

# Transcript Profiling of Functionally Related Groups of Genes During Conditional Differentiation of a Mammalian Cochlear Hair Cell Line

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We have used Affymetrix high-density gene arrays to generate a temporal profile of gene expression during differentiation of UB/OC-1, a conditionally immortal cell line derived from the mouse cochlea. Gene expression was assessed daily for 14 days under differentiating conditions. The experiment was replicated in two separate populations of cells. Profiles for selected genes were correlated with those obtained by RT-PCR, TaqMan analysis, immunoblotting, and immunofluorescence. The results suggest that UB/OC-1 is derived from a population of nonsensory epithelial cells in the greater epithelial ridge that have the potential to differentiate into a hair-cell-like phenotype, without the intervention of *Math1*. Elements of the Notch signaling cascade were identified, including the receptor *Notch3*, with a transient up-regulation that suggests a role in hair cell differentiation. Several genes showed a profile similar to *Notch3*, including the transcriptional co-repressor *Groucho1*. UB/OC-1 also expressed *Mel1*, a putative partner of *Math1* that may confer competence to differentiate into hair cells. Cluster analysis revealed expression profiles for neural guidance genes associated with *Gata3*. The temporal dimension of this analysis provides a powerful tool to study genetic mechanisms that underlie the conversion of nonsensory epithelial cells into hair cells.

[The entire data set published in this paper has been deposited in the National Center for Biotechnology Information (NCBI) Gene Expression Omnibus Database (<http://www.ncbi.nlm.nih.gov/geo/>) under the series accession no. GSE36 and sample numbers contained therein. Supplementary material is available online at <http://www.genome.org>. The following individuals kindly provided reagents, samples or unpublished information as indicated in the paper: T. Hasson, C. Petit, and P. Matsudaira]

We sought to discover gene expression patterns associated with the early differentiation of mammalian auditory hair cells. We used a conditionally immortal cell line (UB/OC-1) selected from the cochlea of the Immortmouse at embryonic day 13 (E13), to represent the time at which hair cell progenitors cease proliferation and start to differentiate (Rivolta et al. 1998), and Affymetrix oligonucleotide arrays to assess gene expression in the cells during differentiation *in vitro*. The cell line UB/OC-1 was derived from Immortmouse embryos and can be induced to differentiate in a conditional manner. These mice have an immortalizing transgene, the temperature-sensitive variant *tsA58*, regulated by a  $\gamma$ -interferon-inducible promoter. When the cells are cultured at 33°C in the presence of  $\gamma$ -interferon, proliferation is maintained, but following removal of  $\gamma$ -interferon from the culture media and an increase in temperature to 39°C, proliferation ceases and the cells start to differentiate (Jat et al. 1991).

Synthetic oligonucleotide arrays and cDNA arrays have been used to analyze gene expression in a wide range of cells and tissues (Alon et al. 1999; Mochii et al. 1999; Perou et al. 2000; Ross et al. 2000). Although the technique is relatively simple (Lockhart et al. 1996; Kozián and Kirschbaum 1999;

Lipshutz et al. 1999), the experimental design and methods of data analysis are critical. Clonal cell lines are valuable because the genetic programs within a cell are highly ordered and this is reflected in the coordinated expression of functionally related genes (Niehrs and Pollet 1999; Lockhart and Winzler 2000). These 'synexpression' groups can most easily be explored by screening the genetic profile of a single cell type with time following a specific experimental manipulation (Hovland et al. 2001; Schulze et al. 2001). In a small, experimentally inaccessible organ like the mammalian inner ear, cell lines present a unique opportunity to investigate complex cellular processes and are well suited to drug discovery (De-bouck and Goodfellow 1999).

There are only 6600 auditory hair cells in the mouse (Echteler et al. 1994). They are born between E12–E14 (Ruben 1967; Kelley et al. 1993) and differentiation proceeds from the base to the apex of the cochlea with a delay of ~2–3 d (Lim and Anniko 1985; Zine and Romand 1996; Nishida et al. 1998). Cells mature around postnatal days P12–P14, when the ear becomes functional (Rubsamen and Lippe 1997; Kros et al. 1998). Additional hair cells can be recruited during this period (Kelley et al. 1993; Zheng and Gao 2000), but there is no evidence for sensory cell replacement after P14. Progressive loss of hair cells accounts for the largest proportion of hearing loss, which affects more than 10% of the human population (Rubel 1997). One advantage of the experimental approach

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that we have adopted lies in the potential identification of key factors and signaling pathways that may allow therapeutic stimulation of hair cell replacement. As a developmental model, it also allows us to study transient events during differentiation of a single cell type.

We have investigated profiles of selected regulatory genes. These include the zinc finger transcription factor *Gata3*, which is expressed in sensory and nonsensory epithelial cells in the developing cochlea at E13 (Rivolta and Holley 1998; Karis et al. 2001; Lawoko-Kerali et al. 2002), the bHLH transcription factor *Math1* (Birmingham et al. 1999; Lanford et al. 2000), and the associated genes *Hes1* (Zheng et al. 2000; Zine et al. 2001) and *Hes5* (Lanford et al. 2000; Zine et al. 2001). *Math1* can induce hair cell differentiation in nonsensory embryonic epithelial cells from the greater epithelial ridge (GER; Zheng and Gao 2000) and its action appears to be opposed by *Hes1* (Zheng et al. 2000). Hair cells and supporting cells share a common progenitor (Fekete et al. 1998; Lang and Fekete 2001) and cell specification depends upon a mechanism of lateral inhibition mediated by Notch1 and its ligands, *Jagged1* and *Jagged2* (Adam et al. 1998; Haddon et al. 1998; Lanford et al. 1999). We show that UB/OC-1 probably originates from the nonsensory epithelial cell population in the GER and that, when differentiated, closely resembles the hair cell phenotype. Thus it has considerable potential for studying the genetic processes underlying a potential mechanism for recruitment of replacement hair cells.

## RESULTS

### Transition from Proliferation at 33°C to Differentiation at 39°C

The transition from proliferating to differentiating conditions is a functional event that should be reflected in the gene expression profiles, thus presenting an opportunity to test the validity of the system. Genes associated with proliferation, including *cyclins A, B1, B2, E, and F*, were down-regulated at 39°C, whereas those associated with the quiescent state, including *Gadd45, Chop-10, and Gas1*, were up-regulated over 15-fold.

Expression levels for over 200 genes changed by more than fivefold during the experiment (Supplemental Table 1 available online at [www.genome.org](http://www.genome.org)). At day 2, expression levels of 55 genes increased relative to day 0, whereas those for 36 genes decreased. At day 4, 119 genes increased and 36 decreased, whereas at day 9, 101 genes increased and 62 decreased. Twelve and 37 genes increased at day 2 and 4, respectively, and then dropped below the fivefold threshold, showing transient expression early in differentiation.

Several loosely related groups of genes changed dramatically. Those encoding extracellular matrix (ECM) proteins such as decorin, *Col6a1*, *Col3a1*, matrix Gla protein, and OSF-2 (similar to fascilin, not to be confused with *osf2/cbfa*) increased from day 2, suggesting that the cells recondition their environment to undergo differentiation by secreting a different basement membrane. *Col8a1*, *Col6a2*, *laminin*  $\beta$ 2 were detected at day 4 and *laminin*  $\beta$ 3 at day 9. Genes encoding growth factors, including cellular retinol binding protein 1 (*CRBP1*), insulin-like growth factor binding protein 2 (*IGFBP-2*), and a member of the *PDGF/VEGF* family increased at day 2. At day 4, the cells expressed *TGF $\beta$ 3*, whereas at day 9 they also expressed insulin-like growth factor binding protein 5 (*IGFBP-5*). The transcription factor *C/EBP $\delta$*  increased

almost 10-fold at day 2 and remained high during late differentiation. The homeobox gene *Prx2* was also up-regulated early in differentiation, but its level dropped below 5-fold by day 4, whereas *gooseoid*, another homeobox gene, peaked at day 4 with a 6-fold increase.

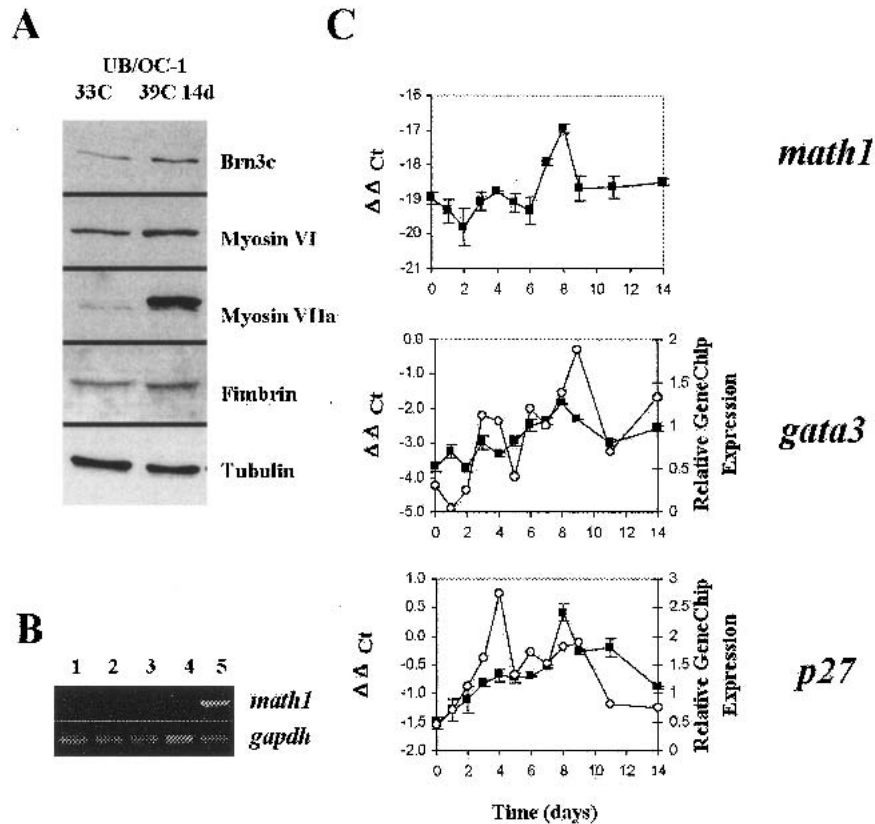
### Characterization of UB/OC-1

UB/OC-1 was originally selected because it expressed key markers for hair cell precursors (Rivolta et al. 1998). These included the transcription factor *Brn3c* (also known as *Brn3.1*), normally expressed from E14 (Ryan 1997) and essential for hair cell survival (Erkman et al. 1996; Xiang et al. 1997, 1998), the  $\alpha$ 9 subunit of the acetylcholine receptor ( $\alpha$ 9AChR), and the cytoskeletal proteins myosin VI, myosin VIIa, and fimbrin (Fig. 1A). The expression of these genes correlated with the developmental stage at which the cells were derived.

One unexpected finding was the absence of the proneural gene *Math1*, which is normally expressed in differentiating hair cells at E12.5–E13 (Birmingham et al. 1999; Lanford et al. 2000). It was recorded absent throughout the microarray analysis and in all samples assayed by RT-PCR (Fig. 1B). Controls with RNA minus reverse transcriptase (RT) and a mock cDNA reaction without RNA were also negative. Furthermore, the TaqMan analysis did not detect any expression above the background of the minus RT controls (Fig. 1C).

There was no obvious evidence for supporting cell markers in differentiated UB/OC-1. Immunoblots were negative for OCP-2, a protein highly expressed by supporting cells in vivo (Yoho et al. 1997). From the connexin family of molecules, only *connexin 31* was detected at 33°C in the microarray, but it disappeared after 2 d at 39°C. The results for *connexins 30, 26, and 43* were negative.  $\alpha$ - and  $\beta$ -*tectorins* are specific to the ear and form part of the tectorial membrane (Legan et al. 1997; Rau et al. 1999). The microarrays were negative for  $\alpha$ -tectorin at all time points. However,  $\beta$ -*tectorin* was detectable at 33°C and quickly down-regulated during differentiation. This ear-specific gene is normally expressed in the sensory epithelium at E12.5, and at E14.5 it is restricted to narrow bands that include the border cells along the inner edge of the inner hair cells, the pillar cells, and the third row of Deiters' cells (Rau et al. 1999). It is not expressed in hair cells.

The profiles obtained for *Gata3* and *p27<sup>Kip1</sup>* in the microarray were broadly similar to those obtained by TaqMan analysis (Fig 1C). These genes are important in inner ear development and their expression in UB/OC1 correlated with that described in vivo. *Gata3* is normally expressed by all cells of the sensory epithelium and is down-regulated initially in differentiating hair cells (Rivolta and Holley 1998; Karis et al. 2001). In UB/OC-1, expression peaked at days 8–9 at 39°C and then dropped later to about half its peak level. For *p27<sup>Kip1</sup>*, there was an initial low level of expression at 33°C and the mRNA increased to peak transiently between days 4 and 8. In vivo, *p27<sup>Kip1</sup>* normally appears around E14 in hair cells and supporting cells, but it is down-regulated in hair cells at ~E16 (Chen and Segil 1999; Lowenheim et al. 1999). Thus the expression profile in the cell line resembled that in vivo, albeit on a slower timescale. This fact, already observed for other genes in this cell line, could prove useful in identifying and dissecting signaling cascade components that would normally have a very short and transient expression in vivo.



**Figure 1** (A) Western blot of protein samples extracted at 33°C and after 14 d at 39°C. (B) RT-PCR for *math1*. Lanes are 1, 33°C; 2, 2 d at 39°C; 3, 4 d at 39°C; 4, 14 d at 39°C; and 5, genomic DNA. (C) Plots comparing the expression profiles for *math1*, *gata3*, and *p27<sup>Kip1</sup>* obtained by TaqMan analysis (filled squares) and GeneChip microarray (open circles). *Math1* plot includes only the TaqMan results because the array values were called absent and are below the normalized threshold. Scales for TaqMan are on the left axis ( $\Delta\Delta$  Ct), whereas those for relative gene chip expression are plotted on the right axis. Standard error bars are shown for the TaqMan assays. The absolute levels of *math1* detected by TaqMan were extremely low. Compare the differences in scale for *math1* and *gata3*. Each unit represents a difference of a power of 2, so at time point 0, *gata3* expression is  $\sim 2^{14}$  greater than *math1*. Moreover, the values obtained for *math1* were not different from those of the minus reverse transcriptase controls.

### Global Cluster Analysis

Global analysis of gene expression can reveal genes with clustered expression profiles that are likely to be functionally related. To establish such groups we employed self-organizing maps (SOMs). SOMs allow easy visualization of changing profiles through time and they place genes with similar profiles in adjacent groups. This produces a matrix with progressive, gradual transitions through the rows and columns. The 1675 genes that displayed a  $>2.5$ -fold change at any time point were clustered into a  $6 \times 6$  lattice (Fig. 2; Supplemental Tables 2–37 available online).

### The Cluster Containing Transcription Factor *Gata3*

In our SOM analysis, *Gata3* located to cluster 1,4 in a group of 130 genes (Supplemental Table 5 available online). Potentially related patterns of gene expression from this set include those of Semaphorins A, E, and F; MEP (a member of the Eph family of receptors that lacks protein tyrosine kinase activity); the cellular retinol binding protein I (CRBPI); the putative regulator of transcription AES1; the homeobox gene *Pbx1*; and several ESTs similar to zinc finger genes.

### Expression Profiles of Elements of the Notch Signaling System in UB/OCI

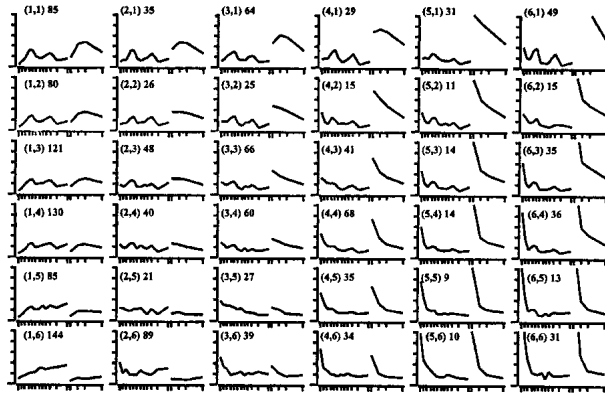
The notch signaling system plays an important role in the determination of cell fates in the inner ear (Adam et al. 1998; Haddon et al. 1998; Lanford et al. 1999) and four members of the notch receptor family, Notch1–4, have been described in the mouse (Weinmaster 1997). UB/OC-1 did not express *Notch2* or *Notch4* at any time point. *Notch1* was present throughout the experiment, although at low levels under proliferative conditions, and it reached a peak after 3–4 d at 39°C (Fig. 3A). *Notch3* was not expressed at 33°C but was detectable after 2 d at 39°C and reached a clear peak at days 3–4 (Fig. 3A). The expression of the notch genes on the arrays correlated well with results from immunoblotting. Notch3 expression was only detected from day 2 onward, whereas Notch1 was detectable at 33°C at low levels, increasing its expression at days 2–3 (Fig. 3B). The expression of Notch1 protein is reduced during late differentiation, being barely detectable after 14 d (not shown). This is in agreement with the expression profile described for differentiating hair cells in vivo (Zine et al. 2000).

*Hes* genes are downstream effectors of the notch cascade. *Hes1* was present at 33°C, increased sharply soon after the temperature switch, and then dropped at days 3–4 to lower levels, at which it remained constant until day 9 (Fig. 3A). Its expression was undetectable by day 14. In contrast, *Hes3* steadily increased during differentiation and reached its peak at day 11 (Fig. 3A). *Hes5* was not detected.

Of the notch ligands, *Dll1* was not detected. *Jagged2* expression peaked at day 11 (Fig. 3A) and on Western blots it was detected from day 2 (Fig. 3B). *Jagged1* probe sets were not included in the MU11K microarray, so we explored its expression profile by Western blot. The anti-jag1 antibody identified a band of  $\sim 190$  Kilodalton (Kd) at 33°C that was up-regulated immediately after the temperature shift. A second band of  $\sim 130$  Kd was also detected. This smaller band was more obvious at 33°C but was progressively down-regulated and virtually undetectable after 6 d at 39°C (Fig. 3B). Proteolytic processing of the full-length ligand may have produced the 130-Kd band. The fact that the extracts were prepared in the presence of several protease inhibitors and that this band was systematically down-regulated with differentiation, suggests that the result reflected a real process within the cells rather than degradation during sample preparation.

### Immunolocalization of Notch3

The pattern of distribution of Notch3 was investigated by im-



**Figure 2** A  $6 \times 6$  matrix displaying the clustered profiles of genes changing over 2.5-fold at any given point. Cluster number is enclosed by parentheses; total number of genes per cluster are outside parentheses. The Y-axis represents normalized relative expression, the X-axis shows both time courses (time course 1 following differentiation over a 14 d period containing 12 data points and time course 2 following differentiation over a 9 d period containing 4 data points). The average profile for each cluster is shown. Genes included in each cluster are listed in Supplemental Tables 2–37 available online.

munolabeling on the cells and also on sections of the developing inner ear. In situ hybridization shows that it is expressed at E11.5 and E13.5 but is no longer detectable at E15.5 (Lardelli et al. 1994; Lindsell et al. 1996; Lewis et al. 1998). In our study, Notch3 was visualized in the epithelium of the inner ear at E14.5 (Fig. 3C). The receptor did not segregate to any particular region or cell type, being homogeneously expressed by the entire epithelial lining, but it was absent from the underlying mesenchyme. At E16.5, its expression was no longer detectable. In UB/OC-1 Notch3 was localized to the cell nuclei after 3 d at 39°C (Fig. 3D). This subcellular distribution suggests that at this time point it was activated, translocated to the nucleus, and engaged in signaling.

**Genes Associated with Notch3**

On the  $6 \times 6$  SOM analysis, Notch3 clustered to the 1,2 group (Fig. 2). This cluster contained a total of 79 genes (Supplemental Table 3 available online.), including 32 ESTs with no significant homology in the database. The annotated genes included the transcription factors *gooseoid*, *AREC3/six4*, and *Sox4*, the cyclin-dependent kinase inhibitor p15ink4b and the vesicle trafficking gene *Munc-18*.

**Other bHLH Genes**

The list of genes expressed by UB/OC-1 was further explored for genes encoding a bHLH protein. *Me1* (also known as *tcf12-hnf4-heb*) was expressed at steady levels from day 0 and *Stra13*, a bHLH-gene inducible by retinoic acid, peaked late during differentiation at days 9–11.

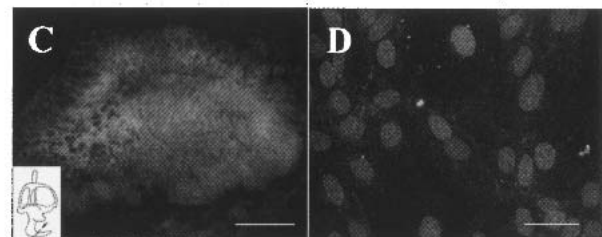
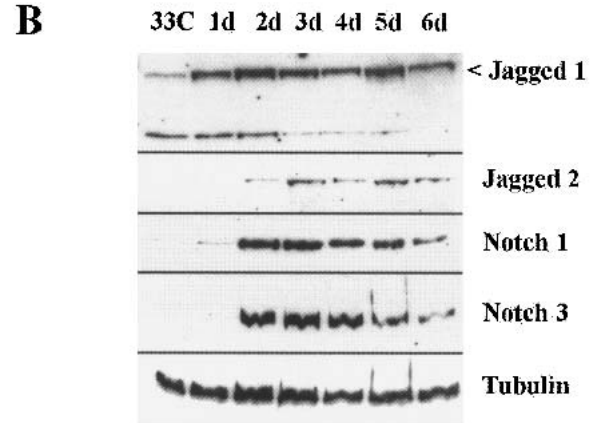
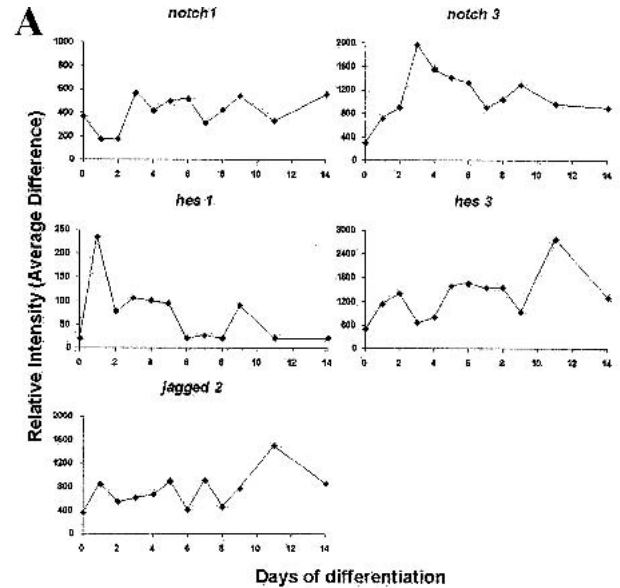
**DISCUSSION**

One of the main applications of global analysis of gene expression using microarrays is to establish hypotheses that can then be tested experimentally. The data presented here have enabled us to characterize further the origin of the cell line UB/OC-1 and then to explore its properties in the context of normal development. Cloned cell lines of known origin allow us not only to identify genes not previously implicated in ear

development, but also to propose potential synexpression groups with or without reference to known genes.

**Identity and Origin of UB/OC-1**

The results strongly suggest that UB/OC-1 was derived from epithelial cells within the GER that have the potential to differentiate into hair cells. Under differentiating conditions, they adopt a hair-cell-like phenotype. None of the supporting



**Figure 3** Notch signaling cascade in UB/OC-1. (A) Relative gene expression for *Notch1*, *Notch3*, *Jagged2*, *hes1*, and *hes3*. (B) Western blot for notch and jagged proteins on protein samples taken at daily intervals. (C) Cross-section of an E14.5 cochlear duct, showing immunolabeling for Notch3. (D) UB/OC1 at 39°C for 3 d, showing nuclear location of Notch3. Bars are 50  $\mu$ m.

cell markers explored is expressed and the cells appear to commit uniformly to the same phenotype, as shown by the homogenous expression of *brn3c* (Rivolta et al. 1998; and this manuscript).

In earlier experiments designed to identify and target proliferating precursors, whole organs of Corti from E13 Immortomice were cultured at 33°C with 100 U/mL  $\gamma$ -interferon to activate the immortalizing T-antigen. However, in the developing sensory epithelium the cell nuclei remained unlabeled by both BrdU and an antibody against the T-antigen. Furthermore, cuticular plates and hair bundles differentiated more or less normally (N. Grix, M. Rivolta and M. Holley, unpubl. results). In the adjacent nonsensory regions, including the GER and the lesser epithelial ridge (LER), the cell nuclei did express the T-antigen and labeled positively for BrdU. A possible interpretation is that the T-antigen is unable to override the controls imposed by intercellular interactions in at least some regions of the intact epithelia. Consequently, cell lines were derived from dissociated cultures and the exact origin of UB/OC-1 is thus unknown. However, the genetic profile suggests that UB/OC-1 originates from the GER. UB/OC-1 expresses *Gata3* and cytokeratins, which are also expressed by sensory and nonsensory epithelial cells in the cochlear duct at the age of derivation (Rivolta and Holley 1998; Rivolta et al. 1998; Karis et al. 2001). It also expresses *Hes1* and  $\beta$ -*tectorin* at 33°C. *Hes1* is present mainly in the GER and the LER (Zheng et al. 2000; Zine et al. 2001), whereas  $\beta$ -*tectorin* appears only in the GER and the pillar cells during development (Rau et al. 1999). On the other hand, *Hes5*, which is normally expressed in the LER (Zheng et al. 2000; Zine et al. 2001), is not expressed by UB/OC-1 at any time point. Finally, *Math1* is expressed in the prosensory cell population but not in the GER.

### Certain Hair Cell Characteristics Are Independent of *Math1* Expression

Our conclusion about the origin of UB/OC-1 is strengthened by the discovery that cells in the GER can become hair cells when transfected with *Math1* (Zheng and Gao 2000). However, UB/OC-1 appears to achieve the expression of numerous hair cell markers without expressing *Math1*. *Math1*-null mice fail to generate hair cells (Bermingham et al 1999), implying that this gene is essential for hair cell production, but in chimeric mice where *Math1*<sup>-/-</sup> embryonic stem cells were introduced into wild type blastocysts, *Math1*<sup>-/-</sup> hair cells have been observed (Du et al. 2000). It thus is conceivable that the pro-hair-cell function assigned to *Math1* is redundant and that other bHLH genes could compensate in its absence. *Math1* may require an inductive signal that is missing in our in vitro system. In UB/OC-1 it is clear that *Math1* is not directly linked to the up-regulation of *myo7a*, *brn3c*, or the  $\alpha$ 9 AChR. However, *Math1* could act by blocking the effect of negative regulators such as *Hes5*, which could directly repress the expression of hair cell genes in supporting cells.

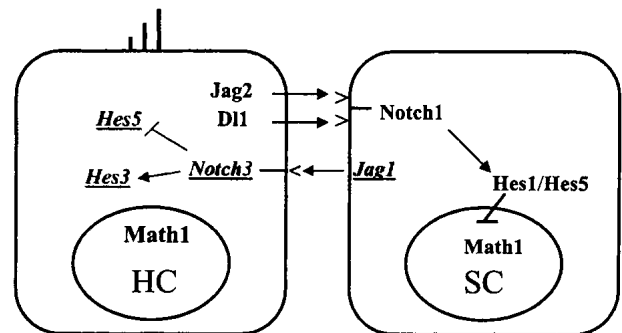
The competence of cells in the GER to form hair cells must be intrinsic because the transfection of other cell types in the same preparations (such as non-neuronal ganglia cells and connective tissue) does not lead to the generation of new hair cells (Zheng and Gao 2000). It is known that *Math1* requires a partner protein, namely members of the class A of HLH proteins (Akazawa et al 1995). The competence of the GER can then be interpreted as the ability of these cells to express such cofactors. With this aim we searched the expres-

sion profiles in the microarray for class A bHLH proteins that were present at any given point. We identified *Me1*, a putative partner of *math1* that colocalizes with it during development of the granular cells of the cerebellum (Uittebogaard and Chiamarello 1999) and that is highly expressed in brain regions that display neuronal plasticity (Chiamarello et al. 1995). In UB/OC-1, *Me1* is expressed at steady levels throughout differentiation.

### Notch3 Could Mediate an Instructive Signal for Hair Cell Differentiation

The expression of *Notch3* could resolve an outstanding issue regarding the current model for notch signaling in the ear. In this model, *Notch1* activation, mediated by *Jagged2*, inhibits the hair cell fate (Lanford et al. 1999; Zhang et al. 2000). The in vivo expression pattern of the notch ligand *Jagged1* is not consistent with a simple model for lateral inhibition. Early in development, *Jagged1* is expressed in the sensory patches (Morrison et al. 1999; Zine et al. 2000), whereas later its expression is segregated to the supporting cells, where it colocalizes with *Notch1*. The mouse mutants *Htu* and *slalom*, produced by missense mutations in *Jagged1*, show a reduction in the numbers of outer hair cells (Kiernan et al. 2001; Tsai et al. 2001), as opposed to the increases seen in loss-of-function studies for *Jagged2*, *Notch1*, and *Hes1*. Thus, *Jagged1* appears to be signaling in the opposite direction to *Jagged2*. It has been proposed from gene reporter studies that *Notch3* acts by interfering with *Notch1* signaling, possibly by blocking *Notch1*-mediated *Hes1* transcription (Beatus et al. 1999, 2001). In UB/OC-1, the level of *Hes1* mRNA drops when *Notch3* increases. It has been shown that activated *Notch3* can induce astroglial differentiation (Tanigaki et al. 2001). Thus it is possible that *Notch3* may act as an antagonist of *Notch1*, mediating an instructive rather than inhibitory signal for hair cell differentiation. In conclusion, we propose that *Jagged1* could have an inductive effect on hair cell differentiation mediated by *Notch3* (Fig. 4).

Several related genes cluster together with *Notch3* or appear in nearby clusters in the SOM analysis, showing a peak of expression early during differentiation. *Groucho1/tle1* is a transcriptional corepressor that interacts with HLH proteins (Fisher and Caudy 1998). It is up-regulated during early stages



**Figure 4** A model for the proposed role of *Notch3* in hair cell differentiation. The conventional lateral inhibition cascade includes *Dll1* and *Jag2*, the ligands expressed by the hair cell that would activate *Notch1* in the adjacent supporting cells. On the other hand, *Jag1* expressed by the supporting cell could be activating *Notch3* in the hair cells and oppose the effects of *Notch1* in this cell type, either by repressing the transcription of *Hes5* or by promoting differentiation through *Hes3*.

of neuronal differentiation in P19 cells exposed to retinoic acid (Husain et al. 1996; Yao et al. 1998), as well as in CFK2 cells, which are a model for chondrocytic determination (Yao et al. 1998). It is also transiently expressed during epithelial differentiation (Liu et al. 1996). *Gooseoid* (*Gsc*) is a homeobox-containing transcription factor that is expressed in the middle ear, as well as in the cochlear epithelia (Yamada et al. 1995; Rivera-Perez et al. 1999). Null-mutant mice have defects mainly in the malleus and tympanic membrane, as well as in the external canal (Yamada et al. 1995). In vitro, expression of *Gsc* can enhance neuronal differentiation (Sawada et al. 2000), whereas in *Drosophila* *Gsc* can interact with *groucho* (Jimenez et al. 1999). *AREC3/Six4* is also a homeobox-containing gene, homolog of the *Drosophila* gene *sine oculis* (Kawakami et al. 1996). It is up-regulated during muscle differentiation of C2C12 cells. In the chicken ear, *cSix4* has been identified at the otic placodes and later to the ventral region of the otocyst that will develop into the cochlea (Esteve and Bovolenta 1999).

Another HLH gene that is up-regulated during differentiation in UB/OC-1 is *Hes3*. During development in vivo, *Hes3* is expressed in the cochlear epithelium (Lobe 1997). Although there is no clear evidence that will support *Hes3* as a positive instructor of differentiation, it has been proposed that the alternatively spliced variant *Hes3a* could inhibit the activity of negative repressors like *Hes1* by competing for a cofactor, therefore promoting differentiation (Hirata et al. 2000).

### *Gata3* and Semaphorins

*Gata3* has a broad pattern of expression in the ear (Rivolta and Holley 1998; Karis et al. 2001; Lawoko-Kerali et al. 2002), but there is strong evidence that it is involved in neural pathfinding (Karis et al. 2001). In this context it may be important to note that three semaphorin genes (*semaA*, *semaE*, and *semaF*) clustered to the same group. Semaphorins are signaling molecules involved in axon pathfinding, mainly acting as a repelling cue for the guidance of axons during development (Tammagnone and Comoglio 2000). During innervation of the organ of Corti, *Gata3* is down-regulated in hair cells but remains expressed by the surrounding supporting cells. If these semaphorin genes are under *Gata3* regulation, it could be hypothesized that they are preventing the axons from establishing synaptic contacts with the supporting cells.

An understanding of the genetic programs that underlie the conversion of nonsensory to sensory cell phenotypes could be essential for developing therapeutic approaches to sensory regeneration in the inner ear. We have shown that the combination of a characterized cell line and gene array technology can yield robust insights into such programs. The data allows us to form sound hypotheses for the functions of developmentally important genes, illustrated here by *Math1*, *Gata3*, and *Notch3*, that can be tested experimentally either in vivo or in vitro.

## METHODS

### Cell Culture

UB/OC-1 and culture conditions have been described previously (Rivolta et al. 1998). Proliferation was maintained by culturing cells in Minimal Essential Medium with Earle's Salts and Glutamax I (MEM, Life Technologies), 10% fetal calf serum (FCS, Life Technologies), and 50 U/mL of  $\gamma$ -Interferon ( $\gamma$ -IFN, Life Technologies) at 33°C. Differentiation was induced by replacing the media without  $\gamma$ -IFN and then trans-

ferring the dishes to 39°C. RNA samples were collected at day 0 (before transfer to differentiating conditions) and then at 24-h intervals up to 14 d after the temperature shift. Two independent time courses were performed and two separate sets of RNA samples were collected.

### RNA Isolation and Array Hybridization

Total RNA was prepared with a Qiagen RNeasy mini-kit according to the manufacturer's instructions. cRNA was prepared as described in the Affymetrix GeneChip Expression Analysis Technical Manual. GeneChip oligonucleotide arrays (Affymetrix), composed of 13,179 probe sets representing over 11,000 genes and ESTs (MU11K; chips A and B) were used for hybridization. A comprehensive list of genes present in the chips and search tools to scan through their contents are available at <http://www.netaffx.com>. Hybridizations were performed according to the Affymetrix GeneChip Expression Analysis Technical Manual. Arrays were scanned using an Affymetrix confocal scanner. Array hybridization was performed at the time points 0, 1, 2, 3, 4, 5, 6, 7, 8, 9, 11, and 14 d for the first experiment (Time Course 1) and at 0, 2, 4, and 9 d for the second experiment (Time Course 2).

### Analysis of GeneChip Data

Scanned output files were visually inspected for hybridization artifacts and then analyzed with GeneChip v. 3 (Affymetrix). Arrays were scaled to an average intensity of 300 and analyzed independently. The expression data for each array was saved as an Excel file containing expression values (Average Difference) and detection of RNA species (Absence Call). These parameters were calculated by GeneChip according to standard Affymetrix procedures. Comparison files were generated with GeneChip for each differentiating time course, using day 0 as a baseline. These comparison files contained a parameter that measured whether gene expression had changed between two time points (Difference Call). For further data mining, clustering, and presentation, tab delimited Affymetrix Excel files were imported into GeneSpring (v. 3.2.11–4.0.0, Silicon Genetics) after values for Average Difference of <20 were given an arbitrary value of 20. In GeneSpring, expression profiles were normalized per time point, making the median of all measurements 1, and normalized per gene to give a median value of 1 for each gene expression profile. These normalizations reduced the influence of outliers and allowed the visualization of expression profiles for different genes on the same axis.

### Criteria for Gene Selection

The following criteria were set for determining which genes were responsive to differentiation in UB/OC-1. Gene lists were generated for genes with Difference Calls that represented expression changes in at least one time point with respect to day 0 in Time Course 1. This gene list was further filtered in GeneSpring by applying a 2.5-fold change restriction in at least one time point for Time Course 1. However, lists were also scanned for potentially interesting genes based on their Presence/Absence Call, despite their fold change.

### Gene Clustering

Clustering was performed using a SOM algorithm in GeneSpring. SOMs have the benefit of being a well-studied method of clustering and have been shown to be able to extract physiologically relevant patterns of gene expression in an experimental setting (Tamayo et al. 1999). All genes modulated by at least 2.5-fold in Time Course 1 were clustered according to the relative behavior of each gene in both time courses.

## Fold Change Lists

Restrictions were applied to Time Courses 1 and 2 and days 0, 2, 4, and 9 using the Affymetrix generated metrics Fold Change, Absolute Call, and Difference Call. Lists were generated of genes with expression that changed measurably (Difference Call) by at least 2.5-fold (Fold Change) compared to day 0, were detected above background in at least one of the time points (called Present in the Absolute Call), and were maintained in both experiments for each day. These lists were further restricted by including only those genes that had an Average Fold Change  $\geq 5$ -fold.

## TaqMan Assay

TaqMan 5' nuclease fluorogenic quantitative PCR assays were performed using the TaqMan Gold RT-PCR kit (PE Applied Biosystems) according to manufacturer's instructions. Total mRNA was prepared from UB/OC-1 grown and harvested as described above. Reverse transcription was performed with 2  $\mu$ g of total RNA using random hexamer primers and identical reactions were performed in the absence of reverse transcriptase. Assays were performed in triplicate using an ABI Prism 7700 Sequence Detector (PE Applied Biosystems), VIC-labeled ribosomal RNA control reagents (PE Applied Biosystems), and the following oligonucleotides (5' to 3'; MWG Biotech UK):

forward primer CTGAAAACCTGAGACAACCAATGC  
reverse primer AAGGGTGCAGGGATATTTGTCA  
FAM-labeled probe CTAGCGCGCGGAAGCCCC.  
GATA-3:

forward primer GAGTCTCCAAGTGTGCGAAGAGT  
reverse primer TCGGGCTTCATGATACTGCTC  
FAM-labeled probe TCCGACCCCTTCTACTTGCGTTTTTCG.  
G1 cyclin-Cdk protein kinase inhibitor p27:  
forward primer CTCCGCCTGCAGAAATCTC  
reverse primer CAGTGCTTCTCCAAGTCCCC  
FAM-labeled probe TCGGCCCGGTCAATCATGAAGAACT

## RT-PCR

End-point RT-PCR was performed on samples taken at 33°C and after 2, 4, and 14 d at 39°C. Controls included genomic DNA and samples without reverse transcriptase. *Math1* primers and cycling conditions were taken from Ben-Arie et al. (1996). *Gapdh* primers used as a normalizing control were as previously described (Rivolta et al. 1998).

## Immunoblotting and Immunolabeling

These procedures were performed as previously described (Rivolta et al. 1998; Lawlor et al. 1999). Primary antibodies used were as follows: brn3c (rabbit polyclonal, Babco); myosin VI (rabbit polyclonal, a kind gift from T. Hasson, UCSF), myosin VIIa (rabbit polyclonal, a kind gift from C. Petit, Institut Pasteur), fimbrin (rabbit polyclonal, a kind gift from P. Matsu-daira, Whitehead Institute for Biomedical Research). Polyclonal antibodies to Jagged1, Jagged2, Notch1, and Notch3 were from Santa Cruz Biotech and the tubulin Ab-4 antibody (Clone DM1A + DM1B) was from Neomarkers.

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