The phosphatidylinositol transfer protein PITP-1 facilitates fast recovery of eating behavior after hypoxia in the nematode Caenorhabditis elegans

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Abstract
Among the fascinating adaptations to limiting oxygen conditions (hypoxia) is the suppression of food intake and weight loss. In humans, this phenomenon is called high-altitude anorexia and is observed in people suffering from acute mountain syndrome. The high-altitude anorexia appears to be conserved in evolution and has been seen in species across the animal kingdom. However, the mechanism underlying the recovery of eating behavior after hypoxia is still not known. Here, we show that the phosphatidylinositol transfer protein PITP-1 is essential for the fast recovery of eating behavior after hypoxia in the nematode Caenorhabditis elegans. Unlike the neuroglobin GLB-5 that accelerates the recovery of eating behavior through its function in the oxygen (O2)-sensing neurons, PITP-1 appears to act downstream, in neurons that express the mod-1 serotonin receptor. Indeed, pitp-1 mutants display wild-type-like O2-evoked-calcium responses in the URX O2-sensing neuron. Intriguingly, loss-of-function of protein kinase C 1 (PKC-1) rescues pitp-1 mutants’ recovery after hypoxia. Increased diacylglycerol (DAG), which activates PKC-1, attenuates the recovery of wild-type worms. Together, these data suggest that PITP-1 enables rapid recovery of eating behavior after hypoxia by limiting DAG’s availability, thereby limiting PKC activity in mod-1-expressing neurons.

KEYWORDS
C elegans, GLOBIN 5, hypoxia, oxygen sensing neurons, PITP-1, PKC-1

Abbreviations: AMS, acute mountain syndrome; C, elegans; Caenorhabditis elegans; DAG, diacylglycerol; GLB, globin; IP3, inositol 1,4,5-trisphosphate; NPR-1, neuropeptide receptor 1; PITP, phosphatidylinositol transfer protein; PLCβ, phospholipase C beta; PMA, phorbol 12-myristate 13-acetate; PKC-1, protein kinase C 1; PtdIn, phosphatidylinositol; PtdOH, phosphatidic acid; sGC, soluble guanylate cyclases.

Zohar Abergel, Maayan Shaked, and Virendra Shukla contributed equally to the work.

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1 INTRODUCTION

Eating behavior influences our food choice, meal timing, and quantity, and therefore, determines our health and disease state. The factors affecting eating behavior are numerous and diverse and include internal and external cues, one of which being oxygen (O2). O2 is essential for energy production in all aerobic animals, and therefore, insufficient O2 supply (hypoxia) is a significant stress condition in most animals, including humans. Among the fascinating responses to hypoxia is the suppression of food intake and, consequently, body mass loss. This phenomenon is called high-altitude anorexia (or hypoxia-anorexia) and is observed in people suffering from acute mountain syndrome (AMS). AMS occurs in response to a rapid exposure to moderate or high altitudes (above 4000 meters), where the partial O2 pressure is low. Results from Operation Everest III (Comex-‘97) demonstrated that in a controlled environment that mimicked the decrease in O2 concentration while climbing on Everest at 5000 to 8000 meters, hypoxia was the stimulus that changed the eating behavior of volunteers and decreased food intake due to a loss in appetite. Intriguingly, the hypoxic-anorexia phenomenon appears to be conserved in evolution and has been observed in crabs, flies, gastropods, fish, rodents, and in the nematode Caenorhabditis elegans (C. elegans). The hypoxic-anorexia effect is reversible. Indeed, data from Comex-‘97 show that upon returning to ambient O2-levels, the volunteers’ body mass increases. However, the mechanism underlying the recovery of eating behavior after hypoxia is still poorly understood.

The eating behavior of C. elegans is modulated by neuronal O2-sensors and neuropeptide signaling. Animals bearing the dominant neuropeptide G-protein-coupled receptor npr-1(215V) allele (e.g., N2 laboratory strain) move slowly on bacteria (the C. elegans food source) and do not clump together on the bacterial lawn border at 21% O2. By contrast, animals bearing either the recessive npr-1(215F) allele (found in many wild C. elegans isolates, e.g., the Hawaii strain CB4856) or the strong loss-of-function allele npr-1(ad609), hereafter referred to as npr-1(-) worms), move fast on food and clump together on the bacterial lawn border (bordering eating behavior) at 21% O2. The bordering behavior is dependent on the activity of the atypical soluble guanylate cyclases (sGCs) O2-sensors GCY-35 and GCY-36. Upon O2 binding, the GCY-35/GCY-36 functional complex is thought to generate cGMP that triggers the opening of the TAX-2/TAX-4 cyclic nucleotide-gated channel complex. The opening of TAX-2/TAX-4 facilitates calcium (Ca2+) entry, and thus, the depolarization of the O2-sensing neurons AQR, PQR, and URX. This cascade activates an escape response from hypoxia (O2 > 12%), which is suppressed by the activity of NPR-1(215V) in the presence of food. Intriguingly, the clumping of worms on the bacteria lawn border appears to decrease the O2 level that the worm experience. Indeed, a previous study demonstrated that the O2 level inside the clump ranges between 6.4% and 13.7% O2, and this is the range of concentrations to which the worm is attracted.

Hypoxia (1% O2) suppresses the bordering eating behavior and results in worms accumulating outside the bacterial lawn border (Figure 1A). This food-leaving behavior is reversible upon returning to 21% O2 and is regulated by the neuroglobin GLB-5. Similar to npr-1, glb-5 is a polymorphic gene that affects the worm bordering behavior and foraging speed. In the case of the npr-1-gene, N2 worms bear the dominant npr-1(215V) gain-of-function allele and the Hawaii strain CB4856 bears the recessive npr-1(215F) allele. In contrast, with the glb-5 gene, N2 worms bear the Bristol allele, hereafter referred to as glb-5(-), which encodes a truncated neuroglobin that appears to be non-functional. By contrast, CB4856 worms bear the Hawaii allele of glb-5, hereafter referred to as glb-5(+), which encodes a full-length functional neuroglobin that appears to represent the ancestral glb-5 allele, as it is found in most C. elegans wild-isolates. GLB-5(+) fine-tunes the O2-responses of worms close to 21% O2. Indeed, npr-1(-) worms that bear the glb-5(-) allele (hereafter referred to as glb-5(-);npr-1(-) worms) are more sensitive to subtle changes in O2 level and sharply decrease their foraging speed in response to a 21% to 17% O2 shift, whereas npr-1(-) worms (bearing the glb-5(-) allele) display only a mild decrease in response to this stimulus. Like the mammalian cytoglobin and neuroglobin, GLB-5 is a hexacoordinated globin. Hypotheses for hexacoordinated globins' in vivo functions include regulation of cellular signaling and protection against oxidative damage.

GLB-5 controls worms' speed and recovery from hypoxia through its activity in the O2-sensing neurons AQR, PQR, and URX. The recovery of the bordering behavior in glb-5(++; npr-1(-)) worms, is rapid and occurs within minutes after returning to 21% O2. By contrast, the recovery of npr-1(-) worms bearing the glb-5(-) allele is slow and is completed only after 4 hours. This significant difference in recovery time has facilitated the isolation of several suppressor mutations that attenuate bordering behavior recovery by suppressing GLB-5 activity in the O2-sensing neurons. That is, glb-5(++; npr-1(-)) animals bearing these mutations do not slow down in response to a 21% to 17% O2 shift and display high URX-Ca2+ levels at both 14% and 17% O2. Here, we discover a new type of suppressor mutation in the gene encoding for the phosphatidylinositol (PtdIn) transfer protein PITP-1, which attenuates the recovery of bordering eating behavior without affecting the function of GLB-5 in the O2-sensing neurons. Our data suggest that PITP-1 facilitates fast recovery of eating behavior after hypoxia by restricting the activation of protein kinase C 1 (PKC-1) by diacylglycerol in mod-1-expressing neurons. This function is distinct from the role of PITP-1 in salt chemotaxis.
2 | MATERIALS AND METHODS

2.1 | Strains

**EVG009** glb-5(Haw) V; npr-1(ad609) X (the parental strain is AX1891)

**EVG034** npr-1(ad609) X (the CGC name is DA609)

**EVG155** glb-5 (Bri) V; npr-1(ad609) X; Ex[Pgly-37::YC2.60 + ccRFP]

**EVG159** glb-5(Haw) V; npr-1(ad609) X; Ex[Pgly-37::YC2.60 + ccRFP]

**EVG270** pitp-1(heb2) III; glb-5(Haw) V; npr-1 (ad609) X

**EVG419**, pitp-1(tm1500) III; glb-5(Haw) V; npr-1(ad609) X; heb21Ex[Ppitp-1::pitp-1(cDNA)::polycismCherry;P-f15e11.1::GFP]

**EVG430** pitp-1(tm1500) III; glb-5(Haw) V; npr-1(ad609) X; heb20Ex[Ppitp-1::pitp-1(gDNA)::polycismCherry;P-f15e11.1::GFP]

**EVG451** pitp-1(tm1500) III; glb-5(Haw) V; npr-1(ad609) X; heb22Ex[Pgly-37::pitp-1(cDNA)::polycismCherry;P-f15e11.1::GFP]

**EVG453** pitp-1(tm1500) III; glb-5(Haw) V; npr-1(ad609) X; heb23Ex[Pfpl-6::pitp-1(cDNA)::polycismCherry;P-f15e11.1::GFP].
Worm synchronization

To generate synchronized worms, we collected eggs from gravid hermaphrodites using hypochlorite solution. In brief, we collected gravid hermaphrodites into 15 mL tubes by washing the NGM plates three times with M9 buffer (22 mM KH2PO4, 42 mM Na2HPO4, 86 mM NaCl, and 1 mM MgSO4). We centrifuged the tubes for 1 minute (1690 g, 1 min), and removed the supernatant until 1 mL of volume remained. Then, we added 1 mL of hypochlorite solution (0.5 N NaOH, 1.25% NaOCl) to each tube and inverted it five times. To assist the release of embryos, we aspirated the worm suspensions (back and forth several times) using a syringe with a 21-gauge needle, for 3 minutes. Then, we sedimented the embryos by centrifugation (1690 g for 2 minutes) and removed most of the hypochlorite solution. Each tube was washed three times with 5 mL of M9 buffer. Next, we removed most of the M9 buffer (without disturbing the embryos pellet) and added 2 mL of fresh M9 buffer to each tube. We rotated the tubes for 16 hours at room temperature (RT, 21°C). The hatched L1 larvae were collected by centrifugation (1690 g for 3 minutes), counted, and placed into NGM plates seeded with OP50 E.coli bacteria and grown until the desired developmental stage.

Bordering behavior assays

Bordering experiments were performed, as described previously. In brief, 40 L4 hermaphrodites, grown in 21% O2/21°C, were transferred into a 3.5 cm NGM plate that was seeded 2 days before with 50 µL OP50 bacteria (OD600 ~ 0.6). The assay plates were put in 1% O2 (Coy hypoxia chamber, COY Lab Products Inc, Grass Lake, MI, USA, RT) for 24 hours, and then, brought back to 21% O2 (control plates were maintained in 21% O2 at 21°C). We calculated the bordering index (BI) after 30 minutes recovery at 21% O2. BI (%) represents the fraction of worms found on the bacterial lawn border divided by the total amount of worms on the plate, multiplied by 100%.

Speed measurements

All speed measurements were done in the presence of food as described in detail in Abergel et al. In brief, 2 days before the experiment, we put 20 µL OP50 bacteria (OD600 ~ 0.6)
on each low-NGM plate (0.13 g/L bacto-peptone) and incubated the plates at RT. For each speed experiment, we trapped 8-10 young hermaphrodites inside 500 µm deep rectangular PDMS (12 mm wide, 0.5 mm deep, and 17 mm long). We used a PHD 2000 syringe pump (Harvard Apparatus) to deliver humidified gases to the microfluidic chambers at a flow rate of 0.5 mL/min. We used Teflon valves, controlled by a ValveBank Controller (Automate Scientific), to rapidly switch between different O₂/N₂ gas mixtures. We recorded the movies using a Q-Imaging MicroPublisher 5.0 RTV Microscope Camera (QImaging, RHos) mounted onto an Olympus SZ61 stereomicroscope (Olympus). The videos were taken at 0.5 frames/s. We used custom MATLAB software (The MathWorks) for video analysis. Notably, in our analysis, we did not track worms that leave the bacterial lawn, and thus, only calculated the speed of worms on food.

2.5 | Salt chemotaxis

Salt (NaCl) chemotaxis assays were performed as previously described, with some modifications. In brief, a day before the experiment, we generated a NaCl gradient on a 9-cm chemotaxis assay plate containing 10 mL of 2% agar, 5 mM potassium phosphate (pH 6.0), 1 mM CaCl₂, and 1 mM MgSO₄. To generate the gradient, we placed a 1.2 cm-diameter agar plug, containing 100 mM of NaCl, 1 centimeter from the edge of the plate. We put one microliter of 0.5 M NaN₃ to the peak of NaCl gradient (the NaCl agar plug) and the equivalent place on the opposite side of the plate. We placed ~200 synchronized young adult hermaphrodites at the center of the assay plate. After 30 minutes, we counted the worms at the NaCl peak and the control spot (1 cm radius from the NaN₃ drop). A chemotaxis index was calculated as followed: [(# of animals at the NaCl peak)—(# of animals at the control spot)]/[(total # of animals)—(# of animals at the center of the plate)]. Each data point represents at least six biological repeats.

2.6 | Phorbol 12-myristate 13-acetate treatment

Two days before the experiment, we prepared NGM plates containing 100 µg/mL of ampicillin and 0.1 µg/mL of phorbol 12-myristate 13-acetate (PMA) (dissolved in DMSO, p8139 Sigma Aldrich) or 0.001998% of DMSO as a control. After 24 hours at room temperature (RT), we seeded the plates with 50 µL ampicillin-resistant-OP50 bacteria (OD600 ~ 0.6) and let them dry for 24 hours at RT. At the day of the experiment, we placed 40 L4 staged hermaphrodites on the plates and transferred them to 1% O₂ (Coy hypoxia chamber, COY Lab Products Inc, Grass Lake, MI, USA, RT) for 24 hours; control plates remained at 21% O₂. Bordering recovery was measured as described above.

2.7 | Ethyl methanesulfonate mutagenesis screen

The ethyl methanesulfonate (EMS) screen was performed as described in detail in Abergel et al.

2.8 | Mapping

We mapped the heb2 mutation to a 1.6 Mb genomic interval (on chromosomes III) using SNPs (single nucleotide polymorphisms). The list of additional SNPs primers we used is presented in Table 1. The molecular identity of heb2 was determined by next-generation sequencing (NGS) at the Technion Genome Center.

2.9 | Molecular biology

Genotyping: The glb-5(Bri) duplication was followed by examining the PCR products amplified using primers that flank the duplication in this allele. The npr-1(215V) and npr-1(ad609) alleles were followed by amplifying the interval containing the mutated allele, and MwoI digestion that distinguishes between the 215V and ad609 alleles. The pitp-1(tm1500), pkc-1 (ok563), and the dgk-1(ok1462) deletion were followed by PCR using primers that flank the deletion region.

2.10 | Transgenes

All rescuing plasmids were generated by either restriction enzyme or Gibson assembly cloning using modified polycistronic mCherry pPD95.75 expression vector, as described previously.

For the pitp-1(genomic) rescuing construct [PPitp-1::pitp-1(gDNA)::polycismCherry], we amplified the pitp-1 promoter (~4.2 Kb), and genomic region (~8.2 kb) using N2 genomic DNA. The promoter region was amplified with 5′ GTTATCCGCTCCAATCCATCCTTCCAT 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CCTCTTCTTTTTTCGACTCGACAACCCAG 3′ and inserted using BspEI and AvrII. The genomic sequence of pitp-1 was cloned in two steps. The first genomic part was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and
inserted using PacI and NotI. To create the pitp-1 cDNA (−3.1 kb) [Ppitp-1::pitp-1(cDNA)::polycismCherry] rescuing construct, the genomic part of the pitp-1 gene was replaced with pitp-1 cDNA. The coding region of pitp-1 was amplified using 5′ ATGCTGATCAAAGAATACCGTATCCTGTTAC 3′ and 5′ CTACCGACGATCTTTTTCATTTTCGAATT 3′ and inserted using AvrII and NotI. To express the pitp-1 cDNA in the oxygen sensing neurons we cut the gcy-37 and glb-5 promoters regions from an existing plasmids using BspEI and AvrII, these promoters regions (~1.3 kb, ~3.1 kb, respectively) replace the pitp-1 promoter, and thus, generate the [Pgcy-37::pitp-1(cDNA)::polycismCherry] and [Pglb-5::pitp-1(cDNA)::polycismCherry] expression constructs. To express the pitp-1 cDNA in other specific neurons, we amplified the different promoters using N2 genomic DNA and replaced the promoter region of pitp-1. To clone the promoters, we introduced a NgoMIV restriction site to the plasmid backbone (using QuikChange II XL, Agilent Technologies). flp-6 promoter region (~2.8 kb) was amplified with 5′ CAATTGGATAGGAAATGTCACACGAC 3′ and 5′ ATTCTGGAATAATCATATTGTTTTC 3′ and inserted using NgoMIV and AvrII; [Pflp-6::pitp-1(cDNA)::polycismCherry]. The ceh-36 promoter region (~3.1 kb) was amplified with 5′ GAACTCCCGCAGAATGCCAAGTTTTC 3′ and 5′ TGTGCATGCGGGGGCAGGCG 3′ and inserted using NgoMIV and AvrII; [Pceh-36::pitp-1(cDNA)::polycismCherry]. The rgef-1 promoter region (~3.6 kb) was amplified with 5′ CATCGAATGGCAATTTCGAGCACT GCC 3′ and 5′ CGTCGTCGTGAGCAACTGTC 3′ and inserted using NgoMIV and AvrII; [Prgef-1::pitp-1(cDNA)::polycismCherry]. The tax-2 promoter region (~1.8 kb upstream from ATG+ the first 215 amino acid) was amplified with 5′ CAATGACATCAAACTCGGTCGAATGGCG 3′ and 5′ ATGAAATGTTCCTTTCAATGAAAACAGC 3′ and inserted using NgoMIV and AvrII; [Ptax-2::pitp-1(cDNA)::polycismCherry]. The ocr-2 promoter region (~3.8 kb upstream from ATG+ the first 98 amino acid) was amplified with 5′ GTATGGGAATGTCAGCATGGGAACG 3′ and 5′ CTTTCGTGAGAATGCTCACCATTTC 3′ and inserted using NgoMIV and AvrII; [Pocr-2::pitp-1(cDNA)::polycismCherry]. The gpa-14 promoter region (~3.3 kb) was amplified with 5′ GGTTTCGATGATGTGGTACTTTGGC 3′ and 5′ AAGCTCTCTCAGTTTTGCAAGCTC 3′ and inserted using NgoMIV and AvrII; [Pgpa-14::pitp-1(cDNA)::polycismCherry]. The cng-3 promoter region (~760 bp) was amplified with 5′ CACCCCGGATCCGGAGAATCGGAGC 3′ and 5′ ATTCTGGAATAATCATATTGTTTTC 3′ and inserted using NgoMIV and AvrII; [Pcng-3::pitp-1(cDNA)::polycismCherry]. The mod-1 promoter region (~760 bp) was amplified with 5′ CACCTCGTGAAATCGGACCTGAAA 3′ and 5′ GTATGGGAATGTCAGCATGGGACG 3′ and inserted using NgoMIV and AvrII; [Pmod-1::pitp-1(cDNA)::polycismCherry].

### Table 1

| Genetic location | Genomic location | Clone | N2 digest | CB4856 digest | Enzyme | Primers |
|------------------|------------------|-------|-----------|---------------|--------|---------|
| III, -11         | 2629011-2631354  | Y71H2B| 355       | 174, 181      | HpyCH4V| F: CCAGGGTGTTCTCTTGAGAACACGTCCTTCTTC R: GCTTCTGACGTTTTCGAACGTCCTTCTTC |
| III, -10.1       | 2691329-2835114  | Y71H2AM| 418       | 27, 124, 267  | Hphi   | F: GATCTGGACGCTCTTGGAAGCG R: GTGCGAGTCATCTGAAACGTCCTTC |
| III, -9          | 2850762-2893560  | F56F11a| 251, 304  | 555           | Accl   | F: GTGTCACCTGAAAGCTTCTAACAA R: TTTCCCGGAATCCTGGAATGTG |
| III, -8.9        | 2891762-2892784  | F56F11| 236, 332  | 568           | Aval II| F: CAGAGCTTTAGTATGAGGGCCAA R: GTATCGCACTACGTTAAGGCAAGC |
| III, -7.9        | 3052479-3100358  | H06I04| 513       | 234, 279      | SpeI   | F: CTGTAATCCTAGTACCCATTGCCAAGCG R: CAGTCTTTAGTCCCTGAGTTC |
| III, -6.6        | 3452186-3486946  | C34C12| 54, 96, 182| 332           | NalI III| F: GCAACCGGTCGCTCTTTCACACAGG R: TTGTGTATATGGTACGTAGGTC |
| III, -5.4        | 3558492-3596862  | C48G5 | 641       | 259, 382      | BglII  | F: GGGGTCTTCTATCGCTCGCAGAC R: TTTTCTGAGCAGCCATTTCT |
| III, -3.8        | 4112743-4142803  | C30D11| 117, 473  | 590           | RsaI   | F: CATCGGGAAGGGCCAAAGTCGCTGCA R: GCCCTCGAAGGCTGTCGCA |

| SNP primers, location, band sizes (observed after restriction enzyme digestion) and the respective restriction enzyme. The primer sequences are displayed from 5′ to 3′ |
|---------------------------------------------------|---------------------|---------------|-----------------|-----------------|-----------------|
| III, -11                                           | 2629011-2631354     | Y71H2B       | 355             | 174, 181        | HpyCH4V        |
| II, -10.1                                          | 2691329-2835114     | Y71H2AM      | 418             | 27, 124, 267    | Hphi           |
| II, -9                                             | 2850762-2893560     | F56F11a      | 251, 304        | 555             | Accl           |
| II, -8.9                                           | 2891762-2892784     | F56F11       | 236, 332        | 568             | Aval II        |
| II, -7.9                                           | 3052479-3100358     | H06I04       | 513             | 234, 279        | SpeI           |
| II, -6.6                                           | 3452186-3486946     | C34C12       | 54, 96, 182     | 332             | NalI III       |
| II, -5.4                                           | 3558492-3596862     | C48G5        | 641             | 259, 382        | BglII          |
| II, -3.8                                           | 4112743-4142803     | C30D11       | 117, 473        | 590             | RsaI           |
The ttx-7 promoter region (~0.6 kb) was amplified with 5′-CCAATTGTTAACCTTGAGTATGTCTTCAC-3′ and 5′-CTAAAACTTAAAATCAATGAATAAAGAGCTGAAG-3′ and inserted using NgoMIV and AvrII; [Pttx-7::pitp-1(cDNA)::polycismCherry]. The constructs were injected into pitp-1(tm1500);glb-5(Haw);npr-1(ad609) worms (0.5-2.5 ng/µL) with the PF15E11.1::GFP co-injection marker (47.5-49.5 ng/µL).

For the pkc-1 cDNA rescuing constructs, we cut the pitp-1 cDNA under rgef-1 and mod-1 promoters with AvrII and NotI and inserted the pkc-1 cDNA with AvrII and NotI; [Prgef-1:pkc-1(cDNA)::polycismCherry] and [Pmod-1-1:pkc-1(cDNA)::polycismCherry], respectively. The pkc-1 cDNA (isoform c) region (~2.3 kb) was amplified with 5′-ATGAAATTCTTCAGTAGTCGGACAATATCATCTG-3′ and 5′-TTAGTAGGTAAAATGCGGATTGATAAATGAAAAACC-3′. The constructs were injected into pitp-1(t-mtm1500);glb-5(Haw);pkc-1(ok563);npr-1(ad609) worms (2.5-25 ng/µL) with the PF15E11.1::GFP co-injection marker (25-47.5 ng/µL).

We used Gibson assembly cloning to generate the [Punc-25::pkc-1C::polycismCherry], [Pmod-1::pkc-1b(gf)::polycismCherry], and [Punc-25::pitp-1(cDNA) polycismCherry] expression constructs. The three constructs share the polycistronic mCherry pPD95.75 expression vector that was amplified with 5′-CCACGATGCCTGTAGCAATGGCAAC-3′ and 5′-ATTTCACTTTTCCAAGTTGTTAGCGTATCCATCG-3′ (~1.6 kb) and with 5′-GTTGCCATTGCTACAGGCATCGTGG-3′ and 5′-GCTGTCTCATCCATTTTTCACCTAG-3′ (~2.8 kb).

For the [Pmod-1::pkc-1b(gf)::polycismCherry] rescuing construct, we amplified mod-1 promoter (~5.5 kb) with 5′-CGATGGATACGCTAACAACTTGGAAATGAAATGTGTTTTCTCGTAATCCCGTTGTACTTC-3′ and 5′-CATATTCATCGTATACCAAACAATCAAAACTCATATATTTTAATTTTCTTTCACCGCATTGGCACCTGG-3′ and also amplified pkc-1b (~1.9 kb)26) using wild-type N2 cDNA as a template with 5′-CAGATGATATTGTCCGACTACTGAAGAATTTCATatatattttTTTTGGCGGTGAACTGAGCTTTTCC-3′ and 5′-CTAGGTGAAAGTAGGATGAGACAGCGTCGACTTAGTAGGTAAAATGCGGATTGATAAATGAAAAACC-3′ primer pair. We used Q5 Site-Directed Mutagenesis Kit (NEB) to introduce the A160E gain-of-function mutation into pkc-1 using 5′-AGACGTGGTGagATGCGACGGA-3′ and 5′-TTGTCGATCATTGAAAGCATTTG-3′ primers. The resulting expression construct was injected into glb-5 (Haw); npr-1 (ad609) worms (25-100 ng/µL) with the PF15E11.1::GFP co-injection marker (25-100 ng/µL).

To generate the GABAergic pkc-1 rescue construct, we amplified the promoter region of unc-25 (~2 kb) with 5′-cgatggatacgctaacaacttggaataaatatCATCATCGGCCCTGTGTTATCAG-3′ and 5′-CTAGGTGAAAGTAGGATGAGACAGCGTCGACTTAGTAGGTAAAATGCGGATTGATAAATGAAAAACC-3′ primer pair. We used Q5 Site-Directed Mutagenesis Kit (NEB) to introduce the A160E gain-of-function mutation into pkc-1 using 5′-AGACGTGGTGagATGCGACGGA-3′ and 5′-TTGTCGATCATTGAAAGCATTTG-3′ primers. The resulting expression construct was injected into glb-5 (Haw); npr-1 (ad609) worms (25-100 ng/µL) with the PF15E11.1::GFP co-injection marker (25-100 ng/µL).

To generate the GABAergic pitp-1 rescue construct, we amplified the promoter region of unc-25 (~2 kb) with 5′-cgatggatacgctaacaacttggaataaatatCATCATCGGCCCTGTGTTATCAG-3′ and 5′-CTAGGTGAAAGTAGGATGAGACAGCGTCGACTTAGTAGGTAAAATGCGGATTGATAAATGAAAAACC-3′ primer pair. We used Q5 Site-Directed Mutagenesis Kit (NEB) to introduce the A160E gain-of-function mutation into pkc-1 using 5′-AGACGTGGTGagATGCGACGGA-3′ and 5′-TTGTCGATCATTGAAAGCATTTG-3′ primers. The resulting expression construct was injected into glb-5 (Haw); npr-1 (ad609) worms (25-100 ng/µL) with the PF15E11.1::GFP co-injection marker (25-100 ng/µL).

Imaging strains: To image pitp-1 neurons localization we used the modified plasmid pEnter (95.75 MCS) GFP. We amplified the pitp-1 promoter (~4.2 kb) using N2 genomic Forward (F) and Reverse (R) sequences.

| Primer | Forward (F) | Reverse (R) |
|--------|-------------|-------------|
| pitp-1(tm1500) | F: CATGGAGATTTCGACCCGGACACACC4534 | R: CTGAAAAGCTGATCGTCTCGATCCAAACTCTG6033 |
| pkc-1(ok563) | F: GACTCGGCGTTGATTAGACGCTTCGG | R: GATCAAACGCTATTTGCGACAGCTAG |
| dgk-1(ok1462) | F: CCTCGGGATTATCTGATCGTCTGG | R: GACCTGGAACACTTCTTTGTCATAG |

Imaging strains: To image pitp-1 neurons localization we used the modified plasmid pEnter (95.75 MCS) GFP. We amplified the pitp-1 promoter (~4.2 Kb) using N2 genomic

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**Table 2** Primers used for deletion strains

| Primer | Forward (F) | Reverse (R) |
|--------|-------------|-------------|
| pitp-1(tm1500) | F: CATGGAGATTTCGACCCGGACACACC4534 | R: CTGAAAAGCTGATCGTCTCGATCCAAACTCTG6033 |
| pkc-1(ok563) | F: GACTCGGCGTTGATTAGACGCTTCGG | R: GATCAAACGCTATTTGCGACAGCTAG |
| dgk-1(ok1462) | F: CCTCGGGATTATCTGATCGTCTGG | R: GACCTGGAACACTTCTTTGTCATAG |
DNA. The promoter was inserted with NheI and Xbal. The constructs were injected into pitp-1(tm1500);glb-5(Haw);n- pr-1(ad609) worms. To validate the expression of pitp-1 in the O2-sensing neurons, we crossed this strain with the AX1846 strain, which expressed mCherry under the promoter region of glb-5.13 The Ca2+ imaging strains were generated as described previously.6,10 The Pgy-37::YC2.60 extrachromosomal array was introduced to pitp-1(tm1500);glb-5(Haw);n- pr-1(ad609) worms by crossing with the EVG159 parental strain.10

2.11 | Ca2+ imaging

We performed the Ca2+ imaging experiments as described in.10 We recorded the Ca2+ responses at two frames per second, using an Olympus IX71S1F-3-5 inverted microscope equipped with UAPON40X Universal apochromatic water immersion objective (Olympus, Tokyo, Japan), DV2 2-channel imager (Photometrics), Rolera EM-C2 (Qimaging), and Meta Morph software (Molecular Devices). To immobilize worms, we glued them to agarose pads (2% agarose in M9) using Dermabond topical skin adhesive (Ethicon). These worms were trapped inside a 500 μm deep rectangular PDMS chamber (12 mm wide, 0.5 mm deep, and 17 mm long). In all of our imaging experiments, humidified gases were delivered to the microfluidic chambers using a PHD 2000 syringe pump (Harvard Apparatus, Holliston, MA, United States) at a flow rate of 0.5 ml per minute. Teflon valves, regulated by a ValveBank Controller (Automate Scientific, Berkeley, California, USA), were used to rapidly switch between gas mixtures. Image analysis was performed using custom-written MATLAB software.

2.12 | Aldicarb assays

We adapted the aldicarb assay protocol described in.26 In brief, a day before the experiment, we prepared experimental NGM plates containing 1.5 mM of aldicarb (33386-100MG Sigma Aldrich; stock solution: 50 mM in 70% ethanol). After 5 hours at RT, we seeded the plates with 100 μL of OP50 bacteria (OD600 ~ 0.6) and let them dry for 24 hours at RT. On these plates, we placed young adult hermaphrodites that were either exposed to hypoxia (1% O2) for 24 hours as described above or remained at 21% O2. We measured paralysis every 20 minutes for 2 hours. Paralysis is defined by the absence of movement when prodded three times with an eyelash on the head and tail. The 1% and 21% O2-aldicarb assays included five biological replicates, in which at least 120 worms were examined.

2.13 | Statistical analysis

For comparison between two groups, we used an unpaired t test with Welch’s correction. For comparison between more than two groups, when two parameters are explored (ie, the effect of O2 level and genetic background), we used two-way ANOVA with Bonferroni posttest. For the aldicarb assays, we used the log-rank (Mantel-Cox) test. Data are presented as mean ± SEM (standard error of the mean) or as a Kaplan-Meier curve (ie, in the case of the aldicarb experiments). All statistical analysis was performed using GraphPad Prism version 8.0.0 for Windows, GraphPad Software, San Diego, California USA, www.graphpad.com.

3 | RESULTS

3.1 | PITP-1 is required for fast recovery of the bordering eating behavior after hypoxia

GLB-5 accelerates the rate of bordering feeding behavior recovery after hypoxia.6,10 To explore the underlying mechanism, we performed an EMS mutagenesis screen and isolated glb-5(+); npr-1(−) mutants that fail to display wild-type bordering behavior (after 24 hours exposure at 1% O2, followed by 30 min recovery at 21% O2). To avoid isolating mutants that are generally defective in their O2-sensing and/or bordering behavior, we verified that these mutants resume wild-type bordering behavior after 4 hours recovery at 21% O2. We used single nucleotide polymorphism (SNP) mapping23 to locate one of these mutations, heb2, to a 1.6 Mb genomic interval on chromosome III. By whole-genome sequencing (Illumina), we identified 10 point mutations in this genomic region. Among these mutations, a G > A transition mutation disrupts a splice-acceptor point mutations in this genomic region. The bordering behavior of the heb2 mutant was similar to the rapid recovery of bordering behavior by generating transgenic pitp-1(−); glb-5(+); npr-1(−) worms that express either the genomic or coding region of pitp-1 under its promoter region (Figure 1D).
3.2 | PITP-1 is not generally required for GLB-5-dependent O₂ responses

So far, all the characterized GLB-5 suppressors have been found to affect the O₂-sensing pathway in the AQR, PQR, and URX neurons. This occurs by directly changing the function of the O₂-sensors GCY-35 and GCY-36 or by interfering with their transport to the dendritic endings of URX.6 To explore the mechanism by which PITP-1 regulates GLB-5 activity, we first set out to determine whether it is co-expressed with glb-5 in the AQR, PQR, and URX neurons. For this, we generated transgenic worms co-expressing a GFP and a mCherry fluorescent tags under the promoter regions of pitp-1 and glb-5, respectively. As previously reported, pitp-1 is expressed in many neurons.27 We observed a co-localization of the GFP and mCherry fluorescence in the AQR, PQR, and URX neurons (Figure 2A), as well as in the BAG O₂-sensing neurons (the BAG neurons act reciprocally to the URX neurons and are activated at low O₂ levels), indicating that pitp-1 is co-expressed with glb-5 in O₂-sensing neurons.

Next, we asked whether pitp-1 activity in these neurons is sufficient to rescue the fast bordering recovery after hypoxia. To answer this, we generated two transgenic pitp-1(-); glb-5(+) and npr-1(-); glb-5(+) worm strains expressing the coding region of pitp-1 just in AQR, PQR, and URXL/R (under the promoter region of gcy-37) or in all six O₂-sensing neurons (i.e., URX L/R, AQR, PQR, and BAG L/R) under the promoter region of glb-5.13,30 The expression of pitp-1, either in the four or the six neurons, did not restore fast bordering after hypoxia (Figure 2B), suggesting the pitp-1 acts in other neurons to accelerate the recovery of feeding behavior after hypoxia.

To further explore pitp-1 function, we asked whether it is generally important for glb-5 activity. To explore this, we performed speed assay experiments in which we exposed npr-1(-); glb-5(+) worms to a 21%-14% O₂ shift. We generated transgenic worms expressing pitp-1(-); glb-5(+) worms to a 21%-17% O₂ shift. We use this O₂ shift as an indicator for glb-5 activity because it induces a significant decrease in the foraging speed of glb-5(+) worms, however, only a mild decrease in npr-1(-) worms that bear a nonactive glb-5.10,13 As expected, the shift from 21% to 17% O₂ induced a robust decrease in the speed of glb-5(+) worms, and a mild decrease in npr-1(-) worms (Figure 2C). The speed-shift observed in pitp-1(-); glb-5(+) worms was similar to that seen in glb-5(+) npr-1(-) controls, suggesting that pitp-1 is not required for glb-5 activity in this behavioral paradigm.

3.3 | O₂-evoked Ca²⁺ responses in URX in pitp-1 mutants

To directly explore whether pitp-1 is important for glb-5-mediated O₂ responses, we performed Ca²⁺ imaging experiments. For this, we expressed the ratiometric YC2.60 Ca²⁺ sensor in the URX neurons (as described in) of npr-1(-); glb-5(+) npr-1(-) and pitp-1(-); glb-5(+) worms. These worm strains were exposed to 1% O₂ for 24 hours or remained at 21% O₂ as a control (i.e., naïve worms). We chose the 21% to 14% shift because it mimics the O₂-concentration change that the worms experience when entering the bacterial lawn border, where the O₂ level is lower than the lawn center and its surroundings.8 The 21% to 14% O₂ shift elicited a significant drop in Ca²⁺ concentration in naïve worms of the three strains (Figure 3A-C). After 24 hours exposure to hypoxia, the 21%-14% O₂ shift did not induce a Ca²⁺ drop in npr-1 worms (Figure 3A). This result is in line with previous studies that show that a functional glb-5 is essential for decreasing the sensitivity of the URX neurons after exposure to hypoxia.10 By contrast, the shift from 21% to 14% O₂ induced a significant decrease in the Ca²⁺ level of both glb-5(+) npr-1(-) and pitp-1(-); glb-5(+) npr-1(-) worms (Figure 3B,C). The magnitude of the Ca²⁺ drop was similar between the two strains.

Together, these results suggest that pitp-1 is not required for glb-5-dependent O₂ responses in the O₂-sensing neurons.

3.4 | PITP-1 roles in salt/benzaldehyde chemotaxis and recovery from hypoxia are distinct

Where does PITP-1 act to facilitate fast recovery from hypoxia? A previous study by Iino and colleagues showed that PITP-1 acts in the ASEL/R neurons to regulate salt (NaCl) chemotaxis/chemorepulsion and in the AWC and the ASH neurons to promote, respectively, olfaction and osmotic avoidance.27 Therefore, we explored whether pitp-1 acts in the ASE and the AWC neurons to facilitate fast recovery from hypoxia. We generated transgenic pitp-1(-); glb-5(+) worms expressing pitp-1 under the flp-6 promoter region (ASE neurons) or under the ceh-36 promoter region (ASE and AWC neurons). Expression of pitp-1 in just the ASE or the ASE/AWC neurons did not rescue the fast recovery of bordering behavior after 24 hours exposure to hypoxia (Figure 4A). By contrast, the expression of pitp-1 in these neurons (just in ASE or in both ASE and AWC) restored the chemotaxis attraction to salt (Figure 4B), albeit to a lower level than wild type. Together, these results indicate that the place of activity of PITP-1 in the control of salt chemotaxis is different from the place of its activity in the regulation of feeding behavior after hypoxia.

3.5 | PITP-1 acts in MOD-1-expressing neurons to promote fast recovery from hypoxia

To determine where pitp-1 activity is required for the fast bordering recovery after hypoxia, we restricted the expression
FIGURE 2  Expression of pitp-1 in the O₂-sensing neurons AQR, PQR, URX, and BAG does not rescue the fast recovery of bordering behavior after hypoxia. A, Bright-field and fluorescence images of transgenic worms co-expressing Pglb-5::glb-5::mCherry and Ppitp-1::pitp-1::GFP. Arrows indicate the AQR, BAG, PQR, and URX neurons in the merge images. Scale bars: 25 μm. B, A bar graph presenting the recovery of bordering after 24 hours in hypoxia. Asterisks indicate significance for comparisons with glb-5(+) npr-1(−) animals after 24 hours at 1% O₂. C, Speed measurements. The speed of worms was measured at 21% and 17% O₂ in the presence of bacteria. Asterisks indicate significance for comparisons with glb-5(+) npr-1(−) worm speed at 17% O₂. Data represent the average of at least six independent experiments. Two-way ANOVA with Bonferroni posttest. ****P < .0001, NS, nonsignificant. Error bars represent SEM.
of pitp-1 cDNA to a subset of neurons using the following promoters: cng-3 [AFD, ASE, ASI, AWB, and AWC], gpa-14 [ADE, ALA, ASH, ASI, ASJ, ASK, AVA, CAN, DVA, PHA, PHB, PVQ, and RIA], mod-1 [AIA, AIB, AIY, AIZ, DD, RIC, RID, RIM, RME, and VD], ocr-2 [expressed in PHA, PHB, ASH, ADL, ADF and AWA], tax-2 [expressed in the AWC, AFD, ASE, ASG, ASJ, AQR, BAG, ASK, ASI, AWB, and PQR neurons], and ttx-7 [ADF,
3.6 Impairment of PKC-1 activity restores rapid bordering recovery after hypoxia in pitp-1 mutants

A previous study showed that the function of PITP-1 in salt chemotaxis plasticity is mediated, in part, by diacylglycerol (DAG) signaling. A key target of DAG is the protein kinase C1 (PKC-1), which is predicted to require DAG for activation. To explore the function of PKC-1 in the PITP-1-dependent recovery from hypoxia, we first asked whether PKC-1 itself is required for the fast recovery. To answer this, we generated glb-5(+) npr-1(−) worms bearing the pkc-1(ok563) deletion allele (hereafter referred to as pkc-1(−)), which removes ~1.3 Kb from the 5′ UTR of the pkc-1B splice isoform (including the ATG translation initiation site). The bordering recovery of glb-5(+) pkc-1(−); npr-1(−) worms from hypoxia was similar to glb-5(+) npr-1(−) controls (P > .9999; Figure 6A), indicating that PKC-1 is not required for the fast recovery of bordering behavior.

We next introduced pkc-1(ok563) into pitp-1(−); glb-5(+) npr-1(−) worms and examined their bordering recovery after hypoxia. Intriguingly, pkc-1 loss-of-function rescued the fast bordering recovery after hypoxia (Figure 6A), albeit to a lesser degree in comparison to glb-5(+) npr-1(−) controls. This result suggests that PKC-1 activity inhibits the fast recovery from hypoxia in pitp-1(−); glb-5(+) npr-1(−) worms.

PITP-1 controls the recovery from hypoxia through its activity in mod-1-expressing neurons (Figure 5). Therefore, we asked whether pkc-1 acts in these neurons to inhibit the fast bordering recovery of pitp-1(−); glb-5(+) pkc-1(−); npr-1(−) worms after hypoxia. To explore this, we generated transgenic pitp-1(−); glb-5(+) pkc-1(−); npr-1(−) worms expressing pkc-1B (wt) under the mod-1 promoter region. Moreover, we generated two additional rescuing strains. One expresses pkc-1B (wt) in all neurons, under the promoter region of rgef-1, and thus, serves as a positive control for neuronal pkc-1 activity. The second expresses pkc-1B (wt) just in GABAergic neurons, under the promoter region of unc-25. Expression of pkc-1B (wt) in all neurons or only in the mod-1-expressing neurons resulted in a significant and a similar degree of bordering-inhibition (P mod vs Pmod = .6755), suggesting that no other neurons, besides the mod-1-neurons, are important for this inhibitory activity of PKC-1. Unexpectedly, restoring PKC-1 activity under the unc-25 promoter resulted in a small, yet, a significant decrease in the recovery of bordering after hypoxia (Figure S1B), suggesting that the activity of PKC-1 in one or more of the GABAergic neurons DD, RME, and VD contributes to the inhibition of bordering after hypoxia. Together, these results indicate that PKC-1 activity in the mod-1-expressing neurons decreases bordering recovery after hypoxia. Furthermore, they suggest that the function of PITP-1 is to attenuate PKC-1 activity in these neurons. Finally, although pitp-1 expression in DD, RME, and VD is not sufficient for rescuing the fast recovery of bordering.
after hypoxia, the above \( \text{pkc-1} \)-rescuing experiments suggest that one or more of these neurons play a role in the recovery of bordering after hypoxia.

To further explore PKC-1 function, we asked whether constitutive PKC-1 activity in the \( \text{mod-1} \)-expressing neurons would interfere with the recovery of bordering after hypoxia. For this, we overexpressed a gain of function mutation of \( \text{pkc-1} \) in the \( \text{mod-1} \)-expressing neurons of \( \text{glb-5}(+)\); \( \text{npr-1}(-) \) worms. This \( \text{pkc-1} \) mutation, hereafter referred to as \( \text{pkc-1}(gf) \), substitutes alanine 160 for glutamate in the pseudosubstrate domain of
PKC-1, resulting in constitutive enzymatic activity.\textsuperscript{51} The bordering index of transgenic \textit{pck-1(\textit{gf})} worms after hypoxia was significantly lower compared to the parental \textit{glb-5(+)\textit{npr-1(−)}} worms (−54% compared to −96%, after hypoxia, Figure 6B). In fact, the bordering index of transgenic \textit{pck-1(\textit{gf})} worms was low even in control worms that did not experience hypoxia, indicating that constitutive PKC-1 activity in the \textit{mod-1}-expressing neurons inhibits the bordering behavior of \textit{glb-5(+)\textit{npr-1(−)}} worms irrespective of the exposure to hypoxia. These data support the hypothesis that excessive PKC-1 activity in the \textit{mod-1}-expressing neurons suppresses bordering after hypoxia.

### 3.7 PMA inhibits fast recovery of bordering behavior after hypoxia

How does PITP-1 attenuate PKC-1 activity? Since PKC-1 is thought to be activated by DAG,\textsuperscript{51} and PITP-1 can potentially affect the level of DAG,\textsuperscript{29} we hypothesized that excess of DAG would attenuate the recovery of bordering behavior of \textit{glb-5(+)\textit{npr-1(−)}} worms. To test this, we put \textit{glb-5(+)\textit{npr-1(−)}} worms on NGM-plates containing PMA, a DAG analog that binds and activates PKC,\textsuperscript{50} and measured the recovery of bordering behavior after hypoxia.

In addition, we measured the effect of PMA on the bordering behavior of control worms that were not exposed to hypoxia and in worms exposed to 0.002% DMSO (the PMA vehicle). PMA significantly decreased the bordering recovery of both \textit{glb-5(+)\textit{npr-1(−)}} and \textit{pma-1(−)} worms after hypoxia (Figure 6C); however, it did not affect the bordering behavior of control worms.

To further explore whether increased DAG signaling can inhibit the recovery of bordering behavior after hypoxia, we introduced the \textit{dgk-1(\textit{ok1462})} null mutation (hereafter referred to as \textit{dgk-1(−)}) into the genetic background of both \textit{glb-5(+)\textit{npr-1(−)}} and \textit{pma-1(−)}; \textit{glb-5(+)\textit{npr-1(−)}} worms. The \textit{dgk-1} gene encodes a diacylglycerol (DAG) kinase\textsuperscript{52} which decreases DAG level by converting it to phosphatidic acid (PA).\textsuperscript{53-55} Therefore, DAG signaling is increased in \textit{dgk-1\textit{mutants}}. DGK-1 was not important for the bordering behavior of control worms that did not experience hypoxia (Figure 6D). However, the \textit{dgk-1} mutation significantly decreased the bordering recovery of \textit{glb-5(+)\textit{npr-1(−)}} worms after hypoxia. Together, these results suggest that excess of DAG attenuates the recovery of bordering behavior after hypoxia. Moreover, this inhibitory effect is specific to the recovery period and does not apply to worms that were not exposed to hypoxia.

### 3.8 PITP-1 effect on aldicarb resistance

So far, our results are consistent with a working model in which PITP-1 restricts PKC-1 activity during the recovery of worms from hypoxia by limiting DAG availability. Previous studies show that increased PKC-1 activity and exogenous PMA sensitize worms to the paralytic effect of the acetylcholinesterase inhibitor aldicarb.\textsuperscript{51,56,57} Therefore, we hypothesized that \textit{pma-1} loss-of-function would result in increased sensitivity of worms to aldicarb after hypoxia. To test this, we measured aldicarb-induced paralysis in \textit{pma-1(−), glb-5(+)\textit{npr-1(−)}}, and \textit{pma-1(−)}; \textit{glb-5(+)\textit{npr-1(−)}} worms after exposure to hypoxia or in worms that were not exposed to hypoxia (as a control). \textit{pma-1(−)}; \textit{glb-5(+)\textit{npr-1(−)}} worms were significantly more sensitive to aldicarb than \textit{glb-5(+)\textit{npr-1(−)}} worms in both experimental conditions (Figure 6E; \textit{P} = .0020, and \textit{P} = .0008, after hypoxia and control conditions, respectively), suggesting that \textit{pma-1} is generally important for restricting neurotransmission in \textit{C elegans}. Moreover, \textit{pma-1(−)} worms were significantly more sensitive to aldicarb than \textit{glb-5(+)\textit{npr-1(−)}} animals (\textit{P} < .0001 for both experimental conditions). Notably, the exposure to hypoxia increased the aldicarb resistant of the three strain; Hazard ratios (after 24 hours exposure to hypoxia vs controls) were \textit{pma-1(−)} \[0.4564\]; \textit{glb-5(+)\textit{npr-1(−)}} \[0.4095\]; and \textit{pma-1(−)}; \textit{glb-5(+)\textit{npr-1(−)}} \[0.3958\], suggesting that...
hypoaxia attenuates neurotransmitter release in a glb-5 and pitp-1 independent manner.

Our data suggest that PITP-1 acts in mod-1-expressing neurons to facilitate the fast recovery of bordering after hypoxia (Figure 5). Therefore, we hypothesized that restoring pitp-1 function in these neurons would rescue pitp-1(−); glb-5(+) worms from the hypersensitivity to aldicarb. To test this, we repeated the above aldicarb experiments with transgenic Pmod-1::pitp-1(cDNA) worms (these worms display rescued bordering phenotype after hypoxia, Figure 5). As predicted, Pmod-1::pitp-1(cDNA) animals were significantly more resistant to aldicarb compared with their pitp-1(−); glb-5(+) animals (P < .0001 for 21% O2-aldicarb assays and P = .0002 for the 1% O2 assays). Since resistance to aldicarb can arise from impaired pre- or postsynaptic activity,26 the hyper-resistance of transgenic Pmod-1::pitp-1 worms to aldicarb may reflect the consequence of over-PITP-1 activity in these neurons, which decreases synaptic transmission to a suboptimal level. In conclusion, this set of experiments further supports the hypothesis that PITP-1 facilitates the fast recovery of bordering after hypoxia by restricting neurotransmission in mod-1-expressing neurons.

4 | DISCUSSION

4.1 | PITP-1 acts downstream to O2-sensing neurons to facilitate fast bordering recovery from hypoxia

We previously isolated the pdl-1(db508), gcy-35(heb1), and gcy-36(heb3) mutations that suppress the recovery of bordering eating behavior after hypoxia in glb-5(+)npr-1(−) worms.6,10 These mutations directly interfere with the O2-sensing machinery in the AQR, PQR, and URX neurons. The pdl-1 mutation prevents the targeting of the soluble guanylate cyclase GYC-35 O2-sensor to URX endings, while the gcy-35/gcy-36 mutations make GCY-35 and GCY-36 insensitive to GLB-5. As a result, pdl-1(db508), gcy-35(heb1), and gcy-36(heb3) mutants are generally defective for glb-5-dependent-O2 responses. That is, they do not display a robust decrease in speed when shifted from 21% to 17% O2 and display high URX Ca2+ level at 14% and 17% O2 after hypoxia.6,10 Notably, the expression of wild-type pdl-1, gcy-35, and gcy-36 in the AQR, PQR, and URX neurons of pdl-1(db508), gcy-35(heb1), and gcy-36(heb3) mutants, respectively, rescues both the bordering behavior after hypoxia and the slowing response phenotypes, indicating that the activities of these genes in AQR, PQR, and URX are both necessary and sufficient for GLB-5-dependent O2 responses.

By contrast, pitp-1 mutants display a robust slowing response when shifted from 21% to 17% O2 and a sharp drop in URX Ca2+ level upon a 21% to 14% O2 transition, after hypoxia (Figures 2C and 3C, respectively). Moreover, the recovery of bordering behavior after hypoxia cannot be rescued by restoring wild-type PITP-1 activity in AQR, PQR, and URX or even by expressing pitp-1 under the promoter region of glb-5 (Figure 2B), which is expressed in additional neurons, including ADF, ASG, and BAG.30 Together, these results indicate that PITP-1 is not required for the function of GLB-5 in O2-sensing per se, and suggest that it acts in other neurons to facilitate the fast recovery from hypoxia. In this respect, it is worth emphasizing that our previous studies suggest that the recovery of bordering behavior after hypoxia depends on the desensitization of the URX neurons by GLB-5.10 The results of this study imply that URX-desensitization is not sufficient for the fast recovery phenotype and that downstream neurosignaling pathways are also required.

Indeed, our results suggest that the recovery process is mediated by PITP-1-dependent neurosignaling in mod-1-expressing neurons. Interestingly, some of the mod-1-expression neurons are involved in dietary choice behavior in C. elegans and have chemical-synapse connections with the GLB-5-expressing neurons. For example, the AIY interneurons, which have chemical synapses with BAG,58 appear to be important for finding high-quality food in diverse environments59 and for regulating food-leaving behavior.60 Moreover, a recent paper suggests that the transcription factor DAF-16 regulates salt avoidance by controlling neurotransmission between ASER, AIA, and AIY61; the same neurons in which PITP-1 acts to regulate salt chemotaxis. Intriguingly, DAF-16 regulates the aversive response to pheromone signals,62 which could be affected by the O2 concentration that the worm experience and potentially change the attraction/avoidance of worms to food. In the future, it will be exciting to explore whether DAF-16 controls the recovery of bordering behavior after hypoxia and whether it affects the functions of PITP-1 in both salt and recovery from hypoxia responses.

4.2 | PITP-1 functions in salt chemotaxis and recovery of bordering after hypoxia

A previous study by Iino and colleagues showed that PITP-1 acts in the ASEL/R neurons to facilitate salt chemotaxis.27 Intriguingly, pitp-1-loss-of-function did not affect the Ca2+ response to salt in ASER, suggesting that PITP-1 is not important for salt sensing per se but rather for neurotransmission from ASER. Indeed, salt-induced Ca2+ responses in the AIB interneurons, which are directly connected through
chemical synapses to the ASER neurons, were significantly decreased in pitp-1 mutants, further supporting the function of PITP-1 in neurotransmission. Our results show that PITP-1 function in the ASEL/R neurons is not important for the fast recovery of bordering behavior after hypoxia (Figure 4A), suggesting that PITP-1 roles in salt chemotaxis and recovery of eating behavior after hypoxia are distinct. In fact, we show that the expression of wild-type pitp-1 under the promoter region of mod-1 results in near-to-wild-type recovery of the bordering behavior after hypoxia (Figure 5). Intriguingly, one of the interneurons in which mod-1 is expressed is AIB (36).

In future studies, we will investigate the effect of PITP-1 on O₂-evoked-Ca²⁺ responses (after hypoxia) in AIB and other interneurons that expresses mod-1 (e.g., AIY).

4.3 | Is PITP-1 the “GLB-5” of mod-1-expressing neuron(s)?

Our previous studies suggest a model in which GLB-5 accelerates the recovery of bordering behavior after hypoxia by decreasing the sensitivity of URX neurons to O₂.¹⁰ In this

![Diagram of PITP-1 function in mod-1-expressing neurons following exposure to 21% O₂ after 24 h in hypoxia.](image)

**FIGURE 7** An illustrated working model explaining the function of PITP-1 in the mod-1-expressing neuron(s) following the exposure of worms to 21% O₂ after 24 hours at 1% O₂. The figure describes the recovery of wild-type (WT) animals (i.e., glb-5(+) ; npr-1(−) worms) vs pitp-1(−) ; glb-5(+) ; npr-1(−) mutants; left of the dash line are the mod-1 neurons of WT animals and right of the dash line are the mutant neurons. In WT, PITP-1 inhibits PKC-1 by limiting DAG availability. It does so by limiting PIP2 synthesis and PtdOH dephosphorylation. As a result, PKC-1 is overactivated after hypoxia, a situation that triggers the release of a yet-to-be-identified neurotransmitter(s)/neuropeptide(s), which inhibits the recovery of bordering eating behavior. The larger fonts and arrows, the pitp-1-mutants panel, indicate excessive substrate concentration and PKC-1 activity.
way, the URX neurons are fine-tuned to respond to subtle changes in the ambient O₂ level. For example, the 21% to ~14% shift that the worm experience when it re-enters the bacterial lawn after leaving it in hypoxia. Here, we would like to extend this model and propose that PITP-1’s function in the mod-1-expression neurons is analogous to GLB-5 in the O₂-sensing neurons. That is, PITP-1 acts to decrease PKC-1 activity in mod-1-expression neurons, and thus, prevent excessive neurotransmission that attenuates the recovery from hypoxia. Below, we present the experimental evidence and logic supporting this working model (Figure 7).

(1) pck-1 loss-of-function rescues the recovery phenotype of pitp-1 mutants (Figure 6A). (2) Restoring the activity of wild-type PKC-1 in the mod-1-expressing neurons of pitp-1(−); glb-5(+); pkc-1(−); npr-1(−) worms attenuates the recovery of bordering behavior after hypoxia (Figure 6A). Moreover, expression of constitutively active PKC-1 in these neurons (pkc-1(gf)) significantly inhibits the bordering behavior of the worms (Figure 6B). (3) PKC-1 activation increases worms’ sensitivity to the acetylcholinesterase inhibitor aldicarb. Indeed, pitp-1 mutants are more sensitive to aldicarb compared with their parental glb-5(+) npr-1(−) strain (Figure 6E). Moreover, rescuing PITP-1 function in the mod-1-expression neurons of pitp-1 mutants increases the resistance to aldicarb (Figure 6F). Together, these observations support the hypothesis that PITP-1 facilitates the fast recovery of bordering after hypoxia by decreasing PKC-1 activity in the mod-1-expressing neurons.

PKC-1 is thought to be activated by diacylglycerol (DAG), which can be generated by the following processes: (a) Through the cleavage of phosphatidylinositol 4,5-bisphosphate (PIP2) to DAG and inositol 1,4,5-trisphosphate (IP₃) by phospholipase Cβ (PLCβ)⁶⁴; (b) Through the dephosphorylation of phosphatidic acid (PtdOH) to DAG by PtdOH phosphatase.⁶⁵ The C. elegans PITP-1 is a member of the class II phosphatidylinositol transfer proteins, which can control DAG availability by binding & shuttling phosphatidylinositol-4-phosphate (PIP2 precursor)⁶⁶ and PtdOH.⁶⁵,⁶⁷ Therefore, we suggest that PITP-1 decreases the activity of PKC-1 by restricting DAG availability.

Two observations support this claim. First, exogenous PMA (DAG analog) decreases the bordering recovery of wild-type glb-5(+) npr-1(−) worms after hypoxia (Figure 6C). Second, dkg-1 loss of function, which increases DAG level, also decreases the recovery of glb-5(+) npr-1(−) worms from hypoxia. PKC-1 regulates the secretion of neuropeptides⁵¹ and may function in neurotransmitter release.⁵⁰ We hypothesize that excessive secretion of a yet to be identified neuropeptide(s) and/or neurotransmitter(s) inhibit(s) the recovery of bordering behavior after hypoxia.

In this respect it is interesting to mention that amphetamine-induced anorexia in the rat is associated with the expression of several PKC isotypes,⁶⁸ including the eta isotype that is structurally similar to the C. elegans PKC-1.⁶⁹ The anorectic effect is decreased by PKC alpha-antisense knockdown, suggesting that PKC suppresses food consumption.⁶⁸ Finally, the amphetamine treatment decreases the level of neuropeptide Y (NPY) mRNA in the rat hypothalamus.⁶⁸ This is interesting because NPY is known to regulate food-consumption and is the ligand of the NPY receptor that resembles the C. elegans NPR-1, which is also involved in eating-behavior control.⁹ In conclusion, despite the significant difference in the complexity of controlling eating behavior between the C. elegans worm and the rat and other mammals, we would like to suggest (with due caution) that the link between PKC, neuropeptides, and suppression of eating behavior may be ancient and preserved in evolution. In this context, it would be fascinating to investigate whether orthologs of PITP-1 in the rat are involved in the suppression of eating behavior due to amphetamines and to explore whether mammalian PITP and PKC are involved in high-altitude anorexia.

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CONFLICT OF INTEREST
The authors declare that they have no conflict of interest.

AUTHOR CONTRIBUTIONS
Z. Abergel, M. Shaked, V. Shukla, and E. Gross designed and performed the experiments, contributed new reagents, analyzed the data, and wrote the paper; Z.X. Wu Contributed to paper writing; all authors read and approved the final manuscript.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.