Column I	Column2	Column3	Column4	Column5			
Function	Library	databaseUsed	Input (demonstrated examples)	Sample Configuration			
Enrich functions							
enricher	clusterProfiler	msigdbr/wiki- download/others	vectorOfGenes	enricher(gene, TERM2GENE = wpid2gene, TERM2NAME = wpid2name)			
enrichKEGG	clusterProfiler	organism='hsa'	vectorOfGenes	enrichKEGG(gene=gene,organism='hsa',pvalueCutoff=0.05)			
enrichMKEGG	clusterProfiler	organism='hsa'	vectorOfGenes	enrichMKEGG(gene = gene,organism = 'hsa')			
	clusterProfiler	OrgDb=org.Hs.eg.db	vectorOfGenes	enrichGO(gene=gene,universe=names(geneList),OrgDb=org .Hs.eg.db,ont="CC",pAdjustMethod="BH",pvalueCutoff=0.0 I,qvalueCutoff=0.05,readable=TRUE)			
enrichPathway	ReactomePA	automatic	vectorOfGenes				
enrichMeSH	ReactomePA	MeSHDb="Mesh.Hsa. eg.db"	vectorOfGenes	enrichMeSH(de, MeSHDb = **"MeSH.Hsa.eg.db"**, database='gendoo', category = 'C')			
enrichDO	DOSE	ont="DO'	vectorOfGenes	enrichDO(gene=gene,ont="DO",pvalueCutoff=0.05,pAdjust Method="BH",universe=names(geneList),minGSSize=5,max GSSize=500,qvalueCutoff=0.05,readable=FALSE)			
enrichNCG	DOSE	automatic	vectorOfGenes				
enrichDGN	DOSE	automatic	vectorOfGenes	enrichDGN(gene)			
enrichDGNv	DOSE	automatic	vectorOfGenes	enrichDGNv(snp)			
groupGO	clusterProfiler	OrgDb=org.Hs.eg.db	vectorOfGenes	groupGO(gene= gene,OrgDb= org.Hs.eg.db,ont= "CC",level= 3,universe= names(geneList),readable = TRUE)			
GSEA functions							
GSEA	clusterProfiler	msigdbr/wiki- download/others	NamedVectorOfFoldchange	GSEA(geneList, TERM2GENE = wpid2gene, TERM2NAME = wpid2name, verbose=FALSE)			
gseKEGG	clusterProfiler	organism='hsa'	NamedVectorOfFoldchange	gseKEGG(geneList=geneList,organism='mcc',nPerm=1000, minGSSize=120,pvalueCutoff=0.05,verbose=FALSE)			
gseMKEGG	clusterProfiler	organism='hsa'	NamedVectorOfFoldchange				
gseGO	clusterProfiler	OrgDb=org.Hs.eg.db	NamedVectorOfFoldchange	gseGO(geneList=geneList,OrgDb=org.Hs.eg.db,ont="CC",n Perm=1000,minGSSize=100,maxGSSize=500,pvalueCutoff =0.05,verbose=FALSE)			
gsePathway	ReactomePA	?	NamedVectorOfFoldchange				
	ReactomePA	MeSHDb="Mesh.Hsa. eg.db"	NamedVectorOfFoldchange	gseMeSH(geneList, MeSHDb = "MeSH.Hsa.eg.db", database = 'gene2pubmed', category = "G")			
gseDO	DOSE	ont="DO'	NamedVectorOfFoldchange	gseDO(geneList,nPerm=100,minGSSize=120,pvalueCutoff= 0.2,pAdjustMethod="BH",verbose=FALSE)			

gseNCG	DOSE	automatic	NamedVectorOfFoldchange	gseNCG(geneList,nPerm=100,minGSSize=120,pvalueCutoff =0.2,pAdjustMethod="BH",verbose=FALSE)			
gseDGN	DOSE	automatic	NamedVectorOfFoldchange	gseDGN(geneList,nPerm=100,minGSSize=120,pvalueCutoff			
8002 011		uutoiiiutit		=0.2,pAdjustMethod="BH",verbose=FALSE)			
			Database & ID conversi	on			
get .gmt files	msigdbr	no	species & category	msigdbr(species = "Homo sapiens", category = "C2") %>%			
				dplyr::select(gs_name, gene_symbol) or			
				read.gmt("./wikipathways-20200510-gmt-			
				Homo_sapiens.gmt")			
				bitr(wpid2gene\$gene,fromType="ENTREZID",toType=c("E			
bitr	clusterProfiler	no	avector of gene ids				
NSEMBL","SYMBOL"),OrgDb=org.Hs.eg.db) Visulization							
barplot	graphics	no	enrichResult	barplot(enricherRes, showCategory=20)			
dotplot	enrichplot	no	enrichRseult, gseaResult &	dotplot(edo2, showCategory=30) + ggtitle("dotplot for			
чосьюе	emiempioe		compareClusterResult	GSEA") && plot_grid(p1, p2, ncol=2)			
				options(repr.plot.width=16, repr.plot.height=24) &&			
cnetplot	enrichplot	no	enrichResult	<pre>cnetplot(edox, foldChange=geneList, showCategory =</pre>			
				I 0,colorEdge=TRUE,circular=TRUE, node_label="all")			
upsetplot	enrichplot	no	enrichRseult and gseaResult	upsetplot(kk2)			
ridgeplot	enrichplot	no	gseaResult	ridgeplot(edo2)			
	enrichplot	no	gseaResult				
gseaplot				gseaplot2(edo2, geneSetID = 2, title = edo2\$Description[2])			
Soombroo				# geneSetID are the row number in the edo2 object			
lultiple category							
idicipie edeego.	/			formula_res <- compareCluster(Entrez~group+othergroup,			
ompareCluste	clusterProfiler	automatic	list of gene sets/ genes with different factor labels	data=mydf, fun="enrichKEGG"); ck <-			
				• •			
				compareCluster(geneCluster = gcSample, fun =			
				"enrichKEGG")			
emapplot	enrichplot	no	compareClusterResult	options(repr.plot.width=8, repr.plot.height=6) &&			
				emapplot(xx,pie="count", pie_scale=2,			
				layout="kk",legend_n=4)ridgeplot(edo2) # show the fold			
				change of core genes in the GSEA analysis.			
	ı .			1 2			