CoA-dependent activation of mitochondrial acyl carrier protein links four neurodegenerative diseases

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Abstract

PKAN, CoPAN, MePAN, and PDH-E2 deficiency share key phenotypic features but harbor defects in distinct metabolic processes. Selective damage to the globus pallidus occurs in these genetic neurodegenerative diseases, which arise from defects in CoA biosynthesis (PKAN, CoPAN), protein lipoylation (MePAN), and pyruvate dehydrogenase activity (PDH-E2 deficiency). Overlap of their clinical features suggests a common molecular etiology, the identification of which is required to understand their pathophysiology and design treatment strategies. We provide evidence that CoA-dependent activation of mitochondrial acyl carrier protein (mtACP) is a possible process linking these diseases through its effect on PDH activity. CoA is the source for the 4'-phosphopantetheine moiety required for the posttranslational 4'-phosphopantetheinylation needed to activate specific proteins. We show that impaired CoA homeostasis leads to decreased 4'-phosphopantetheinylation of mtACP. This results in a decrease of the active form of mtACP, and in turn a decrease in lipoylation with reduced activity of lipoylated proteins, including PDH. Defects in the steps of a linked CoA-mtACP-PDH pathway cause similar phenotypic abnormalities. By chemically and genetically re-activating PDH, these phenotypes can be rescued, suggesting possible treatment strategies for these diseases.

Introduction

Coenzyme A (CoA) is an essential cofactor participating in approximately 9% of all cellular metabolic reactions, such as the tricarboxylic acid (TCA) cycle, and fatty acid synthesis and degradation (Leonardi et al., 2005; Strauss, 2010). CoA is synthesized de novo in cells, utilizing vitamin B5 as a starting molecule and requiring five enzymatic reactions. These are carried out by pantothenate kinase (PANK), phosphopantothenoylcysteine synthetase (PPCS), phosphopantothenoylcysteine decarboxylase (PPCDC), phosphopantetheine adenylyltransferase (PPAT), and dephospho-CoA kinase (DPCK), respectively (Leonardi et al., 2005; Strauss, 2010). In some organisms, including Drosophila melanogaster, mice, and humans, PPAT and DPCK enzyme activities are performed by a single bifunctional protein, referred to as CoA synthase or COASY (Fig 1). The intermediate products that are being sequentially formed from vitamin B5 during the CoA de novo biosynthesis pathway are as follows: 4'-phosphopantothenate, 4'-phosphopantothenoylcysteine, 4'-phosphopantetheine, dephospho-CoA, and CoA (Fig 1). Enzymes of the CoA de novo biosynthesis pathway are evolutionarily conserved, further underscoring the importance of this pathway for all living organisms.

Two autosomal recessive neurodegenerative diseases are caused by mutations in genes encoding enzymes of the CoA pathway. Pathogenic variants in PANK2 and COASY lead to two early-onset neurodegenerative diseases: pantothenate kinase-associated neurodegeneration (PKAN) and CoA synthase protein-associated neurodegeneration (CoPAN). The human genome contains four genes encoding pantothenate kinase homologs, PANK1-4, and only mutations in PANK2 are associated with PKAN. PKAN and CoPAN patients accumulate iron in the globus pallidus, a basal ganglia structure in the brain (Hayflick et al., 2003; Dusi et al., 2014). Iron accumulation is visible on T2-weighted imaging as a hypointense signal on MRI. In PKAN and CoPAN areas, T2-hyperintense signals are also seen at the globus pallidus, indicating edema and tissue damage (Kruer et al., 2012). These CoA-linked
Figure 1.
diseases are characterized by progressive motor dysfunction and severe dystonia. Damage to the globus pallidus occurs in other inborn errors of metabolism as well. MePAN, a third childhood-onset neurodegenerative disorder, manifests with damage to the globus pallidus, which is also visible on brain MRI as hyperintense signal on T2-weighted imaging although without the iron-associated signal abnormalities (Heimer et al., 2016). MePAN patients carry mutations in the gene encoding mitochondrial fatty acyl-acyl-coenzyme A reductase (MECR), one of four enzymes involved in the elongation of fatty acids in mitochondria to form octanoic acid, a precursor of lipoic acid (Brody et al., 1997; Wada et al., 1997; Feng et al., 2009). Finally, mutations causing impairment of a component of the pyruvate dehydrogenase complex, PDH-E2, lead to PDH-E2 deficiency and cause a form of Leigh disease, in which neuroradiographic abnormalities are again observed specifically in the globus pallidus (Head et al., 2005; McWilliam et al., 2010; Leoni et al., 2012). PDH catalyzes the oxidative decarboxylation of pyruvate to produce acetyl-CoA, thereby linking glycolysis to the TCA cycle (Fig 1). The symptoms, signs, and MRI characteristics of PKAN and PDH-E2 deficiency can be similar. In patients with a clinical suspicion of PKAN, PDH-E2 deficiency should also be considered in the differential diagnosis and vice versa (Head et al., 2005; McWilliam et al., 2010; Leoni et al., 2012). The clinical and neuroradiographic overlapping features of PKAN, CoPAN, MePAN, and PDH-E2 deficiency suggest a common element in their pathogeneses.

The pathways of CoA biosynthesis, mitochondrial fatty acid synthesis, and glycolysis/TCA cycle show interdependency and interconnectivity (Leonardi et al., 2005; Strauss, 2010; Beld et al., 2014; Kastaniotis et al., 2017; Fig 1), but a specific molecular etiology common to these four disorders has been lacking.

Herein, we propose a common underlying pathway directly connecting these four diseases. In most CoA-dependent metabolic reactions, CoA acts as an acyl carrier; the acyl moiety is transferred between CoA and another molecule (e.g., carnitine), leaving CoA intact and available for further transfer reactions. CoA-dependent acyl transfer occurs during the TCA cycle, fatty acid synthesis, and fatty acid degradation (Fig 1). Because CoA is re-used as acyl carrier component, these reactions do not lead to reduced levels of total CoA. In contrast, one specific form of posttranslational modification “consumes” CoA. 4'-phosphopantetheinylation adds a 4'-phosphopantetheine moiety to specific proteins resulting in their activation. For this modification, the 4'-phosphopantetheine moiety is derived from CoA (Beld et al., 2014). Therefore, this reaction forms a 4'-phosphopantetheinylated protein, and adenine 3',5'-bisphosphate is released and thereby causes a net loss of CoA (Elovson & Vagelos, 1968). In humans, a select group of proteins requires this 4'-phosphopantetheine moiety in order to function, e.g. 10-formyltetrahydrofolate dehydrogenase, an enzyme of folate metabolism (Strickland et al., 2010), cytosolic fatty acid synthase, and mitochondrial acyl carrier protein (mtACP; Joshi et al., 2003; Beld et al., 2014). We focussed on mtACP because human PANK2, COASY, and PDH-E2 are mitochondrial proteins (Kotzbauer et al., 2005; Dusi et al., 2014) and mitochondria are defective in various PKAN animal models (Rana et al., 2010; Brunetti et al., 2014; Orellana et al., 2016; Jeong et al., 2019). The active 4'-phosphopantetheinylated form of mtACP is referred to as holo-mtACP. mtACP, known in humans as NDUFAB1, is one of the subunits of the respiratory chain complex I and plays a central role in mitochondrial fatty acid synthesis (Brody et al., 1997; Feng et al., 2009). In this latter process, the thiol group of the 4'-phosphopantetheine prosthetic group forms a covalent bond with a constituent of complex I, the a subunit of complex I, leading to a net loss of CoA (Elovson & Vagelos, 1968). In eukaryotic cells, this process of 4'-phosphopantetheinylation is catalyzed by pyruvate dehydrogenase (PDH) catalyzes the oxidative decarboxylation of pyruvate to produce acetyl that is coupled to CoA to produce acetyl-CoA. Impaired function of this enzyme leads to PDH-E deficiency. The product of the PDH reaction, acetyl-CoA, is the fuel for the TCA cycle which is also a CoA re-using pathway as indicated. OAA: oxaloacetate.

Figure 1. Metabolic pathways in which coenzyme A is formed, re-used, or consumed and their interconnections.

A De novo biosynthesis pathway of coenzyme A (CoA) is a pathway during which CoA is produced. Vitamin B5 is taken up by cells and converted into CoA by the action of five enzyme complexes (Leonardi et al., 2005; Strauss, 2010). These are carried out by pantothenate kinase (PANK), phosphopantetheinylcysteine synthetase (PPCS), phosphopantetheinylcysteine decarboxylase (PPCD), phosphopantetheinyltransferase (PPAT), and dephospho-CoA kinase (DPCK), respectively. In Dro sophila melanogaster, mice, and humans, PPAT and DPCK enzyme activities are carried out by a single bifunctional protein, CoA synthase or COASY. Abbreviations of the enzymes are provided. The starting product vitamin B5/pantothenate, the intermediate, 4'-phosphopantetheine, and the final product CoA are depicted. PKAN and CoPAN are inherited recessive diseases caused by homozgyous mutations in PANK2 and COASY, respectively.

B Formation of holo-mtACP (active form of mitochondrial acyl carrier protein) is a CoA consuming metabolic reaction (Beld et al., 2014). 4'-phosphopantetheinylation of inactive apo-mtACP is required to produce the active holo-mtACP form. During this process, CoA serves as the source for the 4'-phosphopantetheinylation, and hereby, a CoA molecule is degraded and adenine 3',5'-bisphosphate is released. Holo-mtACP plays a key role in mitochondrial fatty acid synthesis, which can be visualized in (C).

C Fatty acid synthesis is a metabolic pathway in which CoA is re-used (Kastaniotis et al., 2017). Mitochondrial fatty acid synthesis (mtFASII) is required for the synthesis of lipoic acid, a pathway that is dependent on the activity of holo-mtACP. In this pathway, malonic acid is converted to malonyl-CoA, which requires holo-mtACP (or NDUFAB1 in humans) for the formation of malonyl-holo-ACP as well as for subsequent downstream steps. Mitochondrial Enoyl-[acyl-carrier-protein] reductase (MECR, defective enzyme in MEPAN) is required for the formation of acyl-holo-mtACP as indicated. Via this pathway lipoic acid is formed from octanoyl-holo-mtACP by lipoic acid synthetase (LASS). This product is then required for the formation of lipoylated pyruvate dehydrogenase (PDH), lipoylated α-ketoglutarate dehydrogenase (αKGDH), lipoylated branched-chain α-keto acid dehydrogenase (BCKDH), and lipoylated glycine cleavage system (GCV). Lipoylation of the above-mentioned proteins is necessary for catalysis of their respective reactions to occur.

D Fatty acid (FA) degradation or β-oxidation is an example of a degradation pathway in which CoA is re-used. The starting mitochondrial precursor acyl-carnitine, the intermediates, and the end product of this pathway are indicated, as well as reactions in which CoA is required (but not consumed) and released.

E In the glycolysis and TCA cycle, CoA is re-used. Pyruvate dehydrogenase (PDH) catalyzes the oxidative decarboxylation of pyruvate to produce acetyl that is coupled to CoA to produce acetyl-CoA. Impaired function of this enzyme leads to PDH-E2 deficiency. The product of the PDH reaction, acetyl-CoA, is the fuel for the TCA cycle which is also a CoA re-using pathway as indicated. OAA: oxaloacetate.
LIPT2 and LIPT1. By the action of the conserved enzyme lipoic acid synthase (LIAS in humans), protein-bound octanoate is transformed into lipoic acid (LA) by the insertion of two sulfhydryl groups (Booker, 2004; Hiltunen et al., 2010; Solmonson & DeBerardinis, 2018), enabling the now-lipoylated proteins to function (Fig 1; Hiltunen et al., 2010). Based on these reports, we hypothesized that, because 4'-phosphopantetheinylation of mtACP consumes CoA, this reaction may be most sensitive to impaired CoA biosynthesis. Consequently, downstream processes are predicted to be affected as well, including decreased lipoylation and activity of PDH (Fig 2).

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**Figure 2.** *De novo* CoA biosynthesis pathway and key downstream steps to link PKAN, CoPAN, MePAN, and PDH-E2 deficiency. **Left part:** Proposed linear pathway linking CoA-mtACP-PDH. From top to bottom: The *de novo* CoA biosynthesis pathway starts with the cellular uptake of pantothenate (Vitamin B5). Pantothenate kinase (PANK), phosphopantothenoylcysteine synthetase (PPCS), phosphopantothenoylcysteine decarboxylase (PPCDC), and coenzyme A synthase (COASY) are enzymes required for the *de novo* biosynthesis of CoA. Mitochondrial acyl carrier protein (mtACP) undergoes a posttranslational modification and active *holo*-mtACP is formed. This posttranslational modification consists of 4'-phosphopantetheine, which is derived from CoA. *Holo*-mtACP in turn is required for lipoylation of PDH-E2, a modification necessary for activation of the PDH complex. It is hypothesized that a decrease in CoA biosynthesis leads to decreased amounts of *holo*-ACP, decreased lipoylation of PDH-E2, and decreased activity of PDH. **Right part:** Steps that are affected in PKAN, CoPAN, MePAN, and PDH-E2 deficiency. Primary affected steps caused by the genetic defects are depicted as the most upstream steps as well as the hypothesized downstream steps for CoPAN, PKAN, and MePAN.
This hypothesis could explain how defects in PANK2 and COASY might lead to a phenotype similar to that arising from defects in MECR and PDH (Fig 2). A central tenet of this hypothesis is that impaired de novo biosynthesis of CoA leads to decreased levels of holo-mtACP. Here, we investigated this hypothesis.

The de novo CoA biosynthesis pathway and the metabolic reactions presented in Figs 1 and 2) are highly conserved between human and Drosophila melanogaster (Leonardi et al., 2005; Bosveld et al., 2008). Our combined approach employing the versatile genetic tools of Drosophila melanogaster and mammalian cells enabled us to demonstrate that impaired CoA biosynthesis leads to decreased levels of active, 4'-phosphopantetheinylated mtACP. This observation was associated with decreased lipoylation of PDH-E2 and decreased PDH activity. Our results revealed the presence of a CoA-mtACP-PDH pathway in which the 4'-phosphopantetheinylation of mtACP is a key step. Next, we showed that stimulation of PDH rescued phenotypes caused by impaired CoA biosynthesis, highlighting PDH as a possible common target for ameliorating diseases induced by defects in the CoA-mtACP-PDH pathway. Our findings combined with those reported by Jeong et al suggest therapeutic approaches for PKAN, CoPAN, MePAN, and PDH-E2 deficiency.

Results

holo-mtACP levels are reduced by impeding CoA biosynthesis

PKAN and CoPAN patients carry mutations in genes coding for pantothenate kinase 2 and COASY, enzymes required for CoA biosynthesis (Fig 1). To test our hypothesis (presented in Fig 2), we first investigated consequences of impaired CoA biosynthesis on 4'-phosphopantetheinylated mtACP. For this, we chose Drosophila melanogaster because of its conserved metabolic steps and genes and its versatile genetic tools. mtACP requires activation in order to function; the active holo form is generated by enzymatic transfer of a negatively charged 4'-phosphopantetheine moiety to a conserved serine residue of the inactive apo form (Elovson & Vagelos, 1968; Jung et al., 2016; Fig 3A). For our study we manipulated mtacp, the Drosophila melanogaster gene encoding mtACP, containing Ser-99, which is predicted to bind 4-phosphopantetheine (Ragone et al., 1999). In order to be able to identify and distinguish the two forms of mtACP (holo versus apo), we generated constructs encoding mutant proteins that would be refractory to 4-phosphopantetheinylation and observed their mobility differences using gel electrophoresis. We mutated the crucial serine residue: one to mimic the uncharged apo form (S99A) and two negatively charged forms (S99D and S99E) to mimic the charged holo form of mtACP (Fig 3A). Overexpression of wild-type mtACP constructs in Drosophila S2 Schneider cells enabled the visualization of protein bands that correspond to endogenous apo- and holo-mtACP forms. By comparing these bands to the apo-mimetic S99A and holo-mimetics S99D and S99E, we were able to prove the identity of the bands visualized under control and mtACP wild-type overexpressing conditions. Under physiologic conditions, endogenous holo-mtACP was detected (Fig 3B and C). In contrast, no endogenous apo-mtACP protein was visible, consistent with previous observations in other organisms that the inactive apo-mtACP form is not stable (Jackowski & Rock, 1983; Post-Beittenmiller et al., 1989).

We proceeded to investigate whether levels of active holo-mtACP would decrease upon CoA deprivation, a key assumption of our hypothesis. Treating S2 cells with the PANK inhibitor hopantenate (HoPan) leads to reduced CoA biosynthesis and to reduced levels of total CoA (Rana et al., 2010; Siudeja et al., 2011; Srinivasan et al., 2015). Under these conditions, we observed reduced levels of endogenous holo-mtACP (Fig 3C), and addition of CoA to the medium of HoPan-treated cells reverted this phenotype, consistent with previous studies in which administration of extracellular CoA is able to rescue phenotypes associated with reduced intracellular CoA biosynthesis (Rana et al., 2010; Siudeja et al., 2011; Srinivasan et al., 2015). These results demonstrate that impaired CoA biosynthesis is associated with decreased levels of 4'-phosphopantetheinylation of mtACP.

Protein lipoylation is reduced by impeding CoA biosynthesis

To further investigate the consequences of impaired CoA biosynthesis and reduced levels of 4'-phosphopantetheinylated mtACP, we examined mtACP-dependent processes, specifically those linked to MePAN and PDH-E2 deficiency. holo-mtACP and MECR are required for protein lipoylation (Fig 1), a process that is therefore affected in MePAN patients (Heimer et al., 2016). Four evolutionary conserved lipoylated enzyme complexes have been identified: PDH, α-KGDH, BCKDH, and GCV. To assess whether lipoylation was affected under conditions of impaired CoA biosynthesis, total protein lipoylation was analyzed using Western blot analysis. Incubation with an antibody that detects protein-bound lipoic acid revealed decreased levels of various protein bands under conditions of reduced CoA levels, an effect that was rescued when CoA was supplemented to the medium (Fig 3D, Appendix Fig S1). The availability of an antibody recognizing the Drosophila PDH-E2 subunit allowed the analysis of lipoylated PDH-E2 specifically, demonstrating that lipoylation of PDH-E2 was compromised in a CoA reduced background compared to control levels, an effect that was rescued by replenishing CoA (Fig 3D and E). Specific antibodies for the other lipoylated fly enzymes are lacking, but the observation that total protein lipoylation is reduced under conditions of impaired CoA biosynthesis suggests that the other lipoylated fly enzymes are similarly affected.

Pyruvate dehydrogenase activity decreases upon CoA deprivation

Because we were able to specifically document decreased PDH lipoylation under reduced CoA biosynthesis conditions, we predicted that PDH enzyme activity would be decreased. Lipoylation of PDH-E2 is essential for the subunit and complex to function, as the lipoil moiety binds and oxidizes the hydroxyl-ethyl compound derived from pyruvate and subsequently transfers it to its acceptor CoA to generate acetyl-CoA (Patel et al., 2014). To determine whether CoA deprivation indeed decreases PDH activity, we quantified PDH activity in control S2 cells and cells treated with HoPan (Fig 3F). We observed a significant reduction in PDH activity upon HoPan treatment compared to control cells, an effect rescued by addition of CoA to the HoPan-treated cells (Fig 3F). We assumed that, in the HoPan-treated cells, a fraction of the remaining pool of lipoylated PDH is partly inactivated by the presence of the endogenous PDH-inhibitor pyruvate dehydrogenase kinase (PDK). Therefore, we predicted that residual PDH activity could be increased by...
Figure 3.
interfering with the action of this inhibitory enzyme. To test this, we added dichloroacetate (DCA) to the medium of the HoPan-treated cells. DCA is a clinically used drug (Stacpoole et al, 1983, 1992) that inhibits the PDH inhibitor, PDK (Walsh et al, 1976; Patel & Korotchkin, 2006), and as such leads to activation of PDH (Whitehouse & Korotchkina, 2006), and as such leads to activation of PDH.

Our next step was to investigate a causal connection between CoA dyshomeostasis and PDH activity in a multicellular organism, Drosophila melanogaster, by asking whether phenotypes induced by impaired CoA biosynthesis could be rescued by recovery of PDH activity. To test this in whole organisms and complement our studies in Drosophila S2 cells, we used a genetic approach in flies, employing the binary UAS/GAL4 system (Brand & Perrimon, 1993; Dietzl et al, 2007) to knockdown dPANK/fbl using RNAi. dPANK/fbl is the Drosophila ortholog of human PANK2 (Afshar et al, 2001; Bosveld et al, 2008). In contrast to mammals, Drosophila possesses only one pank gene (Afshar et al, 2001). However, this gene gives rise to several pank isoforms, among which is one that localizes to mitochondria (Wu et al, 2009). Ubiquitous expression of a dPANK/fbl-RNAi construct in all tissues of the fly using an actin promotor resulted in significantly decreased dPANK/fbl mRNA and protein in two independent lines (Fig 4A, Appendix Fig S2), demonstrating the effectiveness of the RNAi constructs. Downregulation of dPANK/fbl resulted in fewer flies that reached the adult stage compared to control flies (Fig 4B, Appendix Fig S3). Impairment of dPANK/fbl in Drosophila results in decreased total CoA levels and reduced viability (Rana et al, 2010; Srinivasan et al, 2015). Addition of pantethine to the fly food, which restores total CoA levels in a dPANK/fbl-deprived background (Rana et al, 2010), fully restored their viability, further confirming the CoA dependency of this phenotype (Fig 4B). Next we investigated whether stimulation of PDH in the dPANK/fbl knockdown flies increased their viability as well. For this, we added DCA to the fly food and consistent with the results obtained from S2 cells, a dose-dependent restoration of viability was observed (Fig 4B). These results suggest that viability can be recovered by boosting the activity of PDH.

Downregulation of key steps of the CoA-PDH pathway during development causes a common abnormal morphogenetic phenotype in Drosophila wings

The dose-dependent rescue by DCA of the dPANK/fbl knockdown phenotype provides evidence that deleterious phenotypes associated with CoA deprivation are at least partially mediated via decreased activity of PDH. The proposed pathway (Fig 2) further predicts that impairment of individual components of this pathway would cause a phenotype with overlapping characteristics. To investigate this, we created a genetically and phenotypically tractable system. Using tissue-specific RNAi-mediated knockdown of various genes along the CoA-mtACP-PDH pathway, we selected the wing as our target tissue. Visible details of wing structure and morphogenesis are well-delineated and provide a robust and quantifiable phenotypic readout. Importantly, disruption of metabolism in specific cells of the developing wing does not cause lethality of the organism, as is common when critical metabolic pathways are disrupted in other organ systems or in whole organisms. During early Drosophila development causes a common abnormal morphogenetic pathway in Drosophila wings.

Figure 3. Decreased levels of CoA are associated with decreased levels of holo-mtACP and lipoylated PDH-E2.

A Schematic presentation of endogenous active and inactive forms of mtACP and synthesized mutant forms of mtACP. For the endogenous form, the S represents the serine residue that is 4'-phosphopantetheinylated, while D and L represent the flanking amino acids. Inactive form of mtACP (apo-mtACP) is indicated with 1. CoA is the source for 4'-phosphopantetheine and is required for 4'-phosphopantetheinylation of mtACP occurring on the serine residue. This posttranslational modification results in an active form of mtACP, holo-mtACP, which is negatively charged and indicated with 2. Three mtACP constructs were generated: One in which serine 99 was modified to an alanine, indicated with a 3 and indicated as S99A (non-4'-phosphopantetheinylatable form); one in which serine 99 was modified into aspartate, indicated with a 4 and indicated as S99D (phosphomimetic); one in which serine 99 was modified into glutamate, indicated with a 5 and indicated as S99E (phosphomimetic). S99D and S99E are negatively charged, mimicking the negatively charged holo-mtACP. The red circle indicates the presence of a negative charge.

B Western blot analysis of S2 cells overexpressing wild-type constructs of mtACP or the various mutant forms. First lane: lysates of control cells, resulting in the detection of holo-mtACP visible after long exposure. Second lane: overexpression (OE) of mtACP WT results in the detection of an apo-mtACP form and a holo-mtACP form, indicated with 1 and 2, respectively. Third lane: overexpression of mtACP S99A mutant form resulted in the detection of an apo-mtACP (non-4'-phosphopantetheinylatable) band only, indicated with 3. Fourth lane: overexpression of mtACP S99D resulted in the detection of a phosphomimetic form of mtACP only, migrating at the same mobility as holo-mtACP, indicated with 4. Fifth lane: overexpression of mtACP S99E results in the detection of a phosphomimetic form of mtACP only, migrating at the same mobility as holo-mtACP, indicated with 5. For visualization, a low exposure and high exposure blot are shown. Note that overexpressed bands required to visualize endogenous forms of mtACP.

C Western blot analysis of mtACP forms under control conditions, under conditions of HoPan treatment, and under conditions of HoPan + CoA treatment. Lanes showing overexpression of mtACP WT, mtACP S99A, and mtACP S99D were used to allow identification of the holo- and the apo-forms of mtACP. α-Tubulin was used as a loading control. Various exposure times of the blots are presented to allow identification of mtACP under all conditions.

D Western blot analysis showing lipoylated proteins under control conditions, after HoPan treatment and after HoPan + CoA treatment. S2 cells were treated with HoPan or HoPan + CoA for 4 days; non-treated cells were used as control. Antibodies specifically recognizing lipoylated proteins or PDH-E2 were used. Arrow heads indicate lipoylated PDH-E2 (left panel) or total PDH-E2 (right panel).

E Quantification of lipoylated PDH-E2 from (D). Mean ± SD is given. n = 3 for all samples.

F PDH activity was measured in control cells, HoPan-treated cells or HoPan-treated cells rescued with CoA or DCA. Mean ± SD of five biological replicates, each composed of three technical replicates and corrected for protein concentration.

Data information: For (E and F), two-tailed Student’s t-test was performed to calculate statistical significance for the indicated subsets. Source data are available online for this figure.
embryogenesis wing imaginal discs are formed, growing in size during later larval stages and finally forming adult wings during metamorphosis (Appendix Fig S4). Specific parts of the imaginal wing discs develop into specific parts of the adult wing. The larval hinge develops into the adult hinge and the larval wing pouch develops into the adult wing blade forming the majority of the wing structure (Appendix Fig S4; Diaz de la Loza & Thompson, 2017). Disruption of specific genes in parts of the wing during development leads to macroscopically visible wing abnormalities in the adult fly that can be readily scored. A plethora of mutant adult wing phenotypes have been reported, including holes, notches, vein abnormalities, blisters, increased/decreased size, and several others of the wing blade, depending on the specific gene that has been targeted by the RNAi (Bejarano et al., 2012). The severity of the blister phenotype varied between the knockdowns, most likely because of variations in effectiveness of the individual RNAi constructs. This is further substantiated because co-expression of the RNAi-enhancing element dcr2 with the RNAi construct of each of the three genes resulted in a stronger phenotype (Fig 5B, D, F and I). Combined RNAi-mediated knockdown of both dPANK/fbl and dPPCDC/ppcdc resulted in an increase of the blistering phenotype (Fig 5G) compared to that arising from downregulation of the single constructs alone (Fig 5C and E). This observation is consistent with the idea of incomplete RNAi-mediated knockdown of either component and suggests an additive phenotype. Together, these experiments validate the genetic knockdown system and implicate a common underlying cellular defect along the CoA-mtACP-PDH pathway leading to a similar phenotype, when the individual steps are compromised.

Genetic and pharmacologic stimulation of the pyruvate dehydrogenase complex rescues wing phenotypes caused by impaired CoA synthesis

The observation that knockdown of the individual genes dPANK/fbl, dPPCDC/ppcdc, and mtacp cause a phenotype with overt similarities is suggestive for the presence of an underlying common cellular defect. To substantiate this further, we investigated whether boosting PDH activity would rescue the wing-blisters phenotype induced

Figure 4. Downregulation of dPANK/fbl induces decreased viability of adult flies, which is rescued by pantetheine and DCA treatment.
A Decreased dPANK/Fbl protein levels were determined in two independent dPANK/fbl-RNAi strains (1 and 2), in which dPANK/fbl-RNAi was ubiquitously expressed. A wild-type strain was used as a control. An anti-dPANK/Fbl antibody was used to detect protein levels with Western blot analysis on whole fly extracts. Tubulin was used as a loading control.
B Survival rate of adult flies was determined for dPANK/fbl-RNAi flies, compared to control flies under normal food conditions and after the supplementation of dichloroacetate (DCA) or pantetheine to the food. The assay is explained and visualized in Appendix Fig S3. dPANK/fbl-RNAi flies show a decreased survival rate compared to control flies. DCA and pantetheine when added to the food increased viability of dPANK/fbl-RNAi flies. Total number of flies used for each group is indicated above each bar. Statistical analyses were carried out using Fisher’s exact test.

Source data are available online for this figure.
Figure 5.
by impaired CoA biosynthesis, comparable to the improved viability we showed in the dPANK/fl-RNAi flies. To investigate this we used the Drosophila line with the wing-specific downregulation of dPPCDC/ppcdc, in which the blistering phenotype is highly penetrant (in approximately 85% of the adult flies; Fig 6A). We assumed that, in the affected wing, comparable to the HoPAN-treated cells, a fraction of the remaining pool of lipoylated PDH is partly inactivated by the presence of the endogenous PDH-inhibitor pyruvate dehydrogenase kinase (PDK). In addition, PDH can be inactivated by endogenous SIRT4, a lipase that also inhibits PDH (Mathias et al., 2014). Therefore, we predicted that residual PDH activity could be increased by interfering chemically or genetically with the action of these inhibitory enzymes (Fig 6B and C). Indeed, we found that feeding the larvae DCA (PDK inhibitor) resulted in a dose-dependent decrease of wing blisters in the adult flies (Fig 6A). In addition, a genetic approach was used (Fig 6C, right part) to decrease the expression of PDK and SIRT4, by RNAi-mediated knockdown using two different lines per target. The knockdown efficacy of these constructs was verified and they were confirmed to cause no apparent phenotype in control flies (Appendix Fig S5). As predicted, downregulation of these PDH-inhibitors by RNAi in the dPPCDC/ppcdc-depleted background partly resolved the wing-blistering phenotype (Fig 6D). Non-specific genetic effects were excluded by using an RNAi line, unrelated to this pathway and from the same Drosophila RNAi library. This unrelated RNAi line in the dPPCDC/ppcdc-depleted background did not affect the blister phenotype (Appendix Fig S6). These results showing that boosting PDH activity can dampen the deleterious effects of impaired CoA biosynthesis and further establish the connection between CoA production and PDH activity.

**Downregulation of PANK2, the defective gene in PKAN, leads to a decrease in mtACP in human cells**

Our results in Drosophila show that defects in CoA biosynthesis lead to a decrease in holo-mtACP levels and decreased lipoylation and activity of PDH, suggesting that a final common pathway through PDH may explain the overlapping phenotypic features of PKAN, CoPAN, MePAN, and PDH-E2 deficiency. Several findings in mammalian systems further support our hypothesis: decreased PDH activity was recently demonstrated in a true PKAN mouse model by Jeong et al. (2019) and decreased lipoylation of PDH was demonstrated in MePAN patient fibroblasts (Heimer et al., 2016). While these findings support our hypothesis, a key player in this pathway, mtACP has not been investigated in mammalian systems with impaired PANK2 activity. Therefore, we generated HEK293T (human embryonal kidney, Appendix Fig S7) and SH-SYSY (human neuroblastoma, Appendix Fig S8) cells in which PANK2 protein levels were downregulated. We generated two independent clones per cell line in which, PANK2-RNAi expression was inducible by doxycycline and this led to decreased PANK2 protein. Efficiency of PANK2 downregulation was verified using Western blot and qPCR analysis. This demonstrated that PANK2 protein levels were undetectable in both cell lines, whereas PANK1, PANK3, and PANK4 RNA levels were unchanged, establishing the specificity of the PANK2 RNAi constructs (Fig 7A and C, Appendix Figs S9 and S10). Downregulation of the Drosophila dPANK/fl-RNAi gene is associated with an actual decrease in total CoA levels (Rana et al., 2010; Srinivasan et al., 2015), explaining the observed decrease in 4′-phosphopantetheinylation of mtACP. Therefore, we sought to determine whether downregulation of PANK2 in the generated clones led to a measurable decrease in total CoA levels. Total CoA levels were not significantly decreased in any cell lines compared to the controls after 10 days of RNAi treatment despite clear evidence of PANK2 protein loss (Fig 7E and F, Appendix Fig S11). Our results do not exclude the possibility that specific subcellular compartmental CoA levels may be decreased in these cells. Next we determined levels of human mtACP under conditions of downregulation of human PANK2. Levels of mtACP were significantly reduced in both PANK2-depleted neuroblastoma cell lines and in one of the two PANK2-depleted HEK293 cell lines compared to controls (Fig 7A–D). Since apo-mtACP is unstable and not detectable in other systems, the observed signal for mtACP is most likely the holo-(activated form of mtACP; Jackowski & Rock, 1983; Post-Beittenmiller et al., 1989). These results show that impaired CoA biosynthesis induced by downregulation of PANK2 in human cells is associated with a reduction of mtACP which occurs independent of any detectable change in total cellular CoA levels.

**Discussion**

Our results demonstrate that impairment of CoA biosynthesis leads to decreased levels of 4′-phosphopantetheinylated holo-mtACP, which causes decreased levels of lipoylated, activated PDH.
Figure 6.

A. Bar graph showing the percentage of blistered wings across different DCA concentrations.

B. Diagram illustrating the metabolic pathway involving Pyruvate, DCA, PDHc, PDK, and SIRT4.

C. Flowchart depicting pharmacological and genetic rescue strategies for dPPCDC/ppcdc-RNAi.

D. Bar graph comparing blistered wings in different conditions involving dPPCDC/ppcdc-RNAi and PDK RNAi.

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Stimulation of PDH activity in the setting of CoA dyshomeostasis rescued phenotypes ranging from viability to organ development, suggesting that impaired PDH function is at least partly responsible for the observed phenotypes. Our results implicate the linear pathway proposed in Fig 2 to explain why impairment at individual steps along this pathway results in overlapping phenotypes in both the Drosophila wing model and in the human disorders PKAN, CoPAN, MePAN, and PDH-E2 deficiency.

Here we show a specific and direct effect of impaired CoA biosynthesis on PDH activity. Other enzyme complexes (∅KGDH, BCKDH, and GCV) that are regulated by lipoylation (Cronan, 2016) are presumably affected as well, as indicated by decreased levels of pantothenylated protein levels (Fig 3D, Appendix Fig S1). The rescue by DCA and SIRT4 downregulation in Drosophila under CoA-compromised conditions indicates that at least some of the defects can be attributed to reduced PDH activity because boosting PDH activity causes specific defects to revert to wild-type phenotypes. SIRT4 regulates PDH activity via its lipomodiase activity. This activity of SIRT4 cleaves the lipoyl moiety from the E2 component of PDH and inhibits PDH activity (Mathias et al, 2014) explaining the observed rescue. In addition, SIRT4 also interacts with ∅KGDH-E2 and BCKDH-E2 (Mathias et al, 2014). Therefore, it could be possible that SIRT4 can negatively influence lipoylation of these other enzymes as well. Therefore inhibition of SIRT4 would increase lipoylation of ∅KGDH-E2, and BCKDH-E2 as well as of PDH-E2. Therefore recovery of activity of one or more of these enzymes is likely to explain the potent rescue observed after SIRT4 downregulation.

This rescue is somewhat surprising because boosting these enzyme activities still does not resolve decreased CoA levels or other effects of decreased levels of holo-mtACP. mtACP serves numerous functions that are conserved across species. Indeed, NDUFAB1, the human ortholog of mtACP, is a subunit of and required for the assembly of complex I (Vinokthumar et al, 2014; Van Vranken et al, 2018) and is involved in iron-sulfur biogenesis (Van Vranken et al, 2016). It is possible that the majority of the RNAi-induced phenotype in the Drosophila wing arises from reduced PDH activity and not from other affected processes downstream from holo-mtACP.

Our results demonstrate that, under conditions of decreased levels of total CoA such as in dPANK/flb-downregulated S2 cells or HoPan-treated samples, levels of holo-mtACP are decreased. This result is expected because CoA is the source for the 4′-phosphopantetheinylation and, by reducing its levels, the rate of posttranslational modification that depends on this source would be expected to decrease. A decrease in mtACP levels, most likely reflecting a decrease in active 4′-phosphopantetheinylated holo-mtACP, under conditions of normal total CoA levels, as in the PANK2-depleted human cells (Fig 7), is more challenging to explain. It is possible that in subcellular compartments like mitochondria, which contain PANK2 and COASY, CoA levels are decreased, inducing a reduction of holo-mtACP. It also may be that PANK2 depletion “only” lowers the rate of CoA de novo biosynthesis, which could trigger a compensatory mechanism inhibiting the formation of holo-mtACP since this process consumes CoA. The reduction of a consumptive process could explain the decrease in holo-mtACP and also could be reconciled with the results showing that total CoA levels remain constant. The prediction would be that under conditions of normal CoA levels but reduced holo-mtACP, complex I activity and iron-sulfur cluster formation would be decreased, which is in agreement with the results of Jeong et al in their Pank2-depleted mouse model. This further suggests that PKAN, and possibly also CoPAN, is caused by a reduced rate of CoA biosynthesis and not by an actual decrease in CoA levels. This is consistent with the findings that total CoA levels in CoPAN patient fibroblasts are also not reduced (Dusi et al, 2014). It is possible then that addition of 4′-phosphopantetheine to Pank2-depleted cells boosts the rate of CoA biosynthesis, thereby explaining its rescuing potential, as reported by Jeong et al (2019).

Our hypothesis can explain some of the phenotypic similarities and differences between PKAN, CoPAN, MePAN, and PDH-E2 deficiency. All these disorders selectively damage the basal ganglia, with most of the pathology localizing in globus pallidus. In PKAN and CoPAN, iron accumulates in this structure, but this phenomenon is not observed in the other two disorders (Fig 8). This difference can be explained by postulating that the iron accumulation stems from dysregulation of an intermediate downstream of PANK and COASY but upstream of MECR and PDH-E2. A candidate intermediate is again holo-mtACP; it was recently shown in eukaryotic cells that holo-ACP is involved in iron-sulfur cluster biogenesis and stability, highlighting a crucial additional role for the 4′-phosphopantetheine–conjugated mtACP (Van Vranken et al, 2016;
Figure 7.
Cory et al., 2017). In Saccharomyces cerevisiae, loss of mtACP leads to reduced iron-sulfur cluster formation, inactivation of iron-sulfur cluster-dependent enzymes such as aconitate, and activation of iron-responsive factors Ahl1 and Ahl2 (Van Vranken et al., 2016). Consistently decreased iron-sulfur-cluster levels result in mitochondrial iron overload (Chen et al., 2002). Abnormal iron homeostasis and reduced aconitate activity are characteristic of PKAN patient fibroblasts, as well as IPS-derived neurons (Santambrogio et al., 2015; Orellana et al., 2016). Iron dyshomeostasis and reduced aconitate activity are also reported by Jeong et al. in their PKAN mouse model (Jeong et al., 2019). This explains why mitochondrial iron dyshomeostasis and accumulation resulting from mtACP dysfunction is observed in PKAN and CoPAN, diseases associated with steps upstream of holo-mtACP, but not in MePAN and PDH-E2 deficiencies, diseases associated with steps downstream of holo-mtACP. Consistent with this, we predict that MECR deficiency compromises lipoic acid production without affecting mtACP levels or activity (Fig 8). Recently, Klopotstock et al. (2019) reported that iron chelation is effective in reducing the brain iron content in PKAN patients; however, the pathway proposed here would predict that iron chelation therapy alone would be insufficient to fully counteract neurodegeneration in patients with PKAN or CoPAN.

Our results and models complement and agree with the work of Jeong et al. and underscore the clinical relevance of our findings. Jeong et al. demonstrate, in a mouse model of PKAN, the presence of a specific set of perturbations in the globus pallidus. These alterations include: impaired complex I function with decreased oxidative phosphorylation, impaired lipoic acid production with loss of activity of a lipoylated enzyme (PDH), and iron dyshomeostasis with loss of activity of dependent enzymes and processes presumably from impaired iron-sulfur cluster biogenesis. All their reported findings are consistent with a primary defect in 4'-phosphopantetheinylated mtACP, including their observation of rescue of all molecular changes in mouse globus pallidus by administration of 4'-phosphopantetheine.

Despite providing a compelling hypothesis and strong results to support it, we recognize that the Drosophila model does not reflect the full complexity of the mammalian system, especially regarding the metabolic step compromised in PKAN. Downregulation of the single dpANK/β1 gene in fruit flies evokes a more severe phenotype compared to downregulation of only PANK2 in mammalian systems. Alternative hypotheses to explain decreased activity of PDH from a CoA metabolic defect should also be considered. One attractive hypothesis is based on the process of CoAlation, a recently identified posttranslational modification. A range of proteins was identified, including PDK, that can be reversibly modified by covalent attachment of CoA, which influences their activity. PDK activity is inhibited by CoAlation (Tsuchiya et al., 2017). Therefore, an alternative explanation for our results may be that under conditions of impaired CoA biosynthesis, levels of PDK-CoAlation are reduced, leading to increased activity of PDK and therefore decreased activity of PDH.

Our hypothesis does not explain why the globus pallidus is selectively damaged in the four diseases. For PKAN, it may be that the other PANKs compensate for the loss of PANK2 in areas outside the globus pallidus; however, this redundancy does not exist for CoPAN, MePAN, and PDH-E2 deficiency. Our hypothesis suggests that the globus pallidus is vulnerable to decreased activity of PDH; however, an explanation for this sensitivity is still lacking. The PKAN mouse model reported by Jeong et al. (2019) is a powerful tool to further investigate the molecular changes in mouse globus pallidus by administration of 4'-phosphopantetheine.
Figure 8.

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tool that may enable discovery of an answer to this important question.

Our model presented in Fig 8, in combination with the results provided by Jeong et al (2019) and Klopotock et al (2019), provides suggestions for possible therapies for the four neurodegenerative diseases. DCA is a blood-brain barrierpermeable drug (Abemayor et al, 1984; Kuroda et al, 1984) that has been investigated as a candidate therapeutic for metabolic diseases as well as cancer (Stacpoole et al, 1992; Stacpoole et al, 2006; Michelakis et al, 2010; Abdelmalak et al, 2013). Despite achieving biochemical remission of lactic acidosis in a genetically heterogeneous cohort of children with congenital lactic acidosis, DCA failed to slow neurological decline (Stacpoole et al, 2006). Moreover, while DCA rescues the Drosophila wing model, it would most likely not rescue iron dyshomeostasis or other dysfunction arising downstream of holo-mtACP such as of aconitase and complex I activities in PKAN and CoPAN patients. Based on the results by Jeong et al, 4'-phosphopantetheine is the most promising therapeutic strategy for PKAN because it restores an early step in the affected pathway and therefore would be predicted to prevent all downstream impairedd processes. In CoPAN patients, a CoA biosynthesis step is affected downstream of 4'-phosphopantetheine; therefore, we would not expect 4'-phosphopantetheineto prevent disease changes in CoPAN. However, based on our proposed pathway, an iron chelator in combination with DCA or a SIRT4 inhibitor may be beneficial in CoPAN. Finally, DCA or a SIRT4 inhibitor may also be beneficial in MePAN and to some extend in PDH-E2 deficiency, because DCA addition will not rescue severe loss of function mutations. Additional investigations in mammalian models are required to substantiate these ideas.

Materials and Methods

**S2 cell culture, transfection, HOPAN, and CoA treatment**

*Drosophila* Schneider’s S2 cells were maintained at standard conditions as described previously (Srinivasan et al, 2015). Here, cells in their exponential phase of growth were transfected (Effectene, Qiagen) with the mtACP WT or mutant constructs listed below and grown for 2 days.

HoPan and CoA or DCA treatment were done on S2 cells in their exponential phase and treated with 0.5 mM HoPan in the presence or absence of 100 μM CoA or 10 mM DCA. Cells were treated for 2, 4, or 7 days, and untreated S2 cells were used as control.

**Cloning of mtACP (mutant) constructs**

In order to overexpress wild-type or mutant mtACP constructs were created in the following manner. An mtACP cDNA clone for isoform B was obtained from the Drosophila Genomics Resource Center (AT22870; FBlc0033645) and multiplied by PCR using primers flanked by EcoRI and Xhol restriction sites.

The pAcs5.1 vector (Invitrogen) was digested using EcoRI and Xhol and ligated with the mtACP PCR product. Competent cells were transfected with the ligated construct and the purified construct was sequenced to ensure its fidelity. Constructs overexpressing mutant mtACP were subsequently created by site-directed mutagenesis of this construct using mutagenesis primers (Q5 Site-directed Mutagenesis Kit, New England Biolabs). The fidelity of the resulting constructs was verified by sequencing. Primers sequences are presented in Appendix Table S1.

**Lentiviral transductions**

Inducible lentiviral shRNA vectors targeting a non-targeted control (NTC) or hpANK2 were obtained from Dharmacon (for sequences, see supplementary data) and lentivirus was produced as previously described (Schepers et al, 2007). 293T cells (ATCC CRL-3216) and SH-SY5Y cells (ATCC CRL-2266) were transduced in 2 consecutive rounds of 8–12 h with lentiviral supernatant supplemented with 10% FCS and polybrene (4 μg/ml; Sigma). Transduced 293T and SH-SY5Y cells were selected in a medium containing 0.6 μg/ml or 0.2 μg/ml puromycin (Thermo Fischer Scientific), respectively, for 1 week.

**FACS analysis**

After 1 week of puromycin selection, expression of the microRNA-TurboGFP cassette was induced with doxycycline (Sigma, concentrations ranging from 0 to 1.5 μg/ml) for 3 days and analyzed for TurboGFP expression on an LSRII (Becton Dickinson) flow cytometer, and data were analyzed using FlowJo software (FlowJo V10).

**Inducible hPANK2 knockdown cells**

Cells were produced as described above. HEK293T and SH-SY5Y inducible hPank2 KD cells were maintained in Dulbecco’s Modified Eagle’s Medium (DMEM, Sigma) supplemented with 10% fetal bovine serum (FBS, Greiner Bio-one) and antibiotics (penicillin/streptomycin, Invitrogen) in 5% CO2 at 37°C. Induction of the microRNA-TurboGFP cassette in HEK293T was done with 1 μg/ml doxycycline for 14 days, in custom made DMEM without vitamin B5 (Thermo Scientific) supplemented with dialyzed FBS (Thermo Scientific) and antibiotics. HoPan treatment (0.25 mM) was done from day 7 till day 14.

A complete list of vectors used, commercially available cell lines, and lines created in this publication can be found in Appendix Tables S1 and S2. The cell lines were regularly visually inspected and tested negative for mycoplasma contamination.

**Western blot analysis and antibodies**

For Western blot analysis, cells were dissolved in 2× Laemli buffer, sonificated, and boiled for 5 min with 5% β-mercaptoethanol (Sigma). Protein concentration was determined using DC protein assay (Bio-Rad). Equal amounts of protein (10–30 μg) were loaded on a 10, 12, or 4–20% gradient gel (Bio-Rad), and transferred onto PVDF membranes using the Trans Blot Turbo System (Bio-Rad). Membranes were blocked in 5% fat-free milk for 1 h at room temperature, and then rinsed in PBS-Tween 20. Incubations with primary antibodies were done overnight at 4°C followed by incubations with HRP-conjugated secondary antibodies (Amersham 1:5,000) for 1.5 h at room temperature. Detection was performed using ECL reagent (Thermoscientific) and visualized using the
ChemiDoc imager (Bio-Rad). The following primary antibodies were used: anti-mtACP antibody (Abcam, 1:1,000), anti-lipoic acid (Merck, 1:1,000), anti-PDH-E2 (Abcam, 1:1,000), anti-α-Tubulin (Sigma, 1:5,000), anti-GAPDH (Fitzgerald, 1:10,000), anti-dPank/Fbl (Bosveld et al., 2008; 1:1,000), and anti-hPANK2 (Origene 1:500). A complete list of antibodies used can be found in Appendix Table S3.

**PDH activity measurements**

S2 cells were cultured as described above for 4 days in control medium or medium containing 500 μM HoPan. Cells were pelleted at approximately 10^6 cells/pellet and washed once with PBS, and later pyruvate dehydrogenase complex activity was measured using the Pyruvate Dehydrogenase Activity Colorimetric Assay Kit (Cat#K679-100, BioVision) according to manufacturer’s instructions. Three biological replicates were used per measurement, with each biological replicate measured in (technical) triplicate. Protein concentration was determined using BCA Protein Assay Kit (ThermoScientific) according to manufacturer’s instructions. All measurements were recorded using a VarioSkan Lux plate reader; analysis was performed with GraphPad (see section “Statistical analysis”).

**Drosophila maintenance and genetics**

*Drosophila melanogaster* stocks were maintained on standard cornmeal agar fly food (containing water, agar 17 g/l, sugar 54 g/l, yeast extract 26 g/l, and nipagin 1.3 g/l) at 22°C. Crosses were raised at various temperatures as indicated in the text/legends. The stocks were either obtained from the Bloomington Stock Centre (Indiana University, USA) or the VDRC (Vienna *Drosophila* RNAi Collection, Vienna, Austria).

**Stocks used for this project**

A complete list of commercially available fly lines, lines created in the context of this publication, and detailed genotypes can be found in Appendix. Crosses were raised at various temperatures as indicated in the text/legends.

**Validation of UAS-dPANK/fbl-RNAi and UAS-dPPCDC/ppcdc-RNAi lines by meiotic recombination**

Based on previous results (expression with various drivers, data not shown), we predicted that our UAS-dPANK/fbl-RNAi (#101437) and UAS-dPPCDC/ppcdc-RNAi (#104495) KK lines, obtained from the VDRC, might contain offspring at location 30B, but an additional one at 40D, which has been shown to act as a phenotypical enhancer and able to cause false-positive results, especially when combined with ubiquitous or wing drivers (Green et al., 2014; Vissers et al., 2016).

To test those lines for RNAi transgene integration site occupancy, we used genomic DNA preparation, followed by PCR analysis, as described previously (Vissers et al., 2016). When both lines were confirmed to carry transgenes at integration sites 30B and 40D, we employed meiotic recombination to remove the one at 40D. Lines with the remaining insertion at 30B were confirmed by PCR (Appendix Fig S1A) and tested for knockdown by immunohistochemistry (anti-Fbl, Fig 4A) and qPCR (fbl and PPCDC, Appendix Fig S1B and C). The “30B-only” cleaned-up lines were used for all experiments presented in this publication unless stated specifically otherwise (also see stock list in the Appendix for details).

**Immunofluorescence analysis of *Drosophila* wing imaginal discs**

For immunofluorescence of *Drosophila* wing discs, the crosses were raised at 29°C (MS1096-GAL4; UAS-GFP × UAS-GFP as control or x UAS-dPANK/fbl-RNAi) or 22°C (MS1096-GAL4; UAS-GFP × UAS-GFP as control or x UAS-mtACP-RNAi #29528). Wandering L3 larvae (day 5) were collected and their wing discs dissected in ice-cold phosphate-buffered saline (PBS). The discs were fixed with 4% formaldehyde (Thermo Scientific Pierce) for 30 min, washed for 3 × 20 min with phosphate-buffered saline (PBS) + 0.1% Triton X-100 (Sigma Aldrich), and afterward incubated in primary antibody rabbit anti-Fbl (Bosveld et al., 2008; 1:500) or rabbit anti-mtACP (ThermoFisher PA5-22191, 1:500) in PBS + 0.1% Triton X-100 overnight to visualize the presence/absence/localization of Fbl or mtACP. After an additional washing step of 3 × 20 min in PBS + 0.1% Triton X-100, the discs were incubated in secondary goat anti-rabbit-Alexa488 antibody (Molecular Probes) for 2 h at room temperature. DAPI (0.2 μg/ml; Thermo Scientific) was used to visualize DNA. Finally, the samples were mounted in 80% glycerol and analyzed using a Zeiss LSM780 NLO confocal microscope with Zeiss software. Adobe Photoshop and Illustrator (Adobe Systems Incorporated, San Jose, California, USA) were used for image assembly. A complete list of antibodies used can be found in Appendix Table S3.

**Drosophila survival rate experiments**

Virgin female Actin-GAL4/CyO flies were crossed with males of either UAS-GFP or UAS-dPANK/fbl-RNAi at 29°C on standard fly food, or food supplemented with either sodium dichloroacetate (DCA, Sigma) or pantethine (Pan, Sigma) at indicated concentrations. The flies were allowed to lay eggs for 5 days, after which the adults were discarded. The emerging adult (eclosing) male flies were evaluated for the presence or absence of the CyO dominant marker. Non-CyO flies were selected because only in these flies the dPANK/fbl-RNAi (Actin-GAL4/UAS-dPANK/fbl-RNAi) or the GFP (Actin-GAL4/UAS-GFP, as control) is expressed. Progeny were counted daily over a period of 5 days to evaluate the total survival rate of the adult flies. To determine the survival rate of dPANK/fbl-RNAi flies, which is an indicator for the viability, the number of male Actin-GAL4/UAS-dPANK/fbl-RNAi flies was divided by the total number of male flies (CyO and non-CyO). At least 6 separate vials containing offspring were used per condition.

**Mounting and imaging of adult fly wings**

To image wings of adult flies from various crosses, F1 males or females of the indicated genotypes were collected for a period of 3 days and kept for an additional 2–3 days after eclosion to allow for optimal unfolding and clearance of the wings. Afterward, they were transferred into 70% ethanol and stored for at least 2–3 days. The wings were removed with tweezers, mounted on slides in 80% glycerol, and imaged with a light microscope (Olympus BX-50) at 2× magnification. Adobe Photoshop and Illustrator (Adobe Systems...
Incorporated, San Jose, California, USA) were used for visualization.

Pharmacological and genetic rescue of dPPCDC/ppcdc-RNAi-induced wing blisters

For the pharmacological rescue, MS1096-GAL4, UAS-GFP or MS1096-GAL4, and UAS-dPPCDC/ppcdc-RNAi females were crossed with UAS-GFP males at 29°C on standard fly food, or food supplemented with sodium dichloroacetate (DCA, Sigma) at indicated concentrations.

For the genetic rescue, MS1096-GAL4, UAS-GFP or MS1096-GAL4, and UAS-dPPCDC/ppcdc-RNAi females were crossed with UAS-GFP, UAS-PDK-RNAi (#28635 and #35142), or UAS-SIRT4-RNAi (#33984 and #36588) males at 29°C on standard fly food.

The females were allowed to lay eggs for 5 days, after which the parents were discarded. F1 males were collected and imaged as described above and analyzed for the presence or absence of “blisters”.

The adult Drosophila wing consists of two layers, a dorsal (upper) and a ventral (lower) one, which are tightly connected to each other (by adhesion molecules). “Blistering” of the wing has been described in general as a condition in which dorsal and ventral wing surfaces separate to form a hemolymph-filled blister (Martin et al, 1999). After mounting and imaging, they appear transparent and the remains of the blister are only detectable by their crinkled surface (Fig 5D and E). In blisters that encompass the whole wing, hemolymph (and associated cell debris) is not cleared sufficiently and after mounting and imaging these wings appear inflated with brownish discolorations (Fig SF–I).

Arrow heads were used to mark the perimeter of the blisters, which in some cases does equal the whole wing. In comparison, wings were scored as “not blistered”, if they appeared flat, structurally normal, and/or no discoloration was observed (Fig 5A and B). For visualization and quantification, images were recorded, using Adobe Photoshop and Illustrator (Adobe Systems Incorporated, San Jose, California, USA) and GraphPad Prism software (GraphPad Software, San Diego, CA, USA). For randomly chosen experiments, the amount of blisters was counted by independent researchers. Blinding was not performed.

qPCR

For quantitative real-time PCR (qPCR), we collected 8 specimens from fly samples, in most cases adult flies, from crosses of Actin-GAL4 with UAS-dPANK/fbl-RNAi F10 and F20, UAS-dPPCDC/ppcdc-RNAi P7 and P17, UAS-PDK-RNAi #28635 and #35142, and UAS-SIRT4-RNAi #33984 and #36588. The cross of Actin-GAL4 with both UAS-dPPCDC/ppcdc-RNAi results in mid- to late-pupal lethality, so we collected those samples during pupal development. The samples were crushed with a motor pestle, lysed in Trizol (Life Technologies), and RNA isolation was performed according to standard protocol. The isolated RNA was treated with Turbo DNase (Ambion) followed by addition of random primers and reverse transcription using M-MLV (Invitrogen). Quantitative PCR was performed using Sybr Green (Bio-Rad) on a CFX Connect Real Time System (Bio-Rad). Quantitative PCR was performed using primer sets to detect dPANK/fbl, dPPCDC/ppcdc, PDK, and SIRT4 mRNA expression, and normalized against Rp49 mRNA-loading controls. qPCR was performed in the human cells to determine PANK1-4 levels. cells were collected and lysed in RLT and RNA isolation was performed using the RNeasy kit from Qiagen according to the manufacturer’s instructions. Fly samples were normalized for rp49, and human samples were normalized for B2M and UBC.

A complete list of qPCR primers can be found in Appendix Table S2.

Statistical analysis

Densitometry and statistical analysis

Western blot analyses were measured using Image Studio Lite (Li-Cor, www.licor.com/bio/image-studio-lite). Statistical analyses were performed using Excel 2016 or GraphPad PRISM 5 (San Diego, CA, USA), with either unpaired Student’s t-test or Fisher’s exact test. Statistically significant results are depicted within each figure. Unless stated specifically in the figure legend, all the data were presented as mean ± SD.

For proportional data, power calculations for comparing proportions were used. F-tests were used to test if variances of the tested populations were equal. The number of flies assured a statistical power of 0.8 with a confidence level of 0.95 for Fisher’s exact test and Student’s t-tests (proportional power calculations; Wang & Chow, 2007), taking into account that the expected proportion of responders is around 25–80%.

Expanded View for this article is available online.

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Author contributions

RAL, YY, HS, BMB, MAV-L, MAT, SJH, NAG, and OCMS designed the research studies, analyzed data, and wrote the manuscript. RAL, YY, HS, NAG, MAV-L, KJA, MAT, and MvdZ conducted experiments, and acquired and analyzed data. KJA acquired samples. MvdZ, RAL, and HS generated essential reagents.

Conflict of interest

Dr. Hayflick and Dr. Sibon are co-inventors on a patent application for 4'-phosphopantetheine for use in disorders exclusive of PKAN. Dr. Sibon is a co-inventor on a patent application for acetyl-4'-phosphopantetheine for use in PKAN and in related disorders. Dr. Hayflick is a non-compensated member of the Scientific Advisory Board of BioPontis Alliance, a non-profit organization. Dr. Hayflick is a non-compensated member of the Scientific and Medical Advisory Board of the NBIA Disorders Association, a non-profit lay advocacy organization. Dr. Hayflick is a non-compensated member of the Scientific and Medical Advisory Board of the NBIA Alliance, a non-profit lay advocacy
The paper explained

Problem
PKAN, CoPAN, MePAN, and PDH-E2 deficiency are neurodegenerative diseases caused by mutations in four different genes for enzymes important in distinct metabolic pathways. The four diseases selectively damage the basal ganglia, which suggests that they share a common metabolic pathogenesis. Identifying this shared metabolic defect is essential to understanding their pathophysiology and designing effective therapeutics for all the four diseases.

Results
Fruitfly and mammalian cell systems were used to model the diseases and reveal a shared metabolic pathway. We present evidence that links the pathways of coenzyme A biosynthesis to the activity of pyruvate dehydrogenase (PDH) via mitochondrial acyl carrier protein. Chemical and genetic strategies were employed to boost PDH activity, which is a final common defect in the four disorders, leading to correction of defects observed in our disease models.

Impact
Our results provide an explanation for the overlap in clinical phenotypes of PKAN, CoPAN, MePAN, and PDH-E2 deficiency and suggest treatment strategies for all four diseases.

References

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