**ORIGINAL ARTICLE**

Incubation of 2-methylisoborneol synthase with the intermediate analog 2-methylneryl diphosphate

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Incubation of synthetic 2-methylneryl diphosphate (2-MeNPP, 10) with 2-methylisoborneol synthase (MIBS) gave a mixture of products that differed significantly from that derived from the natural substrate (E)-2-methylgeranyl diphosphate (3, 2-MeGPP). The proportion of (E)-2-methylisoborneol (1) decreased from 89 to 17% while that of 2-methylenebornane (4) increased from 10 to 26%, with the relative yields of the isomeric homo-monoterpenes 2-methyl-2-bornene (5) and 1-methylcamphene (6) remaining essentially unchanged (≤1% each), as determined by chiral GC–MS analysis. The majority of the product mixture resulting from the MIBS-catalyzed cyclization of 2-MeNPP (10) consisted of the anomalous monocyclic homo-monoterpenes (±)-2-methylbornyl diphosphate (15, 19%) and 2-methyl-α-terpineol (13, 10%), as well as the acyclic derivatives 2-methylnerol (11, 7%) and 2-methylinalool (14, <1%). The steady-state kinetic parameters of the MIBS-catalyzed reaction, determined using [1-3H]-2-methylneryl diphosphate (2-MeNPP), were $k_{\text{cat}} = 0.0046 \pm 0.0003 \text{ s}^{-1}$, $K_m = 18 \pm 6 \mu\text{M}$ and $k_{\text{cat}}/K_m = 2.55 \times 10^2 \text{ M}^{-1} \text{s}^{-1}$. In comparison, the natural substrate 2-MeGPP had a $k_{\text{cat}} = 0.105 \pm 0.007 \text{ s}^{-1}$, $K_m = 95 \pm 49 \mu\text{M}$ and $k_{\text{cat}}/K_m = 1.11 \times 10^3 \text{ M}^{-1} \text{s}^{-1}$. Taken together with earlier X-ray crystallographic studies of MIBS, as well as previous investigations of the mechanistically related plant monoterpen cyclase, bornyl diphosphate synthase, these results provide important insights into the binding and cyclization of both native substrates and intermediates and their analogs.

*The Journal of Antibiotics (2017) 70, 625–631; doi:10.1038/ja.2017.24; published online 1 March 2017*

**INTRODUCTION**

Since the discovery of 2-methylisoborneol (2-MIB, 1) by Gerber in 1969, there has been considerable interest in the detection, bioremediation and biosynthesis of this volatile, odiferous homo-monoterpene (Figure 1).1–4 Possessing a musty odor and muddy off-taste, as well as an extremely low threshold of detection by humans (<10 ng l$^{-1}$), 2-MIB is produced by a wide range of Gram-positive actinomycetes, myxobacteria and cyanobacteria, being second in occurrence only to the common odiferous sesquiterpene geosmin.5

The elucidation of the biosynthesis of 2-MIB at the molecular genetic and biochemical level was independently reported in 2008 by two research groups (Figure 1).5–8 The 2-MIB biosynthetic gene cluster in *Streptomyces coelicolor* was shown to harbor a terpene synthase gene (*sco7700*) translationally coupled to the gene for an S-adenosyl-L-methionine-dependent C-methyl transferase, (*sco7701*), as well as a third gene (*sco7699*) encoding a protein of still unknown function, annotated only as a nucleotide-binding protein.7 This three-gene biosynthetic operon is highly conserved across the genome sequences of more than three dozen bacterial species.5,6 The S-adenosyl-L-methionine-dependent geranyl diphosphate C-methyl transferase (GPPMT; *Sco7701*) was shown to catalyze the first step in the biosynthetic pathway in *S. coelicolor*, the electrophilic methylation of the olefinic C-2 position of geranyl diphosphate (GPP, 2) to form (E)-2-methylgeranyl diphosphate (2-MeGPP, 3).7 *Sco7700*, 2-MIB synthase (MIBS), catalyzes the Mg$^{2+}$-dependent multistep cyclization of the acyclic 2-MeGPP precursor to the bicyclic alcohol 2-MIB (1) which is accompanied by small amounts of the bicyclic homo-monoterpenes 2-methylenebornane (4), 2-methylbornene (5) and 1-methylcamphene (6). The only other enzyme known to utilize 2-MeGPP as substrate is the closely related 2-methylenebornene synthase (Pf1841) of *Pseudomonas fluorescens* PFO-1.9 These findings were fully consistent with the results of independent whole-cell precursor incorporation experiments involving the feeding of [methyl-13C]methionine and samples of deuterated mevalonate to the myxobacterium *Nannocystis exedens* and MS analysis of the resulting 2-MIB in the head-space volatiles.4

The proposed 2-MIB synthase mechanism is based on close analogy to the well-documented cyclization mechanisms established for a wide range of monoterpene synthases (Figure 2).10 MIBS initiates the electrophilic cyclization by ionizing the Mg$^{2+}$-complexed substrate 2-MeGPP (3) to generate the corresponding allylic cation–pyrophosphate ion pair which then collapses to the transoid conformer of the allylic isomer (3R)-2-methylinalyl diphosphate (2-MeLPP, 7). Rotation about the C-2,3-bond generates the cisoid conformer of 7, folded in the anti-endo-boat conformation, which then undergoes ionization and cyclization to generate the (4R)-2-methyl-α-terpinyl...
cation (8). Electrophilic attack on the cyclohexenyl double bond of 8 generates the 2-methyl-2-bornyl cation (9), which is then quenched on the exo face by bound water to generate 2-MIB (1). Competing deprotonation of intermediate 9 or its proximal rearrangement product readily accounts for formation of the co-products 4, 5 and 6. The MIBS cyclization mechanism, including the deduced stereochemistry and conformations of the various intermediates, closely resembles that firmly established for (+)-bornyl diphosphate synthase/α-pinene synthase from Salvia officinalis which converts GPP (2) to (+)-bornyl diphosphate (75%) as well as a mixture of the monoterpenes hydrocarbons (+)-α-pinene, (+)-camphene, (±)-limonene and terpinolene (25% total olefins; Figure 3).11–13 There are of course two key differences between the reactions catalyzed by MIBS and by (+)-bornyl diphosphate synthase/α-pinene synthase: (1) MIBS cyclizes 2-MeGPP instead of GPP. Although MIBS can cyclize GPP to a mixture of monocyclic and bicyclic monoterpene hydrocarbons (+)-α-pinene, (+)-camphene, (±)-limonene and terpinolene (25% total olefins; Figure 3).11–13 There are of course two key differences between the reactions catalyzed by MIBS and by (+)-bornyl diphosphate synthase/α-pinene synthase: (1) MIBS cyclizes 2-MeGPP instead of GPP. Although MIBS can cyclize GPP to a mixture of monocyclic and bicyclic monoterpenes, the observed kcat is four orders of magnitude lower than that for the natural methylated substrate 2-MeGPP.7 (2) The final step in the MIBS-catalyzed cyclization cascade is the quenching of the 2-methyl-2-bornyl cation exclusively on the exo face by a bound water (Figure 2), while in the (+)-bornyl diphosphate synthase/α-pinene synthase-catalyzed reaction, the paired pyrophosphate ion is recombined with the bornyl cation exclusively by endo attack (Figure 3).

High-resolution crystal structures have recently been determined for MIBS in both an unliganded state and in complex with a number of substrate or intermediate analogs.14,15 The 1.80–1.95 Å structures of MIBS bound to geranyl-S-thiolodiphosphate and 2-fluorogeranyl diphosphate, respectively, show each analog coordinated with two 2 Mg2+ ions and bound in extended conformations in which the 6,7-double bond of each analog is twisted away from the parallel relationship expected for the postulated anti-endo-boat conformation of the actual cyclizing cisoid (3R)-2-MeLPP intermediate (Figure 2).14 Although this deviation may simply be the result of crystallographic trapping of a thermodynamically stable, rather than a kinetically or mechanistically relevant conformer, more interestingly it reflects the fact that the required 180° rotation of the 2-propenyl substituent about the C-2,3 bond of the anti-endo-boat conformer of the (3R)-2-methylinalyl diphosphate intermediate is sterically forbidden by a clash with the 6,7-double bond that is augmented by the presence of the C-2-methyl substituent. It is thus probable that adoption of the endo-boat conformation takes place subsequent to conversion of the transoid to the cisoid conformer of 2-MeLPP. Such extended 6,7-double bond conformations of bound substrate analogs have previously been observed in the structures of a number of canonical monoterpene synthases.16,17 Also thought-provoking were the unexpected results from co-crystallization of MIBS with racemic
2-fluorolinalyl diphosphate (2-FLPP), which had been predicted to be an unreactive analog of the natural (3R)-2-methylinalyl diphosphate intermediate due to the strongly electron-withdrawing 2-fluoro substituent. The 2.00 Å structure of the resulting complex revealed that the MIBS-bound 2-FLPP had undergone a reverse allylic rearrangement to generate bound 2-fluoroneryl diphosphate in complex with 2 Mg\(^{2+}\) ions.\(^{15}\) Also unexpectedly, it was found that in solution MIBS converted the supposedly unreactive 2-FLPP to camphor, no doubt formed by cyclization of 2-FLPP and spontaneous elimination of hydrogen fluoride from enzymatically generated 2-fluoro-2-isoborneol. The observation of enzyme-generated 2-fluoroneryl diphosphate trapped within the active site of MIBS raises the question whether the cis geometric isomer of 2-FGPP is a true intermediate of the MIBS-catalyzed ionization and cyclization of 2-FLPP, the nominally unreactive analog of the natural intermediate 2-MeLPP (7), or merely an anomalous shunt metabolite of the reverse allylic diphosphate rearrangement. We have therefore now examined the incubation of MIBS with 2-MeNPP (10), the cis isomer of the natural substrate, 2-MeGPP (3), and report the results below.

RESULTS

Preparation of 2-methylneryl diphosphate (10) and (±)-2-methyl-α-terpineol (13)

To prepare 2-MeNPP (11), we used a modification of the previously reported synthesis of the trans isomer 2-MeGPP (3; Figure 4).\(^{7}\) Wadsworth–Horner–Emmons reaction of 6-methyl-5-hepten-2-one with triethyl 2-phosphonopropionate followed by LiAlH\(_4\) reduction of the resulting mixture of \(Z\) and \(E\) conjugated esters gave a 1:1 mixture of \((Z)\)-2-methylenol (11) and \((E)\)-2-methylgeraniol (12) which were cleanly resolved by flash column chromatography (see Supplementary Information for details). The purified 2-methylenol (11) was reacted with triphenylphosphine and CCl\(_4\) in an Appel reaction and the resulting 2-methylenyl chloride was then directly reacted without further purification with tris(tetramethylammonium) hydrogen pyrophosphate to give 2-MeNPP (10). The corresponding [1\(^{-3}\)H]-2-MeNPP ([1\(^{-3}\)H]-10) was similarly prepared by oxidation of the mixture of 2-methylnerol and 2-methylgeraniol to the derived mixture of aldehydes, followed by reduction with NaBH\(_4\) and chromatographic separation of the resulting mixture of [1\(^{-3}\)H]-2-methylnerol and [1\(^{-3}\)H]-2-methylgeraniol using AgNO\(_3\)-impregnated silica gel.

For the planned enzyme incubations, we also needed an authentic reference sample of the homo-monoterpenyl alcohol (±)-2-methyl-α-terpineol (13),\(^{18}\) which was readily prepared by biomimetically-modeled, p-TsOH-catalyzed solvolysis of 2-methylenol (11) in nitromethane (Figure 4).

In vitro incubation of 2-methylneryl diphosphate with 2-methyliso borneol synthase

Before examining the cyclization of 2-MeNPP by methylisoborneol synthase, we carried out individual control incubations in the absence of enzyme with 2-MeGPP and 2-MeNPP and analyzed the pentane-soluble extracts by chiral GC–MS in order to identify and quantify the formation of Mg\(^{2+}\)-catalyzed solvolysis products (Figures 5a and b). The control incubation with 2-MeGPP produced only racemic (±)-2-methyllinalool (14). The exclusive formation of this tertiary allylic alcohol contrasts with the well-known Mg\(^{2+}\)-catalyzed solvolysis of GPP itself, which typically gives a 3:1 mixture of linalool and geraniol.\(^{18}\) The preferred formation of 2-methyllinalool (14) by solvolysis of 2-MeGPP presumably reflects the higher free energy of 2-methylgeraniol due to the increased steric hindrance about the tetra-substituted double bond compared to geraniol. In contrast, the control Mg\(^{2+}\)-catalyzed solvolysis of 2-MeNPP gave a mixture of three racemic products, consisting of acyclic (±)-2-methyllinalool (14) as well as the monomeric (±)-2-methylisoborneol (15), and (±)-2-methyl-α-terpineol (13; Figure 5b). The Mg\(^{2+}\)-catalyzed formation of both (±)-13 and (±)-15 from 2-MeGPP, but not from 2-MeNPP, is consistent with the fact that the \((E)\)-allylic diphosphate 2-MeGPP (3) is geometrically incapable of direct cyclization. As a positive control, incubation of 2-MeGPP (3) with 2-MIB synthase and chiral GC–MS analysis confirmed the formation of \((-\)\()-2\text{-}2\text{-}\text{MIB}\) (I) as the major product (89%) accompanied by minor amounts of enantiomerically pure 2-methylenebornane (4; 10%), 1-methylcamphene (6; <1%) and 2-methylene-2-bornene (5; <1%), as previously observed (Figure 5c and Table 1).\(^{6,7}\)

Incubation of 2-MeNPP (10) with MIBS and analysis by chiral GC–MS revealed the production of the same group of homochiral bicyclic homo-monoterpenes, \((-\)\()-2\text{-}2\text{-}\text{MIB}\) (1, 17%), 2-methylenebornane (13, 26%), 1-methylcamphene (6) (<1%) and 2-methylene-2-bornene (5) (<1%), but in both substantially different proportions and lower overall yield compared to 2-MeGPP (Figures 5d and 6). Significantl, four additional, previously unobserved products...
Figure 5 Chiral GC–MS analysis (total ion current, TIC) of incubations with methylisoborneol synthase (MIBS). (a) Control incubation of 2-MeGPP (3) in Mg$^{2+}$-containing buffer without MIBS. (b) Control incubation of 2-MeNPP (10) in Mg$^{2+}$-containing buffer without MIBS. (c) Incubation of MIBS with 2-MeGPP (3). (d) Incubation of MIBS with 2-MeNPP (10). Previously unobserved product peaks are shown in red. Peaks denoted with * are geraniol internal standard. A full color version of this figure is available at the *The Journal of Antibiotics* journal online.
transformation of 2-methylneryl diphosphate (2-MeNPP; Figure 6)

Mechanism of the methylisoborneol synthase (MIBS)-catalyzed 2-MeNPP (10) with 2-MIB synthase

Table 1 Distribution of products from the incubation of 2-MeGPP (3) and 2-MeNPP (10) with 2-MIB synthase

| Substrate | MIB (1) | 4 | 5 | 6 | 13 | (±)-15 | 11 | 14 |
|-----------|---------|---|---|---|----|-------|----|----|
| 2-MeGPP (3) | 89 | 10 | <1 | <1 | ND | ND | ND | ND |
| 2-MeNPP (10) | 17 | 26 | <1 | <1 | 10 | 39 | 7 | <1 |

(10/0)b (17/22)b (Q/ <1)b

Abbreviations: MeGPP, methylgeranyl disphosphate; MeNPP, methylneryl diphosphate; MIB, methylisoborneol; ND, not detected.

8Percent corrected for background Mg2+-catalyzed solvolysis.

Steady-state kinetics

The steady-state kinetic parameters for the MIBS-catalyzed conversion of [1-3H]-2-MeNPP (10) to total pentane-soluble homo-monoterpene products were determined and directly compared with those re-determined for the natural substrate [1-3H]-2-MeGPP (3) under identical conditions (Table 2). While the turnover number, kₐ⁄⁄, for 2-MeNPP (0.0045 s⁻¹) was ~20-fold lower than that for 2-MeGPP, the observed kₐ⁄⁄Kₘ of 255 M⁻¹ s⁻¹ for 2-MeNPP was only 4-fold lower than that for 2-MeGPP (1105 M⁻¹ s⁻¹), reflecting a partially compensating ~5-fold decrease in the observed Kₘ for 2-MeNPP compared to 2-MeGPP.

Table 2 Steady-state kinetic parameters for total product formation from incubation with methylisoborneol synthase

| Substrate | kₐ⁄⁄ (s⁻¹) | Kₘ (μM) | kₐ⁄⁄Kₘ (M⁻¹ s⁻¹) |
|-----------|------------|--------|-----------------|
| 2-MeGPP (3) | 0.105 ± 0.007 | 95 ± 49 | 1110 |
| 2-MeNPP (10) | 0.0046 ± 0.0003 | 18 ± 6 | 255 |

DISCUSSION

Although (Z)-2-MeNPP (10, 2-MeNPP) shows only a modest fourfold decrease in the specificity constant kₐ⁄⁄Kₘ compared to the natural substrate, the trans geometric isomer (E)-2-MeGPP (3), the substantially altered product distributions resulting from incubation of 2-MeNPP with 2-MIB synthase establish that 2-MeNPP is a poor surrogate for 2-MeGPP. Thus, while 2-MeNPP could, in principle, be ionized to the same cisoid allylic cation–pyrophosphate ion pair as the presumptive intermediate (3R)-2-methylallyl disphosphate (7, 2-MeLPP), in practice incubation of 2-MeNPP with 2-MIB synthase generated a substantially altered array of bicyclic and monocyclic homo-monoterpene products. Thus, although cyclization of 2-MeNPP generated only a single enantiomer of each of the four natural bicyclic natural products, 2-MIB (1), 2-methylbornene (4), 2-methylbornene (5) and 1-methylcamphene (6), the relative proportion of 1 was reduced more than fivefold from 89% to 17% of the total product mixture, only partially offset by an increase in the fraction of 2-methylenebornane (4) which increased from 10 to 26% (Table 1). The majority of the product mixture resulting from the MBS-catalyzed cyclization of 2-MeNPP (10) consisted of the anomalous monocyclic homo-monoterpenes (±)-2-methylbornene (15, 39%) and a single enantiomer of 2-methyl-α-terpineol (13, 10%), as well as the acyclic derivatives 2-methyleneborane (11, 7%) and a single enantiomer of 2-methylallylalool (14, <1%). Interestingly, while 13, 14 and 15 are not detected among the products of the in vitro incubation of 2-MeNPP with 2-MeGPP, all three of these homo-monoterpenes, of unspecified chiral purity, have been reported as minor components of the volatile head-space extract of a small number of 2-MIB-producing actinomycetes.19,20

Substantial changes in product distribution have been reported when the cis isomer neryl diphosphate is incubated with monoterpene synthases in place of the natural trans allylic substrate, geranyl diphosphate. For example, in spite of only minor differences in overall kₐ⁄⁄, incubation (+)-BPP/(+)-pinene synthase with NPP instead of GPP resulted in a decrease in the proportion of the bicyclic monoterpene α-pinene and camphene from 79% to 23%, while the fraction of the monocyclic products limonene and α-terpinolene exhibited a substantial increase from 15 to 71% of total monoterpenes.21 The enhanced formation of abortive cyclization products is almost certainly the consequence of aberrant positioning of 2-MeNPP and NPP in the active sites of the respective homo-monoterpene or monoterpene synthases, compared to the binding of the native substrates 2-MeGPP or GPP.

The fact that cyclization of 2-MeNPP by 2-MIB synthase generates the naturally occurring enantiomers of 2-MIB (1), 2-methyleneborne (4), 2-methylbornene (5) and 1-methylcamphene (6) indicates that at least a portion of this anomalous substrate is bound to MIBS in a position and conformation that is compatible with, and the fact that the derived intermediate (3R)-2-MeLPP (7) in the MIBS active site. Although the 2-methylbornene (15) and 2-methyl-α-terpineol (13) formed from 2-MeNPP are both formally derivable from a common 2-methyl-α-terpinyl cation, the fact that 15 is generated as a racemic mixture while 13 is produced as...
a single enantiomer indicates clearly that they cannot both be derived from a single mononuclear carbocation intermediate. The surrogate 2-MeNPP substrate therefore must be bound in at least two distinct conformations during MIBS-catalyzed cyclization. The fact that only a single enantiomer of each of the bicyclic, monocyclic and acyclic alcohols 1, 13 and 14 is produced by incubation of 2-MeNPP with MIBS also suggests that the same bound water molecule is responsible for quenching the corresponding carbocation intermediates.

**EXPERIMENTAL PROCEDURE**

**Materials**

Reagents and solvents were purchased from Sigma-Aldrich (St Louis, MO, USA) or Fisher Scientific (Waltham, MA, USA), were of the highest quality available, and were used without further purification. NaBH₄ solution (6 μmol, 80 Ci mmol⁻¹) specific activity, 500 mCi total in 1 ml of 0.1% NaOH was purchased from American Radiolabeled Chemicals (St Louis, MO, USA). Isopropylthio-D-galactopyranoside was purchased from Invitrogen (Waltham, MA, USA). Ni-NTA affinity resin was from Qiagen (Valencia, CA, USA). Amicon Ultra Centrifugal Filter Units (Amicon Ultra-15 10 000 MWCO) were purchased from Millipore (St Charles, MO, USA). Puriﬁcation SCO7700 protein was overexpressed and puriﬁed as previously described.7 The purity of Sco7700 protein was overexpressed and purified as previously described.7

**Methods**

GC – MS analyses were performed using an Agilent 5977A Series GC/MSD instrument (70 eV, electron impact), a 1-μl injection volume, and a 3 min solvent delay. Achiral GC – MS conditions used an HP-5 ms capillary column (0.25 mm ID × 30 m length × 0.25 μm film, Agilent Technologies (Santa Clara, CA, USA)) and a temperature program with a 2 min hold at 60 °C, a 20 °C min⁻¹ increment to 280 °C, followed by a 2 min hold at 280 °C.

Chiral GC – MS separations were performed using a CP-ChiraldS-Dex column (0.32 mm ID × 25 m length × 0.25 μm film, Agilent) and a temperature program with a 1 min hold at 50 °C, a 10 °C min⁻¹ increment to 200 °C, followed by a 1 min hold at 200 °C. Compounds detected by GC – MS were directly compared to their authentic standards using the MassFinder 4.2.1 program (http://www.massfinder.com). Retention indices were measured using C₆-C₂₀ and C₁₀-C₄₀ alkane standards. LC-MS analyses were performed using a Finnigan LXQ LC-MS instrument in negative (electrospray ionization ESI) mode, with a Waters Symmetry C₁₈ column (0.25 mm ID × 30 m, 5 μm, Waters) and the mixture was reﬁltered and concentrated to give 310 mg of yellowish liquid (72% crude chemical yield, 375 mCi, 75% radiochemical yield). The residue was puriﬁed by flash chromatography on 10% (w/w) AgNO₃-impregnated silica gel (EtOAc/hexanes (19:4); 4 × 16 cm column) giving 100 mg of [1-³H]-2-methylnerol ([1-³H]-11), 25% chemical yield, 128 mCi, 26% radiochemical yield and 210 mg of [1-³H]-2-methylgeranyl ([1-³H]-12), 48% chemical yield, 247 mCi, 49% radiochemical yield. The puriﬁed [1-³H]-2-methylnerol ([1-³H]-11) and [1-³H]-2-methylgeranyl ([1-³H]-12) were individually converted to the corresponding diphasphates esters using the same method employed for the synthesis of unlabeled 10 and 11 to yield 98 mg of [1-³H]-2-MeNPP ([1-³H]-10), 37% chemical yield, 6.2 mCi total radioactivity, 2.2% radiochemical yield, 28 mCi/mmoll specific activity and 200.1 mCi of [1-³H]2-MeGePP ([1-³H]-5), 36% chemical yield, 105 mCi total radioactivity, 36% radiochemical yield and 232.8 mCi/mmoll specific activity. The NMR spectra of ([1-³H]-10) and of [1-³H]-3 matched those of unlabeled 10 and 3.

2-Methyl-α-terpineol (13)

p-TsOH (40 mg, 0.20 mmol) was added to a solution of 2-methylnerol (200 mg, 1.19 mmol) in 10 ml of MeNO₂. The reaction mixture was stirred at rt for 1 h and monitored by TLC (51:hexanes/ethyl acetate). The crude product was precipitated in vacuo and purified by column chromatography (51:hexanes/ethyl acetate) yielding 44 mg of 2-methyl-α-terpineol (13, 22% yield). The NMR data matched those previously reported for 13 prepared by an alternative method.19 H NMR (400 MHz, CDCl₃) δ 1.98 (m, 2H, H-6), 1.85 (m, 2H, H-3), 1.72 (m, 1H, H-5), 1.65 (s, 6H, H-10, 11), 1.48 (m, 2H, H-4), 1.22 (s, 6H, H-8, 9). 13C NMR (100 MHz, CDCl₃) δ 19.4 (C-9, CH₃), 19.5 (C-8, CH₃), 24.3 (C-3, CH₃), 26.3 (C-6, CH₃), 27.3 (C-4, CH₃), 32.7 (C-11, CH₃), 33.2 (C-10, CH₃), 45.8 (C-5, CH₃), 72.9 (C-7, C), 125.0 (C-2, C), 125.8 (C-1, C). GC-MS (EI) m/z 168.1.
Incubations with 2-methylisoborneol synthase (SCO7700)

Purified 2-MIB synthase (SCO7700; 10 μM) was added to a glass test tube containing 3.0 ml of assay buffer (50 mM PIPES, 15 mM MgCl2, 5 mM β-mercaptoethanol and 20% glycerol, pH 6.7) and 60 μM of either 2-MeNPP (10) or 2-MeGPP (3). The enzymatic reaction mixture was overlaid with 3.0 ml of pentane and incubated at 30 °C for 2 h. The enzymatic products were extracted with 3×3.0 ml of pentane and the organic extracts dried over 

Na2SO4, filtered, concentrated in vacuo to 100 μl and analyzed by GC–MS.

Control reactions, measuring the background Mg2+-catalyzed hydrolysis of each substrate, were conducted similarly as above minus the addition of SCO7700 to the incubation mixture. An internal geraniol standard (25 pmol) of each substrate, were conducted similarly as above minus the addition of 16.5 Ci mol−1 of either [1-3H]-2-MeNPP ([1-3H]-2-MeNPP, 0.9 Ci/mol) or 2-MeGPP ([1-3H]-2-MeGPP, ([1-3H]-3, 16.5 Ci mol−1). The reactions were initiated by the addition of 0.20 μl of SCO7700 protein, overlaid with 1 ml of pentane and incubated at 30 °C for 2 h. The enzymatic products were extracted 3 more times with 1 ml of ether, and all organic extracts were concentrated to 40 μl.

Supplementary Information accompanies the paper on The Journal of Antibiotics website (http://www.nature.com/ja).