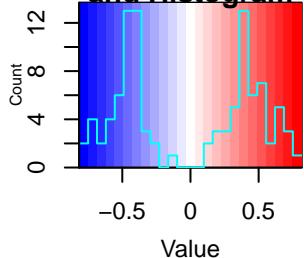
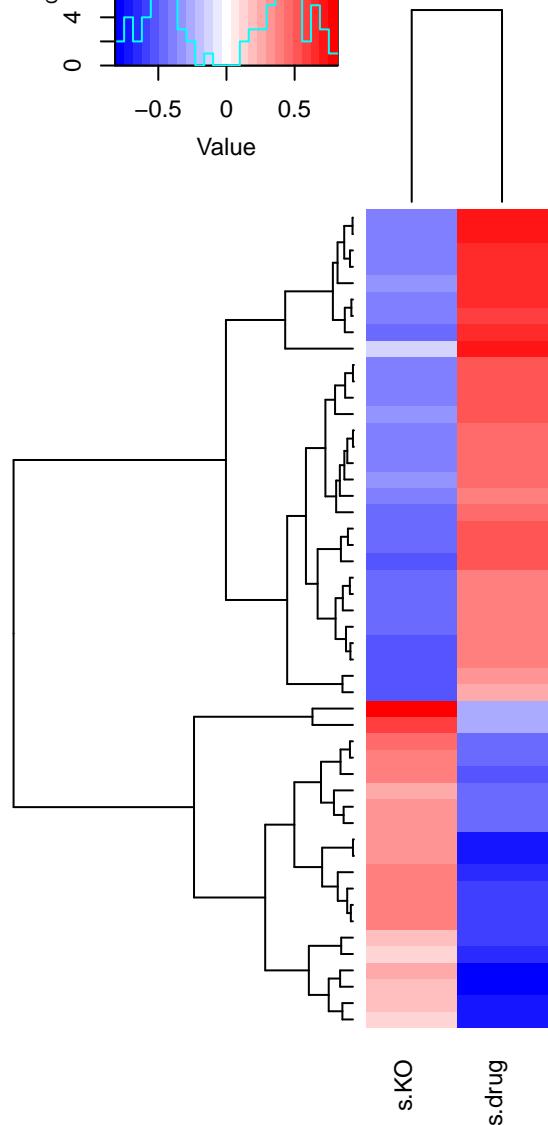


# Color Key and Histogram



# ked Reactomes



Viral mRNA Translation  
 Peptide chain elongation  
 Eukaryotic Translation Termination  
 Eukaryotic Translation Elongation  
 Selenocysteine synthesis  
 Response of EIF2AK4 (GCN2) to amino acid deficiency  
 Nonsense Mediated Decay (NMD) independent of the Exon Junction Complex (EJC)  
 Formation of a pool of free 40S subunits  
 Complex I biogenesis  
 Eukaryotic Translation Initiation  
 Cap-dependent Translation Initiation  
 SRP-dependent cotranslational protein targeting to membrane  
 Selenoamino acid metabolism  
 Nonsense-Mediated Decay (NMD)  
 Nonsense Mediated Decay (NMD) enhanced by the Exon Junction Complex (EJC)  
 rRNA processing  
 Regulation of expression of SLITs and ROBOs  
 Influenza Viral RNA Transcription and Replication  
 Mitochondrial iron-sulfur cluster biogenesis  
 GTP hydrolysis and joining of the 60S ribosomal subunit  
 L13a-mediated translational silencing of Ceruloplasmin expression  
 Formation of the ternary complex, and subsequently, the 43S complex  
 Major pathway of rRNA processing in the nucleolus and cytosol  
 rRNA processing in the nucleus and cytosol  
 ATF4 activates genes in response to endoplasmic reticulum stress  
 Ribosomal scanning and start codon recognition  
 Activation of the mRNA upon binding of the cap-binding complex and eIFs, and subsequent binding to 43S  
 Translation initiation complex formation  
 SLBP independent Processing of Histone Pre-mRNAs  
 SLBP Dependent Processing of Replication-Dependent Histone Pre-mRNAs  
 Mucopolysaccharidoses  
 Biotin transport and metabolism  
 HS-GAG degradation  
 The activation of arylsulfatases  
 Reduction of cytosolic Ca<sup>++</sup> levels  
 Diseases associated with glycosaminoglycan metabolism  
 Chondroitin sulfate biosynthesis  
 A tetrasaccharide linker sequence is required for GAG synthesis  
 Defective EXT2 causes exostoses 2  
 Defective EXT1 causes exostoses 1, TRPS2 and CHDS  
 Defective B4GALT7 causes EDS, progeroid type  
 Other semaphorin interactions  
 Defective B3GALT6 causes EDSP2 and SEMDJL1  
 Defective B3GAT3 causes JDSSDHD  
 CREB1 phosphorylation through the activation of Adenylate Cyclase  
 PKA activation  
 Laminin interactions  
 Syndecan interactions  
 Non-integrin membrane-ECM interactions  
 MET activates PTK2 signaling