Functional analysis of thyroid hormone receptor beta in *Xenopus tropicalis*
founders using CRISPR-Cas

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SUMMARY STATEMENT

Using knocked-out Xenopus tropicalis founders generated by CRISPR-Cas, TRβ was shown to have significant effects on developmental timing but is not required for completion of metamorphosis.

ABSTRACT

Amphibians provide an ideal model to study the actions of thyroid hormone (TH) in animal development because TH signaling via two TH receptors, TRα and TRβ, is indispensable for amphibian metamorphosis. However, specific roles for the TRβ isoform in metamorphosis are poorly understood. To address this issue, we generated trβ-disrupted Xenopus tropicalis tadpoles using the CRISPR-Cas system. We first established a highly efficient and rapid workflow for gene disruption in the founder generation (F0) by injecting sgRNA and Cas9 ribonucleoprotein. Most embryos showed severe mutant phenotypes carrying high somatic mutation rates. Utilizing this founder analysis system, we examined the role of trβ in metamorphosis. trβ-disrupted pre-metamorphic tadpoles exhibited mixed responsiveness to exogenous TH. Specifically, gill resorption and activation of several TH-response genes, including trβ itself and two protease genes, were impaired. On the other hand, hind limb outgrowth and induction of the TH-response genes, klf9 and fra-2, were not affected by loss of trβ. Surprisingly, trβ-disrupted tadpoles were able to undergo spontaneous metamorphosis normally, except for a slight delay in tail resorption. These results indicate TRβ is not required but contributes to the timing of resorptive events of metamorphosis.
INTRODUCTION

Thyroid hormone (TH), acting through TH receptors (TRs), plays critical roles in various biological processes, including development, growth, and metabolism (Yen, 2001; Cheng et al., 2010; Waung et al., 2012; Brent, 2012; Ortiga-Carvalho et al., 2014; Mullur et al., 2014). TRs belong to the nuclear receptor superfamily, and two types of TRs, TR alpha (TRα) and TR beta (TRβ), are coded in separate gene loci in vertebrates (Wahli and Martinez, 1991; Tsai and O'Malley, 1994). TRs are ligand-dependent transcription factors that heterodimerize with the retinoid X receptor (RXR), and the heterodimers bind to thyroid hormone response elements (TREs) in the promoter or enhancer of TH-response genes to regulate their expression (Perlmann et al., 1993; Wu et al., 2001). TRs recruit co-repressors in the absence of TH and co-activators in the presence of TH to TREs to repress and activate target gene expression, respectively, as described by the ‘dual function model’ for the role of TRs during development (Sachs and Shi, 2000; Shi, 2009; Grimaldi et al., 2013).

Amphibian metamorphosis involves unique and dynamic morphological and physiological changes from larva to adult: resorption of gills and tail; remodeling of skin and intestinal tissues; and growth of the limbs (Tata, 1999; Furlow and Neff, 2006; Brown and Cai, 2007). The functions and mechanisms of action of TH and TRs in metamorphosis have been well-studied in various amphibians over several decades. In particular, *Xenopus laevis* and *Xenopus tropicalis* provide excellent models for TH-dependent metamorphosis (Buchholz, 2017). Two TR coding genes, *tra* and *trβ*, are present in the *Xenopus* genome, as in other vertebrates (Yaoita and Brown, 1990). *tra* is widely expressed before metamorphosis, and the level of its expression is maintained
until the end of metamorphosis (Yaoita and Brown, 1990; Wang et al., 2008). In contrast, trβ mRNA is also widely expressed but at much lower levels during pre- and pro-metamorphosis (Yaoita and Brown, 1990; Wang et al., 2008). After TH secretion by the thyroid gland commences (stage 54) (Nieuwkoop and Faber, 1994; Leloup and Buscaglia, 1977), trβ expression levels increase strongly as the TH concentration rises in the plasma (Yaoita and Brown, 1990) until both reach a peak at the climax of metamorphosis (stage 61–62), and then both TH and trβ levels attenuate toward the end of the metamorphosis. The expression of trβ follows the plasma profile of TH due to TREs in its promoter allowing for self-induction in the presence of TH (Tata, 1993; Ranjan et al., 1994; Wang et al., 2008)

Genome-editing techniques have greatly improved the exploration of the roles of TRs in metamorphosis (Choi et al., 2015; Wen and Shi, 2015; Choi et al., 2017; Wen et al., 2017) beyond what was possible using overexpression of dominant negative mutant TRs in transgenic animals (Schreiber et al., 2001; Buchholz et al., 2003). Using TALEN knock-out techniques in X. tropicalis, we and others have demonstrated that trα regulates the timing of amphibian metamorphosis (Choi et al., 2015, Wen and Shi, 2015, Choi et al., 2017, Wen et al., 2017). In general, TRα knock-out phenotypes were mild, such that homozygous TRα knock-out animals have apparently normal survival and fertility. On the other hand, due to its dramatic self-regulation, TRβ is thought to be the master regulator at the top of a gene regulation cascade required for metamorphosis and therefore is expected to play a crucial role in driving radical changes at the climax stage of frog metamorphosis. However, the details of its involvement and degree of functional overlap with TRα during metamorphosis are poorly understood.
The CRISPR-Cas system has several advantages for loss-of-function analysis compared with other genome-editing tools used for *X. tropicalis*, including high efficiency, convenience, and cost-effectiveness (Blitz et al., 2013; Nakayama et al., 2013; Shigeta et al., 2016; Naert et al., 2016). Thus, in this study, we first established a workflow to generate mosaic knock-out founders (crispants) carrying high rates of somatic mutations using injection of the sgRNA/Cas9 ribonucleoprotein (RNP). Next, we generated *trβ* crispants to examine the biological significance of *trβ* in amphibian metamorphosis. In agreement with previous pharmacological studies (Furlow et al., 2004; Denver et al., 2009), exogenous TH did not induce gill resorption in *trβ* crispants at the pre-metamorphic stage. Gene expression analysis indicated that some but not all TH-response genes lacked induction by TH in the crispants. Surprisingly, *trβ* crispants completed metamorphosis, though after a delay. Our results suggest that TRβ, like TRα, is not required to complete metamorphosis but has metamorphic roles distinct from those previously determined for TRα.

RESULTS

**sgRNA/Cas9 ribonucleoprotein effectively produces knocked-out founders in *Xenopus tropicalis***.

In this study, we employed Cas9 protein instead of Cas9 mRNA for injection, to maximize gene disruption efficiency, and we established an effective and convenient protocol for analysis of loss-of-function in *X. tropicalis* founders (Fig. S1). As proof of
concept, we targeted the melanin synthesis-related genes for tyrosinase (tyr) and oculocutaneous albinism 2 (oca2) as model genes. tyr sgRNA was designed within the exon region, whereas oca2 sgRNA was designed across the exon-intron junction (Fig. S2A). Each sgRNA was synthesized by T7 RNA polymerase-based in vitro transcription using PCR templates. sgRNA and Cas9 protein were incubated in vitro to form the RNP complex in 150 mM KCl and 20 mM HEPES, according to previous reports (Burger et al., 2016) The RNPs were then injected into X. tropicalis one-cell-stage fertilized eggs. As expected, injection of tyr or oca2 sgRNA/Cas9 RNP caused albinism phenotypes in almost all cases (Fig. S2B). About 70% of these crispants showed severe complete loss of pigmentation, while the proportion with developmental defects was comparable to uninjected controls, about 15-30% (Fig. S2C). Surprisingly, amplicon sequencing analysis of these on-target sites revealed that somatic mutation rates reached up to 100% in all crispants analyzed, where no wild-type allele reads could be identified (Fig. S2D, E). According to our amplicon sequencing data, we actually detected only 5 to 15 types of mutant alleles and no wild-type allele per animal in tyr and oca2 crispants exhibiting severe albino phenotypes, which suggests that mutagenesis by Cas9 RNP occurred at a very early developmental stage in these animals. Therefore, it seems that the 2,000 reads were saturated, suggesting that most if not all cells of an individual were mutant. Frameshift mutation rates were 89% in tyr crispants which is well above the expected 66% likely because individuals with severe mutations were chosen for sequencing. These results demonstrate that this Cas9 RNP-based protocol enables production of mosaic knocked-out founders (crispants; Burger et al., 2016) with high frequency and low toxicity.
Generation of trβ crispants in X. tropicalis

To investigate the function of TRβ in X. tropicalis metamorphosis, we disrupted the trβ gene using this CRISPR-based system to generate mutant founders. Two sgRNAs were designed in the coding region of the 2nd exon and across the 2nd exon-2nd intron junction to produce frameshift mutations and splicing errors, respectively (Fig. 1A). We injected Cas9 RNP with each of the two sgRNAs, or with mixtures of the two, into one-cell-stage fertilized eggs. On the following day, normally developed embryos were collected and a sub-sample selected for genotyping to examine the efficiency of disruption at the on-target sites. In a heteroduplex mobility assay (HMA), heteroduplex formation was detected in all trβ crispants, indicating that disruption of the target sites occurred in the trβ gene (Fig. 1B). As expected, multiple shorter PCR bands were observed in all embryos injected with the two sgRNAs/Cas9, consistent with these bands being small deletions between the two sgRNA-targeting sites. Injection of both sgRNA/Cas9 RNPs probably induced frameshift mutations and/or small deletions at the exon-intron boundary causing splicing errors (Radev et al., 2015). To examine the possible splicing errors in more detail, we analyzed trβ mRNA from whole bodies of stage 61/62 crispants by RT-PCR using primers designed in the 1st and 3rd exons. In trβ crispants injected with both sgRNAs, 2nd exon deleted bands (101 bp shorter) as well as other non-wild-type bands were present, indicating numerous perturbations of trβ mRNA in the metamorphic climax stage tadpoles (Fig. 1C). We sequenced eight metamorphosed crispants, which showed high somatic mutation rates (48-94%) (Fig. 2).
Limb development but not gill resorption was induced by T3 in pre-metamorphic trβ crispants

No differences in growth and development were observed in embryogenesis and pre-metamorphic development in trβ crispants versus controls, as expected because the expression level of trβ is very low during embryonic and pre-metamorphic stages (Yaoita and Brown, 1990; Wang et al., 2008). To investigate responsiveness to exogenous TH in pre-metamorphic trβ crispants, we treated stage 52–54 tadpoles with or without 10 nM 3,3,5-triiodo-L-thyronine (T3) for 3 d and examined external morphology. As expected, all uninjected control tadpoles responded to the exogenous T3 and exhibited characteristic gill resorption (n = 44/44) (Fig. 3A). In contrast, almost all of the trβ crispants were resistant to T3 treatment and showed no, or slight, morphological changes in the gills (n = 51/54). Gill resorption equivalent to that in the controls was observed in only 5.5% of crispants, which may represent unintentionally uninjected embryos (n = 3/54). However, hind limbs responded to T3 and precociously developed in both uninjected controls and trβ crispants (Fig. 3B). In the absence of T3, gill and limb morphology were unchanged in both groups of tadpoles. These results suggest that the gills exhibit high responsiveness to TH via TRβ-mediated signaling during T3-induced metamorphosis, but the hind limbs do not.

Impaired induction of TH-response genes in trβ crispants

Next, we extracted total RNA from whole bodies of T3-treated and untreated tadpoles and analyzed the induction of the well-known TH-response genes, trβ, krüppel-like
factor 9 (klf9), fos-related antigen-2 (fra-2), matrix metallopeptidase 13 (mmp13), and fibroblast activation protein alpha (fapa) using quantitative RT-PCR. Surprisingly, not all direct TH response genes responded in the same way to loss of trβ (Fig. 4). Two of the direct response genes, key transcription factors related to metamorphosis, klf9 and fra-2, were normally induced by T3 in both uninjected control and trβ crispants. On the other hand, mRNA levels of trβ itself, which is autoregulated by TH, and the two protease genes, mmp13 and fapa, were significantly lower in trβ crispants than in control tadpoles. Therefore, some but not all TH direct response genes require TRβ for activation. The basal expression level of trβ mRNA in the crispants were significantly lower than those of the uninjected controls in the absence of T3 (Fig. 4, S3), possibly due to disruption of trβ function mediated by splicing error and nonsense-mediated mRNA decay (NMD; Popp and Maquat, 2016).

trβ crispants accomplish normal metamorphosis, although slightly delayed

Based on our results from exogenous TH treatments, we expected that trβ crispants would fail to complete metamorphosis, or significantly extend the metamorphic period. To examine the effect of trβ disruption on natural metamorphosis, we observed morphological changes and measured the duration from fertilization to stage 61 immediately after gill resorption and prior to tail resorption, and to stage 66 when tail resorption is complete. There was no statistically significant difference in time from stage 1 to stage 61 between trβ crispants and control tadpoles from the same clutch (Fig. 5A). Unlike in TH-induced metamorphosis, gill resorption occurred normally in trβ crispants at the same time as in controls. However, we found significant differences in
the length of the period from stage 61 to stage 66 (Fig. 5B, C) because tail resorption
was slower in trβ crispants than in uninjected controls. Thus, trβ crispants took longer
to accomplish metamorphosis completely. Metamorphosed trβ crispants started feeding
normally, and the histological structure of the intestine at about one month after
metamorphosis showed that intestinal remodeling was complete (Fig. S4). Interestingly,
tentacles, which are specifically larval organs, persisted in most trβ crispants one week
after metamorphosis but were resorbed completely within the next three weeks (Fig.
S5). Importantly, we mated adult F0 crispants and confirmed that F1 offspring carrying
heterozygous mutant alleles accomplished natural metamorphosis (Fig. 6).

DISCUSSION

We have developed an efficient strategy to achieve mosaic knock-out
founders (crispants) using sgRNA and Cas9 RNP in X. tropicalis. We demonstrated the
effectiveness of this strategy by targeting an early exon and a splice junction in two
melanin synthesis-related genes, tyr and oca2, with easily visible mutant phenotypes.
Under optimized conditions, most injected embryos showed severe (full) albinism with
low levels of mosaicism (<10%) and developmental defects (<15~30%). Significantly,
amplicon-sequencing analysis of the on-targets revealed that somatic mutation rates
were high in severe tyr and oca2 phenotypes, i.e., the wild-type allele reads were not, or
rarely, detected. In the case of tyr sgRNA targeting exon 1, out-of-frame mutation rate
was 89%, consistent with severe albinism in sequenced individuals. On the other hand,
oca2 sgRNA targeting the exon-intron junction induced indels adjacent to the splice
donor site. By targeting splice site, mutant oca2 mRNA likely exhibited exon skipping or intron retention to lose its function (Radev et al., 2015). Weak and moderate phenotypes were hardly observed in oca2 crispants compared to those of tyr potentially because splice targeting mutations have higher frequency of inducing null mutations compared to 66% when targeting coding regions. Combining out-of-frame and splicing site perturbation in trβ caused by co-injection of two sgRNA/Cas9 RNPs is expected to result in a high frequency of loss of protein function. Therefore, our sgRNA/Cas9 RNP-based strategy may rapidly facilitate the examination of the functions of genes of interest in this species.

We used this Cas9 RNP gene disruption strategy to generate trβ mosaic knock-out founders (‘trβ crispants’) and investigated the role of trβ in metamorphosis. A major phenotype in trβ crispants was delayed resorption of some larval tissues (tail and tentacles). The delay of tail resorption and tentacles in trβ crispants may be due to decreased ability to induce metamorphosis-related proteases. Indeed, the activation of two protease genes, mmp13 and fapa by exogenous T3 was impaired in whole bodies of pre-metamorphic trβ crispants compared with wild-type tadpoles treated with T3. These proteases are well-known TH-response genes and are highly expressed during natural metamorphosis in sub-epidermal fibroblasts in the tail, notochord, notochord sheath, and gills, whose expression patterns are coincident with that of trβ (Brown et al., 1996; Berry et al., 1998a; Berry et al., 1998b; Das et al., 2006). These results suggest that TRβ may be the main regulator of TH-induced activation of these protease genes and larval tissue resorption, at least for the tail.
In the gill, resorption was not observed in response to exogenous T3 in pre-
metamorphic trβ crispants, even though it occurred normally in crispants during natural
metamorphosis. It is unclear why loss of trβ showed different phenotypes of gills in T3-
induced versus natural metamorphosis, though TRα expression levels may be sufficient
to compensate for lack of TRβ at the climax of metamorphosis but not during pre-
metamorphosis. Detailed analysis of the TH-regulated transcriptome, including
expression of various proteases in all organs of trβ crispants during T3-induced and
natural metamorphosis is needed for further understanding.

Although the crispants were mosaic including in-frame mutations, trβ
function is expected to be impaired at the organ and tissue level due to the 48-94%
disruption of trβ by amplicon sequencing. Indeed, we observed natural metamorphosis
in crispants, a phenotype confirmed by F1 bi-allelic mutants. Based on these results, we
suggest that additional phenotypes observed in crispants, but not yet examined in non-
mosaic trβ mutant animals, may accurately reflect the effect of loss of trβ. We observed
no effects on limb development and intestinal remodeling in trβ crispants. Exogenous
T3-induced precocious hind limb development occurred in trβ crispants as in wild-type
controls. Also, the transcription factors, klf9, and fra-2, are highly expressed in
developing and remodeling organs such as limb during natural metamorphosis (Wang
and Brown, 1993; Berry et al., 1998b; Das et al., 2006), and normal induction by
exogenous TH of these genes occurred in trβ crispants. These results are consistent with
expectations on the small if any requirement for TRβ in the limb based on the high TRα
to TRβ expression ratio in the limb and results from use of the TRα-selective antagonist
CO23 (Wen et al., 2017; Ocasio and Scanlan, 2006).
At the molecular level, surprising results were obtained regarding the effects of TRβ on TH-response gene expression. Two of the best characterized TH direct response genes are klf9 and trβ itself (Ranjan et al., 1994; Brown et al., 1996; Bagamasbad et al., 2008), but TH-induction of these genes gave contrasting results, namely only trβ and not klf9 induction by exogenous TH was impaired in trβ crispants. Interestingly, in TRα knock-out tadpoles, induction by exogenous TH of both klf9 and trβ was partially impaired (Choi et al., 2015; Wen and Shi, 2015). These results suggest that TRα is able to compensate for loss of TRβ with regard to klf9 but not trβ. Possibilities to explain impaired trβ induction in trβ crispants are that 1) the two TRs exhibit isoform specificity in molecular mechanisms in regulation of trβ, 2) tissue-specific expression of the TR isoforms exists such that trβ and little trα is expressed in tissues that exhibit trβ autoregulation (e.g. red blood cells and brain subventricular cells) (Denver et al., 2009) or 3) trβ mRNA level is decreased by nonsense-mediated mRNA decay (NMD). Similar possible explanations apply to the other TH response genes studied, fra-2, mmp13, and fapa. Further research is required to examine these intriguing possibilities. Although we only analyzed the perturbation of TH-response gene expression using total RNA from whole bodies in this study, various organ and tissue actually showed less responsiveness to TH in the trβ crispants; delay of gill, tail and tentacle resorption during T3-induced and natural metamorphosis. To understand the differences of responsiveness to TH mediated by trβ in each organ, we need to further analyze transcriptome changes in each organ.
Another surprising result from trβ crispants was that they developed normally and were able to complete metamorphosis through tail resorption, albeit with a delay. This result suggests that TRβ is not required for metamorphosis but plays a role in timing of developmental events already fated to occur as specified during development. At least three hypotheses explain why trβ crispants metamorphosed completely. First, TRα may partially compensate for TRβ deficiency so that the metamorphosis program could occur. Second, rather than TH-response gene induction by TRα, derepressed levels (i.e., above basal levels but below TH-induced levels) of TH response gene expression due to lack of TRβ-mediated repression, as detailed by the dual function model (Shi, 2009), may explain the ability to progress through tail resorption. These two hypotheses also explain the delay in metamorphosis in trβ crispants where reduced gene expression levels below that of full induction as occurs in wild-type tadpoles would increase the time required to progress through metamorphosis. Third, unknown mechanisms related to TH carried out metamorphosis (e.g., non-genomic pathway of TH). To address this question, we need to produce and analyze trα/trβ double knock-out frogs in the future.

In conclusion, we show that, despite dramatic contrasting effects on TH response gene expression observed in trβ crispants, TRβ has significant but mild effects on developmental timing that do not prevent complete metamorphosis from occurring. These results in combination with results from TRα knock-out animals provide evidence that TRα and TRβ may at least partly functionally compensate for each other but that TRα and TRβ have separate functions during metamorphosis. Full evaluation of
the role of TRs in amphibian development await analysis of TRα and TRβ double knock-out animals.

MATERIALS AND METHODS

Animal rearing and treatment

*Xenopus tropicalis*, the Golden strain, were provided by the Amphibian Research Center (Hiroshima University) through the National Bio-Resource Project of the AMED, Japan. Eggs and testes were collected from sexually mature adult frogs with an injection of human chorionic gonadotropin (Aska Pharmaceutical, Tokyo, Japan), and *in vitro* fertilization was carried out. To isolate testes, male frogs were treated with 1% MS-222 (tricaine; Sigma-Aldrich, MO, USA) as anesthesia and euthanasia. One-cell-stage embryos were de-jellied with 2% L-cysteine solution. After washing with 0.1 × Marc's modified ringer (MMR), embryos were microinjected in 6% Ficoll (Sigma-Aldrich) in 0.33 × MMR. At the blastula stage, embryos were moved to 0.1 × MMR. At 16–20 h after fertilization, normally developed embryos were collected and counted and represented the initial numbers of individuals in each experiment. Embryos and tadpoles were reared at 25–26 °C. Tadpoles at stage 52–54 were treated with 10 nM 3,3,5-triiodo-L-thyronine (T3; Sigma-Aldrich) for 3 d. Animal rearing and treatments were performed and approved according to the Hiroshima University guidelines for the use and care of experimental animals.
Preparation and microinjection of sgRNA/Cas9 ribonucleoprotein

All sgRNA targeting sequences and oligonucleotide information are listed in Tables S1 and S2. The sgRNA targeting sequence of tyrosinase (tyr) was taken from Blitz et al. (2013), Nakayama et al. (2013). oculocutaneous albinism 2 (oca2) and thyroid hormone receptor beta (trβ) sgRNAs were designed using CRISPR-direct (Naito et al., 2015). tyr and one of the trβ sgRNAs were designed within the exon, whereas oca2 and the other trβ sgRNA was designed across the exon-intron junction to induce splicing errors (Radev et al., 2015). The annealed oligonucleotides for tyr were cloned into the BsmBI site of the pCS2P-gRNA vector, and subsequently amplified for in vitro transcription templates using KOD FX Neo (TOYOBO, Osaka, Japan) and primer sets (IVT-sgRNA-F and R), as described previously (Ota et al., 2014; Shigeta et al., 2016). The other sgRNA templates were assembled by a PCR-based strategy (Nakayama et al., 2014; Sakane et al., 2017). DNA templates were purified with a QIAquick PCR Purification Kit (Qiagen, Hilden, Germany), subsequently, sgRNAs were synthesized using a MEGAscript T7 Kit (Thermo Fisher Scientific, MA, USA) and purified using a MEGAclear Kit (Thermo Fisher Scientific). Before microinjection, recombinant Cas9 protein (Alt-R S.p. Cas9 Nuclease 3NLS; Integrated DNA Technologies, IA, USA) and sgRNA(s) were incubated in 150 mM KCl and 20 mM HEPES buffer to form ribonucleoprotein complexes at room temperature for 10 min. Mixtures of Cas9 1 ng protein and 200 pg of one of the two sgRNAs were injected into one-cell-stage X. tropicalis embryos using a Nanoject II (Drummond, PA, USA).
Preparation of genomic DNA

Uninjected control embryos and tyr-, oca2-, and trβ-sgRNA/Cas9 RNP-injected embryos were individually collected at after 1 d or 3 d of injection. trβ crispants and F1 offspring were individually collected at stage 66. Each genomic DNA was extracted from the whole embryos using a DNeasy Blood and Tissue Kit (Qiagen).

RNA extraction, RT-PCR, and quantitative PCR

For RT-PCR, individual tadpoles were collected at stage 61/62 (n = 6). For quantitative PCR, four or more tadpoles at stage 52-54 were pooled in each sample for each treatment with and without T3 (n = 3), and this experiment was performed three times independently. Tadpoles were homogenized in liquid nitrogen. Total RNA was extracted by RNAiso (TaKaRa, Shiga, Japan) and purified with Direct-zol™ RNA MiniPrep (Zymo Research, CA, USA). The same quantity of total RNA was reverse transcribed with ReverTra Ace qPCR RT Master Mix with a gDNA Remover kit (TOYOBO). RT-PCR was performed using KOD FX Neo (TOYOBO) and analyzed using a microchip electrophoresis system (DNA-500 reagent kit and MCE-202 MultiNA; Shimadzu, Kyoto, Japan). Quantitative PCR (qPCR) was carried out by using KOD SYBR qPCR Mix (TOYOBO) with a Step One real-time PCR system (Thermo Fisher Scientific). Three technical replicates were used per sample and three biological replicates were analyzed in each experiment. Target gene expression levels were normalized by the expression of the housekeeping gene rpl8. RT-PCR and qPCR primer sets are listed in Table S2. All primer sets were designed by Primer 3 (Untergasser et al., 2012)
**HMA and amplicon sequencing**

For HMA, PCR on genomic DNA was performed over 35 cycles using KOD FX Neo (TOYOBO), and PCR products were analyzed using a microchip electrophoresis system (DNA-500 reagent kit and MCE-202 MultiNA; Shimadzu) according to Shigeta et al (2016). An amplicon-sequencing library was prepared based on the Illumina ‘16S Metagenomic Sequencing Library Preparation’. For the first round of PCR, the target regions containing sgRNA targeting sites were amplified from individual genomic DNA of uninjected control embryos ($n = 5$), $t yr$ ($n = 10$), $oca2$ ($n = 5$), and $trβ$ ($n = 8$) crispants using a KAPA HiFi HS ReadyMix (NIPPON Genetics, Tokyo, Japan) with primer sets containing barcode and overhang adaptor sequences. Equal quantities of all PCR products were pooled and purified using a QIAquick PCR Purification Kit (Qiagen). The second round of PCR was performed to construct a sequence library using a Nextra XT index kit (Illumina, CA, USA). The final library was purified and sequenced on the Illumina MiSeq. Library construction and sequencing were performed by Macrogen Japan and Hokkaido System Science. All primers are listed in Table S2.

**Amplicon data analysis**

Sequence data were preprocessed by Scythe (v0.994) (https://github.com/vsbuffalo/scythe) and Sickle programs (Joshi and Fass, 2011, https://github.com/najoshi/sickle) to trim adaptor sequences and remove reads that were shorter than 36 bp. Then, trimmed paired-end reads were joined into single reads using fastq-join (version 1.1.2-537) (Aronesty, 2011,
https://github.com/ExpressionAnalysis/ea-utils). PCR and Illumina sequence error rates were accounted for using wild-type samples, and then wild-type reads and mutant reads for tyr, oca2, and trβ were counted using an in-house script in R (version 3.3.3).

**Histological analysis**

*trβ* crispants were genotyped by HMA using genomic DNA from the tips of digits. Whole body and intestine at about one month after metamorphosis were fixed in 95% ethanol. Isolated intestines were embedded in paraffin, sectioned at 5 μm and stained with hematoxylin and eosin (Choi et al., 2017).

**Statistical analysis**

Welch's *t*-test was used to qPCR analysis and duration from fertilization to adult. When the *p*-value is less than 0.05, we conclude that a significant difference exists.

**Data availability**

The sequencing data have been deposited to DDBJ Sequence Read Archive with the accession number of DRA006344.
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Author contributions


Competing interests

The authors declare no competing or financial interests.

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Fig. 1: trβ sgRNA/Cas9 RNP leads to trβ gene disruption in *X. tropicalis* embryos.

(A) Schematic illustration of the sgRNA targeting sequences on the 2nd exon and 2nd intron of the trβ gene in *X. tropicalis*. One of the sgRNAs was designed in the coding
region (trβ sgRNA-1) and the other was designed across the exon-intron junction (trβ sgRNA-2). Highlights in red and gray denote the protospacer adjacent motif (PAM) and the 20 bp target sequences of sgRNA, respectively. Exon and intron sequences are indicated by capital and small letters, respectively. (B) Genotyping by HMA in uninjected and injected embryos. PCR products encompassing sgRNA target sites were analyzed using a microchip electrophoresis system. Heteroduplex bands and multiple short bands are shown in the sgRNA/Cas9 RNP-injected embryos (crispants). Arrowheads: wild-type original bands; curly brackets: up-shifted heteroduplex bands; square brackets: deleted bands. (C) RT-PCR was performed using total RNA from stage 61/62 tadpoles with primers designed upstream and downstream of the 2\textsuperscript{nd} exon. In addition to the original band (arrowhead), about 101 bp deleted bands were detected in crispants caused by skipping of the 2\textsuperscript{nd} exon (asterisk) in trβ crispants. Heteroduplex bands were also detected in crispants (brackets). Each lane represents an individual embryo. Ui; uninjected control embryos.
Fig. 2: Amplicon sequencing analysis of on-target sites in trβ crispants

(A) Image of trβ crispants at stage 66. Genomic DNA was extracted from whole bodies of eight trβ crispants and the sgRNA target region was PCR-amplified using barcoded primers. PCR amplicons were subjected to amplicon sequencing according to Materials and Methods. (B) The results of wild-type and mutant allele reads in each trβ crispant. All sequenced reads were classified into three groups; wild-type reads, frameshift and/or splice donor site mutation reads, in-frame mutation reads.
Fig. 3: Morphological and gene expression changes induced by T3 treatment in \textit{trb} crispants

(A) Uninjected control tadpoles and \textit{trb} crispants at pre-metamorphic stage (stage 52–54) were treated with or without 10 nM T3 for 3 d and morphological changes
observed. After 3 d of T3 treatment, regression of gills was observed in all uninjected control tadpoles \((n = 44/44)\), whereas these morphological changes were not induced in most of the \(tr\beta\) crispants \((n = 51/54)\). In contrast, morphological change did not occur in the gills of both uninjected control tadpoles and \(tr\beta\) crispants without T3 treatment.

(B) Hind limb development was induced by T3 treatment in all uninjected control tadpoles \((n = 44/44)\) and \(tr\beta\) crispants \((n = 54/54)\). \(N\) values represent the total numbers of tadpoles in three independent experiments.
Fig. 4: Impaired induction of TH-response genes in \( tr\beta \) crispants

mRNA expression of five TH response genes, *thyroid hormone receptor \( \beta \) (\( tr\beta \)), *krüppel-like factor 9* (\( klf9 \)), *fos-related antigen-2* (\( fra-2 \)), *matrix metallopeptidase 13* (\( mmp13 \)), and *fibroblast activation protein alpha* (\( fapa \)), was analyzed by qPCR.

Uninjected control tadpoles and \( tr\beta \) crispants at pre-metamorphic stage (stage 52–54) were treated with or without 10 nM T3 for 3 d and at least four tadpoles were pooled per sample (\( n = 3 \)) in three independent experiments. The expression levels were normalized by \( rpl8 \). Expression changes by T3 induction were calculated relative to those of control animals without T3 treatment. Error bars represent SD (\( n = 9 \)). The three independent experiments exhibited similar trends. Asterisks indicate significant differences between the uninjected control and \( tr\beta \) crispants: *, \( P < 0.005 \); Welch’s t-test.
Fig. 5: Delay of tail resorption in trβ crispants during natural metamorphosis.

(A) The days from fertilization to stage 61 are indicated in a box plot for uninjected control tadpoles (n = 46) and trβ crispants (n = 39). There were no significant differences. (B) The days from stage 61 to stage 66 are indicated in a box plot for uninjected control tadpoles (n = 45) and trβ crispants (n = 38). trβ crispants required significantly more days to reach stage 66 (*; Welch’s t-test, P < 0.05). N values represent the merged total numbers of tadpoles from two independent experiments. (C) Time course of tail resorption in uninjected control tadpoles and trβ crispants.

Uninjected control tadpoles and trβ crispants of the same age were selected from the same clutch at stage 61 and gill resorption observed at 1 d, 3 d, and 5 d after stage 61 (day 0). Uninjected control tadpoles completed tail resorption by day 5. In contrast, the tail remained in trβ crispants on day 5 (an arrowhead in the high magnification image) and a longer time was required to reach the end of metamorphosis.
Fig. 6: F1 trβ mutants accomplished natural metamorphosis

(A) Image of F1 offspring produced by mating of the F0 crispants. (B) Mutation sequences from each F1 offspring. PCR products of trβ target site from six F1 froglets were subcloned and sequenced. Sequences highlighted in red and gray denote protospacer adjacent motif (PAM) and protospacer sequence, respectively. Splice donor site is underlined. The deleted and inserted nucleotides are shown by dashes and blue letters, respectively. Capital and small letters indicate exon and intron, respectively.
Design

Design sgRNA in 5' exon or exon-intron junction.

sgRNA preparation & microinjection

- T7 promoter
- Target-specific sequence
- sgRNA scaffold

PCR

in vitro transcription

sgRNA

RNP complex

Cas9 protein

Microinjection

Analysis

Crispants

(=F0 injected animals)

Genotyping
- HMA
- Amplicon-sequence

Phenotyping
Fig. S1: A workflow for sgRNA/Cas9 RNP injection into *Xenopus tropicalis*

We established an efficient and rapid gene disruption workflow for the loss-of-function analysis using the CRISPR-Cas system in *X. tropicalis* founders. First, we design one or more sgRNAs targeting 5’ exons or exon-intron junctions to induce frameshift mutations and/or splicing errors. Next, sgRNAs are synthesized *in vitro* by T7 RNA polymerase using templates produced by PCR and assembled with Cas9 protein to form an RNP complex. sgRNA/Cas9 RNP is injected into one-cell-stage fertilized eggs, and the developing embryos (crispants) are genotyped by HMA and amplicon-sequencing followed by phenotyping.
A

B

C

D

E

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<tr>
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<th>oca2 (n = 5)</th>
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<td>0 (0 %)</td>
</tr>
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<td>mutant reads (%)</td>
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<tr>
<td>total reads</td>
<td>24936</td>
<td>13309</td>
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Fig. S2: Phenotyping and genotyping of embryos with disrupted melanin synthesis-related genes

(A) Schematic illustration of the gene structure and sgRNA targeting sequences of tyrosinase (tyr) and oculocutaneous albinism 2 (oca2) genes. Coding regions are shown as black boxes. Sequences highlighted in red and gray denote protospacer adjacent motif (PAM) and the 20 bp target sequence of sgRNA, respectively. Capital letters: exon; small letters: intron. (B) Representative severe phenotypes of tyr and oca2 crispants. (C) Phenotypes were classified into four groups: severe, near complete loss of pigmentation in retinal pigmented epithelium; moderate, more than half of pigmentation lost; weak, less than half of pigmentation lost; and normal, no alteration in pigmentation (Shigeta et al., 2016; Sakane et al., 2017). (D) Genotyping of crispants with HMA. Target regions were amplified with 35 cycles of PCR. Genomic PCR products were analyzed using microchip electrophoresis. Arrowheads and brackets indicate homoduplex and heteroduplex bands, respectively. Ui: uninjected control embryos. (E) The numbers and percentages of wild-type and mutant allele reads in each crispant. Reads obtained from next-generation sequencing were classified into each gene and individual by their specific barcode sequences. According to sequence data from uninjected control embryos, the rates of artificial backgrounds such as PCR or sequencing errors were calculated, and we discounted reads below these artificial background rates. Significant reads in ten tyr and five oca2 crispants were counted as total reads and somatic mutation rates were calculated.
Fig. S3: mRNA expression of trβ in the absent of T3

The result of trβ expression in control and crispants without T3. The scale of Y axis is enlarged in Fig. 4. *, P < 0.005; Welch’s t-test.
**Fig. S4: Histological analysis of intestine**

Intestines of uninjected control froglets and *trβ* crispants about one month after metamorphosis were processed for histology and stained with hematoxylin and eosin. Intestinal folds (IF), which develop only in the adult organ, were observed in both control froglets and crispants. Bar: 50 μm.
Fig. S5: Tentacle resorption was delayed in trβ crispants

About one week after metamorphosis, tentacles remained near the mouth of trβ crispants (arrowheads). Images are representative of six uninjected control froglets and eleven trβ crispants. The number of froglets with remaining tentacle(s) is given in the images.
### Supplementary information table

#### Table S1: Sequences of sgRNA targeting sites.

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<th>sgRNA name</th>
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<tr>
<td>oca2</td>
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<tr>
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</tr>
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<td>trβ-2</td>
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PAM sequence is underlined.

#### Table S2: Oligonucleotide information in this study.

**Primers used for HMA.**

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**Oligonucleotides used for PCR assembly for sgRNA template.**

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| trβ-1      | TAATACGACTCTATAGGAGGCTATGGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGCTAGC
Oligonucleotides used for constructing sgRNA vector.

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<th>Sequence (5' to 3')</th>
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| **tyr**    | Sense: TATAGGAACTGCCCCCTGCAAACAG  
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Primers for sgRNA in vitro transcription template.

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Primers for the amplicon sequence.

The overhang adapter and barcode sequences are added to the locus-specific primer. Barcode sequences are indicated by small letters.

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                      | GAAAGGAAACATG |
| tyr-Wt-ampliconseq1-R | GTCTCGTGGGCTCGGAAGATGTGTATAAGAGACAGttaaCAGAGCTG  
                      | GTCAGGACACATC |
| tyr-Wt-ampliconseq2-F | TCGTCGGCAGCGTGTCAGATGTGTATAAGAGACAGaaacgAGCAGCATG  
                      | GAAAGGAAACATG |
| tyr-Wt-ampliconseq2-R | GTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGatcgcCAGAGCTG  
                      | GTCAGGACACATC |
| tyr-Wt-ampliconseq3-F | TCGTCGGCAGCGTGTCAGATGTGTATAAGAGACAGggcgAGCAGCATG  
                      | GAAAGGAAACATG |
| tyr-Wt-ampliconseq3-R | GTCTCGTGGGCTCGGAGATGTGTATAAGAGACAGgtcgcCAGAGCTG  
                      | GTCAGGACACATC |
| tyr-Wt-ampliconseq4-F | TCGTCGGCAGCGTGTCAGATGTGTATAAGAGACAGgcgcAGCAGCATG  
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<td>Primer set</td>
<td>Forward primer sequence (5’ to 3’)</td>
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Primer sequences are shown in lowercase. The forward primer sequences correspond to the 5’ end, and the reverse primer sequences correspond to the 3’ end of the amplicon.
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