Prototypical pacemaker neurons interact with the resident microbiota

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Pacemaker neurons exert control over neuronal circuit function by their intrinsic ability to generate rhythmic bursts of action potential. Recent work has identified rhythmic gut contractions in human, mice, and hydra to be dependent on both neurons and the resident microbiota. However, little is known about the evolutionary origin of these neurons and their interaction with microbes. In this study, we identified and functionally characterized prototypical ANO/SCN/TRPM ion channel-expressing pacemaker cells in the basal metazoan Hydra by using a combination of single-cell transcriptomics, immunochemistry, and functional experiments. Unexpectedly, these prototypical pacemaker neurons express a rich set of immune-related genes mediating their interaction with the microbial environment. Furthermore, functional experiments gave a strong support to a model of the evolutionary emergence of pacemaker cells as neurons using components of innate immunity to interact with the microbial environment and ion channels to generate rhythmic contractions.

The enteric nervous system (ENS) coordinates the major functions of the gastrointestinal tract (1). In all extant animals, the structurally conserved ENS is a diffuse nerve net located within the wall of the gastrointestinal tract. In prebilaterian animals, such as Hydra, the nervous system is structurally simple and thus has great potential to inform us about evolutionary ancient basic structural and functional principles of neural circuits (2) (Fig. L4). The principal function of the ENS is coordination of the rhythmic intestine motility, known as peristalsis, that occurs ubiquitously in the animal kingdom (Fig. L4) and is driven by rhythmic electrical pulses generated by pacemaker cells (3). In mammals, proper functioning of the intestinal cells of Cajal (ICC), that serve as pacemakers in the muscular wall of the gastrointestinal tract (4–6), is essential for normal gut motility (7–9). Dysfunction of the pacemaker system contributes to functional gastrointestinal disorders, such as irritable bowel syndrome (IBS), chronic constipation, and intestinal pseudo-obstruction (9–12).

Here, we discover prototypical pacemaker neurons in the ancient cnidian Hydra and provide evidence for a direct interaction of these neurons with the commensal microbiota. We uncover a remarkable gene-expression program conservation between the Hydra pacemaker neurons and pacemaker cells in Caenorhabditis elegans and the mammalian gut. We suggest that prototypical pacemaker cells emerged as neurons using components of innate immunity to interact with the microbial environment and ion channels to generate rhythmic contractions. The communication of pacemaker neurons with the microbiota represents a mechanistic link between the gut microbiota and gut motility. Our discoveries improve the understanding of the archetypical properties of the enteric nervous systems, which are perturbed in human dysmotility-related conditions.

Significance

The enteric nervous system (ENS) coordinates the major functions of the gastrointestinal tract (1). In all extant animals, the structurally conserved ENS is a diffuse nerve net located within the wall of the gastrointestinal tract. In prebilaterian animals, such as Hydra, the nervous system is structurally simple and thus has great potential to inform us about evolutionary ancient basic structural and functional principles of neural circuits (2) (Fig. L4). The principal function of the ENS is coordination of the rhythmic intestine motility, known as peristalsis, that occurs ubiquitously in the animal kingdom (Fig. L4) and is driven by rhythmic electrical pulses generated by pacemaker cells (3). In mammals, proper functioning of the intestinal cells of Cajal (ICC), that serve as pacemakers in the muscular wall of the gastrointestinal tract (4–6), is essential for normal gut motility (7–9). Dysfunction of the pacemaker system contributes to functional gastrointestinal disorders, such as irritable bowel syndrome (IBS), chronic constipation, and intestinal pseudo-obstruction (9–12).

The Ca2+-activated Cl− channel Anoctamin-1 (encoded by the ANO1 gene) and the voltage-gated Na+ channel Nav1.5 (SCN5A gene) represent molecular markers of the intestinal pacemaker cells in human and mice (13–15), and DNA variants in the corresponding genes have been shown to associate with increased risk of IBS (16–18). Ion channel dysfunction (channelopathy) appears to be a plausible pathogenetic mechanism in functional gastrointestinal disorders (19), as the transient receptor potential cation channel TRPM8 (known as the cold and menthol receptor) and other ion channels have also been implicated in IBS susceptibility and gut dysmotility (20–23). Spontaneous contractile gut activities are not only affected by microbes. In fact, there is evidence that bacterial population dynamics themselves are affected by the periodic stimulation (24). Previous studies in Hydra suggested that the rhythmic peristaltic movements of the body column are dependent on neurons (25) and that they are modulated by the host-associated microbiota since germ-free (GF) animals display reduced and less-regular contraction frequencies (26). Single-cell RNA sequencing (scRNA-seq) uncovered neuron-specific transcription signatures and the presence of distinct neuronal subtypes (27).

Little, however, is known about the nature of the neurons that generate peristaltic movement in a prebilaterian animal and how such prototypical neurons engage with the resident microbiota.
Here, we provide a definition of prototypical pacemaker cells, which integrates marker genes discovered in human dystomyot patients with the recent discovery that spontaneous contractile activities are affected by microbes. The functional experiments connected the rhythm generation and interactions with microbes at the level of this specific neuronal population. These findings shed new light on the evolution of pacemaker neurons, emphasize the role of the microbial environment in dystomyot, and underscore the importance of cross-species comparisons in tracking cell type evolution.

Results

Identification of Pacemaker Cells in Hydra Using Human Orthologous Genes. Previous studies in Hydra suggested that a subpopulation of neurons located in the head region (28–30) might have properties of pacemaker cells to control the regularity of spontaneous body contractions. To gain insight into this specific cell population, we first assessed the molecular and functional diversity of the neuronal populations by scRNA-seq using red fluorescent protein (RFP)-labeled neurons (Fig. 1 B–J and SI Appendix, Figs. S1 and S2). Similar to the analysis of the entire transcriptome, the expression profile of 112 transcripts coding for putative neurotransmitter receptors clearly identified seven distinct clusters of neurons and separated them from the stem cells and nonneuronal cells (Fig. 1 D–F). This indicates that each neuronal subpopulation is characterized by a specific set of neurotransmitter receptors (Fig. 1G and SI Appendix, Fig. S3). For example, most transcripts coding for putative nicotinic acetylcholine receptors (nAChR) were expressed almost exclusively in neuronal subpopulation N2 (Fig. 1G and SI Appendix, Fig. S3), while diverse homologs of muscarinic and N-methyl-D-aspartate glutamate receptors (mGlurRs and NMDARs), and γ-amino butyric acid type-A receptors (GABAARs) were enriched in the neuronal subpopulation N7 (Fig. 1G and SI Appendix, Fig. S3). Similarly, the expression profiles of 431 transcripts coding for ion channels clearly segregated GFP+/RFP− stem cells from the GFP−/RFP+ non-neuronal cells and the seven remarkably distinct clusters of GFP+/RFP+ neurons (Fig. 1H). In addition, each neuronal subpopulation was found to express a unique combination of neuropeptides (Fig. 1I and SI Appendix, Fig. S4). The Hym-355 neuropeptide precursor gene, for example, was exclusively expressed in the neuronal subpopulation N3, while Hym-176A transcripts were discovered predominantly in the N1 subpopulation. Most of RFamide precursor transcripts were found in subpopulation N6 (Fig. 1I and SI Appendix, Fig. S4). Genes coding for some other

Fig. 1. Single-cell transcriptome profiling uncovers the molecular anatomy of Hydra nervous system. (A) Emergence of the first nerve cells preceded the divergence of Cnidaria and Bilateria. Cnidarians possess structurally simple nervous systems and offer a great potential to reveal the fundamental structural and functional principles of neural circuits. Spontaneous rhythmic contractions are ubiquitously observed in Eumetazoa. (B) The Hydra body is made of three cell lineages: the ectodermal and endodermal epithelia separated by the extracellular matrix and the lineage of interstitial cells. The outer surface of the ectoderm is covered by a glyocalyx that serves as a habitat for symbiotic bacteria. The endodermal lining the gastric cavity is free of glyocalyx and stable microbiota. Two nerve nets made of sensory and ganglion neurons are embedded within both epithelia. (C) Genetic construct used to generate transgenic Hydra polyps and differentially label cells within the interstitial lineage by a combination of two fluorescent proteins: GFP expressed under a stem cell-specific nanos promoter (nosP), and RFP driven by the actin promoter (actP) active in terminal differentiated neurons. Both cassettes are flanked by the actin terminator (actT). (D) T-Distributed stochastic neighbor embedding (t-SNE) map constructed by dimensionality-reduction principal component analysis defined by highly covariable genes (Materials and Methods). A total of 928 cells were partitioned in 12 clusters and colored by their cell-type identities inferred from expressed pro- liferation and cell-type-specific marker genes (Datasets S5 and S6). (E) t-SNE map based on analysis of the entire transcriptome made of 116,186 transcripts segregates 12 clusters, including 7 subpopulations of neurons. Cells are color-coded by their phenotype captured by FACS upon sorting. (F) t-SNE map based on expression analysis of 112 transcripts coding for putative neurotransmitter receptors (Dataset S7). Seven neuronal populations are clearly segregated, indicating that each neuronal population is characterized by a specific set of receptors. (G) Heatmap illustrating expression of genes coding for putative nAChR, mGlur and NMDAR, and GABAAR within seven neuronal populations. Expression within the entire interstitial lineage is presented in SI Appendix, Fig. S3. Transcripts specifically up-regulated in the neurons are labeled red; superscript numbers indicate the nerve cell cluster (N1–N7) where the transcripts are significantly (adjusted P < 0.05) enriched. (H) t-SNE map constructed by expression analysis of 431 transcripts coding for putative ion channels (Dataset S8). Seven neuronal populations are clearly segregated, suggesting that each neuronal population is characterized by a specific set of channels. (I) Heatmap illustrates expression of 11 genes coding for main known neuropeptides in Hydra. Each neuronal population expresses a unique combination of neuropeptides. (J) In situ hybridization for marker genes strongly enriched in each of seven nerve cell clusters (N1–N7) (SI Appendix, Fig. S6) reveals that seven neuronal subpopulations reside in spatially restricted domains along the body column of Hydra (Scale bars, 100 μm upper panel, 25 μm lower panel.)
peptides, including Hym-331 and FRamide, were expressed in two or more neuronal subpopulations. Taken together, these observations indicate that the molecular identity of different subpopulations of neurons is determined by the specific expression of ion channels, neurotransmitter receptors, and neuropeptides.

Interestingly, based on the expression profiles of 364 transcript sequences coding for putative transcription factors (TFs), all seven subpopulations of neurons express a common set of TFs, which separates them from stem cells and nonneuronal cell types (SI Appendix, Fig. S5). The neuron-specific TF signature consists mainly of Zn-finger, homeodomain and helix–loop–helix DNA-binding proteins, including the Achaete-scute homologous TFs. Furthermore, each neuronal population is characterized by a combinatorial expression of few genes encoding other TFs, such as the homologs of Aristales, NeuroD, and Orthopedia (SI Appendix, Figs. S5 B and C and Dataset S1).

In situ hybridization for selected marker genes enriched in each of the seven neuronal subpopulations (Fig. 1 and SI Appendix, Fig. S6A) showed that the neuronal subpopulations in Hydra reside in spatially restricted domains along the body column. Neuronal clusters N1 were found confined to the foot region of the polyp (Fig. 1 J and SI Appendix, Fig. S6B). Cluster N2 was represented by a population of neurons located in the base of tentacles. N3-specific neurons were spread in the ectoderm resembling the entire Hydra body, whereas neurons of subpopulations N4 and N5 were found in the endodermal epithelial layer (Fig. 1 J and SI Appendix, Fig. S6B). Neurons from the cluster N6 were confined to the hypostome of a polyp, whereas neurons of the cluster N7 were restricted to the tentacles. Notably, most of the seven spatially restricted neuronal subpopulations contain both sensory and ganglion cells (Fig. 1 J). Analysis of marker gene expression (SI Appendix, Fig. S7) indicated a clear correspondence between the neuronal clusters identified by us and the subtypes reported by Siebert et al. (27).

To identify the pacemaker cells among the neurons in Hydra, we focused on a few human orthologs known to be either restricted in their expression to human ICCs (AN01 and SCNSA) (13, 14) or mechanistically involved in the control of gut motility via circular smooth muscle cell contractions (menthol sensitive Ca2+ channels such as TRPM8) (31) (Fig. 2 A–D). Homology search and phylogenetic analysis uncovered three Hydra genes coding for SCN-like ion channels, six homologs of AN01-like chloride channels, and four homologs encoding TRPM-like cation channels, which are remarkably similar to their human counterparts (Fig. 2 B–D and SI Appendix, Figs. S8–S10). Analysis of the single-cell transcriptome (Fig. 2 E and SI Appendix, Fig. S11) revealed that the expression of genes encoding SCN-, AN01-, and TRPM-like channels overall was very weak, often restricted to only a few cells. However, several of the transcripts coding for SCN- and AN01-homologs were more expressed in neurons (Figs. 2 E and SI Appendix, Fig. S11), with some of the transcripts specific for neuronal subpopulation N2, which is located at the base of tentacles (Figs. 1 J and E and SI Appendix, Fig. S11). Real-time PCR confirmed (Fig. 2 F) that most of the SCN-, AN01-, and TRPM-like ion channel genes are up-regulated in the head region. Transcripts of two of these genes, cluster41630 and cluster33856, coding for AN01-like and SCN-like channels, could be detected by in situ hybridization at the base of tentacles (Fig. 2 G and H). Immunohistochemical analysis using specific antibodies raised against synthetic peptides confirmed the presence of AN01-like and SCN-like channel proteins at the base of the tentacles (Fig. 2 I–K) in the subpopulation N2 domain (Fig. 1 J). High-magnification confocal microscopy identified the cells expressing SCN and AN01 channels as neurons (Fig. 2 L and M). Taken together, these observations indicate that the genes encoding SCN-, AN01-, and TRPM-like channels are expressed in a population of nerve cells resident in the subpopulation N2 at the base of tentacles (Fig. 2 N).

ANO1-, SCN-, and TRPM-like Channels Are Essential for Pacemaker Activity in Hydra. We next tested the role of the neuronal subpopulation N2-specific AN01-, SCN-, and TRPM-like channels in controlling the pacemaker-driven rhythmic spontaneous contractions in Hydra (Fig. 2 O–Q). Exposing polyps to AnIβ, a potent inhibitor of AN01 channels (32), resulted in both a twofold reduction of the contraction frequency (Fig. 2 Q and R) and also in less regular contractions compared to controls (Fig. 2 S). Similar results were obtained upon treatment of polyps with menthol, which activates TRPM8 channels in vertebrates (33), and lidocaine, which interferes with SCN-like ion channels (Fig. 2 Q–S). The results show that modulating the activity of the neuron-specific AN01-, SCN-, and TRPM-like channels in Hydra greatly disturbs the rhythmicity of spontaneous contractions. Since neuronal subpopulation N2 is also characterized by the expression of putative nAChRs (Fig. 1 G and SI Appendix, Fig. S3), we next tested the effects of tubocurarine (DTC), known as a potent antagonist of nAChRs (34, 35), on the frequency and rhythmicity of the spontaneous contractions in Hydra (Fig. 2 Q). We found that the presence of DTC strongly reduced the frequency and affected the regularity of the spontaneous contractions (Fig. 2 R and S). Other reflexes dependent on neural circuits, such as the feeding reflex (36), were not affected by most of the channel-specific inhibitors (SI Appendix, Fig. S12), indicating a specific role of the ion channels expressed in N2 neurons in controlling rhythmic body contractions. Inhibiting GABA_A receptors specifically expressed in neurons of subpopulation N7 and absent in the subpopulation N2 (Figs. 1 G and Q and SI Appendix, Fig. S3) had no effect onto the contraction pattern (Fig. 2 R and S) but strongly influenced the feeding response (SI Appendix, Fig. S12).

Together with the previous pharmacological findings (36), our observations unequivocally identify the pacemaker population N2 as cholinergic, and the neuronal population N7 controlling the feeding response as predominantly GABAergic. Notably, all cells within neuronal population N2 homogeneously expressed high levels of one of the innexin genes, cluster41630 (SI Appendix, Fig. S13). Innexins are the only components of gap-junction complexes known in Hydra (37, 38). Homotypic gap junctions established between neurons of the N2 subpopulation therefore may electrically couple these neurons and allow generation of a neural network with pacemaker properties. In a similar way, a net of neurons present in the peduncle of Hydra is likely coupled by gap junctions (39), and the epithelial cells of both the ectoderm and endoderm are electrically coupled by gap junctions (40–42). In sum, these data suggest that neurons of the N2 subpopulation express marker genes for gut dysmotility (Fig. 2 A–N) and that inhibition of these ion channels disturbs spontaneous and regular body contractions (Fig. 2 O–S). Therefore, neurons of the N2 subpopulation may act as pacemaker cells controlling the spontaneous contraction pattern. This is consistent with earlier electrophysiological recordings in Hydra (30) and the observations that removal of the head region results in loss of spontaneous contractile activity (28).

Pacemaker Neurons in Hydra Are Immunocompetent Cells. We have shown previously that neurons in Hydra secrete antimicrobial peptides (AMPs) to shape the resident microbiome (43). Neurons express a rich repertoire of peptides, including the previously characterized antimicrobial peptide NDA1 and the dual-function neuropeptides RFamide III, Hym-370, and Hym-357, which previously were found to have strong activity against gram-positive bacteria (43) (Fig. 3 A). In addition, neurons express homologs of Kazal2 and Arminin proteins that have been previously characterized (44–46) as antimicrobial peptides in epithelial or gland cells in Hydra (Fig. 3 A). Notably, the N2 pacemaker population also expresses multiple AMP molecules (Fig. 3 A), indicating that these neurons, in addition to governing...
of spontaneous contractions, since the intervals between contractions become longer and less regular. Sampling size:

Experimental design:

Fig. 2. Identification of Hydra pacemaker cells using orthologs of human ion channels. (A) Genome-wide association studies on patients with gut motility disorders, such as IBS identified ion channels SCN5, ANO1, and TRPM8 that are expressed in human pacemaker cells (CCCs) and found to be essential for gut motility control. A BLAST search was used to identify the homologous genes in Hydra. (B–D) Pacemaker-specific ion channels are highly conserved in Hydra. Phylogenetic tree and domain structure of human SCN (B). ANO1 (C), and TRPM (D) channels and the orthologs from Hydra (Hv). Additionally, sequences from other cnidarians, Nematostella vectensis (Nv) and Clytia hemisphaerica (Ch) are included into the phylogenetic analysis. Noncollapsed trees are presented in SI Appendix, Figs. S8–S10. The topology and domain structure of three Hydra SCN-like sodium channels, six ANO1-like chloride channels and four homologs of TRPM-like cation channels are remarkably similar to their human counterparts. (E) Expression levels of genes encoding SCN-, ANO1-, and TRPM-like channels in Hydra single-cell dataset is overall very weak and restricted to only few cells. However, several transcripts coding for SCN and ANO1 homologs are significantly up-regulated in neurons (red) with some of the transcripts specifically enriched in the neuronal subpopulation N2 (red superscript). (F) Expression of genes encoding SCN-, ANO1-, and TRPM-like channels in the neurons of the subpopulation N2 at the base of tentacles. No signal can be detected in the peduncle region of Hydra (K). (G) Expression fold change expression of genes encoding SCN-, ANO1-, and TRPM-like channels in the neurons of the subpopulation N2 at the base of tentacles. No signal can be detected in the peduncle region of Hydra (K).
peristaltic motion, are also directly interacting with the resident microbiota. To determine if interaction with bacteria is a general characteristic of neurons in *Hydra*, we analyzed the seven neuronal subpopulations for expression of immune-related genes (Fig. 3 B–D). Neurons express virtually all components of the Toll-like receptor (TLR/MyD88) pathway (47) (Fig. 3B and SI Appendix, Fig. S14A; see also SI Appendix, Supplementary Results), as well as many C-type lectin (CTL) receptors (Fig. 3C and SI Appendix, Fig. S14C) and intracellular NACHT and NB-ARC domain-containing NOD-like receptors (48) (Fig. 3B). Neurons express virtually all components of the Toll-like receptor (TLR/MyD88) pathway (47) (Fig. 3B and SI Appendix, Fig. S14A; see also SI Appendix, Supplementary Results), as well as many C-type lectin (CTL) receptors (Fig. 3B and SI Appendix, Fig. S14C). Overall, these observations indicate that neurons in *Hydra* are immunocompetent cells, equipped with receptors, signal transducers, and effector molecules to interact with bacteria.

In addition to a conserved toolbox of immune genes (Fig. 3 B–D), Hydra neurons also employ some of their nonconserved, taxonomically restricted genes (TRGs) to interact with bacteria. Nonconserved genes comprise over 70% of the cell-type-specific genes in transcriptomes of the seven neuronal subpopulations (N1–N7) (Fig. 3E, SI Appendix, Fig. S15, and Dataset S2). The majority of the neuron-specific TRGs code for short peptides (<200 aa) with an N-terminal signal peptide sequence, but with no detectable structural domains (Dataset S2). A machine learning-based approach (49) identified that some of these novel genes code for putative peptides with high membrane-stabilizing activity, indicating strong antimicrobial activity (SI Appendix, Fig. S16 and Dataset S3). Surprisingly, the neuronal cluster N2 that contains the pacemakers was one of the populations most enriched in secreted peptides with putative antimicrobial activity (SI Appendix, Fig. S16).

To provide direct evidence for functional relevance of TRGs in immune reactions in neurons, we characterized TRG cluster62692 that is specifically expressed in neuronal subpopulation N7 in the tentacles (Fig. 3 F–H) and encodes a 185-aa-long peptide (Fig. 3F). The N-terminal signaling peptide is followed by a stretch of mostly positively charged residues, which is predicted to have a high membrane-stabilizing activity (Fig. 3F). Screening a *Hydra* peptide database (50) uncovered a 17-aa amidated peptide, referred to as Hym-121, that is identical to amino acids 47 to 63 within the cluster62692 polypeptide (Fig. 3F), thus providing an evidence for translation and proteolytic processing of the preprohormone encoded by TRG cluster62692. To test whether this peptide may serve as an antimicrobial peptide, we synthesized the amidated amino acids 47 to 63 peptide and subjected it to a minimum inhibitory concentration (MIC) assay against diverse bacteria. We have generated a comprehensive molecular profile of the nematode *Caenorhabditis elegans* (53). We surveyed the expression of the *C. elegans* homologs of the genes that comprise the pacemaker signature in both *Hydra* and mouse (coding for SCN, ANO, and TRPM ion channels, nAChR receptors, and gap-junction proteins) in 27 cell types identified by Cao et al. (53) (SI Appendix, Fig. S20). Although this did not lead to an immediate identification of a certain neuronal cell type as the pacemaker, three *C. elegans* cell types drew our attention: The pharyngeal neurons, pharyngeal muscle cells, and interneurons. These three cell types express to some extent all of the pacemaker-specific genes, suggesting that they might function as pacemakers. This is consistent with the observations that the pharyngeal system represents a neuro-muscular complex that, as a whole, possesses a pacemaker activity and drives regular contractions of the pharynx that assist food ingestion and movement along the intestine. A body of functional data provide evidence that both the neurons and epithelio-muscular cells, as well as a number of pacemaker-relevant genes (egl-19, glf-1 and -2, lev-8, eat-2, and ins-6 and -16) (SI Appendix, Fig. S20) are functionally indispensable for the pacemaker function of the pharynx in *C. elegans* (54–58). Taken together, our findings point to the presence of a specific expression program in the pacemakers of the roundworm, and thus support the view that the molecular architecture of pacemaker cells (Fig. 4 C–F) is highly conserved in evolution.

**Discussion**

We have generated a comprehensive molecular profile of the neural subpopulations in *Hydra* and proposed a conceptual framework for phenotypic diversification of one of the simplest nervous systems in the animal kingdom. This framework builds upon seven spatially and functionally segregated neural subpopulations,
Fig. 3. Neurons in Hydra are immunocompetent cells. (A) Neurons express a rich set of peptides that have been previously characterized as antimicrobial peptides or their homologs. *ref. 43; **ref. 44; ***ref. 46. (B) Heatmap illustrates expression of transcripts coding for components of the TLR/MyD88-dependent immune pathway. Most components are present in the neurons, and five of them are significantly (adjusted \( P < 0.05 \)) enriched. (C) Heatmap illustrates expression of some transcripts coding for NACHT and NB-ARC domain-containing NOD-like receptors that have immune function. (D) Multiple C-type lectin receptors that might recognize bacterial products are strongly expressed in the neurons. (E) Over 70% of top 100 transcripts specifically expressed in each of seven neuronal subpopulations (N1 to N7) is represented by genes that have no homologs outside of Cnidaria, and thus are considered as TRGs. In contrast, among the top 100 transcripts specifically enriched in the interstitial stem cells, only 15% are identified as TRGs. (F) Transcripts of TRG cluster62692 are strongly over-regulated in the neuronal subpopulation N7, weakly expressed in other neurons, and absent from nonneuronal cells of the interstitial lineage. (G and H) In situ hybridization provides evidence that the TRG cluster62692 is expressed exclusively in the sensory neurons of the tentacles. (I) Moving-window small-peptide scan prediction map for the peptide encoded by TRG cluster62692 with residue charge and secondary structure annotations. The heat map reflects the peptide's probability (\( \sigma \)-score) of being membrane active as predicted by the machine learning classifier (49). High \( \sigma \)-scores (yellow) suggest that cluster62692 peptide is a potent antimicrobial peptide. N-terminal signal peptide, putative proteolysis sites, and a sequence identical to a previously described peptide Hym-121 (50) are found within the cluster62692 peptide, providing evidence that a preprohormone cluster62692 is processed and gives rise to a secreted active peptide. The 17-aa-long peptide corresponding to amino acids 47 to 63 (SPPWNKFGAFVKSKLAK = Hym-121) with high membrane activity score (\( \sigma = 2.317 \)) and control peptide amino acids 59 to 75 (SKLAKSKREMSNSDGSE) with no membrane activity (\( \sigma = −1.878 \)) were synthesized, C terminally amidated, and tested for antimicrobial activity in a MIC assay. (J) The peptide 47–63 Am is a potent antimicrobial peptide that shows selective growth inhibiting activity against gram-positive and -negative bacteria. Control peptide 59–75 Am demonstrates no antimicrobial activity. Consistently with previous observations (43), dual-function neuropeptides Hym-370 and Hym-357 show some antibacterial activity, yet weaker and more restricted than the peptide 47–63 Am. (K) Representative wells from plates of MIC assay. At concentration 25 \( \mu \)M, the peptide 47–63 Am inhibits growth of Curvibacter sp. and Acidovorax sp. and affects colony morphology of B. megaterium. The growth in the presence of control peptide (59–75 Am, 25 \( \mu \)M) is not different from that in the pure medium (control).
allowing for the emergence of diverse and complex behaviors within a morphologically simple nerve net structure.

Our data establish the neurons of the N2 population as a major contributor to controlling the rhythmicity of spontaneous body contractions in Hydra. These prototypical pacemaker cells reside in the head region and specifically express ANO1-, SCN5-, and TRPM-like ion channels that characterize human gut pacemakers. Consistently, the experimental inhibition of these channels greatly disturbs peristalsis in Hydra, which is consistent with our finding that these neurons are coupled by gap junctions into a network exhibiting pacemaker activity. The high degree of gene-expression program conservation between the Hydra pacemaker subpopulation N2, the pharyngeal pacemaker complex of C. elegans, and murine pacemaker cells (Fig. 4 C–F) supports that peristaltic motor activity of the gut is an evolutionary ancient archetypical property necessary to sustain life (59, 60), and that the cells with recurrent spontaneous electric activity have evidently emerged as early as the nervous system itself.

Finally, we discovered that Hydra pacemaker cells express a rich set of immune genes, including antimicrobial peptides providing a mechanism for direct interference with resident microbes. The finding that many neuron-specific novel genes also encode anti-microbial peptides underlines the important function of neurons in interacting with microbes. Experimental interfering with microbiome in Hydra has a profound effect on the gene-expression program of the pacemaker neurons (Fig. 4B) and disrupts the rhythmic peristaltic activity of the polyps (26). Similarly, disturbances of gut microbiota in humans (61–63) result in changes in pacemaker rhythmicity and abnormal peristalsis. Because the Hydra pacemaker neurons can directly mediate the interaction with the microbiome, it might be plausible that human pacemakers in the gut similarly communicate with microbial communities. In fact, emerging data on mice provide first evidence for direct interactions between the gut microbiota, enteric neurons, and the intestinal motility (64–66). The evolutionary similarity or dissimilarity of the molecular toolkit used for such communication should become a subject of further investigations.

Altogether, our findings will improve the understanding of the archetypical properties of net nerve systems with pacemakers including the human enteric nervous system, which is perturbed in human dysmotility-related conditions affecting a large portion of the general population worldwide. We therefore presume that the principles identified here are relevant far beyond Hydra.

Materials and Methods
Experimental Design. Experiments were carried out using Hydra vulgaris strain AEP. Animals were maintained under constant conditions, including the culture medium, food, and temperature (18 °C) according to standard procedures (67). Experimental animals were chosen randomly from clonally growing asexual Hydra cultures. The animals were typically fed three times a week; however, they were not fed for 24 h prior to pharmacological interference experiments, or for 48 h prior to RNA isolation, immunohistochemical staining, and in situ hybridization.

Generation of Transgenic Hydra Strains. To facilitate the FACS-mediated molecular profiling of single cells, we developed a transgenic Hydra line expressing two reporter proteins. The enhanced GFP (eGFP) was cloned downstream from the previously reported trans promoter (68) and flanked by the actin terminator, and the codon-optimized DsRED2 protein was driven by the actin promoter sequence and flanked by the actin terminator region. After cloning into the LiAg vector (69), the transgenic construct was propagated in E. coli DH5α strain and microinjected into zygotes of H. vulgaris strain AEP, as previously described (67, 69). Founder mosaic transgenic animals were clonally propagated, screened, and enriched for transgenic cells until all interstitial stem cells were transgenic. The transgenic animals show no developmental abnormalities and are maintained in the laboratory for over 5 y.

FACS Isolation of Cells. To isolate cells of the interstitial stem cell lineage from the transgenic Hydra by FACS (SI Appendix, Fig. S1A), the polyps were dissociated into a cell suspension, as previously described (68). The single cells were sorted according to FSC, SSC, eGFP, and RFP fluorescence using FacsAria III cell sorting system (BD Biosciences), directly into 382-well plates containing lysis buffer, RNase inhibitor, oligo-dT20VN primer, and dNTP mix, and snap-frozen. Sorting procedures are described in detail in SI Appendix, Supplementary Methods). In total, 1,152 individual cells were harvested: 384 GFP/RFP neurons, 384 GFP/RFP stem cells, and 384 GFPlow/RFPlow cells.

Smart-Seq2 Library Preparation and Sequencing. To generate cDNA libraries from the isolated cells, a previously described Smart-seq2 protocol (70) was implemented with minor modifications (SI Appendix, Supplementary Methods). The libraries were pooled and paired-end sequenced on Illumina HiSeq2500 instrument. Raw sequences and quality scores for all clusters were extracted using CASAVA software.

scRNA-Seq Data Processing. Quality Control, and Hierarchical Clustering. Raw data from scRNA-seq were processed using a snakemake scRNA-seq pipeline. Cells passing quality control (i.e., 1,016) were analyzed using the R package Seurat2_3.4. To map the reads, we used the previously described (71) reference transcriptome of H. vulgaris strain AEP (accession no. SRP133389). We included genes detected in at least three cells and cells that contained at least 5,000 transcripts. This resulted in a total of 928 cells and 166,186 transcripts. In the initial phase of the project, we generated in total 14 alternative clustering maps that segregated 928 cells into 9 to 22 clusters (SI Appendix, Fig. S21). Since the partitioning into 12 clusters, including 7 populations of neurons, was most consistent with our previous observations and the literature data, and resulted in clusters composed of at least 40 cells each, we further used only this clustering scheme. Specific parameters for the different input gene sets and input genes for each gene set are presented in SI Appendix, Supplementary Methods and Datasets 51, 57, and 58.

In Situ Hybridization. To map the seven neuronal clusters populations in the Hydra body, we performed in situ hybridization with a set of genes strongly enriched in either of the seven neuronal subpopulations. Expression patterns were detected in the whole-mount Hydra preparations by in situ hybridization with antisense digoxigenin-labeled RNA probes, as previously described (72). A DIG-labeled sense-probe was used as a control. Signal was developed using anti-DIG antibodies conjugated to alkaline phosphatase (1:2,000, Roche) and NBT/BCIP staining solution (Roche). Images of the in situ preparations were captured on a Zeiss Axioscope with Axiocam camera.

Pharmacological Interference Assays. To investigate the role of specific ANO1-, SCN-, and TRPM-like channels in the pacemaker activity in Hydra, we exposed normal H. vulgaris AEP polyps to different pharmacological agents, recorded, and quantified their behavior. Polyps were treated with 25 μM An9 (Sigma, Cat. No. SML1813), 200 μM menthol (Sigma, Cat. No. 15785), 100 μM lidocaine (Sigma, Cat. No. LS647), 1 mM tubocurarine (DTC, Sigma, Cat. No. 93750), or 100 μM muscimol (Sigma, Cat. No. M1523) for 1 or 12 h at 18 °C. Control polyps were incubated either in Hydra-medium or in the medium supplemented with 0.16% DMSO (for An9, which has been dissolved in 100% DMSO to stock concentration 15 mM). The spontaneous contractions were video-recorded and quantified, as previously described (26). We recorded the behavior for 90 min with a frequency 20 frames per minute. For further analysis, we excluded first 30 min of the recorded sequence, and quantified number of full-body contractions and their periodicity using a custom ImageJ plugin (26). The contraction frequencies were normalized to the average frequency of contractions in corresponding control polyps. To examine the effects of the modulators on the feeding reflex, Hydra polyps were pretreated with the pharmacological agents, their feeding reflex was elicited by 10 μM reduced glutathione (GSH, Sigma, Cat. No. G4251), and the duration of feeding response was recorded as described by Lenhoff (73).

Statistical Analysis. The sample size (n) reported in the figure legends is the total amount of animals used in each treatment. Each animal employed was assigned to only one treatment and was recorded only once. Treatment of the polyps with pharmacological substances and evaluation of the behavioral parameters (contraction frequency, intervals between contractions, and duration of feeding response) was blinded. Differences in contraction frequency, interval between contractions, and feeding response duration between the treatments (i.e., An9, lidocaine, menthol, DTC, and muscimol) and corresponding control (i.e., Hydra-medium or DMSO) were analyzed using unpaired t test.
Prototypical pacemaker neurons interact with microbiota. (A) To test the immune function of the pacemaker neurons in Hydra, GF animals were generated by antibiotic treatment and then recolonized with the natural microbiota to obtain conventionalized (Conv.) polyps. Polyps treated with DMSO solvent were used as control. Total RNA was extracted from either amino acids 47 to 63 or amino acids 59 to 75 (Sklakskremrsmrsgse) of the prepropeptides encoded by the clusters2692 were synthesized (up to 5 mg). C-terminally amided, and purified to a purity of >95% (GenScript), and their antimicrobial activity was estimated in an MIC assay, as previously described (43). The following bacterial strains were used in MIC assays: B. megaterium ATCC14581, E. coli D31, and four isolates from the natural H. vulgaris strain AEP microbiota: Curvibacter sp., Duganella sp., Acidovorax sp., and Undibacterium sp. (46). MIC determination susceptibility assays were carried out in 96-well microtiter plates that were precoated with sterile 0.1% bovine serum albumin (BSA). After removal of BSA the wells were filled with a twofold dilution series of either amino acids 47 to 63 or amino acids 59 to 75 peptide. We also tested the neuropeptides Hym-370 (KPNAyKgklPiglw-amide) and Hym-357 (KpaFlfgkyp-amide) that has been previously identified as putative antimicrobial substances (43). Lyophilized peptides were dissolved in ultrapure water to stock concentration of 10 mg/mL. Incubation with an inoculum of ~100 CFU per well was performed in phosphate buffered saline (PBS) buffer (pH 6.2) overnight at 37 °C for B. megaterium and E. coli, or in R2A media for 3 to 4 d at 18 °C for four isolates of Hydra bacteria. The MIC was determined as the lowest serial dilution showing absence of a bacterial cell pellet. Experiments were carried out in triplicates.

Phylogenetic Analysis of Ion Channel Genes. To uncover whether Hydra has homologs of the ion channel genes whose expression is known to be either restricted to mammalian ICCs or essential for gut motility, we performed a BLAST search (tblastn) using full-length amino acid sequences of ANO1 (UniProt accession no. Q5XXA6), SCNSA (UniProt accession no. Q14524), and TRPM8 (UniProt accession no. Q722W7) proteins from Homo sapiens against the genome of H. vulgaris (38) (available at https://research.nhgri.nih.gov/hydra/) and the reference transcriptome of H. vulgaris strain AEP. Matches with expectation e-value <1e-10 were considered as signs of homolog presence and were verified by manual domain composition analysis using SMART (74), transmembrane domain prediction with TMHMM (75), and reciprocal BLAST against the UniProt database. Maximum-likelihood phylogenetic trees of ANO1, SCNSA, and TRPM8 homologs from Hydra, human, African clawed frog Xenopus laevis, and zebrafish Danio rerio were built using full-length amino acid sequences aligned using MUSCLE (76) with 1,000 bootstrap iterations.

Generation of GF and Conventionalized Polyps. GF polyps were generated by treating control Hydra polyps for 2 wk with an antibiotic mixture containing rifampicin, ampicillin, streptomycin, and neomycin in final concentrations of 50 μg/mL each and spectinomycin at 60 μg/mL, as previously described (26). Since rifampicin stock is dissolved in DMSO, control polyps were incubated in the corresponding 0.1% DMSO concentration for the same period of time. Antibiotic solution and control medium were replaced every 48 h. After 2 wk of treatment, the animals were transferred into sterile Hydra-medium that was further replaced every 48 h before isolation of polyps. Conventionalized Hydras were generated by incubating GF polyps with tissue homogenates of control animals. Previously, we demonstrated that recolonization of GF polyps in this way results in the establishment of a bacterial community similar to that of an intact control Hydra polyp (26). Generation of GF polyps and recolonization were repeated in triplicates. The GF status of the polyps and success of recolonization were tested by plating a macerated polyp on R2A agar, which supports growth of H. vulgaris strain AEP. Matches with expectation e-value <1e-10 were considered as signs of homolog presence and were verified by manual domain composition analysis using SMART (74), transmembrane domain prediction with TMHMM (75), and reciprocal BLAST against the UniProt database. Maximum-likelihood phylogenetic trees of ANO1, SCNSA, and TRPM8 homologs from Hydra, human, African clawed frog Xenopus laevis, and zebrafish Danio rerio were built using full-length amino acid sequences aligned using MUSCLE (76) with 1,000 bootstrap iterations.

Quantitative Real-Time PCR Gene-Expression Analysis. To test whether the genes coding for ANO1, SCNSA, and TRPM8 homologs in Hydra are differentially expressed in the polyp along the oral-aboral axis, we performed quantitative real-time PCR. We dissected polyps into three body sections: Head (hypostome area with tentacles), body column, and foot (peduncle). Each total RNA was extracted from body fragments obtained from 50

Bacteria-derived products may have profound effects onto the gene-expression program of the pacemakers via immune pathways or might directly target the pacemaker ion channels or neuromediator receptors. (C)
polyps, and converted into the cDNA as previously described (72). For each body condition, we made three to five biological replicates.

To examine whether the gene-expression profile in the pacemaker cells of Hydra is dependent on the presence of specific microbe, we compared expression of genes coding for ANO1, SCNSA, and TRMP8 homologs, as well as nACHR, innexin gap-junction proteins, and three pacemaker-specific transcription factors using real-time PCR. Total RNA was extracted from 100 normal, GF-, or conventionalized polyps and converted into cDNA. For each condition, we made three biological replicates. Real-time PCR was performed using GoTaq qPCR Master Mix (Promega) and oligonucleotide primers specifically designed to amplify the homologs of ANO1, SCNSA, and TRMP8 ion channel genes, nACHR, innexins, and transcription factors, as well as the eIFa (translation elongation factor 1α) and actin genes as equilibration references (3f Appendix, Table S1). The data were collected by ABI 7300 Real-Time PCR System (Applied Biosystems) and analyzed by the conventional ΔΔCt method.

Generation of Antibodies and Immunohistochemistry. To localize the expression of ANO1-like and SCNSA-like ion channels in Hydra using immunocytochemistry, polyclonal antibodies were raised against synthetic peptides in rabbits. The peptides that correspond to intracellular loops located between transmembrane domains of the ion channels (hySCNS-5: SRSPKPMKFDYKYPE; hyANo1 Delta ETRR(DRAQQ)) were synthesized, purified, and N terminally conjugated with KLH prior to injection (GenScript). Polyclonal antibodies were affinity-purified on the antigen and concentrated to 1.5 mg/mL. Serum har

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