Anophele mosquito species have a highly sensitive chemosensory apparatus to detect diverse chemical stimuli that drive host-seeking and blood-feeding behaviors required to vector pathogens for malaria and other diseases. This process incorporates a variety of chemosensory receptors and transduction pathways. We used advanced in vivo gene-editing and -labeling approaches to localize and functionally characterize the ionotropic coreceptor AcIr76b in the malaria mosquito Anopheles coluzzii, where it impacts both olfactory and gustatory systems. AcIr76b has a broad expression pattern in female adult antennal grooved pegs, coeloconic sensilla, and T1 and T2 sensilla on the labellum, stylets, and tarsi, as well as the larval sensory peg. AcIr76b is colocalized with the Orco odorant receptor (OR) coreceptor in a subset of cells across the female antennae and labella. In contrast to Orco and Ir8a, chemosensory coreceptors that appear essential for the activity of their respective sets of chemosensory neurons in mosquitoes, AcIr76b′′ mutants maintain wild-type peripheral responses to volatile amines on the adult palps, labellum, and larval sensory cone. Interestingly, AcIr76b′′ mutants display significantly increased responses to amines in antennal grooved peg sensilla, while coeloconic sensilla reveal significant deficits in responses to several acids and amines. Behaviorally, AcIr76b mutants manifest significantly female-specific insemination deficits, and although AcIr76b′′ mutant females can locate, alight on, and probe artificial blood hosts, they are incapable of blood feeding successfully. Taken together, our findings reveal a multidimensional functionality of Ir76b in anopheline olfactory and gustatory pathways that directly impacts the vectorial capacity of these mosquitoes.

The authors declare no competing interest.

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mutants, highlighting the involvement of other odorant signaling pathways and sensory modalities (21, 22).

In Drosophila, antennal IRs are primarily expressed in the coeloconic sensilla (18). Unlike ORs, which rely on a single coreceptor (Orco), three IRs—Ir8a, Ir25a, and Ir76b—function as IR coreceptors (18, 19). Interestingly, each Drosophila melangaster ionotropic receptor (DmIR) coreceptor is associated with distinct odor preferences; DmIr8a is critical for acid sensitivity, whereas DmIr25a and DmIr76b are both responsible for amine detection (18, 23, 24). In addition to olfactory function, Ir76b and Ir25a are also involved in other sensory modalities and pathways. In Drosophila, DmIr76b has been found to be involved in gustatory responses to salt and amino acids (25, 26), and DmIr25a acts in both thermosensation and hydrosensation (27, 28). Furthermore, while DmIr25a and DmIr76b are both coexpressed with DmIr92a acting as a Drosophila ammonia/amine receptor, DmIr92a is able to function independently of either of these coreceptors (24, 29).

Recently, several studies focusing on mosquito IRs have revealed that the homologs to Drosophila IR coreceptors similarly regulate a range of Orco-independent sensing pathways. In the arbovirus vector Ae. aegypti, AcIr8a null mutants lost the neuronal and behavioral responses to acids (30). In An. coluzzii larvae, all three IR coreceptors are expressed on the larval antennae, and RNA interference (RNAi) knockdown of AcIr76b specifically impacts larval responses to butylamine (31). In adults, AcIr76b is highly expressed in antennal neurons that do not express ORs (11); in Xenopus oocytes, it drives responses to several amines when coexpressed with AcIr25a and either AcIr41a or AcIr41c (11).

To examine the roles and relevance of Ir76b in the olfactory system of An. coluzzii, we utilized CRISPR-Cas9-mediated gene editing to establish an AcrIr76b-QF2 driver and AcrIr76b null mutant lines. Localization studies using the driver within a binary Q system (32) reveal that AcrIr76b is robustly expressed in antennal grooved pegs, coeloconic sensilla, labella, stylets, and tarsi of adult females, as well as in the larval antennae where it specifically innervates the sensory peg. Surprisingly, in adult female AcrIr76b−/− mosquitoes display significantly increased antennal responses to several amines, whereas peripheral responses to acid stimuli are unaffected. Behaviorally, AcrIr76b−/− mutant females display severe mating deficits and, interestingly, have acutely lost the ability to blood feed successfully. These studies demonstrate that AcrIr76b acts in both olfactory and gustatory systems of An. coluzzii, where it impacts the reproductive fitness and ultimately the vectorial capacity of this globally important mosquito.

Materials and Methods
Mosquito Rearing. An. coluzzii (SUAF 2La2La), previously known as Anopheles gambiae sensu stricto “M-form” (1), originating from Suakoko, Liberia, was obtained through the BEI Resources, National Institute of Allergy and Infectious Diseases (NIAID), NIH: Anopheles gambiae, Strain SUAF2La, Eggs, MRA-765, contributed by Alessandra della Torre at University of Rome “La Sapienza”, Rome, Italy. Mosquito lines were reared at 27 °C, 75% humidity under a 12 h light/12 h dark photoperiod and supplied with 10% sucrose water in the Vanderbilt University Insectary (33, 34). For stock propagation, 5 to 7-d-old mated females were blood fed for 30 to 45 min using a membrane feeding system (Hemotek) filled with debrinated sheep blood purchased from Hemostat Laboratories. Mosquito larvae were reared in distilled water at 27 °C under the standard 12 h light/12 h dark cycle, with ~300 larvae per rearing pan in 1 L H2O. The larval food was made from 0.12 g/mL Kaytee Koi food and 0.06 mL yeast in distilled water and subsequently incubated at 4 °C overnight for fermentation. For first and second instar larvae, 0.08 mL larval food was added into the water every 24 h. The An. coluzzii effector line (QUAS-mCD8:GFP) was a generous gift from the laboratory of C. Potter at The Johns Hopkins University School of Medicine, Baltimore, MD (available via BEI Resources, NIAID, NIH: An. coluzzii, Strain QUAS-mCD8:GFP, Eggs, MRA-1301).

Mosquito Mutagenesis. CRISPR-Cas9 gene editing in An. coluzzii was carried out as previously described (35). The CRISPR gene-targeting vector was the generous gift from the laboratory of Andrea Crisanti at Imperial College London, London, UK (36). The single-guide RNA (sgRNA) sequence was designed by CHOPCHOP (37) to target the first exon of AcrIr76b gene (ACM033225). The complementary oligos (Ir76b_sgRNA_F/Ir76b_sgRNA_R; SI Appendix, Table S1) were artificially synthesized (Integrated DNA Technologies) and subcloned into the CRISPR vector by Golden Gate cloning (New England Biolabs). The homologous constructs were built based on the pHD-DsRed vector (Kate O’Connor-Giles, Brown University, Providence, RI; Addgene plasmid #51434; n2t.net/addedge:51434; RRID:Addgene 51434), where the 2-kb homologous arms extending either direction from the double-strand break (DSB) site were PCR amplified with the customized oligonucleotide primers: Ir76b_Aar1/76b_Aar_R; Ir76b_Sal1/Ir76b_Sal_R (SI Appendix, Table S1) and sequentially inserted into the Aa1/Sal restriction sites on the vector.

The microinjection protocol followed a previous study (16, 38). In brief, newly laid (~1-h-old) embryos of the wild-type An. coluzzii were immediately collected and aligned on a filter paper moistened with 25 mM sodium chloride solution. All the embryos were fixed on a coverslip with double-sided tape, and a drop of halocarbon oil 27 was applied to cover the embryos. The coverslip was further fixed on a slide under a Zeiss Axiosvert 35 microscope with a 40x objective. The microinjection was performed using Eppendorf FemtoJet 5247 and quartz needles (Sutter Instrument). The gene-targeting vector and the homologous template were cojected in 300 ng/mL each. Injected embryos were subsequently placed in deionized water with artificial sea salt (0.3 g/L) and thereafter reared under normal Vanderbilt University insectary conditions.

First-generation (G0) injected adults were separated based on sex and crossed to 5x wild-type sex counterparts. Their offspring (F1) were screened for DsRed-derived red eye fluorescence. Red-eyed F1 males were individually crossed to 5x wild-type females to establish a stable mutant line. DNA extraction was performed using DNeasy Blood and Tissue kits following the manufacturer’s instruction (Qia-gen), and genomic DNA was used as templates for PCR analyses of all individuals (after mating) to validate the fluorescence marker insertion using primers that cover DSB sites (Ir76b_F/Ir76b_R; SI Appendix, Table S1). Salient PCR products were sequenced to confirm the accuracy of the genomic insertion. The heterozygous mutant lines were thereafter back-crossed to the wild-type An. coluzzii for at least five generations before putative homozygous individuals were manually screened for DsRed-derived red eye fluorescence intensity. Putative homozygous mutant individuals were mated to each other before being killed for genomic DNA extraction and PCR analyses (as above) to confirm their homozygosity.

To generate the AcrIr76b-QF2 driver line for Q system, T2A-QF2_3xP3:DsRed element was inserted into the Ir76b coding region through CRISPR-mediated homologous recombination. DSB was induced using the CRISPR gene-targeting vector described above. The homologous template contains T2A-QF2_3xP3:DsRed element, which was amplified from the plk301-T2A-QF2 HDR plasmid (39); Leslie Vosshall, Rockefeller University, New York, NY; Addgene plasmid #130667; n2t.net/addedge:130667; RRID:Addgene 130667) and a 2-bp homologous arms, was constructed using NEBuilder HiFi DNA Assembly kit (NEB) (Ir76b_LArm_F/Ir76b_LArm_R); Ir76b_RArm_F/Ir76b_RArm_R; Ir76b_T2A_F/Ir76b_T2A_R; SI Appendix, Table S1). From the left homologous arm immediately preceding the T2A, 2 bp were removed to keep the sequence in-frame. Red-eyed F1 mosquitoes were backcrossed for three generations and then crossed to the effector line to acquire progeny for Ir76b localization studies.

Whole-Mount Imaging. Larval offspring resulting from crosses between the heterozygous AcrIr76b-QF2 driver line and the QUAS-mCD8:GFP effector were screened for eye-specific expression of DsRed. Larval head appendages as well as whole adult antennae, maxillary palps, proboscis, and tarsi from 4- to 6-d-old adult females were dissected into 4% formaldehyde in PBST (0.1% Triton X-100 in phosphate-buffered saline) and fixed on ice for 30 min in the case of the tarsi and larval head appendages or at 4 °C for 24 h for adult antennae. Samples were thereafter washed 3X in PBST for 10 min each and directly transferred onto Superfrost plus slides (VWR Scientific) and mounted in Vectashield fluorescent medium (Vector Laboratories). Confocal microscopy images at 1024 × 1024
pixel resolution were collected with an Olympus FV-1000 instrument equipped with a 100X oil objective at the Vanderbilt University Cell Imaging Shared Resource Core. Laser wavelengths of 405 nm, 488 nm, and 543 nm were used to detect DAPI, green fluorescent protein (GFP), and Cy3, respectively.

Immunohistochemistry. Antibody staining was performed as previously described (10, 16). Antennae, labella, and maxillary palps were dissected into 4% formaldehyde in PBST and fixed on ice for 30 min. Samples were washed 3x in PBST for 10 min each and then embedded in Tissue Freezing Medium (General Data Company Inc.). Cryosections were obtained at −20 °C using a CM1900 cryostat (Leica Microsystems). Samples were sectioned at ~10 μm and transferred onto Superfrost plus slides (VWR Scientific). Slides were air-dried at room temperature (RT) for 30 min and fixed in 4% formaldehyde in PBST for 10 min, followed by 3x rinsing in PBST for 10 min each. Thereafter, 5% normal goat serum (Sigma-Aldrich) in PBST was applied, and the slides were blocked in the dark at RT for 1 h in HybriWell sealing chambers (Grace Bio-Labs). Primary antibody Rabbit α-Orco was diluted 1:500 in 5% normal goat serum in PBST and applied on the slides and incubated overnight at 4 °C.

After primary antibody staining, slides were washed 3x in PBST for 10 min each and stained with secondary antibody Goat α-Rabbit-Cy3 (Jackson ImmunоReaearch) 1:500 in 5% normal goat serum PBST for 2 h at RT and then rinsed 3x. Nuclei were stained with 300 nM DAPI (Invitrogen) at RT for 10 min. Slides were briefly washed and mounted in Vectashield medium (Vector Laboratories). Whole-mount samples were dissected into 4% formaldehyde in PBST and fixed on ice for 30 min. After 3x washing in PBST for 10 min each, the samples were transferred onto slides and mounted in Vectashield medium (Vector Laboratories). Confocal microscopy images were collected as detailed above for the whole mounts.

Transcuticular Electrophysiology/Mass Measurements. Electroantennogram (EAG) and electroladleograph (ELG) recordings were conducted as previously detailed, with modifications (7, 34). Female mosquitoes (4 to 10 after eclosion) with legs and wings removed were dissected on glass slides using double-sided tape. The last segment of antenna with the tip partially cut was subsequently connected to a recording glass electrode filled with Ringer solution [96 mM NaCl, 2 mM KCl, 1 mM MgCl2, 1 mM CaCl2, and 5 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (Hepes), pH 7.5], to which a silver wire was in contact to complete the circuit with another glass reference electrode similarly connected into the compound eye of the female. The antennal preparation was continuously exposed to humidified, charcoal-filtered airflow (1.84 L/min) transferred through a borosilicate glass tube (inner diameter, 0.8 cm) using a stimulus controller (Syntech), and the open end of the glass tube was located 1 cm from the antennal preparation. All chemicals were diluted to 10−3 (vol/vol) working solutions in paraffin oil except for lactic acid, acetic acid, and dimethylamine, which were diluted in ddH2O. For each chemical, a 10-μL aliquot was applied onto a filter paper (3 × 50 mm), which was then inserted into a Fasteur pipette to create the stimulus cartridge. A sample containing the solvent (paraffin oil or water) alone served as the control. The airflow was maintained at a constant 20 mLs throughout the experiment. Purified and humidified air was delivered to the preparation through a glass tube (10-mm inner diameter) perforated by a small hole 10 cm away from the end into which the tip of the Fasteur pipette could be inserted. The stimulus was delivered to the sensilla by inserting the tip of the stimulus cartridge into this hole and diverting a portion of the air stream (0.5 L/min) to flow through the stimulus cartridge for 500 ms using a stimulus controller (Syntech). The distance between the end of the glass tube and the antennae was ≤1 cm.

Mating Bioassay. Newly emerged wild-type females and males were segregated on day 1 to ensure no mating occurred. Three-day-old females (15/bucket) and males (10/bucket) were then placed in a rearing bucket and allowed to mate freely for 5 d. All surviving females were then collected, and their spermatheca were dissected under a compound microscope. The spermathecae were then placed in the buffer (145 mM NaCl, 4 mM KCl, 1 mM MgCl2, 1.3 mM CaCl2, 5 mM D-glucose, and 10 mM Hepes (43) with 300 mM DAPI, and a coverslip was used to gently press and break the spermathecae to release the sperm. As in previous studies (15), the spermathecae were examined under the 1,000X compound microscope to assess the insemination status. The insemination rate was calculated by dividing the number of inseminated females by the total number of females in each bucket.

Blood-Feeding Bioassays. Initial blood-feeding bioassays were carried out with 6- to 8-d-old wild-type and Actλ76b mutant females between ZT11 and ZT2, during which An. coluzzii was shown to be behaviorally active. Mosquitoes were starved, with only water access for 24 h prior to the bioassay. Defibrinated sheep blood was stored at 4 °C and used within 2 wk of purchasing (Hemostat Laboratories). Human foot odorants were provided as cloth strips cut from cotton socks that had been continually worn for 3 to 5 d by a 30-yr-old male volunteer and thereafter incubated overnight at 37 °C in a sealed Ziploc plastic bag (SC Johnson). The acquisition and use of human foot odorants in these studies was reviewed by Vanderbilt’s Institutional Review Board (IRB), which deemed them exempt from requiring human subject research approval. For each replicate, 25 to 30 female mosquitoes were released into a 32-oz container, the top of which was covered with a net through which the mosquito probosces could
penetrate to feed but not to escape. A single membrane feeder (Hemotek) that heated the blood to 37 °C was then placed on the net. A mini camera (GoPro) was set at the bottom of the container to record the landing activity of mosquitoes. A human volunteer gently exhaled into the container for 25 min and then were immediately anesthetized at −20 °C to assess the number with blood-engorged abdomens, indicative of successful blood feeding. The assay containers and videos were also analyzed post hoc by volunteers blinded to the experimental genotypes, who manually counted both blood-engorged mosquitoes and the number of landings onto the feeder during the assay. The total landing count was divided by the total mosquito number in each assay to calculate the landing number per mosquito. Extended blood-feeding assays for wild-type and mixed populations of AcrIr76b hetero/homogzygotic mutants were similarly conducted for 120 min (2 h) using both sheep and human blood obtained from a donor volunteer through the Vanderbilt Center for Immunobiology (VCI). The donor provided informed consent, and this sample was provided by the VCI under IRB approval. Blood samples were de-identified prior to use in our study. Human odorants were obtained from the well-worn incubated cotton socks of a 64-year-old male volunteer (as before, the acquisition and use of human foot odorants was reviewed and deemed exempt from IRB approval for human subject research). Here, mosquitoes were allowed to feed freely for full duration of the assay and then were immediately anesthetized at −20 °C to assess the number with blood-engorged abdomens, indicative of successful blood feeding. The genotypes of individual AcrIr76b hetero/homogzygotic mosquitoes from these assays were validated post hoc by PCR.

Capillary Feeder (CAFE) Bioassay. The CAFE bioassay was conducted following a previous study, with minor modifications (15, 44). Each trial started at ZT12 and ended at ZT18 for 6 h. Four 4- to 8-d-old mosquitoes were provided with water but otherwise fasted for 22 h before being anesthetized on ice briefly and placed into a Drosophila vial (24.5 × 95 mm; Thermo Fisher Scientific). A borosilicate glass capillary (181008-3; World Precision Instruments) was filled with 10% sucrose water and embedded into a cotton plug. The vial opening was then blocked with the cotton plug, and the capillary was placed slightly protruding from the plug into the vial for mosquitoes to feed on. The sugar level in the capillary was compared before and after each trial to generate the initial sugar consumption value. At least four control vials with no mosquitoes inside were used to assess the evaporation at the same time. The initial sugar consumption value. At least four control vials with no mosquitoes

Results

AcrIr76b Expression in Antennal Grooved Pegs. In order to identify the types of antennal sensilla in which AcrIr76b activity is salient, the spatial expression pattern of AcrIr76b was established using the binary expression Q system (16, 45), which provides more sensitivity than fluorescence in-situ hybridization–based methods previously used (11). This system requires genetic crosses between a QUAS-GFP effector line and an AcrIr76b promoter-QF2 (AcrIr76b-QF2) driver line, resulting in GFP fluorescence in AcrIr76b-expressing cells. While previous anopheleine driver lines were developed by integrating promoter–QF2 constructs into predefined or random genomic sites (16, 45), those promoters were presumptive as they were derived from selected regions of 5′ upstream sequences without precise elucidation of the required regulatory sequences. As such, those promoters are potentially error prone and artificial, leading to localizations that are compromised by over-/underexpression from fragmented or otherwise partial regulatory information. To overcome this limitation, we incorporated a newly developed approach using CRISPR-mediated homologous recombination to insert a T2A-QF2-3xP3-DsRed element into the AcrIr76b locus at the first exon (Fig. 1A) (39), and the knock-in was subsequently confirmed by means of genomic PCR (Fig. 1A–C) and sequencing. The T2A peptide induces ribosomal skipping, which facilitates the unbiased expression of QF2 driven by endogenous, fully intact AcrIr76b regulatory sequences (46). For localization studies, F1 progeny derived from crosses between appropriate parental driver and effector lines would therefore express the QF2 transcriptional factor that specifically binds to the QUAS activation sequence to drive expressions of the visual marker GFP in all An. coluzzii cells that normally express AcrIr76b.

In these studies, extensive GFP labeling was observed in antennal grooved pegs of 4- to 6-d-old adult female An. coluzzii, which would be expected to be actively seeking blood meals. Whole-mount examination of the 2nd to 13th antennal flagello-

Generation of AcrIr76b Null Mutants. The AcrIr76b null mutant line was generated using the CRISPR-Cas9 system together with the DSB-specific homology template in order to knock in a 3xP3-DsRed visible marker construct at the AcrIr76b DSB site. The sgRNA was chosen to target the first exon of AcrIr76b to produce an early stop codon that leads to a dysfunctional protein (Fig. 2A). In total, 376 preblastoderm embryos were microinjected with the homologous template and the CRISPR targeting vector, in which Cas9 expression is regulated by the vtx2 promoter to specifically induce germ cell mutagenesis. Of the 17 larvae (4.5%) that survived the injection, two female and two male adults were able to successfully eclose and were designated as G0. These adults were then collectively crossed with the wild-type population, and the progeny (F1) were screened for the presence of red fluorescence in the eye and ventral nerve cord that would be expected if they contained a 3xP3-DsRed insert. At least one of the G0-injected adults produced red-eyed F1 offspring,
which indicates a high mutagenesis efficiency (≥25%). Red-eyed F1 males were individually back-crossed with wild-type females, after which their AcIr76b mutant genotype was confirmed by genomic PCR (Fig. 2A–C) and DNA sequencing. Homozygous mutants were generated by crossing heterozygotes with progeny selected based on their high intensities of red fluorescence, and thereafter genomic PCR (Fig. 2C) and DNA sequencing were carried out to confirm phenotypic assessments.

EAGs Revealed Increased Responses to Amines in AcIr76b Mutants. In order to initially investigate the function of AcIr76b in adult antennal olfactory responses, transcuticular EAG studies were carried out to broadly compare high-concentration (10⁻¹ vol/vol dilutions) responses between AcIr76b⁻/⁻ and wild-type females against an odorant panel comprising acids, alcohols, aldehydes, amines, and esters. Surprisingly, in contrast to response deficits typically seen in other olfactory coreceptor mutants studies (21, 22, 30), the EAG responses to several amines, including 3-pyrroline, pyrrolidine, 3-methylpiperidine, and 1-butylamine were significantly higher in AcIr76b⁻/⁻ females than wild type (Fig. 2D and E). In addition, EAG studies failed to reveal a significant impact on responses of AcIr76b⁻/⁻ females to a panel of acid stimuli (Fig. 2E). Comparative dose–response studies revealed that the enhanced EAG responses observed in AcIr76b⁻/⁻ females were restricted to high-concentration stimulations (SI Appendix, Fig. S3A–F). In addition to depolarization (downward) responses, we occasionally observed hyperpolarization potentials (upward) EAG responses to butyric acid, ammonia, and valeric acid. In total, 1/7 and 2/7 recordings from wild-type females demonstrated hyperpolarization potentials to stimulation with butyric acid and valeric acid, respectively. AcIr76b⁻/⁻ females gave rise to hyperpolarizing EAG responses.

Fig. 1. Localization of AcIr76b in chemosensory appendages. (A) AcIr76b-QF2 driver line schematics. The T2A-QF2-3xP3-DsRed element was inserted into the first exon of AcIr76b via CRISPR-mediated homologous recombination. A pair of primers (Ir76b_F and Ir76b_R) were used for genomic PCR validation. The wild type produces a 442-bp amplicon, whereas the AcIr76b-QF2 driver allele gives rise to a 3,011-bp amplicon. (B) A pair of primers (Ir76b_F2 and DsRed_R) were used to amplify the region outside the homology arm. The AcIr76b-QF2 driver allele gives rise to a 3,947-bp amplicon, whereas the primers are nonspecific for the wild type. (C) Agarose electrophoresis of genomic PCR validation of wild type (WT), heterozygous driver (Ir76b-QF2⁺/−), and homozygous driver (Ir76b-QF2⁻/⁻) with Ir76bF/Ir76bR (F/R) and Ir76bF2/DsRedR (F2DR) primers. (D and E) Representative bright-field image (Left) with GFP fluorescence overlay confocal z-stacks (Right) of whole-mount female antennae indicating AcIr76b is expressed in grooved pegs on sixth flagellomere (D) and 13th flagellomere (E) (highlighted by arrows). (F) A representative confocal z-stack of a whole-mount female antenna showing AcIr76b is expressed in the cell body and dendrites of coeloconic sensilla (highlighted by dashed lines). (G) A representative confocal optical section of the female antennae immunohistochemically labeled with β-Orco antisera (red) and AcIr76b (green), indicating that AcIr76b and Orco are localized in distinct cells. (H) A representative confocal optical section of a female antenna showing AcIr76b and Orco are colocalized in a cell (highlighted by an arrow and enlarged within dashed lines). (I) A representative confocal z-stack of a whole-mount female labellum showing AcIr76b is expressed in dendrites in T1 and T2 sensilla (highlighted by arrows; scale bars, 20 μm). (J) A representative confocal optical section of a female labellum showing AcIr76b and Orco are colocalized in a cell (highlighted by an arrow and enlarged within dashed lines). Nuclei were labeled with DAPI. Scale bars, 10 μm.
to ammonia (1/7 recordings), butyric acid (2/7 recordings), and valeric acid (6/7 recordings). In any case, statistical analyses of both response profiles revealed no significant differences between the wild-type and *AcIr76b*/*AcIr76b* females to these stimuli (SI Appendix, Fig. S3 G and H).

**Elevated Single Sensillum Responses in *AcIr76b*/*AcIr76b* Grooved Pegs.** The localization of antennal *AcIr76b* suggests that the increased EAG responses to amines are most likely due to elevated neuronal responses in populations of grooved pegs. To test this hypothesis, SSR studies to characterize and compare response profiles of *AcIr76b*/*AcIr76b* and wild-type females were carried out across a randomized sampling of 14 antennal grooved pegs across the three distal-most (10th to 13th) antennal flagellomeres where the highest density (~40) of these sensilla are found (14). These data reveal a range of wild-type responses to several amines that include both excitation and inhibition (Fig. 3 A and B). Interestingly, while *AcIr76b*/*AcIr76b* mutants displayed wild-type levels of spontaneous (background) neuronal activity (Fig. 3C) as well as responses to pyridine, 2-methyl-2-thiazoline, and phenylethylamine, we observed significantly elevated excitation elicited by pyrrolidine, 3-pyrroline, 3-methylpiperidine, and 1-butylamine compared with the wild-type responses, with only a single group of spikes as observed in the olfactory responses to amines from antennal trichoid sensilla when comparing *AcIr76b*/*AcIr76b* mutants and similarly aged wild-type female *An. coluzzii* (Fig. 3E).

Randomized SSR responses to acids and amines in peg-in-pit antennal coeloconic sensilla across the second to eighth flagellomeres were also carried out to explore potential *AcIr76b* function (Fig. 4A). While wild-type and *AcIr76b*/*AcIr76b* female coeloconic sensilla displayed similar levels of spontaneous (background) neuronal activity (Fig. 4B), *AcIr76b*/*AcIr76b* females became indifferent to heptanoic acid, phenylacetic acid, pyridine, pyrrolidine, 2-methyl-2-thiazoline, and 1,4-diaminobutane in contrast to robust wild-type responses (Fig. 4C and SI Appendix, Fig. S5A). Noticeably, acetic acid, hexanoic acid, pentanoic acid, 3-methylpiperidine, and spermidine elicited distinct, yet nonsignificant, responses between wild-type and *AcIr76b*/*AcIr76b* females (Fig. 4C and SI Appendix, Fig. S5A). *AcIr76b*/*AcIr76b* mutants showed decreases in poststimulus OFF responses after stimulation with pyridine and phenylacetic acid (Fig. 4D and SI Appendix, Fig. S5B). While *AcIr76b*/*AcIr76b* females exhibited modest, albeit nonsignificant, increases in acetic acid responses compared with wild types (SI Appendix, Fig. S5A), these mutants displayed significantly higher poststimulus OFF responses to acetic acid (SI Appendix, Fig. S5B).

**Peripheral Electrophysiology Across Accessory Chemosensory Appendages.** To further confirm the specificity of the antennal phenotype, we carried out SSR studies on female *An. coluzzii* maxillary palp capitate pegs (cp), which are the sole sensillar class found on that appendage (5). While previous RNA sequencing–based transcriptome profiles of the *An. coluzzii* female maxillary palps uncovered *AcIr76b* transcripts (50, 51), no GFP-labeled cells were identified on female *Ir76b-GFP* progeny, which likely reflects the sensitivity differences between
these two approaches. In that light, it was not surprising that the well-characterized GR- and OR-mediated responses of the cpA neuron to CO₂ and the cpB/C neurons to 1-octen-3-ol, 2,4,5-trimethylthiazole, respectively (5), were not affected in AcIr76b⁻/⁻ mutants (SI Appendix, Fig. S6A).

Lastly, in light of extensive AcIr76b expression across the labellum (Fig. 1G), we also employed the ELG, an EAG-like transcuticular sampling of peripheral neuronal activity on the labellum (7), to assess responses to a panel of amines to determine whether AcIr76b might play a role in olfactory responses on that appendage. Surprisingly, in light of the robust AcIr76b labellum expression, these studies revealed no significant differences between the amine response profiles of the An. coluzzii wild-type and AcIr76b⁻/⁻ female labellum (SI Appendix, Fig. S6B).

**AcIr76b Expression in a Distinct Antennal Organ in Larvae.** AcIr76b expression was also examined across the larval antennae of An. coluzzii, which is the principal sensory appendage of this aquatic preadult life stage (41, 52). In addition to robust labeling of larval antennal neuronal cell bodies, which are clustered within the antennal shaft (SI Appendix, Fig. S7A), the AcIr76b promoter appears to drive GFP labeling of dendrites that specifically innervate the sensory peg organ (SI Appendix, Fig. S7A). The larval sensory peg is a distinctive uniporous apical appendage on the antennae that has been hypothesized to play gustatory roles (SI Appendix, Fig. S7B) (52, 53). AcIr76b localization to the larval sensory peg dendrites is distinct from the dendritic localization of An. coluzzii ORs to the larval sensory cone, where they are associated with olfactory signals (41, 52).

Despite their proximity at the apical tip of the larval antennae, the activity of the sensory peg is distinct from that of the OR-associated sensory cone, which acts as the primary larval olfactory organ in *Anopheles* (41, 52). While technical limitations have thus far precluded direct recording of neuronal activities from the uniporous larval peg, we nevertheless recently carried out a comprehensive electrophysiological analysis of peripheral larval sensory cone neuronal responses to a wide range of volatile stimuli (41). These SSR-based methods were used to examine the functionality of wild-type and AcIr76b⁻/⁻ larval sensory cones to further narrow its role to the sensory peg. As expected, based on the absence of AcIr76b-associated sensory cone dendrites, these recordings failed to reveal any significant differences between wild type and AcIr76b⁻/⁻ mutants insofar as background neuronal activity (SI Appendix, Fig. S7C) or responses to a panel of amines (SI Appendix, Fig. S7D).

**Mating and Blood-Feeding Deficits in AcIr76b Mutants.** In addition to their impact on peripheral electrophysiology in adult females, AcIr76b⁻/⁻ mutants display a range of interesting behavioral and reproduction-related deficits and, as a result, cannot self-propagate under standard laboratory rearing protocols. To investigate this phenotype, we took advantage of a previously developed insemination-based mating bioassay (15) to reveal that self-mated AcIr76b⁻/⁻ mutants display significantly impaired insemination rates when compared with their wild-type counterparts (Fig. 5A). To investigate this phenotype further, mating studies were conducted in which AcIr76b⁻/⁻ females were replaced with their wild-type counterparts. Here, wild-type females successfully mated with AcIr76b⁻/⁻ males to...
fully rescue the mutant mating/insemination deficits. In addition, studies that paired wild-type males with $AcrIr76b^{-/-}$ females had severe mating deficits (Fig. 5A), indicating that $AcrIr76b$ plays a female-specific role in $An. coluzzii$ mating. To further investigate potential causes of these mating deficits, the gross mass of carcasses from individual females of both genotypes were compared. In these studies, $AcrIr76b^{-/-}$ females were found to have significantly less mass than wild-type females (Fig. 5B), perhaps reflective of reduced fitness that, in part, might account for these mating deficits.

We next asked what other aspects of the female reproductive pathway might underlie the mating deficits and sterility of the $AcrIr76b^{-/-}$ mutants. To address this question, we utilized a simple digital video-based blood-feeding bioassay (54) to examine the blood-meal specific (background) neuronal activity. Nonparametric t tests suggest no significant differences between $AcrIr76b$ mutants and wild type. (C) Averaged female coeloconic sensillum SSR responses. Multiple t tests using Holm-Sidak method ($n = 25$ or $26$) suggest responses to specific acids and amines are significantly different in $AcrIr76b$ mutants than in wild type. Significance levels are depicted with asterisks; $P$ value $< 0.05$ (*); $P$ value $< 0.001$ (***). Error bars = SEM, ns, not significant.

As is the case for other mosquitoes, $Anopheles$ blood feeding is a highly specialized behavior that is quite distinct from nectar (sugar) feeding (58). Subsequent to host seeking and landing, another critical aspect of mosquito blood feeding is the process of blood ingestion (uptake), which occurs through the styllet (58). The styllet is a highly specialized component of the labelum that pierces the skin of blood-meal hosts to make direct contact with blood and is used solely for blood feeding. Recent studies in $Ae. aegypti$ have illustrated the roles of $Ir7a$ and $Ir7f$, which are expressed in styllet gustatory neurons where they are specifically responsible for mediating the direct chemosensory (taste) responses to blood-meal components (59). In light of the association of the $DmIr76b$ ortholog in amino acid responses in $Drosophila$ (26), we hypothesized that $AcrIr76b$ also acts as a gustatory IR coreceptor and, in that context, would be expressed on the styllet of $An. coluzzii$. Indeed, crosses between $AcrIr76b$-QF2 driver and QUAS-GFP effector lines revealed robust expression of $AcrIr76b$ in the styllet (Fig. 5F). In order to examine the blood-meal specificity of this feeding deficit
phenotype, we used the well-established CAFE capillary feeding assay (15, 44) to examine sugar feeding. These studies confirmed AcIr76b−/− females can maintain wild-type levels of sugar feeding via their proboscis (Fig. 5G) that are consistent with a highly specialized blood-feeding functionality of AcIr76b on the stylet of An. coluzzii.

Discussion

Acting together or independently, anopheline IRs, GRs, and ORs are critical molecular components of the signal transduction processes that initiate diverse chemosensory processes underlying various elements of the mosquito life cycle. While many of the particulars of these relationships remain unclear, it is apparent that the collective functionality of these elements has direct as well as indirect/modulatory roles in mediating the activation or inhibition of chemosensory neurons. For example, the Ir25a coreceptor is coexpressed with Gr3 and Orco in Aedes aegypti and Drosophila, respectively (47, 48). In Aedes, knocking out Gr3 on the CO2-sensitive cpA maxillary palp neuron removes CO2 sensitivity but importantly does not block responses to amines mediated by Aelr25a in the same neurons (47). In Drosophila, while Orco null mutations completely abolished OSN activity, Dmlr25a mutations resulted in mostly nonaltered or only partially reduced/increased responses, suggesting that Dmlr25a plays a regulatory role when coupled with ORs instead of acting as a canonical coreceptor (48). We initially focused on the role of AcIr76b in adult An. coluzzii females that are actively host seeking for a blood meal. In these insects, robust Ir76b expression occurred in the antennal grooved pegs (Fig. 1), which are considered homologous to Drosophila coeloconic sensilla (60), as well as in T1 and T2 sensilla of the labellum (Fig. 1) and across the tarsi (SI Appendix, Fig. S2). We also examined its expression in the larval chemosensory system, which is housed on the antennae. Here, in contrast to the dendritic localization of An. coluzzii larval ORs, AcIr76b specifically labeled dendrites innervating the sensory peg, which is an apical uniporous structure associated with larval gustatory responses.

The availability of CRISPR-Cas9–mediated gene targeting in An. coluzzii allowed us to examine loss-of-function phenotypes resulting from AcIr76b null mutations. In addition to defective coeloconic SSR responses to amines, consistent with that observed in Drosophila (24), CRISPR-generated AcIr76b null mutants exhibited significantly enhanced antennal responses to amines in both EAG and grooved peg SSR studies in contrast to A. aegypti Aelr8a mutants that entirely abolish olfactory sensitivity to acids (30). These data are similar to previous studies that

Fig. 5. Behavioral deficits of AcIr76b mutants. (A) Averaged insemination rate of females in different mating pairs. Mean values with different grouping letters were significantly different (n = 5; one-way ANOVA; P value < 0.05). (B) Averaged weights of the wild-type (WT) and AcIr76b−/− (Ir76b−/−) adult females. WT females weighed significantly more than Ir76b−/− with P value < 0.01 (**). (C) Average landing counts per female mosquito on the blood feeder during blood feeding. Nonparametric t test suggests no significant differences between AcIr76b mutants and wild type. (D) Averaged blood feeding rate of wild-type (WT), virgin wild-type [WT (Virgin)], and AcIr76b−/− (Ir76b−/−) females in a 25-min blood feeding bioassay. Mean values with different grouping letters were significantly different (n = 4 to 7; one-way ANOVA; P value < 0.05). (E) Averaged blood feeding rate of wild-type (WT), AcIr76b−/− (Ir76b−/−) heterozygotic, and AcIr76b−/− (Ir76b−/−) homzygotic females in a 120-min blood feeding bioassay. Mean values with different grouping letters were significantly different (n = 5 to 7; one-way ANOVA; P value < 0.05). (H) A representative confocal z-stack of female stylet showing AcIr76b expression. Scale bars, 10 μm. (I) Averaged sugar consumptions of female mosquitoes in the CAFE bioassay. Nonparametric t test suggests no significant differences between AcIr76b mutants and wild type. Error bars = SEM. F, female; M, male; ns, not significant.
identified enhanced gustatory responses to sugars in Drosophila Ir76b mutants (61) and suggest that Ir76b’s modulatory function is both context dependent and evolutionarily conserved. In An. coluzzii, this phenotype is restricted to the large population of non–Orco-expressing grooved pegs, where dramatic increases in amine-elicited spike frequencies are seen in AcIr76b null mutants, suggesting that AcIr76b is responsible for inhibition of neuronal activity in those wild-type neurons. The lack of a significant alteration of other SSR spikes (SI Appendix, Fig. S4) suggests this is direct intraneuronal inhibition rather than indirect interaction via ephaptic coupling or other nonsynaptic effects. Furthermore, these data are also consistent with the presence of another IR coreceptor that is primarily responsible for action potential generation. This is likely to be Ir25a in light of its association with Ir76b orthologs in Aedes and Drosophila (18, 47).

Our data provide in vivo evidence for the modulatory role of AcIr76b in antennal responses that is specific to amines and thus extend previous studies emphasizing the role of IRs in the detection and discrimination of amines (11, 62). Importantly, many compounds in this chemical class have been identified as human emanations and are components of odor blends that robustly attract blood-feeding anopheline mosquitoes (63, 64). In contrast to exclusive gustatory functions in Drosophila, mosquito labella also have olfactory capabilities (7, 49). However, despite extensive AcIr76b expression in the labellum of female An. coluzzii mosquitoes, we were unable to uncover any significant electrophysiological differences to volatile odorants between wild-type and mutant mosquitoes in ELG studies. Importantly, our data do not rule out an important gustatory role for AcIr76b in labellum responses to contact cues such as ammonia, carboxylic and amino acids as well as other compounds present in human sweat and in blood meals themselves (65, 66). Future SSR and tip-recording studies investigating the role of labellum expression of AcIr76b in contact cues such as sugars in An. coluzzii will provide validation for this hypothesis. Inasmuch as blood feeding is paramount for An. coluzzii and other mosquitoes to reproduce and to acquire and transmit disease pathogens, the crucial role of Ir76b in blood feeding makes it a provocative target for the development of strategies to reduce mosquito vectorial capacity.

The restriction of larval AcIr76b supports a direct chemosensory role of the larval peg and, moreover, is consistent with our previous RNAi-based gene-silencing studies in An. coluzzii, demonstrating that larval behavioral responses to aqueous butyramine are mediated by AcIr76b (31). While we are currently unable to physiologically examine gustatory responses of the larval sensory peg to aqueous stimuli, it is nevertheless noteworthy that AcIr76b+/– mutant larvae display wild-type responses to a panel of volatile odorants. Taken together, these data indicate that the AcIr76b-associated behavioral sensitivity of An. coluzzii larvae to aqueous butyramine and perhaps other related compounds (31) is a gustatory process mediated through the sensory peg.

Over and above the expected peripheral olfactory impacts of knocking out AcIr76b, we have identified novel roles for Ir76b-based signalling in anopheline reproductive pathways. To begin, female homozygotic AcIr76b+/– mutants displayed significant reductions in insemination rates that persisted when mated with either wild-type or mutant males. In contrast, AcIr76b+/– females maintained wild-type–level insemination rates when paired with wild-type females (Fig. 5A). While these data demonstrate this phenotype is female specific, its mechanistic basis remains unclear. While AcIr76b+/– females are significantly smaller than their wild-type counterparts, which may contribute to their reduced overall fitness and thereby impact mating and other behaviors, we hypothesize that the Ir76b-related mating defect is primarily linked to its impact on direct or volatile-based chemosensory processes on the labella and tarsi of An. coluzzii, where AcIr76b expression is pronounced. Indeed, DmIr76b is involved in multiple gustatory pathways in Drosophila (25, 26), where the labella and tarsi are recognized as gustatory appendages (44, 67, 68) and where multiple studies have identified IRs on those gustatory appendages that directly promote mating (67, 69, 70). While the chemosensory receptors involved in mosquito mating have not been identified molecularly, the tarsi of anopheline mosquitoes are known to be essential for mating where they presumably detect contact mating pheromones (71–73). It is therefore reasonable to suggest that mating deficits observed in AcIr76b+/– females could be caused by disruption of recognition pathways that are dependent on Ir76b-mediated gustatory signaling.

While mating deficits represent a significant AcIr76b+/– phenotype that impedes the propagation of homozygous mutant lines, the total absence of successful blood feeding of AcIr76b+/– females is profound, mandating heterozygotic maintenance of these lines. Despite the ability of these mutants to exhibit wild-type levels of sugar feeding and, importantly, to be fully able to locate, alight on, and even probe blood-containing membrane feeders that are supplemented with human foot-odor blends and CO2, AcIr76b+/– females completely fail to take-up (ingest) blood meals. In Ae. aegypti, the labial stylets, which are essential for gustatory sampling (tasting) of blood, house neurons that respond to whole blood that is rich in amino acids (74) and more specifically to adenosine triphosphate, NaHCO3, and NaCl (59). More importantly, while the cognate IR coreceptors responsible for these responses remain uncharacterized, two Aedes IRs—Ir7a and Ir7f—acting as “tuning” receptors are specifically expressed on the stylenet neurons, where they directly recognize these blood-specific constant cues to activate the uptake of blood to complete blood feeding (59). In light of the robust expression of AcIr76b in the stylenet of adult females, it is reasonable to postulate that AcIr76b is the gustatory IR coreceptor that is directly involved in blood tasting in An. coluzzii. That AcIr76b+/– mutations specifically block the uptake of blood by female mosquitoes that have located, alighted on, and probed membrane feeders provides validation for this hypothesis. Inasmuch as blood feeding is paramount for An. coluzzii and other mosquitoes to reproduce and to acquire and transmit disease pathogens, the crucial role of Ir76b in blood feeding makes it a provocative target for the development of strategies to reduce mosquito vectorial capacity.

Data Availability. All study data are included in the article and/or SI Appendix and therefore fully accessible upon publication.

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10 of 11 https://doi.org/10.1073/pnas.2112385119 pnas.org
