At the genomics level, we use the microarray technology to produce gene expression data for various organisms under many conditions. A desirable advancement is the need to extract from this gene expression data, information that will be useful toward given answers to the questions targeted at the design of the microarray experiment. Analysis of gene expression data of P.f when induced with two anti-malaria drugs (such as chloroquine and choline analogue T4) has shown that the P.f resistance mechanisms may not be elucidate-able at the genomics level. But on a proteomic level, biochemical research has elucidated an increasingly complete image of the metabolic architecture of organisms that included that of P.f. In this work, we sort to use the biochemical network of P.f to deduce its drugs resistance mechanism(s) using the two gene expression data obtained when P.f is treated with chloroquine and choline analogue T4. We do this by mapping these gene expression data onto the enzymatic reaction nodes of the metabolic network. A consecutive ones clustering method was used to derived important clusters. A wavelet and a feature extraction method were used further to study these clusters. These clusters give us important glues toward elucidating the mechanisms that P.f deplores toward resisting these anti-malaria drugs.