KF-1 ubiquitin ligase: an anxiety suppressor

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Anxiety is an instinct that may have developed to promote adaptive survival by evading unnecessary danger. However, excessive anxiety is disruptive and can be a basic disorder of other psychiatric diseases such as depression. The KF-1, a ubiquitin ligase located on the endoplasmic reticulum (ER), may prevent excessive anxiety; $kf\text{-}1^{-/-}$ mice exhibit selectively elevated anxiety-like behavior against light or heights. It is surmised that KF-1 degrades some target proteins, responsible for promoting anxiety, through the ER-associated degradation pathway, similar to Parkin in Parkinson's disease (PD). Parkin, another ER-ubiquitin ligase, prevents the degeneration of dopaminergic neurons by degrading the target proteins responsible for PD. Molecular phylogenetic studies have revealed that the prototype of $kf\text{-}1$ appeared in the very early phase of animal evolution but was lost, unlike $parkin$, in the lineage leading up to $Drosophila$. Therefore, $kf\text{-}1^{-/-}$ mice may be a powerful tool for elucidating the molecular mechanisms involved in emotional regulation, and for screening novel anxiolytic/antidepressant compounds.

Keywords: depression, ERAD pathway, Parkinson's disease, Alzheimer's disease, animal evolution

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INTRODUCTION

Anxiety, or learned fear, is not necessarily harmful to everyday life but, rather, is a natural ability that may have arose to evade unnecessary dangers. However, excessive anxiety is debilitating or disadvantageous for life as it reduces behavioral activities necessary for adaptation. Moreover, anxiety can be a core symptom of various mental/behavioral disorders, such as major depressive disorders, obsessive-compulsive disorders, panic disorder, adaptive disorder, post-traumatic stress disorder, social withdrawal disorder, and various phobias. Patients with anxiety/depression interpret circumstantial incidences, including episodes, comments, and expressions, in a negative way. The interpretation leads individuals to enhanced capture or delusions caused by potential signs of danger. This suggests that a system of negative and positive regulation of the emotional expression may have developed under the evolutionary constraint. Such a system would be most apparent in highly social animals with relatively little reproduction per generation, like humans (Darwin, 1872). Indeed, there is evidence that the amygdala is responsible for the expression of anxiety or fear, and the prefrontal cortex plays a role in fear extinction by regulating the amygdala-mediated expression of fear (see Bishop, 2007). Although the molecular mechanisms underlying negative and positive regulation of the anxiety are not fully understood, many genes have been reported to affect anxiety or fear (Chen et al., 1994; Gogos et al., 1998; Heisler et al., 1998; Holmes et al., 2003; Miyakawa et al., 1994,

2003; Nakajima et al., 2008; Yamasaki et al., 2008). Among these, the genes related to the serotonergic system are seen to be of special interest. For example, human genomic studies of anxiety/depression have focused on genes related to the monoaminergic neurotransmission of serotonin receptors (5-HT$_{1A}$) and transporters (5-HTT) (see Levinson, 2006; Uher and McGuffin, 2008). Genetic studies using gene-targeting techniques have revealed that knockout mice lacking either *5-ht$_{1A}$* or *5-htt* exhibit significantly increased anxiety-like behaviors (Heisler et al., 1998; Holmes et al., 2003). However, proteins that interact directly with neurotransmitters seem to function downstream, rather than upstream, of the serotonergic pathway.

Anxiety/depression is the most common psychiatric disease seen in patients irrespective of nations, societies, and religions. Consequently, pharmaceutical companies have made extensive worldwide efforts to develop anxiolytic/antidepressant drugs, particularly serotonergic compounds such as selective serotonin reuptake inhibitor (SSRI) and serotonin noradrenalin reuptake inhibitor (SNRI) based on the monoamine hypothesis. The hypothesis assumes that the pathogenesis of depression is caused by the depletion of monoamines such as serotonin in the brain. This assumption has however not been proven in the last 60 years. Testing systems have been developed for rodents to measure behavioral despair and to screen serotonergic drugs. These include the forced swim test and the tail suspension test. Compounds that elevate monoamine levels in the brain reduce the despair-like behavior or immobility time of animals under fearful conditions (Porsolt et al., 1978; Steru et al., 1985). However, approximately one-third of the patients with anxiety/depression do not respond to the serotonergic drugs, suggesting that despair-like behavior in rodents does not precisely represent anxiety/depression in humans, from a pharmacological point of view. Therefore, it is desirable to have some genetic animal models that display excessive anxiety-like behavior specifically without affecting despair-like behavior, as observed in the *kf-1$^{-/-}$* mice (Tsujimura et al., 2008), to look for novel anxiolytic/antidepressant drugs that are effective in patients who do not respond to serotonergic drugs.

IDENTIFICATION AND NATURE OF KF-1

The gene for KF-1 was originally discovered in connection with Alzheimer's disease (AD). Genetic studies of familial AD (FAD) made significant progress in 1990s. In this period, three genes encoding β-amyloid precursor protein, presenilin-1 (PS1), and presenilin-2 (PS2) were identified as the causative genes for FAD (see Hashimoto-Gotoh et al., 2006; Lundkvist and Näslund, 2007), and two other genes were found expressed more frequently in the frontal cortex of an AD patient than a non-AD subject (Yasojima et al., 1997). One of the later two genes, *gfap*, was a known gene encoding glial fibrillary acidic protein, and the other was a novel gene, named *kf-1*. The *kf-1* gene is expressed most prominently in the brain and more or less throughout the entire body, but least, if any, in the liver, which implies that *kf-1* may not be a housekeeping gene. Human *kf-1* has four exons ranging over approximately 30 kb, and is mapped to chromosome 2p11.2. The protein structure deduced from the cDNA sequences has revealed that human KF-1 protein (GenBank Acc No. BAA19739) consists of 685 amino acids, and contains a possible leader peptide (amino acid positions 1–19), two membrane-spanning segments (326–345 and 366–380), and a **RING-H2 finger motif** (621–662) close to the C-terminus (**Figure 1A**). KF-1-like proteins have been found in other animals including fish (*Danio rerio*), lancelet (*Blanchistoma floridae*), sea urchin (*Strongylocentrotus purpuratus*), sea anemone (*Nematostella vectensis*), and tablet animals (*Trichoplax adhaerens*) (**Figure 1A**). Molecular phylogenetic studies suggest that genes for the animal and vertebral proteins may be *kf-1* orthologues (**Figures 1B,C,** respectively). Unexpectedly, KF-1 homologues do not exist in insects (*Drosophila melanogaster*) and thread worms (*Caenorhabditis elegans*, and *C. briggsae*), even though it is likely that the prototype gene appeared in the very early phase of animal evolution, before the separation of Placozoa and Eumetazoa (Miller and Ball, 2008; Srivastava et al., 2008). Homologues of the prototype do not exist in prokaryotes, plants, or fungi, but are found first in one of the most primitive multicellular animals, such as tablet animals (Schubert, 1993) (**Figure 1D**). The results may be consistent with the report based on the EST analysis that some genes, formerly thought to be vertebrate inventions, must have been present in the common metazoan ancestor (Kortschak et al., 2003). Choanoflagellates (*Monosiga brevicollis*, and *M. ovata*), supposedly one of the most closely related unicellular protists to animals (King et al., 2008), do not possess *kf-1* (**Figure 1D**).

The KF-1 protein is an E3 ubiquitin ligase that may modulate the cellular protein levels of its unknown substrate(s) (Lorick et al., 1999). The expression of *kf-1* is also increased in the frontal cortex and hippocampus after chronic administration of SSRI in rats (Yamada et al., 2000). Furthermore, rat *kf-1* expression is elevated
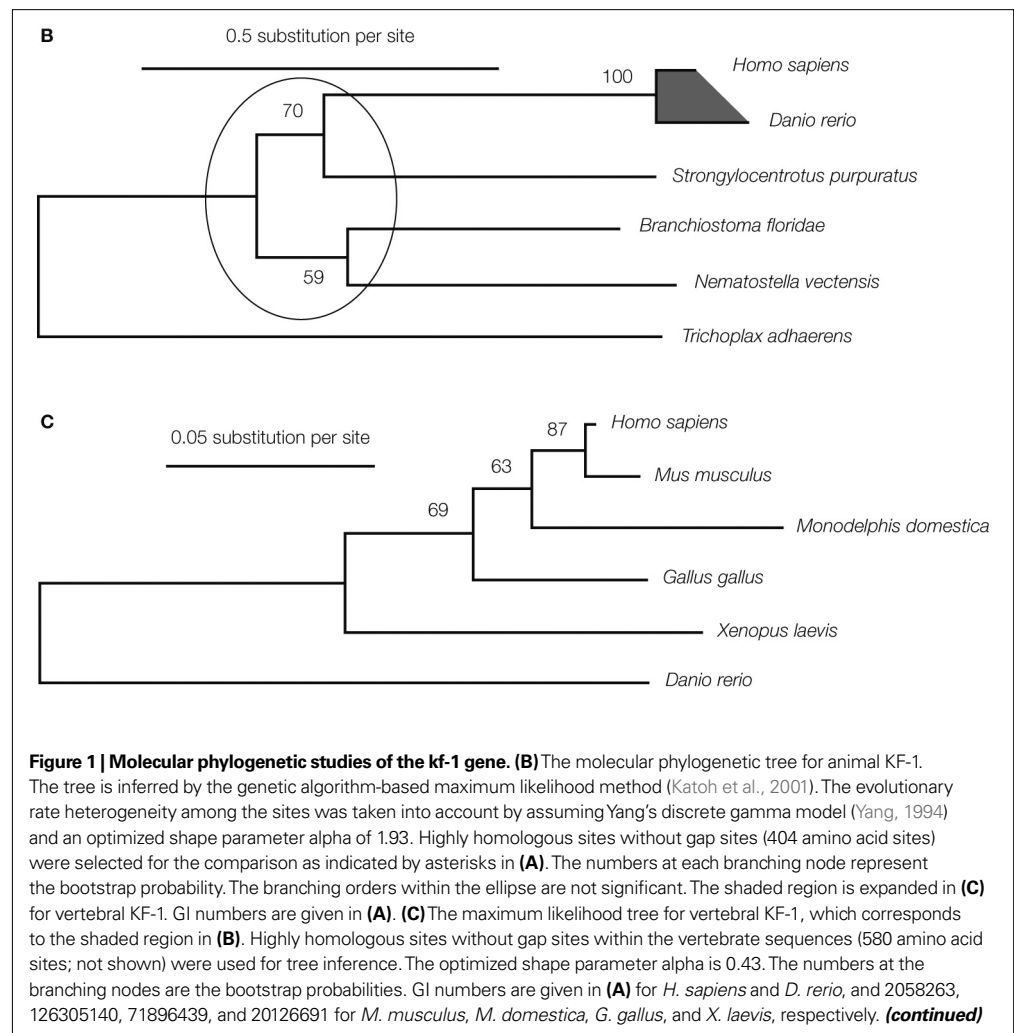
RING-H2 finger motif
A type of zinc binding domains, similar to RING finger motif containing a C3HC4 amino acid motif for binding to two zinc ions. RING-H2 finger motif contains two histidine residues as in C3H2C3 motif.
Many of the RING finger domains function as ubiquitin ligases.

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                                                                                                    120
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Hs  MWL-------KLFFLLLYFLVLFVLARFFE-AIVWYETGIFATQLVDPVALSFKKLKTILECRGLGYSGLPEKKDVRELVEKSGDLMEGELYSALKEE-EASESVSSTNFSGEMHFYELV 111
Dr  MWW-------KLFFLLLYFFILFILARFFE-AIVWYETGIFATQLVDPVTLSFKKLKTILECRGLGYSGLAEKRDVRELVENSGELMQGELYSALKNEKEQAGSDSSTTFSGEMHFYELV 112
Bf  MLV-------RLLLLLVYLCLLLVAVRLLE-AATWFEAGFLAGQVLDPLSISVRRLKMILDSRGISYKGVLEKKELTDLVENSGEPKEGEVLLAAE---DEDTEPTSTNFTGRAHFYEEV 109
Sp  MFM------KKIVFLLVYATLLFLLARMLE-YIPWYQTGLLMMKLIDPVSLSVKKLKSLLDGRGLSYEGVIDKAELTQLVEESGHVMEGEVLMMEQDASEREEEEEPTTTTFSSHA---- 109
Nv  MLT-------KLLLLLVYFFLIFLTTRFLELTASWFEAGCIASQLFDPLSLSVRKLKAILDQRGVSYNGVVEKSELADLVEVSGAVTDPESALTAQGSNDNEQNSDEFTFKGASHFFEEV 113
Ta  MTAGWFLTAIKIILLIVYLICVLLSCKYWN-VQLWKVGDKAARLLLDPATFNLKELIEIIDYRGVSDLNLHDRTNLSYMVNASGLMSEEEKWQSAIIM-AQSKKREAINFTAENYLRAEI 118
CS  M         L  Y          W          DP    L      RG            V    SG          E
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                                                                                                    240
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Hs  EDTKDGIWLVQVIANDRSP--LVGKIHWEKMVKKVSRFGIRTGTFNCSSDPRYCRRRGWVRSTLIMSVPQTSTSKGKVMLKEYSGR--KIEVEHIFKWITAHAASRIK--TIYNAEHLKE 225
Dr  EDTKDGIWLVQVIAQDRNP--LLSTANWGKMVQKVSQFGIRTGTFNCSSDSRYCHKRGWMKSTLIMSVPQTYASKGKVMLKEYNGR--RIETEHIFKWMTAHVASRIK--TVRYSDQLMD 226
Bf  EDTKDSVWLVQVIPEDHIP--LLGPQQWKSLVRKVSRFGIRTGTFKCQLDRKLCWRKGWDRPSLALALPRGHQAKGHISVQVFNT---PSKEQTILDWINQHLSSRTH--SVLSPHQLQT 222
Sp  ------QFSEEVIPRNFGP--LLGKRAWSTVVKKLSRFGIRHGTFDCSIEPSICPRKNWNLPLLLLAMPQGHRHKGQVTMAKFTS---EGKAQQIINWVYLELAKKVN--TERGFGQ-EY 215
Nv  EDTKAGSWLVEVIPENHIP--LLRRKQWSSLKRKMRLFGIRTGSFKCEEDPWLCRKYKWNRPSLVLSMPKGNQPKGNVILQTYQA---KPNVNSVLLWINSELSSKVI--ELDSTNTLNK 226
Ta  EDNTPGIWLLRIDTRNSSKEDILVDNLWDDVVPKLYTLGVQSGTIRCRTCPTICRKNNWTKSDILLLRTTASDSDWSKGILYYPGRLANANSSSILIWMRHHLNSHINHFSTRNFPQLRE 238
CS               W        K        G G     C       C      W                                    W
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                                                                                                    360
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     ***************************** *****************************************************************
Hs  EW---NKSDQYWLKIYLFANLDQPPAFFSALSIKFTGRVEFIFVN---VENWDNKSYMTDIGIYNMPSYILRTPEGIYRYGNHTGEFISLQAMDSFLRSLQPEVNDLFVLSLVLVNLMAW 339
Dr  DW---YQMEKQPVKMFLFARLLQPPAFFSALSIKFTGRIEFIFVD---VRNWDNNTCLEEIGVQQMPSYILKTPEGIYRYGNSTGEFISLHAMDTFLRSVQPEVNDLFILSLVMVNLMAW 340
Bf  DW--LMKNRTHPVQVVFFSRLKQPPMFYSALSVKFTGRVKFGYMR---LNNSRNRLDIS--GREKIPGILVITPERRYWYGTGKGELLNLQSMQTYLRTMQPEVNDIFLVCVVVVNLMAV 335
Sp  DWDKLESNRDVAVKVVLFSRNQEPPVFFSALNLKYSGRVKFVFVS---DSKTFYVLDNR-AQKYWLPSYIIVTPEGKKVYGENNGEYCTYSALDLYLQILSPEANDIFVLTFINVNAICF 331
Nv  IL--QSDRDPNYIYVVYHSTLTEPPMFLSSLSIKFTGRVKFVYCR---SHLKHRKEDIN-FDGFKVPSLFVITPERRVLFGLKKGEIYDYSSLELYLRTLHPEVNDLFLAALVITNLCCM 340
Ta  TI-----KKQSGIHVILFSTLTVAPTFISSLAIKFSGRINFSMVTIKTINDTIQNLFANEFNVEKLPTYRIFTPEKNFTYGNRHGEYYGYHCHHEFLTSLYPATNDIFVAIVIGINLHCL 353
CS               P FSL  K  GR F                         P         TPE    G   GE           L   P ND F       N
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     *************************************
Hs  MDLFITQGATIKRFVVLISTLGTYNSLLIISWLPVLGFLQLPYLDSFYEYSLKLLRYSNTTTLASWVRAD-WMFYSSHPALFLSTYLGHGLLIDYFEKKRRR--NNNNDEVNANNLEWL- 455
Dr  MDLFITQGATIKRFVVLISTLGTYNSLLIISWLPILGFLQLPYLDSFYEYSLKLLRYADTTTIASWVRAD-WTFYSSHPALFLSTYLAHGLLIDYFEKKRRC---SNEDQ-NANNLEWL- 454
Bf  LEVFLCGGQVGVGVLRLLWAVGKYNCVLLMVCLPVVGLFQLPCMEGVVQAGLTALRNISSSGLVAQARQD-WLLYSSHKPFLVGTFLLYSMAVGIVASRWKS-GEEMASETTPTEQAGT- 452
Sp  MGLFLIQGGIAKRICGFLWTIGKYNITLILLWLPLLGIIQLPFLASVQDYSYKMLRLSSQTWLIGKLRQD-WMMYSGHSYIIIGSFLLYCVAVNWVAKKIQA---PTNEDSMTSAM-WLR 446
Nv  LESFLIHGGILRRTFRLLCMLTFYNTSLIMLCLPMVWLFQLPFLQPVLDFTLKCCRCIMSGDIASLLRHD-LMFWMNYDYFVLIGYFVFGFTLGYIRNKYKCYFGVGDDDLEDPNADWL- 458
Ta  LKLLLIDGTFLACLLKFIILFCQYNFAVVLLWFPMYNFFTSPLMHIVYDLTMMVMRCLMGTDFVAAIRRSTHLTHPGCLFMLLTESIIFASVIYYWIECQEG--QISGTESMA-SCNYL- 469
CS         G              YN      P        P        R       R
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                                                                                                    600
                                         ***********************************
Hs  SSLWDWYTSYLFHPIASFQNFPVE--SDWDEDPDLFLERLAFPDLWLHPLIPTDYIKNLPMWRFKCL---------------------------------------------------- 520
Dr  SSLWDWYTSYLVHPIASFQNF-E--SDWDDDPNFLLERLAFPDLWLRPLVPIDYIKNLPTWKFRLL---------------------------------------------------- 517
Bf  NGWWSSFMNYLFQPISTFNHTRPPNLIGLEDGLDYLIERLAVPDLWLHPVIPTDYVRNLPVWLYKGKMPLVRKVCSKCCGWGGSADQAAMVMKLPTVGHTESNRVMSLPPSPRRPGHRPP 572
Sp  ASLVDFR-SLLLRPSPSFRNGLPRN-GHTEEGLDLLIDQMGVPDLWLHPLIRDDYIKDLLTWRYSCQ---------------------------------------------------- 511
Nv  TQDLNYF-SRILQSLSHWQPQIHHTTSGFEDGFEMLVRRLAVPDLWLHPIIPTDYIKQLPTWNFCCK---------------------------------------------------- 524
Ta  --SLSSYQSSLYSILTYRSACERYSREDIEEGIDILIEHFAIPKLYVQPLVPMDYIQSLPTFTYGDK---------------------------------------------------- 534
CS                        P  L  P     DY  L
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                                                                                                    720
Hs  ------------------------GVQSEEEMSEGSQDTENDS-ESENTDTLSSEKEVFEDKQ-SVLHNSPGTASHCDAEACSCANKYCQTSPCERKGRSYGSYNTNEDMEPDWLTWP 612
Dr  ------------------------------------------QPEGSNLA--DRD--KRQK-AMMNQNEDSSGNKVAHQCSSDVLHDHYGCSAA-TKSEESWSGGEEQDTDWSQWP 585
Bf  SPSRHPSQAHKCNKTRSTDDTPLSYSDNHSSDQAKCTCGERSKGASDHGNVYGHDVNVPSKHNNSPSGHGPPPVMPTPSLDKPCTCGNPQQTSSASG---------------------FP 671
Sp  ------WLQNPSSSNDSSDSEKGFMMCRHHTKKCREEKLECPQAEKETETEIETPDASAKSLESKDRENVNCIVASRRHSPKNCTKDAKPKTSSRIKSR------------SPHTDSWP 612
Nv  ------------------------------------------------------DGCLQSNPEHPELNATADCLPCSVGRPS------------------------- 552
Ta  ------------------------------------------------------IRFNHPTWKEKLNWKSDNTEIAQLEENITSEQD-------------------- 567
CS
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                                                                                                    801
        ####################################   ######
        ***********************************   **********
Hs  ADMLHCTECVVCLENFENGCLLMGLPCGHVFHQNCIVMWLAGGR----HCCPVCRWPSYKKKQPYAQHQPLSNDVPS----  685  100.0%
Dr  CGMLHCTECVVCLENFETDCLVMGLPCGHVFHQQCIVVWLAGGR----HCCPVCRWPSYKKRP--VRQRATEQLDPE----  656   69.8%
Bf  LGILPCEDCAICLEEYEVGCSLLGLPCGHSFHERCIMMWLSAGN----HCCPVCRWPAFKFKPA-LHLHSE----------  737   39.1%
Sp  EGILYDAQCAICIEAYTNGAELCGLPCGHAYHQQCIVAWLNNGN----HVCPICRWPAYKKKGSKLSKHME----------  679   37.5%
Nv  -WMIPCGECVICLDEFKPGCTLLGLPCGHSFHQHCIEVWLAGDNTAPHHCCPNCRWPAYRAKSH-VH--------------  617   34.9%
Ta  -YNCDDNQCSICLTNYINDDYLCCLPCSHVFHHDCIVQWLSIGTINT-CRCPLCRWPAYRSYLQPSSACHATSITSNEDSS  646   24.7%
CS        C  C          LPC H  H  CI  WL       CP CRW
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Figure 1 | Molecular phylogenetic studies of the *kf-1* gene. (A) The alignment of KF-1 protein sequences. The sequence identities (%) given at the ends were obtained against human sequences minus the regions after the RING-H2 finger motif, namely after amino acid position 621 in human KF-1. Asterisks show the alignment sites used for inferring the **molecular phylogenetic tree** in **(B)**. The two transmembrane regions of human KF-1 are indicated by '=' and the RING-H2 finger motif for zinc-binding is indicated by '#'. The number given at the right side indicates the amino acid position of the C-terminal amino acid in each line.

The alignment positions, including gaps, are given on the top right of each row. Hs, Dr, Bf, Sp, Nv, and Ta denote *Homo sapiens*, *Danio rerio*, *Blanchistoma floridae*, *Strongylocentrotus purpuratus*, *Nematostella vectensis* and *Trichoplax adhaerens*, respectively. The gene identification (GI) numbers are 1945615, 52219054, 210102052, 115660659, 156406687, and 196009364 in the respective species. Blast *Expect* values (blastp with BLOSUM62 for Matrix) of Dr, Bf, Sp, Nv, and Ta against Hs are $E < 10^{-130}$, $E = 2 \times 10^{-130}$, $E = 3 \times 10^{-117}$, $E = 3 \times 10^{-104}$, and $E = 3 \times 10^{-57}$, respectively. CS denotes consensus amino acid. **(continued)**

Figure 1 | Molecular phylogenetic studies of the kf-1 gene. (B) The molecular phylogenetic tree for animal KF-1. The tree is inferred by the genetic algorithm-based maximum likelihood method (Katoh et al., 2001). The evolutionary rate heterogeneity among the sites was taken into account by assuming Yang's discrete gamma model (Yang, 1994) and an optimized shape parameter alpha of 1.93. Highly homologous sites without gap sites (404 amino acid sites) were selected for the comparison as indicated by asterisks in **(A)**. The numbers at each branching node represent the bootstrap probability. The branching orders within the ellipse are not significant. The shaded region is expanded in **(C)** for vertebral KF-1. GI numbers are given in **(A)**. **(C)** The maximum likelihood tree for vertebral KF-1, which corresponds to the shaded region in **(B)**. Highly homologous sites without gap sites within the vertebrate sequences (580 amino acid sites; not shown) were used for tree inference. The optimized shape parameter alpha is 0.43. The numbers at the branching nodes are the bootstrap probabilities. GI numbers are given in **(A)** for *H. sapiens* and *D. rerio*, and 2058263, 126305140, 71896439, and 20126691 for *M. musculus*, *M. domestica*, *G. gallus*, and *X. laevis*, respectively. **(continued)**

Molecular phylogenetic tree
A diagram showing the evolutionary relationship or history of organisms, genes or proteins. It is inferred by a phylogentic method such as neighbor-joining, maximum parsimony, or maximum likelihood, based on the nucleotide or amino acid sequences of various species.
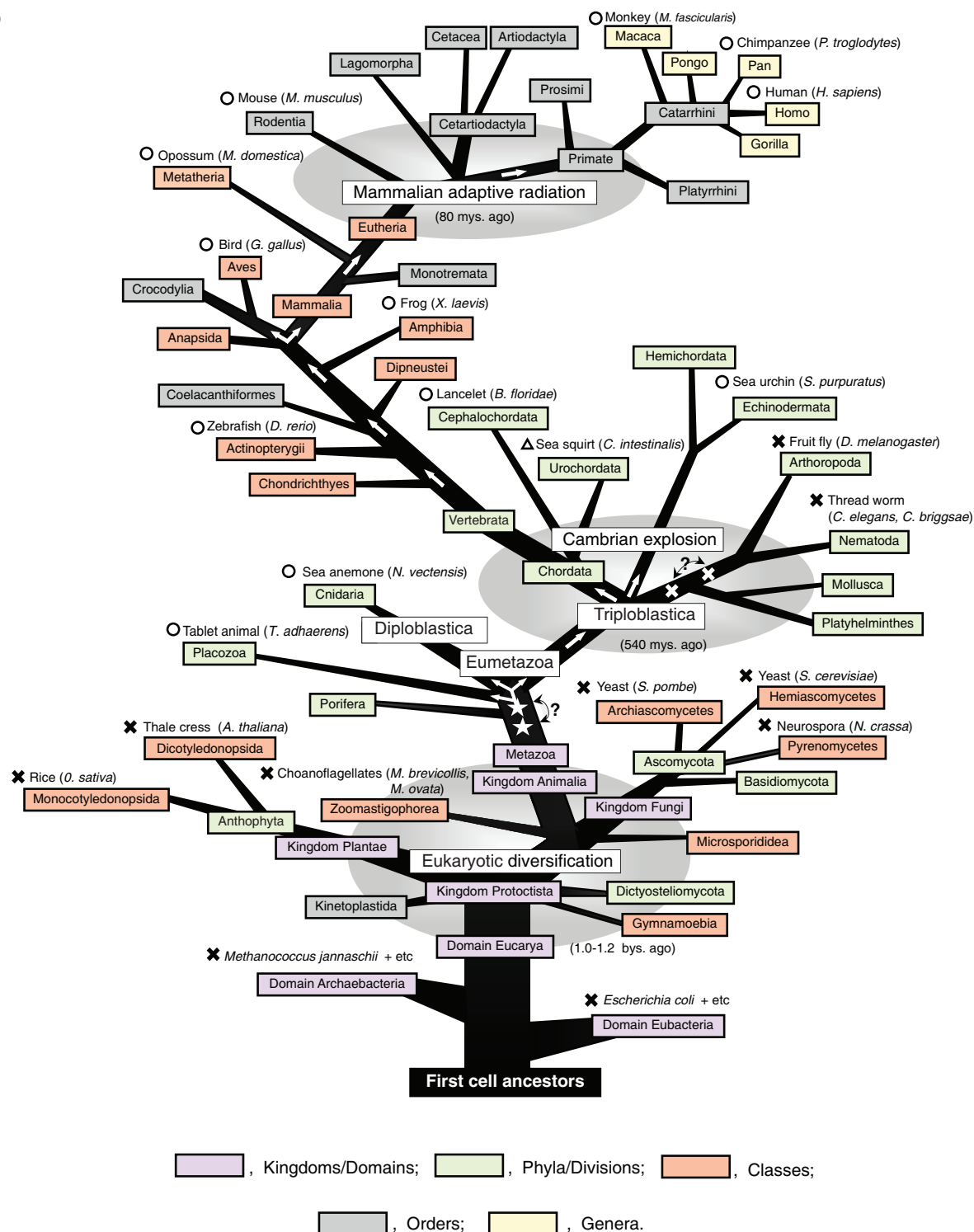
after physical antidepressant treatments, such as electroconvulsive therapy (Nishioka et al., 2003), and repetitive transcranial magnetic stimulation (Kudo et al., 2005). This implies that the up-regulation of *kf-1* expression is associated with some physiological responses to antidepressant treatments rather than being caused by a chemical reaction to serotonergic compounds such as SSRIs. It was, however, not clear whether this was a result, cause, or coincidental side effect of anti-depressive processes.

As a possible correlation between KF-1 and AD has been suggested (Yasojima et al., 1997), and as depression can occur early in the course of AD in *ps1* mutation carriers unaware of their genetic status (Ringman et al., 2004), the first working hypothesis was that presenilins could be KF-1 sub-strates in the endoplasmic reticulum (ER) associated degradation (ERAD) pathway (see Carvalho et al., 2006; Lorick et al., 2006). Therefore, the intracellular localization of KF-1 was examined in comparison to that of presenilins known to be located on ER (Kovacs et al., 1996). The results have revealed that KF-1 is co-localized with both PS1 and PS2 (**Figures 2A,B** and Tsujimura et al., 2008), which implies that KF-1 may in fact be an ER-ubiquitin ligase. The co-localization of KF-1 with other ER markers, such as Der-1 and VCP, has also been observed (Maruyama et al., 2008).

SELECTIVE EXPRESSION OF INCREASED ANXIETY-LIKE BEHAVIORS IN *kf-1*⁻/⁻ MICE

Although mouse *kf-1* is expressed in many tissues, particularly in the brain, the lack of *kf-1* does not lead to any abnormalities in appearance or behaviors, including body weight, reproductive capability, exploratory locomotion, nociception, social behavior, motor coordination, behavioral despair, spatial working memory, and context memory (**Table 1**).

The exception is that *kf-1*⁻/⁻ mice display a pronounced increase in anxiety-like behavior in the light/dark transition test (stay time in light compartment: $p = 0.0004$; number of

D

Cetacea | Artiodactyla

○ Monkey (*M. fascicularis*)
Macaca

Lagomorpha

Pongo

○ Chimpanzee (*P. troglodytes*)
Pan

Prosimi

Cetartiodactyla

Catarrhini

○ Human (*H. sapiens*)
Homo

○ Mouse (*M. musculus*)
Rodentia

Gorilla

Primate

○ Opossum (*M. domestica*)
Metatheria

Platyrrhini

Mammalian adaptive radiation
(80 mys. ago)

Eutheria

○ Bird (*G. gallus*)
Aves

Monotremata

Crocodylia

○ Frog (*X. laevis*)

Mammalia

Amphibia

Anapsida

Hemichordata

Dipneustei

Coelacanthiformes

○ Lancelet (*B. floridae*)
Cephalochordata

○ Sea urchin (*S. purpuratus*)
Echinodermata

○ Zebrafish (*D. rerio*)
Actinopterygii

△ Sea squirt (*C. intestinalis*)
Urochordata

✘ Fruit fly (*D. melanogaster*)
Arthoropoda

Chondrichthyes

✘ Thread worm
(*C. elegans, C. briggsae*)

Vertebrata

Nematoda

○ Sea anemone (*N. vectensis*)
Cnidaria

Cambrian explosion

Chordata

Mollusca

Diploblastica

?

○ Tablet animal (*T. adhaerens*)

Triploblastica
(540 mys. ago)

Platyhelminthes

Placozoa

Eumetazoa

Porifera

✘ Yeast (*S. pombe*)
Archiascomycetes

✘ Yeast (*S. cerevisiae*)
Hemiascomycetes

✘ Thale cress (*A. thaliana*)
Dicotyledonopsida

?

Metazoa

✘ Neurospora (*N. crassa*)
Pyrenomycetes

✘ Rice (*0. sativa*)
Monocotyledonopsida

✘ Choanoflagellates (*M. brevicollis, M. ovata*)

Kingdom Animalia

Ascomycota

Anthophyta

Zoomastigophorea

Kingdom Fungi

Basidiomycota

Kingdom Plantae

Microsporididea

Eukaryotic diversification

Kinetoplastida

Kingdom Protoctista

Dictyosteliomycota

Gymnamoeba

Domain Eucarya
(1.0-1.2 bys. ago)

✘ *Methanococcus jannaschii* + etc

Domain Archaebacteria

✘ *Escherichia coli* + etc

Domain Eubacteria

First cell ancestors

☐ , Kingdoms/Domains;   ☐ , Phyla/Divisions;   ☐ , Classes;

☐ , Orders;   ☐ , Genera.

Figure 1 | Molecular phylogenetic studies of the *kf-1* gene.
(D) Presence/absence, generation, and transmission of *kf-1* on
the evolutionary tree of life. The tree is essentially the same as the one,
that was previously presented (Hashimoto-Gotoh et al., 2003), with minor
modifications and corrections except for the genes concerned. '?'
with bi-arrows and two open stars or crosses denotes uncertainty
of the precise position of generation or loss of *kf-1*, respectively.

Circles or crosses represent species in which the presence or absence
of *kf-1* has been confirmed in the Gen Bank database on January 29, 2009.
In the case of sea squirt (*Ciona intestinalis*), the presence of *kf-1*
is represented by a triangle as only fragmented amino acid sequences
have been found similar to those of KF-1. This is probably due to
an incomplete determination of genomic sequences. Detailed data
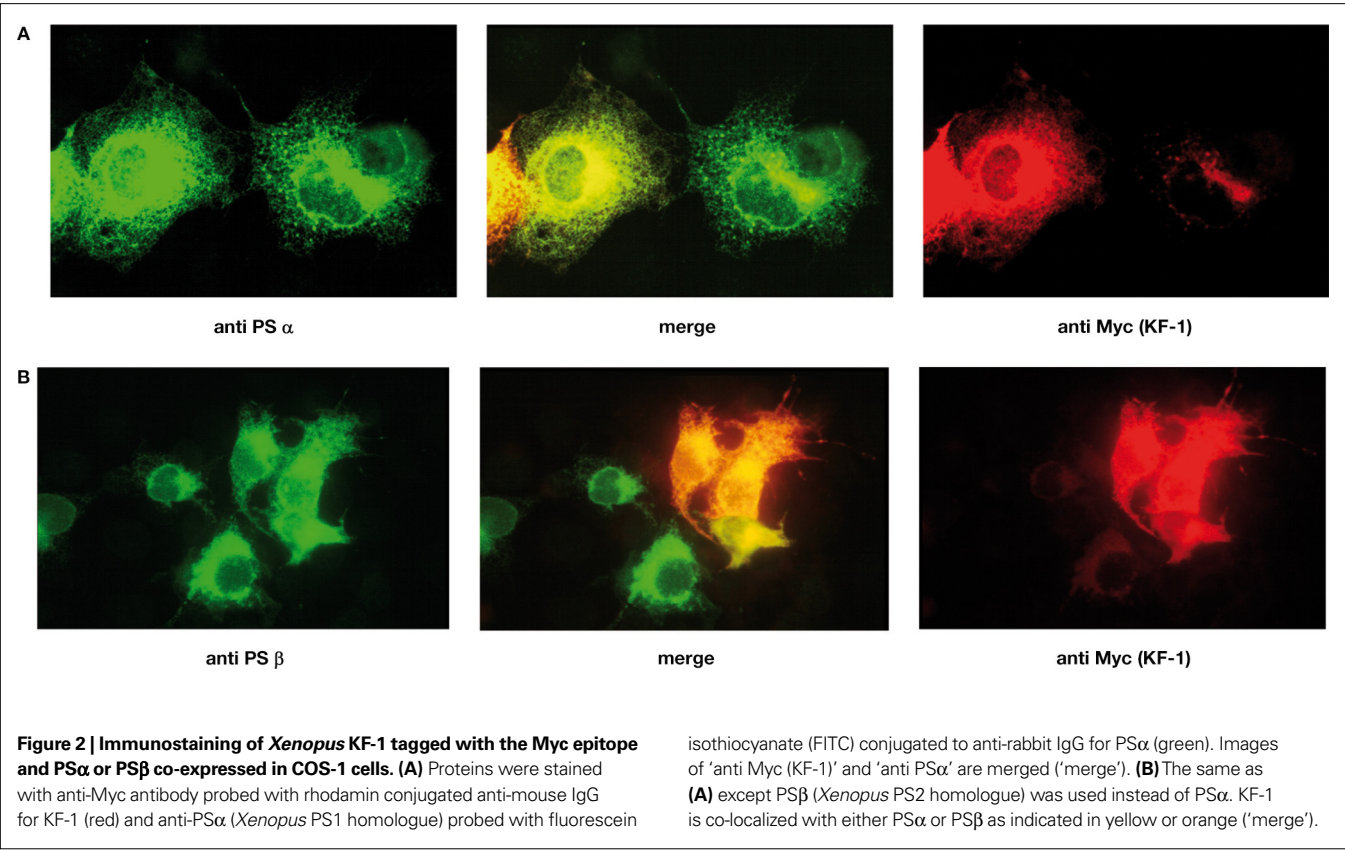on which this illustration is essentially based are given in **(A)**, **(B)**, and **(C)**.

Figure 2 | Immunostaining of *Xenopus* KF-1 tagged with the Myc epitope and PSα or PSβ co-expressed in COS-1 cells. (A) Proteins were stained with anti-Myc antibody probed with rhodamin conjugated anti-mouse IgG for KF-1 (red) and anti-PSα (*Xenopus* PS1 homologue) probed with fluorescein isothiocyanate (FITC) conjugated to anti-rabbit IgG for PSα (green). Images of 'anti Myc (KF-1)' and 'anti PSα' are merged ('merge'). **(B)** The same as **(A)** except PSβ (*Xenopus* PS2 homologue) was used instead of PSα. KF-1 is co-localized with either PSα or PSβ as indicated in yellow or orange ('merge').

Table 1 | Behavioral phenotypes of *kf-1*⁻/⁻ mice compared to the behavioral phenotypes of *kf-1*⁺/⁺ littermates.

Tests	Measurements	Phenotypes[a]
General health examination	Whisker, coat, reflexes	=
Physical test	Body temperature	=
	Body weight	=
	Wire hanging time	↑
	Grip strength	=
	Auditory capacity	=
Light/dark transition test	Anxiety	↑
Open field test	Exploratory locomotion	=
Elevated plus maze test	Anxiety	↑
Hot plate test	Pain sensitivity (latency time)	=
Social interaction test	Total duration of contacts	=
	Number of contacts	=
	Total duration of active contacts	=
	Mean duration/contact	=
	Distance traveled	=
Rotarod test	Motor coordination	=
Prepulse inhibition (PPI) test	Sensorimotor gating	↑
Porsol forced swimming test	Immobility time (behavioral despair)	=
T-maze test	Spatial working memory	=
Cued and contextual fear conditioning test	Immediate freezing during conditioning phase	↑
	Contextual testing conducted after conditioning	=
	Cued test with altered context	=
Tail suspension test	Immobility time (behavioral despair)	=

The testing order is from the top to the bottom of this table. Further details are presented in Tsujimura et al. (2008). *[a] =, no significant difference; ↑, increased in kf-1⁻/⁻ compared to kf-1⁺/⁺. The raw data of the results are available at https://behav.hmro.med.kyoto-u.ac.jp/.*

SNP

DNA sequence variation, called single nucleotide polymorphism, occurring on a single nucleotide in the genome among individual members of a species or between paired chromosomes within an individual. 'SNP (pronounced *snip*)' was defined initially as that with a minor allele frequency of ≥1%, but this definition is rather artificial and not meaningful.

Ubiquitination

Ubiquitination (or ubiquitylation) is the post-translational tagging reaction mediated by E3 ubiquitin ligase to a protein with one or more highly conserved peptide molecules called 'ubiquitin' consisting of 76 amino acids. Ubiquitination targets the substrate protein for proteasomal degradation.

transitions: $p = 0.0392$) (Tsujimura et al., 2008). Consistently, the mice also show 'timidity-like' responses under stressful situations, such as prolonged wire-hanging time, enhanced immediate freezing with an aversive foot shock, and decreased locomotor activity at heights (**Table 1** and Tsujimura et al., 2008). As significant differences are not observed in body weight and grip strength, and in general exploratory locomotion, the 'timidity-like' responses are likely to be related to psychological but not physical alterations in *kf-1*$^{-/-}$ mice. This implies that KF-1 plays a role in emotional control by suppressing anxiety at least in mice. In this context, it is of particular interest to note that a number of **SNP**s are reported in the coding and non-coding exonic regions of human *kf-1* according to the NCBI's SNP database, and some are supposed to substantially modulate KF-1 activity in homozygous or dual heterozygous carriers; for example, rs35921467 resulting in a frame shift at amino acid position 180, rs17857046 and rs17853383 in critical amino acid substitutions (S to P and P to H) at 251 and 502, respectively, and rs11695337 in a termination codon at 626.

As KF-1 is an E3 ubiquitin ligase located on ER, KF-1 may be responsible for conducting the ERAD pathway, in which some factors promoting

anxiety may be targeted. Therefore, the absence or reduction of KF-1 activity may result in 'enhanced anxiety' by increasing putative anxiety promoters (**Figure 3A**). Regarding serotonergic metabolisms, it is not clear at the moment whether or how the ERAD pathway directed by KF-1 is associated with the serotonergic pathway, and it has to be elucidated in future. However, KF-1's targets for proteasomal degradation should be neither the serotonin receptor (5-HT$_{1A}$) nor the transporter (5-HTT), because a lack of these proteins increases anxiety-like behaviors (Heisler et al., 1998; Holmes et al., 2003).

KF-1 AND PARKIN AS ER-BASED E3 UBIQUITIN LIGASES

Ubiquitination plays an essential regulatory role in all critical eukaryotic cellular processes. Proteasomal degradation through the ERAD pathway is not an exception. It has been well established that these processes play an important role in a variety of human somatic diseases, ranging from cancer, viral infection, diabetes, and inflammation to muscle wastage and neurodegenerative disorders (see Kostova et al., 2007; Petroski, 2008). However, there are no reports on the animal specific ubiquitin ligase, which is responsible for emotional control or mental disorders.
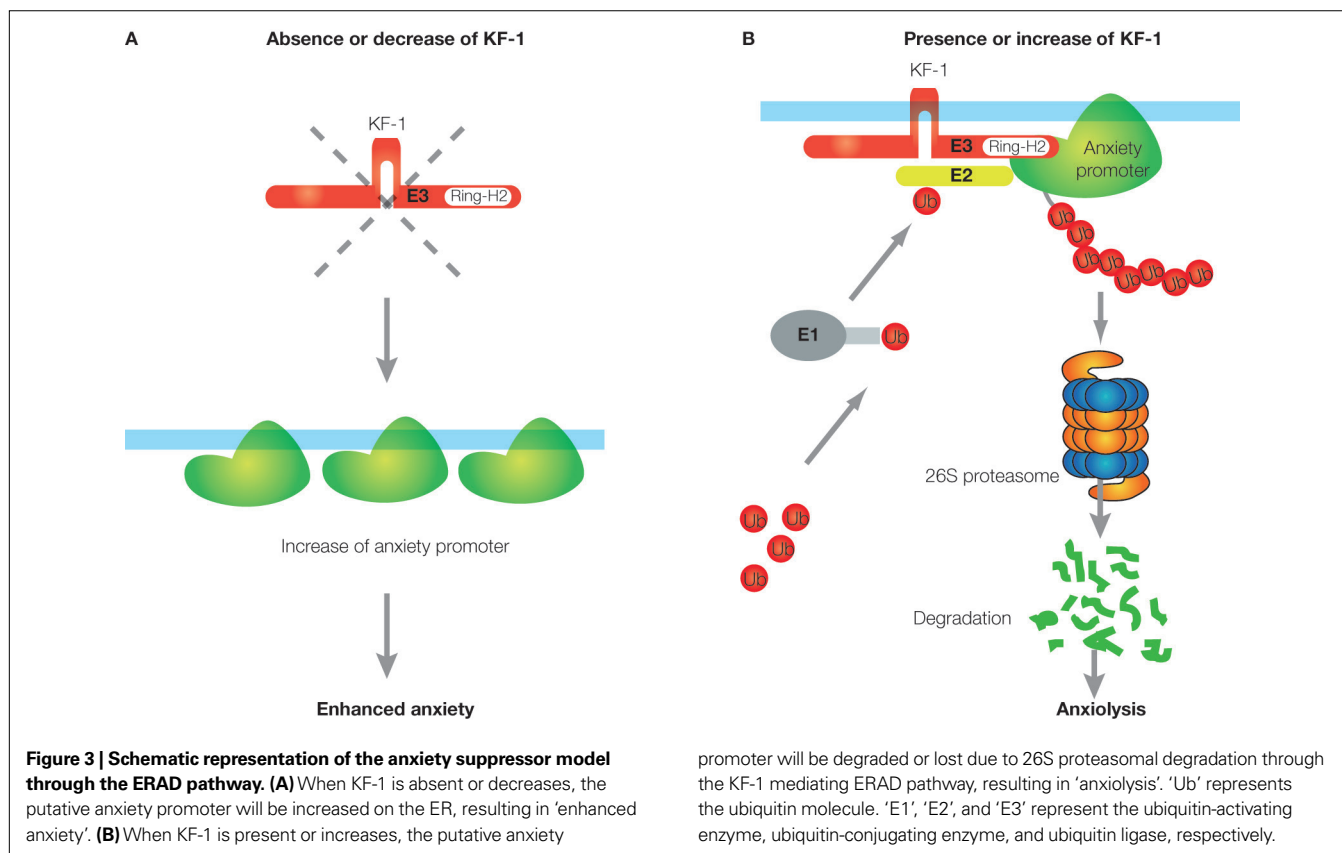
Figure 3 | Schematic representation of the anxiety suppressor model through the ERAD pathway. (A) When KF-1 is absent or decreases, the putative anxiety promoter will be increased on the ER, resulting in 'enhanced anxiety'. **(B)** When KF-1 is present or increases, the putative anxiety promoter will be degraded or lost due to 26S proteasomal degradation through the KF-1 mediating ERAD pathway, resulting in 'anxiolysis'. 'Ub' represents the ubiquitin molecule. 'E1', 'E2', and 'E3' represent the ubiquitin-activating enzyme, ubiquitin-conjugating enzyme, and ubiquitin ligase, respectively.

Concerning neurodegenerative disorders, Parkin, another ER-ubiquitin ligase, plays a key role in preventing the degeneration of dopaminergic neurons in Parkinson's disease (PD). A recessive mutation in *parkin* is responsible for the neurodegenerative disorder known as the autosomal recessive juvenile form of Parkinsonism, the most common form of familial PD (Kitada et al., 1998). Human *parkin* with 12 exons ranging over approximately 1.5 Mb is located on chromosome 6q25.2.27, and encodes an E3 ubiquitin ligase, consisting of 465 amino acids, with two RING finger motifs. Parkin substrates have been identified being degraded through the ERAD pathway, such as Pael-R, CDCrel-1, α-Syn, and synphilin-1 (Yang et al., 2003). The accumulation of these proteins in dopaminergic neurons leads to ER stress-induced apoptosis, resulting in PD (see von Coelln et al., 2004). Promoting their degradation would aid PD patients by preventing neurodegeneration. In fact, the neuronal synthesis of the Parkin substrate Pael-R has been shown to cause a loss of dopaminergic cells in transgenic *Drosophila* models. The co-expression of *parkin* results in the degradation of Pael-R and nearly completely blocks neurodegeneration, whereas interfering with the function of endogenous *Drosophila* Parkin promotes Pael-R accumulation and augments its toxicity (Yang et al., 2003). Hence, the over-expression of *parkin* has emerged as a powerful approach to PD with

complementary effects to approaches described for the use of neurotrophic factors against PD (Ulusoy and Kirik, 2008). An analogy could be made by having such *Drosophila* models to study the role of KF-1 in anxiety/depression. However, the analogy may not be feasible as no evidence is available for the involvement of neuronal cell death in anxiety/depression, and *kf-1* is absent in *Drosophila* (**Figure 1D**). It should be noted that *kf-1*$^{-/-}$ mice have increased **PPI** values (**Table 1**), which implies that at least one of the KF-1 target proteins is involved in regulation of **sensorimotor gating**. In such cases, over-expression of *kf-1* might cause serious side effects as the PPI values are usually reduced in schizophrenic patients. Despite this, one cannot rule out the possibility of *kf-1* up-regulation as an anxiolytic/antidepressant treatment.

Instead, *kf-1*$^{-/-}$ mice provide a powerful tool for some pharmacological approaches to anxiety/depression. For example, by using the mice, one can identify the putative anxiety promoters that are degraded through the ERAD pathway mediated by KF-1 and are responsible for the sensitivity to potential signs of danger or to stressful situations (**Figures 3A,B**). KF-1 substrates acting as anxiety promoter may be found by the computed differential screening in 2-dimensional gel electrophoresis (see Marengo et al., 2008) of cell homogenates derived from the frontal cortex or hippocampus of *kf-1*$^{-/-}$ and *kf-1*$^{+/+}$ mice. As the

Figure 4 | A circuitry presentation of the anxiety suppressor model by the up-regulation of *kf-1* expression.
The circuitry model was constructed by considering the fear extinction neural model (Sotres-Bayon et al., 2006).
The anxiolytic pathway after *kf-1* up-regulation is incorporated into the model.
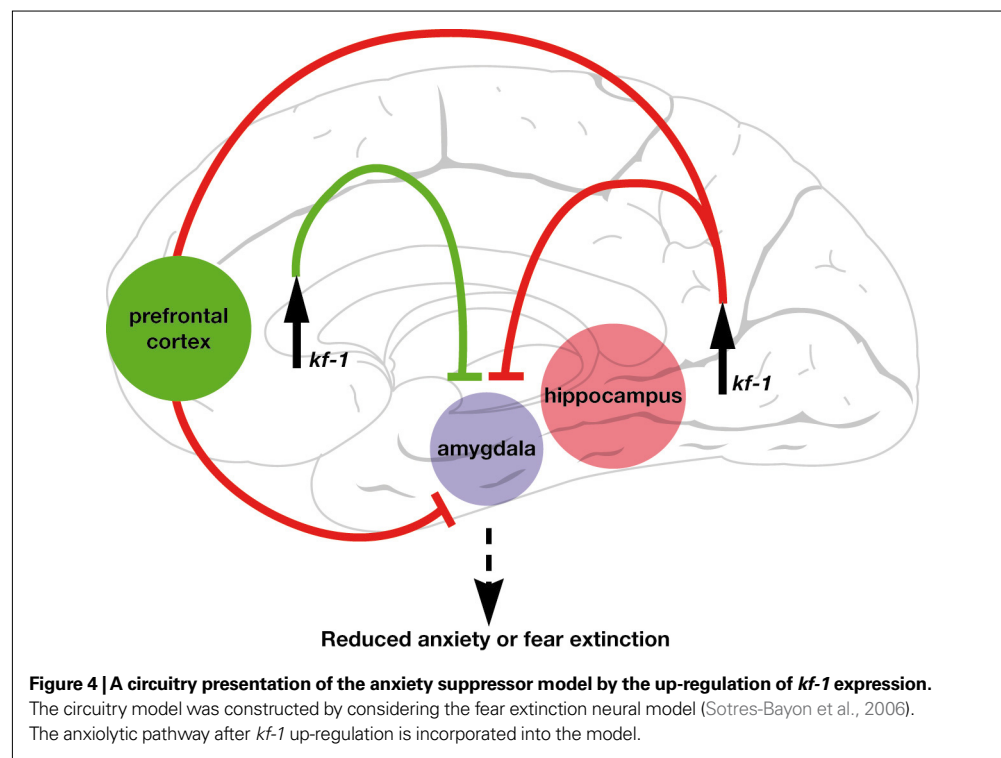
expression of *kf-1* is up-regulated in the frontal cortex and hippocampus after antidepressant treatments, it is likely that the anxiolytic/anti-depressive effect of KF-1 functions primarily in these tissues and that their inhibitory signals are transmitted to the amygdala to prevent the manifestation of anxiety or fear (**Figure 4**). This proposal may be highly hypothetical and hence has to be elucidated in future studies. To conclude, examining *kf-1*⁻/⁻ mice in a simple light/dark transition test is an effective method of screening novel anxiolytic/antidepressant drugs that inhibit the putative anxiety promoters.

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