Comparative transcriptomic analysis reveals gene regulation mediated by caspase activity in a chordate organism

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Abstract

Background: Apoptosis is a caspase regulated cell death present in all metazoans defined by a conserved set of morphological features. A well-described function of apoptosis is the removal of excessive cells during development and homeostasis. Recent studies have shown an unexpected signalling property of apoptotic cells, affecting cell fate and/or behaviour of neighbouring cells. In contrast to the apoptotic function of cell elimination, this new role of apoptosis is not well understood but seems caspase-dependent. To deepen our understanding of apoptotic functions, it is necessary to work on a biological model with a predictable apoptosis pattern affecting cell fate and/or behaviour. The tunicate Ciona intestinalis has a bi-phasic life cycle with swimming larvae which undergo metamorphosis after settlement. Previously, we have shown that the tail regression step during metamorphosis, characterized by a predictable polarized apoptotic wave, ensures elimination of most tail cells and controls primordial germ cells survival and migration.

Results: We performed differential transcriptomic analysis between control metamorphosing larvae and larvae treated with the pan-caspase inhibitor Z-VAD-fmk in order to explore the transcriptional control of apoptotic cells on neighbouring cells that survive and migrate. When caspase activity was impaired, genes known to be involved in metamorphosis were downregulated along with other implicated in cell migration and survival molecular pathways.

Conclusion: We propose these results as a confirmation that apoptotic cells can control surrounding cells fate and as a reference database to explore novel apoptotic functions in animals, including those related to migration and differentiation.

Keywords: Comparative transcriptomics, Caspases, Apoptosis, Metamorphosis, Migration, Ascidians

Background

Apoptosis is a regulated cell death defined by morphological features and depending on caspases [1–3]. Being ubiquitous in metazoans [4–7], the main function of apoptosis is the elimination of unwanted cells [8], such as the interdigital tissues during embryogenesis of tetrapods [9] or larval tissues during metamorphosis of many animals [10]. Apoptosis is also implicated in homeostasis by fine-tuning the balance between cell death and proliferation like in gonad size control in rodents [11].

While the function of cell elimination by apoptosis is highly documented, apoptosis seems to have a broader morphogenetic function that goes beyond cells elimination. Accumulating evidence in many metazoans suggest that apoptotic cells can emit caspase-dependent signals to their neighbours, modulating their fate (survival/death, differentiation or proliferation) or behaviour (migration) [12, 13]. Apoptotic-induced proliferation was...
reported in several organisms, such as in the cnidarian *Hydra*, where apoptosis promotes proliferation of adjacent cells through emission of Wnt3 signalling in a caspase-dependent manner during head regeneration [14]. Similar observations have been made during rodent liver regeneration, where cleavage of iPLA2 by caspases in apoptotic cells led to proliferation of adjacent cells [15]. Apoptotic cells can also induce differentiation, as was reported during the metamorphosis of the cnidarian *Hydractinia echinata* [16]. Finally, apoptotic cells modulate death or survival balance, as was observed in the *Drosophila* imaginal disc, by producing the Tie ligand Pvf1, that induces death resistance of neighbouring cells [17]. Apoptotic-dependent migration is well described in mammals, in which apoptotic cells secrete directional signals such as Lysophophatidylcholine or Sphingosine-1-Phosphate, promoting migration of leukocytes [18, 19]. Furthermore, in *Xenopus laevis* tadpole, an apoptotic-dependent axon guidance was suggested during tail regeneration after injury [20].

While caspase-dependent signals emitted by apoptotic cells start to be characterised, almost no target genes are identified in receptor cells. Evaluating the capacity of caspases to affect gene expression and influence fate and/or behaviour of cells adjacent to apoptosis is the first fundamental step to better characterise this new function in animals.

In this report, we took advantage of the predictable spatiotemporal apoptotic profile of ascidians [21, 22], in particular during the *Ciona intestinalis* metamorphosis, which is concomitant with cell survival and cell migration [21, 23, 24]. *Ciona* belongs to the Urochordata, the sister group of vertebrates [25], and has a biphasic life cycle composed of a pelagic larva and a benthic adult. Embryogenesis generates a tadpole-like swimming larva consisting of an anterior trunk and a posterior tail which, after few hours of swimming, settles and metamorphoses into a sessile juvenile. A hallmark of metamorphosis is the apoptotic-dependent tail regression [21] during which apoptosis starts at the tip of the tail and propagates toward the trunk, making it easily anticipated; a unique feature among chordates. Using TUNEL labelling and electron microscopy, we previously described in detail this apoptotic profile. Apoptosis starts in epidermal cells at the most posterior tail tip, then, it affects the notochord cells and the striated muscle cells in a postero-anterior wave. The tunic cells are also randomly affected. On the opposite, two tissues survive and migrate during, or prior to, tail regression; the ventral endodermal strand and the eight primordial germ cells (PGC) [23, 24, 26]. PGC, located at the postero-ventral side of the tail tip, escape from cell death by moving toward the trunk, and are consequently surrounded by apoptotic cells during the tail regression. Interestingly, we previously demonstrated that the pan-caspase inhibitor Z-VAD-Fmk blocks both apoptotic dependent-tail regression and PGC migration, suggesting that PGC migration is regulated by apoptotic cells [23].

To identify genes controlled by caspases activity during tail regression, we performed a comparative transcriptomic analysis at the beginning and at mid-tail regression between control metamorphosing larvae or larvae exposed to the pan-caspase inhibitor Z-VAD-fmk, already successfully used in *Ciona* [21] (Fig. S1). Based on a de novo transcriptome, we identified 61 genes differentially expressed between the beginning and mid-tail regression in control larvae. Next, we found 65 genes differentially expressed between control and treated larvae (21 and 44 at the beginning and at mid-tail regression, respectively). Interestingly, misregulation of some genes upon caspase inhibition was reported during ascidian metamorphosis, but also reported in cell survival and cell migration in several animals [27–30]. Our results offer a first database of genes transcriptionally modulated by apoptosis during a dynamic morphogenetic process in a chordate, allowing exploration of novel apoptotic functions in animals.

**Results**

**Genes differentially express between the onset and the mid-tail regression**

Globally, the robustness of our analysis is shown by a heat map allowed visualization of result congruence between replicates (Fig. 1). To confirm relevance of our data, we randomly chose 6 genes differentially expressed in our transcriptomic analysis to performed real-time PCR (Fig. 2A). Five of them exhibited an expression profile that was similar for both experiments.

One of them, KH.C7.314, is less expressed at the mid-tail regression in comparison to the beginning of the process in control larvae in our transcriptomic analysis. Similar profile was observed with real-time PCR, but the difference is less pronounced and not statistically significant. Despite this, control by real-time PCR globally validated our transcriptomic data.

First, we have focused on genes differentially expressed in control metamorphosing larve between the beginning and the mid-tail regression. Twenty-five genes (41%) were up-regulated from the beginning to the mid-tail regression with the strongest increase recorded for KH.C9.385 (*smyd3*) (Table 1). Conversely, 36 genes (59%) show decreased expression, with a subset of genes whose expression was no longer detectable such as KH.14.261 (*lysozyme*-like) (Table 2). Among these identified genes, several were already reported during a previous cDNA chip analysis of *Ciona* metamorphosis, such as KH.C7.314 (*MED30*, KH.C11.274 (*p-selectin*, known as *Ci-sushi*) which is expressed in almost all the tail, or...
Fig. 1 (See legend on next page.)
KH.L132.4 (matrilin-2) expressed in epidermal cells at the tip of the tail in premetamorphic larva (Fig. 2B) [29]. In addition, some of the genes identified by our screen had already been detected by substantial RNA sequencing data coupled with in situ hybridization data available on ANISEED database (http://www.aniseed.cnrs.fr). For example, KH.L132.16 (fibrillin-3) starts to be expressed at mid-neurulae with a maximum expression in larva, in which it is detected in all the epidermis including the tail. Furthermore, KH.C10.250 (fox-1, RNA-binding) is expressed in tail muscle at mid-tailbud stage to become ubiquitous in larva [31].

We also observed that some among these were also impacted by Z-VAD-fmk treatment (Tables 3, 4, 5 and 6) such as KH.C6.197 (meta2), KH.C1.10 (LIM), KH.C1.216 (myosin light chain 3), and KH.C11.274 (Csi-sushi) [27–30]. The detection of genes already known to be expressed in the tail and involved in tail regression (i.e. Csi-sushi, Csi-meta2) is consistent with previous studies on ascidians metamorphosis, validates our experimental procedures and supports the relevance of our database.

**Caspase inhibition affects gene expression during the tail regression**

In total, 65 genes were affected by the inhibition of caspsases during Ciona tail regression (Fig. 1). At the beginning of tail regression 12 genes were upregulated by the Z-VAD-fmk treatment (so negatively regulated by caspase activity in physiological conditions) with the greatest expression increase observed for KH.C11.506 (girdin) and KH.S1380.1 (mucin-5 AC) (Table 3), and 9 genes were downregulated (so positively regulated by caspase activity) with extreme effects on KH.C1.66, which is no longer expressed (Table 4).

At mid-tail regression, we identified 38 upregulated genes in the Z-VAD-fmk condition with the highest increase noted for KH.C12.516 (Table 5), and 6 downregulated genes (Table 6) with strongest effect concerning KH.C1.910.

**Transcription is necessary for the PGCs migration**

We previously demonstrated that Z-VAD-fmk blocked both tail regression and PGC migration, and that apoptotic wave propagation and PGC migration speed were correlated [23]. Here, we show that transcription is necessary for PGC survival and migration (Fig. 3). We exposed metamorphosing larvae to the transcription inhibitor actinomycin-D; in the control (devoid of actinomycin-D), a classical apoptotic profile is observed with PGC movement, as already described [23], and in actinomycin-D-treated larvae PGC migration is blocked and PGC nuclei become TUNEL-positive, indicating
destruction as the apoptotic wave progresses (Fig. 3). This result, and our previous studies, argue that survival and migration of PGC depend on transcription that could be partially controlled by caspase activity from apoptotic cells (Fig. 3). The comparative transcriptomic analysis between control and Z-VAD-fmk treated larvae is fundamental to understand this apoptotic dependent migration.

**Discussion**

**Caspase activity modulates expression of genes implicated in cell death/survival**

Interestingly, the 2 genes most inhibited by caspase activity at the beginning of tail regression, *girdin* and *mucin-5 AC*, were both reported to promote survival in mammalian cells [32, 33]. Among the other genes negatively impacted by caspase activity, KH.C1.1084, the thioredoxin homologue in mammals, is able to control the activation of the executioner caspase 3, the initiator caspase 9, and the release of the cytochrome c, all characterising the intrinsic apoptosis [34–37]. In addition, KH.C3.45 (*kinase C alpha*) homologue is known to promote survival and increases anti-apoptotic Bcl-2 efficiency in cultured cells [38]. Taken together, our database highlights that cells expressed a variety of genes regulating cell death/survival fate in a caspase dependent manner, suggesting that apoptotic cells actively participate in apoptosis propagation during tail regression.

**Caspase activity modulates expression of genes implicated in cell migration**

We identified numerous genes modulated by caspase activity, with a majority of them upregulated, previously observed in cell migration regulation. It is the case of KH.C1.216 (*myosin light chain 3*) and KH.C9.494 (*myosin light chain kinase*) for which homologues are key players in mesenchymal migration [32, 39] and KH.C13.153 (*uromodulin*-like) which provides guidance cues for neutrophile migration [40]. Furthermore,
| KyotoGrail KH gene model | Expression ratio | Identity and/or GO function | References in ascidians/reported functions in animals |
|-------------------------|------------------|-----------------------------|---------------------------------------------------|
| KH.C14.261              | 0.023            | putative lysozyme-like protein; PE-PGRS family protein | x |
| KH.L37.67               | 0.025            | TGM1, Transglutaminase      | x |
| KH.C7.501               | 0.026            | ferritin                    | x |
| KH.C1.459               | 0.026            | Sorting nexin-18            | x |
| KH.C4.177               | 0.028            | Glucocorticoid receptor-like, Pinch protein | x |
| KH.C7.314               | 0.050            | mediator of RNA polymerase 2 transcription subunit 30 (med30) | Chambon et al. 2007 (c0100134193) apoptosis |
| KH.C14.281              | 0.050            | glycerol-3-phosphate (1)-acyltransferase | x |
| KH.L142.4               | 0.056            | universal stress protein YxiE | x |
| KH.C1.263               | 0.061            | ribosomal protein L4        | x |
| KH.C10.335              | 0.061            | calmodulin like protein     | migration |
| KH.C11.384              | 0.092            | uncharacterized protein LOC100186758 | x |
| KH.L112.30              | 0.103            | transmembrane matrix receptor MUP-4-like | x |
| KH.C1.1024              | 0.129            | glutamate-rich WD repeat-containing protein 1 | x |
| KH.C4.94                | 0.129            | lipopolysaccharide-induced tumor necrosis factor-alpha factor | x |
| KH.L92.1                | 0.144            | protein DDB_G0283697        | x |
| KH.C2.124               | 0.155            | Glycerol-3-phosphate (1)-acyltransferase | x |
| KH.L108.23              | 0.167            | protein DD3-3               | x |
| KH.C11.93               | 0.174            | N-acetyltransferase 6       | x |
| KH.C10.133              | 0.194            | holine-phosphate cytidlyltransferase A-like | x |
| KH.C3.231               | 0.216            | uncharacterized protein LOC100184171 | x |
| KH.L59.3                | 0.227            | uncharacterized protein LOC100175851 | x |
| KH.C4.450               | 0.235            | sushi, von Willebrand factor type A, EGF and pentraxin domain-containing protein 1 | x |
| KH.C11.274              | 0.243            | P-selectin-like             | Chambon et al. 2007 (c0100139289) apoptosis |
| KH.C4.73                | 0.268            | uncharacterized protein LOC100175287 | x |
| KH.C14.356              | 0.275            | putative lysozyme-like protein | x |
| KH.C11.287              | 0.278            | unc-93 homolog A-like       | x |
| KH.C14.337              | 0.313            | lysozyme-like protein; PE-PGRS family protein PE_PGRS30 | x |
| KH.C14.549              | 0.335            | lysozyme-like protein; PE-PGRS family protein PE_PGRS30 | x |
| KH.L108.30              | 0.344            | uromodulin-like 1 precursor | migration |
| KH.C12.655              | 0.390            | uncharacterized protein LOC113474772 | x |
| KH.L132.16              | 0.428            | fibrillin-3; MATN2          | migration |
| KH.L132.4               | 0.428            | matrilin-2; MEGF6           | Chambon et al. 2007 (c0100138016) migration |
| KH.C2.613               | 0.443            | zinc transporter 9-like     | x |
| KH.C11.454              | 0.446            | uncharacterized protein LOC100184022 | x |
| KH.C8.121               | 0.463            | SCO-spondin                 | x |
| KH.C4.402               | 0.470            | galanin receptor type 3     | x |
integrin subunits and components of the Extra Cellular Matrix, crucial for mesenchymal migration, are well represented in our transcriptomic approach. In addition, KH.C1.10 (lasp1) mammalian homolog is necessary for migration of NIH 3 T3 cells [41, 42].

Furthermore, KH.C12.516 (ceramidase) could be involved in tissue migration regulation, as it has been described as participating in the sphingolipid pathways. Interestingly, sphingosine 1-phosphate signalling leads to tissue migration [43], particularly in germline cells (due to a sphingosine 1-phosphate receptor) in the tunicate Botryllus [44], making this pathway a strong candidate for PGC migration in Ciona. Given that these genes and PGC migration are both controlled by caspases [23], PGC migration can be regulated by apoptotic cells.

**Ascidians as crucial biological models to studies novel apoptotic function**

This study, coupled with our previous work suggested that apoptosis regulates the migration of primordial germ cells [23], confirm Ciona as a perfect biological model to understand apoptotic-induced migration. In addition, a regulation of cell proliferation by apoptosis was reported in this species [45]. Ascidians are well known to have an impressive regenerative capacity for a chordate, in particular in the context of the reconstruction of the distal body parts after amputation, thanks to stem cell reorganization from the branchial sac [46–48].

Apoptosis was reported to occur as a transient event at the site of injury, and precedes the regeneration of the neural complex, the oral siphon, or the atrial siphon [49]. In the case of siphon amputation, it has been shown by caspases inhibition experiment that an apoptotic-dependant Wnt signalling allows the cell proliferation, leading to the successful regeneration [45]. In addition, apoptosis appears crucial to maintain the growth of branchial sac and its inhibition lead to a hazardous organization of this tissue, suggesting a fundamental function in homeostasis maintaining. Importantly, stem cells requisition and migration from the branchial sac toward the wounding site seem also dependant on caspases, making it an other potential

| Table 3 | Genes upregulated by the Z-VAD-fmk at the tail regression beginning |
|-----------------|-------------------|-----------------|-------------------------------|
| KyotoGrail KH gene model | Expression ratio | Identity and/or GO function | References in ascidians/reported functions in animals |
| KH.C11.506 | 9850 | girdin | apoptosis |
| KH.S1380.1 | 7405 | mucin-5 AC | apoptosis |
| KH.C8.46 | 4453 | vesicle-associated membrane protein 4-like | x |
| KH.C1.451 | 4028 | orphan gene | x |
| KH.C14.511 | 3917 | myotubularin-related protein 2-like | x |
| KHL41.62 | 3503 | reticulocalbin-2-like | x |
| KHL152.1 | 3305 | proline-rich protein PRCC | x |
| KH.C11.93 | 2521 | gamma-glutamylycyclotransferase 2; N-acetyltransferase 6 | x |
| KH.C11.218 | 2463 | cytospin-A isoform XI | x |
| KH.C1.1084 | 2290 | thioredoxin-2-like | apoptosis |
| KHL105.13 | 2121 | DENN domain-containing protein 5B-like | x |
| KHL.C12.373 | 2011 | uncharacterized protein LOC100175595 | x |

| Table 4 | Genes downregulated by the Z-VAD-fmk at the tail regression beginning |
|-----------------|-------------------|-----------------|-------------------------------|
| KyotoGrail KH gene model | Expression ratio | Identity and/or GO function | References in ascidians/reported functions in animals |
| KHL9.1.1 | 0.473 | protein DDB_G023697 | x |
| KHL.C1.197 | 0.449 | META2 protein precursor | Nakayama et al., 2001–2002; Chambon et al., 2007 |
| KH.C1.216 | 0.369 | myosin light chain 3, skeletal muscle | Nakayama et al., 2002 |
| KHL24.20 | 0.366 | collagen alpha-6(VI) chain-like | migration |
| KHL154.8 | 0.309 | unnamed protein product | x |
| KHL.C1.153 | 0.308 | uromodulin-like | migration |
| KHL.C3.237 | 0.229 | 605 ribosomal protein L7a | x |
| KHL.C1.103 | 0.098 | nuclear lamin-L (II)-like | x |
| KHL.C1.66 | 0.000 | orphan gene | x |
| KyotoGrail KH gene model | Expression ratio | Identity and/or GO function                          | References in ascidians/reported functions in animals |
|--------------------------|------------------|------------------------------------------------------|-------------------------------------------------------|
| KL.C12.516               | 11,357           | neutral ceramidase-like                              | Kassmer et al. (2015) migration                       |
| KL.C9.170                | 10,571           | protein CNPPD1                                      | x                                                     |
| KL.C2.541                | 7634             | cell cycle control protein 50A                       | x                                                     |
| KL.C3.623                | 7551             | kinesin-like protein 2                               | x                                                     |
| KL.C4.189                | 6713             | ribosomal protein CEP52                              | x                                                     |
| KL.L124.15               | 5285             | vesicular integral-membrane protein VIP36            | x                                                     |
| KL.S522.7                | 5242             | TPA: zinc finger protein                             | x                                                     |
| KL.C14.231               | 4625             | 60S ribosomal protein L12                            | x                                                     |
| KL.S605.3                | 4398             | gamma-crystallin 5                                   | x                                                     |
| KL.C9.791                | 4198             | integrin alpha-2-like                                | migration                                             |
| KL.C4.73                 | 4127             | uncharacterized protein LOC100175287                 | x                                                     |
| KL.L5.23                 | 4122             | uncharacterized protein LOC100179749                 | x                                                     |
| KL.C1.568                | 3255             | uncharacterized protein LOC100176279                 | x                                                     |
| KL.S390.2                | 3227             | RNA-binding motif, single-stranded-interacting protein 1 | x                                                     |
| KL.C9.660                | 3187             | supervillin-like                                     | migration                                             |
| KL.L106.13               | 3179             | DENN domain-containing protein 5B-like               | x                                                     |
| KL.S534.8                | 3075             | coiled-coil domain-containing protein 178-like       | x                                                     |
| KL.L108.23               | 2846             | protein DD3–3                                        | x                                                     |
| KL.C7.514                | 2824             | SLIT and NTRK-like protein 3                         | x                                                     |
| KL.C2.448                | 2822             | plasminogen                                          | x                                                     |
| KL.C7.297                | 2732             | solute carrier family 22 member 21                   | x                                                     |
| KL.L37.43                | 2586             | L-threonine ammonia-lyase                            | x                                                     |
| KL.C11.274               | 2571             | P-selectin-like                                      | Chambon et al., 2007 (ci0100139289) apoptosis         |
| KL.C11.218               | 2518             | cytospin-A                                           | x                                                     |
| KL.L5.93                 | 2488             | uncharacterized protein LOC100175851                 | x                                                     |
| KL.C10.71                | 2418             | 3-phosphoinositide-dependent protein kinase 1        | x                                                     |
| KL.C9.715                | 2397             | neurobeachin-like protein 1                          | x                                                     |
| KL.C3.562                | 2392             | unconventional myosin-X                             | x                                                     |
| KL.C4.94                 | 2374             | lipopolysaccharide-induced tumor necrosis factor-alpha factor | x                                                     |
| KL.S534.2                | 2333             | RB1-inducible coiled-coil protein 1                  | x                                                     |
| KL.C9.496                | 2327             | uncharacterized protein LOC100183253                 | x                                                     |
| KL.C1.10                 | 2314             | LIM and SH3 protein                                  | Terasaki et al., 2008                                 |
| KL.C11.93                | 2252             | N-acetyltransferase 6                                | x                                                     |
| KL.L11.320               | 2179             | transmembrane matrix receptor MUP-4-like             | x                                                     |
| KL.C11.567               | 2162             | serine palmitoyltransferase 2                        | x                                                     |
| KL.L23.74                | 2064             | uncharacterized protein LOC1001758692                | x                                                     |
| KL.C11.637               | 2056             | putative glutathione-specific gamma-glutamylcyclotransferase 2 | x                                                     |
| KL.C7.260                | 2031             | tropomyosin-like protein                             | Nakayama et al., 2002 migration                       |
apoptotic-dependant migration [45]. Taken together, all of these studies make *Ciona* one of the rare biological models where several cell fates and behaviours, controlled by apoptotic cells in a caspases dependent manner, has been put in evidence or suggested in adults as well as in larvae.

In the colonial ascidian *Botryllus schlosseri*, apoptosis is known to play on budlets elimination [50–52]. In this species, blastogenetic cycle (asexual reproduction of adults) give rise to a bud, itself given next four budlets (reviewed in [53, 54]). Usually, two of them growth despite the two other undergo apoptosis and are reabsorbed. In addition to this function in the blastogenetic cycle, apoptosis was also reported in tunic, epidermis, branchial leaflet, vessel epithelium, circulatory system or gonads in adults. With so many tissues presenting apoptosis in physiological condition, it seems clear that apoptosis should have crucial functions in *Botryllus* and likely could participate to regulation of other cell fates and behaviours as it was reported in *Ciona*. All of these tissues are as many opportunities to deepen or understanding of apoptosis functions in a same species.

**Conclusions**

Here, we offer a potential roadmap in order to design future functional studies on *Ciona* and other animals, including mammals, where apoptosis takes place together with other cell fates and behaviours. Identification of molecular actors needing cleavage for their activation, cell receptors receiving signals from apoptotic cells, or

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**Table 6** Genes down regulated by the Z-VAD-fmk at mid-tail regression

| KyotoGrail KH gene model | Expression ratio | Identity and/or GO function | References in ascidians/reported functions in animals |
|--------------------------|------------------|-----------------------------|----------------------------------------------------|
| KH.C11.688               | 0,385            | thioredoxin-related transmembrane protein 1-like | x                                                   |
| KH.C9.494                | 0,319            | myosin light chain kinase, smooth muscle | Nakayama et al., 2002 migration                     |
| KH.C3.865                | 0,270            | orphan gene                 | x                                                   |
| KH.C7.361                | 0,237            | regulator of chromosome condensation 1 (rcc1) | x                                                   |
| KH.C3.45                 | 0,217            | protein kinase C alpha type | Apoptosis                                           |
| KH.C1.910                | 0,109            | low-density lipoprotein receptor-related protein 2 | x                                                   |

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**Fig. 3**

A. Larvae labelled by TUNEL (green) with VASA immunostaining (red) and counterstained with DAPI (blue). PGC of larvae treated with the transcription inhibitor actinomycin-D (1 µg/ml) show impaired migration and undergo apoptosis. B. Representation of apoptotic functions during tail regression. Apoptotic cells emit signals modifying transcription of adjacent cells leading to PGC survival and migration to the trunk.
transcription factors could be promising for further research.

The last decade has brought evidence that the cleavage capacity of caspases is a prerequisite to activate molecular actors promoting migration or differentiation in a non-apoptotic context [55–59]. Consequently, Z-VAD-fmk blocks all processes led by caspases, making it crucial to distinguish the level of implication of apoptotic and non-dying cells for future research.

Finally, our study provides an additional and novel chapter in the research of apoptotic functions beyond their role as a simple cell elimination mechanism, opening opportunities in the field of medical science [60].

Methods
Sample collection
Adult Ciona intestinalis were collected in the field in Brittany (France) by the Centre National Marine Biological Resources, EMBRC-France, Roscoff (Station Biologique de Roscoff, Sorbonne Université, Brittany, France). Individuals were maintained in 35% salinity artificial seawater (ASW) at 18 °C in the UMR7138 Evolution Paris Seine laboratory (Sorbonne University, Paris, France). Oocytes and sperm were obtained by dissection before RNA extraction. Embryos were cultured at 18 °C in 0.2 μm filtered ASW with 0.1 M Hepes (#H4034, Sigma-Aldrich; Merck KGaA). Embryos of each fertilisation were divided into two subsets before the first cell division to obtain one control and one treatment condition. Six fertilisations were performed, divided into twelve petri dishes (6 control and one treatment condition). Six fertilisations were performed, divided into twelve petri dishes (6 control and one treatment condition). After the treatment, larvae were collected at two time points (see below), and compacted by manual centrifugation before RNA extraction.

Pharmacological treatments and TUNEL labelling
The pan-caspase inhibitor Z-VAD-Fmk (#V116, Sigma-Aldrich; Merck KGaA) was stored in DMSO (#D8418, Sigma-Aldrich; Merck KGaA) at −20 °C and used at final concentration of 10 μM. After the settlement of more than 50% of larvae on the dish, and when some of them stopped moving (indicating that metamorphosis will begin soon), supernatant was discarded and replaced with filtered ASW containing DMSO as control, or Z-VAD-Fmk diluted in DMSO as treatment. Z-VAD-Fmk was renewed every two hours. The experiment was divided into two time points (with three biological replicates for each one – itself divided between control and treated larva) following control tail regression stages: first when the tail regression just began in the control (beginning), and the second at mid-progression (mid-tail regression) given three controls with three Z-VAD-Fmk equivalents for each point.

In a separate experiment, 1 μg/ml of actinomycin-D (#A1410, Sigma-Aldrich; Merck KGaA) was added to swimming larvae. After settlement and tail regression initiation, paraformaldehyde fixation, TUNEL labelling and Vasa immunostaining were performed as previously described [23].

RNA preparation and sequencing
RNA was extracted using the RNAqueous™-Micro Total RNA Isolation Kit (#AM1931 – ThermoFisher Scientific) according to the manufacturer’s protocol. DNase treatment was performed using the TURBO DNA-free™ Kit (#AM1907 - ThermoFisher Scientific). RNA was purified and concentrated using the RNasy MinElute Cleanup Kit (#74204 – Qiagen) according to the manufacturer’s protocol. RNA quantity was measured using a NanoDrop and quality was assessed with the Experion RNA HighSens (#700–7155 – BioRad) from 1 μL (0.1 μL of RNA completed with 0.9 μL of Milli-Q water) following standard instructions. Transcriptomes were sequenced on Illumina Hi-Seq 2000 (150 bp reads) at the Sistemas Genomicos plate-form of the Ascires Biomedical Group (Spain).

De novo transcriptome assembly and differential expression
Raw reads of the twelve Ciona intestinalis samples were checked with FastQC and merged to assemble a de novo transcriptome with Oases [61] and Trinity [62]. Best assemblies from each one (best N50) were merged with Cap3 to obtain the de novo transcriptome. De novo transcriptome contains 85,506 contigs (N50 = 1314, minimum contig length = 102, maximum contig length = 13,998). Then, raw reads of each sample were mapped against the de novo transcriptome with Bowtie2 [63] with standard parameters. Low quality mapping reads (MQ > 20) were removed using Samtools [64] and Picard Tools (http://broadinstitute.github.io/picard). Expression quantification was performed using HTSeq [65] and differential expression evaluated with the DESeq2 package from R. We kept the genes with an expression ratio > 2 (upregulated) and lower than 0.5 (downregulated). Then, Gene Ontology (GO) enrichment was performed with Blast2GO 5.2 and sequences were blasted on the Ciona genome to identify their KyotoGrail KH gene model.

Real-time PCR
We synthetised the cDNA from RNA coming from same sample which was sending for sequencing using the SuperScript™ II Reverse Transcriptase (18,064,014 - ThermoFisher Scientific) according to the manufacturer protocol. Real-time PCR was performed with the SYBR
Green Supermix (Biorad) on a Biorad’s thermal cycler with the following profile: 95 °C for 10 min; 40 cycles of amplification with successively 95 °C for 15 s, 60 °C for 10s, and 72 °C for 20s; one cycle for melting curve analysis with an acquisition every 0.5 °C from 65 °C to 95 °C to verify the presence of a single product. Each assay included four successive dilutions as standard to determine the reaction efficiencies and Ct values. Reactions were done in triplicate. All PCR amplicon have a length from 140 to 160 bp. Normalized was made using the reference gene Ci-actin (forward 5′ ATGTGCAAGG CCGTTT 3′); reverse 5′ GACACGGAGTTCGTTGT 3′), already successfully used [66]. Genes of interest and their primer pair are: KH.09.385 with forward 5′ ACTT CTGAAAGAGCGGACG 3′ and reverse 5′ CATTG- CAAACAGACCATCTTGC 3′, KH.07.314 with forward 5′ AGCCAACCTACCGAATGGGA 3′ and reverse 5′ GTTCACGTCTCACAACCTCTC 3′, KH.01.103 with forward 5′ TGGCTGA- GATACCGAAGCTG 3′ and reverse 5′ CATTTGCTCAAGATCGGCG 3′, KH.09.791 with forward 5′ GGATCTAAGCAGTGTCTGGAC 3′ and reverse 5′ CAAGGGCGTTGGTGTTCAGT 3′, KH.09.170 with forward 5′ AATCCCCGCCTTGGAA GAAA 3′ and reverse 5′ GGGGTGATTGATGAAATAC ATGG 3′, and KH.01.910 with forward 5′ ACATGAAG ATTTTGGTACCGAC 3′ and reverse 5′ CAAAAC TTGGCCAAAAAGTGTGG 3′. Differences was evaluated by Wilcoxon Mann-Whitney tests using R i386 4.0.3 software, and considered statistically significant for p-value < 0.05.

In situ hybridization combined with vasa immunodetection
Larvae were fixed 2 h at 4 °C in MEM-PFA (4% paraformaldehyde, 0.1 M MOPS pH 7.4, 0.1 M NaCl, 1 mM EGTA, and 2 mM MgSO4, 0.05% Tween-20). After fixation larvae were washed three times and dehydrated through a graded series of ethanol/PBS baths and stocked in 100% ethanol at −20 °C. Rehydration was made by successive washes of ethanol/PBS solution to a full PBS final solution. Hybridization was made according to a previous protocol [67] except that methanol was replaced by ethanol. After hybridization, immunodetection of VASA was conducted as we previously described [23].

Supplementary Information
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Authors’ contributions
JP managed the project and made in situ hybridization. GK performed Z- VAD-fmk experiments, RNA extractions, real-time PCR, analysis, created figures and wrote the manuscript. AK performed actinomycin and TUNEL experiments. All authors contributed to comments on the manuscript. The authors read and approved the final manuscript.

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Availability of data and materials
Any request can be addressed to the corresponding author. Raw data are available under BioProject PRIN725676.

Declarations
Ethics approval and consent to participate
According to the French legislation (Décret n° 2013-118 du 1er février 2013 relatif à la protection des animaux utilisés à des fins scientifiques), experiments on Ciona intestinalis do not require an approval from Animal Ethics Commission because its not a “live vertebrate animals” and not a “live cephalopods”.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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References
1. Kerr JF, Wyllie AH, Currie AR. Apoptosis: a basic biological phenomenon with wide-ranging implications in tissue kinetics. Br J Cancer. 1972;26(4): 239–57. https://doi.org/10.1038/bjc.1972.33.
2. Hengartner MO. The biochemistry of apoptosis. Nature. 2000;407(6805):770–6. https://doi.org/10.1038/35037710.
3. Galluzzi L, Vitale I, Aaronson SA, Abrams JM, Adam D, Agostinis P, et al. Molecular mechanisms of cell death: recommendations of the nomenclature committee on cell death 2018. Cell Death Differ. 2018;25(3): 486–541. https://doi.org/10.1038/s41418-017-0012-4.
4. Lai M, David CN, Böttger A. Apoptosis in pre-Bilaterians: Hydra as a model. Apoptosis. 2010;15(3):269–78. https://doi.org/10.1007/s10495-009-0442-7.
5. Accordi F, Chimenti C. Programmed cell death in the pancreas of Bufo bufo during metamorphosis. J Anat. 2001;199(1 Pt 4):419–27. https://doi.org/10.1046/j.1469-7580.2001.19940419.x.

6. Kiss T. Apoptosis and its functional significance in molluscs. Apoptosis. 2010;15(3):313–21. https://doi.org/10.1007/s10495-009-0446-3.

7. Jiang C, Barhecke EH, Thummel CS. Steroid regulated programmed cell death during Drosophila metamorphosis. Development. 1997;124(2):4673–83. https://doi.org/10.1242/dev.124.22.4673.

8. Jacobson MD, Well M, Raff MC. Programmed cell death in animal development. Cell. 1997;88(3):347–54. https://doi.org/10.1016/S0092-8674(00)81873-5.

9. Wood W, Turmaine M, Weber R, Camp V, Maki RA, McKercher SR, et al. Mesenchymal cells engulf and clear apoptotic footplate cells in macrophageless Pu1 null mouse embryos. Development. 2000;127(4):5245–52. https://doi.org/10.1242/dev.127.4.354.

10. Nishihava A, Hayashi H. Spatial, temporal and hormonal regulation of apoptotic cell death during metamorphosis of the frog Xenopus laevis. Differentiation. 1995;59(4):407–14. https://doi.org/10.1006/diff.1995.5902.

11. Furuta I, Porpka-Heiskanen T, Scarpelous K, Tapanainen J, Turek FW, Hsueh PJ. Evidence for an instructive role of Wnt in osteoclast and osteoblast development. Dev Biol. 2003;255(1):62–70. https://doi.org/10.1016/S0012-1606(03)00270-6.

12. Chambon J-P, Nakayama A, Takamura K, McDougall A, Satoh N. ERK- and JNK-signalling regulate gene networks that stimulate metamorphosis and apoptosis in tail tissues of ascidian tadpoles. Development. 2007;134(6):1203–19. https://doi.org/10.1242/dev.002220.

13. Fuchs Y, Steller H. Live to die another way: modes of programmed cell death. Genes Dev. 2000;14(3):353–74. https://doi.org/10.1101/gad.14.3.353.

14. Chambon J-P, Soule J, Pomies P, Fort P, Sahuquet A, Alexandre D, et al. Tail regression in Ciona intestinalis: modulation by the TGF-beta and Notch pathways to promote wound healing and tissue regeneration. Dev Cell. 2009;17(2):279–90. https://doi.org/10.1016/j.devcel.2009.07.013.

15. Chambon J-P, Nakayama A, Takamura K, McDougall A, Satoh N. ERK- and JNK-signalling regulate gene networks that stimulate metamorphosis and apoptosis in tail tissues of ascidian tadpoles. Development. 2007;134(6):1203–19. https://doi.org/10.1242/dev.002220.

16. Wittig K, Kasper J, Seipp S, Leitz T. Evidence for an instructive role of apoptosis during the metamorphosis of Hydractinia echinata (Hydrozoa). Dev Biol. 2002;242(2):188–203. https://doi.org/10.1006/dbio.2002.0538.

17. Baehrecke EH, Thummel CS. Steroid regulated programmed cell death. Development. 2002;129(13):3017–30. https://doi.org/10.1242/fj.0210871.

18. Shirae-Kurabayashi M, Nishikata T, Takamura K, Tanaka KJ, Nakamoto C, Nakamura A. Dynamic redistribution of vasa homolog and exclusion of apoptotic-dependent tail re-egression during Ciona larva metamorphosis. Dev Biol. 2009;332(2):304–17. https://doi.org/10.1016/j.ydbio.2009.07.014.

19. Lak B, Chambon J-P, Nakayama A, Takamura K, McDougall A, Satoh N. Isolation and characterization of genes that are regulated by the TGF-beta and Notch pathways to promote wound healing and tissue regeneration. Cell. 2003;113(6):717–31. https://doi.org/10.1016/S0012-1606(03)00270-6.

20. Kiss T. Apoptosis and its functional significance in molluscs. Apoptosis. 2010;15(3):313–21. https://doi.org/10.1007/s10495-009-0446-3.

21. Chambon J-P, Soule J, Pomies P, Fort P, Sahuquet A, Alexandre D, et al. Tail regression in Ciona intestinalis: modulation by the TGF-beta and Notch pathways to promote wound healing and tissue regeneration. Dev Cell. 2009;17(2):279–90. https://doi.org/10.1016/j.devcel.2009.07.013.

22. Jeffery WR. Programmed cell death in the ascidian embryo: modulation by the TGF-beta and Notch pathways to promote wound healing and tissue regeneration. Cell. 2003;113(6):717–31. https://doi.org/10.1016/S0012-1606(03)00270-6.
47. Auger H, Sasakura Y, Joly J-S, Jeffery WR. Regeneration of oral siphon pigment organs in the ascidian Ciona intestinalis. Dev Biol. 2010;339(2):374–89. https://doi.org/10.1016/j.ydbio.2009.12.040.

48. Jeffery WR. Regeneration, stem cells, and aging in the tunicate Ciona: insights from the Oral siphon. Int Rev Cell Mol Biol. 2015;319:255–82. https://doi.org/10.1016/bs.ircmb.2015.06.005.

49. Jeffery WR. Progenitor targeting by adult stem cells in Ciona homeostasis, injury, and regeneration. Dev Biol. 2019;448(2):279–90. https://doi.org/10.1016/j.ydbio.2018.09.005.

50. Tiozzo S, Ballarin L, Burighel P, Zaniolo G. Programmed cell death in vegetative development: apoptosis during the colonial life cycle of the ascidian Botryllus schlosseri. Tissue Cell. 2006;38(3):193–201. https://doi.org/10.1016/j.tice.2006.02.003.

51. Franchi N, Ballin F, Manni L, Schiavon F, Basso G, Ballarin L. Recurrent phagocytosis-induced apoptosis in the cyclical generation change of the compound ascidian Botryllus schlosseri. Dev Comp Immunol. 2016;62:8–16. https://doi.org/10.1016/j.dci.2016.04.011.

52. Ballarin L, Schiavon F, Manni L. Natural apoptosis during the blastogenetic cycle of the colonial ascidian Botryllus schlosseri: a morphological analysis. Zool Sci. 2010;27(2):96–102. https://doi.org/10.2108/zsj.27.96.

53. Manni L, Zaniolo G, Cima F, Burighel P, Ballarin L. Botryllus schlosseri: a model ascidian for the study of asexual reproduction. Dev Dyn. 2007;236(2):335–52. https://doi.org/10.1002/dvdy.21037.

54. Tiozzo S, Ballarin L, Burighel P, Zaniolo G. Programmed cell death in vegetative development: apoptosis during the colonial life cycle of the ascidian Botryllus schlosseri. Tissue Cell. 2006;38(3):193–201. https://doi.org/10.1016/j.tice.2006.02.003.

55. Franchi N, Ballin F, Manni L, Schiavon F, Basso G, Ballarin L. Recurrent phagocytosis-induced apoptosis in the cyclical generation change of the compound ascidian Botryllus schlosseri. Dev Comp Immunol. 2016;62:8–16. https://doi.org/10.1016/j.dci.2016.04.011.

56. Tiozzo S, Ballarin L, Burighel P, Zaniolo G. Programmed cell death in vegetative development: apoptosis during the colonial life cycle of the ascidian Botryllus schlosseri. Tissue Cell. 2006;38(3):193–201. https://doi.org/10.1016/j.tice.2006.02.003.

57. Franchi N, Ballin F, Manni L, Schiavon F, Basso G, Ballarin L. Recurrent phagocytosis-induced apoptosis in the cyclical generation change of the compound ascidian Botryllus schlosseri. Dev Comp Immunol. 2016;62:8–16. https://doi.org/10.1016/j.dci.2016.04.011.

58. Tiozzo S, Ballarin L, Burighel P, Zaniolo G. Programmed cell death in vegetative development: apoptosis during the colonial life cycle of the ascidian Botryllus schlosseri. Tissue Cell. 2006;38(3):193–201. https://doi.org/10.1016/j.tice.2006.02.003.

59. Franchi N, Ballin F, Manni L, Schiavon F, Basso G, Ballarin L. Recurrent phagocytosis-induced apoptosis in the cyclical generation change of the compound ascidian Botryllus schlosseri. Dev Comp Immunol. 2016;62:8–16. https://doi.org/10.1016/j.dci.2016.04.011.

60. Tiozzo S, Ballarin L, Burighel P, Zaniolo G. Programmed cell death in vegetative development: apoptosis during the colonial life cycle of the ascidian Botryllus schlosseri. Tissue Cell. 2006;38(3):193–201. https://doi.org/10.1016/j.tice.2006.02.003.