



## ***sqet4Et* /+(AB);*sqet20Et* /+(AB) (CZRC catalog ID: CZ1379)**

### **Nature of the transgene**

The *sqet4Et ;sqet20Et* allele was generated by integration of an enhancer trap construct carries the EGFP reporter gene controlled by a partial epithelial promoter from the *keratin8* gene. This line express GFP in the, mantle cell and lateral line of the neuromast. In the ET20, the external mantle cells of the neuromast were labeled, whereas in the ET4 line, the expression of the reporter gene was restricted only to the internal hair cells. Crossing of these lines allowed visualizing of both cell types simultaneously, demonstrating the relative positions of these cells.

### **Genotyping assay**

Genotyping of the *sqet4Et ;sqet20Et* allele is based on the fluorescent microscope. As identified by fluorescent microscope, the GFP fluorescence signal is detectable at 72 hpf.

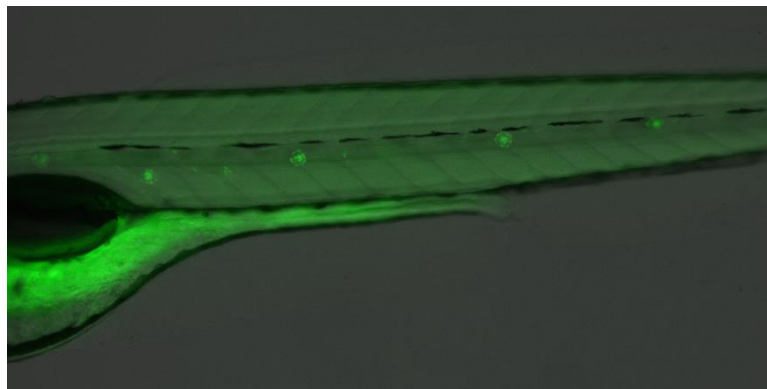


Figure. GFP expression in the lateral line at 5dpf in *sqet4Et ;sqet20Et* line.

The figure shows the lateral view of *sqet4Et ;sqet20Et* embryos at 5dpf.

### **Reference**





国家水生生物种质资源库国家斑马鱼资源中心  
China Zebrafish Resource Center (CZRC)  
National Aquatic Biological Resource Center (NABRC)

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Parinov, S., Kondrichin, I., Korzh, V., and Emelyanov, A. (2004) Tol2 transposon-mediated enhancer trap to identify developmentally regulated zebrafish genes in vivo. *Developmental dynamics* : an official publication of the American Association of Anatomists. 231(2):449-459

地址: 湖北省武汉市东湖南路7号中科院水生所 430072  
邮箱: zebrafish@ihb.ac.cn

电话: (027)68780570  
网址: www.zfish.cn



微信公众号