Wounding response in Porifera (sponges) activates ancestral signaling cascades involved in animal healing, regeneration, and cancer

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Upon injury, the homeostatic balance that ensures tissue function is disrupted. Wound-induced signaling triggers the recovery of tissue integrity and offers a context to understand the molecular mechanisms for restoring tissue homeostasis upon disturbances. Marine sessile animals are particularly vulnerable to chronic wounds caused by grazers that can compromise prey’s health. Yet, in comparison to other stressors like warming or acidification, we know little on how marine animals respond to grazing. Marine sponges (Phylum Porifera) are among the earliest-diverging animals and play key roles in the ecosystem; but they remain largely understudied. Here, we investigated the transcriptomic responses to injury caused by a specialist spongivorous opisthobranch (i.e., grazing treatment) or by clipping with a scalpel (i.e., mechanical damage treatment), in comparison to control sponges. We collected samples 3 h, 1 d, and 6 d post-treatment for differential gene expression analysis on RNA-seq data. Both grazing and mechanical damage activated a similar transcriptomic response, including a clotting-like cascade (e.g., with genes annotated as transglutaminases, metalloproteases, and integrins), calcium signaling, and Wnt and mitogen-activated protein kinase signaling pathways. Wound-induced gene expression signature in sponges resembles the initial steps of whole-body regeneration in other animals. Also, the set of genes responding to wounding in sponges included putative orthologs of cancer-related human genes. Further insights can be gained from taking sponge wound healing as an experimental system to understand how ancient genes and regulatory networks determine healthy animal tissues.

Tissue homeostasis is the capacity to maintain, via feedback loops, the internal conditions that allow the proper functioning of an animal. If the mechanisms of homeostasis fail, diseases like chronic wounds or cancer will develop1–5. In fact, there are intriguing parallels between wound healing and cancer3–6. Wound healing activates mechanisms that are strikingly similar to those that, when dysregulated, trigger tumor initiation and progression3–6. Processes like inflammation, activation of migration, or enhanced proliferation, regulated in wound healing and uncontrolled in cancer6, are largely regulated by ancient genes that evolved at the emergence of multicellularity7. Thus, an evolutionary perspective into the molecular mechanisms of homeostasis can reveal conserved gene interactions and functions in coordinating cellular behavior and sensing the environment. Wounding response and healing research offers a context to understand fundamental processes that control how healthy tissues are maintained and restored upon disturbances.

We can learn about the molecular basis of animal tissue homeostasis, and its implications for health and disease, by investigating wounding responses and regeneration throughout the animal kingdom8–11. Some animals that can reconstruct extensive body parts, like anthozoan cnidaria (Hydra sp., Nematostella vectensis), planaria (Schmidtea mediterranea, Dugesia japonica) or zebrafish (Danio rerio), have been the predominant study systems.

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for understanding the mechanisms of tissue repair, and recent studies suggest that they may share common molecular features at the early points of regeneration, at the time of the recognition and signaling of the wound. However, beyond these classical model systems, we know very little about the molecular mechanisms involved in animal response to wounding.

Marine sessile animals like sponges or corals are particularly susceptible to chronic wounds caused by grazers. These wounds can yield disease and/or magnify the detrimental effects of other stressors. As a result, our understanding on how these sessile animals respond to grazing at the molecular level is scarce. Here we focused our attention on marine sponges (Phylum Porifera) because they play a key role in the ecosystem, they belong to the group of animals with high regenerative capacity, and constitute one of the earliest-diverging animal lineages; but, compared to other animal groups, they have been little studied. Sponge wounding response and healing research mainly focused on the cellular processes involved in reparation of small incisions. This response likely varies among sponge species, but typically lasts several days and engages different cell populations migrating to the wounded area and undergoing differentiation. At molecular level, studies focused on specific genes or lacked enough replicates for differential gene expression analysis at transcriptome level. Thus, molecular processes regulating wounding responses in sponges remain to be characterized.

Here, we investigated the injury-induced transcriptomic response of the sponge Aplysina aerophoba (Nardo, 1833). We applied two different wounding treatments: grazing caused by its primary predator, the opisthobranch Tylodina perversa (Gmelin, 1791), (i.e., grazing treatment), and repeated clipping of the sponge surface with a scalpel (i.e., mechanical damage treatment). The grazing treatment offers an ecological-relevant context of wound repair, while mechanical damage allows us to control wound size and periodicity. In our previous study, we ran three experiments to collect samples at 3 h, 1 d, and 6 d post-treatment, respectively (Fig. 1). We found a

Figure 1. Experimental design. (a) Sponge individuals were divided into three specimens and let acclimate for 1 week. The specimens of each sponge individual were randomly assigned to either control (C), grazing (G), or mechanical damage (M) treatment. Each experiment corresponds to a different time point after stop of treatment: 3 h (3 h); 1 day (1 d) and 6 days (6 d). Number of replicates are shown. One grazed sample collected at 3 h presented low-quality reads and was not included in the differential gene expression analysis. Created with Biorender.com. (b) Example of grazed (left) and mechanically-damaged (right) sponges.
great resemblance in the chemical and cellular response of _A. aerophoba_ to wounding caused by grazing and by mechanical damage, which we defined as a “jack-of-all-trades” defense strategy. Now we performed differential gene expression analysis on RNA-seq data on those two research questions: (i) does mechanical damage cause a similar transcriptomic response as grazing damage, confirming the unspecific response to the specialist sponginous and (ii) does the wound-induced transcriptomic response in the sponge _Tylodina perversa_ have conserved molecular features of animal tissue homeostasis?

**Materials and methods**

**Specimen collection and experimental set-up.** The samples for this study come from the experiments described in Wu et al. In short, the sponge _Aplysina aerophoba_ and the opisthobranch _Tylodina perversa_ were collected by Scuba diving in the Mediterranean coast of Spain (42.29408 N, 3.28944 E in 2016 and 42.1145863 N, 3.168486 E in 2017), at a depth between 2 and 10 m. Animals were transported to the Experimental Aquaria Zone at the Institute of Marine Science (ICM-CSIC) in Barcelona (Spain). There, each sponge individual was divided into 2–3 specimens with its own osculum, by applying a clean cut. Each specimen was placed into individual aquaria (6 L) maintained in a flow-through system with direct intake of seawater. After 1 week acclimation, the specimens from each sponge individual were randomly assigned to one of the following three treatments (Fig. 1), (i) control: no treatment, (ii) grazing: one sea slug, which starved for 24 h, was placed in direct contact to the sponge specimen and allowed to feed freely for 24 h, and (iii) mechanical damage: the sponge specimen was repeatedly clipped with a scalpel for 3 min every half hour at the first 3 h and the last 3 h (in the process, the ectoderm at the wound was removed). All treatments were stopped after 24 h. This design was applied in independent experiments that differ in the sampling points after stop of treatments (and use different sponge individuals). In the first experiment, samples were collected 3 h after stop of the treatment; in the second experiment after 1 d; and in the third experiment, after 6 d. The 6 d experiment was performed in 2016, the 3 h and 1 d experiments were performed in 2017.

**RNA extraction and sequencing.** A total of 40 samples from 3 h, 1 d, and 6 d experiments were collected for RNA-seq (Fig. 1). Samples from grazing and mechanical damage treatments were collected right at the wound, washed with artificial seawater, and fixed in RNAlater® (Invitrogen, Thermo Fisher Scientific, Germany) at 4 °C overnight. Samples were then stored at – 80 °C until RNA extraction. Total RNA was extracted and processed as in43. In short, Total RNA was obtained by using the AllPrep DNA/RNA Mini Kit (Qiagen, Germany) and treated with DNA-free DNase Treatment and Removal Reagent (Ambion, Germany). DNA contamination was further checked by PCR amplification of eukaryotic 18S rRNA gene (Sigma-Aldrich, primers by Stewart et al.44). Total RNA was checked for purity, quantity, and quality by NanoDrop 2000c Spectrophotometer (peola, Germany), Qubit 2.0 (Life Technologies, Carlsband, CA), and Experion™ Electrophoresis Station (Bio-Rad, Hercules, CA). Only the total RNA which was DNA-free and presented OD260/280 > 1.8 (Nanodrop) as well as RIN > 8 (Experion) was subsequently used for RNA-seq. Equal amounts of the total RNA (500 ng) were used for library construction with the TruSeq stranded mRNA library prep kit (Illumina, Inc., USA), including a poly-A enrichment step. Samples were sequenced (150 base paired-end) on the HiSeq 4000 system (Illumina, Inc., USA). Library preparation and sequencing was performed at the IKMB in Kiel, Germany.

**Differential gene expression analysis.** Differential gene expression analysis was performed following the protocol described by Pita et al.45. Raw Illumina RNA-seq reads were qualitatively trimmed and filtered to remove adapters and low-qualitative reads in Trimmomatic-version 0.3846, parameters: TruSeq3-PE-2.fa:2:20:10 LEADING:3 TRAILING:3 SLIDINGWINDOW:4:15 MINLEN:36. Prokaryotic and microbial eukaryotic reads were filtered in the classifier Kaiju-version 1.6.247, in greedy-5 mode (database accessed in February 2018). Because of the lack of a reference genome for _A. aerophoba_, the reads from all samples were then combined to create a de novo reference transcriptomic assembly in Trinity-version 2.6.648. Assembly statistics were obtained in Trinity and TransRate-version 1.0.3.49. Completeness of the de novo reference transcriptome was assessed by comparing the assembly against the Metazoa reference data in BUSCO-version 3.0.2 pipeline50: first, open reading frames (ORFs) were identified in the transcript sequence (Transdecoder) and those ORFs codifying for proteins > 100 amino acids were further characterized in homology searches to publicly available data (BLAST+/SwissProt), protein domain identification (HMMER/Pfam), protein signal peptide and transmembrane domain prediction (signalP/tmHMM), as well as eggNOG, Gene Ontology (GO), and KEGG annotation51. Those contigs with blast or blastp matches to Bacteria, Archaea, or Virus were further removed from the reference assembly.

Gene quantification was estimated based on Trinity component abundances by RSEM bowtie2-based quantification (version 1.2.19). Differential gene expression analysis within each experiment (i.e., 3 h, 1 d, and 6 d) was performed in edgeR as implemented in Trinity-version 2.6.5 (default parameters). Differentially expressed genes (DEGs) in grazing versus control and mechanical damage vs control were defined by False Discovery Rate –corrected (FDR) _p_ value < 0.005 and log2[fold change] ≥ 2 expression (fourfold change).

Based on the set of DEG for each experiment, a Gene Ontology (GO) enrichment analysis was performed in blast2GO-v552. GO-annotated genes in the reference transcriptome were used as reference set. Enrichment was determined by Fisher’s exact test (FDR _p_ value < 0.005). Enriched GO terms were then analyzed in REVIGO53 for reducing redundancy and grouping them based on semantic similarity (semantic similarity measure: SimRel). The results from differential gene expression analysis were visualized with the package ggplot254 in R-version 3.5.1 (R Core Team, 2019) as implemented in R-Studio55. Final layout of the figures was done in Inkscape http://inkscape.org.
Figure 2. Activation of gene expression upon wounding. (a) Number of up- and down-regulated DEGs in grazing (G) and mechanical damage (M) groups compared to control group at 3 h, 1 d, and 6 d post-treatment experiments. Note that none DEG was detected at 6 d. “Up-regulated” refers to those genes with higher expression levels in the corresponding treatment than in control; “down-regulated” refers to those genes with lower expression levels in treatment than in control. DEGs were defined according to FDR p value < 0.005 and log2 |FC|≥ 2 expression, as calculated in edgeR. (b) Percentage of DEGs with annotation per time point and comparison. Annotation was performed following Trinotate pipeline. (c) GO categories enriched in response to each treatment at each time point. The enrichment of a GO term is indicated by a circle. Circle color corresponds to the semantic cluster as defined in REVIGO with circle size reflecting Fisher test’s absolute log2(FDR-corrected p value); that is, a value of 10 means a FDR-corrected p value of 1e−10. In this sense, this bubble chart is equivalent to the tree maps generated by REVIGO. Separate panel refers additional GO terms enriched in mechanical damage after 3 h. Significance threshold for enrichment test: FDR p value < 0.005. REVIGO dispensability threshold < 0.5.

Protein interaction networks in STRING. The translated coding regions of 3 h post-mechanical damage DEGs (Transdecoder; > 100 amino acids) were compared to the proteome of A. queenslandica (UniProt UP000007879_444682) by Blastp (e-value < 1e−5). The best blastp match in STRING v1159 for protein–protein network analysis (medium required interaction score = 0.400). In addition, we highlighted those reciprocal Blastp best hit (rBBH; e-value < 1e−5), identified as in56. rBBH analysis (protocol as in Pritchard et al., 2018) of 3 h post-mechanical damage DEGs was also performed against the human reference proteome (UP000005640_9606). Those human proteins with reciprocal Blastp best hit (e-value < 1e−5) were then used as input for protein–protein network analysis in STRING v1159. The genes coding for those proteins were searched in COSMICS v91, https://cancer.sanger.ac.uk57 and in the set of key cancer regulators identified by Trigos et al.58. Final layout of the figures was done in Inkscape http://inkscape.org.

Results

Overview RNA-seq analysis. We used RNA-seq to characterize the molecular response of the sponge A. aerophoba to wounding by grazing and by mechanical damage compared to control sponges on the samples derived from40. Each of the three experiments corresponds to a different sampling time point (Fig. 1): 3 h, when we expected to detect most of the signaling response based on other transcriptomic studies41; 1 d, concomitant to our observation of cell accumulation at the wound40; and 6 d, when signs of tissue recovery were obvious in all wounded sponges, although scars were still evident. Some sponges showed signs of recovery 3 d post-treatment (Supplementary Fig. S1 online)40.

We sequenced a total of 40 samples of A. aerophoba, corresponding to between 3 and 6 biological replicates per treatment (Fig. 1). The number of paired-end Illumina reads generated in this study is summarized in Supplementary Table S1 online. The de novo reference transcriptomic assembly comprised 459,466 genes (Trinity components45), representing 92.2% of the 902 core Metazoan genes, and 60.6% of these were complete (BUSCO46). The statistics of the resulting reference assembly are summarized in Supplementary Table S2 online. Overall, 72.32 ± 3.69% (average ± standard error) of the reads of each sample aligned to the de novo-assembled reference transcriptome. One grazed sample collected after 3 h (H15407-L1_S56) had a lower alignment rate of 53.53% as well as a lower number of read pairs than the other samples. Therefore, we discarded this sample for downstream analysis. The results from the differential expression analysis in edgeR as well as the annotation of the DEGs are reported in Supplementary Data S1 online (3 h experiment) and Supplementary Data S2 online (1 d experiment).

The response of the sponge upon grazing was similar to the response upon mechanical damage. Each wounding treatment was compared to control sponges, within each time point. We found strong activation of gene expression, both in the 3 h as well as the 1 d post-treatment experiment (Fig. 2). However, 6 d post-treatment, gene expression levels in grazing and mechanical damage samples resembled those in control (all FDR p-values > 0.1) (Fig. 2). At 3 h post-treatment, mechanical damage seemed to activate a stronger gene regulation than grazing, in terms of number of genes and expression levels (Fig. 2; Supplementary Figs. S2 and S3 online). However, most of the genes detected as unique to mechanical damage also showed elevated expression in grazing (Supplementary Figs. S2 and S3 online). Moreover, the functional categories (GO terms) enriched upon mechanical damage and grazing were similar, although genes related to signaling were more prominent 3 h post-mechanical damage treatment (Fig. 2c). For example, in the mechanical-damage response additional transcription factors were activated, including the transcription factors Runt and Sox as well as a NFκß-like gene (Supplementary Data S1 online).

From now on, we present the transcriptomic responses according to the set of annotated DEGs, which comprised ca. 46% of the total of DEGs (FDR p value < 0.005) (Fig. 2b).

Wound-induced signaling pathways. Grazing and mechanical damage treatments induced a clotting-like cascade49-51. One of the most enriched functions both at 3 h and 1 d post-treatment experiments was the molecular function “protein-glutamine gamma-glutamyltransferase activity”, related to the GO terms “keratinization” and “peptide cross-linking” (Fig. 2c). This function corresponds to the activation of genes annotated as transglutaminases (Fig. 3). The transcriptomic response retrieved both in 3 h and 1 d experiments also included many genes involved in calcium signaling, in extracellular matrix rearrangement, and cell adhesion (e.g., integrins, cadherins, fibrillin), (Supplementary Data S1 online). At 3 h we additionally detected the activation of...
Figure 3. Expression changes of differentially expressed genes involved in clotting-like cascades and signaling during sponge wounding response. Heatmaps showing log₂-transformed median-centered TMM-normalized expression changes of selected DEG (rows) in each sample from 3 h post-treatment experiment. Differential gene expression analyses were performed in edgeR following Trinity pipeline. Genes were annotated in Trinotate. The domain annotation (PFAM) is shown in brackets. The full list of DEGs is found in Supplementary Data S1 online.
multiple metalloproteases, which are usually involved in degradation/deposition dynamics of the extracellular matrix, and multiple protein kinases and transcription factors (mainly Ets (PF00178) and Rfx (PF02257)-containing genes) (Fig. 3). Wounding also induced the regulation of receptors, in particular Scavenger receptors cysteine rich (SRCRs) and G-protein coupled receptors (GPCRs) (Fig. 3). A total of 13 and 8 SRCR genes were identified within the 3 h and 1 d DEG dataset, respectively, based on the presence of one or multiple SRCR domains (PF00530) (Fig. 3). GPCRs were identified and classified according to the 7 transmembrane domain (7tm) and included several members of the Adhesion (7tm_2, PF00002) and Rhodopsin (7tm1_PF00001) GPCR families, as well as one gamma-aminobutyric acid type B receptor (7tm_3, PF00003) (Fig. 3). We have summarized the set of genes that were activated in response to grazing and to mechanical-damage in 3 h experiment in a schematic representation (Fig. 4), which, we propose, summarizes the key components of the sponge response to wounding.

The response to both grazing and mechanical damage detected in 1 d experiment resembled the one reported in 3 h experiment, particularly in terms of transglutaminase-like genes and proteases (GO terms “keratinization” and ‘peptide cross-linking’, Fig. 2c). However, after 1 d, most of the genes related with signaling cascades (e.g., kinases, proteases) are not significantly regulated (Fig. 2c; Supplementary Data S1 online; Supplementary Fig. S4 online).

**STRING protein–protein interaction networks.** We created a putative protein–protein interaction network of the DEGs in 3 h post-mechanical damage treatment in STRING55, based on the blastp protein hits to the proteome of *A. queenslandica*, the sponge species of reference in public genomic databases (Fig. 5A). STRING analysis revealed three major signaling cascades in the response to wounding: MAPK signaling pathway, Wnt signaling pathway, and the clotting-like cascade mediated by thrombospondin and other components of the extracellular matrix (Fig. 5A). A member of the Wnt signaling pathway, DVL (disheveled), could potentially mediate the crosstalk with the MAPK pathways via MAPK3 (Fig. 5A). These pathways are also connected to other receptors via a calcium/calmodulin-dependent protein kinase (CAMK, Fig. 5A). Another prominent hub was a CUB and Sushi domain-containing (CSMD3-like) protein, which appeared connected to transmembrane receptors and multiple proteins of the extracellular matrix (latent TGFβ binding protein, Fig. 5A; Supplementary Data S1 online).
Gene function evidence is scarce for sponges; thus, the network generated in STRING relies on information for similar genes/putative orthologs in other organisms. This means that they likely represent conserved gene–gene and protein–protein interactions in animal evolution. In fact, we identified 53 putative human orthologous of the 3 h post-mechanical-damage DEGs by reciprocal best hits search (rbHH, e-value < 1 e\(^{-5}\)); including several GPCRs, GTPases, kinases, and multiple transcription factors (Fig. 5B). 11 out of these 53 human proteins are codified by genes listed in the Catalogue of somatic mutations in cancer (COSMIC v91) (https://cancer.sanger.ac.uk\(^{-57}\)) (Fig. 5B, in bold). Within these, we found eight genes defined as key regulators of ancient gene networks (Fig. 5A). We directly compared grazed vs mechanical-damage, Supplementary Fig. S1 online) and the periodicity of it (i.e., tightly controlled in the mechanical damage compared to grazing), both treatments induced similar transcriptomic responses in terms of activated functions (Fig. 3, Supplementary Fig. S3 online). This was also evident when we directly compared grazed vs mechanical-treated sponges (Supplementary Table S3; Supplementary Fig. S5 online). Thus, these results suggest that the sponge has not evolved a specific tolerance to its specialist grazer. Information of grazing pressure in the T. perversa\(^{-39}\) interaction is limited to a single location\(^{-39}\). There, > 50% sponges in the shallow waters (first 10 m, the preferred habitat of the sponge) hosted 1, occasionally 2 opisthobranchs\(^{-39}\). It could be that this predation pressure is not lethal per se, but may cause chronic wounds with detrimental effects on sponge health. For example, Caribbean Aplysina fistularis looked diseased as a consequence of sustained grazing by cowries\(^{-39}\). Our study shows that wound cues strongly induced gene expression at the wound site (Figs. 2, 3), which could constitute a chronic response and significant metabolic burnt in the field\(^{-39}\), particularly if grazing pressure is high. Overall, we need more studies to evaluate the impact of grazing on sponges as an additional stressor that can compromise sponge health, particularly in times of anthropogenic-driven environmental changes.

In our experiments, sponges were divided into clones 1 week before the experiments in order to achieve replicates. This manipulation may affect sponges in a way that could mask differences in the transcriptomic pattern of control and treated sponges in the 6 d experiment. In our current setup, we cannot completely rule out this possibility. However, we find more likely that the lack of differentially-expressed genes between treated and control samples 6 days post-treatment is due to a "return-to-baseline" situation, based on two main arguments. First, the division of sponges for making the clones is very different from the wounding treatment. The clones are generated by a clean cut between sponge chimneys at the base of the sponge, whereas mechanical damage consists on repeated clipping and affects the first 3–5 mm from the surface to the inside of the sponge. Second, we are collecting the samples at the wounding site, which does not overlap with the place sponges were cut for generating the clones. Nevertheless, we are currently working in characterizing the response of the sponge in the latter time points (6 days post-treatment and beyond) to get a better understanding of the recovery phase.

We defined sponge molecular response to wounding as a "clotting-like cascade"\(^{-19-41}\), because we found multiple DEGs annotated as transglutaminases both in grazing and mechanical-damage treatments and in the 3 h as well
as the 1 d experiment, resembling the molecular cascades involved in clot formation and stabilization in other animals, including vertebrates and invertebrates. We propose a reconstruction of the main genes involved in sponge response to wounding (Fig. 4) and hypothesize that this gene regulation is mediated by cellular migration based on reports in other animals and on the observation of spherulous cell accumulation at the wound in *Aplysia* sponges, particularly 1 d post-treatment. In addition, the observed activation of Ets-domain transcription factors may point to differentiation by increased “pinacocyte-like” expression signature at the wound. Single-cell RNA-seq from the sponge *A. queenslandica* revealed that Ets-domain transcription factors are enriched in pinacocytes, sponge epidermal cells. The family of Ets-domain transcription factors is considered a metazoan novelty, with conserved roles in cellular differentiation and development in animals. It is difficult to unequivocally distinguish different cellular processes (i.e., migration, proliferation, and differentiation) based on gene expression patterns alone. Yet, migration and differentiation prevail at the wound in different sponge species, whereas proliferation seems more relevant in adjacent areas. Further studies are necessary to connect the molecular responses to cellular processes in sponges.

Wounding-induced gene expression patterns in the sponge resembled the early steps of regeneration in sponges and other animal phyla. The signaling pathways involved in the 3 h post-treatment transcriptomic response are similar to those identified at the onset of whole-body regeneration in sea stars (3 h post-treatment), *Hydra* (3 h post-treatment), and planaria (12 h post-treatment), which typically correspond to the wound response phase. In particular, they overlap in the regulation of MAPK and calcium signaling, and transcription factors. Another common player is the Wnt signaling pathway, which determines patterning in whole-body regeneration as well as in animal development (e.g., and references therein). In sponges, Wnt pathway is involved in both development and regeneration. Recently, Soubigou et al. monitored the morphological and gene expression changes involved in the regeneration of adult disassociated cells into a new functional sponge during the first 24 h after disassociation. They found the pathways involved in sponge regeneration and in sponge post-larval development. Despite the differences in the degree of damage and sampling time points, several of the genes and pathways in sponge regeneration identified by Soubigou et al. are also induced by wounding in our study, like the Wnt pathway, the TGFβ signaling, and genes involved in cell adhesion and extracellular matrix rearrangement. Overall, the transcriptomic response of the sponge to wounding agrees with conserved genetic features involved in tissue repair in animals with high regeneration capacity.

Sponge response to wounding revealed key genes in maintaining animal tissue homeostasis, from sponges to humans. In the set of proteins codified by sponge genes responding to mechanical damage, we identified putative orthologs of human proteins involved in G-protein coupled signaling pathway, ubiquitination, and components of the MAP kinase signaling transduction pathway. In humans, these proteins mediate diverse biological processes such as cell adhesion, proliferation, or differentiation (Supplementary Data S3 online). For some of these human genes, their malfunction or dysregulation has been observed during tumors (COSMIC). Finding similar genes in the context of sponge response to wounding adds additional support of their conserved role in regulating homeostasis. It has been previously reported that sponge genomes harbor genes related to those involved in human cancer. As one of the oldest animal groups till extant, sponges offer a unique opportunity for understanding the evolution of the functions and interactions of these cancer-related genes. Validating gene functions in sponges is difficult due to their limited amenability to manipulation, however, wound healing has the potential to provide an experimental context with a phenotype that can be directly linked to molecular responses involved in the fundamental process of tissue homeostasis.

**Conclusion**

Here, we provide the first detailed study of the transcriptomic response to wound healing in sponges. The transcriptomic response of *A. aerophoba* to grazing by *T. pervera* resembles the response to wounding caused by mechanical damage. Both wounding cues induced a strong activation of gene expression that, we hypothesize, may have a detrimental metabolic cost for the sponge. Wounding activated the Wnt signaling, MAPK signaling and a “clotting-like” cascade, which resembles the onset of whole-body regeneration in sponges and other animal phyla. The sponge transcriptomic response to wounding included putative orthologs of human genes that, when altered, yield cancer. Cancer development constitutes a disruption of tissue homeostasis and is largely controlled by ancient molecular pathways. Sponges are a key phylum to understand the emergence of animal life and we propose that sponge wound healing can offer an innovative system to research how these ancient genes and regulatory networks contribute to tissue homeostasis and animal health.

**Data availability**

Raw reads, metadata, and gene quantification matrices generated during this study are available in the ArrayExpress database at EMBL-EBI archive ([www.ebi.ac.uk/arrayexpress](http://www.ebi.ac.uk/arrayexpress)) under accession number E-MTAB-8601. De novo reference transcriptome assembly is available at the European Nucleotide Archive under project PRJEB46965. The annotation of the reference assembly is provided in Supplementary Data S4 online. Further generated/processed data are included in the Supplementary Material online.
Acknowledgements

We thank Dr. Rafel Coma, Manel Bolívar, and Marc Catllà (CEAB-CSIC) for support during sponge collection and experimentation. We thank Prof. Dr. Sebastian Fraune (Heinrich Heine University Düsseldorf) for feedback on an early version of the manuscript and Dr. Tyler Carrier (GEOMAR Helmholtz Centre for Ocean Research Kiel, GEOMAR) for language editing. We thank Prof. Dr. Ute Hentschel for providing access to lab and resources at the Marine Symbiosis Unit of GEOMAR.

Author contributions
L.P. and Y.W. conceived the experiments; L.P., Y.W., and M.R. planned and executed of the experiments; L.P. and Y.W. extracted the RNA; S.F. advised and performed the RNA sequencing; Y.W. performed the transcriptomic assembly; L.P. performed the differential gene expression analysis and further data analyses; L.P. prepared the figures and tables and wrote the paper. All authors edited and approved the final version of the manuscript.

Funding
Open Access funding enabled and organized by Projekt DEAL. L.P. was awarded a postdoctoral fellowship from Alexander von Humboldt Foundation, which was sponsored by The Future Ocean Cluster of Excellence. L.P. is currently at the Institute of Marine Science (ICM-CSIC) thanks to a fellowship supported by “la Caixa” Foundation (ID 10010434) and from the European Union's Horizon 2020 research and innovation programme under the Marie Skłodowska-Curie grant agreement No 847648, fellowship code is 104855. Financial support was provided by the Spanish Government through the Grant (RTI2018-094187-B-100) and the ‘Severo Ochoa Centre of Excellence’ accreditation (CEX2019-000928-S) to MR. This is a contribution from the Marine Biogeochemistry and Global Change group from the Generalitat de Catalunya (2017SGR1011).
Competing interests
The authors declare no competing interests.

Additional information
Supplementary Information The online version contains supplementary material available at https://doi.org/10.1038/s41598-022-05230-x.

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