

1.

What are the major technological advances inspiring the recent boom in Synthetic Biology?

- ☐ Multiplexed and high-throughput Polymerase Chain Reaction (m-PCR/ht-PCR)
- ☐ Autonomous Laboratory Workflows (ALW) and robotics
- ☐ Matrix-Assisted Laser Desorption and Ionization and Electrospray Ionisation (MALDI/ESI)
- ☒ High-performance DNA sequencing and DNA synthesis
- ☐ Homologous recombination and bacterial transformation

2. What

kind of activity or group does the abbreviation *iGEM* refer to in Synthetic Biology?

- ☐ An international non-governmental watchdog organization supervising the ethical conduct of genetic modifications
- ☐ A government task-force, established in the UK, to develop industrial guidelines for engineering microbiology
- ☒ An international competition in which student teams produce genetically engineered machines
- ☐ A lobbying group supporting the introduction of genetically modified organisms to the European market
- ☐ The Institute for Genetic Engineering and Manipulation at Stanford University

3. What does the abbreviation *DBT* stand for in Synthetic Biology?

- ☐ Develop-Balance-Tinker
(the three core concepts underlying the genetic engineering of complex biological systems by Synthetic Biology)
- ☒ Design-Build-Test
(an engineering approach for the iterative improvement of engineered biological systems)
- ☐ Data-driven
Biological Technologies (the combination of computational analysis and post-genomic molecular profiling techniques, such as metabolomics)
- ☐ Directed Botanical Therapeutics
(the production of plant-based drugs in microbial systems)
- ☐ Dicyclobenzotoluene (a key chemical required for high-throughput genome engineering)

4. Why are untargeted metabolomics based experiments usually considered to take a longer time than similar targeted analyses?

- ☐ Highly complex data produced
- ☐ Longer Chromatography Runs
- ☐ Time taken to Identify Unknown Metabolites
- ☒ All of the above

5. Within the TEST platform, spectroscopic based analyses (aside from NMR) are considered useful as:

- ☐ Data can be directly overlaid on to species specific biochemical maps
- ☐ They generate rich multi-dimensional data
- ☒ They are quick and non-destructive
- ☐ Can identify and quantify a large proportion of all chemical species within a sample

6. What are the two main approaches to producing designer genomes?

- ☐ Genome transplantation and mutagenesis
- ☒ Genome editing /reduction and genome synthesis
- ☐ DNA synthesis (writing) and sequencing (reading)
- ☐ Gene knock-outs and knock-ins
- ☐ Genome modelling and synthesis

7. Responsible research and innovation should seek to continuously pursue which of the following steps?

- ☐ Anticipate
economic, social and environmental impacts, intended or otherwise
- ☐ Reflect
on purposes, motivations, potential implications, uncertainties, ignorance, assumptions, questions, and dilemmas
- ☐ Engage
on opening up visions, impacts, and questioning to broader deliberation, dialogue, engagement and debate in an inclusive way
- ☐ Act
– using these processes to influence the direction and trajectory of the research and innovation process itself
- ☒ All
of the above

8. What factor(s) must be considered when designing a microbial cell factory?

- ☐ Genetic differences between the consumers of cell factory products
- ☒ The design of engineered synthetic pathways
- ☒ The natural metabolism of the host organism
- ☒ The feedstock on which the cell factory will grow

9. Constraint-based metabolic modelling can be used to predict...?

- ☒ Metabolic flux through a network
- ☐ The most valuable chemical
- ☐ An average temperature at which to run an experiment

10. Optimisation of the ribosome binding site (RBS) sequence has an affect on:

- ☒ translation
- ☐ neither transcription or translation
- ☐ transcription

11. Why might we wish to control the timing of gene expression in a pathway?

- ☐ To change the product being produced
- ☒ To reduce the metabolic burden on the cell
- ☒ To reduce accumulation of toxic intermediates