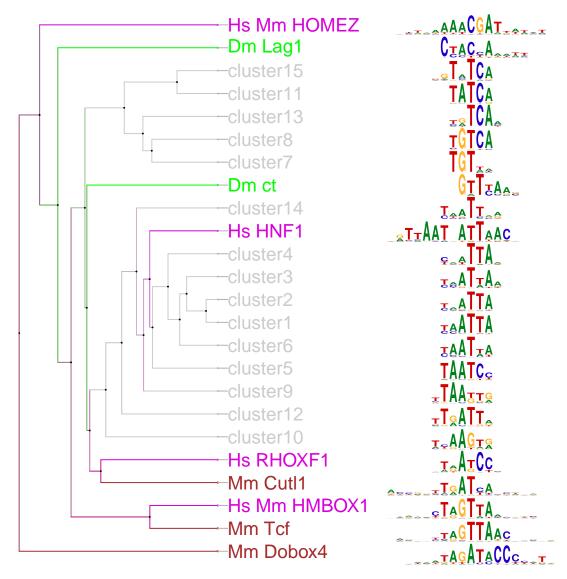
Supplementary Figure 8

Contents

```
##load the library
library(motifStack)
colorSet <- c("Dm"="#00FC00", "b1h"="#00FC00", "sw"="#008080", "bml"="darkgreen",
               "Mm"="brown", "MmDREAM"="blue", "Ms"="#F69156",
               "Hs"="#D900D9")
## function to read example data
readDataDoAna <- function(pcmpath, outpath="output", groupDistance=2.5, trim=0.2){</pre>
    pcms <- readPCM(pcmpath)</pre>
    pfms<-lapply(pcms,pcm2pfm)</pre>
    matalign_path <- "./app/matalign-v4a"</pre>
    neighbor_path <- "./app/neighbor.app/Contents/MacOS/neighbor"</pre>
    system(paste("perl MatAlign2tree.pl --in . --pcmpath", pcmpath, "--out", outpath,
              "--matalign", matalign_path, "--neighbor", neighbor_path, "--tree", "UPGMA"))
    newickstrUPGMA <- readLines(con=file.path(outpath, "NJ.matalign.distMX.nwk"))</pre>
    phylog <- newick2phylog(newickstrUPGMA, FALSE)</pre>
    phylog <- reorderUPGMAtree(phylog, pfms)</pre>
    leaves <- names(phylog$leaves)</pre>
    motifs <- pfms[leaves]</pre>
    if(!is.na(groupDistance)){
        motifSig <- motifSignature(motifs, phylog, groupDistance=groupDistance,</pre>
                                      min.freq=1, trim=trim)
        sig <- signatures(motifSig)</pre>
        gpCol <- sigColor(motifSig)</pre>
    }else{
        motifSig <- NA
        sig <- NA
        gpCol <- NA
    }
    return(list(phylog=phylog, sig=sig, gpCol=gpCol,
                 motifs=DNAmotifAlignment(motifs, minimalConsensus=3),
                 leaves=leaves, unaligned.pfms=motifs))
gpDis <- 2
Fly DNA <- readDataDoAna("pcmsDatasetFly", groupDistance=gpDis)</pre>
unUnique <- Fly_DNA$sig
pcms <- readPCM(file.path("pcmsUni", "14.UniqueCleanCluster"))</pre>
dimmer <- sapply(pcms, motifStack:::isHomoDimer)</pre>
pcms.mono <- lapply(pcms[dimmer], function(.ele){</pre>
    pos <- motifStack:::getHomoDimerCenter(.ele)</pre>
    .ele$mat <- .ele$mat[, 1:as.numeric(pos["pos"])]</pre>
    .ele
})
```

```
pcms.mono <- c(pcms[!dimmer], pcms.mono)</pre>
unUnique <- unUnique[!sapply(unUnique, function(.ele) .ele$name) %in% names(pcms)]
names(unUnique) <- paste("Dm_cluster", 1:length(unUnique), sep="")</pre>
unUnique <- mapply(function(.ele, n) {.ele@name <- n; .ele}, unUnique, names(unUnique))
pfms <- c(lapply(pcms.mono, pcm2pfm), unUnique)</pre>
pfms.cp <- c(lapply(pcms, pcm2pfm), unUnique)</pre>
pcms <- lapply(pfms, function(.ele){</pre>
    mat <- floor(.ele@mat * 1000)</pre>
    new("pcm", mat=mat, name=.ele@name)
})
dir.create("uniMotifTmp")
sta <- lapply(pcms, function(.ele){</pre>
    mat <- cbind(rownames(.ele@mat), "|", .ele@mat)</pre>
    write.table(mat, file=file.path("uniMotifTmp", paste(.ele@name, "pcm", sep=".")),
                 row.names=FALSE, col.names=FALSE, sep="\t", quote=FALSE)
})
uniMotif <- readDataDoAna("uniMotifTmp", groupDistance=NA)</pre>
unlink("uniMotifTmp", recursive = TRUE)
attach(uniMotif)
motifs <- pfms.cp[leaves]</pre>
motifs=DNAmotifAlignment(motifs, minimalConsensus=3, threshold=.5)
leaveCols <- colorSet[gsub("^(Dm|Mm|Hs)_.*$", "\\1", leaves)]</pre>
leaveCols[grepl("cluster", leaves)] <- "gray80"</pre>
leaves <- gsub("Dm_cluster", "cluster", leaves)</pre>
motifPiles(phylog=phylog, motifs,
           labels.leaves=leaves, col.leaves=leaveCols, col.tree=leaveCols,
           plotIndex=FALSE, clabel.leaves=1.5)
detach(uniMotif)
```



Supplementary Figure 8: Species-specific motifs in the HD family