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(54) Titre : DRIMENOL SYNTHASES III
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(57) **Abrégé/Abstract:**

The present invention discloses a method of producing drimenol and/or drimenol derivatives by contacting a polypeptide with farnesyl pyrophosphate. Also provided is an amino acid sequence of a polypeptide useful in the methods of the invention and nucleic acid encoding the polypeptides of the invention. The method further provides host cells or organisms genetically modified to express the polypeptides of the invention and useful to produce drimenol and /or drimenol derivatives.

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(54) **Title:** DRIMENOL SYNTHASES III

(57) **Abstract:** The present invention discloses a method of producing drimenol and/or drimenol derivatives by contacting a polypeptide with farnesyl pyrophosphate. Also provided is an amino acid sequence of a polypeptide useful in the methods of the invention and nucleic acid encoding the polypeptides of the invention. The method further provides host cells or organisms genetically modified to express the polypeptides of the invention and useful to produce drimenol and /or drimenol derivatives.



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DRIMENOL SYNTHASES III

Technical field

The field relates to a method of producing drimenol, said method comprising contacting a polypeptide with farnesyl pyrophosphate (FPP). In particular, said method may be carried out *in vitro* or *in vivo* to produce drimenol, a very useful compound in the field of perfumery. Also provided herein is an amino acid sequence of a polypeptide useful in the methods provided herein. A nucleic acid encoding the polypeptide of an embodiment herein and an expression vector containing said nucleic acid are provided herein. A non-human host organism or a cell transformed to be used in the method of producing drimenol is also provided herein.

Background

Terpenes are found in most organisms (microorganisms, animals and plants). These compounds are made up of five carbon units called isoprene units and are classified by the number of these units present in their structure. Thus monoterpenes, sesquiterpenes and diterpenes are terpenes containing 10, 15 and 20 carbon atoms, respectively. Sesquiterpenes, for example, are widely found in the plant kingdom. Many sesquiterpene molecules are known for their flavor and fragrance properties and their cosmetic, medicinal and antimicrobial effects. Numerous sesquiterpene hydrocarbons and sesquiterpenoids have been identified.

Biosynthetic production of terpenes involves enzymes called terpene synthases. There is virtually an infinity of sesquiterpene synthases present in the plant kingdom, all using the same substrate (farnesyl pyrophosphate, FPP) but having different product profiles. Genes and cDNAs encoding sesquiterpene synthases have been cloned and the corresponding recombinant enzymes characterized.

Currently the main sources for drimenol are plants naturally containing drimenol and the contents of drimenol in these natural sources are low. Chemical synthesis approaches have been developed but are still complex and not cost-effective.

Summary

Provided herein is a method of producing drimenol comprising:

- i) contacting an acyclic farnesyl diphosphate (FPP) precursor with a polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2 or comprising SEQ ID NO:2 to produce the drimenol; and

ii) optionally isolating the drimenol.

Also provided herein is an isolated polypeptide having drimenol synthase activity comprising SEQ ID NO: 2.

Further provided herein is an isolated nucleic acid molecule encoding a polypeptide having
5 drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%,
95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2 or comprising SEQ ID NO:2

Further provided is an isolated nucleic acid molecule encoding a polypeptide provided herein
where the molecule has a nucleotide sequence 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%,
95%, 96%, 97%, 98% or 99% sequence identity to a nucleotide sequence selected from the group
10 consisting of SEQ ID NO: 1 and SEQ ID NO: 5 or comprising the nucleotide sequence selected from
the group consisting of SEQ ID NO: 1 and SEQ ID NO: 5

Description of the drawings.

15 **Figure 1.** Structure of (-)-drimenol

Figure 2. Mass spectrum of authentic (-)-drimenol

Figure 3. ¹³C NMR spectrum of authentic (-)-drimenol.

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Figure 4. X-Ray (Cu K α radiation) structure of authentic (-)-drimenol

Figure 5. Shows GC/MS extracted ion chromatogram (m/z 109) of *Paeonia anomala* root extract
(dichloromethane). The arrow denotes the peak of drimenol.

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Figure 6. Shows GC/MS analysis (extracted ion chromatogram at m/z 109) of transient expression
experiment of the isolated drimenol synthase on tobacco leaves. B. Mass spectrum of the peak at
32.442 min is drimenol. C. Mass spectrum of authentic sample of drimenol.

30 **Detailed Description**

For the descriptions herein and the appended claims, the use of "or" means "and/or" unless
stated otherwise. Similarly, "comprise," "comprises," "comprising", "include," "includes," and
"including" are interchangeable and not intended to be limiting.

It is to be further understood that where descriptions of various embodiments use the term "comprising," those skilled in the art would understand that in some specific instances, an embodiment can be alternatively described using language "consisting essentially of" or "consisting of." In one aspect, provided here is a method of producing drimenol comprising:

5 i) contacting an acyclic terpene pyrophosphate, particularly farnesyl diphosphate (FPP)) with a polypeptide having drimenol synthase activity and comprising SEQ ID NO: 2 to produce drimenol; and

ii) optionally isolating the drimenol.

In one aspect, the drimenol is isolated.

10 In another aspect provided here, the drimenol is produced with greater than or equal to, 60%, 80%, or 90% or even 95% selectivity.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

15 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

20 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

25 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

30 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

5 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

10 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 97%, 98% or 99% sequence identity to SEQ ID NO: 2.

15 In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 98% or 99% sequence identity to SEQ ID NO: 2.

In one embodiment provided herein is an isolated polypeptide a polypeptide having drimenol synthase activity and having at least 99% sequence identity to SEQ ID NO: 2.

Further provided here is an isolated polypeptide having drimenol activity comprising an amino acid of SEQ ID NO: 2.

20 Further provided herein is an isolated nucleic acid molecule encoding a polypeptide comprising an amino acid sequence of SEQ ID NO: 2.

Further provided herein a nucleic acid molecule comprising the sequence SEQ ID NO: 1 or SEQ ID NO: 5.

25 Further provided here is a method as recited in claim 1 comprising the steps of transforming a host cell or non-human organism with a nucleic acid encoding a polypeptide a polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2 or comprising SEQ ID NO: 2 and culturing the host cell or organism under conditions that allow for the production of the polypeptide.

30 Further provided is at least one vector comprising the nucleic acid molecules described herein.

Further provided herein is a vector selected from the group of a prokaryotic vector, viral vector and a eukaryotic vector.

Further provided here is a vector that is an expression vector.

As a “drimenol synthase” or as a “polypeptide having a drimenol synthase activity”, we mean

here a polypeptide capable of catalyzing the synthesis of drimenol, in the form of any of its stereoisomers or a mixture thereof, starting from an acyclic terpene pyrophosphate, particularly FPP. Drimenol may be the only product or may be part of a mixture of sesquiterpenes.

The ability of a polypeptide to catalyze the synthesis of a particular sesquiterpene (for example drimenol) can be simply confirmed by performing the enzyme assay as detailed in Examples 1 to 2.

According to the present invention, polypeptides are also meant to include truncated polypeptides provided that they keep their drimenol synthase activity.

As intended herein below, "a nucleotide sequence obtained by modifying SEQ ID NO: 1 or SEQ ID NO: 5 the complement thereof" encompasses any sequence that has been obtained by changing the sequence of SEQ ID NO: 1 or SEQ ID NO: 5, or of the complement thereof using any method known in the art, for example by introducing any type of mutations such as deletion, insertion or substitution mutations. Examples of such methods are cited in the part of the description relative to the variant polypeptides and the methods to prepare them.

15 Abbreviations used

bp	base pair
kb	kilo base
BSA	bovine serum albumin
20 DNA	deoxyribonucleic acid
cDNA	complementary DNA
DTT	dithiothreitol
FID	flame ionization detector
FPP	farnesyl pyrophosphate
25 GC	gas chromatograph
IPTG	isopropyl-D-thiogalacto-pyranoside
LB	lysogeny broth
MS	mass spectrometer / mass spectrometry
MVA	mevalonic acid
30 PCR	polymerase chain reaction
RMCE	recombinase-mediated cassette exchange
3'-/5'-RACE	3' and 5' rapid amplification of cDNA ends
RNA	ribonucleic acid
mRNA	messenger ribonucleic acid

miRNA	micro RNA
siRNA	small interfering RNA
rRNA	ribosomal RNA
tRNA	transfer RNA

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The term "polypeptide" means an amino acid sequence of consecutively polymerized amino acid residues, for instance, at least 15 residues, at least 30 residues, at least 50 residues. In some embodiments of an embodiment herein, a polypeptide comprises an amino acid sequence that is an enzyme, or a fragment, or a variant thereof.

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The term "isolated" polypeptide refers to an amino acid sequence that is removed from its natural environment by any method or combination of methods known in the art and includes recombinant, biochemical and synthetic methods.

15

The term "protein" refers to an amino acid sequence of any length wherein amino acids are linked by covalent peptide bonds, and includes oligopeptide, peptide, polypeptide and full length protein whether naturally occurring or synthetic.

The terms "drimenol synthase" or "drimenol synthase protein" refer to an enzyme that is capable of converting farnesyl diphosphate (FPP) to drimenol.

The terms "biological function," "function," "biological activity" or "activity" refer to the ability of the drimenol synthase provided herein to catalyze the formation of drimenol from FPP.

20

The terms "nucleic acid sequence," "nucleic acid," and "polynucleotide" are used interchangeably meaning a sequence of nucleotides. A nucleic acid sequence may be a single-stranded or double-stranded deoxyribonucleotide, or ribonucleotide of any length, and include coding and non-coding sequences of a gene, exons, introns, sense and anti-sense complimentary sequences, genomic DNA, cDNA, miRNA, siRNA, mRNA, rRNA, tRNA, recombinant nucleic acid sequences, isolated and purified naturally occurring DNA and/or RNA sequences, synthetic DNA and RNA sequences, fragments, primers and nucleic acid probes. The skilled artisan is aware that the nucleic acid sequences of RNA are identical to the DNA sequences with the difference of thymine (T) being replaced by uracil (U).

25

An "isolated nucleic acid" or "isolated nucleic acid sequence" is defined as a nucleic acid or nucleic acid sequence that is in an environment different from that in which the nucleic acid or nucleic acid sequence naturally occurs. The term "naturally-occurring" as used herein as applied to a nucleic acid refers to a nucleic acid that is found in a cell in nature. For example, a nucleic acid sequence that is present in an organism, for instance in the cells of an organism, that can be isolated

from a source in nature and which has not been intentionally modified by a human in the laboratory, is naturally occurring.

"Recombinant nucleic acid sequences" are nucleic acid sequences that result from the use of laboratory methods (molecular cloning) to bring together genetic material from more than one source, creating a nucleic acid sequence that does not occur naturally and would not be otherwise found in biological organisms.

"Recombinant DNA technology" refers to molecular biology procedures to prepare a recombinant nucleic acid sequence as described, for instance, in Laboratory Manuals edited by Weigel and Glazebrook, 2002 Cold Spring Harbor Lab Press; and Sambrook et al., 1989 Cold Spring Harbor, NY: Cold Spring Harbor Laboratory Press.

The term "gene" means a DNA sequence comprising a region, which is transcribed into a RNA molecule, e.g., an mRNA in a cell, operably linked to suitable regulatory regions, e.g., a promoter. A gene may thus comprise several operably linked sequences, such as a promoter, a 5' leader sequence comprising, e.g., sequences involved in translation initiation, a coding region of cDNA or genomic DNA, introns, exons, and/or a 3' non-translated sequence comprising, e.g., transcription termination sites.

A "chimeric gene" refers to any gene which is not normally found in nature in a species, in particular, a gene in which one or more parts of the nucleic acid sequence are present that are not associated with each other in nature. For example the promoter is not associated in nature with part or all of the transcribed region or with another regulatory region. The term "chimeric gene" is understood to include expression constructs in which a promoter or transcription regulatory sequence is operably linked to one or more coding sequences or to an antisense, i.e., reverse complement of the sense strand, or inverted repeat sequence (sense and antisense, whereby the RNA transcript forms double stranded RNA upon transcription).

A "3' UTR" or "3' non-translated sequence" (also referred to as "3' untranslated region," or "3'end") refers to the nucleic acid sequence found downstream of the coding sequence of a gene, which comprises for example a transcription termination site and (in most, but not all eukaryotic mRNAs) a polyadenylation signal such as AAUAAA or variants thereof. After termination of transcription, the mRNA transcript may be cleaved downstream of the polyadenylation signal and a poly(A) tail may be added, which is involved in the transport of the mRNA to the site of translation, e.g., cytoplasm.

"Expression of a gene" involves transcription of the gene and translation of the mRNA into a protein. Overexpression refers to the production of the gene product as measured by levels of

mRNA, polypeptide and/or enzyme activity in transgenic cells or organisms that exceeds levels of production in non-transformed cells or organisms of a similar genetic background.

"Expression vector" as used herein means a nucleic acid molecule engineered using molecular biology methods and recombinant DNA technology for delivery of foreign or exogenous DNA into a host cell. The expression vector typically includes sequences required for proper transcription of the nucleotide sequence. The coding region usually codes for a protein of interest but may also code for an RNA, e.g., an antisense RNA, siRNA and the like.

An "expression vector" as used herein includes any linear or circular recombinant vector including but not limited to viral vectors, bacteriophages and plasmids. The skilled person is capable of selecting a suitable vector according to the expression system. In one embodiment, the expression vector includes the nucleic acid of an embodiment herein operably linked to at least one regulatory sequence, which controls transcription, translation, initiation and termination, such as a transcriptional promoter, operator or enhancer, or an mRNA ribosomal binding site and, optionally, including at least one selection marker. Nucleotide sequences are "operably linked" when the regulatory sequence functionally relates to the nucleic acid of an embodiment herein.

"Regulatory sequence" refers to a nucleic acid sequence that determines expression level of the nucleic acid sequences of an embodiment herein and is capable of regulating the rate of transcription of the nucleic acid sequence operably linked to the regulatory sequence. Regulatory sequences comprise promoters, enhancers, transcription factors, promoter elements and the like.

"Promoter" refers to a nucleic acid sequence that controls the expression of a coding sequence by providing a binding site for RNA polymerase and other factors required for proper transcription including without limitation transcription factor binding sites, repressor and activator protein binding sites. The meaning of the term promoter also includes the term "promoter regulatory sequence". Promoter regulatory sequences may include upstream and downstream elements that may influence transcription, RNA processing or stability of the associated coding nucleic acid sequence. Promoters include naturally-derived and synthetic sequences. The coding nucleic acid sequences is usually located downstream of the promoter with respect to the direction of the transcription starting at the transcription initiation site.

The term "constitutive promoter" refers to an unregulated promoter that allows for continual transcription of the nucleic acid sequence it is operably linked to.

As used herein, the term "operably linked" refers to a linkage of polynucleotide elements in a functional relationship. A nucleic acid is "operably linked" when it is placed into a functional relationship with another nucleic acid sequence. For instance, a promoter, or rather a transcription regulatory sequence, is operably linked to a coding sequence if it affects the transcription of the

coding sequence. Operably linked means that the DNA sequences being linked are typically contiguous. The nucleotide sequence associated with the promoter sequence may be of homologous or heterologous origin with respect to the plant to be transformed. The sequence also may be entirely or partially synthetic. Regardless of the origin, the nucleic acid sequence associated with the promoter sequence will be expressed or silenced in accordance with promoter properties to which it is linked after binding to the polypeptide of an embodiment herein. The associated nucleic acid may code for a protein that is desired to be expressed or suppressed throughout the organism at all times or, alternatively, at a specific time or in specific tissues, cells, or cell compartment. Such nucleotide sequences particularly encode proteins conferring desirable phenotypic traits to the host cells or organism altered or transformed therewith. More particularly, the associated nucleotide sequence leads to the production of drimenol in the organism. Particularly, the nucleotide sequence encodes drimenol synthase.

"Target peptide" refers to an amino acid sequence which targets a protein, or polypeptide to intracellular organelles, i.e., mitochondria, or plastids, or to the extracellular space (secretion signal peptide). A nucleic acid sequence encoding a target peptide may be fused to the nucleic acid sequence encoding the amino terminal end, e.g., N-terminal end, of the protein or polypeptide, or may be used to replace a native targeting polypeptide.

The term "primer" refers to a short nucleic acid sequence that is hybridized to a template nucleic acid sequence and is used for polymerization of a nucleic acid sequence complementary to the template.

As used herein, the term "host cell" or "transformed cell" refers to a cell (or organism) altered to harbor at least one nucleic acid molecule, for instance, a recombinant gene encoding a desired protein or nucleic acid sequence which upon transcription yields a drimenol synthase protein useful to produce drimenol. The host cell is particularly a bacterial cell, a fungal cell or a plant cell. The host cell may contain a recombinant gene according to the present invention which has been integrated into the nuclear or organelle genomes of the host cell. Alternatively, the host may contain the recombinant gene extra-chromosomally. Homologous sequences include orthologous or paralogous sequences. Methods of identifying orthologs or paralogs including phylogenetic methods, sequence similarity and hybridization methods are known in the art and are described herein.

Paralogs result from gene duplication that gives rise to two or more genes with similar sequences and similar functions. Paralogs typically cluster together and are formed by duplications of genes within related plant species. Paralogs are found in groups of similar genes using pair-wise Blast analysis or during phylogenetic analysis of gene families using programs such as CLUSTAL.

In paralogs, consensus sequences can be identified characteristic to sequences within related genes and having similar functions of the genes.

Orthologs, or orthologous sequences, are sequences similar to each other because they are found in species that descended from a common ancestor. For instance, plant species that have common ancestors are known to contain many enzymes that have similar sequences and functions. The skilled artisan can identify orthologous sequences and predict the functions of the orthologs, for example, by constructing a phylogenetic tree for a gene family of one species using CLUSTAL or BLAST programs. A method for identifying or confirming similar functions among homologous sequences is by comparing of the transcript profiles in plants overexpressing or lacking (in knockouts/knockdowns) related polypeptides. The skilled person will understand that genes having similar transcript profiles, with greater than 50% regulated transcripts in common, or with greater than 70% regulated transcripts in common, or greater than 90% regulated transcripts in common will have similar functions. Homologs, paralogs, orthologs and any other variants of the sequences herein are expected to function in a similar manner by making plants producing drimenol synthase proteins.

An embodiment provided herein provides amino acid sequences of drimenol synthase proteins including orthologs and paralogs as well as methods for identifying and isolating orthologs and paralogs of the drimenol synthases in other organisms. Particularly, so identified orthologs and paralogs of the drimenol synthase retain drimenol synthase activity and are capable of producing drimenol starting from FPP precursors.

The term "selectable marker" refers to any gene which upon expression may be used to select a cell or cells that include the selectable marker. Examples of selectable markers are described below. The skilled artisan will know that different antibiotic, fungicide, auxotrophic or herbicide selectable markers are applicable to different target species.

"Drimenol" for purposes of this application refers to (-)-drimenol (CAS: 468-68-8).

The term "organism" refers to any non-human multicellular or unicellular organisms such as a plant, or a microorganism. Particularly, a micro-organism is a bacterium, a yeast, an algae or a fungus.

The term "plant" is used interchangeably to include plant cells including plant protoplasts, plant tissues, plant cell tissue cultures giving rise to regenerated plants, or parts of plants, or plant organs such as roots, stems, leaves, flowers, pollen, ovules, embryos, fruits and the like. Any plant can be used to carry out the methods of an embodiment herein.

The polypeptide to be contacted with an acyclic pyrophosphate, e.g. FPP, *in vitro* can be obtained by extraction from any organism expressing it, using standard protein or enzyme extraction technologies. If the host organism is an unicellular organism or cell releasing the polypeptide of an

embodiment herein into the culture medium, the polypeptide may simply be collected from the culture medium, for example by centrifugation, optionally followed by washing steps and re-suspension in suitable buffer solutions. If the organism or cell accumulates the polypeptide within its cells, the polypeptide may be obtained by disruption or lysis of the cells and further extraction of the polypeptide from the cell lysate.

The polypeptide having a drimenol synthase activity, either in an isolated form or together with other proteins, for example in a crude protein extract obtained from cultured cells or microorganisms, may then be suspended in a buffer solution at optimal pH. If adequate, salts, DTT, inorganic cations and other kinds of enzymatic co-factors, may be added in order to optimize enzyme activity. The precursor FPP is added to the polypeptide suspension, which is then incubated at optimal temperature, for example between 15 and 40°C, particularly between 25 and 35°C, more particularly at 30°C. After incubation, the drimenol produced may be isolated from the incubated solution by standard isolation procedures, such as solvent extraction and distillation, optionally after removal of polypeptides from the solution.

According to another particularly embodiment, the method of any of the above-described embodiments is carried out *in vivo*. In this case, step a) comprises cultivating a non-human host organism or cell capable of producing FPP and transformed to express at least one polypeptide comprising an amino acid comprising SEQ ID NO: 2 and having a drimenol synthase activity, under conditions conducive to the production of drimenol.

According to a more particular embodiment, the method further comprises, prior to step a), transforming a non-human organism or cell capable of producing FPP with at least one nucleic acid encoding a polypeptide comprising an amino acid comprising SEQ ID NO: 2 and having a drimenol synthase activity, so that said organism expresses said polypeptide.

These embodiments of an embodiment herein are particularly advantageous since it is possible to carry out the method *in vivo* without previously isolating the polypeptide. The reaction occurs directly within the organism or cell transformed to express said polypeptide.

According to a more particular embodiment at least one nucleic acid used in any of the above embodiments comprises a nucleotide sequence that has been obtained by modifying SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof. According to another embodiment, the at least one nucleic acid is isolated from a plant of the Paeoniaceae family, particularly from *Paeonia anomala*. The organism or cell is meant to “express” a polypeptide, provided that the organism or cell is transformed to harbor a nucleic acid encoding said polypeptide, this nucleic acid is transcribed to mRNA and the polypeptide is found in the host organism or cell. The term “express” encompasses “heterologously express” and “over-express”, the latter referring to levels of mRNA, polypeptide

and/or enzyme activity over and above what is measured in a non-transformed organism or cell. A more detailed description of suitable methods to transform a non-human host organism or cell will be described later on in the part of the specification that is dedicated to such transformed non-human host organisms or cells as specific objects provided herein and in the examples.

5 A particular organism or cell is meant to be “capable of producing FPP” when it produces FPP naturally or when it does not produce FPP naturally but is transformed to produce FPP, either prior to the transformation with a nucleic acid as described herein or together with said nucleic acid. Organisms or cells transformed to produce a higher amount of FPP than the naturally occurring organism or cell are also encompassed by the “organisms or cells capable of producing FPP”.

10 Methods to transform organisms, for example microorganisms, so that they produce FPP are already known in the art.

To carry out an embodiment herein *in vivo*, the host organism or cell is cultivated under conditions conducive to the production of drimenol. Accordingly, if the host is a transgenic plant, optimal growth conditions are provided, such as optimal light, water and nutrient conditions, for
15 example. If the host is a unicellular organism, conditions conducive to the production of drimenol may comprise addition of suitable cofactors to the culture medium of the host. In addition, a culture medium may be selected, so as to maximize drimenol synthesis. Optimal culture conditions are described in a more detailed manner in the following Examples.

Non-human host organisms suitable to carry out the method of an embodiment herein *in vivo*
20 may be any non-human multicellular or unicellular organisms. In a particular embodiment, the non-human host organism used to carry out an embodiment herein *in vivo* is a plant, a prokaryote or a fungus. Any plant, prokaryote or fungus can be used. Particularly useful plants are those that naturally produce high amounts of terpenes. In a more particular embodiment the non-human host organism used to carry out the method of an embodiment herein *in vivo* is a microorganism. Any
25 microorganism can be used but according to an even more particular embodiment said microorganism is a bacteria or yeast. Most particularly, said bacteria is *Escherichia coli* and said yeast is *Saccharomyces cerevisiae*.

Some of these organisms do not produce FPP naturally. To be suitable to carry out the method of an embodiment herein, these organisms have to be transformed to produce said precursor.
30 They can be so transformed either before the modification with the nucleic acid described according to any of the above embodiments or simultaneously, as explained above.

Isolated higher eukaryotic cells can also be used, instead of complete organisms, as hosts to carry out the method of an embodiment herein *in vivo*. Suitable eukaryotic cells may be any non-human cell, but are particularly plant or fungal cells.

In another particular embodiment, the polypeptide comprises SEQ ID NO: 2.

According to another particular embodiment, the at least one polypeptide having a drimenol synthase activity used in any of the above-described embodiments or encoded by the nucleic acid used in any of the above-described embodiments comprises an amino acid sequence that is a variant of SEQ ID NO: 2, obtained by genetic engineering, provided that said variant keeps its Drimenol synthase activity, as defined above and has the required percentage of identity to SEQ ID NO: 2. In other terms, said polypeptide particularly comprises an amino acid sequence encoded by a nucleotide sequence that has been obtained by modifying SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof. According to a more particular embodiment, the at least one polypeptide having a drimenol synthase activity used in any of the above-described embodiments or encoded by the nucleic acid used in any of the above-described embodiments consists of an amino acid sequence that is a variant of SEQ ID NO: 2, obtained by genetic engineering, i.e. an amino acid sequence encoded by a nucleotide sequence that has been obtained by modifying SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof.

According to another particular embodiment, the at least one polypeptide having a Drimenol synthase activity used in any of the above-described embodiments or encoded by the nucleic acid used in any of the above-described embodiments is a variant of SEQ ID NO: 2 that can be found naturally in other organisms, such as other plant species, provided that it keeps its Drimenol synthase activity. As used herein, the polypeptide is intended as a polypeptide or peptide fragment that encompasses the amino acid sequences identified herein, as well as truncated or variant polypeptides, provided that they keep their drimenol synthase activity as defined above and that they share at least the defined percentage of identity with the corresponding fragment of SEQ ID NO: 2.

Examples of variant polypeptides are naturally occurring 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 of an embodiment herein. Polypeptides encoded by a nucleic acid obtained by natural or artificial mutation of a nucleic acid of an embodiment herein, as described thereafter, are also encompassed by an embodiment herein.

Polypeptide variants resulting from a fusion of additional peptide sequences at the amino and carboxyl terminal ends can also be used in the methods of an embodiment herein. In particular such a fusion can enhance expression of the polypeptides, be useful in the purification of the protein or improve the enzymatic activity of the polypeptide in a desired environment or expression system. Such additional peptide sequences may be signal peptides, for example. Accordingly, the present

invention encompasses methods using variant polypeptides, such as those obtained by fusion with other oligo- or polypeptides and/or those which are linked to signal peptides. Polypeptides resulting from a fusion with another functional protein, such as another protein from the terpene biosynthesis pathway, can also be advantageously be used in the methods of an embodiment herein.

5 According to another embodiment, the at least one polypeptide having a drimenol synthase activity used in any of the above-described embodiments or encoded by the nucleic acid used in any of the above-described embodiments is isolated from a plant of the Paeoniaceae family, particularly from *Paeonia anomala*. An important tool to carry out the method of an embodiment herein is the polypeptide itself. A polypeptide having a drimenol synthase activity and comprising an amino acid
10 sequence of SEQ ID NO: 2 is therefore provided herein.

 According to a particular embodiment, the polypeptide is capable of producing a mixture of sesquiterpenes wherein drimenol represents at least 20%, particularly at least 30%, particularly at least 35%, particularly at least 90%, particularly at least 95%, more particularly at least 98% of the sesquiterpenes produced. In another aspect provided here, the drimenol is produced with greater than
15 or equal to 95%, more particularly 98% selectivity.

 According to a particular embodiment, the polypeptide comprises an amino acid sequence of SEQ ID NO: 2.

 According to another particular embodiment, the polypeptide consists of SEQ ID NO: 2.

20 The at least one polypeptide comprises an amino acid sequence that is a variant of SEQ ID NO: 2, either obtained by genetic engineering or found naturally in *Paeonia* plants or in other plant species. In other terms, when the variant polypeptide is obtained by genetic engineering, said polypeptide comprises an amino acid sequence encoded by a nucleotide sequence that has been obtained by modifying SEQ ID NO: 5 or the complement thereof. According to a more particular
25 embodiment, the at least one polypeptide having a drimenol synthase activity consists of an amino acid sequence that is a variant of SEQ ID NO: 2 obtained by genetic engineering, i.e. an amino acid sequence encoded by a nucleotide sequence that has been obtained by modifying SEQ ID NO: 5.

 According to another embodiment, the polypeptide is isolated from a plant of the Paeoniaceae family, particularly from *Paeonia anomala*. As used herein, the polypeptide is intended as a
30 polypeptide or peptide fragment that encompasses the amino acid sequence identified herein, as well as truncated or variant polypeptides, provided that they keep their activity as defined above and that they share at least the defined percentage of identity with the corresponding fragment of SEQ ID NO: 2.

 As mentioned above, the nucleic acid encoding the polypeptide of an embodiment herein is a

useful tool to modify non-human host organisms or cells intended to be used when the method is carried out in vivo.

A nucleic acid encoding a polypeptide according to any of the above-described embodiments is therefore also provided herein.

5 According to a more particular embodiment, the nucleic acid comprises SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof.

According to another particular embodiment, the nucleic acid consists of a nucleotide sequence SEQ ID NO: 5 or the complement thereof.

10 The nucleic acid of an embodiment herein can be defined as including deoxyribonucleotide or ribonucleotide polymers in either single- or double-stranded form (DNA and/or RNA). The terms "nucleotide sequence" should also be understood as comprising a polynucleotide molecule or an oligonucleotide molecule in the form of a separate fragment or as a component of a larger nucleic acid. Nucleic acids of an embodiment herein also encompass certain isolated nucleotide sequences including those that are substantially free from contaminating endogenous material. The nucleic acid
15 of an embodiment herein may be truncated, provided that it encodes a polypeptide encompassed by the present invention, as described above.

In one embodiment, the nucleic acid of an embodiment herein can be either present naturally in plants of the *Paeonia* species or other species, or be obtained by modifying SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof.

20 The nucleic acids comprising a sequence obtained by mutation of SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof are encompassed by an embodiment herein, provided that the sequences they comprise share at least the defined sequence of SEQ ID NO: 1 or SEQ ID NO: 5 or the complement thereof and provided that they encode a polypeptide having a drimenol synthase activity, as defined in any of the above embodiments. Mutations may be any kind of mutations of
25 these nucleic acids, such as point mutations, deletion mutations, insertion mutations and/or frame shift mutations. A variant nucleic acid may be prepared in order to adapt its nucleotide sequence to a specific expression system. For example, bacterial expression systems are known to more efficiently express polypeptides if amino acids are encoded by particular codons.

30 Due to the degeneracy of the genetic code, more than one codon may encode the same amino acid sequence, multiple nucleic acid sequences can code for the same protein or polypeptide, all these DNA sequences being encompassed by an embodiment herein. Where appropriate, the nucleic acid sequences encoding the drimenol synthase may be optimized for increased expression in the host cell. For example, nucleotides of an embodiment herein may be synthesized using codons particular by a host for improved expression.

Another important tool for transforming host organisms or cells suitable to carry out the method of an embodiment herein in vivo is an expression vector comprising a nucleic acid according to any embodiment of an embodiment herein. Such a vector is therefore also provided herein.

5 The expression vectors provided herein may be used in the methods for preparing a genetically transformed host organism and/or cell, in host organisms and/or cells harboring the nucleic acids of an embodiment herein and in the methods for making polypeptides having a drimenol synthase activity, as disclosed further below.

10 Recombinant non-human host organisms and cells transformed to harbor at least one nucleic acid of an embodiment herein so that it heterologously expresses or over-expresses at least one polypeptide of an embodiment herein are also very useful tools to carry out the method of an embodiment herein. Such non-human host organisms and cells are therefore provided herein.

A nucleic acid according to any of the above-described embodiments can be used to transform the non-human host organisms and cells and the expressed polypeptide can be any of the above-described polypeptides.

15 Non-human host organisms of an embodiment herein may be any non-human multicellular or unicellular organisms. In a particular embodiment, the non-human host organism is a plant, a prokaryote or a fungus. Any plant, prokaryote or fungus is suitable to be transformed according to the methods described herein. Particularly useful plants are those that naturally produce high amounts of terpenes.

20 In a more particular embodiment the non-human host organism is a microorganism. Any microorganism is suitable as a non-human host, but according to an even more particular embodiment said microorganism is a bacterium or yeast. Most particularly, said bacterium is *E. coli* and said yeast is *Saccharomyces cerevisiae*.

25 Isolated higher eukaryotic cells can also be transformed, instead of complete organisms. As higher eukaryotic cells, we mean here any non-human eukaryotic cell except yeast cells. Particular higher eukaryotic cells are plant cells or fungal cells.

A variant may also differ from the polypeptide of an embodiment herein by attachment of modifying groups which are covalently or non-covalently linked to the polypeptide backbone. The variant also includes a polypeptide which differs from the polypeptide provided herein by introduced
30 N-linked or O-linked glycosylation sites, and/or an addition of cysteine residues. The skilled artisan will recognize how to modify an amino acid sequence and preserve biological activity.

The functionality or activity of any drimenol synthase protein, variant or fragment, may be determined using various methods. For example, transient or stable overexpression in plant, bacterial or yeast cells can be used to test whether the protein has activity, i.e., produces drimenol from FPP

precursors. Drimenol synthase activity may be assessed in a microbial expression system, such as the assay described in Example 2 herein on the production of drimenol, indicating functionality. A variant or derivative of a drimenol synthase polypeptide of an embodiment herein retains an ability to produce drimenol from FPP precursors. Amino acid sequence variants of the drimenol synthases provided herein may have additional desirable biological functions including, e.g., altered substrate utilization, reaction kinetics, product distribution or other alterations.

An embodiment herein provides polypeptides of an embodiment herein to be used in a method to produce drimenol contacting an FPP precursor with the polypeptides of an embodiment herein either in vitro or in vivo.

Provided herein is also an isolated, recombinant or synthetic polynucleotide encoding a polypeptide or variant polypeptide provided herein. An embodiment of an embodiment herein provides an isolated, recombinant or synthetic nucleic acid sequence of SEQ ID NO: 1 or SEQ ID NO: 5 encoding for a drimenol synthase having the amino acid sequence of SEQ ID NO: 2. or fragments thereof that catalyze production of drimenol in a cell from a FPP precursor. Provided herein are also cDNA, genomic DNA and RNA sequences. Any nucleic acid sequence encoding the drimenol synthase or variants thereof is referred herein as a drimenol synthase encoding sequence.

According to a particular embodiment, the nucleic acid of SEQ ID NO: 1 or SEQ ID NO: 5 is the coding sequence of a drimenol synthase gene encoding the drimenol synthase obtained as described in the Examples.

A fragment of a polynucleotide of SEQ ID NO: 1 or SEQ ID NO: 5 refers to contiguous nucleotides that is particularly at least 15 bp, at least 30 bp, at least 40 bp, at least 50 bp and/or at least 60 bp in length of the polynucleotide of an embodiment herein. Particularly the fragment of a polynucleotide comprises at least 25, more particularly at least 50, more particularly at least 75, more particularly at least 100, more particularly at least 150, more particularly at least 200, more particularly at least 300, more particularly at least 400, more particularly at least 500, more particularly at least 600, more particularly at least 700, more particularly at least 800, more particularly at least 900, more particularly at least 1000 contiguous nucleotides of the polynucleotide of an embodiment herein. Without being limited, the fragment of the polynucleotides herein may be used as a PCR primer, and/or as a probe, or for anti-sense gene silencing or RNAi.

It is clear to the person skilled in the art that genes, including the polynucleotides of an embodiment herein, can be cloned on basis of the available nucleotide sequence information, such as found in the attached sequence listing, by methods known in the art. These include e.g. the design of DNA primers representing the flanking sequences of such gene of which one is generated in sense orientations and which initiates synthesis of the sense strand and the other is created in reverse

complementary fashion and generates the antisense strand. Thermo stable DNA polymerases such as those used in polymerase chain reaction are commonly used to carry out such experiments.

Alternatively, DNA sequences representing genes can be chemically synthesized and subsequently introduced in DNA vector molecules that can be multiplied by e.g. compatible bacteria such as e.g. E. coli.

In a related embodiment provided herein, PCR primers and/or probes for detecting nucleic acid sequences encoding a drimenol synthase are provided. The skilled artisan will be aware of methods to synthesize degenerate or specific PCR primer pairs to amplify a nucleic acid sequence encoding the drimenol synthase or fragments thereof, based on SEQ ID NO: 1 or SEQ ID NO: 5. A detection kit for nucleic acid sequences encoding the drimenol synthase may include primers and/or probes specific for nucleic acid sequences encoding the drimenol synthase, and an associated protocol to use the primers and/or probes to detect nucleic acid sequences encoding the drimenol synthase in a sample. Such detection kits may be used to determine whether a plant has been modified, i.e., transformed with a sequence encoding the drimenol synthase.

The nucleic acid sequences obtained by mutations of SEQ ID NO: 1 or SEQ ID NO: 5 can be routinely made and are also within embodiments provided herein. It is clear to the skilled artisan that mutations, deletions, insertions, and/or substitutions of one or more nucleotides can be introduced into the DNA sequence of SEQ ID NO: 5. Generally, a mutation is a change in the DNA sequence of a gene that can alter the amino acid sequence of the polypeptide produced.

To test a function of variant DNA sequences according to an embodiment herein, the sequence of interest is operably linked to a selectable or screenable marker gene and expression of the reporter gene is tested in transient expression assays with protoplasts or in stably transformed plants. The skilled artisan will recognize that DNA sequences capable of driving expression are built as modules. Accordingly, expression levels from shorter DNA fragments may be different than the one from the longest fragment and may be different from each other. Provided herein are also functional equivalents of the nucleic acid sequence coding the drimenol synthase proteins provided herein, i.e., nucleotide sequences that hybridize under stringent conditions to the nucleic acid sequence of SEQ ID NO: 1 or SEQ ID NO: 5 .

The skilled artisan will be aware of methods to identify homologous sequences in other organisms and methods (identified in the Definition section herein) to determine the percentage of sequence identity between homologous sequences. Such newly identified DNA molecules then can be sequenced and the sequence can be compared with the nucleic acid sequence of SEQ ID NO: 1 or SEQ ID NO: 5.

A related embodiment provided herein provides a nucleic acid sequence which is complementary to the nucleic acid sequence according to SEQ ID NO: 1 or SEQ ID NO: 5 such as inhibitory RNAs, or nucleic acid sequence which hybridizes under stringent conditions to at least part of the nucleotide sequence according to SEQ ID NO: 1 or SEQ ID NO: 5. An alternative
5 embodiment of an embodiment herein provides a method to alter gene expression in a host cell. For instance, the polynucleotide of an embodiment herein may be enhanced or overexpressed or induced in certain contexts (e.g. following insect bites or stings or upon exposure to a certain temperature) in a host cell or host organism.

Alteration of expression of a polynucleotide provided herein also results in "ectopic
10 expression" which is a different expression pattern in an altered and in a control or wild-type organism. Alteration of expression occurs from interactions of polypeptide of an embodiment herein with exogenous or endogenous modulators, or as a result of chemical modification of the polypeptide. The term also refers to an altered expression pattern of the polynucleotide of an
15 embodiment herein which is altered below the detection level or completely suppressed activity.

In one embodiment, several drimenol synthases encoding nucleic acid sequences are co-
20 expressed in a single host, particularly under control of different promoters. Alternatively, several drimenol synthase proteins encoding nucleic acid sequences can be present on a single transformation vector or be co-transformed at the same time using separate vectors and selecting transformants comprising both chimeric genes. Similarly, one or more drimenol synthase encoding
25 genes may be expressed in a single plant together with other chimeric genes, for example encoding other proteins which enhance insect pest resistance, or others.

The nucleic acid sequences of an embodiment herein encoding drimenol synthase proteins can be inserted in expression vectors and/or be contained in chimeric genes inserted in expression
30 vectors, to produce drimenol synthase proteins in a host cell or host organism. The vectors for inserting transgenes into the genome of host cells are well known in the art and include plasmids, viruses, cosmids and artificial chromosomes. Binary or co-integration vectors into which a chimeric gene is inserted are also used for transforming host cells.

An embodiment provided herein provides recombinant expression vectors comprising a
35 nucleic acid sequence of a drimenol synthase gene, or a chimeric gene comprising a nucleic acid sequence of a drimenol synthase gene, operably linked to associated nucleic acid sequences such as, for instance, promoter sequences. For example, a chimeric gene comprising a nucleic acid sequence of SEQ ID NO: 1 or SEQ ID NO: 5 or may be operably linked to a promoter sequence suitable for expression in plant cells, bacterial cells or fungal cells, optionally linked to a 3' non-translated
nucleic acid sequence.

Alternatively, the promoter sequence may already be present in a vector so that the nucleic acid sequence which is to be transcribed is inserted into the vector downstream of the promoter sequence. Vectors are typically engineered to have an origin of replication, a multiple cloning site, and a selectable marker.

5 The following examples are illustrative only and are not intended to limit the scope of the claims and embodiments described herein.

Examples

10 Example 1

***Paeonia anomala* plant material sourcing and root transcriptome sequencing.**

Paeonia anomala plant material was obtained from Datong in Qinghai, China. To establish if *Paeonia anomala* contained drimenol, the roots were collected, dried in the shade and extracted with dichloromethane. The extract was analyzed by GC-MS, the parameters of GC-MS analysis were
15 described as below: An Agilent 6890 series GC system equipped with a DB1-ms column 30m × 0.25mm × 0.25 μm film thickness, P/N 122-0132 (J&W scientific Inc, Folsom, CA) and coupled with a 5975 series mass spectrometer was used. The carrier gas was helium at a constant flow of 0.7 mL/min. Injection was in split (1:5) mode with the injector temperature set at 250°C. The oven temperature was programmed from 50°C (5 min hold) to 300°C at 5°C/min, then to 340°C at
20 50°C/min and held for 3 min. Identification of products was based on comparison to data recorded from an authentic standard. The roots of *Paeonia anomala* contained a small amount of drimenol (Figure 5).

Fresh roots of *Paeonia anomala* were used for transcriptome analysis. Total RNA was extracted using the Column Plant RNA out (TIANDZ, China). This total RNA was processed using
25 the Illumina Total RNA-Seq technique and sequenced on Illumina MiSeq sequencer. A total of 9 million of paired-end reads of 2×251bp were generated. The reads were assembled using the Trinity (<http://trinityrnaseq.sf.net/>) software. 26457 unigenes with an average size of 1109 bp were obtained. The unigenes were annotated by NCBI Blast (<http://www.ncbi.nlm.nih.gov/>) as well as InterProScan software (<http://www.ebi.ac.uk/Tools/pfa/iprscan/>). This approach provided the sequences for 7 new
30 putative sesquiterpene synthases including PaTPS1. The enzymatic activity of PaTPS1 was evaluated as described in the following examples.

Example 2

Functional expression and characterization of PaTPS1 from *P. anomala*.

The total RNA extracted by Column Plant RNA out kit was first reverse transcribed into cDNA using the SuperScriptIII First-Strand Synthesis kit (Invitrogen, Shanghai, China). And then the product was used as the template, forward primer (5'-GGGGTACCATGTCTCTTCCCGTCTCAGTAG-3') and reverse primer (5'-GCTCTAGATCATATTGGGATGGGATCAATT-3') were used to amplify the gene from the cDNA library of *P. anomala*, and restriction site of *KpnI* was added to the 5' end of PaTPS1 while *XbaI* was added to the 3' end. PaTPS1 was then sub-cloned into a modified binary vector pCAMBIA2300-35S-OCS-35S-GgFPSopt to form pCAMBIA2300-35S-OCS-35S-GgFPSopt-PaTPS1 (pGgFPS-PaTPS1 in short).

pCAMBIA2300-35S-OCS-35S-GgFPSopt is an engineered vector based on the commercial plasmid pCAMBIA2300 (Cambia). Cauliflower Mosaic Virus promoter (CaMV35S) was inserted between the *EcoRI* and *KpnI* restriction sites, the octopine synthase (OCS) terminator was inserted between the *PstI* and *HindIII*, and the kanamycin resistance gene inside the right and the left borders of transfer DNA (T-Border) of pCAMBIA2300 plasmid was replaced by the codon optimized *Gallus gallus* farnesyldiphosphate synthase (GgFPS) using *XhoI* restriction enzyme followed by direction confirmation. The binary vector pCAMBIA2300-35S-OCS-35S-GgFPSopt was thus formed to produce the precursor of sesquiterpene synthase, Farnesylpyrophosphate (FPP).

1 micro gram of pGgFPS-PaTPS1 in short plasmid was transformed into agrobacteria EHA105. Transformed cells were selected on kanamycin (50 µg/mL) and rifampicin (25 µg/mL) LB-agarose plates at 28°C for 2 days. Single colony was inoculated into 25 mL LB with the same antibiotics and incubated at 28°C in a shaker with 200 rpm speed overnight. When the OD600 reached 1.0, the culture was centrifuged at 5000 rpm for 10 min at room temperature. The precipitates were re-suspended with 20 mL MgSO₄ (10 mM). The suspension was then centrifuged again and the precipitates were re-suspended with 5 mL Acetosyringone (AS) solution (765 µM), then the OD600 was adjusted to 1.0 with AS solution.

The agrobacteria strain transformed with the tomato bushy stunt virus p19 which could suppress gene silencing was mixed with the strain transformed with pGgFPS-PaTPS1 with volume ratio 1:1. The mixture was infiltrated into intercellular space from the abaxial side of 4-week old tobacco leaves grown at 28°C with 12 hour light/ 12 hour dark photoperiod. 3 days later, the infected leaves were fed with mevalonate (25 mM). After 8 hours the leaf samples were harvested and about 100-300 mg of the leaf was weighed into 10 mL Eppendorf tube with 2 beads and frozen in liquid nitrogen immediately. The samples were ground to powder by shaking and extracted by 2 mL ethyl

acetate with 20 $\mu\text{g/mL}$ dodecane as internal standards overnight. The sample was filtered and the filtrate was dried over anhydrous Na_2SO_4 and then analyzed by GC/MS. The parameters of GC/MS were the same as described in Example 1.

In this tobacco transient transformation experiment, PaTPS1 produced drimenol as the only
5 detectable product based on comparison to data recorded from an authentic standard (Figure 6). The yield of drimenol in this experiment was calculated based on the internal standard as 3.9 μg drimenol per gram of fresh tobacco leaves.

Sequence Listings

SEQ ID NO: 1

Coding DNA sequence of *PaTPS1*:

ATGTCTCTTCCCGTCTCAGTAGCTAATCAGCCACCGGCGAAATGTAAACAAGAGGTTAT
 5 TCGCAATACAGCAAATTTCCCCCGTGGTATTTCCGCTCATCAGTTCATCACTTACACTCC
 TCAAGATGAGGAAACTCGTGCGCATATAGTACAAGAGATTGAAGAACTGAAAGAAATT
 GTAAGAACAGAGGTGATGACACTTGTTGGTATACCTTCACAACAACACTCAAGCTTGTTGA
 TGCAATCCAACGCCTTGGTGTGGGATACCACTTTCAGAAGGAGATAGATGAAGCCTTAC
 ACCAACTGTATGATACATATGGTCATGGCCAAGATGAAGATCTCTTCACAGTTGCTCTTT
 10 GGTTTCAGACTTCTAAGACAACAAGGGTATAATGTTTCATGTGACATATTCAACAAATTC
 ACCGACGACAAGGGAAACTTCAAGGAATGCTTGGTTCAAAATGTGGAGGGGCATGGTAG
 CCTTGTACGAAGCAACGCATCTCAGAGTGCATGGAGAAGATGTACTTGAAGCAGCACTC
 ACTTTTACAACCATTACCTTAAGGCCTTGGCAACTCATCTTGCAAGCCATCCCCTTAGA
 GCACTAGTGAATCGTGCCCTAGAACAGCCTACCTATAAGGGTGTACCTAGGCTGGAGGC
 15 AAGACATTATATATCTTTCTATCAACAAGAGCAATTGCATGATAAAGCTTTACTGAGAC
 TTGCCAAGTTAGATTTTAACCTACTACAATCATTGCACAAAAAGGAGCTATCGGAGGTC
 GCTAGATGGTGGACAAAAGTAAACTTTGAAAACAAGTTACCTTTCGTGAGAGACAGGGT
 GGTGGAGAGTTACTTTTGGGGATTGGGAGATTATTTTGAGCCTGAGCACTCCATTGCTA
 GAATGATATTAAGCAAAAATAATCGCCCTAGTAACGGTTATGGATGATATTTATGACGCA
 20 TATGGTACACTGGAAGAACTCGAGCTATTTACAGATGCAGTTCAAAGGTGGGATATCAA
 CTGCACCCAACAACACTCCCAGAATACATTAAAGTGTTTTTTTCAGGCAATGTTAGATGCAT
 ACGAAGAGATTGAAGAAGAATTATCTAACGAACCTGGACGAACATATCGTGTTTCATTAT
 GCAATAGAAGCCATGAAAATACAAGCCCAAGCATAACCTTGCTGAAACAAAATGGTCTA
 ATGAAAAGTATGTACCGACATTTGAGGAGTATATGGATAATGCACGATTAAGCGCAGGT
 25 TACTTCATGCTTACAGTCATATCTTTTCTTTTTATGGGGGAAGACGCGACAAAAGATTCA
 TTTGATTGGCTGTTCAACGACCCTAAGATTCTTAGAGCCTCATCAATCATTACCAGGCTC
 ATGGATGACATAGTTTCTCATAAGTTTGAGCAAGAGAGAGGACATGTTGCATCATCCGT
 TGAGTGTTACATGAAGCAACACAATGTTTCGGAGCAACAAACATATCAAGAGTTTCAA
 TGAAAATTGTGGAGGGATGGAAAGATCTAAATCAGGCATTACTCATACCTACTGATGCA
 30 TCGATTCTCTCCTTACTCGTATCCTTAATTTTACACGCTTTATGGATGTTAACTACAAGG
 AACGAGATGAATTCACACATGTTGGAGACGTTTTTAAAAGATCGAATTGCATTGTTACTA
 ATTGATCCCATCCCAATATGA

SEQ ID NO: 2

Amino acid sequences of *PaTPS1*:

MSLPVSVANQPPAKCKQEVRNTANFPRGISAHQFITYTPQDEETRAHIVQEIEELKEIVRTEV
 MTLVGIPSQQLKLVDAIQRLGVGYHFQKEIDEALHQLYDTYGHGQDEDLFTVALWFRLLRQ
 5 QGYNVSCDIFNKFTDDKGNFKECLVQNVEGMVALYEATHLRVHGEDVLEAALTFTTIHLK
 ALATHLASHPLRALVNRALAQPTYKGVPRLEARHYISFYQQEQLHDKALLRLAKLDFNLLQ
 SLHKKELSEVARWWTKVNFENKLPFVRDRVVESYFWGLGDYFEPEHSIARMILSKIIALVTV
 MDDIYDAYGTLEELFTDAVQRWDINCTQQLPEYIKVFFQAMLDAYEEIEEELSNEPGRTY
 RVHYAIEAMKIQAQAYLAETKWSNEKYVPTFEEYMDNARLSAGYFMLTVISFLFMGEDAT
 10 KDSFDWLFNDPKILRASSITRLMDDIVSHKFEQERGHVASSVECYMKQHNVSEQQTYQEFQ
 MKIVEGWKDLNQALLIPTDASIPLLTRILNFTRFMDVNYKERDEFTHVGDVVKDRIALLLIDP
 IPI

SEQ ID NO: 3

15 The DNA sequence of *CaMV35S* promoter

GAATTCCCATGGAGTCAAAGATTCAAATAGAGGACCTAACAGAACTCGCCGTAAAGAC
 TGGCGAACAGTTCATACAGAGTCTCTTACGACTCAATGACAAGAAGAAAATCTTCGTCA
 ACATGGTGGAGCACGACACGCTTGTCTACTCCAAAAATATCAAAGATACAGTCTCAGAA
 GACCAAAGGGCAATTGAGACTTTTCAACAAAGGGTAATATCCGGAAACCTCCTCGGATT
 20 CCATTGCCAGCTATCTGTCACTTTATTGTGAAGATAGTGGAAAAGGAAGGTGGCTCCT
 ACAAATGCCATCATTGCGATAAAGGAAAGGCCATCGTTGAAGATGCCTCTGCCGACAGT
 GGTCCCAAAGATGGACCCCCACCCACGAGGAGCATCGTGGAAAAGAAGACGTTCCAA
 CCACGTCTTCAAAGCAAGTGGATTGATGTGATATCTCCACTGACGTAAGGGATGACGCA
 CAATCCCACTATCCTTCGCAAGACCCTCCTCTATATAAGGAAGTTCATTTTCAATTTGGAG
 25 AGGACAGGGTACC

SEQ ID NO: 4

The DNA sequence of *OCS* terminator

CTGCAGGCATGCCCTGCTTTAATGAGATATGCGAGACGCCTATGATCGCATGATATTTG
 30 CTTTCAATTCTGTTGTGCACGTTGTAAAAAACCTGAGCATGTGTAGCTCAGATCCTTACC
 GCCGGTTTCGGTTCATTCTAATGAATATATCACCCGTTACTATCGTATTTTTATGAATAA
 TATTCTCCGTTCAATTTACTGATTGTCCAAGCTT

SEQ ID NO: 5

The optimized DNA sequence of *GgFPS*

ATGCAACCACACCATCATCACAAAGAAGGAAGAATGCACAAGTTTACTGGAGTTAACG
CAAAGTTTCAACAGCCTGCTCTCAGAAATCTTTCTCCTGTTGTGGTTGAAAGAGAGAGG
5 GAAGAGTTTGTGGGATTTTTCCCTCAAATTGTTAGAGATTTGACTGAAGATGGAATCGG
TCATCCAGAAGTGGGAGATGCAGTTGCTAGGCTCAAGGAAGTTTTACAGTATAATGCTC
CTGGAGGTAAATGTAACAGAGGACTCACAGTGGTTGCTGCATACAGGGAATTATCTGGA
CCAGGTCAAAGGATGCTGAGTCACTTAGATGTGCATTGGCTGTGGGATGGTGCATCGA
GCTTTTCCAGGCATTTTTCTTGGTTGCTGATGATATTATGGATCAATCACTCACCAGAAG
10 GGGACAGTTATGTTGGTATAAGAAAGAAGGAGTTGGTCTTGATGCTATTAATGATAGTT
TTCTTTTGGAGTCTTCAGTGTACAGAGTTTTGAAGAAATATTGTAGACAAAGGCCTTACT
ACGTTTCATCTCTTAGAACTTTTTCTTGCAAACCTGCATACCAGACAGAGTTGGGACAGATG
CTCGATCTTATTACAGCTCCAGTGTCAAAGGTTGATCTCTCACACTTTAGTGAAGAGAGA
TATAAGGCAATCGTTAAGTACAAAACCGCTTTCTACAGTTTCTACCTTCCTGTGGCTGCT
15 GCTATGTATATGGTTGGTATAGATTCTAAGGAAGAGCATGAAAACGCAAAGCTATACT
TTTGGAAATGGGAGAGTACTTCAAATACAGGATGATTACCTTGATTGCTTCGGAGATC
CTGCTTTGACCGGAAAGGTTGGTACTGATATCCAAGATAATAAGTGTTCTTGGCTCGTG
GTTCAATGCTTACAGAGAGTTACACCTGAACAAAGGCAGCTCTTAGAGGATAACTATGG
AAGAAAGGAACCAGAGAAAGTGGCAAAGGTTAAAGAATTGTACGAGGCTGTTGGTATG
20 AGAGCAGCTTTCCAACAGTATGAAGAGAGTTCTTACAGAAGGCTTCAAGAATTGATCGA
GAAACACTCAAACAGACTCCCTAAAGAAATCTTCCTCGGACTCGCTCAGAAAATCTACA
AAAGGCAAAAATGATAA

Claims

What is claimed is:

- 5 1. A method of producing drimenol comprising:
 - i) contacting an acyclic farnesyl diphosphate (FPP) precursor with a polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2 or comprising SEQ ID NO:2 to produce the drimenol; and
 - 10 ii) optionally isolating the drimenol.
2. The method as recited in Claim 1 wherein the drimenol is isolated
3. The method as recited in any one of claims 1-3 comprising contacting the drimenol with at least one enzyme to produce a drimenol derivative.
4. The method as recited in any one of claims claim 1-3 comprising converting the drimenol to a
15 drimenol derivative using a chemical synthesis
5. An isolated polypeptide having drimenol activity comprising SEQ ID NO: 2.
6. An isolated polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2.
- 20 7. An isolated nucleic acid molecule encoding a polypeptide recited in claims 5 and 6 selected from the group consisting of SEQ ID NO: 1 and SEQ ID NO: 5.
8. An isolated nucleic acid molecule encoding a polypeptide recited in claims 5 and 6 having a nucleotide sequence 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to a nucleotide sequence selected from the group consisting of
25 SEQ ID NO: 1 and SEQ ID NO: 5.
9. The method as recited in claim 1 comprising the steps of transforming a host cell or non-human organism with a nucleic acid encoding a polypeptide having drimenol synthase activity and having at least 50%, 55%, 60%, 65%, 70%, 75%, 80%, 85%, 90%, 95%, 96%, 97%, 98% or 99% sequence identity to SEQ ID NO: 2 or comprising SEQ ID NO:2 and
30 culturing the host cell or organism under conditions that allow for the production of the polypeptide.
10. A vector comprising the nucleic acid molecule of claims 7 and 8.

11. The vector of claim 10 wherein the vector is a prokaryotic vector, viral vector or a eukaryotic vector.
12. The vector of claim 10 or 11 that is an expression vector.
13. The method recited in claim 9 wherein the cell is a prokaryotic cell.
- 5 14. The method as recited in claim 9 wherein the cell is a bacterial cell.
15. The method as recited in claim 9 wherein the cell is an eukaryotic cell.
16. The method as recited in claim 9 wherein the eukaryotic cell is a yeast cell or a plant cell.

FIGURES

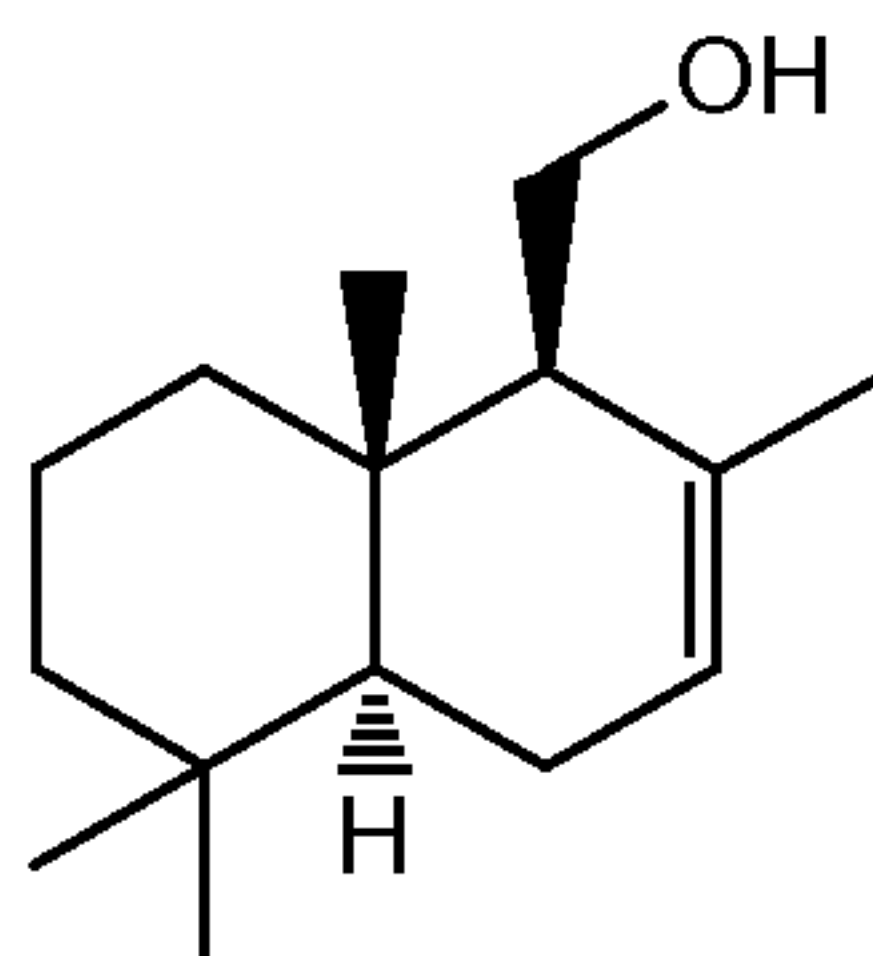


Figure 1

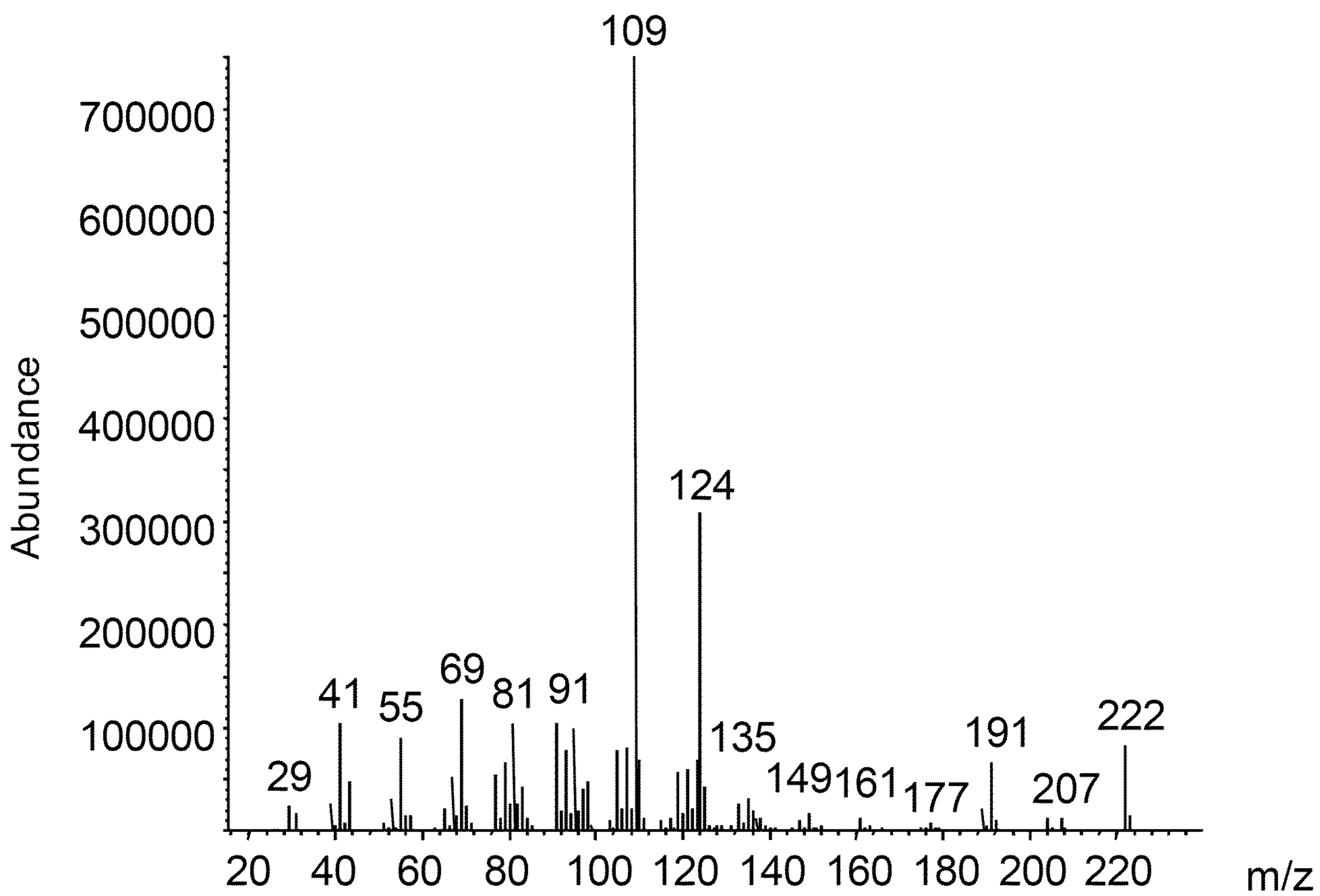


Figure 2.

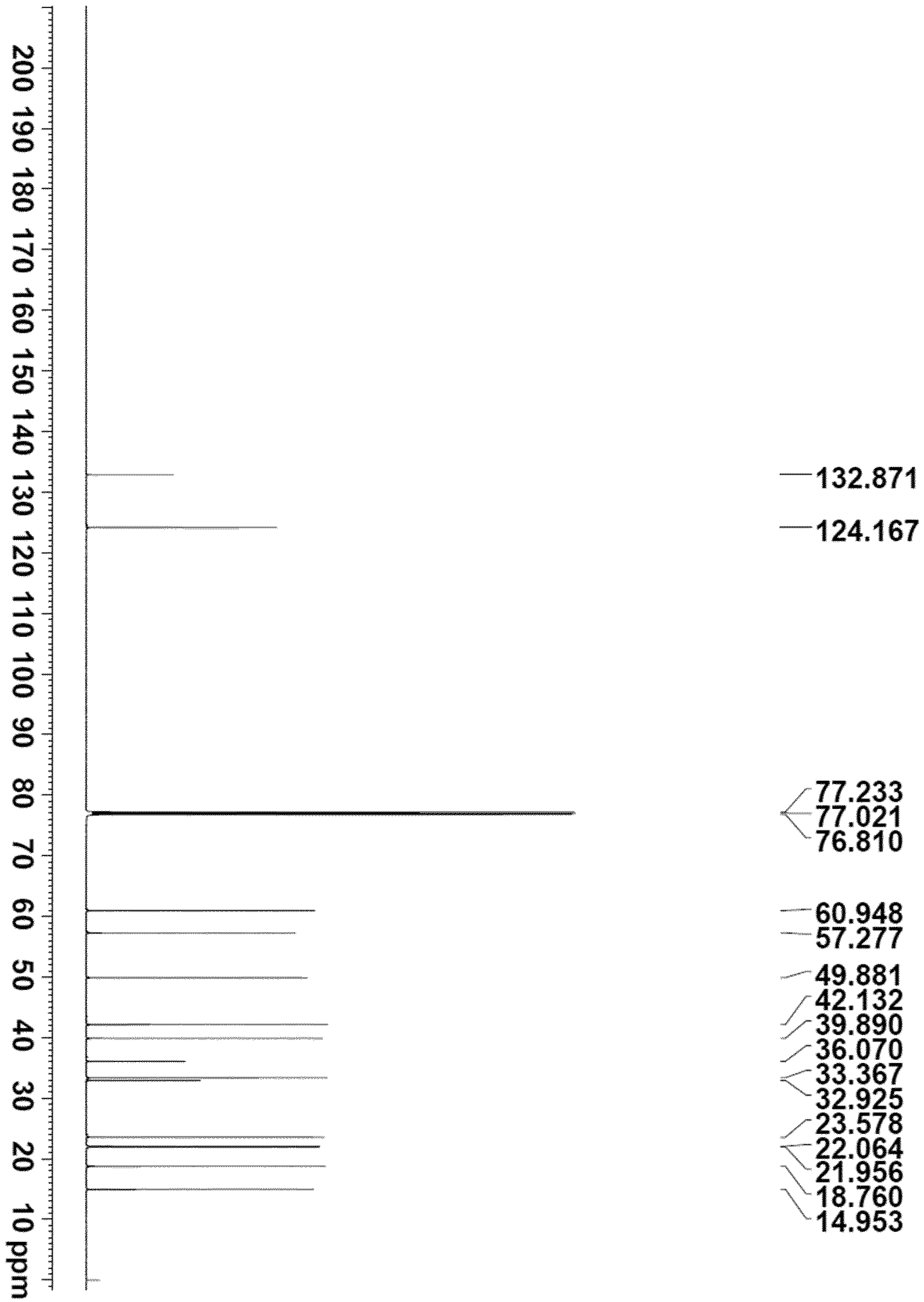


Figure 3

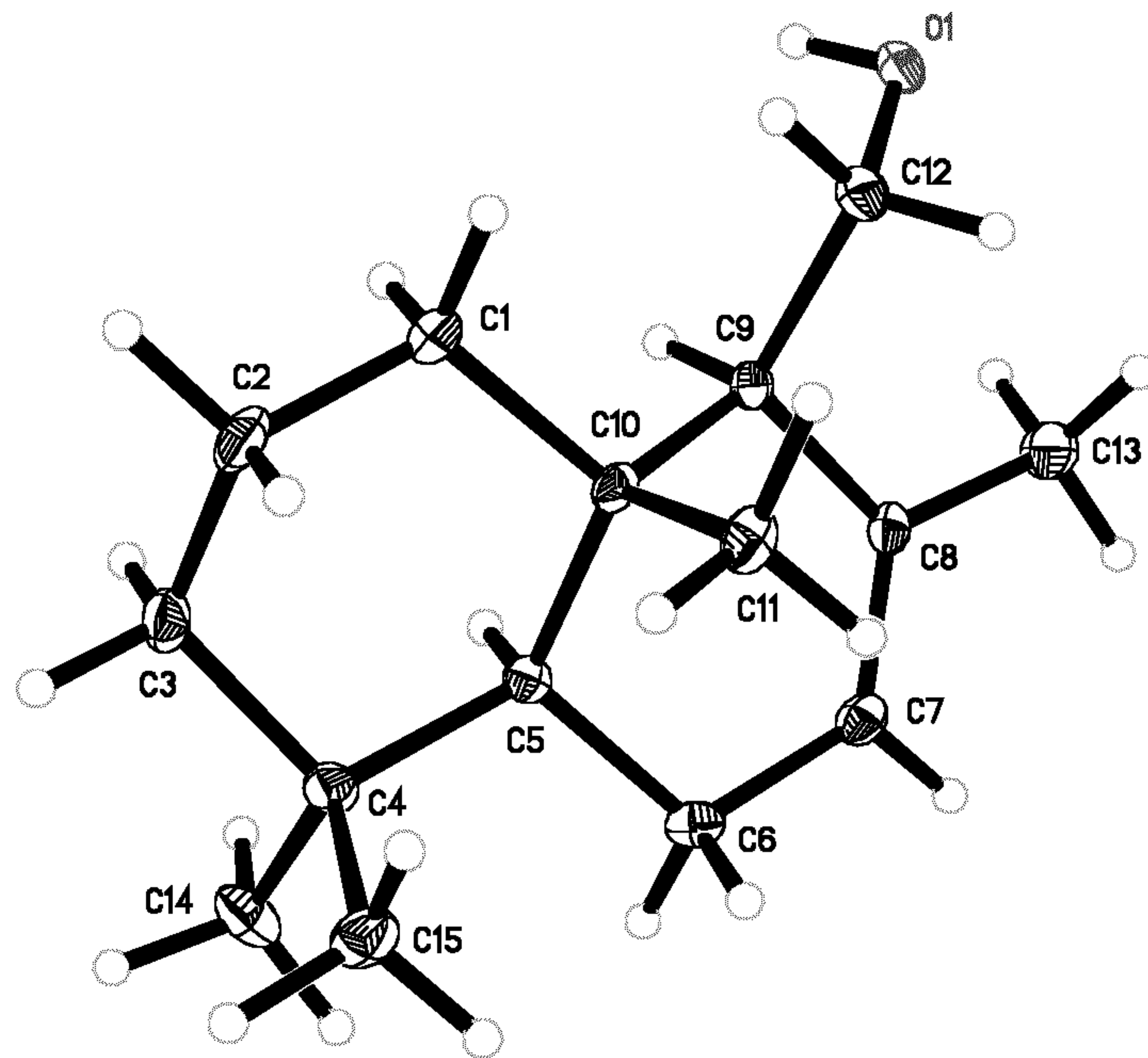


Figure 4.

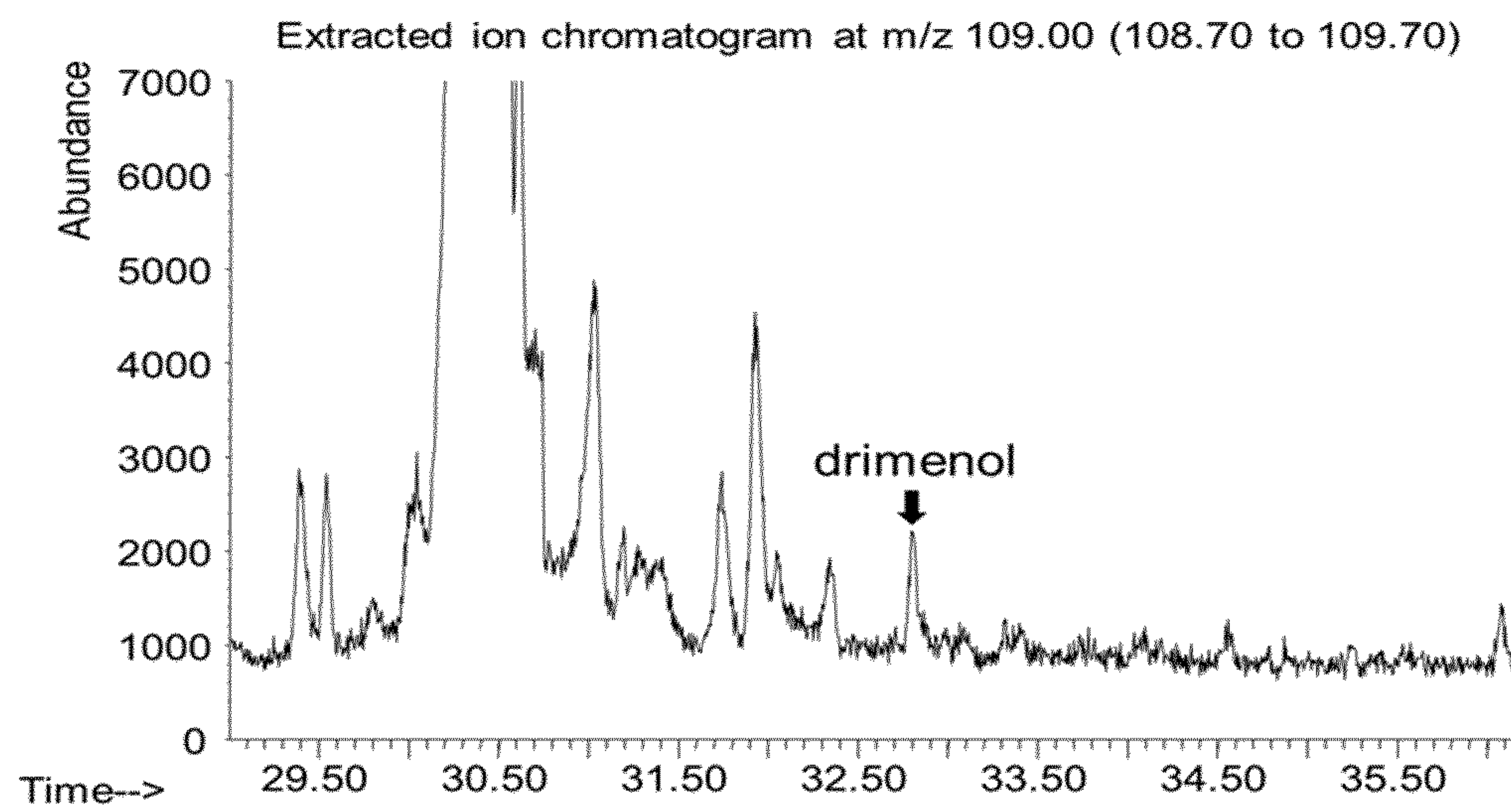


Figure 5

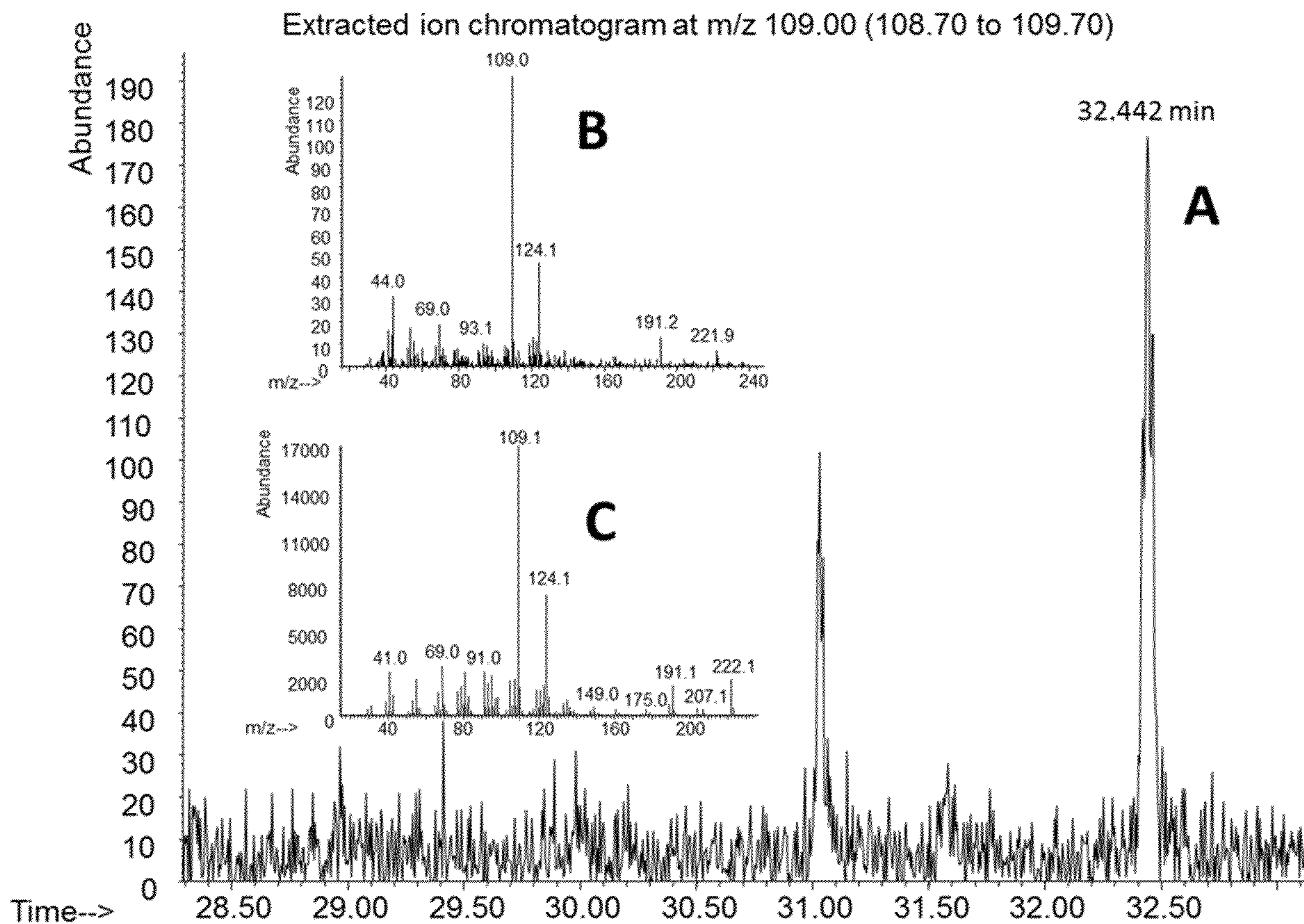


Figure 6