## **RESEARCH REPORTS**

**Biological** 

J.W. Kim<sup>1,4†</sup>, H. Choi<sup>2,4†</sup>, B.C. Jeong<sup>2,4</sup>, S.H. Oh<sup>2,4</sup>, S.W. Hur<sup>2,4</sup>, B.N. Lee<sup>1,4</sup>, S.H. Kim<sup>4</sup>, J.E. Nör<sup>3</sup>, J.T. Koh<sup>2,4\*</sup>, and Y.C. Hwang<sup>1,4\*</sup>

<sup>1</sup>Department of Conservative Dentistry, School of Dentistry, Dental Science Research Institute, Chonnam National University, Gwangju, Korea; 
<sup>2</sup>Department of Pharmacology and Dental Therapeutics, Dental Science Research Institute, School of Dentistry, Chonnam National University, Gwangju, Korea; 
<sup>3</sup>Angiogenesis Research Laboratory, Department of Cariology, Restorative Sciences, Endodontics, University of Michigan, School of Dentistry, Ann Arbor, MI 48109-1078, USA; 
<sup>4</sup>Research Center for Biomineralization Disorders, Chonnam National University, Gwangju, Korea; and 
<sup>†</sup>authors contributing equally as first authors; 
<sup>\*</sup>corresponding authors, jtkoh@chonnam.ac.kr, ychwang@chonnam.ac.kr

J Dent Res DOI: 10.1177/0022034514525199

## Transcriptional Factor ATF6 is Involved in Odontoblastic Differentiation

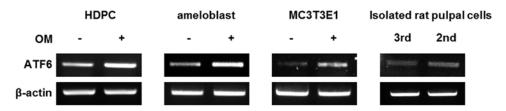
## **APPENDIX**

Appendix Table 1. Primer Sequences Used in RT-PCR of Rat Tooth Germs

Gene	Primer Sequence (5'-3')	Temperature (°C)	Product (bp)
ΑΤΕ6α	F: GGATTTGATGCCTTGGGAGT	55	192
	R: ATTTTTTCTTTGGAGTCAG		
DMP1	F: GGAGCAAGGTGACAGCGAGT	60	104
	R: GAGACTGGAGGCCTTCCTGG		
DSPP	F: GGGAAGGTGCTGGTTTGGAT	55	89
	R: TCCATCTCCTGCGTCTGCAC		
β-Actin	F: GCTGACAGGATGCAGAAGGA	55	124
	R: TGGACAGTGAGGCCAGGATA		

## Appendix Table 2. Primer Sequences Used in RT-PCR of Immortalized Human Dental Pulp Cells (HDPCs)

Gene	Primer Sequence (5'-3')	Temperature (°C)	Product (bp)
ΑΤΓ6α	F: TCCTCGGTCAGTGGACTCTTA	55	212
	R: CTTGGGCTGAATTGAAGGTTTTG		
DMP1	F: GCTAGCTGGTGGCTTCTCCA	62	522
	R: CAGCAATTGGCTGCCACCTG		
DSPP	F: CAGCCAAAGAATAGAGGAC	52	133
	R: GGGACCCTTGATTTCTAT		
β-Actin	F: ACCCACACTGTGCCCATCTAC	55	206
	R: GCCATCTCCTGCTCGAAGTC		



**Appendix Figure.** Expression of ATF6 during the differentiation process. HDPC, ameloblast, and MC3T3E1 cells were cultured with 50  $\mu$ g/mL ascorbic acid and 5 mM  $\beta$ -glycerophosphate ( $\beta$ -GP) and harvested for total RNA isolation. RT-PCR was performed. During the differentiation process in each cell type, ATF6 expression was increased.