

Demonstration of the ethylmalonyl-CoA pathway by using ^{13}C metabolomics

Rémi Peyraud^a, Patrick Kiefer^a, Philipp Christen^a, Stephane Massou^b, Jean-Charles Portais^{b,c}, and Julia A. Vorholt^{a,1}

^aInstitute of Microbiology, Eidgenössische Technische Hochschule Zurich, 8093 Zurich, Switzerland; ^bUnité Mixte de Recherche 5504, Unité Mixte de Recherche 792, Ingénierie des Systèmes Biologiques et des Procédés, Centre National de la Recherche Scientifique, Institut National de la Recherche Agronomique, Institut National des Sciences Appliquées, 31400 Toulouse, France; and ^cFaculté Sciences de la Vie Université Paul Sabatier, 31062 Toulouse, France

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The assimilation of one-carbon (C1) compounds, such as methanol, by serine cycle methylotrophs requires the continuous regeneration of glyoxylate. Instead of the glyoxylate cycle, this process is achieved by a not yet established pathway where CoA thioesters are known to play a key role. We applied state-of-the-art metabolomics and ^{13}C metabolomics strategies to demonstrate how glyoxylate is generated during methylotrophic growth in the isocitrate lyase-negative methylotroph *Methylobacterium extorquens* AM1. High-resolution mass spectrometry showed the presence of CoA thioesters specific to the recently proposed ethylmalonyl-CoA pathway. The operation of this pathway was demonstrated by short-term ^{13}C -labeling experiments, which allowed determination of the sequence of reactions from the order of label incorporation into the different CoA derivatives. Analysis of ^{13}C positional enrichment in glycine by NMR was consistent with the predicted labeling pattern as a result of the operation of the ethylmalonyl-CoA pathway and the unique operation of the latter for glyoxylate generation during growth on methanol. The results also revealed that 2 molecules of glyoxylate were regenerated in this process. This work provides a complete pathway for methanol assimilation in the model methylotroph *M. extorquens* AM1 and represents an important step toward the determination of the overall topology of its metabolic network. The operation of the ethylmalonyl-CoA pathway in *M. extorquens* AM1 has major implications for the physiology of these methylotrophs and their role in nature, and it also provides a common ground for C1 and C2 compound assimilation in isocitrate lyase-negative bacteria.

^{13}C labeling | CoA ester | methylotroph | one-carbon metabolism | glyoxylate regeneration

Methylotrophic bacteria are organisms capable of using reduced carbon compounds, such as methanol or methane, as sole sources of carbon and energy, and they play a key role in carbon cycling in their environment. They also represent promising organisms in biotechnology for the conversion of one-carbon (C1) substrates to value-added products (1). The elucidation of the mechanisms enabling growth on reduced C1 compounds of *Methylobacterium extorquens* AM1, one of the most studied methylotrophs, has been a longstanding goal, and although great progress has been made (2–5), it is still not fully achieved. A key point has been to understand how the bacterium incorporates C1 units into cell material. The serine cycle was elucidated in this organism during the early 1960s by Quayle and coworkers (6–9). The assimilation of C1 units by this pathway requires continuous regeneration of glyoxylate from acetyl-CoA and can be achieved, in principle, via the well-known glyoxylate cycle (10). However, Dunstan and coworkers (11–14) showed in 1972 and 1973 that *M. extorquens* AM1 lacks the key enzyme of the glyoxylate cycle, isocitrate lyase, but has an alternative route involving oxidation of acetate to glyoxylate that functions during growth on both C1 and C2 compounds. Also, other organisms, including the photosynthetic *Rhodospira rubra* are known to require an alternative to the glyoxylate cycle when growing on C2 substrates or on substrates that are converted into acetyl-CoA to enter central metabolism (15–18).

Recent studies, including mutant analyses, gene predictions, enzyme assays, and metabolite studies in *M. extorquens* AM1, have led to the observation that a complex sequence of CoA thioester derivatives is involved in glyoxylate regeneration, resulting in the hypothesis of the so-called glyoxylate regeneration cycle (GRC) (19, 20) [Fig. 1 and supporting information (SI) Table S1]. According to this pathway, a C5 compound, methylsuccinyl-CoA, is formed from the condensation of 2 acetyl-CoA molecules plus 1 CO_2 and is decarboxylated twice in a process similar to valine degradation. The specific intermediates of the GRC are isobutyryl-CoA, metacrylyl-CoA, and hydroxyisobutyryl-CoA, and the result is the formation of propionyl-CoA. Subsequently, propionyl-CoA is transformed to malate, from which 1 glyoxylate and 1 acetyl-CoA are generated (20). More recently, a second hypothesis, the ethylmalonyl-CoA pathway (EMCP), was proposed from studies of C2 assimilation pathways in *R. sphaeroides* (21–23). This pathway (Fig. 1 and Table S1) includes the formation of methylsuccinyl-CoA, which is further converted to methylmalyl-CoA, from which both glyoxylate and propionyl-CoA are released by cleavage (22). The propionyl-CoA can then be converted to C4 compounds and assimilated as cell material (23).

The 2 pathways mentioned above are still hypothetical, and none has been firmly demonstrated to operate *in vivo*. They differ strikingly in terms of carbon balance and, therefore, overall carbon yield for methylotrophic growth. The GRC includes a net decarboxylation step, whereas the ethylmalonyl-CoA pathway includes net carboxylation steps. This makes the second pathway more efficient in terms of carbon assimilation and has important implications with regard to the physiology of these methylotrophs and their actual biotechnological potential.

In this work, we combined state-of-the-art metabolomics and ^{13}C metabolomics strategies to examine the pathway of glyoxylate regeneration occurring in *M. extorquens* AM1 under methylotrophic conditions. The development of an original liquid chromatography high-resolution mass spectrometry (LC-HRMS) method allowed for the identification of almost all intermediates of the ethylmalonyl-CoA pathway. Detailed and conclusive information regarding the operation of the ethylmalonyl-CoA pathway as the predominant process for glyoxylate formation was obtained from ^{13}C -labeling experiments in which kinetic isotopomer profiles collected by LC-HRMS during short-term ^{13}C -labeling experiments were combined with steady-state isotopomer distributions measured by NMR.

Results

Identification of CoA Thioesters by Liquid Chromatography-Mass Spectrometry (LC-MS). The pathways proposed for the conversion of acetyl-CoA to glyoxylate in isocitrate lyase-negative bacteria involve

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¹To whom correspondence should be addressed. E-mail: vorholt@micro.biol.ethz.ch.

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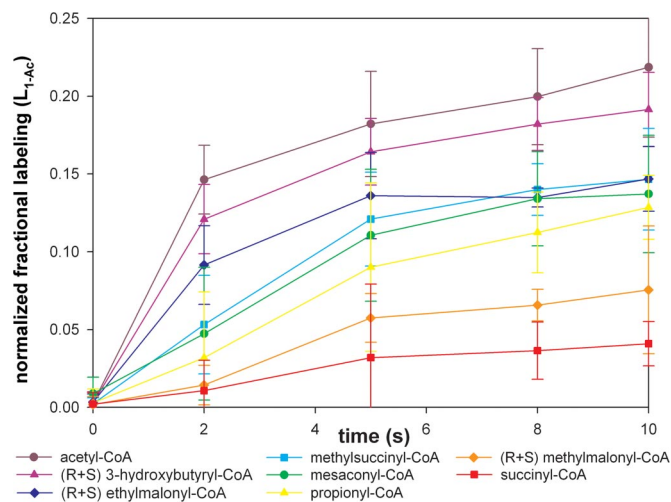


Fig. 4. Kinetics of ^{13}C label incorporation in CoA thioesters after addition of $[1-^{13}\text{C}]$ acetate to $[^{12}\text{C}]$ methanol-grown *M. extorquens* AM1 cells. $L_{1-\text{Ac}}$ represents the percent of ^{13}C label incorporated in a given metabolite, normalized to the maximal number of carbon atoms received from the first carbon of acetate. Results are mean values + SDs from 3 independent biological replicates.

propionyl-CoA was essentially indistinguishable from that of the 2 C5 CoA thioesters, methylsuccinyl-CoA and mesaconyl-CoA, indicates that no loss of labeled carbon occurred. After propionyl-CoA, methylmalonyl-CoA was found to be labeled, followed by succinyl-CoA. Their isotopomer profiles were similar to those of metabolites in the early steps of the pathway, indicating that they are all connected through the same sequence of reactions.

After 10 s of incubation, the percentage of ^{13}C label incorporated into ethylmalonyl-CoA did not increase further, unlike other CoA metabolites (Fig. 4). Further investigations are needed to understand this observation. However, the mass isotopomer distribution (relative proportions of M0, M+1, M+2, etc.) of ethylmalonyl-CoA remained constant over time, suggesting that the process by which the label was transferred from $[^{13}\text{C}]$ acetate to this metabolite did not change during the incubation period.

The mass isotopomers of few CoA intermediates were difficult to quantify because of small pool sizes. Therefore, replicate samples collected at the same time points were pooled and concentrated. The concentrated samples indicated that crotonyl-CoA was labeled similarly to 3-hydroxybutyryl-CoA and before butyryl-CoA and ethylmalonyl-CoA. Butyryl-CoA was found to be labeled more than ethylmalonyl-CoA and similarly to methylsuccinyl-CoA. β -methylmalyl-CoA was labeled before the other C5 compounds mentioned above and with a high level of M+2, which supposes an incorporation of label from mesaconyl-CoA plus an entry of label from glyoxylate due to the reversibility of the reaction catalyzed by L-malyl-CoA/ β -methylmalyl-CoA lyase (22, 24).

Taken together, the label incorporation in CoA thioesters is in agreement with the sequence of reactions shown in Fig. 1 and suggests the operation of the ethylmalonyl-CoA pathway (23).

Mass Isotopomers of Propionyl-CoA. Because the fate of carbon atoms derived from $[1-^{13}\text{C}]$ acetate is different in the 2 proposed pathways, the generated propionyl-CoA molecules do not receive the same number of carbon atoms; i.e., doubly labeled M+2 propionyl-CoA would be formed according to the ethylmalonyl-CoA pathway (Fig. 3), whereas only singly labeled M+1 propionyl-CoA would be formed according to the GRC as a result of carbon loss as CO_2 . Examination of the evolution of the mass isotopomer distribution of propionyl-CoA during $[1-^{13}\text{C}]$ acetate-labeling experiments showed increases of M+2 until $24.1\% \pm 2.6\%$ at 90 s of

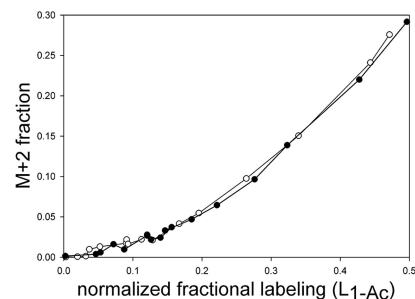


Fig. 5. Comparison of the time course evolution of M+2 isotopomeric fractions in propionyl-CoA (\circ) and methylsuccinyl-CoA (\bullet) during $[1-^{13}\text{C}]$ acetate labeling experiments carried out with methanol-grown *M. extorquens* AM1 cells. The parallel development of the M+2 fraction in the 2 metabolites indicated that no loss of carbon occurred in the process by which methylsuccinyl-CoA was converted to propionyl-CoA.

incubation time. To determine whether a decarboxylation step operates between methylsuccinyl-CoA and propionyl-CoA, the proportion of M+2 isotopomers in methylsuccinyl-CoA and propionyl-CoA during the entire labeling experiment was compared (Fig. 5). The proportion was found to be the same, indicating that no decarboxylation reaction occurred, as would have been observed if the GRC was operating under our experimental conditions.

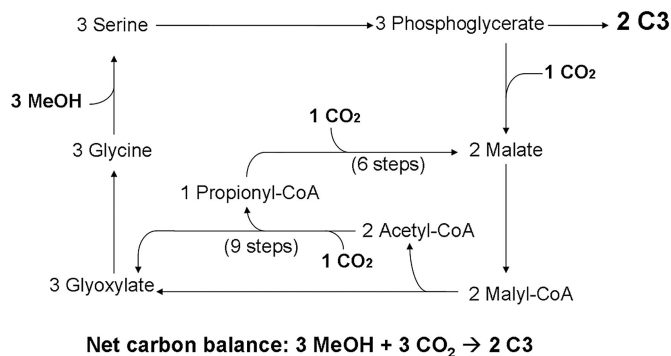
Analysis of Glycine Isotopomers Generated from $[^{13}\text{C}]$ Methanol Under Pure Methylothetic Conditions.

To examine the contribution of the ethylmalonyl-CoA pathway to glyoxylate regeneration under pure methylothetic conditions, a model of glyoxylate metabolism was built up to simulate the theoretical fate of carbon in the central metabolic network (serine cycle, glyoxylate regeneration, and citric acid cycle) of *M. extorquens* AM1 from $[^{13}\text{C}]$ methanol and CO_2 at natural abundance. This model was used to calculate the steady-state isotopomer distribution in glyoxylate expected to occur when each pathway (either GRC or EMCP) was considered separately (for information on underlying considerations and results, see Fig. S1). By operation of the GRC (20), the 2 major isotopomers are expected to be $^{12}\text{C}_1-^{13}\text{C}_2$ and $^{13}\text{C}_1-^{12}\text{C}_2$, and they should represent 41.1% and 33.3%, respectively, of all glyoxylate isotopomers (Table 1). According to the ethylmalonyl-CoA pathway (23), the 2 major isotopomers are expected to be $^{12}\text{C}_1-^{13}\text{C}_2$ and $^{12}\text{C}_1-^{12}\text{C}_2$, and their steady-state proportions should represent 57.1% and 39.1%, respectively, of all glyoxylate isotopomers (Table 1). Taken together, the results of these simulations revealed that the 2 pathways result in strongly discriminative labeling patterns in glyoxylate.

Notably, Large et al. (7) in 1962 used ^{14}C -labeling strategies to investigate the serine cycle in *M. extorquens*. They found disproportionate labeling, in which 92.5% of the label from $[^{14}\text{C}]$ methanol incorporated into glycine was recovered in the C2 position. Although these results found almost half a century ago were not interpreted in that way, they are consistent with the operation of the ethylmalonyl-CoA pathway (23) (Table 1). This seminal study, however, did not provide detailed measurement of the various isotopomers from which conclusive information regarding the operation of the 2 pathways could be obtained.

To obtain more detailed labeling information, we analyzed by NMR the positional isotopomers of glycine. Data were generated during steady-state ^{13}C -labeling experiments carried out with ^{13}C -enriched methanol and CO_2 at natural abundance, so that the labeling of free glycine—and glyoxylate—could be measured from the more abundant proteinogenic glycine. The 4 positional isotopomers of glycine were measured by combining 2D ZQF-TOCSY and 2D-HSQC experiments (25, 26) (Table S3 and Fig. S2). The results obtained for 3 biological replicates are shown in Table 1. The fractional enrichments (percentages of label in the carbon position)

A ICL⁻, Serine cycle methylotrophy



B ICL⁺, Serine cycle methylotrophy

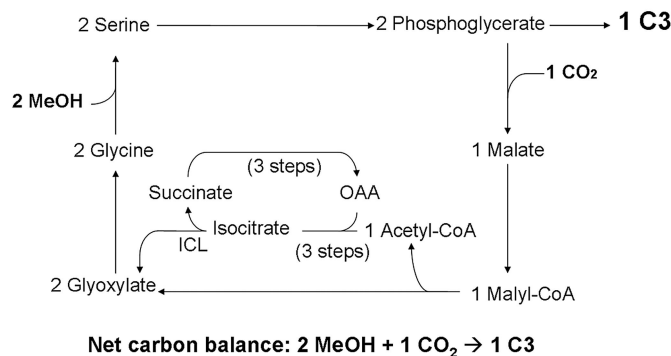
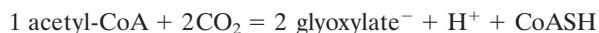


Fig. 7. Comparison of C1 assimilation pathways in isocitrate lyase-negative (ICL⁻; A), proposed from this study, and positive (ICL⁺; B) serine cycle methylotrophs. Note that CO₂ is derived from methanol, and therefore the overall carbon balance is the same in the 2 organisms: 3 methanol → 1 C₃.

This balance highlights the occurrence of 2 net carboxylation steps in the ethylmalonyl-CoA pathway, which is different not only from the GRC, where carbon loss would occur, but also from the classical glyoxylate cycle (10), where no carboxylation occurs.

The operation of the ethylmalonyl-CoA pathway has major implications for C1 assimilation in *M. extorquens* AM1 in terms of carbon balance and metabolic organization. The carbon balance of the whole process of C1 assimilation depends on the behavior of the succinyl-CoA molecule generated along with glyoxylate in the ethylmalonyl-CoA pathway. Erb et al. (23) proposed that succinyl-CoA was directly incorporated into cell material; however, the labeling data collected here for pure methylotrophic growth conditions indicate that succinyl-CoA is used to regenerate glyoxylate. Moreover, the positional isotopomers of glycine measured by NMR were consistent with a significant contribution of this process to glyoxylate regeneration, because the proportion of glyoxylate molecules regenerated from propionyl-CoA was calculated to be similar to that released by the cleavage of methylmalyl-CoA. These results indicate that not only 1 but 2 molecules of glyoxylate are regenerated by the ethylmalonyl-CoA pathway in *M. extorquens* AM1, with the following carbon balance:



The demonstration of the ethylmalonyl-CoA pathway closes the serine cycle during methylotrophic growth, a problem that has been unsolved since 1963 (9). The mechanism of C1 assimilation resulting from our observations (Fig. 6) shows that 2 molecules of glyoxylate can be regenerated at the same time. It also provides the molecular basis for the explanation of labeling data showing that roughly half the biomass carbon comes from CO₂ during methylotrophic growth conditions under laboratory conditions (6, 30). Therefore, the oxidation of methanol into CO₂ occurring in the initial steps of methanol utilization appears to be critical not only for energy purposes but also for carbon assimilation. A comparison of the carbon balance in ICL-negative and ICL-positive serine cycle methylotrophs (Fig. 7) reveals that the former organisms have a higher CO₂ fixation ability than the latter, although the energetic cost is likely to be higher as well. The high efficiency of carbon recovery during methylotrophic growth has major implications for the physiology of these widespread organisms and for their role in carbon cycling in their environment, and it poses questions as to what extent CO₂ is recycled from endogenously formed methanol or the atmosphere under natural conditions.

Materials and Methods

Reagents, Medium Composition, and Culture Conditions. [¹³C]methanol (99%) and [¹²C]methanol (99.9%) were purchased from Cambridge Isotope Laboratories; D₂O (99.8% and 99.97%) were purchased from Eurisotop. All other chemicals

were purchased from Sigma. Acetonitrile, formic acid, and ammonium used for HPLC solvents were of LC-MS degree. *M. extorquens* AM1 was grown on minimal medium containing 1.62 g/L NH₄Cl, 0.2 g/L MgSO₄, 2.21 g/L K₂HPO₄, 1.25 g/L NaH₂PO₄·2H₂O, and the following trace elements: 15 mg/L Na₂EDTA·2H₂O, 4.5 mg/L ZnSO₄·7H₂O, 0.3 mg/L CoCl₂·6H₂O, 1 mg/L MnCl₂·4H₂O, 1 mg/L H₃BO₃, 2.5 mg/L CaCl₂, 0.4 mg/L of Na₂MoO₄·2H₂O, 3 mg/L FeSO₄·7H₂O, and 0.3 mg/L CuSO₄·5H₂O. All batch cultures were carried out in a 500-mL bioreactor (Infors-HT) at 28 °C and at 1000 rpm. The pH was kept constant at 7.0 by addition of 1 M NH₄OH. For the purpose of short-term ¹³C-labeling experiments, cells were grown in 300 mL of medium containing 0.5% ¹³C-depleted methanol, and were aerated with compressed air at 0.15 L/min. Initial optical density (OD₆₀₀) was 0.2, and sampling was performed between ODs 2.8 and 3.0. Cultures carried out for the purpose of steady-state labeling experiments were grown in 400 mL of medium containing [¹³C]methanol and aerated with synthetic air containing 5% natural labeled CO₂. To keep the fraction of dissolved ¹³C-CO₂ produced from methanol below 2%, aeration rate was increased as follows: 0.2 L/min until OD 0.6, 0.4 L/min until OD 1, and 0.6 L/min until sampling. Initial OD was 0.01, and cells were harvested at around OD 1.5.

Sampling and Extraction of CoA Thioesters. A total of 1 mL of culture corresponding to 0.6 mg of cell dry weight was directly transferred into 4.5 mL of 95% acetonitrile at -20 °C containing 25 mM formic acid for quenching. To provide instantaneous quenching of metabolic activity, the sample was added into the quenching solution on a Vortex. Cells were disrupted by 3 sonication steps (30 s, 23 kHz) by using a Soniprep 150 device (Sanyo) and carried out in a cooling bath (T < -10 °C), with 30 s between each treatment. After the addition of 20 mL of ice-cold H₂O, the sample was chilled with liquid nitrogen. Frozen samples were stored at -20 °C until freeze drying. Subsequently, 300 μL of an ice-cold, 25 mM ammonium formate buffer (pH 3.5, 2% MeOH) was added. The suspension was centrifuged (14,000 × g, 2 min, -5 °C), and the supernatant was filtered through a Sartorius Minisart filter (pore size 0.2 μm) before analysis.

LC-MS Analysis. Analyses were performed with a Rheos 2200 HPLC system (Flux Instruments) coupled to an LTQ Orbitrap mass spectrometer (Thermo Fisher Scientific), equipped with an electrospray ionization probe. CoA esters were separated with a C₁₈ analytical column (Gemini 150 × 2.0 mm, particle size 3 μm; Phenomenex) at a flow rate of 220 μL min⁻¹. Injection volume was 10 μL. Solvent A was 50 mM formic acid adjusted to pH 8.1 with NH₄OH, and solvent B was methanol. The following gradient of B was applied: 0 min, 5%; 1 min, 5%; 10 min, 23%; 20 min, 80%; 22 min, 80%. MS analysis was done in the negative FTMS mode at a resolution of 15,000 (*m/z* = 400) to determine mass isotopomer distribution patterns and at a resolution of 60,000 (*m/z* = 400) to identify CoA thioesters and to detect potential mass peak overlapping problems. Sheath gas flow rate was 40, aux gas flow rate was 10, tube lens was -90 V, capillary voltage was -4 V, and ion spray voltage was -4.7 kV.

For the identification of CoA thioesters in cell extracts, chromatograms were analyzed for the presence of [M-H]⁻ ions corresponding to the exact mass expected from the 15 CoA thioesters potentially involved in glyoxylate regeneration, with a mass tolerance of 5 ppm. The number of carbon atoms was validated by using additional extracts from *M. extorquens* cells grown on [¹³C]methanol.

Short-Term ¹³C-Labeling Experiments. Incubation of cells with labeled acetate was realized in 2-mL Eppendorf tubes containing 50 μL of 1.05 M [1-¹³C]acetate.

To start, 1 mL of culture growing on [¹²C]methanol was quickly added with a syringe in the Eppendorf tube, and the sample was vortexed. A final acetate concentration of 50 mM was chosen to reach the same concentration as methanol. The acetate homogenization time in the final sample was found to be 1.1 ± 0.2 seconds. After various incubation times, the solution was added to the quenching solution, and the CoA thioesters were analyzed as explained above. The efficiency of the quenching process was evaluated by examining label incorporation into CoA thioesters in a culture sample injected in the quenching solution containing [¹⁻¹³C]acetate without an incubation period and revealing no incorporation of label.

Calculations of Normalized ¹³C-Label Incorporation. The incorporation of ¹³C label in each CoA ester during short-term ¹³C-labeling experiments was calculated from the analysis of the corresponding isotopic cluster in the mass spectra. The data were corrected for naturally occurring isotopes, the contribution of which was determined from the analysis of samples collected just before the addition of [¹⁻¹³C]acetate. Results are expressed as percent of ¹³C atoms incorporated in the molecule. Because the different CoA esters do not receive the same number of ¹³C atoms from [¹⁻¹³C]acetate, results were normalized according to the maximum number of ¹³C atoms that can be received from [¹⁻¹³C]acetate in the considered molecule ("normalized fractional labeling," L_{1-Ac}):

$$L_{1-Ac} = \frac{\left(\sum_{i=1}^n Mi \times i \right)}{n}$$

Where M_i is the proportion of the mass isotopomer corresponding to molecules having incorporated i ¹³C atoms from [¹⁻¹³C]acetate; n is the maximum number of [¹³C]carbon that can be incorporated into the molecule from [¹⁻¹³C]acetate (note that for some compounds, n is different according to GRC or EMCP).

NMR Analyses. All 1D and 2D NMR spectra were recorded on a Bruker Avance II 500-MHz spectrometer using a 5-mm z-gradient BBI probe head. The data were

acquired and processed by using TOPSPIN 1.3 software (Bruker). The temperature was 298 K. The 1D ¹H spectra were acquired by using a 30° pulse, 5,000-Hz sweep width, and 3.27-s acquisition times. A total of 128 scans were recorded, and relaxation delay between scans was 10 s. Proteinogenic amino acid sample was prepared as described previously (25) and modified as explained in the *SI Text*. Positional isotopomers of proteinogenic glycine were measured from (i) the analysis of carbon-carbon couplings in 2D-[¹H-¹³C]HSQC spectra and (ii) the analysis of heteronuclear ¹H,¹³C couplings in 2D ZQF-TOCSY spectra (25, 26). The peak deconvolution was realized with the software GOSA-fit (www.biol-log.biz/). The value of the glycine ²J_{H2C1} coupling constant was determined from the 1D ¹H analysis of 0.45 M [^{U-13}C, ¹⁵N]glycine performed with and without COOH decoupling.

Flux Calculations. For the purpose of flux calculations, a reaction network describing C1 metabolism in *M. extorquens* AM1, including simplified reactions for biomass formation, was designed. This model described the stoichiometry of the reactions as well as the transitions of carbon atoms. Biomass requirements were obtained from Van Dien and Lidstrom (31). Relative flux distributions were calculated from the positional isotopomers of glycine collected by NMR using both the 13C-Flux software developed by Wiechert (32). Results are expressed as molar fluxes relative to the rate of methanol uptake (set arbitrarily to 1.0).

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Supporting Information

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SI Text

Theoretical Fate of Carbon in the Central Metabolism of *Methylobacterium extorquens* AM1. The design of the steady-state ^{13}C -labeling experiment was done by simulation of the theoretical labeling of glycine obtained through the model of the central metabolic network (serine cycle, glyoxylate regeneration, and citric acid cycle) of *M. extorquens* AM1 from [^{13}C]methanol and CO_2 at natural abundance. Simplification of the discriminative process by which the glyoxylate regeneration pathway and the ethylmalonyl-CoA pathway differ is explained below and is illustrated in Fig. S1.

According to the glyoxylate regeneration cycle (1), 1 succinyl-CoA is generated from 2 acetyl-CoA molecules plus 1 CO_2 and contains predominantly 2 carbon atoms coming from methanol and 2 from CO_2 . The conversion of succinyl-CoA into malate and its cleavage into glyoxylate plus acetyl-CoA occurs via the formation of a symmetrical intermediate, namely succinate, which distributes the label equally between the C1 and C2 positions of glyoxylate. By operation of the GRC (1), the 2 major isotopomers are expected to be $^{12}\text{C}_1\text{-}^{13}\text{C}_2$ and $^{13}\text{C}_1\text{-}^{12}\text{C}_2$, and they should represent 41.1% and 33.3%, respectively, of all glyoxylate isotopomers (Table 1).

According to the ethylmalonyl-CoA pathway (2), glyoxylate is generated by cleavage of methylmalyl-CoA into propionyl-CoA and glyoxylate, but a second glyoxylate molecule can be generated potentially from succinyl-CoA. The glyoxylate directly generated by cleavage of methylmalyl-CoA is predominantly

singly labeled (expected isotopomer: $^{12}\text{C}_1\text{-}^{13}\text{C}_2$). The glyoxylate molecule generated from succinyl-CoA can have 2 different isotopic patterns—one ($^{12}\text{C}_1\text{-}^{13}\text{C}_2$) is singly labeled, and the other ($^{12}\text{C}_1\text{-}^{12}\text{C}_2$) is unlabeled. By operation of the EMCP, the 2 major isotopomers are expected to be $^{12}\text{C}_1\text{-}^{13}\text{C}_2$ and $^{12}\text{C}_1\text{-}^{12}\text{C}_2$, and their steady-state proportions should represent 57.1% and 39.1%, respectively, of all glyoxylate isotopomers.

Extraction of Proteinogenic Amino Acids. Extraction of proteinogenic amino acids was performed as described previously (3) for *Escherichia coli* and was adapted to *M. extorquens* AM1 as follows. The totality of a steady-state culture (400 mL) was centrifuged at $4,600 \times g$ and washed with 0.1 M NaCl solution and centrifuged. The cells were disrupted by 3 cycles of freezing (liquid N_2 during 10 s) and thawing (10°C during 2 min). The treated cells were resuspended in 20 mM Tris-HCl (pH 7.6), and disrupted again by bead-beating with 0.1-mm beads (zirconium/silice). After removal of beads by short centrifugation, the final extraction step was realized by sonification at 4°C : 3 times (30 s, 23 kHz) with breaks of 30 s. The cell debris was removed by ultracentrifugation ($33,000 \times g$, 30 min, 4°C), and the proteins were precipitated by ethanol (70% final) and centrifuged ($33,000 \times g$, 30 min, 4°C). The pellet was suspended in 6 M HCl and hydrolyzed for 14 h at 107°C . The acid was removed by evaporation, and labile protons were exchanged 3 times with deuterium by successive resuspensions in 2 mL of D_2O 99.8% followed by lyophilization. The sample was finally resuspended in 600 μL of D_2O 99.97%.

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Table S1. Summary of enzymes and their corresponding genes proposed for glyoxylate formation according to the GRC and the EMCP

No.	Reaction	Gene			Protein		Reaction	
	Name	Identified	Ref.	Essential	Identified	Ref.	Measured	Ref.
A	β -Ketothiolase	phaA*	(1)	–	–		+*	(1, 2)
B	Acetoacetyl-CoA reductase	phaB*	(1)	+	–		+*	(1, 2)
C	(R)3-hydroxybutyryl-CoA dehydratase	croR*	(3)	+	–		+*	(1, 3)
D	Crotonyl-CoA reductase	ccr*,†	(3, 4)	+	+†	(4)	+†	(4, 5)
E	Butyryl-CoA carboxylase	pccA*, pccB*	(3)	+	–		+*	(3)
F	Crotonyl-CoA reductase/carboxylase	ccr†	(3, 4)	+	+†	(4)	+†	(4, 5)
G	Ethylmalonyl-CoA epimerase	epm (epi)*,†	(6, 7)	+	+†	(7)	+†	(7)
H	Ethylmalonyl-CoA mutase	meaA (ecm)*,†	(6, 7)	+	+†	(7)	+†	(7)
I	Methylsuccinyl-CoA decarboxylase	–			–		–	
J	Isobutyryl-CoA dehydrogenase	ibd2*	(3)	+	–		–	
K	Metacrylyl-CoA hydratase	meaC*	(8)	+	–		–	
L	(2S)methylsuccinyl-CoA dehydrogenase	ibd2†	(2, 3, 8)	+	+†	(2)	–	
M	Mesaconyl-CoA hydratase	meaC*,†	(2, 8, 9)	+	+†	(2, 9)	+†	(9)
N	β -Methylmalyl-CoA lyase	mcl1*,†	(6, 10)	+	+*,†	(10, 11)	+*,†	(10, 11)
O	Propionyl-CoA carboxylase	pccA*, pccB*	(3)	+	–		+*	(3)
P	Methylmalonyl-CoA epimerase	epm (epi)*	(6, 7)	+	†	(7)	†	(7)
Q	Methylmalonyl-CoA mutase	mcmA*, mcmB*	(12)	+	+*,†	(2, 12)	+*	(12)

Letters indicating the different reactions are according to Fig. 1.

*Data from *M. extorquens* AM1.

†data from *R. sphaeroides*.

(Gene name) synonym in *R. sphaeroides*.

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Table S2. Identification of CoA thioesters by stable isotope assignment and spiking of standards using LC-HRMS

CoA ester	Formula	Identification by LC-HRMS					Occurrence in pathway			Detection in MeOH-grown cells
		[M0-H]-		[MU-H]-		Cn	Confirmation by spiking	GRC	EMCP	
		Theoretical mass	Observed mass	Theoretical mass	Observed mass					
Acetyl-	C ₂₃ H ₃₈ N ₇ O ₁₇ P ₃ S	808.1185	808.1182	831.1957	831.1946	23	+	+	+	+
Propionyl-	C ₂₄ H ₄₀ N ₇ O ₁₇ P ₃ S	822.1341	822.1350	846.2147	846.2127	24	+	+	+	+
Crotonyl-	C ₂₅ H ₄₀ N ₇ O ₁₇ P ₃ S	834.1341	834.1338	859.2180	859.2143	25	+	+	+	+
Methacrylyl-	C ₂₅ H ₄₀ N ₇ O ₁₇ P ₃ S	834.1341	834.1324	859.2158	n.d.	25	+	+	-	-
Butyryl-	C ₂₅ H ₄₂ N ₇ O ₁₇ P ₃ S	836.1498	836.1501	861.2337	861.2291	25	+	+	-	+
Isobutyryl-	C ₂₅ H ₄₂ N ₇ O ₁₇ P ₃ S	836.1498	836.1502	861.2337	n.d.	25	+	+	-	(+)*
Acetoacetyl-	C ₂₅ H ₄₀ N ₇ O ₁₈ P ₃ S	850.1291	850.1292	875.2129	n.d.	25	+	+	+	-
β-Hydroxybutyryl-	C ₂₅ H ₄₂ N ₇ O ₁₈ P ₃ S	852.1447	852.1451	877.2286	877.2266	25	+	+	+	+
β-Hydroxisobutyryl-	C ₂₅ H ₄₂ N ₇ O ₁₈ P ₃ S	852.1447	n.d.	877.2286	n.d.	25	-†	+	-	-
Succinyl-	C ₂₅ H ₄₀ N ₇ O ₁₉ P ₃ S	866.1240	866.1239	891.2078	891.2058	25	+	+	+	+
Methylmalonyl-	C ₂₅ H ₄₀ N ₇ O ₁₉ P ₃ S	866.1240	866.1246	891.2078	891.2049	25	+	+	+	+
Malyl-	C ₂₅ H ₄₀ N ₇ O ₂₀ P ₃ S	882.1189	882.1155	907.2028	907.2024	25	+	+	+	+
Mesaconyl-	C ₂₆ H ₄₀ N ₇ O ₁₉ P ₃ S	878.1240	878.1234	904.2112	904.2109	26	+	-	+	+
Ethylmalonyl-	C ₂₆ H ₄₂ N ₇ O ₁₉ P ₃ S	880.1396	880.1397	906.2269	906.2260	26	+	+	+	+
Methylsuccinyl-	C ₂₆ H ₄₂ N ₇ O ₁₉ P ₃ S	880.1396	880.1366	906.2269	906.2234	26	+	+	+	+
β-Methylmalyl-	C ₂₆ H ₄₂ N ₇ O ₂₀ P ₃ S	896.1334	896.1315	922.2218	922.2209	26	+	-	+	+

CoA thioesters proposed to be involved in the glyoxylate regeneration pathway of *M. extorquens* AM1 according to the GRC and the EMCP. Cn indicates number of carbon atoms in molecule; M0, monoisotopic mass peak of natural labeled compound; MU, uniformly ¹³C-labeled mass peak of ¹³C-labeled compound; n.d., not detected.

*Not always present.

†No standard available.

Table S3. 2D-NMR analysis of positional isotopomers of proteinogenic glycine extracted from *M. extorquens* AM1 cells grown on [¹³C]methanol plus 5% CO₂

NMR parameters of glycine in D₂O at pH 1.0

	δ H (ppm)	δ C (ppm)	Heteronuclear coupling constants	Homonuclear coupling constant
COO(D)	Not observable	Not measured	$J_{H_2C_1} = 5.96 \pm 0.04$	$J_{C_1C_2} = 59.40 \pm 0.26$
CH ₃	3.78	40.02	$J_{H_2C_2} = 145.26 \pm 0.11$	
TOCSY data collected on labeled proteinogenic glycine				
Resonance	Multiplet component	Isotopomer(s) at the origin of the multiplet component	Relative area	Value \pm SD
H2	Singlet	¹² C ¹ - ¹² C ²	I _s	32.3 \pm 0.7
	Doublet 1 ($J_{H_2C_2}$)	¹² C ¹ - ¹³ C ²	I _{d1}	60.1 \pm 2.9
	Doublet 2 ($J_{H_2C_1}$)	¹³ C ¹ - ¹² C ²	I _{d2}	2.0 \pm 1.0
	Doublet of doublet ($J_{H_2C_2} + J_{H_2C_1}$)	¹³ C ¹ - ¹³ C ²	I _{dd}	5.6 \pm 2.0
HSQC data collected on labeled proteinogenic glycine				
Resonance	Multiplet component	Isotopomer(s) at the origin of the multiplet component	Relative area	Value \pm SD
C2	Singlet	¹² C ¹ - ¹³ C ²	I _s	91.6 \pm 2.3
	Doublet ($J_{C_1C_2}$)	¹³ C ¹ - ¹³ C ²	I _d	8.4 \pm 2.3

TOCSY data: multiplet components of the glycine H2 resonance in 2D ZQF-TOCSY spectra, allowing the discrimination of 4 glycine isotopomers. HSQC data: multiplets observed in the C2 resonance of glycine in 2D ¹H-¹³C HSQC spectra, allowing the discrimination of 2 glycine isotopomers.