Ataxia-telangiectasia
A new remitting form with a peculiar transcriptome signature

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Abstract

Objective
Ataxia-telangiectasia (AT) is a rare, severe, and ineluctably progressive multisystemic neurodegenerative disease. Variant AT phenotypes have been described in patients with mild- and late-onset neurologic deterioration and atypical features (dystonia and myoclonus). We report on the clinical characteristics and transcriptome profile of patients with a typical AT presentation and genotype who experienced an unexpected favorable course.

Methods
A 24-year-old woman developed, by the age of 3 years, all the classic symptoms of AT associated with increased alpha-fetoprotein levels, a compound AT-mutated (ATM) genotype with an inframe deletion c.2250G>A (p.Glu709_Lys750del42) and a missense mutation c.8122G>A (p.Asp2708Gln), and no residual ATM protein expression. By the age of 12 years, ataxia slowly disappeared, and a very mild choreic disorder was the only neurologic feature in adulthood. Brain MRI was normal. The blood transcriptome profile was assessed and compared with that of healthy controls and patients with the classic AT phenotype.

Results
The atypical clinical course of the patient was associated with a transitional transcriptome profile: while 90% of transcripts were expressed as in patients with the classic AT presentation, 10% of transcripts were expressed as in healthy controls.

Conclusions
The unexpected mild clinical outcome and transcriptome profile of this patient with AT suggest the existence of individual resilience to the altered ATM synthesis. Because of their possible prognostic and therapeutic implications, the identification of modifier factors affecting the phenotype would deserve further studies.
Ataxia-telangiectasia (AT; OMIM#208900) is a rare genetic disease caused by mutations in the AT-mutated (ATM) gene encoding PI3 kinase, which controls the cell cycle and DNA repair. Patients with classic AT present with early-onset progressive cerebellar ataxia, oculocutaneous telangiectasias, immunodeficiency, late-onset peripheral neuropathy, and higher incidence of infections and tumors. Patients usually use a wheelchair by the age of 10 years. Exitus occurs by the second or third decade of life because of chronic lung disease or malignancies. Biomarkers of the disease are increased alpha-fetoprotein (AFP) (95% of patients), chromosomal radiosensitivity, and undetectable ATM protein (98% of patients). Milder variants (late-onset, slowly progressive ataxia/dyskinesia syndrome) have been associated not only with missense ATM mutations and residual ATM kinase activity but also with the classic severe genotype and absent ATM protein.

Here, we report on a new clinical variant with an early progressive course, late remission, and stable neurologic status until adulthood, despite the classic ATM genotype and absent ATM protein. Accordingly, blood transcriptome in this case showed a pattern of alteration intermediate between healthy controls and severely affected patients.

Case report
A 24-year-old woman was born after a normal pregnancy and delivery from healthy unrelated parents. Psychomotor development was normal until the age of 16 months when trunk swaying was noticed by the parents. On examination, she was found to have mild trunk ataxia, external beating nystagmus, mild conjunctival angioectasias, and subcutaneous angiomas in the lumbar and calf regions. Brain MRI and sensory and motor nerve conduction velocities were normal. Blood immunoglobulin A (IgA) levels were marginally decreased (78 mg/100 mL; r.v. 85–450), whereas the AFP level was increased (37.5 ng/mL; r.v. 0–10). Chromosome analysis revealed a 7;14 rearrangement, whereas the serum IgA level was normal.

In the following years, trunk ataxia worsened and other neurologic features, such as motor impersistence and chorea of the trunk and limbs, slurred speech, and hypometric saccades, emerged. Nevertheless, the ability to walk autonomously was preserved, and she could attend a normal school and conduct a normal life. Mental development was normal: at the age of 7 years, the Wechsler Intelligence Scale for Children-Revised IQ score was 92. Starting from the age of 12 years, trunk ataxia progressively disappeared. On examination, at the age of 24 years, she presented with only mild clumsiness associated with nondisabling choreic movements of the limbs (video, links. lww.com/NXG/A47). The International Cooperative Ataxia Rating Scale score was 2 (normal 0). Brain MRI was normal. She maintained normal mental functioning (Wechsler Adult Intelligence Scale IQ score 110, Verbal IQ score 104, and performance IQ score 79) with adequate personal and social skills. She never had pulmonary infections; MRI of the lung performed at the age of 21 years was normal. Her menses were irregular, and she had polycystic ovarian syndrome. She underwent surgical removal of ameloblastoma of the mandible. Recently, ultrasonography has revealed a fatty liver disease in the absence of dyslipidemia or abnormal liver enzymes. At the age of 22 and 24 years, the AFP level was 115 and 190 ng/mL, respectively; the serum IgA level was normal.

Transcriptome analysis: the study of a whole gene expression signature (appendix e-1, links.lww.com/NXG/A46) showed that 1,326 probes were differentially expressed (upregulated and downregulated, table e-1, links.lww.com/NXG/A43) in 5 patients with classic AT compared with healthy controls. Using these probes in hierarchical clustering computation (figure 1), our case resulted as a nonclustered sample: 90% of probes were expressed as in patients with classic AT (cutoff 1.2 FC), whereas 10% of them were expressed as in healthy controls. Of 654 gene symbols derived from the differently expressed probes, 43 gene transcripts had an expression level similar to the healthy controls (table e-2, links.lww.com/NXG/A44). Two of them, such as ZCRB1 (a zinc finger RNA binding protein) and THOC3 (a protein involved in splicing) showed a biological and molecular functional interaction when analyzed by Reactome FI [1] (plugin for Cytoscape).

Patient consent
The patient provided informed consent for participating in the research and publishing the resulting data. She also provided consent to disclose of any recognizable person in the video.

Discussion
We describe an unusual AT phenotype, characterized by a clinical presentation mimicking the classic severe form with the typical biomarkers of this condition, such as absent ATM protein, chromosome rearrangement, and increasing levels of AFP, with an unexpected favorable course of neurologic disorders during teenage years.

The ATM protein level and residual kinase activity seem to influence the severity of the clinical phenotype in patients with AT. Variant AT has been documented in patients

**Glossary**

AFP = alpha-fetoprotein; AT = ataxia-telangiectasia; ATM = ataxia-telangiectasia mutated; IgA = immunoglobulin A.
presenting with a mild neurologic phenotype, often normal brain MRI, and less frequent extraneurologic involvement. In these cases, regulatory, missense, or leaky splicing ATM mutations and a residual ATM activity were detected. The genotype of our patient is characterized by an inframe deletion and a missense mutation without residual ATM activity. Accordingly, a classic presentation and outcome should have been expected. Recent reports showed that absent ATM activity and high AFP may be associated with a late-onset or atypical neurologic presentation (table e-3, links.lww.com/NXG/A45). Of interest, 3 patients belonging to the Canadian Mennonites with late dystonia/myoclonus-dystonia syndrome associated with 6200C>A mutation had transient ataxia in early childhood. The same genotype has been reported in a patient with early-onset myoclonus-dystonia, high blood AFP, and progressive cerebellar atrophy.

In contrast to these cases, our patient experienced the classic presentation and course of the disease until the end of the first decade of life when progressively ataxia vanished, leaving a mild and stable choreic disorder. Nevertheless, AFP and some systemic manifestations of AT (such as ovarian polycystic
The 2 most important genes expressed were *scrb1* and *thoc3*, which are involved in wound reparation and genome stability, respectively. Thoc3 is part of the THO complex, which plays a role in transcriptional elongations, nuclear RNA export, and genome stability.

The finding of a transitional transcriptome profile suggests the occurrence of modifying factors possibly influencing individual vulnerability and resilience to the altered ATM synthesis resulting in an unexpected mild outcome.

**Author contributions**

Vincenzo Leuzzi and Daniela D’Agnano: drafting/revising the manuscript, analysis and interpretation of clinical and biochemical data, and acquisition of data. Michele Menotta: drafting/revising the manuscript, analysis and interpretation of biochemical data, and acquisition of data. Caterina Caputi: drafting/revising the manuscript, analysis and interpretation of clinical and biochemical data, and acquisition of data. Luciana Chessa: drafting/revising the manuscript, analysis and interpretation of clinical and biochemical data, and molecular analysis interpretation. Mauro Magnani: drafting/revising the manuscript, analysis and interpretation of clinical and biochemical data, and acquisition of data.

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**Disclosure**

V. Leuzzi, D. D’Agnano, M. Menotta, C. Caputi, and L. Chessa report no disclosures. M. Magnani has served on the scientific advisory board of, holds stock/stock options in, and/or receives board of directors’ compensation from EryDel SpA. holds a patent regarding a method for the encapsulation of agents within erythrocytes; and has received research support from EryDel SpA and Fondazione Cassa di Risparmio di Fano. Full disclosure form information provided by the authors is available with the full text of this article at Neurology.org/NG.

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**References**

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