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Transcriptome sequencing and development of an expression microarray platform for liver infection in adenovirus type 5-infected Syrian golden hamsters

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ABSTRACT

The Syrian golden hamster is an attractive animal for research on infectious diseases and other diseases. We report here the sequencing, assembly, and annotation of the Syrian hamster transcriptome. We include transcripts from ten pooled tissues from a naïve hamster and one stimulated with lipopoly-saccharide. Our data set identified 42,707 non-redundant transcripts, representing 34,191 unique genes. Based on the transcriptome data, we generated a custom microarray and used this new platform to investigate the transcriptional response in the Syrian hamster liver following intravenous adenovirus type 5 (Ad5) infection. We found that Ad5 infection caused a massive change in regulation of liver transcripts, with robust up-regulation of genes involved in the antiviral response, indicating that the innate immune response functions in the host defense against Ad5 infection of the liver. The data and novel platforms developed in this study will facilitate further development of this important animal model.

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Introduction

The Syrian golden hamster has been used broadly in biomedical research for decades, especially as a model for cancer development. More recently, it has been served as models in research on diabetes (Bhathena et al., 2011), atherosclerosis (Dillard et al., 2010; Jove et al., 2013), and neural plasticity (Stafford and Meisel, 2012). Also, the Syrian hamster is frequently employed in studies of infectious disease, as it is highly susceptible to infection with a wide range of viruses, bacteria, and parasites (Espitia et al., 2010; Zivcec et al., 2011). The Syrian hamster has been used to study emerging and highly pathogenic RNA viral infections and virus-induced diseases, including flaviviruses (West Nile virus, Japanese encephalitis virus, St. Louis encephalitis virus, Yellow fever virus), alphaviruses (Venezuelan-, Western-, Eastern equine encephalitis virus), henipavirus (Nipah virus), arenaviruses (Pirital virus, Pichinde virus), bunyaviruses (Rift valley fever virus, Punta Toro virus, Andes virus), filoviruses (Ebola virus, Marburg virus), and SARS-coronavirus (Ebihara et al., 2013; Gowen and Holbrook, 2008; Holbrook and Gowen, 2008; Nakayama and Saijo, 2013; Roberts et al., 2005). Further, the Syrian hamster has proved to be a useful model to evaluate vaccines and anti-viral drugs against both RNA and DNA viruses (Julander et al., 2007, 2011; Monath

et al., 2010; Morrey et al., 2004, 2008; Toth et al., 2008; Wold and Toth, 2012).

Still further, the Syrian hamster has served as a model to study human adenovirus (Ad) pathogenesis and to evaluate oncolytic Ad vectors in cancer gene therapy (Bortolanza et al., 2007; Cerullo et al., 2012; Dhar et al., 2009a, 2009b, 2012, 2014; Lichtenstein et al., 2009; Spencer et al., 2009; Thomas et al., 2006, 2007, 2008; Toth et al., 2007; Wold and Toth, 2012; Ying et al., 2009a; Young et al., 2013a, 2013b). Human Ads are double-stranded DNA viruses with a non-enveloped icosahedral capsid. There are approximately 60 types (previously referred to as serotypes) that are classified into seven species from A to G (Berk, 2013). In immunocompetent healthy individuals, most Ad infections are generally mild and self-limiting, resulting in infections in the upper respiratory tract, gastrointestinal, and urinary tracts (Wold and Ison, 2013). Some serotypes can cause serious ocular illness such as epidemic keratoconjunctivitis. In immunocompromised individuals, especially in pediatric hematopoietic stem cell transplant patients, Ads can cause severe disseminated multi-organ infections which are sometimes lethal (Echavarria, 2008; Ison, 2006; Lion, 2014; Matthes-Martin et al., 2012, 2013; Sandkovsky et al., 2014; Stercz et al., 2012; Wold and Ison, 2013).

Syrian hamsters are permissive for human Ads (reviewed in Wold and Toth (2012)). Ad5 infection following intravenous (i.v.) administration results in the hamster's body weight loss, elevated serum liver enzyme levels, and virus replication in the liver, lungs, and other

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organs (Lichtenstein et al., 2009; Ying et al., 2009). When the hamster is immunosuppressed using high-dose cyclophosphamide, Ad5 replication continues for extended periods in multiple organs, especially the liver, resulting in increased mortality (Thomas et al., 2008; Toth et al., 2008). This extended replication in hamsters recapitulates human Ad infection in immunocompromised patients.

A major limitation for using the Syrian hamster as an animal model for infectious disease is the lack of information on the hamster genome and its transcription. The Syrian hamster genome has just been sequenced and released but has not been annotated (NCBI Bio Project 77669). The transcriptome of Chinese hamster ovary and Syrian baby hamster kidney cell lines as well as hamster brain and liver tissue have been analyzed (Johnson et al., 2014; Schmucki et al., 2013; Vishwanathan et al., 2015). More recently, the Syrian hamster transcriptome with additional tissues from brain, lung, spleen, kidney, heart, and liver has been published (Tchitchek et al., 2014), but the complete global transcriptome representing whole hamster transcripts, especially genes engaging in immune responses, has not been documented. Quantitative reverse transcriptase PCR (qRT-PCR) to detect and quantify some cytokines and chemokines has been published, but to our knowledge, microarrays specific to global hamster genes have not been developed due to the insufficiency of genetic data.

We report here the sequencing, assembly, and annotation of the Syrian hamster whole transcriptome, together with the generation of hamster microarrays. By using these new platforms, we describe the first global transcriptional analysis of gene expression in the liver of Ad5-infected Syrian hamsters. Our results provide unique insights into acute Ad5 infection of the liver, an organ often infected in disseminated Ad disease in humans.

Results

Hamster transcriptome sequencing and annotation

Our goal was to produce a comprehensive catalog of transcripts expressed in the adult hamster. To this end, we isolated RNA from 10 organs from a naïve animal and a hamster treated with LPS. The organs were chosen because they are the targets of many viruses (e.g. liver) or these organs are complex (e.g. brain) in that they are composed of many cells types and thus express many genes. LPS treatment was included to increase the transcription of genes involved in immune responses, as would be seen in microbiological infections. Pooled RNA was subjected to library construction for deep sequencing.

Illumina HiSeq sequencing produced 5.1×10^{10} raw 100 bp paired-end reads (Table 1). Raw Illumina RNA-Seq data were adapter trimmed, validated, and filtered based on quality. Sequences that passed validation and quality checks were assembled via the *de novo* RNA-seq assembler Trinity (Grabherr et al., 2011; Haas et al., 2013). Trinity assembled contigs were validated via ESTscan and custom software, resulting in a set of 49,176 high-confidence transcripts. These high-confidence transcripts denote the assembled transcriptome. Of the 49,176 transcripts in the assembled transcriptome, 44,705 sequences (91%) had at least one BLASTP hit, and 30,956 (63%) had at least one InterProScan hit. The raw reads, as well as the assembled transcripts, have been deposited in GenBank under BioProject ID PRJNA285471.

Generation of custom hamster microarray and analysis

To assess the genome-wide response to Ad5 infections of the liver, two groups of hamsters were administered i.v. with either 10^{10} pfu of Ad5 or PBS (mock-infection). Ad5 replicates preferentially in the liver

Table 1
Analysis of transcriptome by RNA-seq.

A. Tissue sampled	B. Transcriptome	
Blood	Total raw read (100bp paired end)	5.1×10^{10}
Bone Marrow	Total number of validated	49,176
Brain	Transcripts	
Kidney	Total number of unique transcripts	42,707
Liver	Total number of unique genes	34,191
Cervical lymph node		
Lung		
Muscle		
Spleen		
Thymus		

A. Each tissue was sampled from two hamsters; one treated with LPS and the second was untreated. B. Pooled RNA from both hamsters was sequenced on Illumina HiSeq platform. Summary statistics for raw reads, number of validated transcripts, unique transcript (eliminating transcripts with alternative start site or 5' end) and total number of genes (based on BLASTP to rat genome).

of Syrian hamsters following i.v. administration (Ying et al., 2009). Liver RNAs were extracted 18 h post-infection (p.i.) and gene expression was analyzed by a custom designed microarray. Two biological replicates (independent infections) were performed and each replicate was analyzed in triplicate (Ad5-infected was labeled with Cy3 and mock was labeled with Cy5). A dye-swap experiment was also performed to correct for any dye bias, for a total of four technical replicates per biological replicate (one slide). The data were analyzed using an ANOVA model as previously described (Churchill, 2004; Kiesel et al., 2007).

A total of 7110 genes in the liver were differentially expressed with a $p \leq 0.01$. Of these, 3788 were upregulated and 3322 were down-regulated in response to Ad5 infection. All significantly differentially expressed genes were further subjected to Gene Ontology analysis (GO) to identify functional categories of change in gene expression. As shown in Tables 2A and 2B, transcripts in a number of critical function categories are over-represented. Most notably, genes involved in immune response (57 out of 64 genes (57/64)), defense response to virus (16/17), and antigen processing and presentation (27/31) were largely upregulated in the liver, although some were downregulated (Table 2A). Also largely upregulated were genes associated with regulating DNA-dependent transcription (82/119), signal transduction (76/109), cell adhesion (18/18), and cell communication (16/21) (Table 2A). In contrast, when the most down-regulated genes were investigated for GO function, there was over-representation of genes associated with oxidation–reduction process (214/249), metabolic process (82/101), carbohydrate metabolic process (45/69), DNA topological change (16/21), protein translation (15/18), and tricarboxylic acid cycle (7/7) (Table 2B). Differentially regulated genes are also overrepresented in the GO categories of peptidoglycan catabolic process and transmembrane transport. In the GO category of peptidoglycan catabolic process, 4 genes were upregulated and 4 genes were downregulated (p value = $1.17e-9$). In the GO category of transmembrane transport, 11 genes were upregulated and 12 genes were downregulated (p value = $1.10e-08$).

Table 3 shows a more detailed depiction of differentially expressed genes in the GO category of anti-viral responses: 16 out of 17 genes were upregulated, including toll-like receptor 7 and 2, IL-6, and interferon-inducible proteins such as OAS family members and PKR.

Validation of microarray data by qRT-PCR

To validate the differential gene expression profiles obtained by microarray analysis, qRT-PCR was performed. First, we confirmed the suitability of four housekeeping genes, namely PPI, RPS6KB1, HBP2, and GAPDH. These genes range from low, medium–low, medium, and

Table 2A

Gene Ontology categories with genes largely up-regulated.

GO Category	Number	# of Genes Up/Down regulated	Frac.	P value	Go term
GO:0006355	119	82/37	2.14	2.20E–16	Regulation of transcription, DNA-dependent
GO:0007165	109	76/33	1.96	2.20E–16	Signal transduction
GO:0006955	64	57/7	1.15	2.20E–16	Immune response
GO:0019882	31	27/4	0.56	2.20E–16	Antigen processing and presentation
GO:0051607	17	16/1	0.31	4.58E–16	Defense response to virus
GO:0006952	9	8/1	0.16	8.30E–10	Defense response
GO:0007155	18	18/0	0.32	1.16E–09	Cell adhesion
GO:0007154	21	16/5	0.38	1.91E–08	Cell communication
GO:0016192	20	14/6	0.36	2.45E–08	Vesicle-mediated transport

Table 2B

Gene Ontology categories with genes largely down-regulated.

GO Category	Number	# of Genes Up/Down regulated	Frac.	P value	Go term
GO:0055114	249	35/214	4.47	2.20E–16	Oxidation–reduction process
GO:0008152	101	19/82	1.81	2.20E–16	Metabolic process
GO:0005975	69	24/45	1.24	2.20E–16	Carbohydrate metabolic process
GO:0006265	21	5/16	0.38	2.20E–16	DNA topological change
GO:0006418	18	3/15	0.32	3.36E–11	tRNA aminoacylation for protein translation
GO:0046854	11	4/7	0.2	4.17E–09	Phosphatidylinositol phosphorylation
GO:0048015	10	3/7	0.18	1.43E–08	Phosphatidylinositol-mediated signaling
GO:0006099	7	0/7	0.13	1.14E–07	Tricarboxylic acid cycle

Table 3

Regulation of genes involved in viral response.

Locus ID	Gene	Fold change (mean ± std. dev.)
locus_03211	Toll-like receptor 7	10.3 ± 0.197
locus_05992	Toll-like receptor 2	6.5 ± 0.126
locus_11815	Mitochondrial antiviral-signaling protein-like	–1.5 ± 0.056
locus_06466	2'-5'-oligoadenylate synthase 2-like	11.9 ± 0.226
locus_11667	2'-5'-oligoadenylate synthase 1A-like	11.9 ± 0.074
locus_11853	59 kDa 2'-5'-oligoadenylate synthase-like protein-like	4.1 ± 0.067
locus_16812	2'-5'-oligoadenylate synthetase 3-like	22.6 ± 0.146
locus_29063	2'-5'-oligoadenylate synthetase 3	7.6 ± 0.174
locus_41602	2'-5'-oligoadenylate synthase 1-like	23.3 ± 0.017
locus_41730	2'-5'-oligoadenylate synthase 3-like	5.3 ± 0.151
locus_48794	2'-5'-oligoadenylate synthase 1-like	1.5 ± 0.033
locus_32403	Interferon-inducible protein 10	35.9 ± 0.042
locus_15775	Interferon-activable protein 204	22.2 ± 0.034
locus_25036	IL-6	6.1 ± 0.104
locus_06991	Antigen-presenting glycoprotein CD1d1-like	6.5 ± 0.126
locus_23284	Proteoglycan 3-like	6.1 ± 0.094
locus_11801	Interferon-induced, double-stranded RNA-activated protein kinase	26.5 ± 0.093

high regarding the abundance of their expression, and their expression was unaffected by Ad5 infection. Therefore, these genes were appropriate to serve as endogenous controls. We then examined the relative expression of a few selected target genes which are differentially expressed as found in our microarray analysis. The selected target genes for validation covered a full expression abundance spectrum, ranging from low (prostaglandin G/H synthase 1 precursor, PTGS; 2'-5'-oligoadenylate synthase 3, OAS3); medium–low (eukaryotic translation initiation factor 2 alpha kinase2, PKR); medium (interferon-inducible protein 10, ICP10; prostaglandin reductase 1, PTGR1), and high (interferon-induced protein with tetratricopeptide repeat3-like, IFIT3; MHC-II β chain, MHCII β). These selected targets also represent genes involved in the inflammatory response (PTGR1, PTGS1), the immune response (MHC II), interferon-inducible genes (IFP10, IFIT3), and interferon-inducible antiviral effector genes (PKR, OAS3).

The overall results of qRT-PCR were in agreement with the microarray analysis (Fig. 1); less than two-fold differences were obtained between the microarray and qRT-PCR data in five out of

seven genes tested. Two genes (OAS3, IFIT3) showed 8–9 fold differences between these two types of assays, which probably reflects the intrinsic sensitivity differences between these two techniques.

The number of Class II Histocompatibility Antigen (MHC II) positive cells is increased in the liver of Ad5-infected hamsters

To assess whether the increase for the mRNA levels seen for MHC II translated into increases at the protein levels, we assayed by flow cytometry single-cell suspensions of livers collected from mock-infected and Ad5-infected animals. We found that there was a significant elevation in the number of MHC II positive cells in the liver of Ad5-infected hamsters compared to mock-infected animals (Fig. 2). The mean fluorescence intensity of the MHC II positive cells also increased approximately 3-fold. These data indicate that the elevation in the MHC II mRNA levels detected by the microarray and qRT-PCR were a good indication of the actual protein levels of MHC II in the sample.

Discussion

Global transcriptome representing whole Syrian golden hamster transcripts

In this report, we extend previously described Syrian hamster transcriptome data (see Introduction) and we provide novel information on the pathogenesis of Ad5 in a permissive animal model. We present the sequence of the hamster transcriptome, the construction of a unique hamster microarray, and the characterization of the global transcriptional response in the liver of hamsters to i.v. infection with Ad5. Our transcriptome covers hamster transcripts derived from ten pooled tissues. Our study has increased the number of non-redundant transcripts for the Syrian hamster from 11,648 to 42,707, representing 34,191 unique genes, resulting in 4 × increased gene discovery in our data set. The larger tissue sampling combined with LPS treatment and deeper sequencing all contributed to the increased gene discovery.

Systemic adenovirus infection causes a massive change in regulation of the hamster liver transcriptome

Investigations into interactions among Ad and specific cellular proteins and pathways in cell culture have provided insights about Ad and host responses. Genome-wide gene expression analyses have been performed by several laboratories using different cell lines, cell culture conditions, cells (non-cancerous *versus* cancer; proliferating or quiescent), wild-type Ad, various replication-competent and replication-defective vectors, and different conditions such as time post-infection and multiplicity of infection. Several microarray studies were conducted with Ad5 or the closely related Ad2 in non-cancerous human cells, namely in IMR-90 cells (a normal diploid cell derived from lungs of a 16 week female fetus) (Granberg et al., 2006; Stilwell and Samulski, 2004; Zhao et al., 2007), WI-38 cells (diploid cell line derived from normal embryonic lung culture) (Rao et al., 2006), synchronized human foreskin fibroblasts (Miller et al., 2007), or normal human bronchial epithelial cells (Dorer et al., 2011). Then, more recently, Ad2-induced transcriptome changes were studied in detail in IMR-90 cells using a sensitive cDNA deep sequencing strategy (Zhao et al., 2012). In general, these studies depict a struggle between the virus and the host cell in which the virus assumes control of the cell and the cell fights back with anti-viral defenses. The response begins within minutes of the virus interacting with the cell receptors and continues throughout the periods of virus early gene expression, viral DNA replication and early-late gene expression, and very late gene expression and virus assembly. Three of the microarray studies identified 988 (Zhao et al., 2007), about 2000 (Miller et al., 2007), or 943 (424 up, 519 down) (Dorer et al., 2011) cellular genes that were differentially up- or down-regulated by a factor of 2, and the RNA sequencing approach found 3791 genes that were differentially regulated (Zhao et al., 2012). Of these genes, it's not clear whether the changes are caused by the action of the virus directly, e.g. through the activity of the E1A transcriptional co-regulator, or by the cell "sensing" that it has been infected. Altogether, these studies revealed that Ad infection leads to induction of genes involved in cell cycle regulation (activation), cell proliferation, chromatin reorganization, DNA replication, RNA transcription and processing, nucleotide metabolism, and the Wnt signaling pathway. Some but not all genes whose promoters contain binding sites for the E2F transcription factor were conspicuously up-regulated (Miller et al., 2007; Zhao et al., 2012). When quiescent human foreskin fibroblasts were infected, there was a reversal of the quiescent state similar to that seen in the cellular response to serum (Miller et al., 2007). In contrast to the up-regulated genes, there is repression of genes involved in cell differentiation, various signaling pathways (TGFβ, Rho, G-protein, Map kinase, STAT, NFκB), cell death regulation, and metabolism (Dorer et al., 2011; Zhao

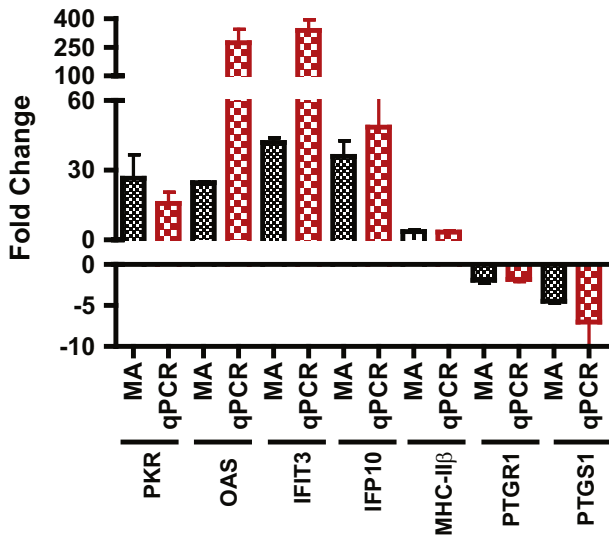


Fig. 1. Validation of Microarray (MA) data of selected genes by qRT-PCR. The mRNA expression levels of various genes were determined by qRT-PCR to validate the expression data of the microarray analysis. The fold change represents the mRNA expression level in Ad5-infected hamsters over that of untreated hamsters. For qRT-PCR (red bar), the value represents mean \pm SD of three biological replicates without pooling RNA. For microarray (MA, black bar), the value represents the average of two biological replicates.

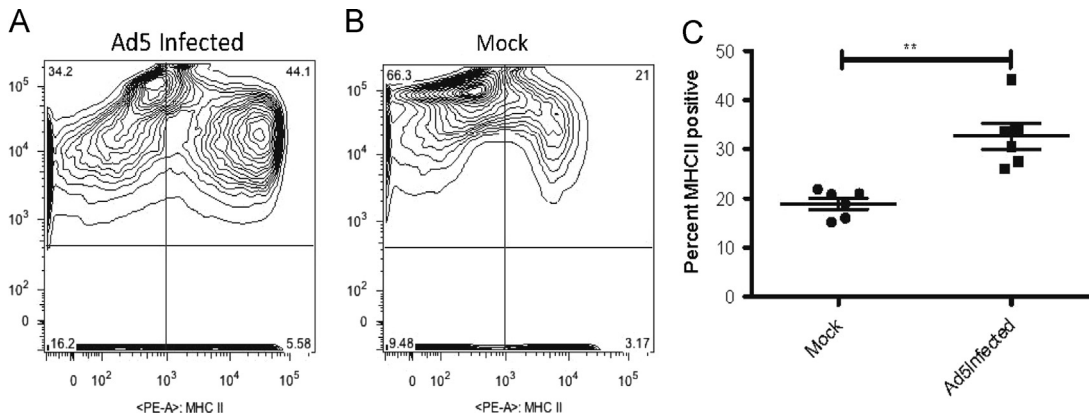


Fig. 2. The number of MHC II positive cells is elevated in the liver of Ad5-infected hamsters. Combined results from two independent experiments are shown. A, B: Flow cytometry data from two individual animals. C: The symbols represent data from individual animals, horizontal bars symbolizes the mean, and the whiskers signify the standard deviation. ** $p < 0.01$.

et al., 2012). Transcriptional activity associated with the p53 transcription factor was suppressed (Miller et al., 2007).

Another study in IMR-90 cells employed chromatin immunoprecipitation combined with microarray (ChIP–chip) technology to analyze global changes in gene expression at early (6 h), intermediate (12 h), and late times p.i. with h5dl1500, an Ad5 mutant that expresses only the 243 amino acid (243R) version of E1A protein (Ferrari et al., 2008), see review by Ferrari et al. (2009). Interaction of the conserved region 1 (CR1) and conserved region 2 (CR2) domains of the E1A 243R protein with RB family proteins leads to the release of free E2F family members to activate (or, with some E2F members, to repress) transcription of target genes. Also, interaction of the 243R N-terminus with histone acetyltransferase proteins (p300/CBP) regulates chromatin structure by altering the global pattern of histone modification, in particular H2K18ac, an epigenetic marker for actively transcribed genes (Ferrari et al., 2012; Horwitz et al., 2008). Ferrari et al. (2008) found that transcription of ~70% of cellular genes was altered in cells infected by H5dl1500 and that these changes were linked to E1A binding to RB members and p300/CBP. In brief, at early times p.i., E1A caused induction of genes that function in cell cycle regulation, and at late times (24 h p.i.) E1A activity led to repression of genes that function in development, differentiation, and cell signaling. Cell genes involved in defense against virus infection were induced at 6 h but repressed by 24 h p.i. The results of Ferrari et al. (2008) are generally consistent with the gene profiling data discussed earlier and they appear to make sense: cell cycle and DNA replication genes needed by the virus are induced and antiviral defense genes and cell differentiation genes that are harmful or not needed by the virus are repressed.

In addition to non-cancerous cell lines, microarray gene profiling studies have been conducted in several cancer cell lines infected by wild type Ad or Ad vectors, namely HeLa (Granberg et al., 2005; Zhao et al., 2003), Huh7 (Martina et al., 2007), and melanoma (Dorer et al., 2011; Volk et al., 2005) cells. In general, in these cells Ad infection affected genes involved in cell growth arrest, cell metabolism, and antiviral defense. Interestingly, fewer genes were dysregulated in cancer cells than in normal cells, consistent with the idea that infection of normal cells by Ad induces a cancer cell-like state (Ferrari et al., 2009, 2008; Horwitz et al., 2008).

A very recent study used microarray technology to examine cellular transcription in response to induction of the Ad2-coded E4orf3 protein in cells stably transfected with the gene for E4orf3 (Vink et al., 2015). E4orf3 functions in Ad replication by causing cellular proteins to re-localize into filamentous nuclear inclusions termed “tracks”. Multiple cellular proteins that are re-localized into these tracks function in the DNA damage response, the interferon response, and cellular transcription. Upon induction of E4orf3 in the transfected cells, more than 400 cellular genes were differentially regulated; 70% of the genes were upregulated and 30% were downregulated. With at least 8 of these genes, dysregulation was linked to nuclear track formation, i.e. physical relocation of cellular proteins into the tracks (Vink et al., 2015). Many of the genes with altered expression influence cellular growth properties and a variety of the proteins coded by these genes localize to the extracellular space or plasma membrane. Some genes affected by E4orf3 are also involved in antiviral activity.

Microarray based gene expression analysis of Ad-induced transcriptome changes have also been documented in a mouse cell culture system (mouse embryo fibroblasts, MEFs) as well as in non-permissive C57BL/6 J mice following systemic infection with a replication-defective E1- and E3-minus Ad5-based vector (Hartman et al., 2007, 2008). While these studies revealed differences in the expression of genes belonging to particular function groups, these studies only

analyzed the host transcriptome response in a non-permissive host for Ad infection, and did not address the role of viral replication and viral late gene expression in modulating host responses.

As discussed earlier, the liver is the major site of Ad5 replication in Syrian hamsters following i.v. administration of Ad5 (Lichtenstein et al., 2009; Thomas et al., 2008; Toth et al., 2008; Ying et al., 2009). Accordingly, we developed a hamster expression microarray to investigate the transcriptional response in the liver to systemic Ad5 infection. Our analysis revealed that approximately 16% of total unique transcripts were differentially expressed, indicating an extensive change in liver transcription. Functional analysis using GO categories revealed that genes involved in the immune response, defense response to virus, and antigen processing and presentation were largely upregulated (Table 2A). The same is true with genes associated with regulating DNA-dependent transcription, signal transduction, cell adhesion, and cell communication (Table 2A). Cell adhesion and cell-cell communication are critical in immune response networks where various lymphocytes, antigen-presenting cells, and antigen interact actively. Up-regulation of genes in these categories is further indication of an activated immune response to virus infection. The most down-regulated genes were associated with oxidation–reduction process, metabolic process, carbohydrate metabolic process, DNA topological change, protein translation, and tricarboxylic acid cycle (Tables 2B). Our results are in good agreement with the microarray analysis of gene expression in the liver of C57BL/6 J mice infected i.v. with a replication-defective Ad5 vector (Hartman et al., 2007). The latter study found that up-regulated genes are over-represented in categories involving the defense response, immune response, antigen presentation, signal transduction, and RNA processing, whereas down-regulated genes are over-represented in categories associated with metabolism, oxidoreduction, chromatin maintenance, and remodeling.

Robust up-regulation of genes involved in antiviral response

Taking a closer examination of differentially expressed genes involved in anti-viral responses, receptors engaging in the inflammatory immune response such as toll-like receptor 2 and 7 were upregulated (Table 3). Other genes associated with the inflammatory response such as proteoglycan 3 and cytokine IL-6 were also upregulated. Most notably, a cluster of genes including PKR and oligoadenylate synthase family members OAS1-3 were upregulated. PKR and OAS are two major IFN-induced effectors demonstrated to possess antiviral activities against various viruses through different mechanisms of action. Following activation by viral double-stranded RNA (dsRNA), OAS synthesizes 2',5'-oligoadenylates that in turn activate RNase L, which in turn triggers cellular and viral RNA cleavage. IFN-induced PKR, following binding of dsRNA, inhibits cellular and viral protein translation by phosphorylation of eukaryotic translation initiation factor 2 α (Randall and Goodbourn, 2008; Sadler and Williams, 2008).

Our data imply that innate immune responses mediated by activation of toll-like receptors, induction of proinflammatory cytokines and IFN pathways provide a first line of host defense against systemic Ad5 in the liver at the early encounter of infection, and that innate immune responses play a crucial role in controlling virus infection, likely mediated through the action of Kupffer cells. We have shown that the number of MHC II positive cells increased in the liver of Ad5 infected hamsters, and that these cells up-regulate the surface expression of MHC II, indicating up-regulation of the antigen presentation process. These flow cytometry data corroborate data obtained using microarray and qRT-PCR. This elevation in the number of MHC II positive cells in the liver is somewhat unexpected, inasmuch as Ad5 is known to preferentially infect and kill Kupffer cells, the major MHC II positive cell population in the liver, within minutes of intravenous injection (Smith et al., 2008). Further, the Ad E1 A protein

down-regulates the expression of MHC II RNA (Kretsovali et al., 1998). Clearly, further investigation is needed to discover the nature of the MHC II positive cells in the liver of Ad5-infected hamsters.

In summary, our current study significantly expanded the number of non-redundant transcripts of the Syrian hamster transcriptome to 42,707, representing 34,191 unique genes, and generated a microarray platform for global transcription analysis. By using these new platforms, we describe the first global transcriptional analysis of gene expression in the liver of Ad5-infected Syrian hamsters. We anticipate that the novel platforms developed in this study will facilitate further development of new tools for this important animal model, and will be beneficial to the whole hamster research community.

Materials and methods

Animals

Female Syrian hamsters (*Mesocricetus auratus*) were obtained from Harlan Laboratories, Indianapolis, IN. All studies were approved by the Institutional Animal Care and Use Committee of Saint Louis University and were conducted according to federal and institutional regulations.

RNA extraction

To extract RNA for transcriptome sequencing, two female Syrian hamsters (5–6 weeks old), one untreated and one treated intraperitoneally with lipopolysaccharide (LPS) for 18 h, were anesthetized with CO₂ and then sacrificed. Whole blood was collected by cardiac puncture, and bone marrow was flushed from the femur. Both blood and bone marrow samples were placed into RNA Protect Animal Blood tubes (Qiagen, Valencia, CA). Eight different organs were collected and frozen immediately in liquid nitrogen (Table 1). Total RNAs from blood and bone marrow were extracted using RNeasy Protect Animal Blood kit (Qiagen). RNA was extracted from all organs except liver by homogenizing the samples in Trizol reagent and then extracting the RNA from the homogenate. To extract liver RNA the sample was homogenized in RNALater buffer and the homogenate was processed using the RNeasy mini kit (Qiagen). All RNA samples were treated with RNase-free DNase followed by RNA cleanup to eliminate DNA contamination. RNA integrity was determined using a 2100 Bioanalyzer (Agilent Technologies, Santa Clara, CA), while the RNA yield was determined on a NanoDrop-2000 spectrophotometer. To obtain a diverse representation of the whole transcriptome, total RNAs from all the organs were pooled from an untreated and a LPS-treated hamster (Table 1). The pooled RNA quality, as assessed by Bioanalyzer (Agilent), had a RNA integrity number (RIN) of 8.5.

To extract RNA for microarray analysis, four female hamsters were divided into two groups; two hamsters in each group were administered i.v. with either phosphate buffered saline (PBS) or 10¹⁰ plaque forming units (pfu) of Ad5 in a total volume of 200 μ l. Hamsters were sacrificed 18 h later and liver RNA from each hamster was extracted using the RNeasy mini kit.

To extract RNA for microarray validation, two groups of three hamsters each were treated with PBS or Ad5 i.v. and then liver RNA was extracted as described as above.

Library construction and sequencing

Approximately 450 ng (quantified by Qubit; Life Technologies) of pooled RNA was used as input for library construction. For sequencing, a library was generated using Illumina TruSeq RNA preparation kit version 2 following the manufacturer's

directions. The library was sequenced on Illumina HiSeq to obtain 100 base pair (bp) paired-end reads.

Sequence assembly and annotation

Raw Illumina RNAseq data were adapter trimmed, validated, and filtered based on quality. Quality-passed data were assembled via the *de novo* RNA-seq assembler Trinity (Grabherr et al., 2011; Haas et al., 2013). Trinity assembled contigs were validated via ESTscan and custom scripts, resulting in a set of 49,176 high-confidence transcripts. These high-confidence transcripts denote the assembled transcriptome. Each protein sequence from the assembled transcriptome was annotated using BLASTP versus the NCBI/Genbank NR protein database, and using the protein domain/motif prediction program InterProScan. BLASTP matching against protein database typically allows for better annotations because wobble positions and sequencing errors are eliminated in the alignment. The sequencing and summary statistics are shown in Table 1.

Microarray design and analysis

While RNA-Seq is likely to replace microarrays eventually, microarrays are still used because they are highly time and cost-effective and because methods of normalization and analysis of data are more robust. In our hands two-color microarrays perform as well, if not better than RNA-Seq. Therefore, we designed microarrays based on the hamster transcriptome. The 60-mer probes were designed using the high-confidence transcripts. Briefly, each transcript was divided into non-overlapping (contiguous) 60 nucleotide oligonucleotide (oligos). We calculated four parameters for each oligo: uniqueness, sequence complexity, self-annealing and melting temperature (T_M). The uniqueness parameter is the binding energy (Allawi and SantaLucia, 1998) calculated using the sequence of the oligo to the reverse-complementary sequence and to its closest match identified by BLASTN against the high-confidence transcripts. Sequence complexity is calculated using the Ziv and Lempel algorithm (Ziv and Lempel, 1977). The T_M is calculated using equation and parameters as previously described (Bolton and Mc Carthy, 1962). The designed probes were submitted to Agilent for printing of the arrays in the 8 \times 60K format, where each slide has eight identical arrays with 61,657 60-mer-oligonucleotide probes.

Quantitative reverse transcriptase RT-PCR (qRT-PCR)

For qRT-PCR, 2 μ g of each RNA and 50 pmol of oligo (dT) primer were used for *in vitro* reverse transcription (RT) using the High Capacity cDNA Reverse Transcription kit (ABI, Foster city, CA). SYBR Green based qPCR was used to specifically detect target gene mRNA. The primers were designed using Primer 3 software and synthesized by Integrated DNA Technologies (Coralville, IA). The sequence of the primers is shown in Supplementary Table S1. The PCR was set up in a 20 μ l volume containing 1x SYBR[®] select master mix (ABI), 250 nM forward and reverse primers, and 2 μ l of the diluted RT template. Quantification was done in triplicate for each sample using an ABI model 7500 genetic analyzer with the following cycling parameters: 1 cycle at 50 °C for 2 min, 1 cycle at 95 °C for 10 min, followed by 40 cycles of 95 °C for 15 s and 60 °C for 1 min. Dissociation curve analysis was performed following PCR to verify the specificity of the amplified product.

The data were analyzed using the $\Delta\Delta C_t$ method. Four house-keeping genes were used as endogenous controls for normalization, namely peptidyl-prolyl cis-trans isomerase-like 1 (PPI), ribosomal protein S6 kinase polypeptide 1 (RPS6KB1), histone-binding protein RBBP7 isoform2 (HBP2), and glyceraldehyde-3-

phosphate dehydrogenase (GAPDH). Briefly, the Ct of each target gene in a treated hamster was first normalized to the Ct of the endogenous control (Δ Ct) and then compared to the same normalized gene in a mock-treated (calibrator) hamster to determine the $\Delta\Delta$ Ct. The final value is displayed as the relative fold change between the Ad5-infected and mock-treated hamsters.

Flow cytometry

Syrian hamsters were mock-infected or infected i.v. with 1.2×10^{10} pfu of Ad5 ($n=3$ for both groups; the experiment was repeated once). At 18 h post injection, the animals were sacrificed, and the liver was perfused with PBS. The liver was harvested, and the left lateral lobe was homogenized by forcing it through a 100 μ m cell strainer (Fisher Scientific, Waltham, MA). The cell suspensions were washed using PBS. The cells were incubated with Live/Dead Fixable Aqua cell stain (Life Technologies, Grand Island, NY), washed twice with FACS buffer (PBS containing 2% fetal bovine serum), and then re-suspended in FACS buffer containing a 1:2000 dilution of PE-conjugated anti-mouse/rat MHC class II (I-Ek; clone 14-4-4 S). After 1 h incubation, the cells were washed twice with PBS, fixed in 2% paraformaldehyde solution for 5 min, washed in PBS, re-suspended in FACS buffer, and analyzed on a BD FACS LSRII analyzer.

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Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.virol.2015.07.024>.

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