

A systematic framework for gene–transcript–protein–reaction associations of human metabolism

김예지, 김현욱, 류재용, 이상엽[†]
한국과학기술원

Alternative splicing plays a significant role in producing different transcripts from a gene, and therefore resulting in various protein isoforms. Here, we show a systematic framework for the production of gene–transcript–protein–reaction associations (GeTPRA) of the human metabolism. This framework generated 11,415 GeTPRAs for 1,106 metabolic genes. The generated GeTPRAs were further evaluated, using a updated human genome–scale metabolic model (GEM), which is biochemically consistent and transcript–level data compatible. Both principal and nonprincipal transcripts of metabolic genes were considered in the framework based on 446 personal RNA–Seq data and 1,784 personal GEMs. The framework and the GeTPRAs are expected to provide better understanding of human metabolism and enable medical applications.