



MAKÜ FEBED  
ISSN Online: 1309-2243  
<http://febed.mehmetakif.edu.tr>

*Mehmet Akif Ersoy Üniversitesi Fen Bilimleri Enstitüsü Dergisi* 5 (2): 1-4 (2014)

Research Paper / Araştırma Makalesi

## Telomerase Activity in Different Tissues of Zebrafish (*Danio rerio*)

Ayşe Gul Mutlu<sup>\*</sup>, Hulya Yıldız, Abdülkerim Bilginer

Mehmet Akif Ersoy University, Faculty of Arts and Sciences, Department of Biology, Burdur, Turkey

Geliş Tarihi (Received): 21.07.2014, Kabul Tarihi (Accepted): 15.10.2014

✉ Yazışmalardan Sorumlu Yazar (Corresponding author): [agmutlu@gmail.com](mailto:agmutlu@gmail.com) (A.G. Mutlu)

☎ +90 248 213 3035 📠 +90 248 213 30 99

### ABSTRACT

Zebrafish (*Danio rerio*) is a powerful model for aging, cancer, developmental biology and some aging-related diseases. It is critical to investigate complementary vertebrate models to understand telomere and telomerase function in biological systems. One of the best model organisms for telomerase studies is zebrafish. Zebrafish have different telomerase activity in different tissues. These differences may depend on the tissue's function or regeneration capacity of organs. The aim of this study is to compare the telomerase activity of liver, heart and spleen tissues in juvenile and adult zebrafish. RTA (relative telomerase activity) values of all tissues were different from each other statistically in the adult group (6 months of age). In the juvenile group (2 months of age) there is no difference between heart and spleen values but the other groups are statistically different from each other.

**Key Words:** Zebrafish (*Danio rerio*), telomerase, spleen, heart, liver

### Zebrabalığının (*Danio rerio*) Farklı Dokularında Telomeraz Aktivitesi

### ÖZET

Zebra balığı (*Danio rerio*) yaşlanma, kanser, gelişim biyolojisi ve bazı yaşlanma ile ilgili hastalıklar için önemli bir model organizmadır. Biyolojik sistemlerde telomer ve telomeraz fonksiyonunu anlamak için bu tür omurgalı modellerini araştırmak kritik öneme sahiptir. Zebra balığı, telomeraz çalışmaları için en iyi organizmalarından biri olarak kabul edilir. Zebra balığının farklı dokularındaki telomeraz aktiviteleri de farklıdır. Bu farklılıklar, dokuların işlevine ya da organların yenilenme kapasitesine bağlı olabilir. Bu çalışmanın amacı, yavru ve yetişkin Zebra balıklarının karaciğer, kalp ve dalak dokularında telomeraz aktivitelerini karşılaştırmaktır. Yetişkin grupta (6 aylık) tüm dokuların RTA (relatif telomeraz aktivitesi) değerleri istatistiksel olarak birbirinden farklı bulunmuştur. Yavru grubun (2 aylık) kalp ve dalak değerleri arasında bir farklılık gözlenmemiştir fakat bu dokuların telomeraz aktivitesi karaciğer dokusundan istatistiksel olarak farklı bulunmuştur.

**Anahtar Kelimeler:** Zebra balığı (*Danio rerio*), telomeraz, dalak, kalp, karaciğer

## INTRODUCTION

Telomeres are specialized functional complexes that protect the ends of eukaryotic chromosomes. The essential telomeric DNA sequences at each end of the eukaryotic linear chromosomes are, in most species, tandem repeats of a short sequence unit (Blackburn, 2001). Greider and Blackburn identified a telomere terminal transferase involved in the addition of telomeric repeats necessary for the replication of chromosome ends in eukaryotes (Greider and Blackburn, 1985). Telomerase adds multiple copies of this DNA unit to the terminal portion of one strand of the repeat tract (Blackburn, 2001).

Telomerase is a very important enzyme for aging process and carcinogenesis. Primary human cells exhibited limited replicative potential but the cancer lines divided indefinitely with passage in culture (Artandi and DePinho, 2010). To grow indefinitely, human cancer cells must counteract the progressive loss of telomeric DNA that universally accompanies cell division (Shay et al., 2012). This immortality is a result of telomerase activity mainly. Telomerase is expressed in more than 85% of cancer cells (Buseman et al., 2012), but the telomere length can be maintained in the absence of telomerase. It was deduced that one or more alternative telomerase-independent mechanisms exist in human cells (Shay et al., 2012).

Telomerase shortening may cause aging and death. Some evidence suggests that the progressive loss of telomeric repeats of chromosomes may function as an important timing mechanism during the aging process. Numerous epidemiological studies show that shorter telomeres in humans are associated with many age related diseases (Boccardi and Paolisso, 2014). Telomerase gene therapy in adult and old mice delays aging and increases longevity (de Jesus et al., 2012; de Jesus and Blasco, 2013).

Zebrafish (*Danio rerio*) has been developed as a powerful model for aging, cancer, developmental biology and some aging-related diseases (Kishi et al., 2003). It is critical to investigate complementary vertebrate models to understand telomere and telomerase function in biological systems. Contrary to the inbred laboratory mouse, zebrafish have heterogeneous telomeres of human-like length and shorten with age. Like humans, telomerase expression in zebrafish somatic cells is not sufficient to prevent telomere shortening (Henriques et al., 2013).

Different tissues have different telomerase activity. Zebrafish also have different telomerase activity in different tissues. These differences may depend on the tissue's function or regeneration capacity of organs. There is little information about telomerase activity differences in zebrafish tissues in the literature. The aim

of this study is to compare the telomerase activity of liver, heart and spleen tissues in juvenile and adult zebrafish. This data will provide a source for zebrafish researchers who study cancer, aging and regeneration in particular.

## MATERIALS AND METHODS

### Maintenance of Zebrafish

Zebrafish (*Danio rerio*) were maintained at  $24 \pm 2^\circ\text{C}$  with a light/dark cycle of 14:10 hours and were fed dry flake food. The fish were anesthetized with ice before organs were excised. Thirty zebrafishes were used for this research as 15 adults and 15 juveniles. All zebrafish applications were approved by the Ethical Committee of the Mehmet Akif Ersoy University (93773921-65).

### Telomerase Assay

Telomerase activities were measured using the Roche TeloTAGGG Telomerase PCR ELISA kit according to manufacturer's instructions. This kit allows highly specific amplification of telomerase-mediated elongation products combined with non-radioactive detection following an ELISA protocol. Relative Telomerase Activity (RTA) values were calculated for mg/ml of protein. Protein values were determined by the Bradford method (Bradford Reagent SIGMA B 6916). Minitab Release 13.0 statistical software was used for analysis. The results were estimated with the Mann-Whitney Test.

## RESULTS

RTA values of all tissues were different from each other statistically in the Adult group (6 months of age) ( $p < 0.05$ ). In the Juvenile group (2 months of age) there is no difference between heart and spleen values but the other groups are statistically different from each other ( $p < 0.05$ ) (Table 1 and Figure 1).

Table 1. Relative Telomerase Activity (RTA)  $\pm$  standard error (SE) values of tissues.

Tissues	Relative Telomerase Activity (RTA) $\pm$ SE
Heart (6 months)	0.301 $\pm$ 0.058
Spleen (6 months)	1.122 $\pm$ 0.261
Liver (6 months)	0.131 $\pm$ 0.050
Heart (2 months)	1.132 $\pm$ 0.507
Spleen (2 months)	0.572 $\pm$ 0.206
Liver (2 months)	0.130 $\pm$ 0.032

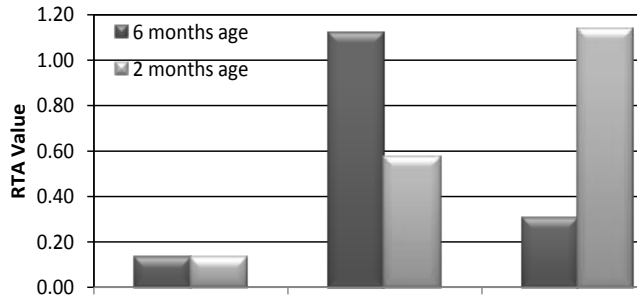


Figure 1. Comparison of the telomerase activity of different tissues

## DISCUSSION

Research about telomere and telomerase in vertebrates has become progressively more important due to the association of these two biological endpoints with cellular aging and cancer in humans (Downs et al., 2012). Activation of telomerase is proposed as an essential step in cancer cell immortalization and cancer progression (Lund et al., 2009). Telomerase is expressed in low levels in a variety of normal tissues in humans (Greider, 1998). Somatic stem and progenitor cells express relatively higher levels of telomerase. Telomere shortening limits the replicative life span of telomerase-negative cells, but telomerase activity in somatic stem cells likely contribute to the prolonged proliferative capacity of these cells compared to differentiated somatic cells (Günes and Rudolph, 2013). However, the inability to maintain the ends of the chromosomes in somatic cells contributes to the lack of true regenerative ability in human tissues (Lund et al., 2009).

Expression of telomerase in aquatic animals is likely not related to longevity but to their ability to regenerate injured tissue (Elmore et al., 2008). Although human organs have a limited ability to heal and regenerate damaged or lost tissue, the zebrafish retains remarkable regenerative abilities in most tissues. Moreover, the zebrafish has constitutively abundant telomerase activity in somatic tissues from embryos to adults (Anchelin et al., 2011). According to Lund and colleagues (2009), telomere lengths are similar in Zebrafish liver, heart, kidney, intestine and gill tissues (Lund et al., 2009). Telomerase activity present in the zebrafish organs may maintain the similar telomere lengths.

We detected highest telomerase activity in the adult spleen. Hematopoietic tissue is located in the stroma of the spleen and the interstitium of the kidney in Zebrafish (Menke et al., 2011). High telomerase activity in the spleen tissue may depend on the hematopoietic functions of the organ. Telomerase activity was higher in adult spleen tissue than juvenile. Anchelin and colleagues found that both telomerase expression and telomere length increased from embryo to adulthood

stages, but they drastically declined in aged fish despite the telomerase activity detected in different tissues of old fish (Anchelin et al., 2011).

The teleost liver plays an important role in the metabolic homeostasis of the body (Menke et al., 2011). Zebrafish have high regeneration capacity in the liver (Sadler et al., 2007; Kan et al., 2009). We measured very similar RTA values in juvenile and adult livers. However the zebrafish liver had the lowest mean RTA values compared to the spleen and the heart.

According to our research, both juvenile and adult fish have high telomerase activity in heart tissue. In another study, 20% of the heart tissue of zebrafish was surgically removed by Poss and colleagues (2002) and they observed full regeneration of the heart within 60 days. Some other studies also confirm the regeneration ability of the zebrafish heart (Kikuchi, 2014; Major and Poss, 2007). High telomerase activity in the zebrafish heart may be connected with the regeneration ability. Investigation of the regeneration ability of the zebrafish heart is important to cure some human heart diseases like myocardial infarction.

Telomerase enzyme is especially important for cancer, aging and regeneration researches. One of the best model organisms for telomerase studies is the zebrafish, but there is limited information in the literature for telomerase activity in different tissues of the zebrafish. Current research contributes to the scientific literature in that respect. This study is also important in terms of telomerase activity evaluated in adult and juvenile fish.

## ACKNOWLEDGEMENTS

This study was supported by the Mehmet Akif Ersoy University Scientific Research Projects Unit with a project number of 0202-YL-13.

## REFERENCES

- Anchelin, M., Murcia, L., Perez, F.A., Navarro, E.M.G., Cayuela, M.L. (2011) Behaviour of telomere and telomerase during aging and regeneration in Zebrafish. *PLOS One* 6(2): e16955.
- Artandi, S.E., Depinho, R.A. (2010) Telomeres and telomerase in cancer. *Carcinogenesis* 31(1): 9-18.
- Blackburn, E.H. (2001) Switching and signaling at the telomere. *Cell* 106: 661-673.
- Boccardi, V., Paolisso G. (2014) Telomerase activation: a potential key modulator for human healthspan and longevity. *Ageing Res Rev* 15: 1-5.
- Buseman, C.M., Wright, W.E., Shay, J.W. (2012) Is telomerase viable target in cancer. *Mutat. Res.*, 730 (1-2), 90-97.
- De Jesus, B.B., Vera, E., Schneeberger, K., Tejera, A.M., Ayuso, E., Bosch, F., Blasco, M.A. (2012)

- Telomerase gene therapy in adult and old mice delays aging and increases longevity without increasing cancer. *EMBO Mol Med* 4: 691-704.
- De Jesus, B.B., Blasco, M.A. (2013) Telomerase at the intersection of cancer and aging. *Trends Genet* 29(9): 513-520.
- Downs, P.D., Shen, Y., Pasquali, A., Beldorth, I., Savage, M., Gallier, K., Garcia, T., Booth, R.E., Walter, R.B. (2012) Characterization of telomeres and telomerase expression in *Xiphophorus*. *Comp Biochem Physiol C Toxicol Pharmacol* 155(1): 89–94.
- Elmore, L.W., Norris, M.W., Sircar, S., Bright, A.T., McChesney, P.A., Winn, R.N., Holt, S.E. (2008) Upregulation of telomerase function during tissue regeneration. *Exp Biol Med* (Maywood) 233(8): 958-967.
- Greider, C.W., Blackburn, E.H. (1985) Identification of a specific telomere terminal transferase activity in Tetrahymena extracts. *Cell* 43: 405-413.
- Greider, C.W. (1998) Telomerase activity, cell proliferation and cancer. *Proc. Natl. Acad. Sci. USA*, 95, 90-92.
- Günes, C., Rudolph, K.L. (2013) The role of telomeres in stem cells and cancer. *Cell* 152: 390-393.
- Henriques, C.M., Carneiro, M.C., Tenente, I.M., Jacinto, A., Ferreira, M.G. (2013) Telomerase is required for Zebrafish lifespan. *PLOS Genet* 9(1):e1003214.
- Kan, N. G., Junghans, D., Izpisua Belmonte, J. C. (2009) Compensatory growth mechanisms regulated by BMP and FGF signaling mediate liver regeneration in zebrafish after partial hepatectomy. *Faseb* 23: 3516-3525.
- Kikuchi, K. (2014) Advances in understanding the mechanism of zebrafish heart regeneration. *Stem Cell Research* 13: 542–555.
- Kishi, S., Uchiyama, J., Baughman, A.M., Goto, T., Lin, M.C., Tsai, S.B. (2003) The zebrafish as a vertebrate model of functional aging and very gradual senescence. *Experimental Gerontology* 38: 777-786.
- Lund, T.C., Glas, T.J., Tolar, J., Blazar, B.R. (2009) Expression of telomerase and telomere length are unaffected by either age or limb regeneration in Danio Rerio. *PLOS One* 4(11): e7688.
- Major, R.J., Poss, K. D. (2007) Zebrafish heart regeneration as a model for cardiac tissue repair. *Drug Discov Today Dis Models* 4(4): 219–225.
- Menke, A.L., Spitsbergen, J.M., Woterbeek, A.P.M., Woutersen, R.A. (2011) Normal anatomy and histology of the adult Zebrafish. *Toxicologic Pathology* 39: 759-775.
- Poss, K. D., Wilson, L. G., Keating, M. T. (2002) Heart regeneration in zebrafish. *Science* 298: 2188-2190.
- Sadler, K. C., Krahn, K. N., Gaur, N. A., Ukomadu, C. (2007) Liver growth in the embryo and during liver regeneration in zebrafish requires the cell cycle regulator. *Proc Natl Acad Sci USA* 104: 1570-1575.
- Shay, J.W., Reddel, R.R., Wright, W.E. (2012) Cancer and telomeres an alternative to telomerase. *Science* 336: 1388-1390.
-