Research Article

Osteoponin Promoter Controlled by DNA Methylation: Aberrant Methylation in Cloned Porcine Genome

Chih-Jie Shen,1,2 Yung-An Tsou,1,2,3 Hsiao-Ling Chen,4 Hung-Jin Huang,5 Shinn-Chih Wu,6 Winston T. K. Cheng,7 Calvin Yu-Chian Chen,2,8,9,10 and Chuan-Mu Chen1,11

1 Department of Life Sciences, National Chung Hsing University, Taichung 402, Taiwan
2 School of Medicine, College of Medicine, China Medical University, Taichung 40402, Taiwan
3 Department of Otolaryngology Head and Neck Surgery, China Medical University, Taichung 40402, Taiwan
4 Department of Bioresources, Da-Yeh University, Changhua 515, Taiwan
5 Department of Chinese Pharmaceutical Sciences and Chinese Medicine Resources, College of Pharmacy, China Medical University, Taichung 40402, Taiwan
6 Department of Animal Science and Technology, National Taiwan University, Taipei 106, Taiwan
7 Department of Animal Science and Biotechnology, Tung Hai University, Taichung 407, Taiwan
8 Department of Biomedical Informatics, Asia University, Taichung 41354, Taiwan
9 Research Center for Chinese Medicine & Acupuncture, China Medical University, Taichung 40402, Taiwan
10 Rong Hsing Research Center for Translational Medicine and the iEGG Center, National Chung Hsing University, Taichung 402, Taiwan

Correspondence should be addressed to Calvin Yu-Chian Chen; ycc929@mit.edu and Chuan-Mu Chen; cmchen7010978@gmail.com

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1. Introduction

Nowadays, many of pathogenesis of diseases have been determined [1–3]. Methylation in the 5’ cytosine in the CpG dinucleotides is crucial a mechanism that regulates gene expression without changing DNA sequence and can be inherited to the offspring [4]. The promoter region contains various transcription factor binding motifs with numerous CpG dinucleotides. Some transcription factors are blocked by methylated CpG island resulting in inhibition of gene expression [5]. Somatic cell nuclear transfer (SCNT) technique is used to generate an identical genetic background offspring [6, 7]. However, SCNT cloning animals usually showed low survival rate and inappropriate methylation reprogramming.
process [8]. This dilemma of SCNT animal may be caused by methylation controlled genes, such as imprinting genes [9].

OPN is an extracellular matrix protein and hydrophilic glycoprotein identified firstly in the bone as a sialoprotein. It contains a thrombin and transglutaminase cutting site, and the molecular weight is about 25 kDa to 75 kDa; in pig, the molecular is about 67 kDa; it contains numerous isoforms [10]. OPN has a hydrophobic N terminal; thus, it can be secreted out of cell membrane; the amino sequence of OPN is full of Asp, Thr, and Ser that can elevate the binding activity with calcium, glycosylation, and phosphorylation, respectively [11]. Thus, OPN plays numerous roles in many aspects, such as bone remodeling, cell migration, INOS regulation, repairment, and leucocyte recruitment [12]. And acquired OPN expression has been found in a variety of cancer cell types, especially in the liver, lung, breast, colon, brain, and spleen [13, 14].

Thus, OPN plays a crucial role in the regulation of many essential physiological processes such as bone remodeling, cell migration, iNOS regulation, and retardation of growth of bones. Therefore, this study focuses on the methylation change of OPN promoter that may be disrupted by inappropriate reprogramming process. Consequently, aberrant methylation of promoter could lead to aberrant expression of OPN. In the previous studies, OPN expression was induced with TSA (trichostatin A) in mouse undifferentiated mesenchymal cell line by API site [22]. The TSA is a histone deacetylase inhibitor. It can lose the chromatin structure in order to let gene restore its expression. 5-aza-dc is also an analog with the same structure of cytosine without methyl group adding in the 5’ C end [23].

Thus, 5-aza-dc addition leads to low methylation percentage in the CpG sites rich region. The hypomethylation status in the promoter may contribute its gene transcription activity. Porcine fetal fibroblasts in 5th passage cultures were treated with 0.5, 1.0, 2.0, and 3.0 μM 5-aza-dc for 96 h; 5-aza-dc inhibited the growth of cell at all concentrations. 5-aza-dc induced a reduction of transcripts level in OPN promoter that may be disrupted by inappropriate reprogramming process. Consequently, aberrant methylation of promoter could lead to aberrant expression of OPN. In the previous studies, OPN expression was induced with TSA (trichostatin A) in mouse undifferentiated mesenchymal cell line by API site [22]. The TSA is a histone deacetylase inhibitor. It can lose the chromatin structure in order to let gene restore its expression. 5-aza-dc is also an analog with the same structure of cytosine without methyl group adding in the 5’ C end [23].

2. Materials and Methods

2.1. CpG Island Prediction. The sequence of a putative CpG island in OPN promoter was analysed by using MethPrimer software (http://www.urogene.org/methprimer/index1.html).

2.2. Cell Culture. The porcine fibroblast cell line was grown in Dulbecco’s modified Eagle’s medium (DMEM; Gibco-BRL, Gaithersburg, MD, USA) supplemented with 10% fetal bovine serum (FBS, Gibco BRL) and containing 100 U/mL penicillin and streptomycin. The cells were incubated at 37°C in humidified incubator with 5% CO2.

2.3. 5-aza-dc Demethylation Drug Treatment. For 5-aza-dc treatment, porcine fibroblast cell in 5th passage cultures was treated with 5-aza-dc (sigma) at various concentrations, that is, 0 (control), 0.5, 1.5, and 2.0 μM, for 72 h. Medium was changed every 24 h and then cells were collected for RNA and DNA extraction and stored at −80°C [24].

2.4. Quantitative Real Time-PCR. 2 μg RNA of ear fibroblast cell was used to be transformed to cDNA. 0.5 μL of cDNA was performed for quantitative real time-PCR with Rotor-Gene 6000 (Corbett). β-actin was the internal control to normalize target gene, OPN. The calculated gene expression fold from CT value was according to the previous study. P value less than 0.5 exhibited the obviously significant difference.

2.5. Methylation Analysis by Combined Bisulfite Restriction Analysis (COBRA). For amplification of porcine OPN promoter methylation analysis site, PCR was performed using 2 μL of bisulfite-converted genomic DNA as template. The primer sets of COBRA were OPN-C sense 5’-TTTTTTGAGGGAGATTAGTTTTTG-3’ and antisense 5’-ATTCCTACTAAAACTCAAACCCACCC-3’. The COBRA-PCR products were purified by phenol/chloroform, followed by ethanol precipitation. The DNA was resuspended in 8.5 μL of distilled deionized water. Purified PCR products were then digested with 10 U BstUI restriction enzyme (New England Biolabs, MA, USA) at 65°C. Products were electrophoresed on 6% native acrylamide gel, stained with 200 g/mL ethidium bromide, and visualized using a Kodak 1D software.

2.6. Methylation Specific-PCR. Genomic DNA (0.5 μg) was treated with sodium bisulfite according to the manufacturer’s recommendations (EZ DNA Methylation Kit; Zymo research, CA, USA) and amplified with specific primers for methylated or unmethylated DNA. The primer sets of MS-PCR were OPN-M sense 5’-AAGCGGGAAGGAGTATTACGT-3’, antisense 5’-TCCGACAAAAACGAAAACATACACA-3’, OPN-U sense 5’-GAAGTGGGAGAGTATTATGTG-3’, and antisense 5’-CAATAACTCCAAAACAACAACTTC-3’. All PCR reactions were performed on PTC 200 thermocyclers (MJ Research, MA, USA) and in 25 μL volume using the PlatinumTaq DNA polymerase system (Invitrogen, CA, USA). PCR products were separated on 1.5% agarose gels. The M-set primers contained at least three
CpG sites to distinguish the methylation status of investigated region. And U-set primers overlapping the M-set primers were used to amplify the unmethylated region.

2.7. Plasmid Constructs. A full length pig OPN promoter (−2615-luc) was amplified from wild-type pig heart tissue cDNA. This fragment was cloned into a luciferase fusion plasmid, pGL3-Enhancer vector (Promega), to generate pOPN-full-luc. HindIII and NcoI cutting sites were used for cloning. Three truncated forms of pOPN promoter were prepared by PCR using the pOPN-full-luc as a template and using synthesized oligonucleotides as follows: pOPN-full-luc: sense, 5'-AAGCTTGAATTCCGTCTTCTTCTGGAGA-3' , and antisense, 5'-CCATGGGCTGACAGCTGGACCTCCC-3'; −2239-luc: sense, 5'-AAGCTTCTTACTAATGTACTAC- GTTCAATATTAGC-3', and antisense, 5'-CCATGGGCTGACAGCTGGACCTCCC-3'; −1505-luc: sense, 5'-AAGCTTATTTCTAATTACATTTAAGTAACCAACTTTATATATC-3', and antisense, 5'-CCATGGGCTGACAGCTGGACCTCCC-3'. The sequence of constructs was confirmed by DNA sequencing. There were two-point mutation different from NCBI: one is 287A to T and the other is 957T to A.

2.8. Transient Transfection and Luciferase Assay. Pig ear fibroblasts were transfected using the Lipofectamine 2000 (Invitrogen). Fibroblasts cells were incubated at a density of 8 × 10^3 cells into 35 mm diameter dishes. After 24 h when cell was adherent to the dishes, 3 μg of reporter plasmid DNA was transfected for 6 h in Lipofectamine mixture (Invitrogen). 24 h after the transfection, cell lysates were collected for a luciferase assay. The luciferase activity of the cell lysates was detected by Dual-light system (Applied biosystems). The activity data was measured with PARADIGM Detection Platforms (Beckman Coulter). Luciferase activity was normalized with 1μg β-gal plasmid. All luciferase assays were carried out in triplicate.

2.9. In Vitro Methylation of the OPN Promoter Region. The OPN reporter construct −495-luc was methylated by incubation with SssI methyltransferase (New England BioLabs). The −2615-luc construct was methylated by HhaI and HpaII methyltransferase (New England BioLabs) for 16 h at 37°C. The methylation status was also verified by digested with HhaI and HpaII enzyme.

2.10. Electrophoresis Mobile Shift Assays. Nuclear extracts were prepared from HEK293T cells. Two probes were designed for methylation binding activity test. Two probes containing either 6–8 CpG sites or 13th CpG site in the −2615 to −2239 of the OPN promoter were generated by annealing two complementary oligonucleotides (OPN 6–8: 5'-TGCGATGTACGGTCCTCCGCCCC-GTGCCGGAGTACGTACGGGAAACCAGAGGTG-3'; 5'-ACCTCCGATCCTCCCCGATGGCAGGGCCGACCTGGACGATTGCA-3', the predicted core

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**Figure 1:** (a) (b) The semiquantitative RNA expression of OPN. C1–C4 indicated four different cloned pigs. The black arrows represent the contrary expression in the cloned pig tissues relative to WT tissues. Br: brain, Ea: ear, He: heart, Ki: kidney, Li: liver, Lu: lung, Mu: muscle, Sk: skin, In: intestine, Sp: spleen, Pl: placenta, Um: umbilical cord, B: blood; S: blood treated with SsSI, and W: ddH2O.

**Figure 2:** MS-PCR analysis of OPN promoter. (a) The distribution of CpG sites in the pig OPN promoter. The two arrows indicated the MS-PCR primer sets. (b) The MS-PCR results of OPN promoter in cloned pigs. The arrows represent both existence of methylation and unmethylation DNA element. Br: brain, Ea: ear, He: heart, Li: liver, Lu: lung, Mu: muscle, Pl: placenta, B: blood; S: blood treated with SsSI, and W: ddH2O.
sequence of the API binding site is underlined; OPN13th: 5'-CCTCCGTGGTCCTGTTAATGTTGATGCCTGGAAATAGTTCC3'; 5'-GAACTATTTCCCAACAACGCAGCCTACTAACATTTAACGGAGGAGG3'; the predicted core sequence of the ADRI binding site is underlined). The transcription factor prediction software is TFSEARCH 3.0 version. The probes were labeled with γ-32P-ATP by using T4 kinase (Promega). Annealing probes also were methylated with SssI methyltransferase (NEB). Nuclear extracts containing 5.6 μg of the protein were preincubated in 20 μL of binding buffer (50 mM Tris-HCl (pH 8.0), 750 mM KCl, 2.5 mM EDTA, 0.5% Triton-X 100, 62.5% glycerol (v/v), and 1 mM DTT) with or without unlabeled competitor (10-fold molar excess). For supershift assay, antibody of AP1 was added to the preincubation buffer. After 10 min of preincubation on ice, the DNA probe labeled with [γ-32P]-ATP was added, and the mixtures were incubated at room temperature for 30 min. The reaction mixtures were resolved on 6% polyacrylamide gels. The gels were dried and subjected to PhosphorImager analysis using a Typhoon system and ImageQuant TL software (Amersham Biosciences, Sunnyvale, CA, USA).

2.11. Molecular Modelling. The molecular docking was then further analyzed for proving the further mechanism of our findings; we further surveyed the interaction of API (c-Jun) and TFIIB by computational biology. The API (c-Jun) is possible higher spot for hypermethylation in OPN promoter area and provides the binding domain for RNA-polymerase II initial binding transcription factor (TFIIB) in this study. Therefore, we first utilized the Z-DOCK program to simulate the structures of c-Jun and TFIIB. After that, we further used molecular dynamics (MD) to validate the stability of c-Jun and TFIIB complex under the GROMACS 4.5.5 program [25] with charmm27 force field. The model is set in the TIP3P water modeling in 1.2 nm distance of box for water box setting. Na and Cl ions in the concentration of 0.145 M NaCl model are used for system neutralization. All bonds are fixed by linear constraint solver (LINCS) algorithm to constrain all bonds lengths in the simulation system. Newton's Law is utilized for calculating the motion of molecular dynamics as follows:

$$\frac{d^2 r}{dt^2} = M^{-1}F.$$  \hspace{1cm} (1)

The Particle mesh Ewald (PME) is also used for calculate the coulomb type of electrostatics. The Van der Waals (VDW) interactions are set as 1.4 nm cut-off distance for nonbond interaction. The first step is set on the 5,000 cycle steps performed in the manner of Steepest Descent algorithm for energy minimization. And then, equilibration was performed.
Figure 4: The bisulfite sequencing of OPN\textsuperscript{−2610\textendash 2400} nt upstream the promoter in cloned pigs’ tissues. The closed circles represent the methylation CpG sites. The hollow circles represent the unmethylated CpG sites. The bottom number indicated the methylation percentage of each sample. The range of square showed that the region may be the methylation controlled region of OPN promoter.

Figure 5: Methylation and deletion analysis of OPN promoter in 293T cells. The match-like bar with black circle represents the methylation CpG site; white circle of match bar indicated the unmethylation CpG sites. PGL3 vector as standard; in 293T cell line; PGL3-enhance vector as negative control; cell lysate as the background; pCMV-b-gal as internal control. The relative value is adjusted by cell lysate; −495 M-luc indicated the methylation in PGL3 backbone with \textit{HhaI} and \textit{HpaII} methyltransferase. (**\textit{P} < 0.01); −2615-luc and −2615 M-luc: \(n = 3\); −495-luc and −495 M-luc: \(n = 4\). The experiments were repeated three times and the results were analyzed and presented as the mean ± SE.
Figure 6: The sequence and CpG sites distribution of pig OPN promoter. 20 CpG sites exist in the front region of OPN promoter. The underline indicates the predicted AP1-like binding site. The gray marker indicates two methyltransferase sites of HhaI (GCGC) and HpaII (CCGG).

Figure 7: Electrophoresis mobile shift assay in porcine OPN promoter CpG sites. (a) Using CpG1 contained DNA element binding with SH-SY5Y nuclear extract. Methylated and unmethylated competitors were used as 2X, 5X, and 10X concentration than isotope labeled probe. (b) CpG13th probe of OPN promoter binding with HEK-293 nuclear extract. (c) CpG20th of OPN promoter element binding with HEK-293 nuclear extract. There was no difference between methylated or unmethylated competitors in competition experiment.

Figure 8: The CpG 13 and CpG 20 showed competition binding activity characteristics with c-Jun and c-Fos. The competition reduced the binding activity between transcription factors and labeled OPN EMSA probe. The amount of EMSA c-Jun and c-Fos antibody is 3 μg. The EMSA probe is 30 μg. Adding the Ab with a premix way can reduce the binding effects with the transcription factors. It is suggested that the premix with antibody blocks the access to its binding site (antibody-transcription factor-DNA).

2.12. Molecular Dynamics Analysis. First we survey the stability of all atoms performed by using the GROMACS 4.5.5 software though the commands of g_rms and g_gyrate to calculate root mean square deviation (RMSD) and radius of gyration (Rg), respectively. Secondly, we calculate the total energy for all the systems by the command of g_energy. Thirdly, we further calculate root mean squared fluctuation (RMSF) for each protein residue by commands of g_rmsf. Fourthly, the distance between c-Jun and TFIIB and movement analysis are calculated by the g_dist program. Fifthly, the migration of dock protein (c-Jun and TFIIB) is
3. Results

3.1. Distribution of Porcine OPN CpG Island. There are dense CpG sites existing in the front of the OPN promoter region. One putative CpG island was found (CpG island size > 100, GC Percent > 50.0, Obs/Exp > 0.6) by MethPrimer program.

3.2. Methylation and Expression Analysis of OPN in Cloned Pig Various Tissues. Firstly, the OPN mRNA expression was investigated in the WT pig tissues and cloned pig tissues. Data showed the various expression levels in different tissues. Particularly, Copy1 brain overexpressed the OPN and Copy2 ear with no expression of OPN relative to its wild-type tissue, respectively (Figure 1). The unique aberrant
expression patterns exhibited the different control way of the OPN expression. We proposed that OPN expression may be a tissue-specific manner. MS-PCR primers were designed to estimate the methylation status of OPN promoter. Hypomethylation generally appeared in the various tissues of cloned pigs. However, there were still some tissues that showed the methylated region in OPN promoter (Figure 2).

3.3. 5-aza-dc Increases OPN mRNA and Decreases Methylation of OPN Promoter in Pig Ear Fibroblast Cell. In order to realize whether the methylated OPN promoter affects the activity of OPN promoter. The 5-aza-dc treated porcine ear fibroblast cells showed that when the concentration of 5-aza-dc level increased, it will decrease the methylation of OPN promoter and restore the OPN RNA expression at 0.5 to 2.0 μM (Figure 3). The results suggested that the activity of OPN promoter can be affected by DNA methylation directly or indirectly. COBRA assay was also used to investigate the methylation status of OPN promoter in WT tissues. Brain, ear, liver, and lung tissues exhibited little part methylation of the OPN promoter (data not show). This is thought that the methylation of OPN promoter in the aforementioned tissues may be involved with OPN transcript regulation mechanism.
3.4. Bisulfite Sequencing Analysis of the Whole CpG Sites Methylation Profile in Cloned Pigs. To investigate which region of the OPN promoter is affected by methylation in the CpG site, bisulfite sequencing was performed to dissect the methylation status of CpG sites in the OPN promoter. In the brain tissue, bisulfite sequencing of Copy5 and Copy1 brain exhibited the saturated status in their methylated region while WT brain exhibited fragmentary methylated CpG sites. The 18.06% methylation percentage of Copy5 brain was more than the WT brain, 14.25% (Figure 4). In the heart tissues, bisulfite sequencing showed that the Copy2 heart had extremely hypermethylated percentage with 13.33% more than WT heart, 2.94% (Figure 4). Particularly, the data showed the inhibition of Copy2 heart mRNA (Figure 1). In the liver, the different methylation pattern also appeared in the Copy4 liver relative to WT liver. Copy2 ear with 18.05% methylation in the OPN analyzed region was higher than the WT ear, 11.67%; the result was proved by the experiment of OPN mRNA expression (Figure 1). These bisulfite sequencing results matched our previous hypothesis that methylation in
the OPN promoter region regulates the activity of OPN promoter. Moreover, the hypermethylated OPN promoter may directly affect the activity of OPN transcription, especially in heart and ear tissues.

3.5. Analysis of Methylation Implication of OPN Transcription by Promoter Assay. The analyzed OPN promoter region may involved with methylated control of gene transcription. Promoter assay was designed to explain the direct inhibition of OPN promoter activity by methylation on the OPN CpG sites. Four different truncated forms of OPN full length (2.6 kb) were used to prove the hypothesis (Figure 5). Particularly, the truncated form 2.2 kb deleted the 377 bp promoter region (−2615−−2239 nt). This region is the analyzed region for bisulfite sequencing profile. And this region is thought to be the most possible element that regulates the OPN transcription. Moreover, four CpG sites were in vitro added to the methyl group by methyltransferase that can provide important evidence how methylation affect the OPN transcription. Figure 5(b) shows that the OPN promoter activity was significantly decreased in methylated −2615-luc plasmid. However, −2239-luc that deleted the 377 bp containing methylated characteristic DNA element leads to less inhibition of promoter activity than the methylated vector −2615 M-luc (Figures 5(a) and 5(b)). The results indicated that methylation in the front OPN promoter is not only decreasing the promoter activity to the basal level but also recruiting the inhibition factors to enhance the inhibition ability. In order to avoid the effects of CpG sites in PGL3-enhancer backbone, −495 M-luc that have no methylated CpG sites in the OPN promoter but it can be methylated in the vector backbone CpG sites compared to −495-luc. Data showed that there is no difference in the promoter activity between methylation or unmethylation in the PGL3-enhancer backbone CpG sites (Figure 5(b)). It is suggested that methylation in the critical region, such as OPN promoter front end, may lead to the rearrangement of chromatin structure. Otherwise, deletion of the OPN promoter to the 495 bp with significant promotion of the promoter activity indicated that in the middle part of promoter DNA element may able to inhibit the activity of OPN promoter.

3.6. Methylation in CpG 13th and CpG 1st of OPN Promoter Blocks the Binding Access of Transcription Factors. We investigate that the methylated CpG sites in the −2615−−2239 nt of the OPN promoter region affect the transcription factor binding activity. Electrophoresis mobile shift assay was performed with nuclear extracts from human HEK293T and SH-SY5Y cell line. Four EMSA probes that contain the CpG 1, CpG 3-4, CpG 6–8, CpG 11–15, and CpG 19–20 were designed according to the CpG sites in our analyzed region (Figure 6). EMSA data suggested that CpG 13th and 1st sites showed methylation noncompetition phenomenon which had influence on binding with transcription factor (Figure 7). 19-20th CpG sites containing EMSA probe showed no competition ability in the methylated or unmethylated status. The premix with antibody and nuclear extracts by EMSA assay indicated that the c-Jun and c-Fos were involved in the binding to CpG sites 13 and 20 (Figure 8). However, the adding of antibody in the mixture of probe and nuclear extracts showed no significant shift bands. It is indicated that other transcription factors may also participate in the transcription activity OPN promoter. Thus, the c-Jun and c-Fos could be involved in the partial OPN transcriptional activity in a competition way. In addition, the c-Jun had higher binding affinity than c-Fos in the EMSA probe analysis. We therefore did followed computational survey to see why c-Jun affects the consequent OPN transcription.

3.7. The Computational Biology Results. After surveying the possible zone for OPN promoter hypermethylation, we found that the API (c-Jun) sequence frequently appeared in our
molecular laboratory study. Then we further validated that the c-Jun methylation will cease the further mRNA production by inhibiting the binding of RNA-polymerase II initiation factor TFIIB. We do the further computational modelling for mechanism survey between the c-Jun and TFIIB. By the Z-DOCK analysis, we found the c-Jun and TFIIB could combine tightly (Figure 9). We chose the highest docking pose (dock score = 18.04) for further MD analysis.

Then the disorder predication was employed to observe the protein folding analysis and the result is shown in Figure 10. We found that the c-Jun has relative high disorder in folding than TFIIB; we suppose that the flexibility of c-Jun structure could easier bound to TFIIB. This finding could be an explanation why the c-Jun bound to TFIIB by Z-DOCK program. To confirm the stability of the c-Jun and TFIIB complex, the series of molecular dynamic studies further visualize their interactions.

Protein complex RMSD analysis proved that the c-Jun and TFIIB were stable from 3000 ps to 5000 ps. In addition, we found the TFIIB are easier to be stable during the molecular dynamics (Figure 11(a)). We also found the radius of gyration tend to be stable for all simulation times with average of 2.05 nm (Figure 11(b)), suggesting that the two protein structures are compact after binding together. Figure 11(c) also shows the binding complex in a stable fluctuation and the energy of the binding complex is stable around −9.15 × 10^5 (kJ/mol).

3.8. Stability Analysis of Residues on the Major Binding Region during MD Simulation. To analyze the flexibility of residues on protein structure, the RMSF calculation was used to observe the flexibility of each residue; Figure 12(a) shows that the chain A of c-Jun had high frequency of fluctuation (binding site 200–210 binding resides). However, the chain B of the c-Jun has relative fewer frequency of fluctuation and the major binding region (from 228 to 240 residues) showed a less fluctuation as shown in Figure 12(b). Figure 12(c) reveals the binding regions (200–210 binding residues) for chain A of c-Jun that is more fluctuated and unstable compared to the binding regions (228–240) for chain B of c-Jun binding site in the TFIIB binding region. Figure 13 is the result for secondary structure variation calculated by DSSP analysis. Most of the main scaffold belong to alpha Helix; there are no significant changes during the whole MD simulation. All helices of the secondary structure for c-Jun and TFIIB binding remained stable during a 5,000 ps simulation time (Figure 13). We thereafter surveyed the distance between each residue of c-Jun for 5,000 ps. The variation of distances between residues in c-Jun chain is wider than the distances of residues in c-Jun chain B. Therefore, the chain B of c-Jun is more stable for TFIIB binding (Figure 14).

The hydrophobic area was then calculated by SASA in Figure 15(a); the value of hydrophobic area decreased during the last 1000 ps. This indicated the compactness of the c-Jun and TFIIB binding increased by the MD time period in our study. It is worthy to know that the distance between centrales of masses of c-Jun and TFIIB was decreasing more and more after time goes by in the 5000 ps survey (Figure 15(b)). In the migration analysis of c-Jun and TFIIB, the MSD was employed to count the migration of c-Jun and TFIIB. The c-Jun is more unstable than TFIIB during the binding interaction throughout the whole MD simulation in 5000 ps period (Figure 15(c)). Besides, we further utilized the principal component analysis (PCA) to measure all MD frames over all simulation times. The first two eigenvectors (PC1 and PC2) were shown in Figure 18; most of frames are ranged in the short range of eigenvalues −10 and 10 in PC1 (Figure 16(a)), and arranged in eigenvalues from −5 to 5 in PC2 (Figure 16(b)). The phase space comparing for PC1 and PC2 was shown in Figure 17; we found that each frame
could be grouped into two clusters. This suggests that the motion of each frame was not changed significantly over all simulation times. In order to select the most representative structure for snapshot investigation, we did cluster analysis (Figure 18). We found that the last group (cluster 14) is the predominant cluster and is also displayed in the time range from 4000 to 5000 ps; the cluster 14 also appears most predominant in frame numbers (Figure 18), and the middle structure (4260 ps) of cluster 14 is regarded as representative frame. For snapshot analysis, the comparison of initial and representative frames is shown in Figure 19; we found that the chain A of c-Jun is more encompassed by TFIIB at 4260 ps through the inward rotation of TFIIB. This made the bindings between chain A of c-Jun and TFIIB more compacted through 0 ps to 4260 ps along with time. The elevated activation between TFIIB and chain A of c-Jun is also confirmed in RMSF analysis. Therefore, we supposed the initiated transcription factor on RNA polymerase II (TFIIB) is closed interaction to the chain A of c-Jun (AP1) from 0 ps to 4260 ps. This hypothesis was also confirmed by Figure 20. There were more spaces between chain B and TFIIB than chain A. Hence, we could see that the TFIIB acts more close to
Figure 16: The number of MD frames of the first two eigenvectors (PC1 and PC2) by PCA analysis during simulation time of 5000 ps. The higher range of eigenvalue denotes the wider motion of protein structure over all simulation times.

Figure 17: Phase space analysis of comparing the first two eigenvectors (PC1 and PC2) for principle component analysis. The eigenvalues of the two eigenvectors are projected into one phase space; small motion of protein structure could be grouped into clusters.

4. Discussion

Previous study has shown that DNA element (GGGT-CATATGGTTCA) located in osteopontin promoter −2245 to −2259 nt can be regulated by vitamin D3 [27]. This DNA regulation region can easily be affected by the change of calcium concentration. The promoter region of porcine OPN was analyzed in transcription factor binding sites except the region −2615 to −2239. Interestingly, this region of porcine OPN promoter is rich in CpG sites compared to human, mouse, and bovine genome. Sakata also proved that OPN promoter transcription activity is regulated by some specific DNA modification mechanism of rearrangement of chromatin structure [22].

In the present study, four cloned pigs were surrounded by many defects. For example, Copy1 pig had a retardation of limb bone growth. Copy2 heart organ showed a pericarditis and copy3 heart had valvular heart disease. This physiology defects appeared aberrant development, especially in bone or heart, may involved in the initially fetus stage with inappropriate organ differentiate. Thus, our data suggested that the consequent result in aberrant OPN expression or incompletely epigenetic modification in OPN promoter (Figure 5). These aberrant molecular data of OPN are correlated with the defects of bone and heart in cloned pigs. Semiquantitative PCR of OPN mRNA showed that discrepant expression pattern was identified in several cloned pig tissues, especially in brain (99.75% up-regulation), heart (11.5% downregulation), and ear (18.03% downregulation) (Figure 2). OPN mRNA has different expression in brain development in different embryonic stages [28]. The overexpression of OPN in brain tissue may cause some unexpected brain damage or neuron development. OPN can induce myocardial fibrosis and repair tissue after inflammation. Lacking OPN will cause faulty wound healing after myocardial infarction [29, 30]. Silent expression of OPN in cloned pig heart tissue may also be the main cause of heart disease.

Recent studies indicated that OPN gene expression may be affected by treatment of TSA (a histone deacetylase inhibitor). The results applied that OPN promoter could be regulated by epigenetic mechanism [22]. In this study, we investigated OPN methylation profile after 5-aza-dc treatment. The results indicated that mRNA expression of OPN is
Figure 18: Cluster analysis of all MD frames of c-Jun and TFIIB complex during simulation time of 5 ns for identifying representative structure. All MD frames were grouped into fourteen clusters by linkage method; the RMSD cut-off distance between each neighbor frame is 0.14 nm. The most predominant group is the cluster 14, which is displaced in the region of simulation time from 4000 to 5000 ps, and the MD frames in cluster 14 are the most number among all clusters. The middle frame of cluster 14 is displaced in simulation time of 4260 ps.

Figure 19: The structural comparison between the first frame (0 ps) and representative structure (4260 ps). The structures of c-Jun and TFIIB are represented by blue ribbon and red solid phase, respectively. The chain A of c-Jun was surrounded more compactly by TFIIB at 4260 ps by the structural inward rotation to make more compactness between TFIIB and c-Jun through 0 ps to 4260 ps.

Figure 20: The possible space prediction between c-Jun and TFIIB among all simulation times. The predicted channels are colored in red and green. The structures of c-Jun and TFIIB are colored in blue and orange, respectively. Each possible space is represented by channels; each channel was generated by Caver 3.0 program.

directly affected by adding methyltransferase inhibitor 5-aza-dc (Figure 4). COBRA was performed to study the methylation of OPN front end promoter in wild-type and cloned pig different tissues. Sodium bisulfite sequencing analysis also revealed that the methylation of CpG sites concentrated in front of the 20 CpG sites in front of OPN promoter (Figure 5). Discrepancy methylation in this promoter region
also happened in brain, heart, ear, and liver tissues between wild-type and cloned pigs (Figure 5). It revealed that DNA methylation of OPN promoter may be involved with regulation of expression of OPN mRNA. In order to characterise which promoter DNA element is important, four constructs of OPN promoter (−2615-luc, −2239-luc, −1505-luc, and −495-luc) were used for analysis. Obvious downregulation in methylated −2615-luc (Hpall and Hhal methyltransferase) was observed. Compared to the deletion of this control region (−390 nt) in front of OPN promoter, the decreasing level of promoter activity is not as obvious as −2615 M-luc construct (Figure 7). It means that methylation in the front of OPN promoter caused some silent mechanism that make chromosome structure more compact or block some promotion transcription factors. The EMSA data indicated that 13th CpG site of our analyzed region could bind to API transcription factor and binding activity is affected by methylation in this CpG site (Figure 7). Taken together, all these findings correlated with DNA methylation in tissue- or cell-specific gene expression. OPN promoter region was densely methylated in some low expression (Figures 2, 4, and 5).

Our data revealed that DNA methylation of CpG sites in OPN promoter was the main mechanism through specific transcription factor that makes the tissue-specific expression. In previous study, API-like binding site (TGAGCGA) was identified as a methylated insulator region in human blastoma cell line [31]. Analyzed region in front of porcine OPN promoter showed that CpG 1st binding site contained the specific binding site sequence. While in our interesting DNA regulation region range from −2615 to −2239 nt of OPN promoter also exhibited little block access in the competition of probe. It is suggested that CpG 1st and CpG 13th play an important role in methylation controlled mechanism to regulate gene expression. We finally utilize Z-dock program [32] to analyze the interaction between API (PDB code: 1NJM) and RNA polymerase II initial transcription factor (TFIIB) (PDB code: IVOL) [33] to see if they had stable binding. From the docking result of Z-dock (Figure 8), we proved the API is significantly bound to TFIIB. We also found the API can autoregulate the HDAC-1 in promoter region and lead to significant higher degrees of hypermethylation in the OPN promoter region and cause API to be hypermethylated consequently ceasing the OPN mRNA expression [34].

Further mechanical studies by the computational biology also pointed out that the DNA sequence for hypermethylation of OPN promoter binding sites is c-Jun. The chain A of c-Jun could be encompassed more tightly by inward rotational structure change of TFIIB during the MD process (Figure 19). Therefore, we found c-Jun had crucial role for interaction of initiating transcription by RNA polymerase II. The methylation of c-Jun leads to of hyper-condense helix structural change and makes transcription termination which stops OPN mRNA production. Therefore, the MD docking results reconfirm the c-Jun partake the crucial roles in consequent OPN transcription that matches our wet laboratory studies. We suppose this will cause the problems in the embryonic development and lead to threatened conditions. Therefore, adjusting OPN promoter c-Jun (API) methylation will affect transcription binding and could be the treatment for genetic developing errors in the future.

In conclusion, aberrant methylation of porcine OPN gene was frequently found in different tissues of somatic nuclear transferred cloning pigs and bisulfite sequence data suggested that the OPN promoter region of −2615 to −2239 nt may be a crucial regulation DNA element. In pig ear fibroblast cell culture study, the demethylation of OPN promoter was found in dose-dependent response of 5-aza-dc treatment and followed the OPN mRNA reexpression. In cloned pig study, discrepant expression pattern was identified in several cloned pig tissues, especially in brain (99.75% up-regulation), heart (11.5% down-regulation), and ear (18.03% downregulation). Promoter assay data revealed that four methylated CpG sites presenting in the −2615 to −2239 nt region cause significant downregulation (approximately 75%) of OPN promoter activity (P < 0.001). EMSA data also suggested that CpG 13th and 1st sites showed methylation noncompetition phenomenon which had influence on binding with transcription factor.

Conflict of Interests

The authors declare that there is no conflict of interests regarding the publication of this paper.

Authors’ Contribution

Chih-Jie Shen, Yung-An Tsou, and Hsiao-Ling Chen contributed equally to this work.

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