Serum Contactin-1 in CIDP

A Cross-Sectional Study

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Abstract

Objective

To investigate whether serum levels of contactin-1, a paranodal protein, correlate with paranodal injury as seen in patients with CIDP with antibodies targeting the paranodal region.

Methods

Serum contactin-1 levels were measured in 187 patients with CIDP and 222 healthy controls. Paranodal antibodies were investigated in all patients.

Results

Serum contactin-1 levels were lower in patients (N = 41) with paranodal antibodies compared with patients (N = 146) without paranodal antibodies ($p < 0.01$) and showed good discrimination between these groups (area under the curve 0.84; 95% CI: 0.76–0.93).

Conclusions

These findings suggest that serum contactin-1 levels have the potential to serve as a possible diagnostic biomarker of paranodal injury in CIDP.

Classification of Evidence

This study provides class II evidence that serum contactin-1 levels can discriminate between patients with CIDP with or without paranodal antibodies with a sensitivity of 71% (95% CI: 56%–85%) and a specificity of 97% (95% CI: 83%–100%).

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Class of Evidence
Criteria for rating therapeutic and diagnostic studies
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Glossary

AUC = area under the curve; **CNTN1** = contactin-1; **IVIg** = IV immunoglobulin; **ROC** = receiver operating characteristic; **sCNTN1** = serum levels of contactin-1.

Most patients with CIDP show a good response to first-line treatments like corticosteroids or IV immunoglobulin (IVIg). However, several reports have emphasized that patients with CIDP and paranodal antibodies have poor response to first-line treatments. Early identification of these patients is important to guide treatment decisions and prevent long-term axonal damage.

Contactin-1 (CNTN1) is an axonal protein that anchors paranodal myelin in complex with contactin-associated protein 1 (Caspr1) and neurofascin-155 (NF-155).[1] Pathogenic IgG4 antibodies targeting these paranodal proteins are found in up to 10%.[2] The CNTN1 (protein) exists in a soluble form making it good biomarker candidate for paranodal damage.[3] Decreased CNTN1 levels have been found in other demyelinating disorders such as MS and neuromyelitis optica.[4,5] We hypothesize that paranodal injury in CIDP leads to altered serum levels of contactin-1 (sCNTN1) in patients with CIDP with paranodal antibodies compared to CIDP patients without.

Methods

Patients were selected from cohorts from 3 CIDP tertiary referral centers in the Netherlands (Amsterdam), Spain (Barcelona), and the United Kingdom (Oxford). The Amsterdam cohort comprised patients who were included in ongoing prospective CIDP cohort studies (N = 103). The Barcelona (N = 55) and Oxford (N = 30) cohorts comprised nonconsecutive patients who were referred because of suspected antibody-mediated CIDP. All patients fulfilled the definite or probable EFNS/PNS criteria.[6] Samples were collected during different disease stages. In addition, 222 healthy controls were included.

Standard Protocol Approvals, Registrations, and Patient Consents

The study was approved by the local medical ethical committees of participating centers. All patients provided signed informed consent.

Serum Measurements of CNTN1 and Antibodies

Isolated serum was stored at −80°C in each center. sCNTN1 levels were measured centrally in the Neurochemistry Laboratory at Amsterdam UMC on Bio-Plex 200 system (Bio-Rad Laboratories, Veenendaal, The Netherlands) using the Human Magnetic Luminex Assay (LXSAHM; R&D Systems, Minneapolis, MN) according to the manufacturer's instructions. Samples were randomized and analyzed in duplicate in a 6-fold dilution, blinded for the presence of paranodal antibodies. Assay validation is described elsewhere.[4] The intra-assay CV was 3.2%, and measurements with an intra-assay CV

> 15% and outliers were repeated and were excluded if the CV remained >15% (N =1). For sCNTN1 measurements below the LLOQ, a value was assigned of half of the manufacturer's reported LLOQ (i.e., 1.8 pg/mL).

The presence of paranodal antibodies was determined at the time of sampling in Barcelona (for the Amsterdam and Barcelona cohort) and Oxford (for the Oxford cohort) using dedicated cell-based assays and ELISAs against NF-155, NF-186, CNTN1, and Caspr1.[7]

Statistical Analysis

A receiver operating characteristic (ROC) curve with area under the curve (AUC) was used to investigate the discriminatory potential of sCNTN1 for the presence or absence of paranodal antibodies. The Youden index was used to select the optimal cutoff point for the ROC curve. Data were analyzed using R, version 3.6.2.

Data Availability

The data that support the findings of this study are available from the corresponding author on reasonable request.

Results

A total of 188 patients with CIDP and 222 healthy controls were included. One measurement in a patient with CIDP was excluded due to high CV. Clinical data for patients with CIDP can be found in the table. Paranodal antibodies were found in 41 (22%) patients, NF-155 antibodies in 18 patients, CNTN1 antibodies in 13, Caspr1 antibodies in 6, and NF-140 antibodies in 4. Treatment status at sampling is shown in the table.

sCNTN1 in Healthy Controls

In 222 healthy controls (mean age 46 years; SD 14; range 19–98 years), median sCNTN1 levels were 12,470 pg/mL (IQR 10,160–14,440 pg/mL). sCNTN1 levels were not associated with age (r: −0.07 95% CI: −0.20 to 0.06).

sCNTN1 in Patients With and Without Paranodal Antibodies

The figure shows sCNTN1 levels for patients with or without paranodal antibodies. Median sCNTN1 levels were lower in patients with paranodal antibodies compared with patients without antibodies ($p < 0.01$) and lowest in patients with CNTN1 antibodies. All 6 samples that were below the LLOQ were present in this subgroup (figure). Compared with healthy controls, median sCNTN1 levels were lower in patients with CIDP, both in patients without (p:0.04) and in patients with paranodal antibodies ($p < 0.01$). There was no correlation

Table Clinical Data of Patients With CIDP

	Patients without paranodal antibodies (N = 146)	Patients with paranodal antibodies (N = 41)
Age (y), mean (±SD)[a]	60 (±15)	55 (±16)
Male, n (%)[b]	101 (69%)	32 (80%)
Clinical phenotype[c]		
Classic, n (%)	121 (84%)	26 (74%)
Acute onset classic, n (%)	14 (10%)	10 (29%)
Asymmetric, n (%)	15 (10%)	0
Sensory predominant, n (%)	3 (2%)	3 (9%)
Motor predominant, n (%)	4 (3%)	2 (6%)
Distal predominant, n (%)	2 (1%)	4 (11%)
Presence of paranodal antibodies		
NF-155 +, n (%)		18 (44%)
CNTN1 +, n (%)		13 (32%)
Caspr1 +, n (%)		6 (14%)
NF-186 +, n (%)		4 (10%)
Treatment status at moment of sampling[d]		
Untreated	53 (37%)	5 (13%)
Receiving treatment	92 (63%)	33 (87%)

Abbreviations: CNTN1 = contactin-1; Caspr1 = contactin-1 associated protein 1; NF-155 = neurofascin-155; NF-186 = neurofascin-186.
[a] Missing for 6 patients.
[b] Missing for 1 patient.
[c] Missing for 7 patients.
[d] Missing for 4 patients.

between sCNTN1 levels and antibody titers at the time of measurement, including in those patients with CNTN1 antibodies (data not shown). For discriminating between the presence and absence of paranodal antibodies, the AUC of the ROC curve was 0.84 (95% CI: 0.76–0.93). The associated optimal cutoff value was 5,810 pg/mL, indicating that lower values had a sensitivity 71% (95% CI: 56%–85%) and a specificity of 97% (95% CI: 83%–100%) to discriminate between CIDP with or without paranodal antibodies.

Classification of Evidence

This study provides class II evidence that serum contactin-1 levels can discriminate between patients with CIDP with or without paranodal antibodies with a sensitivity of 71% (95% CI: 56%–85%) and a specificity of 97% (95% CI: 83%–100%).

Discussion

We found that sCNTN1 levels were distinctly lower in patients with CIDP with paranodal antibodies and were highly specific

for the presence of paranodal antibodies in patients with CIDP. Serum measurements of CNTN1 may therefore aid in diagnosing paranodal CIDP mediated by IgG4 antibodies. Testing for CIDP antibodies is not yet widely available, which can delay diagnosis, whereas sCNTN1 level testing is simple, cheap, and fast to perform. The high specificity of sCNTN1 in combination with the low overall prevalence of paranodal antibodies (estimated at 10%) indicates that normal sCNTN1 values may have a high negative predictive value and may therefore by used as a screening assay preceding focused assays testing for the presence of specific paranodal antibodies in patients with CIDP with a clinical picture suggesting the presence of paranodal antibodies. Also, low or undetectable sCNTN1 may guide early treatment choices as traditional first-line treatments are frequently ineffective in paranodal CIDP mediated by IgG4 antibodies.[8]

Other studies in demyelinating disorders of the CNS also found decreased sCNTN1 levels in serum, although not as low as seen in patients with CIDP with paranodal antibodies.[4,5] Although we found a significant difference on group level between patients with CIDP without paranodal antibodies and healthy controls, the difference was small with considerable overlap in sCNTN1 levels between patients with CIDP without paranodal antibodies and healthy controls. It is therefore unlikely that sCNTN1 can be used to confirm the diagnosis of CIDP in general. Longitudinal studies during various disease stages are needed to further study the potential role of sCNTN1 as biomarker of disease activity in patients with CIDP with and without paranodal antibodies.

In this study, there was uneven recruitment of patients across different states of disease activity reflected that could partly contribute to the results. The lack of patients with acute neuropathies such as the Guillain-Barre syndrome can also be seen as limitation of this study. Also, in the subgroup of patients with paranodal antibodies, there is a risk of selection bias due to nonconsecutive recruitment in some of the cohorts. We cannot rule out interference with our assay by CNTN1 antibodies binding to the antigen at the same epitope as the assay antibodies as an explanation for the low and unmeasurable sCNTN1 levels seen in this subgroup. We presume that assay interference is not a major factor influencing our results because sCNTN1 levels were also distinctly reduced in patients with other paranodal antibodies and because we did not find a correlation between sCNTN1 and antibody titers.
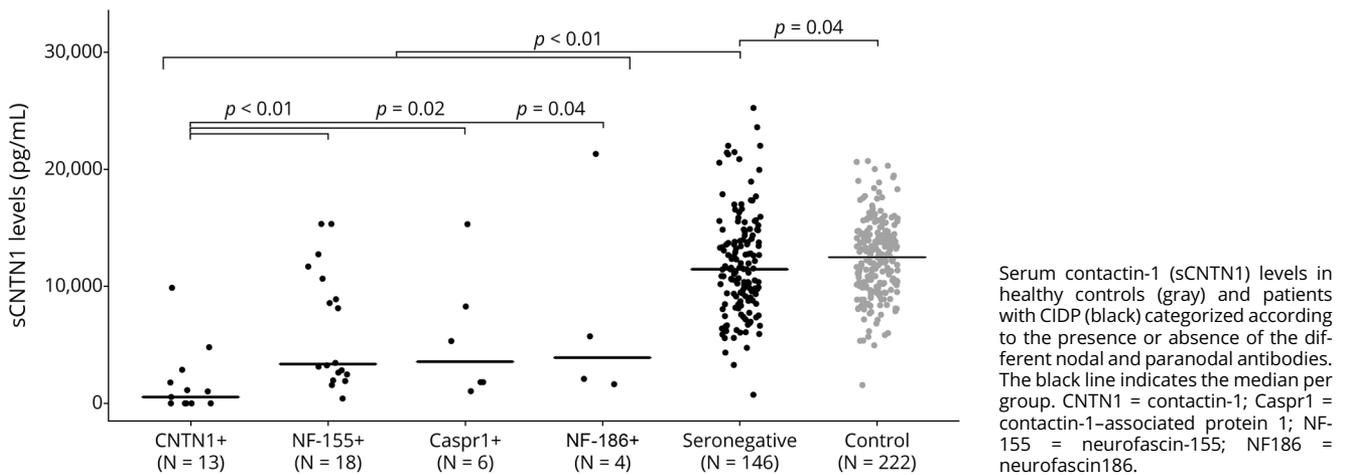
Conclusion

These findings indicate that serum contactin-1 level is a promising new diagnostic biomarker of paranodal injury in CIDP.

Study Funding

Figure Serum Contactin-1 Levels in Patients With CIDP With or Without Paranodal Antibodies

Serum contactin-1 (sCNTN1) levels in healthy controls (gray) and patients with CIDP (black) categorized according to the presence or absence of the different nodal and paranodal antibodies. The black line indicates the median per group. CNTN1 = contactin-1; Caspr1 = contactin-1–associated protein 1; NF-155 = neurofascin-155; NF186 = neurofascin186.

Disclosure

Publication History

Appendix Authors

Name	Location	Contribution
Luuk Wieske, MD, PhD	Amsterdam UMC, location AMC, Amsterdam, the Netherlands	Designed and conceptualized the study; major role in the acquisition of data; analyzed the data; and drafted the manuscript for intellectual content

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