Phenotypic and genetic exploration of severe demyelinating and secondary axonal neuropathies resulting from GDAP1 nonsense and splicing mutations

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We identified two novel GDAP1 homozygous mutations in children affected with severe demyelinating peripheral neuropathies and born to consanguinous parents. A 9 year old Lebanese girl carried a nonsense mutation in exon 5 and two Algerian brothers aged 10 and 8 years carried a mutation at the intron 3 acceptor splicing site. The clinical, electrophysiological, and neuropathological explorations showed common features consistent with a severe demyelinating peripheral neuropathy associated with loss of major fibres. Our findings, supported by the first GDAP1 expression study in patients, show further evidence that mutations in this gene cause an autosomal recessive severe demyelinating peripheral phenotype (CMT4A) associated with axon loss.

Charcot-Marie-Tooth disease (CMT) (also called hereditary motor and sensory neuropathy (HMSN)) is clinically, electrophysiologically, and genetically extremely heterogeneous with more than 40 loci and 16 genes identified to date. The most common inherited peripheral neuropathies are CMT type 1 and CMT type 2, which are characterised by progressive weakness and atrophy, initially of the peroneal muscles and later of the distal muscles of the arms. Myelination is mainly affected in CMT1, with neuropathological findings of demyelination and remyelination with onion bulb formation, clusters of regeneration, and Schwann cell proliferation. CMT2 is an axonal neuropathy characterised by the reduction of fibre density in the absence of clusters of myelin regeneration or proliferation of Schwann cells. Very often, neuropathological analysis of patients carrying demyelinating neuropathies shows concomitant axon fibre loss. A motor nerve conduction velocity (MNCV) threshold value of 38 ms\(^{-1}\) at the median nerve is used to classify patients with CMT as being either CMT1 (MNCV<38 ms\(^{-1}\)) or CMT2 (MNCV>38 ms\(^{-1}\)). CMT4A defines a particularly severe polyneuropathy of the demyelinating type, associated with distal weakness and atrophy of the limbs, with early onset. The muscular atrophy rapidly progresses, extending proximally and leading to inability to walk by the end of the first decade. Skeletal deformities and scoliosis are frequent. Mutations in the ganglioside induced differentiation associated protein 1 gene (GDAP1) at chromosome 8q21.3 were originally reported in families carrying typical CMT4A and in families affected with autosomal recessive axonal CMT associated with vocal cord paresis. Other GDAP1 mutations in patients affected with demyelinating and axonal peripheral neuropathies have recently been reported. The 12 published mutations are listed in table 1.

Here we report on two families, of Lebanese and Algerian origin, in which two new mutations in GDAP1 have been found. All affected persons had clinical, electrophysiological,

### Key points
- We studied two consanguineous families, one Lebanese and one Algerian, affected with early onset, severe peripheral neuropathies.
- Electrophysiological and histopathological characterisation showed prominent, rapidly progressing demyelinating features associated with loss of major fibres.
- We sequenced GDAP1 exons and intron-exon boundaries in the two families and found two unreported homozygous mutations segregating with the disease, one in exon 5 (c.668T>A) and the other in the intron 3 acceptor splicing site (IVS3-2A>G).
- On the one hand, the c.668T>A mutation is predicted to create a stop codon interrupting the protein at residue 223; on the other hand, expression analyses showed that the intron 3 homozygous mutation has a deleterious effect on transcription, causing the skipping of exon 4 in all the detectable transcribed units.
- Whatever the physiological functions of GDAP1, the truncated proteins do not seem to be able to ensure normal peripheral nervous system development, giving rise to the severe, early onset CMT4 phenotype found in the patients reported here.

### Table 1

<table>
<thead>
<tr>
<th>Exon/intron</th>
<th>Mutation</th>
<th>Effect</th>
<th>References</th>
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<tr>
<td>Exon 1</td>
<td>c.92G&gt;A</td>
<td>W31X</td>
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<tr>
<td>Exon 3</td>
<td>c.349_350insT</td>
<td>T117fs</td>
<td>7</td>
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<td>Exon 3</td>
<td>c.359G&gt;A</td>
<td>R120Q</td>
<td>8</td>
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<td>c.482G&gt;A</td>
<td>R161H</td>
<td>4</td>
</tr>
<tr>
<td>Intron 3</td>
<td>IVS3-2A&gt;G</td>
<td>r.485_579del</td>
<td>This study</td>
</tr>
<tr>
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<td>5, 8</td>
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<td>Intron 4</td>
<td>IVS4+1G&gt;A</td>
<td>?</td>
<td>7</td>
</tr>
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</tr>
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<td>Exon 6</td>
<td>c.786delG</td>
<td>G262fs</td>
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Abbreviations: CMT, Charcot-Marie-Tooth disease; EMG, electromyography; GDAP1, ganglioside induced differentiation associated protein 1 gene; MNCV, motor nerve conduction velocity; NCV, nerve conduction velocity; OFC, occipitofrontal circumference; RT-PCR, reverse transcriptase PCR; SNCV, sensory nerve conduction velocity.
and neuropathological features consistent with a severe demyelinating phenotype associated with major fibre loss. The affected Lebanese girl carried a mutation in exon 5 (c.668T>A) leading to the substitution of the leucine 223 residue to a stop codon (p.L223X), and the two affected Algerian brothers carried a mutation of the third intron acceptor splicing site (IVS3–2A>G), which caused the skipping of the fourth exon in all the transcribed units. This report, supported by the first GDAP1 expression study in patients, shows further evidence that mutations in GDAP1 cause autosomal recessive, severe, demyelinating peripheral phenotypes (CMT4A) associated with axonal loss.

**PATIENTS AND METHODS**

**AR-CMT families**

We studied two consanguineous families (AR-CMT LIB284 and AR-CMT ALG384, fig 1) including children affected with early onset, severe peripheral neuropathy. In both families, consanguinity suggested that the neuropathic phenotype segregated as an autosomal recessive trait.

**AR-CMT LIB284**

The affected girl (LIB284-IV.2, fig 1) was the second child of healthy, first cousin, Lebanese parents. Pregnancy was uneventful with no history of exposure to prenatal or perinatal environmental toxins, and delivery was at term. Birth weight, length, and occipitofrontal circumference (OFC) measurements were not available but were considered normal by the parents. Psychomotor developmental milestones were in the normal range. Disease onset was around 1 year of age, when walking difficulties became evident with a wide based and ataxic gait associated with steppage/foot dropping. By the age of 5 years, weakness of the upper limbs, particularly affecting the hands, and finger retraction, appeared (fig 2E, F). Since then, the progression of the disease has been slow.

**AR-CMT ALG384**

ALG384-II.1 and his sister ALG384-II.2 (fig 1) were respectively the first and second (2 years younger) children of healthy, consanguineous, Algerian parents. Pregnancies and birth were both uneventful and at term. Psychomotor developmental milestones were in the normal range. Disease onset was around 1 year of age, when walking difficulties became evident with a wide based and ataxic gait associated with steppage/foot dropping. By the age of 5 years, weakness of the upper limbs, particularly affecting the hands, and finger retraction, appeared (fig 2E, F). Since then, the progression of the disease has been slow.

The aim of this study was explained to affected families by the primary doctor or neurologist they were referred to and informed consent was obtained from participating people. The study complies with the ethical guidelines of the institutions involved.

**Electrophysiology**

Motor and sensory nerve conduction velocities (MNCVs and SNCVs) as well as electromyographic features were examined in the upper and lower limbs. The MNCVs were recorded at medial and peroneal nerves. Median (elbow-wrist), peroneal (head of fibula-ankle) MNCVs, and muscle action potential amplitudes were registered with surface electrodes. Sensory action potentials were recorded antidromically from the median nerve. Needle electromyography (EMG) examination was performed on distal and proximal muscles.

**Ultrastructural characterisation**

For analysis of nerve biopsies, fascicles of the superficial peroneal nerve were fixed in formaldehyde (10%) and embedded in paraffin. Routine sections were stained using conventional methods. Others fascicles were fixed and buffered in glutaraldehyde, processed, and embedded in epon.

**Genotyping and sequence analyses**

Involvement of MPZ (1q22), PMP22 (17p11.2), EGR2 (10q21-q22), and NDRG1 (8q24) was excluded by direct sequencing. For the other genes reported as being involved in demyelinating AR-CMT (MTMR2, PRX, and GDAP1), a segregation analysis was performed using microsatellite markers respectively on chromosomes 11p12 (AFM119XH4, D11S4134/AFMB021ZG9), 19q13.3 (D19S881/AFMA284Y9, D19S420/AFM326XH9), and 8q21.3 (D8S551/AFM308ZE5, D8S548/AFM301ZC9). Polymerase chain reactions (PCRs) were performed under standard conditions, using IRD-800 labelled forward primers; analysis of PCR products was obtained with a Li-Cor 4200 automated sequencer. The six GDAP1 coding exons and splicing sites (Genbank accession number NM_018972.1) from the affected patients were amplified and then sequenced using a BigDye terminator cycle sequencing procedure (Applied Biosystems). All the sequence variations found were explored in the same conditions from the unaffected members of the families, to confirm the segregation pattern. Sequencing reactions were run on an ABI automated capillary DNA sequencer 310 (Applied Biosystems).
Data were collected and analysed using Sequencher (GeneCodes). All sequence mutations are described according to den Dunnen and Antonarakis.

Effect on splicing of the IVS3–2A>G mutation

Predicted effect

Wild type and mutated IVS3–2A>G genomic sequences were analysed at the web site http://www.charite.de/ch/medgen/img_files/Forschung/Splice/splice.html using the splice score algorithm of Shapiro and Senapathy.

Reverse transcription (RT)-PCR

Total RNA was extracted from EBV immortalised lymphoblastoid cell lines of patients ALG384-II.1 and ALG384-II.2 and an age matched control, with TRIZOL (Invitrogen-Life Technologies). Reverse transcription was performed with Superscript II reverse transcriptase (Invitrogen-Life Technologies) following company recommendations. To obtain enough DNA for sequencing, nested PCRs were performed on cDNAs with primers lying in GDAP1 exons 3 and 5 (see fig 5A, C, D, E). The expected amplicon sizes were 203 bp for controls and 108 bp for a transcript missing exon 4 (r.485_579del); with those primers (fig 5E), eventual genomic contaminants would have been amplified at a 2736 bp size. A PCR with primers lying in glyceraldehyde-3-phosphate dehydrogenase (GADPH) exons 7 and 8 (amplicon size 298 bp) was used as a control of cDNA quality, as previously described (see fig 5B). The products of reverse transcriptase PCR (RT-PCR) were subsequently sequenced (see genotyping and sequence analysis).

RESULTS

Clinical follow up

AR-CMT LIB284

During the follow up, our first clinical examination of the girl, when she was 9 years old, showed normal psychological development, and she was registered in a public school. On examination, head circumference and stature were normal. There were no dysmorphic features except a pectum excavatum. Neurological examination showed important sensory ataxia, bilateral steppage gait, mild kyphoscoliosis, symmetrical and marked atrophy of intrinsic hand muscles and lower leg muscles, and retraction of the fingers (fig 2E, F). Muscle strength was very diminished distally in the upper and lower limbs. Tendon reflexes were absent in the four limbs, and distal sensory abnormalities including deficits of position, vibration, pain, and temperature were registered in both upper and lower limbs in a glove/stocking distribution. No seizures or facial weakness were noted. Abdominal ultrasound, echocardiography, electroencephalography, and ophthalmological evaluation were all unremarkable. Results of routine laboratory tests were within the normal range.

AR-CMT ALG384

Affected patients ALG384-II.1 and ALG384-II.2 were examined at the ages of 9 and 7 years respectively. They both showed a distal pronounced atrophy of the upper and lower limbs with finger retraction and claw hands (fig 2C, D, and A, B respectively). Motor deficit was predominantly distal with foot drop and severe steppage. Hypotonia and areflexia

Figure 2 Distal muscular atrophy and finger retraction in patients (A, B) CMT-ALG384.II.2, (C, D) CMT-ALG384.II.1, and (E, F) CMT-LIB.IV.2.
were seen in the four limbs. Lumbar kyphoscoliosis was discrete. No sensitive or vasomotor disturbances were found. Subsequent neurological examination at the ages of 12 and 10 respectively showed extension of atrophy and motor deficit in proximal muscles of the lower limbs, causing difficulties in standing up. Vasomotor disturbances had appeared, with increased sweating and cold extremities. Distal sensory abnormalities were mild. No cranial nerve involvement was apparent, and abdominal ultrasound, echocardiography, electroencephalography, and ophthalmological evaluation were all unremarkable. Results of routine laboratory tests were within the normal range.

**Electrophysiology**

**LIB284-IV.2**

Electromyography performed at the age of 18 months showed the absence of evoked sensory action potentials at median nerves. Evoked motor action potentials were also absent in the tibial and peroneal nerves, and in the cubital and median nerves they had low amplitudes (0.4 and 1.4 mV respectively) with normal conduction velocities (45 and 40 ms⁻¹ respectively). No potentials were obtained through voluntary contraction at tibial nerves whereas a low amplitude tracing with an accelerated pattern was obtained at the lateral right gastrocnemius. Nerve conduction studies and EMG performed on the unaffected parents and brothers were normal.

**ALG384-II.1**

Electromyography performed at the age of 5 was compatible with axonal loss associated with demyelination. Potential amplitudes (0.48 mV), and MNCVs at the median nerve were very low (20 ms⁻¹). Also, SNCVs at the sural nerve were 34.8 ms⁻¹ with amplitude 5.4 mV. Sensory potential amplitudes were 1.7 and 1.9 mV respectively in the median and ulnar nerves. A second EMG, performed at the age of 10, indicated worsening of the neuropathy with progressive axonal loss.

**Electrophysiology**

**LIB284-IV.2**

Electromyography performed at the age of 3 showed MNCVs of 39 ms⁻¹ at the left ulnar nerve, with a 6 mV potential amplitude, indicating a demyelinating process. A second EMG at the age of 8 showed an associated axonal involvement at the sural nerve, with a 31 ms⁻¹ SNCV and a 2.7 mV potential amplitude. Sensory potentials were undetectable in the ulnar nerve. Thus, the EMG results indicated an association of demyelinating neuropathological processes with axonal loss. Nerve conduction studies and EMG performed on the unaffected parents were unremarkable.

**Nerve biopsy results**

Ultrastructural analysis of all patients' nerve biopsies showed prominent, likely chronic, demyelinating features, with a severe reduction of myelinated fibres (fig 3A, C). The large diameter fibres were primarily lacking. The myelin sheaths of most of the few remaining myelinated fibres were abnormally thin although compaction was not modified; some Schwann cells contained one naked axon typical of complete demyelination (fig 3B). Many large onion bulbs were seen around unmyelinated and myelinated fibres (fig 3B, D). They were made of numerous, concentrically arranged, flattened processes of Schwann cell cytoplasm. Unmyelinated fibres were also significantly decreased. These anomalies were characteristic of a typical chronic “demyelinating-remyelinating” process associated with a large reduction of axonal density.

**Molecular genetic analysis**

Involvement of *MPZ* (1q22), *PMP22* (17p11.2), *EGR2* (10q21-q22), and *NDRG1* (8q24) was excluded by direct sequencing. Because the affected patients were born to consanguineous parents, we assumed identity by descent and used a homozygosity mapping approach for the other genes reported as being involved in demyelinating AR-CMT: *MTMR2* (11q22), *PRX* (19q13.3), and *GADAP1* (8q21.3). The
segregation analysis excluded linkage to chromosomes 11q22 and 19q13.3 whereas the segregation of haplotypes at chromosome 8q21.3 was compatible with linkage of the disorder to this region in both family CMT-LIB284 and family CMT-ALG384. Genotyping analysis showed positive although not significant linkage to chromosome 8q21.3, from D8S551 to D8S548 (Zmax = 0.25 at θ = 0 for both markers) in family CMT-LIB284 (data not shown). Only the affected people were homozygous for these markers, with alleles that were not shared by the two families; we thus continued screening for mutations in the coding region and splicing sites of the \textit{GDAP1} gene. Sequencing showed a single nonsense mutation lying in \textit{GDAP1} exon 5 carried by patient LIB284-IV.2 (c.668T>A, fig 4) and a mutation in the acceptor splicing site of intron 3 (IVS3–2A>G) carried by the sibs ALG384-II.1 and ALG384-II.2 (fig 4). The c.668T>A change is predicted to lead to the transition from leucine to a premature stop codon at position 223 of the protein product (p.L223X). The splice score consensus values of the IVS3–2A>G genomic sequence versus the wild type were, respectively, 78.66 and 94.69, with a 0.83 mutant/wild type score ratio, indicating a reduced splicing efficiency for the IVS3–2A>G mutated site. Transcripts of patient ALG384-II.1 were subsequently analysed at the mutated splicing site. RT-PCR amplification showed a 108 bp fragment in patient ALG384-II.1 instead of the 203 bp expected, which was present in the normal control and in patient LIB284-IV.2 (fig 5A). Sequencing of the RT-PCR products showed that patient ALG384-II.1 expressed a unique, aberrant transcript missing exon 4 (c.485_579del) (fig 5D).

In both families the segregation of the mutated alleles was consistent with the autosomal recessive inheritance of the disorder.

**DISCUSSION**

\textit{gdap1} was first identified as one of 10 cDNAs (\textit{gdap1–10}) highly expressed in a differentiated Neuro2a mouse neuroblastoma cell line, in which cholinergic differentiation with neurite sprouting had been induced by transfection of GD3 synthase (α2,8 sialyltransferase) cDNA. Transfection of GD3 synthase cDNA in Neuro2a cells has been shown to induce the expression of GD3 as well as GQ1b and GD1b, which are b series complex gangliosides (glycosphingolipids containing sialic acid). Ganglioside specific sialyltransferases as well as gangliosides are highly enriched and specifically localised in

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**Figure 4** Genomic normal (+/+), heterozygous (+/-), and homozygous (-/-) mutated sequences in families CMT-ALG384 and CMT-LIB284.
the vertebrate nervous system,\textsuperscript{20–22} where they have been involved in signal transduction pathways leading to neural differentiation.\textsuperscript{20–22}

gdap1 expression increases greatly during Neuro2a neural differentiation induced by GD3 synthase; in parallel, its expression pattern seems to be developmentally regulated in mice.\textsuperscript{19} being maximal and specific in adult mouse brain. \textit{GDAP1}, the human orthologue, is also mostly expressed in central and peripheral nervous tissues.\textsuperscript{1} It is expressed in neurons or in Schwann cells, as it has been isolated from human sural nerve, mouse sciatic nerve,\textsuperscript{5} and cauda equina,\textsuperscript{4} and is likely to play a fundamental part during development and myelination of the peripheral nervous system.

\textit{GDAP1} encodes a 358 aa protein, containing two predicted transmembrane helices (aa 292–311 and 319–343). In parallel, phylogenetic and secondary structure analysis has shown that GDAP1 belongs to a new and probably monophylegetic group of glutathione S-transferases. Amino acid residues 26–119 (N-terminal subdomain) show homology with the thioredoxin-like fold domain binding glutathione and amino acid residues 210–287 (C-terminal subdomain) show homologies with the domain recognising xenobiotic substrates.\textsuperscript{5}

Structure and expression patterns of GDAP1 locate this protein at the “crossroads” between gangliosides and glutathione S-transferases in terms of diverse and possibly linked functions.

Because of its primary structure, GDAP1 is likely to be localised at the cellular membrane, where it might participate in the complex and permanent interactions existing between Schwann cells and neurons.\textsuperscript{26–27} It should be investigated, for instance, whether GDAP1 is included in the membrane glycosphingolipid enriched signalling domain of Neuro2a cells. This domain contains the signal transducer molecules c-Src, Lyn, Rho A, and Ha-Ras and is likely to participate in signalling pathways leading to differentiation and neurogenesis through the enhancement of MAPK activity.\textsuperscript{27}

In parallel, proteins belonging to the GST multigenic family are continuing to be discovered as having many diverse functions besides the detoxification of electrophilic compounds through glutathione conjugation and maintenance of the cellular redox state, for example, transactivation of transcription factors,\textsuperscript{30} composition and regulation of ionic channels,\textsuperscript{31} and activity as recognition molecules.\textsuperscript{32}

GDAP1 might thus have diverse possible functions in the development of the peripheral nervous system, which remain to be elucidated.

Whether GDAP1 is physiologically involved in signal transduction, cell-cell recognition, neurotransmitter release modulation, or cellular detoxification, the mutated proteins (that of LIB284-IV.2 is predicted to be truncated at aa 223, which is localised in the GST-xenobiotic recognising domain, and those of ALG384-II.1 and ALG384-II.2 are predicted to be missing amino acids 162–193, which are GDAP1 specific) are likely not able to ensure the proper communication between Schwann cells and axons, giving rise to the severe, early onset CMT phenotype found in the patients reported on here.

To date, only GDAP1 mutations causing recessive CMT have been reported; the mutations thus do not seem to act in a dominant negative fashion and half gene dosage seems sufficient to preserve a normal peripheral nerve function. It is worth noting that all patients carrying a mutation in \textit{GDAP1} do not show central nervous system involvement,\textsuperscript{45–46} making it

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**Figure 5** Exon 4 skipping in \textit{GDAP1} transcripts from patient ALG384-II.1. Agarose gel electrophoresis of RT-PCR fragments performed between [A, B] \textit{GDAP1} exons 3 and 5 and \textit{GAPDH} exons 7 and 8 as a control. (C) Sequence of exons 3–4 and 4–5 junction fragments in control LIB284-IV.2 and (D) of exons 3–5 aberrant junction fragment in patient ALG384-II.1. (E) Sequence of the sense and antisense primers used to amplify cDNAs between exons 3 and 5.
likely that a certain redundancy of function exists between GDAP1 and other proteins in this tissue. Several findings suggest that different signalling pathways are likely to be involved in myelin processes in the peripheral versus the central nervous system."

Peripheral nerve ultrastructure, together with EMG results in the patients described here, suggest that GDAP1 mutations affect myelin sheaths as well as axons. Other reports have shown that demyelination and axonal loss often coexist in Charcot-Marie-Tooth disease, axonal degeneration probably being secondary to the disruption of Schwann cell-axon interactions and probably being the main cause of clinical disability.

Animal models, along with more detailed functional studies in humans, will help to understand further the role of GDAP1 and its interaction partners in the complex molecular interactions underlying the physiological development as well as the pathophysiology of the peripheral nervous system.

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