V16 The Double Description method:
Theoretical framework behind EFM and EP

Double Description Method Revisited

Komei Fukuda\(^1\) and Alain Prodon\(^2\)

\(^1\) Institute for Operations Research, ETHZ, CH-8092 Zürich, Switzerland
\(^2\) Department of Mathematics, EPFL, CH-1015 Lausanne, Switzerland

Double Description Method (1953)

All known algorithms for computing EMs are variants of the Double Description Method.

- derive simple & efficient algorithm for extreme ray enumeration, the so-called Double Description Method.

- show that it serves as a framework to the popular EM computation methods.

Analogy with Computer Graphics problem: How can one efficiently describe the space in a dark room that is lighted by a torch shining through the open door?
Duality of Matrices

Left: all points above the dividing line (the shaded area) fulfill the condition $x \geq 0$.
Middle: the points in the grey area fulfill the conditions $x_1 \geq 0$ and $x_2 \geq 0$.

But how could we describe the points in the grey area on the right side in a correspondingly simple manner?
Obviously, we could define a new coordinate system $(r_1, r_2)$ as a new set of generating vectors.
But we could also try to transform this area back into the grey area of the middle panel and use the old axes $x_1$ and $x_2$.

In 2D, this transformation can be obviously best performed by multiplying all vectors inside the grey area by a two-dimensional rotation matrix.
The Double Description Method

A pair \((A, R)\) of real matrices \(A\) and \(R\) is said to be a **double description pair** or simply a **DD pair** if the relationship

\[
A \mathbf{x} \geq 0 \text{ if and only if } \mathbf{x} = R \lambda \text{ for some } \lambda \geq 0
\]

holds. Clearly, for a pair \((A, R)\) to be a DD pair, the column size of \(A\) has to equal the row size of \(R\), say \(d\).

For such a pair, the set \(P(A)\) represented by \(A\) as

\[
P(A) = \{\mathbf{x} \in \mathbb{R}^d : A\mathbf{x} \geq 0\}
\]

is simultaneously represented by \(R\) as

\[
\{\mathbf{x} \in \mathbb{R}^d : \mathbf{x} = R\lambda \text{ for some } \lambda \geq 0\}
\]

A subset \(P\) of \(\mathbb{R}^d\) is called **polyhedral cone** if \(P = P(A)\) for some matrix \(A\), and \(A\) is called a **representation matrix** of the polyhedral cone \(P(A)\).

Then, we say \(R\) is a **generating matrix** for \(P\). Clearly, each column vector of a generating matrix \(R\) lies in the cone \(P\) and every vector in \(P\) is a nonnegative combination of some columns of \(R\).
The Double Description Method

**Theorem 1** (Minkowski’s Theorem for Polyhedral Cones)
For any $m \times n$ real matrix $A$, there exists some $d \times m$ real matrix $R$ such that $(A, R)$ is a DD pair, or in other words, the cone $P(A)$ is generated by $R$.

The theorem states that every polyhedral cone admits a generating matrix. The nontriviality comes from the fact that the row size of $R$ is finite. If we allow an infinite size, there is a trivial generating matrix consisting of all vectors in the cone.

Also the converse is true:

**Theorem 2** (Weyl’s Theorem for Polyhedral Cones)
For any $d \times n$ real matrix $R$, there exists some $m \times d$ real matrix $A$ such that $(A, R)$ is a DD pair, or in other words, the set generated by $R$ is the cone $P(A)$. 
The Double Description Method

**Task:** how does one construct a matrix $R$ from a given matrix $A$, and the converse?

These two problems are computationally equivalent. Farkas' Lemma shows that $(A,R)$ is a $DD$ pair if and only if $(R^T,A^T)$ is a $DD$ pair.

A more appropriate formulation of the problem is to require the minimality of $R$: find a matrix $R$ such that no proper submatrix is generating $P(A)$. A minimal set of generators is unique up to positive scaling when we assume the regularity condition that the cone is **pointed**, i.e. the origin is an extreme point of $P(A)$.

Geometrically, the columns of a minimal generating matrix are in 1-to-1 correspondence with the **extreme rays** of $P$.

Thus the problem is also known as the **extreme ray enumeration problem**.

No efficient (polynomial) algorithm is known for the general problem.
Double Description Method: primitive form

Suppose that the $m \times d$ matrix $A$ is given and let $P(A) = \{x : Ax \geq 0\}$
(This is equivalent to the situation at the beginning of constructing EPs or EFM: we only know $S$.)

The $DD$ method is an incremental algorithm to construct a $d \times m$ matrix $R$
such that $(A, R)$ is a $DD$ pair.

Let us assume for simplicity that the cone $P(A)$ is pointed.

Let $K$ be a subset of the row indices $\{1, 2, \ldots, m\}$ of $A$ and let $A_K$ denote the
submatrix of $A$ consisting of rows indexed by $K$.
Suppose we already found a generating matrix $R$ for $A_K$, or equivalently,
$(A_K, R)$ is a $DD$ pair. If $A = A_K$, we are done.

Otherwise we select any row index $i$ not in $K$ and try to construct a $DD$ pair
$(A_{K+i}, R')$ using the information of the $DD$ pair $(A_K, R)$.

Once this basic procedure is described, we have an algorithm to construct a
generating matrix $R$ for $P(A)$.  

Bioinformatics III  

16. Lecture WS 2014/15
Geometric version of iteration step

The procedure can be understood geometrically by looking at the cut-section $C$ of the cone $P(A_k)$ with some appropriate hyperplane $h$ in $\mathbb{R}^d$ which intersects with every extreme ray of $P(A_k)$ at a single point.

Let us assume that the cone is pointed and thus $C$ is bounded. Having a generating matrix $R$ means that all extreme rays (i.e. extreme points of the cut-section) of the cone are represented by columns of $R$. Such a cutsection is illustrated in the Fig.

Here, $C$ is the cube $abcdefgh$. 
Geometric version of iteration step

The newly introduced inequality \( A_i \cdot x \geq 0 \) partitions the space \( \mathbb{R}^d \) into three parts:

\[
\begin{align*}
H_i^+ &= \{ x \in \mathbb{R}^d : A_i \cdot x > 0 \} \\
H_i^0 &= \{ x \in \mathbb{R}^d : A_i \cdot x = 0 \} \\
H_i^- &= \{ x \in \mathbb{R}^d : A_i \cdot x < 0 \}
\end{align*}
\]

The intersection of \( H_i^0 \) with \( P \) and the new extreme points \( i \) and \( j \) in the cut-section \( C \) are shown in bold in the Fig.

Let \( J \) be the set of column indices of \( \mathbf{R} \). The rays \( r_j \ (j \in J) \) are then partitioned into three parts accordingly:

\[
\begin{align*}
J^+ &= \{ j \in J : r_j \in H_i^+ \} \\
J^0 &= \{ j \in J : r_j \in H_i^0 \} \\
J^- &= \{ j \in J : r_j \in H_i^- \}
\end{align*}
\]

We call the rays indexed by \( J^+ \), \( J^0 \), \( J^- \) the \textbf{positive}, \textbf{zero}, \textbf{negative} rays with respect to \( i \), respectively.

To construct a matrix \( \mathbf{R}' \) from \( \mathbf{R} \), we generate new \( |J^+| \times |J^-| \) rays lying on the \textit{ith} hyperplane \( H_i^0 \) by taking an appropriate positive combination of each positive ray \( r_j \) and each negative ray \( r_j' \), and by discarding all negative rays.
Geometric version of iteration step

The following lemma ensures that we have a $DD$ pair $(A_{K+i}, R')$, and provides the key procedure for the most primitive version of the $DD$ method.

**Lemma 3** Let $(A_K, R)$ be a $DD$ pair and let $i$ be a row index of $A$ not in $K$. Then the pair $(A_{K+i}, R')$ is a $DD$ pair, where $R'$ is the $d \times |J'|$ matrix with column vectors $r_j$ ($j \in J'$) defined by

$$J' = J^+ \cup J^0 \cup (J^+ \times J^-),$$

and

$$r_{jj'} = (A_i \cdot r_j) \cdot r_{j'} - (A_i \cdot r_{j'}) \cdot r_j$$

for each $(j,j') \in J^+ \times J^-$. 

**Proof** omitted.
Finding seed $DD$ pair

It is quite simple to find a $DD$ pair $(A_K, R)$ when $|K| = 1$, which can serve as the initial $DD$ pair.

Another simple (and perhaps the most efficient) way to obtain an initial $DD$ form of $P$ is by selecting a maximal submatrix $A_K$ of $A$ consisting of linearly independent rows of $A$.

The vectors $r_j$'s are obtained by solving the system of equations

$$A_K R = I$$

where $I$ is the identity matrix of size $|K|$, $R$ is a matrix of unknown column vectors $r_j$, $j \in J$.

As we have assumed $\text{rank}(A) = d$, i.e. $R = A_K^{-1}$, the pair $(A_K, R)$ is clearly a $DD$ pair, since $A_K \cdot x \geq 0 \iff x = A_K^{-1}\lambda$, $\lambda \geq 0$. 

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Primitive algorithm for DoubleDescriptionMethod

procedure DoubleDescriptionMethod(A);
begin
  Obtain any initial DD pair \((A_K, R)\);
  while \(K \neq \{1, 2, \ldots, m\}\) do
  begin
    Select any index \(i\) from \(\{1, 2, \ldots, m\} \setminus K\);
    Construct a DD pair \((A_{K+i}, R')\) from \((A_K, R)\);
    \(R := R';\quad K := K + i;\)
  end;
  Output \(R;\)
end.

This algorithm is very primitive, and the straightforward implementation will be quite useless, because the size of \(J\) increases extremely fast.

This is because many vectors \(r_{jj'}\) generated by the algorithm (defined in Lemma 3) are unnecessary. We need to avoid generating redundant vectors.

For this, we will use the zero set or active set \(Z(x)\) which is the set of inequality indices satisfied by \(x\) in \(P(A)\) with equality.

Noting \(A_i\), the \(i\)th row of \(A\), \(Z(x) = \{i : A_i \cdot x = 0\}\)
Towards the standard implementation

Two distinct extreme rays $r$ and $r'$ of $P$ are adjacent if the minimal face of $P$ containing both contains no other extreme rays.

**Proposition 7.** Let $r$ and $r'$ be distinct rays of $P$. Then the following statements are equivalent

(a) $r$ and $r'$ are adjacent extreme rays,
(b) $r$ and $r'$ are extreme rays and the rank of the matrix $A_{Z(r) \cap Z(r')}$ is $d - 2$,
(c) if $r''$ is a ray with $Z(r'') \supseteq Z(r) \cap Z(r')$ then either $r'' \simeq r$ or $r'' \simeq r'$.

**Lemma 8.** Let $(A_K, R)$ be a DD pair such that $\text{rank}(A_K) = d$ and let $i$ be a row index of $A$ not in $K$. Then the pair $(A_{K+i}, R')$ is a DD pair, where $R'$ is the $d \times |J'|$ matrix with column vectors $r_j$ ($j \in J'$) defined by

$J' = J^+ \cup J^0 \cup \text{Adj}$
$\text{Adj} = \{(j,j') \in J^+ \times J^- : r_j \text{ and } r_j' \text{ are adjacent in } P(A_K)\}$
$r = (A_i r_j) r_j' - (A_i r_{j'}) r_j$ for each $(j,j') \in \text{Adj}$.

Furthermore, if $R$ is a minimal generating matrix for $P(A_K)$ then $R'$ is a minimal generating matrix for $P(A_{K+i})$. 
Algorithm for standard form of double description method

Hence we can write a straightforward variation of the DD method which produces a minimal generating set for $P$:

```
procedure DDMethodStandard(A) begin
    Obtain any initial DD pair $(A_K, R);$ such that $R$ is minimal
    while $K \neq \{1, 2, \ldots, m\}$ do
        begin
            Select any index $i$ from $\{1, 2, \ldots, m\} \setminus K$;
            Construct a DD pair $(A_{K+i}, R')$ from $(A_K, R);$ /* by using Lemma 8 */
            $R := R'$; $K := K + i$;
        end;
    Output $R$;
end.
```

To implement $\text{DDMethodStandard}$, we must check for each pair of extreme rays $r$ and $r'$ of $P(A_K)$ with $A_i r > 0$ and $A_i r' < 0$ whether they are adjacent in $P(A_K)$. 

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16. Lecture WS 2014/15
Tools for analyzing network states

The two steps that are used to form a solution space — reconstruction and the imposition of governing constraints — are illustrated in the centre of the figure. Several methods are being developed at various laboratories to analyse the solution space. $C_i$ and $C_j$ concentrations of compounds $i$ and $j$; EP, extreme pathway; $v_i$ and $v_j$ fluxes through reactions $i$ and $j$; $v_1 - v_3$ flux through reactions 1-3; $v_{\text{net}}$, net flux through loop.

Application of elementary modes

Metabolic network structure of *E. coli* determines key aspects of functionality and regulation

Compute EFMs for central metabolism of *E. coli*.

Catabolic part: substrate uptake reactions, glycolysis, pentose phosphate pathway, TCA cycle, excretion of by-products (acetate, formate, lactate, ethanol)

Anabolic part: conversions of precursors into building blocks like amino acids, to macromolecules, and to biomass.

**Metabolic network topology and phenotype**

**Idea:**
Can the total number of EFMs for given conditions be used as quantitative measure of metabolic flexibility?

**a.** Relative number of EFMs $N$ enabling deletion mutants of gene $i$ ($\Delta i$) in *E. coli* to grow (abbreviated by $\mu$) for 90 different combinations of mutation and carbon source.

Shown are results for 90 deletions of different individual genes.

Answer: Yes, the # of EFMs for mutant strain allows correct prediction of growth phenotype in more than 90% of the cases.

Robustness analysis

The # of EFMs qualitatively indicates whether a mutant is viable or not, but does not describe quantitatively how well a mutant grows.

Define maximal biomass yield $Y_{\text{mass}}$ as the optimum of:

$$Y_{i,X/S_i} = \frac{e_i^\mu}{e_i^{S_k}}$$

$e_i$ is the single reaction rate (growth and substrate uptake) in EFM $i$ selected for utilization of substrate $S_k$.

Robustness Analysis

Dependency of the mutants' maximal growth yield $Y_{\text{max}}(\Delta i)$ (open circles) and the network diameter $D(\Delta i)$ (open squares) on the share of elementary modes operational in the mutants.


Central metabolism of *E.coli* behaves in a highly robust manner because mutants with significantly reduced metabolic flexibility show a growth yield similar to wild type.
Current metabolomics

Review:
(1) recent work on metabolic networks required revising the picture of separate biochemical pathways into a **densely-woven metabolic network**

(2) **Connectivity** of substrates in this network follows a **power-law** (Yeong&Barabasi).

(3) **Constraint-based modeling approaches** (FBA) were successful in analyzing the **capabilities** of cellular metabolism including
   - its capacity to predict **deletion phenotypes**
   - the ability to calculate the **relative flux values** of metabolic reactions, and
   - the capability to identify properties of **alternate optimal growth states**
   in a wide range of simulated environmental conditions

Open questions
- what parts of metabolism are involved in adaptation to environmental conditions?
- is there a central essential metabolic core?
- what role does transcriptional regulation play?
Distribution of fluxes in *E. coli*

**Global organization of metabolic fluxes in the bacterium* Escherichia coli***

E. Almaas¹, B. Kovács¹, T. Vicsek², Z. N. Oltvai³ & A.-L. Barabási¹

¹Department of Physics, University of Notre Dame, Notre Dame, Indiana 46556, USA
²Biological Physics Department and Research Group of HAS, Eötvös University, H-1117 Budapest, Hungary
³Department of Pathology, Northwestern University, Chicago, Illinois 60611, USA


Stoichiometric matrix for *E. coli* strain MG1655 containing 537 metabolites and 739 reactions taken from Palsson *et al.*

Apply flux balance analysis to characterize solution space (all possible flux states under a given condition).

\[
\frac{d}{dt}[A_i] = \sum_j S_{ij} v_j = 0
\]

\(v_j\) is the flux of reaction \(j\) and \(S_{ij}\) is the stoichiometric coefficient of reaction \(j\).
Optimal states

Denote the mass carried by reaction \( j \) producing (consuming) metabolite \( i \) by

\[
\hat{v}_{ij} = |S_{ij}| v_j
\]

Observation:

**Fluxes vary widely**: e.g. dimensionless flux of succinyl coenzyme A synthetase reaction is 0.185, whereas the flux of the aspartate oxidase reaction is 10.000 times smaller, \( 2.2 \times 10^{-5} \).

Using linear programming and adapting constraints for each reaction flux \( v_i \) of the form \( \beta_i^{min} \leq v_i \leq \beta_i^{max} \), the flux states were calculated that optimize cell growth on various substrates.

Plot the flux distribution for active (non-zero flux) reactions of \( E.coli \) grown in a glutamate- or succinate-rich substrate.
Overall flux organization of *E.coli* metabolic network

**a.** Flux distribution for optimized biomass production on succinate (black) and glutamate (red) substrates.

The solid line corresponds to the power-law fit that a reaction has flux $\nu$

$$P(\nu) \propto (\nu + \nu_0)^{-\alpha}, \text{ with } \nu_0 = 0.0003 \text{ and } \alpha = 1.5.$$ 

**d.** The distribution of experimentally determined fluxes from the central metabolism of *E. coli* shows power-law behaviour as well, with a best fit to

$$P(\nu) \propto \nu^{-\alpha} \text{ with } \alpha = 1.$$ 

Both computed and experimental flux distribution show wide spectrum of fluxes.


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Response to different environmental conditions

Is the flux distribution independent of environmental conditions?

b, Flux distribution for optimized biomass on succinate substrate (black) with an additional 10% (red), 50% (green) and 80% (blue) randomly chosen subsets of the 96 input channels (substrates) turned on.

The flux distribution was averaged over 5,000 independent random choices of uptake metabolites.

→ Yes, the flux distribution is independent of the external conditions.

Use scaling behavior to determine local connectivity

The observed flux distribution is compatible with two different potential local flux structures:
(a) a **homogenous local organization** would imply that all reactions producing (consuming) a given metabolite have comparable fluxes
(b) a more delocalized „**high-flux backbone (HFB)“** is expected if the local flux organisation is heterogenous such that each metabolite has a dominant source (consuming) reaction.

![Diagram](image)

\[
Y(k, i) = \sum_{j=1}^{k} \left[ \frac{\hat{v}_{ij}}{\sum_{l=1}^{k} \hat{v}_{il}} \right]^2
\]

Characterizing the local inhomogeneity of the flux net

\( kY(k) \) shown as a function of \( k \) for incoming and outgoing reactions, averaged over all metabolites, indicates that \( k \times Y(k) \propto k^{0.73} \).

Inset shows non-zero mass flows, \( v_{ij}^\wedge \), producing (consuming) FAD on a glutamate-rich substrate.

\( \rightarrow \) an intermediate behavior is found between the two extreme cases.

\( \rightarrow \) the large-scale inhomogeneity observed in the overall flux distribution is also increasingly valid at the level of the individual metabolites.

The more reactions that consume (produce) a given metabolite, the more likely it is that a single reaction carries most of the flux, see FAD.

Clean up metabolic network

Use simple algorithm that removes for each metabolite systematically all reactions but the one providing the largest incoming (outgoing) flux distribution.

The algorithm uncovers the „high-flux-backbone“ of the metabolism, a distinct structure of linked reactions that form a giant component with a star-like topology.
Maximal flow networks

glutamate rich

Directed links: Two metabolites (e.g. A and B) are connected with a directed link pointing from A to B only if the reaction with maximal flux consuming A is the reaction with maximal flux producing B.

Shown are all metabolites that have at least one neighbour after completing this procedure. The background colours denote different known biochemical pathways.

succinate rich substrates
FBA-optimized network on glutamate-rich substrate

High-flux backbone for FBA-optimized metabolic network of \textit{E. coli} on a glutamate-rich substrate. Metabolites (vertices) coloured blue have at least one neighbour in common in glutamate- and succinate-rich substrates, and those coloured red have none. Reactions (lines) are coloured blue if they are identical in glutamate- and succinate-rich substrates, green if a different reaction connects the same neighbour pair, and red if this is a new neighbour pair. Black dotted lines indicate where the disconnected pathways, for example, folate biosynthesis, would connect to the cluster through a link that is not part of the HFB. Thus, the red nodes and links highlight the predicted changes in the HFB when shifting \textit{E. coli} from glutamate- to succinate-rich media. Dashed lines indicate links to the biomass growth reaction.

(1) Pentose Phospate  
(2) Purine Biosynthesis  
(3) Aromatic Amino Acids  
(4) Folate Biosynthesis  
(5) Serine Biosynthesis  
(6) Cysteine Biosynthesis  
(7) Riboflavin Biosynthesis  
(8) Vitamin B6 Biosynthesis  
(9) Coenzyme A Biosynthesis  
(10) TCA Cycle  
(11) Respiration  
(12) Glutamate Biosynthesis  
(13) NAD Biosynthesis  
(14) Threonine, Lysine and Methionine Biosynthesis  
(15) Branched Chain Amino Acid Biosynthesis  
(16) Spermidine Biosynthesis  
(17) Salvage Pathways  
(18) Murein Biosynthesis  
(19) Cell Envelope Biosynthesis  
(20) Histidine Biosynthesis  
(21) Pyrimidine Biosynthesis  
(22) Membrane Lipid Biosynthesis  
(23) Arginine Biosynthesis  
(24) Pyruvate Metabolism  
(25) Glycolysis

**Interpretation**

Only a few pathways appear disconnected indicating that although these pathways are part of the HFB, their end product is only the second-most important source for another HFB metabolite.

Groups of individual HFB reactions largely overlap with traditional biochemical partitioning of cellular metabolism.

How sensitive is the HFB to changes in the environment?

**b.** Fluxes of individual reactions for glutamate-rich and succinate-rich conditions. Reactions with negligible flux changes follow the diagonal (solid line). Some reactions are turned off in only one of the conditions (shown close to the coordinate axes). Reactions belonging to the HFB are indicated by black squares, the rest are indicated by blue dots. Reactions in which the direction of the flux is reversed are coloured green.

Only reactions in the high-flux territory undergo noticeable differences!

**Type I:** reactions turned on in one conditions and off in the other (symbols).

**Type II:** reactions remain active but show an orders-in-magnitude shift in flux under the two different growth conditions.

Flux distributions for individual reactions

Shown is the flux distribution for four selected *E. coli* reactions in a 50% random environment.

a Triosphosphate isomerase;

b carbon dioxide transport;

c NAD kinase;

d guanosine kinase.

Reactions on the $\sigma \propto v$ curve (small fluxes) have **unimodal/gaussian distributions** (a and c). Shifts in growth-conditions only lead to small changes of their flux values.

Reactions off this curve have **multimodal distributions** (b and d), showing several discrete flux values under diverse conditions. Under different growth conditions they show several discrete and distinct flux values.

Summary

Metabolic network use is highly uneven (power-law distribution) at the global level and at the level of the individual metabolites.

Whereas most metabolic reactions have low fluxes, the overall activity of the metabolism is dominated by several reactions with very high fluxes.

*E. coli* responds to changes in growth conditions by reorganizing the rates of selected fluxes predominantly within this high-flux backbone. Apart from minor changes, the use of the other pathways remains unaltered. These reorganizations result in large, discrete changes in the fluxes of the HFB reactions.
The same authors as before used Flux Balance Analysis to examine utilization and relative flux rate of each metabolite in a wide range of simulated environmental conditions for *E. coli*, *H. pylori* and *S. cerevisae*:

For each system they considered 30,000 randomly chosen combinations where each uptake reaction is assigned a random value between 0 and 20 mmol/g/h.

→ adaptation to different conditions occurs by 2 mechanisms:

(a) flux plasticity: changes in the fluxes of already active reactions.

E.g. changing from glucose- to succinate-rich conditions alters the flux of 264 *E. coli* reactions by more than 20%

(b) less commonly, adaptation includes structural plasticity, turning on previously zero-flux reactions or switching off active pathways.
The two adaptation method mechanisms allow for the possibility of a group of reactions not subject to structural plasticity being active under all environmental conditions.

Assume that active reactions were randomly distributed.

If typically a $q$ fraction of the metabolic reactions are active under a specific growth condition, we expect for $n$ distinct conditions an overlap of at least $q^n$ reactions. This converges quickly to 0.
Emergence of the Metabolic Core

(a–c) The average relative size of the number of reactions that are always active as a function of the number of sampled conditions (black line).

(d and e) The number of metabolic reactions (d) and the number of metabolic core reactions (e) in the three studied organisms.

In a–c, as the number of conditions increases, the curve converges to a constant enoted by the dashed line, identifying the metabolic core of an organism.

Red line: number of reactions that are always active if activity is randomly distributed in the metabolic network. The fact that it converges to zero indicates that the real core represents a collective network effect, forcing a group of reactions to be active in all conditions.
Shown are all reactions that are found to be active in each of the 30,000 investigated external conditions.

**Blue**: Metabolites that contribute directly to biomass formation,

**Red (green)**: core reactions (links) catalyzed by essential (or nonessential) enzymes.

Black-colored links: enzymes with unknown deletion phenotype.

**Blue dashed lines**: multiple appearances of a metabolite,

Links with arrows: unidirectional reactions.

Note that 20 out of the 51 metabolites necessary for biomass synthesis are not present in the core, indicating that they are produced (or consumed) in a growth-condition-specific manner.

**Blue and brown shading**: folate and peptidoglycan biosynthesis pathways

White numbered arrows denote current antibiotic targets inhibited by: (1) sulfonamides, (2) trimethoprim, (3) cycloserine, and (4) fosfomycin.

A few reactions appear disconnected since we have omitted the drawing of cofactors.
The metabolic cores contain 2 types of reactions:

(a) reactions that are essential for biomass production under all environment conditions (81 of 90 in E.coli)

(b) reactions that assure optimal metabolic performance.
Characterizing the Metabolic Cores

(A) The number of overlapping metabolic reactions in the metabolic core of *H. pylori*, *E. coli*, and *S. cerevisiae*. The metabolic cores of simple organisms (*H. pylori* and *E. coli*) overlap to a large extent. The largest organism (*S. cerevisiae*) has a much larger reaction network that allows more flexibility → the relative size of the metabolic core is much lower.

(B) The fraction of metabolic reactions catalyzed by essential enzymes in the cores (black) and outside the core in *E. coli* and *S. cerevisiae*. → Reactions of the metabolic core are mostly essential ones.

(C) One could assume that the core represents a subset of high-flux reactions. This is apparently not the case. The distributions of average metabolic fluxes for the core and the noncore reactions in *E. coli* are very similar.
Summary

- Adaptation to environmental conditions occurs via structural plasticity and/or flux plasticity. Here: identification of a surprisingly stable **metabolic core** of reactions that are tightly connected to each other.

- The reactions belonging to this core represent **potential targets** for antimicrobial intervention.