

# StressChip as a High-Throughput Tool for Assessing Microbial Community Responses to Environmental Stresses

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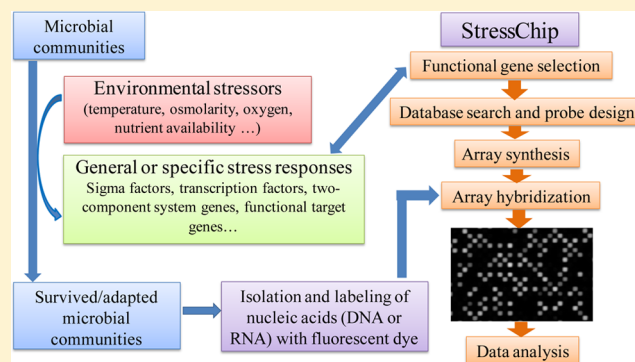
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## Supporting Information

**ABSTRACT:** Microbial community responses to environmental stresses are critical for microbial growth, survival, and adaptation. To fill major gaps in our ability to discern the influence of environmental changes on microbial communities from engineered and natural environments, a functional gene-based microarray, termed StressChip, has been developed. First, 46 functional genes involved in microbial responses to environmental stresses such as changes to temperature, osmolarity, oxidative status, nutrient limitation, or general stress response were selected and curated. A total of 22,855 probes were designed, covering 79,628 coding sequences from 985 bacterial, 76 archaeal, and 59 eukaryotic species/strains. Probe specificity was computationally verified. Second, the

usefulness of functional genes as indicators of stress response was examined by surveying their distribution in metagenome data sets. The abundance of individual stress response genes is consistent with expected distributions based on respective habitats. Third, the StressChip was used to analyze marine microbial communities from the Deepwater Horizon oil spill. That functional stress response genes were detected in higher abundance ( $p < 0.05$ ) in oil plume compared to nonplume samples indicated shifts in community composition and structure, consistent with previous results. In summary, StressChip provides a new tool for accessing microbial community functional structure and responses to environmental changes.



## 1. INTRODUCTION

Microorganisms are the most diverse group of life presently known, inhabiting almost every imaginable environment on Earth. They play important roles in ecosystem functioning, such as biogeochemical cycling of carbon (C), nitrogen (N), sulfur (S), phosphorus (P), and various metals. The composition, structure, and functions of microbial communities are dynamic and sensitive to environmental changes. Given the complexity of microbial communities and divergent molecular mechanisms of microbial responses to environmental stress, it remains a great challenge to monitor stress responses of microbial communities to environmental changes. Therefore, development of rapid, specific, sensitive, and high-throughput quantitative tools for detecting the responses of microbial communities to environmental stresses is greatly needed.

The adaptation of microbial communities to environmental changes involves the regulation of microbial populations, gene expression and activity, and/or genetic/evolutionary changes. Extensive studies in model microorganisms such as *Escherichia coli* and *Bacillus subtilis* have demonstrated that a complex global regulatory network including regulation at the transcriptional or post-transcriptional levels are important aspects of stress responses.<sup>1–3</sup> For example, two-component systems, such as *desK-desR*, control the cold induction of a desaturase gene, *des*, which is important for keeping proper membrane fluidity at low temperature;<sup>4</sup> regulatory gene *hrcA* and chaperon machinery

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genes *dnaK*, *grpE*, *groES*, and *groEL* are involved in heat shock responses.<sup>5</sup> The ProU system genes *proX*, *proV*, and *proW* play important roles in osmotic stress response.<sup>6,7</sup> Due to the difficulty of monitoring post-transcriptional changes, regulatory and/or functional genes that have been demonstrated to be involved in stress response in pure cultures are targets for analyzing microbial communities in response to environmental changes.

Key functional genes involved in microbial functional processes have been shown to be very useful markers for examining functional diversity, composition, structure, and metabolic potential/activity of microbial communities with microarray platforms, namely, functional gene arrays (FGAs).<sup>8–10</sup> Comprehensive FGAs, such as GeoChip, have been developed and applied to analyze microbial communities from different habitats,<sup>11–16</sup> and experimental results demonstrate GeoChips are specific, sensitive, and quantitative tools to rapidly profile microbial communities and link these communities to ecosystem functioning.<sup>8</sup> Great effort has been undertaken to optimize the probe design and improve the hybridization specificity and sensitivity of GeoChip.<sup>11,13,14,17–23</sup> Whole-genome amplification protocols for microbial community DNA/RNA samples were established to overcome limitations due to low biomass or low DNA/RNA yield from community samples.<sup>22,24</sup> However, high-throughput cultivation-independent tools for monitoring microbial community responses to environmental stresses are still lacking.

In this study, we have developed a stress genes-based microarray, termed StressChip, for the assessment of microbial responses to environmental stresses. Functional genes involved in general and specific stress responses to changes of temperature, osmolarity, oxygen level, and nutrient limitation were included. We primarily focus on the selection of the stress response genes, computational evaluation of the probes, analysis of the distribution of the functional genes in metagenomes from various environmental microbial communities, and application of the StressChip to analyze seawater microbial communities. To our knowledge, this is the first metagenomic tool developed for comprehensive examination of microbial community responses to environmental stresses.

## 2. MATERIALS AND METHODS

**2.1. Selection of Key Functional Genes.** A total of 46 functional genes have been selected to target general and specific stress responses (Table 1).

(i). *Sigma Factor, General Stress Response, and Stringent Response.* We selected four sigma factors,<sup>25</sup> including  $\sigma^{70}$  (RpoD, the “housekeeping” sigma factor or primary sigma factor),  $\sigma^{38}$  (RpoS, induced during entry into stationary phase and controls the general stress response in Gram-negative bacteria),<sup>26</sup>  $\sigma^{32}$  (RpoH, the heat shock sigma factor), and  $\sigma^{24}$  (RpoE, the extracytoplasmic/extreme heat stress sigma factor), *katE* (haem-catalase)<sup>27</sup> for general stress response, and *obgE* (encoding a GTPase involved in keeping low intracellular concentrations of ppGpp) for stringent response.<sup>28–31</sup>

(ii). *Temperature Stresses.* Fluctuations in temperature are often encountered by environmental microorganisms. Most of the heat shock proteins are molecular chaperones or ATP-dependent proteases which promote renaturation or degradation of partially unfolded proteins.<sup>32</sup> The microbial cold shock response involves an increase in the proportion of unsaturated fatty acids (UFAs) in membrane lipids and the induction of cold shock proteins (Csp) as chaperones to eliminate secondary

**Table 1. Summary of the StressChip probes**

stress response	gene name	no. of probes	sequence-specific probes	group-specific probes	covered CDS
sigma factor	$\sigma^{24}$	475	57	418	1008
	$\sigma^{32}$	1990	225	1765	4284
	$\sigma^{38}$	371	26	345	1026
	$\sigma^{70}$	1881	106	1775	5249
general stress response	<i>katE</i>	961	172	789	2326
stringent response	<i>obgE</i>	1264	54	1210	3990
heat shock	<i>hrcA</i>	644	130	514	1467
	<i>dnaK</i>	262	0	262	715
	<i>GroES</i>	33	10	23	69
	<i>groEL</i>	153	51	102	302
	<i>grpE</i>	563	24	539	1295
cold shock	<i>desR</i>	9	0	9	18
	<i>desK</i>	7	0	7	14
	<i>cspA</i>	36	0	36	88
	<i>cspB</i>	18	0	18	50
osmotic stress	<i>opuE</i>	25	1	24	65
	<i>proX</i>	50	23	27	80
	<i>proV</i>	364	14	350	1106
	<i>proW</i>	18	3	15	35
oxidative stress	<i>perR</i>	32	3	29	69
	<i>oxyR</i>	522	15	507	1454
	<i>ahpC</i>	903	42	861	2437
	<i>ahpF</i>	369	31	338	1182
	<i>katA</i>	90	13	77	204
oxygen limitation	<i>fnr</i>	1925	2	1923	10160
	<i>narG</i>	1278	502	776	3501
	<i>narH</i>	157	22	135	410
	<i>narJ</i>	99	20	79	198
	<i>narI</i>	323	36	287	1004
	<i>cydA</i>	131	3	128	333
	<i>cydB</i>	159	5	154	430
	<i>arcA</i>	67	5	62	207
	<i>arcB</i>	58	30	28	86
	<i>bglP</i>	44	2	42	88
glucose limitation	<i>bglH</i>	42	1	41	93
	<i>phoA</i>	239	40	199	623
phosphate limitation	<i>phoB</i>	597	27	570	1455
	<i>pstS</i>	322	38	284	698
	<i>pstA</i>	803	80	723	1915
	<i>pstB</i>	723	0	723	3522
	<i>pstC</i>	2800	0	2800	16660
	<i>glnA</i>	1296	7	1289	7079
nitrogen limitation	<i>glnR</i>	158	4	154	894
	<i>tnrA</i>	7	0	7	21
	<i>ctsR</i>	159	2	157	647
protein stress	<i>clpC</i>	428	12	416	1071
	<b>total</b>	<b>46</b>	<b>22855</b>	<b>1838</b>	<b>21017</b>
					<b>79628</b>

structures and restore RNA functionality.<sup>33</sup> Functional genes targeting heat shock such as chaperon machinery genes *dnaK*, *grpE*, *groES*, and *groEL* and the regulatory gene *hrcA*<sup>5</sup> or cold shock such as cold shock protein genes *cspA* and *cspB*<sup>34,35</sup> and two-component system genes *desK-desR*<sup>4</sup> exclusively controlling the cold-induction of a desaturase gene *des* were included in the StressChip.

(iii). *Osmolarity*. Cells respond to osmotic shock by adjusting the cellular concentration of osmolytes or compatible solutes.<sup>36</sup> We selected *opuE* (osmoprotectant uptake, single component sodium/solute symporter with high affinity for proline<sup>37,38</sup>) and the ProU system genes *proX*, *proV*, and *proW* (with a broad substrate specificity for osmoprotectants but a clear preference for glycine betaine and proline betaine<sup>6,7</sup>) for monitoring microbial responses to osmotic stress.

(iv). *Oxidative Stress and Oxygen Limitation*. Oxidative stress represents an imbalance between the production and scavenging of reactive oxygen species (ROS). We selected the functional genes *ahpCF* (alkyl hydroperoxide reductase) and *kata* (catalase) involved in detoxification of ROS and regulatory genes *perR* and *oxyR* for oxidative stress response.<sup>39–41</sup> When the oxygen level decreases, microorganisms adjust their energy metabolism for survival. Cytochrome genes (*cydA* and *cydB*), activated when oxygen becomes limiting, regulatory genes for *cydAB* expression such as *fnr* and two-component system genes (*arcA* and *arcB*),<sup>42,43</sup> and nitrate reductase genes (*narG*, *narH*, *narJ*, and *narI*) allowing nitrate as an electron acceptor grown anaerobically<sup>44</sup> were chosen to target oxygen limitation.

(v). *Nutrition Limitation*. Limited nutrition such as glucose, phosphate, and nitrogen is a common stressor faced by microbes in the natural environment. We selected *bglP* (aryl-beta-glucoside-specific enzyme II) and *bglH* (phospho-beta-glucosidase) for utilization of aryl- $\beta$ -glucosides as alternative carbon sources<sup>45</sup> to target glucose limitation. Phosphate is an essential nutrient for microbial metabolism because the majority of the building blocks of cellular functional structures are phosphorylated. Phosphate-specific transport system (Pst, the main inorganic phosphate (Pi) uptake system under phosphate limiting conditions) genes *pstS*, *pstA*, *pstB*, and *pstC*, alkaline phosphatase gene *phoA* liberating free Pi from external sources, and their regulator gene *phoB* (response regulator) were selected for phosphate limitation.<sup>46–52</sup> The regulation of nitrogen (N) metabolism upon N limitation is quite diverse among different microbial organisms. Glutamine serves as a sensor of external nitrogen availability in enteric bacteria.<sup>53</sup> Glutamine synthase gene *glnA* and regulatory genes *tnrA* and *glnR* were selected for nitrogen limitation.<sup>54</sup>

(vi). *Protein Stress*. Overproduction of recombinant proteins in microbial cells stimulates protein stress.<sup>55,56</sup> Accumulation of recombinant protein induces the activation of heat shock sigma factor  $\sigma^{32}$  and  $\sigma^{32}$ -dependent genes.<sup>57</sup> We selected *clpC* (ATPase subunit in the Clp machinery) and regulator gene *ctsR* to target protein stress.<sup>58–60</sup>

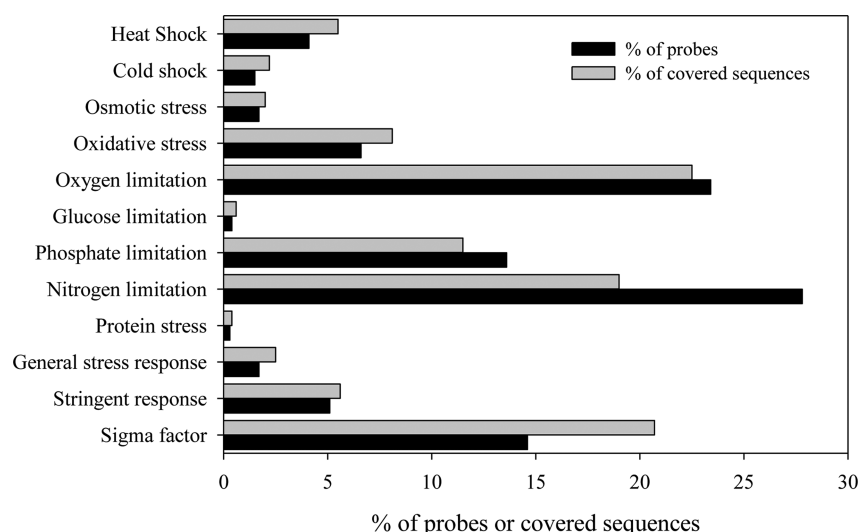
**2.2. Retrieval and Verification of Functional Gene Sequences.** Sequence retrieval was performed using the GeoChip design pipeline.<sup>14</sup> Briefly, for each functional gene, keywords such as the name of the target gene or protein, its abbreviation or enzyme commission number (EC), and affiliated domains (bacteria, archaea, and fungi) were listed as a query and submitted to the GenBank Protein Database to fetch amino acid sequences. Then, the sequences were downloaded and verified by HMMER 2.3.2 (Ashburn, VA, USA) with seed sequences, whose functions have been experimentally characterized in certain microorganisms. Finally, the nucleic acid sequences for the confirmed protein sequences were obtained from GenBank and used for probe design. All sequences were downloaded from the GenBank database before Feb. 5 2010.

**2.3. Oligonucleotide Probe Design, Computational Evaluation of Probes, and Construction of The StressChip.** Oligonucleotide probes were designed with an improved version of CommOligo,<sup>20</sup> which has group-specific probe design

**Table 2. Microbial Domain and Phylum Covered by StressChip**

domain	phylum	no. of species	no. of probes	no. of covered CDS
archaea		76	709	2417
	Crenarchaeota	19	99	411
	Euryarchaeota	52	573	1904
	Korarchaeota	1	7	24
	Nanoarchaeota	1	1	2
	Thaumarchaeota	2	8	28
	unclassified	1	21	48
bacteria		985	21689	76324
	Acidobacteria	4	145	340
	Actinobacteria	124	1931	6381
	Aquificae	5	102	302
	Bacteroidetes	62	1069	2957
	Candidatus Poribacteria	1	4	4
	Chlamydiae	9	66	331
	Chlorobi	11	265	658
	Chloroflexi	8	297	888
	Cyanobacteria	23	1215	3618
	Deferribacteres	1	24	56
	Deinococcus-Thermus	9	145	417
	Dictyoglomi	2	21	62
	Elusimicrobia	2	18	45
	Fibrobacteres	1	22	54
	Firmicutes	221	4063	18397
	Fusobacteria	12	78	188
	Gemmatimonadetes	1	37	94
	Lentisphaerae	2	31	72
	Nitrospirae	2	35	85
	Planctomycetes	8	276	531
	Proteobacteria	415	9931	35403
	Spirochaetes	17	122	472
	Synergistetes	4	65	153
	Tenericutes	19	98	378
	Thermotogae	12	101	440
	Verrucomicrobia	7	256	577
	unclassified bacterium	3	1272	3421
eukaryota		59	457	887
	Ascomycota	46	386	753
	Basidiomycota	9	64	123
	Glomeromycota	1	2	3
	Microsporidia	2	3	6
	Neocallimastigomycota	1	2	2
total		1120	22855	79628

features. Oligonucleotide probes were chosen on the basis of the following criteria: (i) gene-specific probes: targeting a single sequence and having  $\leq 90\%$  sequence identity,  $\leq 20$ -base continuous stretch, and  $\geq -35$  kcal/mol free energy with their nontarget sequences;<sup>21</sup> (ii) group-specific probes: targeting a group of highly homologous sequences exclusively and having  $\geq 96\%$  of sequence identity,  $\geq 35$ -base continuous stretch, and  $\leq -60$  kcal/mol of free energy within the group.<sup>19</sup> All designed probes (50-mer) were verified by the in-house software ProbeChecker. GenBank (NR) nucleic acid database was used for computational specificity evaluation of the probes. The StressChip was manufactured by NimbleGen (Madison, WI, USA). The StressChip design has been integrated into the more comprehensive GeoChip 4.0,<sup>61,62</sup> the latter being used to analyze the samples presented in this study.



**Figure 1.** Distribution of probes and targets for different stress responses in the StressChip.

## 2.4. Analysis of the Distribution of Stress Functional Genes in Metagenomes from Different Environments.

Seven metagenomes from different environments were downloaded from the Joint Genome Institute Integrated Microbial Genomes (IMG) database (<http://img.jgi.doe.gov/cgi-bin/m/main.cgi>). These samples represent four ecosystem types including marine, freshwater, nonmarine saline and alkaline, and desert soil. Detailed information of the metadata is listed in Table S1, Supporting Information. Distribution of 46 stress functional genes in these metagenomes was selected from the abundance profiles of COG function in these metagenomes. The hierarchical linkages of the metagenomes were generated with Gene Cluster 3.0.

## 2.5. Target DNA Preparation, Hybridization, and StressChip Data Analysis.

Seawater samples were collected from the Gulf of Mexico between May 27 and June 2, 2010 after the Deepwater Horizon oil spill occurred.<sup>61</sup> Eight samples (BM053, BM054, BM057, BM058, BM064, OV201, OV401, and OV501) from the MC252 dispersed oil plume and five samples (OV003, OV004, OV009, OV013, OV014) from a nonplume area at a depth of 1099–1219 m<sup>62</sup> were analyzed in this study.

Approximately 100 ng of DNA isolated from 0.22  $\mu$ m polyethylenesulfone membranes filters (MO BIO Laboratories, Inc., Carlsbad, CA)<sup>61</sup> was amplified with the Templiphi kit (GE Healthcare; Piscataway, NJ).<sup>22</sup> Cy3 labeling of the amplified DNA (2  $\mu$ g) and hybridization were performed as described previously.<sup>62</sup> After a 16 h hybridization at 42 °C with 40% formamide on a MAUI hybridization station (BioMicro Systems, Salt Lake City, UT, USA), the arrays were scanned with a NimbleGen MS200 microarray scanner (RocheNimbleGen, Madison, WI, USA) at a laser power of 100% and 100% PMT (photomultiplier tube).

The raw data from hybridization was preprocessed by removing spots with signal-to-noise ratio (SNR) less than 2.0. SNR was calculated as (average of signal pixels in the probe space – average of background pixels in four void spots)/standard deviation of background pixels. Genes detected in only one sample were removed. Processed StressChip data (Table S2) is available in the Supporting Information. Canonical correspondence analysis (CCA) was used to link the stress responses of the microbial communities to environmental variables.

## 3. RESULTS AND DISCUSSION

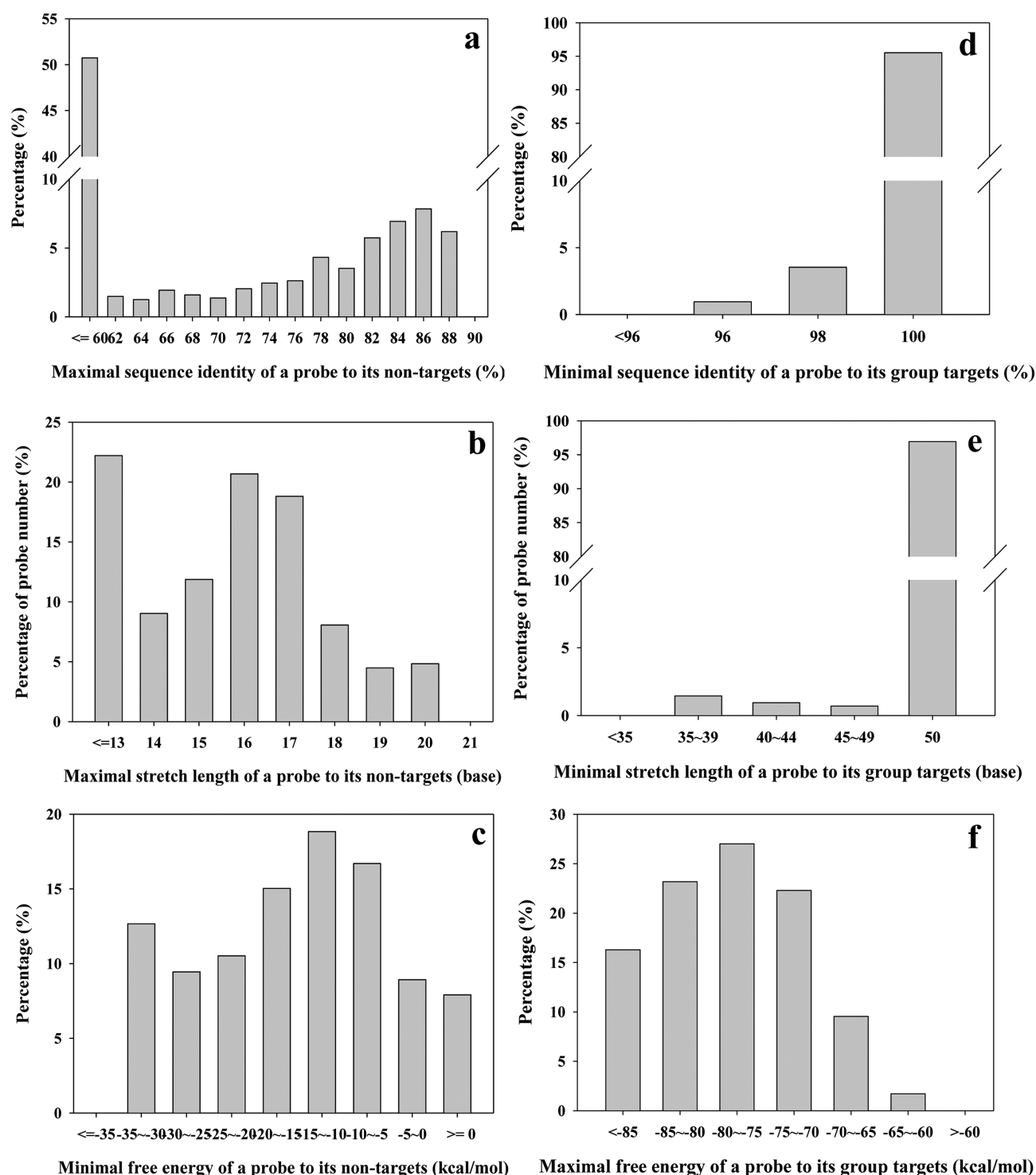
### 3.1. Features of the StressChip.

A total of 46 functional gene families with 40 targeting specific stress response pathways including nine for temperature shift, four for osmotic stress, 14 for oxidative stress/oxygen limitation, 11 for nutrient limitation, and two for protein stress were covered by the StressChip (Table 1). Also, four sigma factors, one general stress response gene, and one stringent response gene were included in the StressChip (Table 1). A total of 22,855 probes covering 79,628 gene sequences with about 8.0% (1838 probes) as sequence-specific probes and about 92.0% (21,017 probes) as group-specific probes (Table 1) were designed. Of these, 21,689 probes (94.9%) target 985 bacterial, 709 probes (3.1%) target 76 archaeal, and 457 probes (2.0%) target 59 eukaryotic species/strains (Table 2). Percentages of probes and covered CDS sequences in the StressChip are 27.8% and 19.0% for nitrogen limitation, 23.4% and 22.5% for oxygen limitation, 13.6% and 11.5% for phosphate limitation, 14.6% and 20.7% for sigma factors (Figure 1). In addition, internal controls, including degenerate probes for 16S rRNA gene as positive controls and probes for hypothetical genes of hyperthermophiles as negative controls,<sup>14</sup> and 50-mer common oligonucleotide reference standard (CORS)<sup>23</sup> were randomly localized in each array for quality control and data normalization, respectively.

### 3.2. Computational Evaluation of the StressChip.

A computational analysis of the specificity of all probes in terms of sequence identity, stretch length, and free energy of a probe to its nontargets was performed. All probes fell in the desired ranges of sequence identity, stretch length, or free energy with only a small portion of the designed probes close to the thresholds. For gene-specific probes, 6.2% of the probes had 88% identities with their closest nontargets (Figure 2a), 4.8% of the probes with 20 bases of continuous match with nontargets (Figure 2b), and 12.7% of the probes with free energy of –35 to –30 kcal/mol with nontargets (Figure 2c). For group-specific probes, about 99% of the probes had more than 98% sequence identities with group targets including about 95.2% of the probes with 100% sequence identities to group targets (Figure 2d), about 96.9% of the probes had 50-base stretches with the group targets and only 1.4% of the probes had 35–39 continuous match with group targets (Figure 2e), and 98.3% of the probes had free energy less than –65 kcal/mol with group targets and only 1.7% of the probes had



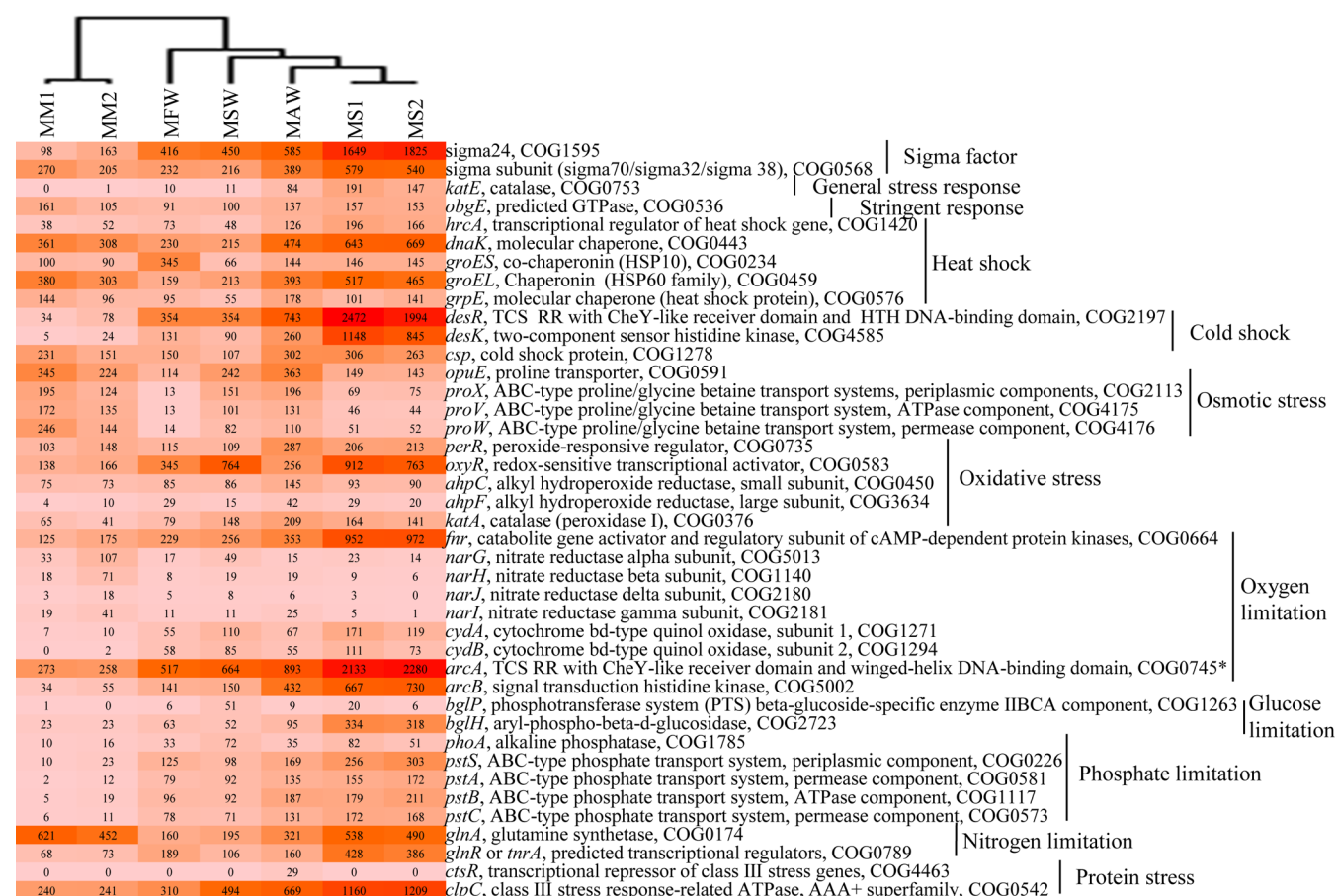


**Figure 2.** Computational evaluation of all probes on the StressChip. Distribution of maximal sequence identity (a), maximal stretch length (b), and minimal free energy (c) of gene-specific probes to their nontargets; minimal sequence identity (d), minimal stretch length (e), and maximal free energy (f) of group-specific probes to their group targets are demonstrated.

free energy of  $-65$  to  $-60$  kcal/mol with group targets (Figure 2f). The results indicate all designed probes are specific to their corresponding targets. Previous experiments demonstrated that all designed probes were highly specific as evaluated with synthesized oligonucleotides and genomic DNA samples.<sup>13,14,17–19,21</sup>

**3.3. Distribution of the Functional Genes in Different Environment Microbial Communities.** Comprehensive analysis of prokaryotic genome sequences has demonstrated

that gene expression regulation and signal integration have been strongly selected to enable rapid adaptation to environmental conditions, and larger genomes harbor more regulation genes.<sup>63,64</sup> To examine whether the stress related functional genes in StressChip are good indicators of stress, the distribution of these genes in microbial communities from different environments was analyzed with seven online available metagenome data sets. Among these samples, marine samples

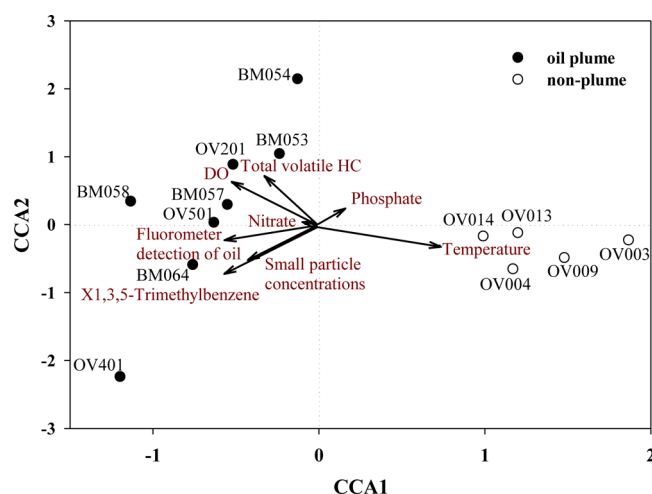


**Figure 3.** Distribution of stress functional genes in metagenome data sets from seven different environmental microbial communities. Numbers of gene counts for each functional gene are shown. Color scale: maximum: red; minimum: pink. Hierarchical cluster of seven microbial communities is generated with Gene Cluster 3.0. MM1, marine community sample line P from station P12 (Pacific Ocean) at a depth of 500 m sampled in June 2009; MM2, marine community sample F 10 from line P, station S103 (Pacific Ocean) at a depth of 100 m sampled in February 2010; MFW, freshwater community from lentic Trout Bog Lake (Epilimnion) sampled on June 7, 2007, Wisconsin; MSW, saline water community from Great Salt Lake South Arm Stromatolite, Utah; MAW, saline and alkaline water sediment community from lentic Soda Lake (CA); MS1, desert soil community from FACE test site NTS\_067, Nevada; and MS2, desert soil community from Dark Crust, Green Butte site, Colorado Plateau.

(MM1 and MM2) represent a relatively stable (not stressful) environment, and nonmarine saline and alkaline water samples (MSW and MAW) and desert soil samples (MS1 and MS2) represent relatively unstable (stressful) environments; a freshwater sample (MFW) may represent a less stressful environment than desert and nonmarine saline and alkaline water samples. As shown in Figure 3, the two marine samples cluster together in one clade; all other samples are in one clade, and two desert samples cluster together, suggesting the distribution of stress functional genes well represents the stressfulness of the environmental communities. Regulatory genes are enriched in communities in more stressful environments. Osmotic stress functional genes are enriched in saline water communities and marine microbial communities, which is consistent with the potential osmotic stress from the high salinity. Nutrient limitation genes are enriched in microbial communities from the barren desert. The abundances of stress functional genes are less in marine microbial communities (MM1, 1.08%; MM2, 0.82%) than others (MFW, 1.50%; MSW, 1.54%; MAW, 1.35%; MS1, 1.66%; MS2, 1.55%). Stress responses have great potential as a controlling factor in shaping microbial communities. We might expect a highly stochastic environment to favor conservation of stress response genes whereas in a stable environment stress response genes tend to be lost. Therefore, the StressChip

has the potential to assess the stress responses of microbial communities from different environments and/or their stability.

**3.4. Application of StressChip on Microbial Community Samples Affected by Oil Contamination.** The Deepwater Horizon oil leak, starting on April 20, 2010 at a depth of 1544 m, released large amounts of oil and methane to the ocean.<sup>65</sup> Previous studies on seawater samples taken from the vicinity of the wellhead after the oil spill demonstrated significant temporal community composition changes.<sup>61,65–67</sup> With StressChip, a total of 1744 probes were detected. More genes (1117 ± 83) were detected in oil plume samples than nonplume samples (852 ± 30). The overall similarities of the stress response functional genes in plume and nonplume samples were low with two clearly separated clusters in the canonical correspondence analysis (CCA) plot (Figure 4,  $p = 0.05$ ). Result of the CCA also indicated that the functional structure of seawater microbial communities was shaped by environmental variables, such as total volatile hydrocarbon (HC), dissolved oxygen (DO), fluorometer detection of oil, 1,3,5-trimethylbenzene, small particle concentrations, and nitrate, which were closely correlated with the plume samples (Figure 4). A similar CCA plot was obtained in a previous GeoChip study of these microbial communities;<sup>62</sup> however, significantly correlated environmental variables were different. For example, several variables such as



**Figure 4.** Canonical correspondence analysis of the relationship between the stress response genes of seawater microbial communities (symbols) and environmental factors (arrows).

1,3,5-trimethylbenzene, small particle concentrations, and nitrate did not appear in the CCA plot of GeoChip data; correlation of total extractable petroleum HC and GeoChip functional genes was not observed here. Significantly more probes for sigma factors, the stringent response gene *obgE*, general stress response *katE* and functional genes involved in heat shock, oxidative stress, oxygen limitation, phosphate limitation, or nitrogen limitation were detected in plume than in nonplume samples (Table S2 and Figure S1, Supporting Information). Such increases may be largely due to a sudden carbon input into the aquatic system, which could stimulate a microbial community shift toward copiotrophs and rapid response to changing environmental conditions by utilizing a range of carbon sources.<sup>68</sup>

A metagenomics study of plume (BM058 and OV201) and nonplume (OV003) communities<sup>66</sup> demonstrated obvious community composition changes within 6 h (OV201) or 39 h (BM058) of exposure to hydrocarbons. *Oceanospirillales* became the dominant species, indicating the rapid response of the marine community to oil spills. In the StressChip data, a total of 14 *Oceanospirillales* probes from 7 species (*Marinomonas* sp. MED121, *Chromohalobacter salexigens* DSM 3043, *Alcanivorax borkumensis* SK2, *Kangiella koreensis* DSM 16069, *Alcanivorax* sp. DG881, *Hahella chejuensis* KCTC 2396, and *Halomonas halodenitrificans*) were detected with  $10 \pm 2$  probes in plume and  $5 \pm 2$  probes in nonplume communities. *Halomonas halodenitrificans* was detected in plume communities only. In contrast to data from 454 pyro-sequencing or clone library sequencing of 16S rRNA genes from plume communities, the relative abundances of the detected probes at the phyla level were similar between plume and nonplume communities except for slight changes in *Actinobacteria* and *Firmicutes* (Figure S2, Supporting Information). Methylotrophs were found in previous metatranscriptomic studies of plume (BM058 and OV201) communities taken in August 2010 and later.<sup>65,66</sup> The probes of four methylotrophs from *Methylococcaceae*, *Methylophaga*, and *Methylophilaceae* were also detected by StressChip in both plume and nonplume communities.

In summary, the consistency of the abundance of stress response genes with habitat conditions and the application of the StressChip to oil spill samples indicated that the StressChip is a powerful, rapid, and cost-effective high-throughput tool for assessing the responses of microbial communities to environmental

stresses. It is well-known that the global environment is changing; information provided by the StressChip could be valuable for decision-making within the context of ecosystem management, environmental cleanup, bioremediation and restoration, environmental engineering, and applied microbiology. Although omics analysis of microbial communities offers the possibility of understanding “who is there” and “what they are doing”,<sup>69</sup> the challenges in data mining and issues associated with random sampling/under sampling still remain.<sup>70</sup> A combination of microarray (e.g., StressChip) and high throughput sequencing (e.g., metagenomics, metatranscriptomics) will allow a comprehensive examination of the complex microbial community responses to environmental changes.

## ■ ASSOCIATED CONTENT

### Supporting Information

Details of the metagenomes used for analysis of distributions of stress function genes and StressChip data of deepwater horizon oil spill plume samples. This information is available free of charge via the Internet at <http://pubs.acs.org>.

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### Notes

The authors declare no competing financial interest.

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