; and (e) removing the aqueous orange stripper essence phase.

In U.S. Pat. No. 5,260,086 Downton et al. describe a process for making an aseptic citrus sensible pulp/juice slurry by extracting and removing juice from citrus juice containing sensible pulp. After this process is complete, flavorants, such as valencene are added to make up for those that are lost during this extraction process.

Hiramoto et al., in U.S. Pat. No. 6,495,193 prepares a citrus flavor from a low-boiling part of a cold pressed oil by a hydrate alcohol solvent extraction. To maintain the stability of the flavor, a stabilizing coumarin analogue component is added.

In a US Patent Application, publication number US 20030185956, Gradley claims a separation method for extracting desired sesquiterpene aroma compounds, such as valencene and nootkatone, from an aqueous phase by separating the aqueous mixture from a water-immiscible hydrophobic phase by means of a hydrophilic membrane and allowing the desired components to move out off the aqueous phase through the membrane and into the hydrophobic phase.

Kotachi et al., in US Patent Application, publication number 20030203090, teaches of a process for preparing orange oil useful as fragrance or flavor material, by mixing raw material oil containing valencene with a high-boiling solvent having a boiling point exceeding 240° C. under normal pressure, to give a mixture, and fractionally distilling the mixture obtained.

Nootkatone is a high demand, high value flavorant added to many of the commercial soft drinks sold worldwide. Currently, the practice of extracting nootkatone from citrus pulp and rind is considered an expensive and somewhat unreliable process. Nootkatone can be synthesized by the oxidation of valencene. The valencene starting material is expensive and is easily degraded during evaporative heat concentration processes typically used to remove the bulk of water from the feed juice. Thus, current methods to purify valencene from citrus fruits are costly, difficult, and are limited by what the fruit can deliver. Moreover, such methods are vulnerable to interruptions in the supply of citrus fruits, which is dependent on the weather. A frost or hailstorm in a major citrus fruit growing region such as Florida can interrupt the supply. Furthermore, methods to produce nootkatone that consume valencene are quite costly, and thus not commercially desirable. Therefore, there is a need for an alternative means for preparing valencene and nootkatone.

SUMMARY OF THE INVENTION

65 In one embodiment, the invention relates to isolated nucleic acids that encode a sesquiterpene synthase. The invention provides an isolated nucleic acid selected from: (a)

a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO: 1; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 4; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by said nucleic acid is a sesquiterpene synthase. Other embodiments include: a polypeptide encoded by a nucleic acid of the invention; a host cell comprising a nucleic acid of the invention; a non-human organism modified to harbor a nucleic acid of the invention; and methods of producing a polypeptide comprising culturing host cells of the invention.

In another embodiment, the invention provides an isolated polypeptide comprising an amino acid sequence substantially as set out in SEQ ID NO: 4.

In a further embodiment, the invention provides a vector comprising at least one nucleic acid chosen from (a) a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO: 1; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 4; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by said nucleic acid is a sesquiterpene synthase. Other embodiments include, methods of making a recombinant host cell comprising introducing a vector of the invention into a host cell.

In a further embodiment, the invention relates to isolated nucleic acids that encode a sesquiterpene synthase. The invention provides an isolated nucleic acid selected from: (a) a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO: 5 or SEQ ID NO.: 6; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 7 or SEQ ID NO.: 8; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by said nucleic acid is a sesquiterpene synthase. Other embodiments include: a polypeptide encoded by a nucleic acid of the invention; a host cell comprising a nucleic acid of the invention; a non-human organism modified to harbor a nucleic acid of the invention; and methods of producing a polypeptide comprising culturing host cells of the invention.

In another embodiment, the invention provides an isolated polypeptide comprising an amino acid sequence substantially as set out in SEQ ID NO: 7 or SEQ ID NO.: 8.

In a further embodiment, the invention provides a vector comprising nucleic acid chosen from (a) a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO: 5 or SEQ ID NO.: 6; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 7 or SEQ ID NO.: 8; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by said nucleic acid is a sesquiterpene synthase. Other embodiments include, methods of making a recombinant host cell comprising introducing a vector of the invention into a host cell.

In one embodiment, the invention provides a method of making at least one sesquiterpene synthase comprising culturing a host cell modified to contain at least one nucleic acid sequence under conditions conducive to the production of said at least one sesquiterpene synthase. In one embodiment, the at least one nucleic acid is chosen from (a) a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO:1, SEQ ID NO.: 5 or SEQ ID NO.: 6; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 4, SEQ ID NO.: 7 or SEQ ID NO.: 8; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by

said nucleic acid is a sesquiterpene synthase. The host may be chosen from, for example, plants, microorganisms, bacterial cells, yeast cells, plant cells, and animal cells.

In another embodiment the invention provides a method of making at least one terpenoid comprising:

- 1) contacting at least one acyclic pyrophosphate terpene precursor with at least one polypeptide encoded by a nucleic acid of the current invention. In one embodiment, the nucleic acid is chosen from (a) a nucleic acid comprising the nucleotide sequence substantially as set out in SEQ ID NO.: 1, SEQ ID NO.: 5 or SEQ ID NO.: 6; (b) a nucleic acid encoding the polypeptide substantially set out in SEQ ID NO: 4, SEQ ID NO.: 7 or SEQ ID NO.: 8; and (c) a nucleic acid that hybridizes to the nucleic acid of (a) or (b) under low stringency conditions, wherein the polypeptide encoded by said nucleic acid is a sesquiterpene synthase,
- 2) isolating at least one terpenoid produced in 1).

In one embodiment, the at least one terpenoid is chosen from the group consisting of sesquiterpene.

In a further embodiment, the at least one acyclic pyrophosphate terpene precursor is farnesyl-diphosphate (FPP). The sesquiterpene produced by the methods of the invention include, but are not limited to, valencene, valencene derivatives, valencene fragments, and compounds having the citrus valencene carbon skeleton.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 shows the conversion of FPP to the sesquiterpene reaction products 5-epi-aristolochene, premnaspirodiene and valencene catalyzed by *Nicotiana tabacum* epi-aristolochene synthase (TEAS), *Hyoscyamus muticus* premnaspirodiene synthase (HPS), and *Citrus paradisi* valencene synthase (CVS), respectively.

FIG. 2 illustrates a mechanism for the enzyme catalyzed development of regio- and enantio-specificity in eremophilanes. Upon loss of the diphosphate from C1 and generation of the initial carbocation, a chiral center arises from the first attack on either the si or re face of sp² hybridized C10 yielding either (10R)-germacrene A or (10S)-germacrene A. Two additional chiral centers develop upon formation of an internal bond between C2 and C7. This process of proton-initiated, internal cyclization (illustrated as transition states) requires that the p-orbitals of sp² hybridized C2 and C7 face and align with each other. The 4 possible combinations of overlapping orbitals are dependent on the orientation of the respective p-orbitals either above or below the plane of the eudesmallyl carbocation intermediate. The possible re and si orientations of the C2 p-orbitals are given on the vertical axis of the diagram and those for C7 depicted on the horizontal axis. These orientations direct the stereochemistry of the ensuing methyl migrations and hydride shifts (not illustrated) and define, as a consequence, the final chirality of the methyl substitutions at C2 and C3. Subsequent to these rearrangements, a tertiary carbocation centered at C7 is formed (an eremophilyl cation, not shown) and alternate elimination of a proton from C6 or C8 give two subclasses of double bond regioisomers, illustrated as layers. Known eremophilanes are noted by their common name.

FIG. 3 illustrates a proposed pathway for the biosynthesis of nootkatone in citrus. The scheme suggests at least two steps, the first step is catalyzed by the sesquiterpene synthase of the current invention, denoted as "1" in the figure, which leads to the production of valencene and the second step consisting of a regio-specific hydroxylation, followed by oxidation. The second step could be catalyzed by a single mul-

tifunctional hydroxylase or could involve sequential enzyme mediated reactions, and which are denoted as “2” in the figure.

FIG. 4 illustrates a sequence alignment of amino acids lining the active site (1st tier) and those within 3 Å of the active site residues (2nd tier) of TEAS with the corresponding positions of CVS. For uniformity, the TEAS amino acids numbering was used, and thus the corresponding CVS amino acids renumbered—termed comparative numbering. The 1st tier residues lie within 3 Å of a substrate analog co-crystallized within the TEAS enzyme and includes residues making up the J/K loop which clamps down over the active site upon substrate binding. The corresponding residues within CVS (valencene synthase) were initially identified by primary sequence alignment, then visual inspection of the relevant CVS sequences overlaid on the TEAS 3-dimensional structure. Residues different from TEAS are highlighted.

FIG. 5 shows SEQ ID NO.: 1, which is the DNA sequence for the citrus valencene synthase gene from *Citrus paradisi* isolated in the current invention.

FIG. 6 shows SEQ ID NO.: 4, which is the protein sequence for the citrus valencene synthase from *Citrus paradisi*.

FIG. 7 illustrates reaction product analysis of citrus valencene synthase (CVS) incubated at pH 7.5. Lysate of *E. coli* expressing the CVS cDNA was incubated with FIT at pH 7.5 and total pentane extractable products evaluated by GC-MS (upper panel). The mass spectrum of the reaction product corresponding to peak A (middle panel) is compared to that of authentic valencene in the lower panel. The mass spectrum of peak B is identical to that for beta-elemene, the thermal-induced rearrangement product of germacrene A.

FIG. 8 illustrates a mass spectrum for the peak with a retention time of 7.38 minutes from FIG. 7 (top) compared to the spectra for beta-elemene published by the NIST library (bottom).

FIG. 9 illustrates mass spectrum for the peak with a retention time of 8.89 in FIG. 7 (top) compared to that for valencene purchased from Fluka Chemical Company (bottom).

FIG. 10 illustrates that the reaction product specificity of citrus valencene synthase (CVS) is pH dependent. Partially purified synthase isolated from *E. coli* expressing the CVS cDNA was incubated with FPP at the indicated pH values and ethylacetate extracts were examined directly by GC-MS. Absolute values for valencene (solid symbols) and germacrene A (measured as the thermally rearranged product β -elemene) (open symbols) are reported and represent greater than 95% of the total reaction products at all pHs.

FIG. 11 is a sequence alignment of amino acids lining the active site (1st tier) of TEAS with the corresponding positions of HPS and CVS. The 1st tier residues lie within 3 Å of a substrate analog co-crystallized within the TEAS enzyme and includes residues making up the J/K loop which clamps down over the active site upon substrate binding. The corresponding residues within the other terpene synthases were initially identified by primary sequence alignment, then visual inspection of the relevant sequences overlaid on the TEAS 3-dimensional structure. Residues in CVS differing from TEAS are highlighted.

FIG. 12 shows SEQ ID NO.: 5, which is the DNA sequence from SEQ ID NO.: 1 corresponding to the amino acids forming Tier 1. Comparative numbering from Tier 1 amino acid residues is also shown (SEQ ID NO.: 7).

FIG. 13 shows SEQ ID NO.: 6, which is the DNA sequence from SEQ ID NO.: 1 corresponding to the amino acids forming Tier 2. Comparative numbering from Tier 2 amino acid residues is also shown (SEQ ID NO.: 8).

FIG. 14a shows the absolute amino acid sequence from SEQ ID NO.: 4 corresponding to the Tier 1 amino acid residues. FIG. 14b shows the comparative amino acid sequence of the Tier 1 amino acid residues (SEQ ID NO.: 7)

FIG. 15a shows the absolute amino acid sequence from SEQ ID NO.: 4 corresponding to Tier 2 amino acid residues. FIG. 15b shows the comparative amino acid sequence of the Tier 2 amino acid residues (SEQ ID NO.: 8).

FIG. 16 is a sequence alignment of the amino acid sequence of TEAS active site compared to the amino acid sequence of the CVS active site. The alignment maximizes residue similarities and introduce gaps where necessary. Absolute amino acid numbering is shown for CVS, thus amino acid numbering from SEQ ID NO.: 4 is shown for CVS.

FIG. 17 is a chart showing the TEAS Tier 1 and Tier 2 amino acid residues in column 1; the CVS Tier 1 and Tier 2 amino acid residues with absolute numbering in column 2; and the CVS Tier 1 and Tier 2 amino acid residues with comparative numbering in column 3.

DETAILED DESCRIPTION OF THE INVENTION

Abbreviations and Terms

In accordance with the present invention and as used herein, the following terms and abbreviations are defined with the following meanings, unless explicitly stated otherwise. These explanations are intended to be exemplary only. They are not intended to limit the terms as they are described or referred to throughout the specification. Rather, these explanations are meant to include any additional aspects and/or examples of the terms as described and claimed herein.

The following abbreviations are used herein:

As used herein, a “derivative” is any compound obtained from a known or hypothetical compound and containing essential elements of the parent substance.

The phrase “substantially identical” means that a relevant sequence is at least 70%, 75%, 80%, 85%, 90%, 92%, 95%, 96%, 97%, 98%, or 99% identical to a given sequence. By way of example, such sequences may be allelic variants, sequences derived from various species, or they may be derived from the given sequence by truncation, deletion, amino acid substitution or addition. Percent identity between two sequences is determined by standard alignment algorithms such as ClustalX when the two sequences are in best alignment according to the alignment algorithm.

As used herein, the term “hybridization” or “hybridizes” under certain conditions is intended to describe conditions for hybridization and washes under which nucleotide sequences that are significantly identical or homologous to each other remain bound to each other. Appropriate hybridization conditions can be selected by those skilled in the art with minimal experimentation as exemplified in Ausubel, F. A., et al., eds., Current Protocols in Molecular Biology Vol. 2, John Wiley and Sons, Inc., New York (1995). Additionally, stringency conditions are described in Sambrook et al. Molecular Cloning: A Laboratory Manual, 2nd ed., Cold Spring Harbor Laboratory Press, New York (1989). Variations on the conditions for low, moderate, and high stringency are well known in the art and may be used with the current invention.

The terms “nucleic acid” or “nucleic acid molecule” refer to a deoxyribonucleotide or ribonucleotide polymer in either single- or double-stranded form, and unless otherwise limited, would encompass known analogs of natural nucleotides that can function in a similar manner as naturally occurring nucleotides. A “nucleotide sequence” also refers to a poly-

nucleotide molecule or oligonucleotide molecule in the form of a separate fragment or as a component of a larger nucleic acid. The nucleotide sequence or molecule may also be referred to as a "nucleotide probe." Some of the nucleic acid molecules of the invention are derived from DNA or RNA isolated at least once in substantially pure form and in a quantity or concentration enabling identification, manipulation, and recovery of its component nucleotide sequence by standard biochemical methods. Examples of such methods, including methods for PCR protocols that may be used herein, are disclosed in Sambrook et al. *Molecular Cloning: A Laboratory Manual*, 2nd ed., Cold Spring Harbor Laboratory Press, New York (1989), Ausubel, F. A., et al., eds., *Current Protocols in Molecular Biology*, John Wiley and Sons, Inc., New York (1987), and Innis, M., et al. (Eds.) *PCR Protocols: A Guide to Methods and Applications*, Academic Press, San Diego, Calif. (1990). Reference to a nucleic acid molecule also includes its complement as determined by the standard Watson-Crick base-pairing rules, with uracil (U) in RNA replacing thymine (T) in DNA, unless the complement is specifically excluded.

As described herein, the nucleic acid molecules of the invention include DNA in both single-stranded and double-stranded form, as well as the DNA or RNA complement thereof. DNA includes, for example, DNA, genomic DNA, chemically synthesized DNA, DNA amplified by PCR, and combinations thereof. Genomic DNA, including translated, non-translated and control regions, may be isolated by conventional techniques, e.g., using any one of the cDNAs of the invention, or suitable fragments thereof, as a probe, to identify a piece of genomic DNA which can then be cloned using methods commonly known in the art.

Polypeptides encoded by the nucleic acids of the invention are encompassed by the invention. As used herein, reference to a nucleic acid "encoding" a protein or polypeptide encompasses not only cDNAs and other intronless nucleic acids, but also DNAs, such as genomic DNA, with introns, on the assumption that the introns included have appropriate splice donor and acceptor sites that will ensure that the introns are spliced out of the corresponding transcript when the transcript is processed in a eukaryotic cell. Due to the degeneracy of the genetic code wherein more than one codon can encode the same amino acid, multiple DNA sequences can code for the same polypeptide. Such variant DNA sequences can result from genetic drift or artificial manipulation (e.g., occurring during PCR amplification or as the product of deliberate mutagenesis of a native sequence). Deliberate mutagenesis of a native sequence can be carried out using numerous techniques well known in the art. For example, oligonucleotide-directed site-specific mutagenesis procedures can be employed, particularly where it is desired to mutate a gene such that predetermined restriction nucleotides or codons are altered by substitution, deletion or insertion. Exemplary methods of making such alterations are disclosed by Walder et al. (*Gene* 42:133, 1986); Bauer et al. (*Gene* 37:73, 1985); Craik (*BioTechniques*, Jan. 12-19, 1985); Smith et al. (*Genetic Engineering: Principles and Methods*, Plenum Press, 1981); Kunkel (*Proc. Natl. Acad. Sci. USA* 82:488, 1985); Kunkel et al. (*Methods in Enzymol.* 154:367, 1987). The present invention thus encompasses any nucleic acid capable of encoding a protein of the current invention.

The current invention provides for isolated polypeptides. As used herein, the term "polypeptides" refers to a genus of polypeptide or peptide fragments that encompass the amino acid sequences identified herein, as well as smaller fragments. Alternatively, a polypeptide may be defined in terms of its antigenic relatedness to any peptide encoded by the nucleic

acid sequences of the invention. Thus, in one embodiment, a polypeptide within the scope of the invention is defined as an amino acid sequence comprising a linear or 3-dimensional epitope shared with any peptide encoded by the nucleic acid sequences of the invention. Alternatively, a polypeptide within the scope of the invention is recognized by an antibody that specifically recognizes any peptide encoded by the nucleic acid sequences of the invention. Antibodies are defined to be specifically binding if they bind polypeptides of the invention with a K_d of greater than or equal to about 10^7 M^{-1} , such as greater than or equal to 10^8 M^{-1} . As used herein, the term "isolated," in reference to polypeptides or proteins, means that the polypeptide or protein is substantially removed from polypeptides, proteins, nucleic acids, or other macromolecules with which it, or its analogues, occurs in nature. Although the term "isolated" is not intended to require a specific degree of purity, typically, the protein will be at least about 75% pure, more typically at least about 90% pure, preferably at least about 95% pure, and more preferably at least about 99% pure.

A polypeptide "variant" as referred to herein means a polypeptide substantially homologous to a native polypeptide, but which has an amino acid sequence different from that encoded by any of the nucleic acid sequences of the invention because of one or more deletions, insertions or substitutions. Variants can comprise conservatively substituted sequences, meaning that a given amino acid residue is replaced by a residue having similar physiochemical characteristics. See Zubay, *Biochemistry*, Addison-Wesley Pub. Co., (1983). It is a well-established principle of protein and peptide chemistry that certain amino acid substitutions, entitled "conservative" amino acid substitutions, can frequently be made in a protein or a peptide without altering either the conformation or the function of the protein or peptide. Such changes include substituting any of isoleucine (I), valine (V), and leucine (L) for any other of these amino acids; aspartic acid (D) for glutamic acid (E) and vice versa; glutamine (Q) for asparagine (N) and vice versa; and serine (S) for threonine (T) and vice versa.

The above-mentioned substitutions are not the only amino acid substitutions that can be considered "conservative." Other substitutions can also be considered conservative, depending on the environment of the particular amino acid. For example, glycine (G) and alanine (A) can frequently be interchangeable, as can be alanine and valine (V). Methionine (M), which is relatively hydrophobic, can frequently be interchanged with leucine and isoleucine, and sometimes with valine. Lysine (K) and arginine (R) are frequently interchangeable in locations in which the significant feature of the amino acid residue is its charge and the differing pK 's of these two amino acid residues are not significant. Still other changes can be considered "conservative" in particular environments.

The effects of such substitutions can be calculated using substitution score matrices such PAM120, PAM-200, and PAM-250 as discussed in Altschul, (*J. Mol. Biol.* 219:55565 (1991)). Other such conservative substitutions, for example, substitutions of entire regions having similar hydrophobicity characteristics, are well known.

Naturally-occurring peptide variants are also encompassed by the invention. Examples of such variants are proteins that result from alternate mRNA splicing events or from proteolytic cleavage of the polypeptides described herein. Variations attributable to proteolysis include, for example, differences in the N- or C-termini upon expression in different

types of host cells, due to proteolytic removal of one or more terminal amino acids from the polypeptides encoded by the sequences of the invention.

Variants of the valencene synthase of the invention may be used to attain desired enhanced or reduced enzymatic activity, modified regiochemistry or stereochemistry, or altered substrate utilization or product distribution. A variant or site direct mutant may be made by any methods known in the art. Variants and derivatives of native polypeptides can be obtained by isolating naturally-occurring variants, or the nucleotide sequence of variants, of other or same plant lines or species, or by artificially programming mutations of nucleotide sequences coding for native citrus polypeptides.

In one embodiment, the invention contemplates: vectors comprising the nucleic acids of the invention. A vector as used herein includes any recombinant vector including but not limited to viral vectors, bacteriophages and plasmids.

Expression vectors containing a nucleic acid sequence of the invention can be prepared using well known methods and include a cDNA sequence encoding the polypeptide operably linked to suitable transcriptional or translational regulatory nucleotide sequences. Examples of regulatory sequences include transcriptional promoters, operators, or enhancers, mRNA ribosomal binding sites, and appropriate sequences which control transcription and translation initiation and termination. Nucleotide sequences are "operably linked" when the regulatory sequence functionally relates to the cDNA sequence of the invention. Expression vectors, regulatory elements and the construction thereof are well known in the art, and therefore are not limited to those recited above.

In addition, sequences encoding appropriate signal peptides that are not naturally associated with the polypeptides of the invention can be incorporated into expression vectors. For example, a DNA sequence for a signal peptide (secretory) leader can be fused in-frame to a nucleotide sequence of the invention so that the polypeptide of the invention is initially translated as a fusion protein comprising the signal peptide. A signal peptide that is functional in the intended host cells enhances extracellular secretion of the expressed polypeptide. The signal peptide can be cleaved from the polypeptide upon secretion from the cell. In some cases, signal peptides are cleaved in two or more stages; this is also within the scope of the invention where appropriate.

Fusions of additional peptide sequences at the amino and carboxyl terminal ends of the polypeptides of the invention can be used with the current invention.

In one embodiment, the invention includes a host cell comprising a nucleic acid of the invention. Another embodiment of the invention is a method of making a recombinant host cell comprising introducing the vectors of the invention, into a host cell. In a further embodiment, a method of producing a polypeptide comprising culturing the host cells of the invention under conditions to produce the polypeptide is contemplated. In one embodiment the polypeptide is recovered. The methods of the invention include methods of making at least one valencene synthase of the invention comprising culturing a host cell comprising a nucleic acid of the invention, and recovering the sesquiterpene synthase accumulated.

Suitable host cells for expression of polypeptides of the invention are well known in the art, and include, but are not limited to, prokaryotes, yeast, higher eukaryotic cells, or combinations thereof. (See for example, Pouwels et al. *Cloning Vectors: A Laboratory Manual*, Elsevier, N.Y. (1985)). Cell-free translation systems, also well known in the art, could also be employed to produce the disclosed polypeptides using RNAs derived from DNA constructs disclosed herein.

Host cells may be modified by any methods known in the art for gene transfer including, for example, the use of delivery devices such as lipids and viral vectors, naked DNA, electroporation and particle-mediated gene transfer.

In one embodiment, the cDNAs of the invention may be expressed in such a way as to produce either sense or antisense RNA. The expression of antisense RNA can be used to down-modulate the expression of the protein encoded by the mRNA to which the antisense RNA is complementary.

A further embodiment of the invention is methods of making terpenoids and sesquiterpene compounds, for example, using the nucleotides and polypeptides of the invention.

As used herein an acyclic pyrophosphate terpene precursor is any acyclic pyrophosphate compound that is a precursor to the production of at least one terpene including but not limited to geranyl-pyrophosphate (GPP), farnesyl-diphosphate (FPP) and geranylgeranyl-pyrophosphate (GGPP).

In one embodiment, the distribution of products or the actual products formed may be altered by varying the pH at which the synthase contacts the acyclic pyrophosphate terpene precursor.

Also within the practice of the invention is an organism (e.g., microorganism or plant) that is used to construct a platform for high level production of a substrate of sesquiterpene synthases (e.g., FPP) and the introduction of a nucleic acid of the invention into the organism.

Unless otherwise indicated, nucleic acids of the invention that are DNA encompass both cDNA (DNA reverse transcribed from mRNA and lacking introns) and isolated genomic DNA (DNA that can contain introns.)

In one embodiment, the nucleic acids of the invention are used to create other nucleic acids coding for sesquiterpene synthases. For example, the invention provides for a method of identifying a sesquiterpene synthases comprising constructing a DNA library using the nucleic acids of the invention, screening the library for nucleic acids which encode for at least one sesquiterpene synthase. The DNA library using the nucleic acids of the invention may be constructed by any process known in the art where DNA sequences are created using the nucleic acids of the invention as a starting point, including but not limited to DNA shuffling. In such a method, the library may be screened for sesquiterpene synthases using a functional assay to find a target nucleic acid that encodes a sesquiterpene synthase. The activity of a sesquiterpene synthase may be analyzed using, for example, the methods described herein. In one embodiment, high through put screening is utilized to analyze the activity of the encoded polypeptides.

As used herein a "nucleotide probe" is defined as an oligonucleotide or polynucleotide capable of binding to a target nucleic acid of complementary sequence through one or more types of chemical bonds, through complementary base pairing, or through hydrogen bond formation.

A "target nucleic acid" herein refers to a nucleic acid to which the nucleotide probe or molecule can specifically hybridize. The probe is designed to determine the presence or absence of the target nucleic acid, and the amount of target nucleic acid. The target nucleic acid has a sequence that is significantly complementary to the nucleic acid sequence of the corresponding probe directed to the target so that the probe and the target nucleic acid can hybridize. Preferably, the hybridization conditions are such that hybridization of the probe is specific for the target nucleic acid. As recognized by one of skill in the art, the probe may also contain additional nucleic acids or other moieties, such as labels, which may not specifically hybridize to the target. The term target nucleic acid may refer to the specific nucleotide sequence of a larger

nucleic acid to which the probe is directed or to the overall sequence (e.g., gene or mRNA). One skilled in the art will recognize the full utility under various conditions.

Other than in the operating example, or where otherwise indicated, all numbers expressing quantities of ingredients, reaction conditions, and so forth used in the specification and claims are to be understood as being modified in all instances by the term "about." Accordingly, unless indicated to the contrary, the numerical parameters set forth in the specification and claims are approximations that may vary depending upon the desired properties sought to be obtained by the present invention. At the very least, and not as an attempt to limit the application of the doctrine of equivalents to the scope of the claims, each numerical parameter should be construed in light of the number of significant digits and ordinary rounding approaches.

Notwithstanding that the numerical ranges and parameters setting forth the broad scope of the invention are approximations, the numerical values set forth in the specific examples are reported as precisely as possible. Any numerical value; however, inherently contains certain errors necessarily resulting from the standard deviation found in their respective testing measurements.

Hundreds of genes with similarity to terpene synthases are readily observed in protein sequence or keyword searches of GenBank with a significant percentage of these currently annotated as terpene synthase-like. In order to isolate synthase genes coding for enzymes that share catalytic features in common with tobacco 5-epi-aristolochene synthase (TEAS) or *Hyoscyamus muticus* premnaspirodiene synthase (HPS) (FIG. 1), Inventors considered possible biosynthetic routes for all isomeric forms of the eremophilane-type sesquiterpenes (FIG. 2). (+)Valencene and its oxygenated derivative nootkatone (FIG. 3) were readily recognized as high value natural products isolated from citrus used as flavor enhancers. To identify the terpene synthase responsible for the biosynthesis of valencene, Inventors screened sequences from several citrus (*Citrus x paradisi* and *Citrus junos*) EST sequencing projects deposited in GenBank (www.ncbi.nlm.nih.gov/Genbank/index.html) for sequences similar to TEAS and HPS. Two sequences were observed, one from *Citrus junos* (accession AF288465) and the other from *Citrus paradisi* (accession AF411120). The predicted proteins from both of these cDNAs were between about 40% identical and about 60% similar to TEAS, but the *C. junos* cDNA predicted a protein that was missing segments of 10 and 12 amino acids relative to TEAS. The isolated protein from *Citrus paradisi* was 45% homologous with TEAS (see below). Inventors, therefore, focused on the isolation and characterization of a full-length cDNA corresponding to AF411120 from grapefruit (*C. paradisi*).

The cDNA isolated by Inventors is a terpene synthase gene, the protein product of which is citrus valencene synthase (CVS). The following non-limiting examples describe Inventors isolation, and expression of the CVS gene, and also further characterize the CVS protein via enzyme activity assays and pH assays. Applicants have further characterized the key amino acid residues comprising the active domain of the CVS protein via these enzyme activity assays and via a comparison to other eremophilane-type sesquiterpenes, in particular TEAS, HPS and delta-cadinene synthase from cotton.

Sequence comparisons between terpene synthases have suggested that simple sequence alignments are not sufficient to identify amino acids or protein regions contributing to catalytic specificity. Inventors aligned the residues forming the active region of the isolated CVS with the residues form-

ing active region of TEAS. In order to perform a comparative analysis of the two active regions (CVS and TEAS), the absolute amino acid number for CVS was shifted to correspond with the TEAS amino acid numbers. FIG. 16 illustrates the alignment of CVS amino acid residues with those of TEAS and maintains the numbering of TEAS and the numbering for CVS. Inventors then adjusted the amino acid numbering of CVS to match that of TEAS (comparative numbering). Tier 1 and Tier 2 amino acid numbering follows this comparative numbering, thus, Tier 1 and Tier 2 amino acids would also have the comparative numbering shift (SEQ ID NOS.: 7 and 8, respectively). FIG. 17. Thus, as is used herein the term "absolute numbering" refers to the sequence number of the CVS amino acid residues in SEQ ID NO.: 1 and the nucleotide numbering in SEQ ID NO.: 4. As used herein, the term "comparative numbering" refers to the sequence numbering of the amino acid residues based on the TEAS numbering, which were designated as a means to compare TEAS and CVS active site amino acid residues, and neighboring residues (Tiers 1 and 2). Nucleic acid residues coding for the amino acids comprising Tier 1 (SEQ ID NO.: 5) and Tier 2 (SEQ ID NO.: 6) are based on the nucleotide sequence of SEQ ID NO.: 1, and are absolute numbers.

As is seen in FIG. 4 and FIG. 11 wherein Inventors compared the amino acids of TEAS, HPS and CVS, only a single amino acid difference was observed within the 1st tier residues of TEAS and CVS; however 8 amino acid differences were observed within the second tier residues of TEAS and CVS. CVS exhibits an alanine at position 403 rather than a threonine as is found in TEAS. Using this discovery, Inventors have further compared amino acid residues surrounding the active site and discovered that these amino acids, termed Tier 2 amino acids and discussed below, influence the catalytic outcome of these enzymes.

For clarity in this disclosure, Tier 1 amino acids are the amino acid residues that form the catalytic pocket in a sesquiterpene synthase, and Tier 2 amino acids are those amino acids that are within a few angstroms of the Tier 1 residues, preferably between the range of about 1 angstrom and about 5 angstroms, more preferably between the range of about 2 angstrom and about 4 angstrom and most preferably 3 angstrom. In this disclosure, the amino acid residue sequence for Tier 1 and Tier 2, based on the amino acid residue sequence of SEQ ID NO.: 4, but adjusted for a comparative analysis with TEAS, are SEQ ID NO.: 7, shown in FIG. 14b, and SEQ ID NO.: 8, shown in FIG. 15b. The corresponding nucleotide sequence for the Tier 1 and Tier 2 amino acid residues, therefore, are based on the nucleotide sequence of SEQ ID NO.: 1, and are SEQ ID NO.: 5, shown in FIG. 12, and SEQ ID NO.: 6, shown in FIG. 13. The recited amino acid residue sequences, recited position number, corresponding nucleotide and nucleotide position number are based on the full protein and nucleotide sequences recited in SEQ ID NO. 1 and SEQ ID NO.: 4, respectively. Because the current invention accounts for frame shift mutations, fusion proteins, fragments, degeneracy of the genetic code and other events that can change these recited sequences and positions without losing the spirit of the current invention, the sequences and positions of these events are included in the current invention.

Inventors then analyzed the second tier amino acid residues for the TEAS and CVS protein sequences. Second tier residues, those amino acid R groups within a few Å of the 1st tier amino acids, were examined for differences between TEAS and CVS (FIG. 4). Three positions within the 2nd tier exhibited common differences between TEAS and CVS (positions 402, 436 and 516), as well as substitutions unique to CVS (270, 401, 407, 448, 515) relative to TEAS. A role for 2nd tier

residues in catalysis, then, includes determining whether a synthase produces a C7-C8 double bond or a C6-C7 double bond. This arises because the final proton abstraction occurred at C6 rather than C8, a spatial distance of 3-4 Å or approximately the diameter of a methyl group. Equally important, regio-specific abstraction necessarily arises from proper geometric positioning of C8 or C6 of the eremophyl cation in proximity to an active site residue or activated water molecule capable of the final abstraction. Positioning of the final reaction intermediate therefore comes about by simple spatial allowances created by the presence or absence of slightly bulkier R-groups on one side of the active site pocket balanced by the converse on the opposite site of the pocket. For instance, the poly-threonine sequence at positions 401-403 of TEAS is replaced by the small amino acids serine, cysteine and alanine in CVS, which appear balanced by a smaller amino acid (valine 516) on the opposite side of the active site pocket in TEAS and the larger amino acid isoleucine in CVS. Without being bound by any theory, differences between sesquiterpene synthases, particularly TEAS and CVS may be due to alterations of the active site architecture affecting the proton donor positioning for proton donation; and/or a dynamic mechanism wherein the intermediate compounds are brought into proper stereoelectronic alignment for the proton donation and cyclization geometry.

There now follows a description of the isolation, cloning, sequencing and functional characterization of citrus valencene synthase from *Citrus paradisi*. These examples are provided for the purpose of illustrating the invention, and are not to be considered as limiting.

Accordingly, one aspect of the present invention is an isolated nucleic acid comprising the nucleic acid sequence of SEQ ID NO 1. The nucleic acid is typically DNA, but can be RNA. If it is DNA, it is typically double-stranded, but can alternatively be single-stranded. Alternatively, it can be an RNA-DNA hybrid, i.e., either a complete or partial hybrid.

Another aspect of the present invention is an isolated nucleic acid that encodes the amino acid sequence of SEQ ID NO: 4. The nucleic acid is typically DNA, but can be RNA. If it is DNA, it is typically double-stranded, but can alternatively be single-stranded. Alternatively, it can be an RNA-DNA hybrid, i.e., either a complete or partial hybrid.

Another aspect of the present invention is an isolated nucleic acid encoding a protein that has the amino acid sequence of SEQ ID NO. 4 with 0 to 20 conservative amino acid substitutions, with the proviso that a conservative amino acid substitution is not made at amino acids 264, 273, 403, 404, 440, 444, 520, 527, and 528 in the protein that has the amino acid sequence of SEQ ID NO: 4, wherein the encoded protein specifically binds farnesyl pyrophosphate in a substantially hydrophobic pocket and has sesquiterpene synthase activity. Conservative amino acid substitutions are defined as above. Preferably, the nucleic acid sequence encodes a protein with 0 to 10 conservative amino acid substitutions. More preferably, the nucleic acid sequence encodes a protein with 0 to 5 conservative amino acid substitutions. Typically, the nucleic acid is DNA.

Yet another aspect of the present invention is an isolated nucleic acid encoding a protein that has the amino acid sequence of SEQ ID NO. 4 with 0 to 20 conservative amino acid substitutions, with the proviso that a conservative amino acid substitution is not made at amino acids 20, 264, 266, 270, 273, 298, 302, 372, 376, 401, 402, 403, 404, 407, 436, 440, 441, 444, 445, 448, 512, 515, 516, 519, 520, 527, 528, and 529 in the protein that has the amino acid sequence of SEQ ID NO: 4, wherein the encoded protein specifically binds farnesyl pyrophosphate in a substantially hydrophobic pocket and

has sesquiterpene synthase activity. Preferably, the nucleic acid sequence encodes a protein with 0 to 10 conservative amino acid substitutions. More preferably, the nucleic acid sequence encodes a protein with 0 to 5 conservative amino acid substitutions. Typically, the nucleic acid is DNA.

Yet another aspect of the invention is an isolated nucleic acid that hybridizes to the nucleic acid of SEQ ID NO: 1 under stringent conditions with no more than about a 5% mismatch. Preferably, there is no more than about a 2% mismatch. More preferably, there is no more than about a 1% mismatch. Typically, the nucleic acid is DNA.

Yet another aspect of the invention is an isolated nucleic acid that encodes a protein that is at least 500 amino acids in length, that specifically binds farnesyl pyrophosphate in a hydrophobic pocket, and that has sesquiterpene synthase activity, wherein the nucleic acid includes SEQ ID NO. 5 and SEQ ID NO. 6. Typically, the nucleic acid is DNA.

Still another aspect of the invention is an isolated nucleic acid that encodes a protein that is at least 500 amino acids in length, that specifically binds farnesyl pyrophosphate in a hydrophobic pocket, and that has sesquiterpene synthase activity, wherein the nucleic acid includes SEQ ID NO. 5. Typically, the nucleic acid is DNA.

Still another aspect of the invention is an isolated nucleic acid that encodes a protein that includes amino acid residues 264 to 528 of SEQ ID NO. 4, wherein the protein specifically binds farnesyl pyrophosphate in a hydrophobic pocket and has sesquiterpene synthase activity. Typically, the nucleic acid is DNA.

As defined above, another embodiment of the invention is an isolated polypeptide having appropriate enzymatic activity, namely sesquiterpene synthase activity. These isolated polypeptides include, but are not necessary limited to:

- (1) the polypeptide of SEQ ID NO: 4;
- (2) a polypeptide that has the amino acid sequence of SEQ ID NO. 4 with 0 to 20 conservative amino acid substitutions, with the proviso that a conservative amino acid substitution is not made at amino acids 264, 273, 403, 404, 440, 444, 520, 527, and 528 in the protein that has the amino acid sequence of SEQ ID NO: 4, wherein the encoded protein specifically binds farnesyl pyrophosphate in a substantially hydrophobic pocket and has sesquiterpene synthase activity, preferably with 0 to 10 conservative amino acid substitutions, more preferably with 0 to 5 conservative amino acid substitutions;
- (3) a polypeptide that has the amino acid sequence of SEQ ID NO. 4 with 0 to 20 conservative amino acid substitutions, with the proviso that a conservative amino acid substitution is not made at amino acids 20, 264, 266, 270, 273, 298, 302, 372, 376, 401, 402, 403, 404, 407, 436, 440, 441, 444, 445, 448, 512, 515, 516, 519, 520, 527, 528, and 529 in the protein that has the amino acid sequence of SEQ ID NO: 4, wherein the encoded protein specifically binds farnesyl pyrophosphate in a substantially hydrophobic pocket and has sesquiterpene synthase activity, preferably with 0 to 10 conservative amino acid substitutions, more preferably with 0 to 5 conservative amino acid substitutions;
- (4) a polypeptide encoded by an isolated nucleic acid that hybridizes to the nucleic acid of SEQ ID NO: 1 under stringent conditions with no more than about a 5% mismatch, preferably no more than about a 2% mismatch, more preferably no more than about a 1% mismatch;
- (5) a polypeptide that is at least 500 amino acids in length, that specifically binds farnesyl pyrophosphate in a hydrophobic pocket, and that has sesquiterpene syn-

these activity, wherein the sequence of the protein includes SEQ ID NO. 7 and SEQ ID NO. 8;

(6) a polypeptide that is at least 500 amino acids in length, that specifically binds farnesyl pyrophosphate in a hydrophobic pocket, and that has sesquiterpene synthase activity, wherein the sequence of the protein includes SEQ ID NO. 7; and

(7) a polypeptide that includes amino acid residues 264 to 528 of SEQ ID NO. 4, wherein the protein specifically binds farnesyl pyrophosphate in a hydrophobic pocket and has sesquiterpene synthase activity.

Typically, the sesquiterpene synthase activity of proteins or polypeptides within the scope of the present invention is valencene synthase activity, but other sesquiterpenes can be synthesized by proteins or polypeptides within the scope of the present invention, either as the major reaction product or in side reactions.

Another embodiment of the present invention is a vector comprising a nucleic acid of the present invention, as described above, operably linked to at least one control sequence. Typically, the control sequence is a promoter or enhancer. As described above, such vectors are well known in the art.

Yet another embodiment of the present invention is a host cell transformed or transfected with a vector of the present invention. The host cell can be a prokaryotic cell or a eukaryotic cell, such as a bacterial cell, a yeast cell, a plant cell, or an animal cell, such as an insect cell or a mammalian cell. Typically, the host cell transformed or transfected with the vector is capable of expressing the protein or polypeptide encoded by the vector. Suitable host cells for expression of polypeptides of the invention include prokaryotes, yeast or higher eukaryotic cells. Appropriate cloning and expression vectors for use with bacterial, fungal, yeast, and mammalian cellular hosts are described, for example, in Pouwels et al., *Cloning Vectors. A Laboratory Manual*, Elsevier, N.Y., (1985). Cell-free translation systems could also be employed to produce the disclosed polypeptides using RNAs derived from DNA constructs disclosed herein.

Prokaryotes include gram negative or gram positive organisms, for example, *E. coli* or Bacilli. Suitable prokaryotic host cells for transformation include, for example, *E. coli*, *Bacillus subtilis*, *Salmonella typhimurium*, and various other species within the genera *Pseudomonas*, *Streptomyces*, and *Staphylococcus*. In a prokaryotic host cell, such as *E. coli*, the polypeptides can include a N-terminal methionine residue to facilitate expression of the 16 recombinant polypeptide in the prokaryotic host cell. The N-terminal methionine can be cleaved from the expressed recombinant polypeptide.

Examples of useful expression vectors for prokaryotic host cells include those derived from commercially available plasmids such as the cloning vector pET plasmids (Novagen, Madison, Wis., USA) or yet pBR322 (ATCC 37017). pBR322 contains genes for ampicillin and tetracycline resistance and thus provides simple means for identifying transformed cells. To construct an expression vector using pBR322, an appropriate promoter and a DNA sequence encoding one or more of the polypeptides of the invention are inserted into the pBR322 vector. Other commercially available vectors include, for example, pKK223-3 (Pharmacia Fine Chemicals, Uppsala, Sweden) and pGEM-1 (Promega Biotec, Madison, Wis., USA). Other commercially available vectors include those that are specifically designed for the expression of proteins; these would include pMAL-p2 and pMAL-c2 vectors that are used for the expression of proteins fused to maltose binding protein (New England Biolabs, Beverly, Mass., USA).

Promoter sequences commonly used for recombinant prokaryotic host cell expression vectors include bacteriophage T7 promoter (Studier F. W. and Moffatt B. A., *J. Mol. Biol.* 189:113, 1986), β -lactamase (penicillinase), lactose promoter system (Chang et al., *Nature* 275:615, 1978; and Goeddel et al., *Nature* 281:544, 1979), tryptophan (trp) promoter system (Goeddel et al., *Nucl. Acids Res.* 8:4057, 1980; and EP-A-36776), and tac promoter (Maniatis, *Molecular Cloning: A Laboratory Manual*, Cold Spring Harbor Laboratory, p. 412, 1982). A particularly useful prokaryotic host cell expression system employs a phage λ PL promoter and a c1857ts thermolabile repressor sequence. Plasmid vectors available from the American Type Culture Collection ("ATCC"), which incorporate derivatives of the PL promoter, include plasmid pHUB2 (resident in *E. coli* strain JMB9 (ATCC 37092)) and pPLc28 (resident in *E. coli* RR1 (ATCC 53082)).

Polypeptides of the invention can also be expressed in yeast host cells, preferably from the *Saccharomyces* genus (e.g., *S. cerevisiae*). Other genera of yeast, such as *Pichia* or *Kluyveromyces* (e.g. *K. locus*), can also be employed. Yeast vectors will often contain an origin of replication sequence from a 2 μ yeast plasmid, an autonomously replicating sequence (ARS), a promoter region, sequences for polyadenylation, sequences for transcription termination, and a selectable marker gene. Suitable promoter sequences for yeast vectors include, among others, promoters for metallothionine, 3-phosphoglycerate kinase (Hitzeman et al., *J. Biol. Chem.* 255:2073, 1980), or other glycolytic enzymes (Hess et al., *J. Adv. Enzyme Reg.* 7:149, 1968; and Holland et al., *Biochem.* 17:4900, 1978), such as enolase, glyceraldehyde phosphate dehydrogenase, hexokinase, pyruvate decarboxylase, phosphofructokinase, glucose phosphate isomerase, 3-phosphoglycerate mutase, pyruvate kinase, triosephosphate isomerase, phosphoglucose isomerase, and glucokinase. Other suitable vectors and promoters for use in yeast expression are further described in Hitzeman, EPA-73, 657 or in Fleer et al., *Gene*, 107:285-195 (1991); and van den Berg et al., *Bio/Technology*, 8:135-139 (1990). Another alternative is the glucose-repressible ADH2 promoter described by Russell et al. (*J. Biol. Chem.* 258:2674, 1982) and Beier et al. (*Nature* 300:724, 1982). Shuttle vectors replicable in both yeast and *E. coli* can be constructed by inserting DNA sequences from pBR322 for selection and replication in *E. coli* (Ampr gene and origin of replication) into the above-described yeast vectors.

In one embodiment, mammalian or insect host cell culture systems are employed to express recombinant polypeptides of the invention. Baculovirus systems for production of heterologous proteins in insect cells are reviewed by Luckow and Summers, *Bio/Technology* 6:47 (1988). Established cell lines of mammalian origin also can be employed. Examples of suitable mammalian host cell lines include the COS-7 line of monkey kidney cells (ATCC CRL 1651) (Gluzman et al., *Cell* 23:175, 1981), L cells, C127 cells, 3T3 cells (ATCC CCL 163), Chinese hamster ovary (CHO) cells, HeLa cells, and BHK (ATCC CRL 10) cell lines, and the CV-1/EBNA-1 cell line (ATCC CRL 10478) derived from the African green monkey kidney cell line CV1 (ATCC CCL 70) as described by McMahan et al. (*EMBO J.* 10: 2821, 1991).

In one embodiment, transfected DNA is integrated into a chromosome of a non-human organism such that a stable recombinant system results. Any chromosomal integration method known in the art may be used in the practice of the invention, including but not limited to, recombinase-mediated cassette exchange (RMCE), viral site specific chromosomal insertion, adenovirus, and pronuclear injection.

Yet another embodiment of the present invention is a non-human multicellular organism that is modified to harbor a nucleic acid of the present invention such that the nucleic acid can be expressed in the non-human multicellular organism. The non-human multicellular organism can be, but is not necessarily limited to, a plant or insect. The plant can be the tobacco plant. The non-human organism can be modified by any of the methods known in the art for gene transfer and the production of transgenic organisms, including, but not necessarily limited to, the use of delivery devices such as lipids and viral vectors, naked DNA, electroporation, and particle-mediated gene transfer.

Established methods for introducing DNA into mammalian cells have been described (Kaufman, R. J., *Large Scale Mammalian Cell Culture*, 1990, pp. 15-69). Additional protocols using commercially available reagents, such as Lipofectamine (Gibco/BRL) or Lipofectamine-Plus, can be used to transfect cells (Felgner et al., *Proc. Natl. Acad. Sci. USA* 84:7413-7417, 1987). In addition, electroporation can be used to transfect mammalian cells using conventional procedures, such as those in Sambrook et al. *Molecular Cloning: A Laboratory Manual*, 2 ed. Vol. 1-3, Cold Spring Harbor Laboratory Press, 1989). Selection of stable transformants can be performed using resistance to cytotoxic drugs as a selection method. Kaufman et al., *Meth. in Enzymology* 185: 487-511, 1990, describes several selection schemes, such as dihydrofolate reductase (DHFR) resistance. A suitable host strain for DHFR selection can be CHO strain DX-131 1, which is deficient in DHFR (Urlaub and Chasin, *Proc. Natl. Acad. Sci. USA* 77:4216-4220, 1980). A plasmid expressing the DHFR cDNA can be introduced into strain DX-131 1, and only cells that contain the plasmid can grow in the appropriate selective media.

Transcriptional and translational control sequences for mammalian host cell expression vectors can be excised from viral genomes. Commonly used promoter sequences and enhancer sequences are derived from polyoma virus, adenovirus 2, simian virus 40 (SV40), and human cytomegalovirus. DNA sequences derived from the SV40 viral genome, for example, SV40 origin, early and later promoter, enhancer, splice, and polyadenylation sites can be used to provide other genetic elements for expression of a structural gene sequence in a mammalian host cell. Viral early and late promoters are particularly useful because both are easily obtained from a viral genome as a fragment, which can also contain a viral origin of replication (Fiers et al., *Nature* 273:113, 1978; Kaufman, *Meth. in Enzymology*, 1990).

There are several methods known in the art for the creation of transgenic plants. These include, but are not limited to: electroporation of plant protoplasts, liposome-mediated transformation, polyethylene-glycol-mediated transformation, microinjection of plant cells, and transformation using viruses. In one embodiment, direct gene transfer by particle bombardment is utilized.

Direct gene transfer by particle bombardment provides an example for transforming plant tissue. In this technique a particle, or microprojectile, coated with DNA is shot through the physical barriers of the cell. Particle bombardment can be used to introduce DNA into any target tissue that is penetrable by DNA coated particles, but for stable transformation, it is imperative that regenerable cells be used. Typically, the particles are made of gold or tungsten. The particles are coated with DNA using either CaCl_2 or ethanol precipitation methods which are commonly known in the art.

DNA coated particles are shot out of a particle gun. A suitable particle gun can be purchased from Bio-Rad Laboratories (Hercules, Calif.). Particle penetration is controlled

by varying parameters such as the intensity of the explosive burst, the size of the particles, or the distance particles must travel to reach the target tissue.

The DNA used for coating the particles may comprise an expression cassette suitable for driving the expression of the gene of interest that will comprise a promoter operably linked to the gene of interest.

Methods for performing direct gene transfer by particle bombardment are disclosed in U.S. Pat. No. 5,990,387 to Tomes et al., incorporated herein by this reference.

In one embodiment, the cDNAs of the invention may be expressed in such a way as to produce either sense or antisense RNA. Antisense RNA is RNA that has a sequence which is the reverse complement of the mRNA (sense RNA) encoded by a gene. A vector that will drive the expression of antisense RNA is one in which the ONA is placed in "reverse orientation" with respect to the promoter such that the non-coding strand (rather than the coding strand) is transcribed. The expression of antisense RNA can be used to down-modulate the expression of the protein encoded by the mRNA to which the antisense RNA is complementary. Vectors producing antisense RNAs could be used to make transgenic plants, as described above.

In one embodiment, transfected DNA is integrated into a chromosome of a non-human organism such that a stable recombinant system results. Any chromosomal integration method known in the art may be used in the practice of the invention, including but not limited to, recombinase-mediated cassette exchange (RMCE), viral site specific chromosomal insertion, adenovirus, and pronuclear injection.

Still another embodiment of the invention is a method of producing a sesquiterpene synthase comprising the steps of:

- (1) culturing a host cell transformed or transfected by a vector of the present invention under conditions in which the sesquiterpene synthase encoded by the vector is expressed; and
- (2) isolating the sesquiterpene synthase expressed by the host cell.

Typically, the sesquiterpene synthase is valencene synthase, as described above.

Yet another embodiment of the invention is a method of producing a sesquiterpene comprising the steps of:

- (1) reacting a protein or polypeptide of the present invention having sesquiterpene synthase enzymatic activity with farnesyl pyrophosphate under conditions in which the enzymatic activity catalyzes the formation of a sesquiterpene; and
- (2) isolating the sesquiterpene formed in step (1).

Typically, the sesquiterpene synthase is a valencene synthase, in which case the sesquiterpene formed is valencene.

Still another embodiment of the present invention is a method of producing nootkatone. In one alternative of this embodiment, the method comprises the steps of:

- (1) reacting a protein or polypeptide of the present invention having valencene synthase enzymatic activity with farnesyl pyrophosphate under conditions in which the enzymatic activity catalyzes the formation of valencene;
- (2) reacting the valencene formed in step (1) by regio-specific hydroxylation and then oxidation to form nootkatone; and
- (3) isolating the nootkatone produced.

The second step could be catalyzed by a single multifunctional hydroxylase or could be catalyzed by sequential enzyme mediated reactions.

Yet another embodiment of the invention is an antisense nucleic acid that is the complement of a nucleic acid according to the present invention as described above.

EXAMPLES

A cDNA of 1710 bp was prepared from fresh grapefruit via RT-PCR using non-degenerate primers designed to amplify from the translation start to stop sites of AF411120, then inserted into appropriate vectors for DNA sequencing and bacterial expression. For enzymological studies, the cDNA was inserted into a pET expression vector in-frame with an amino terminal hexa-histidine tag, and lysates of appropriately grown bacterial cultures used for nickel-affinity purification of the citrus cDNA encoded protein. The isolated citrus protein was approximately 30-40% pure as determined by Coomassie blue staining after SDS-PAGE and migrated as a 64 kD polypeptide (data not shown). The conceptual translation of the isolated cDNA predicted a protein of 548 amino acids having a molecular size of 63,646 daltons.

Example 1

Molecular Cloning of a *Citrus sesquiterpene*
Synthase cDNA

RNA was isolated from the juice vesicles of freshly harvested red grapefruit using TRIZOL reagent and following the manufacturer's protocol (Invitrogen Corp., Carlsbad, Calif.). Reverse transcription of the isolated RNA also followed manufacturer's protocol, and 10 µg of total RNA was reverse transcribed using Superscript II RNase H (Stratagene, La Jolla Calif.) and a reverse primer (18 nucleotides in length) complementary to the 3' end including the stop codon of the AF411120 sequence reported in Genbank. A full-length cDNA was then amplified using Pfu turbo Taq polymerase (Stratagene, La Jolla Calif.), more of the initial reverse primer, a forward primer (18 nucleotides in length) complementary to the 5' end initiating at the start codon, and an aliquot of the first strand cDNA using standard PCR conditions. An amplification fragment approximating the expected size of 1,800 bp was observed by agarose gel electrophoresis, T/A cloned into the pGem T-easy vector (Promega, Madison, Wis.) and then subjected to automated DNA sequencing using the BigDye terminator system with an ABI 310 sequencer. The sequence, which is shown in FIG. 5, was obtained from start to stop codons in duplicate and rectified with the reported sequence for the *Citrus x paradisi* putative terpene synthase mRNA. (Seq ID: 1). One nucleotide was found to be altered in the recovered cDNA, 1544 bp 5' to the ATG start site an A instead of a G was observed, converting codon 492 from specifying an aspartate to an arginine. Sequence analysis was performed using software tools available from the NCBI web site (www.ncbi.nlm.nih.gov) or using ClustalX.

Example 2

Expression of *Citrus valencene* Synthase in *E. coli*

The *Citrus valencene* synthase (CVS) cDNA was inserted into an appropriate expression vector, pGEM to provide an amino-terminal hexa-histidyl tag for protein purification after expression of the putative valencene synthase cDNA in *E. coli*. The cDNA was re-amplified using PCR primers designed to amplify from the ATG start codon (5'-GGG-GAATTCATCTGGTCTGGAGAAACATTTCTGCC-3' (SEQ ID NO.: 2) to the TGA stop codon (5'-CCGCTCGAG-GAAGTATAGAACTAGTCGTCAAAATGG-3' (SEQ ID NO.:3)) and to provide restriction sites EcoRI and XhoI, (NEB, Beverly Mass.) respectively, using the pGEM-CVS

plasmid as template under standard PCR conditions. The PCR product was digested with EcoRI and XhoI, purified using a QIAquick PCR purification kit (Qiagen, Valencia, Calif.), and ligated into a pET-28a(+) expression vector (Novagen, San Diego, Calif.) that had been digested with corresponding enzymes (NEB, Beverly Mass.), dephosphorylated with calf intestine alkaline phosphatase (Invitrogen, Carlsbad, Calif.), and purified with a QIAquick kit. Proper construction of the resulting expression vector was verified by DNA sequencing and subsequently referred to as the pET-28a(+)-CVS expression vector. The expression vector was transformed into BL21 (DE3) cells (Stratagene, La Jolla, Calif.), and grown in a 10 mL inoculate of LB Growth Media, using well known techniques. The cells were grown to an OD₆₀₀=1. IPTG (Sigma-Aldrich, St. Louis, Mo.) was used to induce expression of the putative valencene synthase gene. The transformed cells were incubated at 28° C. for four hours, centrifuged to form a pellet and the pellet was collected and resuspended in 1 mL of cyclase buffer (200 mM Tris-HCl, pH 7.5, 40 mM MgCl₂). Cells were then sonicated with three 10-sec bursts using a microtip ultrasonicator at 40% power and the lysate was centrifuged at 10,000×g for 10 minutes.

Samples of the lysate were divided and some samples were assayed for protein purity, while other samples were used in Example 3 below. For the protein purity assay, protein was purified by nickel affinity chromatography according to Mathis et al. (J. R. Mathis, K. Back, C. Starks, J. Noel, C. D. Poulter, J. Chappell, Biochem. 36 (1997) 8340-8348). Isolated CVS protein was approximately 30 to 40% pure based on the intensity of Coomassie blue staining of samples analyzed by SDS-PAGE. A protein sequencing reaction was performed via the Edman Degradation reaction performed using a Perkin Elmer Applied Biosystems Model 494 Precise protein/peptide sequencer with an on-line Perkin Elmer Applied Biosystems Model 140C PTH Amino Acid Analyzer. The protein sequence for the expressed CVS protein is shown in FIG. 6. (SEQ ID NO.: 4)

Partially purified protein was subsequently incubated with FPP at pH 7.5 using typical sesquiterpene synthase reaction conditions and the pentane extractable products examined by GC-MS (FIG. 7). Two compounds accounting for greater than 95% of the total reaction products dominated the GC profiles and were identified as beta-elemene (30%) and valencene (65%) on the basis of MS matches with authentic standards. The beta-elemene peak most certainly represents a thermal rearrangement product of germacrene A resulting from high temperature injection into the GC.

Example 3

Terpene Synthase Activity Assays and Reaction
Product Identification

Small scale reactions of 50 µL were used for screening purposes and rate determinations. Reactions typically contained 200 mM Tris-HCl, pH 7.5, 40 mM MgCl₂, 0.5 µCi [1-3H]FPP, 25-30 µM FPP and 160 nM enzyme. For kinetic determinations, 10 µL aliquots of FPP (giving final concentrations of 0.7-23 µM) were rapidly mixed with 40 µL of enzyme solution at room temperature (23° C.) and allowed to incubate for 1 minute. The reactions were terminated by addition of 150 µL of a 100 mM KOH, 0.5 M EDTA stop solution. Reactions were extracted with 500 µL of hexane and an aliquot was taken for determination of radiolabeled hydrophobic product via liquid scintillation counting. Hexane

extracted samples were not subject to silica chromatography prior to counting because background was minimal and synthase mutants could possibly produce reaction products containing alcohols, which would bind to silica. Kinetic constants were determined from direct fits of the Michaelis-Menton equation to the data using Graphpad Prism 2.01 software.

Initial synthase reaction products were examined by GC-MS. Preparative reactions were performed similarly to the reactions described above except they were scaled to 2.5 mL and employed 2 μ M of purified enzyme and 80 μ M of unlabeled FPP. The reactions were incubated for 1 hr and then extracted twice with 2 mL of pentane. Pooled extracts were dried to 50 μ L under a stream of nitrogen gas and 1 μ L aliquots of this organic extract were analyzed by GC-MS using an HP-GCDplus (Hewlett-Packard, Palo Alto, Calif.) equipped with a DB-5 ms column, an injector temperature of 250° C. and the mass selective detector set to scan for ions within the range from 45 to 250 m/z. The GC was programmed to hold for 1 minute at 100° C., followed by a temperature ramp of 8° C./min to a final temperature of 270° C. The results are shown in FIG. 7. Spectra from this analysis was compared to the NIST library standard for beta-elemene (FIG. 8) and to that for a sample of valencene purchased from Fluka Chemical Company (FIG. 9).

The dominance of valencene as a reaction product under these conditions was sufficient to classify the *C. paradisi* cDNA as citrus valencene synthase, CVS. However, the amount of germacrene A, a putative reaction intermediate (FIG. 1), generated by the CVS enzyme was atypical relative to previous studies with TEAS and HPS. Neither of these enzymes release appreciable amounts of reaction intermediates. Further consideration of the acidic conditions likely to exist within juice vesicles/sacs where sesquiterpenes accumulate in citrus fruit suggested that the CVS enzyme might have a pH optimal different from other terpene synthases.

As shown in FIG. 10, virtually no germacrene A was detected as a reaction product between pH 6.0 to 7.0, with optimal valencene biosynthesis occurring at pH 7.0 (FIG. 10). In contrast, germacrene A biosynthesis was optimal at pH 8.5 with a relatively sharp transition point from valence as the dominate reaction product to germacrene A at pH 8.0.

Example 4

pH Dependence Assays

Reactions to determine pH dependence were performed in glass GC-vials with 100 nM of purified enzyme, 20 mM MgCl₂, 50 μ M FPP (Echelon; Salt Lake City, Utah), and 100 mM buffer at various pH values; total reaction volume 500 μ L. The buffers were chosen within one pH unit of the buffer's pKa (pH 5-5.5, acetate; pH 6-7, MES; pH 7.5-9, Tris; pH 9.5-10, ethanolamine; pH 10.5, CAPS). Reactions were allowed to proceed at room temperature for 30 min prior to overlaying with 500 μ L of ethylacetate, vortexing for 10 sec, then direct analysis of the reaction products using an HP 6890 gas chromatograph with a 5973 mass spectrometer (Agilent Technologies, Palo Alto, Calif.) with an auto-sampler programmed to remove 1 μ L samples only from the organic phase. GC separations were performed on a 5%-phenyl-me-

thylpolysiloxane column (J&W Scientific, Folsom, Calif.) of 30 m \times 0.25 mm i.d. \times 0.25 m thickness with He as the carrier gas at 2 mL/min and a temperature gradient of 10° C./min from 50° C. (5-min hold) to 180° C. (4-min hold). GC/MS data was analyzed using HP-Chemstation software (version B.01.00) and integration of terpene peaks used for quantification.

As seen in FIG. 10, a pH dependent transition between valencene and germacrene A biosynthesis occurs at approximately pH 8.2. Inventors interpreted this as a titration point for the protonation of germacrene A. This pH value is close to the pKa value of 8.3-8.5 for free cysteine, which is involved in the second protonation step of the chemical cascade catalyzed by TEAS. Those results demonstrated that when C440 of TEAS was mutated to alanine, a robust germacrene A synthetic activity resulted. In combination with the sequence data provided in SEQ ID NO.: 7 and SEQ ID NO.: 8, it is apparent that germacrene synthases consistently differ from CVS, TEAS and HPS by the absence of cysteine at position 440 in their putative active site while Y520, previously implicated in the same protonation step, is conserved. The observation of germacrene A as a reaction product and the pH dependence for its synthesis are consistent with its intermediacy in the reaction catalyzed by CVS.

The inventions illustratively described herein can suitably be practiced in the absence of any element or elements, limitation or limitations, not specifically disclosed herein. Thus, for example, the terms "comprising," "including," "containing," etc. shall be read expansively and without limitation. Additionally, the terms and expressions employed herein have been used as terms of description and not of limitation, and there is no intention in the use of such terms and expressions of excluding any equivalents of the future shown and described or any portion thereof, and it is recognized that various modifications are possible within the scope of the invention claimed. Thus, it should be understood that although the present invention has been specifically disclosed by preferred embodiments and optional features, modification and variation of the inventions herein disclosed can be resorted to by those skilled in the art, and that such modifications and variations are considered to be within the scope of the inventions disclosed herein. The inventions have been described broadly and generically herein. Each of the narrower species and subgeneric groupings falling within the scope of the generic disclosure also form part of these inventions. This includes the generic description of each invention with a proviso or negative limitation removing any subject matter from the genus, regardless of whether or not the excised materials specifically resided therein.

In addition, where features or aspects of an invention are described in terms of the Markush group, those schooled in the art will recognize that the invention is also thereby described in terms of any individual member or subgroup of members of the Markush group. It is also to be understood that the above description is intended to be illustrative and not restrictive. Many embodiments will be apparent to those of in the art upon reviewing the above description. The scope of the invention should therefore, be determined not with reference to the above description, but should instead be determined with reference to the appended claims, along with the full scope of equivalents to which such claims are entitled. The disclosures of all articles and references, including patent publications, are incorporated herein by reference.

SEQUENCE LISTING

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545																

We claim:

1. A method of producing valencene, comprising:
 - a) reacting a heterologous polypeptide having valencene synthase enzymatic activity with farnesyl pyrophosphate under conditions in which the enzymatic activity catalyzes the formation of a valencene, wherein: the reaction of the polypeptide with farnesyl pyrophosphate is performed at pH between about 6.0 and 7.0; and the polypeptide comprises the sequence of amino acids set forth in SEQ ID NO: 4; and
 - (b) isolating the valencene formed in step (a).
2. The method of claim 1, wherein the reaction of said polypeptide with farnesyl pyrophosphate is conducted at about pH 7.
3. The method of claim 1, further comprising producing a sesquiterpene comprising the steps of:
 - (c) reacting the valencene in step (b) by regiospecific hydroxylation and then oxidation to form the sesquiterpene; and
 - (d) isolating the sesquiterpene produced in step (c), wherein the sesquiterpene is nootkatone.
4. The method of claim 3, wherein step (c) is catalyzed by a single multifunctional hydroxylase.

5. The method of claim 3, wherein step (c) is catalyzed by sequential enzyme mediated reactions.
6. A method of producing valencene comprising the steps of:
 - (a) reacting a heterologous polypeptide having valencene synthase enzymatic activity with farnesyl pyrophosphate under conditions in which the enzymatic activity catalyzes the formation of a valencene, wherein the polypeptide comprises the sequence of amino acids set forth in SEQ ID NO:4; and
 - (b) isolating the valencene formed in step (a).
7. The method of claim 6, comprising further producing a sesquiterpene comprising the steps of:
 - (c) reacting the valencene in step (b) by regiospecific hydroxylation and then oxidation to form the sesquiterpene; and
 - (d) isolating the sesquiterpene produced in step (c), wherein the sesquiterpene is nootkatone.
8. The method of claim 7, wherein step (c) is catalyzed by a single multifunctional hydroxylase.
9. The method of claim 7, wherein step (c) is catalyzed by sequential enzyme mediated reactions.

* * * * *

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 8,192,950 B2
APPLICATION NO. : 12/259497
DATED : June 5, 2012
INVENTOR(S) : Chappell et al.

Page 1 of 1

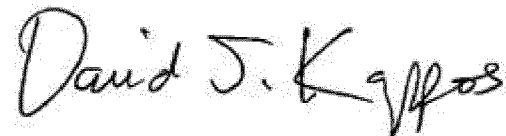
It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

On the Title Page:

The first or sole Notice should read --

Subject to any disclaimer, the term of this patent is extended or adjusted under 35 U.S.C. 154(b)
by 532 days.

Signed and Sealed this
Second Day of October, 2012

A handwritten signature in black ink that reads "David J. Kappos". The signature is written in a cursive, slightly slanted style.

David J. Kappos
Director of the United States Patent and Trademark Office

UNITED STATES PATENT AND TRADEMARK OFFICE
CERTIFICATE OF CORRECTION

PATENT NO. : 8,192,950 B2
APPLICATION NO. : 12/259497
DATED : June 5, 2012
INVENTOR(S) : Chappell et al.

Page 1 of 1

It is certified that error appears in the above-identified patent and that said Letters Patent is hereby corrected as shown below:

IN THE SPECIFICATION:

Column 1, line 8-11, please replace “benefit of priority under 35 U.S.C. 120 is claimed herein to U.S. Utility Application No. 10/899,356, filed on July 26, 2006, now U.S. Patent No. 7,442,758, issued October 28, 2008” should read --benefit of priority under 35 U.S.C. 120 is claimed herein to U.S. Utility Application No. 10/899,356, filed on July 26, 2004, now U.S. Patent No. 7,442,785, issued October 28, 2008--.

Signed and Sealed this
Eleventh Day of March, 2014



Michelle K. Lee
Deputy Director of the United States Patent and Trademark Office