The occurrence of neurological symptoms and developmental delay in patients affected by congenital hypothyroidism (CH) has been attributed to the lack of thyroid hormone in the developing CNS. Accordingly, after the introduction of neonatal screening programs for CH, which allowed early and adequate treatment, an almost normal outcome for most CH patients could be achieved. However, a few patients did not reach this favorable outcome despite early and adequate treatment. Here we describe five patients with variable degrees of CH who suffered from choreoathetosis, muscular hypotonia, and pulmonary problems, an association of symptoms that had not been described before this study. Since this clinical picture matched the phenotype of mice targeted for deletion of the transcription factor gene Nkx2-1, we investigated the human NKX2-1 gene in these five patients. We found heterozygous loss of function mutations in each of these five patients, e.g., one complete gene deletion, one missense mutation (G2626T), and three nonsense mutations (2595insGG, C2519A, C1302A). Therefore, the unfavorable outcome in patients with CH, especially those with choreoathetosis and pulmonary symptoms, can be explained by mutations in the NKX2-1 gene rather than by hypothyroidism. Moreover, the association of symptoms in the patients with NKX2-1 mutations points to an important role of human NKX2-1 in the development and function of thyroid, basal ganglia, and lung, as already described for […]

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Introduction
Choreoathetosis is defined as rapid involuntary and slow writhing movements due to congenital or acquired defects of the basal ganglia. Huntington disease represents a late-onset progressive variant of chorea with neurodegeneration of the striatum caused by inherited amplification of a CAG repeat in exon 1 of the huntingtin gene (1) (MIM 143100). In contrast, the early onset and nonprogressive chorea in few familial cases of “benign hereditary chorea” (MIM118700) suggest a molecular defect of basal ganglia development. Recently, benign hereditary chorea was linked to a region on chromosome 14 including the NKX2-1 gene, known to be expressed during basal ganglia development in rodents (2).

The occurrence of neurological symptoms and developmental delay in patients affected by congenital hypothyroidism (CH) has been attributed to the lack of thyroid hormone in the developing CNS. Accordingly, after the introduction of neonatal screening programs for CH, which allowed early and adequate treatment, an almost normal outcome for most CH patients could be achieved. However, a few patients did not reach this favorable outcome despite early and adequate treatment. Here we describe five patients with variable degrees of CH who suffered from choreoathetosis, muscular hypotonia, and pulmonary problems, an association of symptoms that had not been described before this study. Since this clinical picture matched the phenotype of mice targeted for deletion of the transcription factor gene Nkx2-1, we investigated the human NNX2-1 gene in these five patients. We found heterozygous loss of function mutations in each of these five patients, e.g., one complete gene deletion, one missense mutation (G2626T), and three nonsense mutations (2595insGG, C2519A, C1302A). Therefore, the unfavorable outcome in patients with CH, especially those with choreoathetosis and pulmonary symptoms, can be explained by mutations in the NNX2-1 gene rather than by hypothyroidism. Moreover, the association of symptoms in the patients with NNX2-1 mutations points to an important role of human NNX2-1 in the development and function of thyroid, basal ganglia, and lung, as already described for rodents.

This article was published online in advance of the print edition.
The date of publication is available from the JCI website, http://www.jci.org.

Choreoathetosis, hypothyroidism, and pulmonary alterations due to human NNX2-1 haploinsufficiency

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Received for publication on October 10, 2001, and accepted in revised form on January 8, 2002.

The Journal of Clinical Investigation | February 2002 | Volume 109 | Number 4

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failed to show NKX2-1 mutations in patients with various types of thyroid dysgenesis (7–9). In contrast, two patients with a large chromosomal deletion spanning the NKX2-1 gene were found to be affected by a variety of symptoms in the CNS, thyroid, and lung (10, 11). Therefore, it could be hypothesized that mutations in the human NKX2-1 gene might result in a complex disease including neurological, thyroid, and respiratory problems.

We searched for mutations in the NKX2-1 gene in patients with symptoms matching this predicted NKX2-1–deficiency phenotype.

**Methods**

**Mutational analysis.** The entire coding region of the NKX2-1 gene was amplified in one fragment from genomic DNA (forward primer: 5´ CGGGAGGCAGTGCACTCCTACCTCAG; reverse primer: 5´ AGGGCCGGCCGCGTCCTCTCTCACC). Exon 1, 2, and 3 were sequenced with an ABI prism 377 sequencer (Applied Biosystems, Foster City, California, USA) using appropriate primers. Identified mutations were confirmed by sequencing of both alleles after cloning PCR fragments into a TOPO cloning vector (Invitrogen, Groningen, The Netherlands) or by restriction analysis of PCR fragments using MseI, BfaI, and HpyCH4v (NEB, England Biolabs).

**Immunohistochemistry.** Histological studies were performed using standard techniques for peroxidase staining using a NKX2-1/TTF-1–specific Ab generated against rat recombinant antigen (DAKO Corp., Carpinteria, California, USA). The human tissues were from autopsy sections. The rat tissues were prepared without prior extended fixation.

**Electromobility shift assay.** NKX2-1 HD was cloned from genomic DNA of patient 2 into the pCRT7/CT-TOPO expression, and the correct sequence was confirmed by direct sequencing of the constructs. BL21I plys cells were chemically transformed and stimulated with isoprpyl-β-D-thiogalactopyranosid (IPTG) to produce an HD/Flag fusion protein according to standard protocols. Total protein extracts were then purified using a Bio-Rad Econo-Pac S Cartridge (Bio-Rad, München, Germany). An electromobility shift assay (EMSA) was performed using 32P-labeled oligo-C (5´-CAC TGC CCA GTC AAG TGT TCT TGA-3´) as a probe and mutated oligo-C (5´-CAC TGC CCA GTC ACG CGT TCT TGA-3´) as a nonspecific control, as described previously (3). To ensure equivalent amounts of wild-type and mutant HD protein, Western blot analysis was performed using a monoclonal M2 anti-Flag Ab (Sigma, Deisenhofen, Germany) according to the improved manufacturer’s recommendations. Equivalent amounts of total protein were ascertained by toping the samples with purified extract from bacteria transformed with the empty vector alone and measurement of the correct protein load with the Bio-Rad protein assay (Bio-Rad).

**Results**

In a cohort of 150 children identified in a neonatal screening program for CH, we observed choreoathetosis, neonatal respiratory distress, and frequent severe pulmonary infections in two patients (Table 1).

**Cytogenetic studies** identified an interstitial deletion of chromosome 14 affecting the chromosomal region (14; q11.2q13.3) in patient 1 (Figure 1), including the NKX2-1 locus confirmed by comparative genomic hybridization (data not shown). In patient 2, direct sequencing of PCR products spanning the entire coding region of the NKX2-1 gene revealed a heterozygous mutation (G2626T) in exon 3 (Figure 1). This missense mutation changes a highly conserved valine to amino acid (aa) position 45 of the DNA binding HD to phenylalanine (Figure 2a).

Based on these results, we extended the NKX2-1 mutational analysis to other patients with CH and mild thyroid abnormalities associated with neurological symptoms including choreoathetosis, muscular hypotonia, ataxia, and retarded development, as well as pulmonary symptoms diagnosed in several other centers. Within this group of 13 patients we detected a boy (patient 3, Table 1) with a heterozygous GG insertion at nucleotide

<table>
<thead>
<tr>
<th>Patient</th>
<th>Current age (years)</th>
<th>Max. TSH* (µIU/l) (0.5–5)</th>
<th>Min. T4 (µg/dl) (6.5–15)</th>
<th>Thyroid gland imaging</th>
<th>Respiratory distress</th>
<th>Pulmonary infection</th>
<th>Choreoathetosis</th>
<th>Mutation</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>16</td>
<td>44</td>
<td>5.9</td>
<td>hypoplasia</td>
<td>severe</td>
<td>severe–frequent</td>
<td>severe</td>
<td>deletion</td>
</tr>
<tr>
<td>2</td>
<td>15</td>
<td>122</td>
<td>2.9</td>
<td>hypoplasia</td>
<td>moderate</td>
<td>severe–frequent</td>
<td>severe</td>
<td>G2626T</td>
</tr>
<tr>
<td>3</td>
<td>3</td>
<td>12</td>
<td>1.1</td>
<td>normal</td>
<td>no</td>
<td>few–mild</td>
<td>no</td>
<td>259SinsGG</td>
</tr>
<tr>
<td>4</td>
<td>13</td>
<td>55</td>
<td>3.9</td>
<td>hypoplasia</td>
<td>no</td>
<td>no</td>
<td>severe</td>
<td>C2519A</td>
</tr>
<tr>
<td>5</td>
<td>1.5</td>
<td>825</td>
<td>&lt;1</td>
<td>agenesis</td>
<td>severe</td>
<td>no</td>
<td>no</td>
<td>C1302A</td>
</tr>
</tbody>
</table>

* TSH concentrations of patients 1, 2, 4, and 5 are from the initial diagnosis in the neonatal period or from the off-treatment period at an age of 2 years. Patient 2 was diagnosed with euthyroid hyperthyreotropinemia at an age of 15 months. All TSH concentrations during the period of l-thyroxine treatment were normal, indicating adequate substitution. *No apparent thyroid tissue seen in scintigraphy performed in neonatal period; hypoplasia at reinvestigation by ultrasound. *So far only scintigraphy performed in neonatal period. *Muscular hypotonia represents the initial symptom of infants developing choreoathetosis later in childhood.
position 2595 in exon 3 (Figure 1). He was affected predominantly by choreoathetosis and had only mild thyroid dysfunction with elevated thyroid-stimulating hormone (TSH) and normal serum thyroid hormone concentrations. The mutation results in a frameshift that leads to a truncated protein lacking the entire third helix of the HD (Figure 2). In a fourth male patient with CH and choreoathetosis, a heterozygous point mutation (C2519A) in exon 3 changed Ser at position 9 of the HD to a premature stop mutation. The fifth patient was identified with severe CH associated with muscular hypotonia and was affected by severe respiratory distress after birth, requiring oxygen administration for several weeks. A nonsense mutation in exon 2 was found (C13302A) changing Cys at position −75 relative to the HD into a premature stop codon (Figure 2). 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Discussion

We identified the first five patients with heterozygous NKX2-1 mutations who are affected by choreoathetosis, hypothyroidism, and respiratory problems, an association of symptoms as yet not described. Since all five patients were adequately treated with L-thyroxine, the unfavorable neurological outcome is most likely related to NKX2-1 deficiency in the thyroid and the brain rather than to the consequence of hypothyroidism. Therefore, the observation of a developmental delay or neurological symptoms in CH patients diagnosed in neonatal screening programs should lead to a molecular diagnosis of genes expressed in the developing thyroid and brain.

The finding of dysgenesis of the pallidum in patient 2 corresponds to developmental defects of the basal ganglia in Nkx2-1–deleted mice (14). In addition, the structural alteration of the basal ganglia in patient 2 seems to explain the clinical finding of choreoathetosis. Since immunohistochemical studies revealed that the NKX2-1 gene product is not expressed in adult neurons of the basal ganglia, NKX2-1 seems to be of importance in the developmental regulation of basal ganglia formation, but does not play a role in continuous basal ganglia function. This predominant role for CNS organogenesis rather than CNS function was also suggested by the finding of a cystic diencephalic mass in patient 1 and 2. Nkx2-1 is expressed in the diencephalic endoderm leading to subsequent expression of Fgf8, which in turn promotes pituitary development (15). In addition, homozygous targeted disruption of Nkx2-1 results in ablation of the pituitary gland (6). This functional role of Nkx2-1 at the interphase of diencephalic and pituitary development in rodents correlates with the formation of a cystic mass exactly at the same position in affected patients. Since in the heterozygous patients the morphology of the pituitary appeared to be normal and pituitary hormone secretion was not altered, these findings suggest that the cystic mass reflects an alteration of diencephalic development rather than pituitary function.

The variable spectrum of thyroid abnormalities in the NKX2-1–mutant patients includes functional as well as developmental defects. While in patient 3 the mild elevation of TSH with normal thyroid hormone secretion clearly points to a functional defect only, the finding of thyroid dysgenesis in the other four patients also reflects a developmental defect of the thyroid gland. A functional thyroid defect can be explained by the reduced activation of the well-known NKX2-1 target genes in thyroid follicular cells, e.g., TSH receptor, thyroid peroxidase, and thyroglobulin gene (16). The molecular defects leading to thyroid dysgenesis in the NKX2-1 mutation carriers remains elusive since no target genes of NKX2-1 during thyroid development have been described.

Nkx2-1 was shown to participate both in lung morphogenesis and in respiratory epithelial cell gene regulation, especially of surfactant protein (SP) genes (17). It has been shown that SP-B is critical for reduction of alveolar resistance leading to respiratory distress in SP-B–/– mice (18). SP-A and -D play an important role in pulmonary host defense (19). Variable pulmonary problems were found in all patients, including neonatal respiratory distress requiring mechanical ventilation (for a short time in patients 1 and 2 and for several weeks in patient 5). In addition, frequent and sometimes severe pulmonary infections in the first years of life were present (patient 1 and 2). In patient 1,
a developmental defect appearing as lung sequestration was diagnosed and resected within the first year of life. Therefore, alteration of NNX2-1 function during development could have resulted in the lung sequestration of patient 1, whereas the predisposition to pulmonary infections and neonatal respiratory distress can be explained on a molecular level by the reduced expression of SPs.

The detailed description of defects in homozygous Nkx2-1–null mice show a more severe phenotype compared with the symptoms observed in the heterozygous human patients. Moreover heterozygous mice were not reported to have any abnormalities (6). The same differences in human versus mice phenotypes were found in another member of the NK2 gene family, e.g., NNX2-5. In humans carrying a heterozygous NNX2-5 mutation an atrioventricular conduction block associated with mild to severe morphological cardiac defects was described (20). In contrast, heterozygous Nkx2-5+/− mice were reported to be normal while in NNX2-5−/− mice an arrest of heart development at an early stage was reported (21). Subsequent reinvestigation of heterozygous Nkx2-5 mice revealed subtle functional and morphological cardiac defects influenced by the genetic background (22). Similar findings in heterozygous NNX2-1 mice have now been reported (23). This dominant inheritance of transcription factor mutations with species-specific sensitivity to gene dosage has already been described for members of other transcription factor gene families involved in early development, such as PAX genes (24).

Our data show that NNX2-1 mutations result in a complex phenotype resembling the tissue expression pattern of the NNX2-1 gene, affecting brain, thyroid, and lung. This pattern of organ manifestation of NNX2-1 mutations appears to be similar in the affected patients. This clinical similarity in patients affected by large deletion of chromosome 1q4 and by loss of function mutations within the NNX2-1 gene implies that this phenotype is entirely due to NNX2-1 haploinsufficiency. The
spectrum of developmental and functional defects further point to a dual role of NKX2-1 in organogenesis as well as in gene expression in adult tissues. The finding of dysplasia of the pallidum associated with choreoathetosis in NKX2-1-deficient patients represents, to our knowledge, the first monogenetic defect leading to a developmental disorder of basal ganglia. The prominent finding of nonprogressive choreoathetosis in NKX2-1-deficient patients may suggest that other congenital extrapyramidal movement disorders could be caused by NKX2-1 mutations. This is supported by recent link of benign hereditary chorea to a region on chromosome 14 including the NKX2-1 gene locus (2).

Acknowledgments
We thank K. Huhne, P. Ambrugger, and A. Gerlach for technical assistance, and D. Lehmann for providing MRI images of patients 1 and 2. This work was supported by a grant of the Deutsche Forschungsgemeinschaft (SFB 577 A5) and the Sonnenfeld-Stiftung.