ORIGINAL ARTICLE

Niemann–Pick type C disease as proof-of-concept for intelligent biomarker panel selection in neurometabolic disorders

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Funding information
This research was supported, and funded, by the National Institute for Health Research Biomedical Research Centre at Great Ormond Street Hospital for Children NHS Foundation Trust and University College London. A.P. and P.G. received funding from Actelion for the study of undiagnosed neurodegenerative disorders. This work was also supported by a Clinical Research Training Fellowship (Action Medical Research and the British Paediatric Neurology Association, GN2465) to A.P. The authors have stated that they had no interests which might be perceived as posing a conflict or bias.

Abstract

Aim: Using Niemann–Pick type C disease (NPC) as a paradigm, we aimed to improve biomarker discovery in patients with neurometabolic disorders.

Method: Using a multiplexed liquid chromatography tandem mass spectrometry dried bloodspot assay, we developed a selective intelligent biomarker panel to monitor known biomarkers N-palmitoyl-O-phosphocholineserine and 3β,5α,6β-trihydroxycholanoyl-glycine as well as compounds predicted to be affected in NPC pathology. We applied this panel to a clinically relevant paediatric patient cohort (n = 75; 35 males, 40 females; mean age 7 years 6 months, range 4 days–19 years 8 months) presenting with neurodevelopmental and/or neurodegenerative pathology, similar to that observed in NPC.

Results: The panel had a far superior performance compared with individual biomarkers. Namely, NPC-related established biomarkers used individually had 91% to 97% specificity but the combined panel had 100% specificity. Moreover, multivariate analysis revealed long-chain isoforms of glucosylceramide were elevated and very specific for patients with NPC.

Interpretation: Despite advancements in next-generation sequencing and precision medicine, neurological non-enzymatic disorders remain difficult to diagnose and lack robust biomarkers or routine functional testing for genetic variants of unknown significance. Biomarker panels may have better diagnostic accuracy than individual biomarkers in neurometabolic disorders, hence they can facilitate more prompt disease identification and implementation of emerging targeted, disease-specific therapies.

Abbreviations: CDH, ceramide dihexoside; DBS, dried blood spot; GBA, glucocerebrosidase; GlcCer, glucosylceramide; NPC, Niemann–Pick type C disease; PPCS, N-palmitoyl-O-phosphocholineserine.

Apostolos Papandreou and Ivan Doykov contributed equally to this work; Kevin Mills, Paul Gissen and Wendy Heywood contributed equally to this work.

Members of the clinical cohort recruitment and characterization group are listed in the Acknowledgements.

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Dev Med Child Neurol. 2022;00:1–8.
Despite advances made in next-generation sequencing, timely diagnosis and subsequent initiation of appropriate treatments remain problematic in many neurological conditions. This is particularly true in the field of rare disorders, where clinical presentations are often insidious, and expertise is only available in a few specialist centres. Genomic DNA sequencing can detect most disease-causing variants, but functional assays and relevant biomarkers are often essential to confirm pathogenicity of genetic changes. Additionally, early detection is paramount for patients with disorders affecting the developing brain or resulting in neurodegeneration in order to achieve the best clinical outcomes. This is more pertinent in the current era of personalized medicine, where an ever-increasing number of novel, targeted, and disease-specific therapeutic approaches are becoming available for neurological diseases that in the past were considered untreatable.

Disease-relevant biomarkers can greatly facilitate timely diagnosis and disease monitoring. However, in cases lacking a single ultra-sensitive and ultra-specific marker (which is the norm for most disorders), intelligent design of a biochemical marker panel could be considered as a way to increase diagnostic specificity and accuracy. In this project we used Niemann–Pick type C disease (NPC) as such an example of a clinically heterogeneous neurodegenerative disorder.

NPC is caused by biallelic mutations in the NPC1 gene (in most cases) or, rarely, NPC2. Its prevalence is reportedly 0.35 to 2.2 per 100,000 live births, depending on the country, but this might be an underestimate in view of failure to reach diagnosis in many cases. At a cellular level, NPC results in accumulation of various lipids including cholesterol and sphingolipids in the endo-lysosomal system. Biomarkers recently established to improve NPC detection include cholesterol oxidation products (oxysterols), the 3β,5α,6β-trihydroxy-cholanoyl-glycine bile acid, lyso-sphingomyelin 509 (now identified as N-palmitoyl-O-phosphocholineserine [PPCS]), and lyso-sphingomyelin. So far, the sensitivity and specificity for most of these biomarkers have been assessed in patients with NPC against typically developing comparison individuals, rather than against a relevant age-matched symptomatic patient cohort with similar manifestations. Hence, for our study we recruited a cohort of paediatric patients with neurodevelopmental and neurodegenerative symptoms and signs, in which NPC was part of the differential diagnosis.

To test the hypothesis that a panel of intelligently selected biomarkers could improve detection of NPC, we combined currently available biomarkers and additional glycosphingolipids, sphingomyelins, and bile acids (predicted to be abnormal on the basis of disease pathology) into one multiplexed dried blood spot (DBS) assay.

What this paper adds

- Intelligent biomarker panel design can help expedite diagnosis in neurometabolic disorders.
- In Niemann–Pick type C disease, such a panel performed better than individual biomarkers.
- Biomarker panels are easy to implement and widely applicable to neurometabolic conditions.

METHOD

Ethics

This study was approved by the National Research Ethics Service in the UK (National Research Ethics Service Committee: London – Bloomsbury, REC reference 13/LO/0168, IRAS project 95005) and Great Ormond Street Hospital Research and Development Audit Department (reference 12CM29). Written informed consent from patients or guardians and family members was obtained in all cases.

Patient ascertainment

Undiagnosed paediatric patients from a single UK paediatric hospital’s neurology and metabolic disease clinics, presenting between March 2015 and September 2016, with a possible neurometabolic disorder, were prospectively recruited. Inclusion criteria firstly included disease of childhood onset (0–18 years). Moreover, patients with the presence of one or more neurological feature (such as developmental delay, neurological regression, movement disorder, gaze palsy, epilepsy, autistic features), with or without other pointers towards an underlying inborn error of metabolism (such as suggestive biochemical markers, neuro-imaging abnormalities, or organomegaly), as identified through routine clinical care, were eligible to participate. Characterization of the clinical phenotype was undertaken by direct clinical examination and case note review. For comparison, a cohort of population norm adults was also prospectively established.

Sample collection

All patients underwent blood sampling for research purposes after informed consent. Blood samples in ethylenediaminetetraacetic acid-containing vials were collected for genomic DNA analysis and sequencing. DBS samples from patients and comparison individuals were also collected for biomarker analysis and stored at −80°C.
Genetic sample analysis

In cases of very suggestive clinical, biochemical, and/or radiological presentations, targeted gene testing was performed in the first instance. If this was negative, or in the absence of features pointing towards a specific underlying genetic diagnosis, gene panel testing, whole-exome, clinical exome, and/or whole-genome sequencing analysis was done, either as part of NHS clinical care or on a research basis. Research next-generation sequencing largely occurred through sample contribution to larger studies such as Deciphering Developmental Disorders or the Genomics England 100,000 Genomes Project. Copy number variants (deletions, duplications) were also investigated by comparative genomic hybridization microarray analysis.

Multiplex bloodspot panel

A complete method description is given in Appendix S1, S2, Tables S1–S5, and Figure S1). In summary, a 6 mm punch out of each DBS sample was added to 100 μL of extraction solution containing internal standards. Samples were briefly incubated and centrifuged at 16000 g to fully submerge. Samples were extracted for 10 minutes in a sonication bath at room temperature and centrifuged for 5 minutes at 16000 g to remove any particulate. Each sample was diluted in a 1:1 ratio with methanol and subjected to liquid chromatography with tandem mass spectrometry analysis. All compounds were standardized to an internal standard (Tables S1 and S5). Kuchar et al. have demonstrated the use of both PPCS and lyso-sphingomyelin help discriminate NPC,21 therefore, we have shown our data as a ratio of PPCS to lyso-sphingomyelin.

Data analysis

Raw liquid chromatography with tandem mass spectrometry data were analysed using TargetLynx version 4.0 (Waters, Milford, MA, USA). Standardized data were exported. GraphPad Prism version 6 (GraphPad Software, CA, USA) was used for comparative analysis. Kruskal–Wallis non-parametric analysis of variance was used to determine statistical significance. Multivariate analysis was done using SIMCA version 15 (Sartorius, Goettingen, Germany) (Table S6).

RESULTS

Patient cohort characteristics

Seventy-five patients were recruited, presenting with a heterogeneous range of symptoms and signs suggestive of neurogenetic, neurometabolic, or neurodegenerative conditions. Additionally, 10 population norm adults (age range 23–48 years) were recruited as a comparison group. Mean and median ages were 7 years 6 months and 5 years 2 months respectively (age range 4 days–19 years 8 months). Both sexes were almost equally represented (35 males, 40 females, ratio 0.88:1). Clinical manifestations are summarized in Figure 1. The most common presenting features were neurodevelopmental delay (61 out of 75), non-epileptic movement disorders such as ataxia or dystonia (38 out of 75), epilepsy (24 out of 75), and neurological regression (14 out of 75).

All patients underwent extensive genetic, biochemical, and/or neuroimaging studies as part of their clinical care. Genetic investigations included whole-exome, whole-genome, or gene panel analysis in all cases (75 out of 75). Eventually a definitive genetic diagnosis was established in 43 cases. Of those, five were diagnosed with NPC (NPC1 mutations), while NPC was sufficiently excluded in the 32 out of 75 undiagnosed patients. Other diagnoses established are detailed in Table S7.

Univariate analysis of multiplex assay biomarkers

Compounds identified in the multivariate analysis were subsequently also subjected to univariate analysis (Figure 3).
Known NPC biomarkers PPCS/lyso-sphingomyelin and $3\beta,5\alpha,6\beta$-trihydroxy-cholanoyl-glycine, when analysed individually, had a specificity of 91% and 96.7% respectively (values between the 2.5th and 97.5th centiles were used as the reference range for each group; Table S8). The PPCS:lyso-sphingomyelin ratio, previously identified as a very specific marker, showed a significant increase in the NPC group compared with population norm adults and paediatric patients, both diagnosed and undiagnosed (Figure 3a). However, three patients from the known mutation group (harbouring mutations in SCN1A, KMT2B, and ABA-T) also demonstrated a PPCS:lyso-sphingomyelin ratio in the NPC range. The $3\beta,5\alpha,6\beta$-trihydroxy-cholanoyl-glycine bile acid was also elevated in the NPC cohort, by 30- to 50-fold compared with the other groups (Figure 3a). However, one patient (patient 509, with a chromosome 7 paternal uniparental disomy, presenting with cholestatic disease in infancy and developmental delay)
also had a level in the NPC range. New biomarkers, used to increase the specificity of NPC diagnosis, included long-chain C24 isoforms of GlcCer. These were increased in NPC-positive cases (Figure 3b); however, some patients from the known mutation group also demonstrated levels within the NPC range. Finally, the multivariate analysis had indicated reductions in isoforms of CDH. As these were likely to represent the natural precursor of the elevated GlcCer isoforms (Figure S1), the ratio of the corresponding isoform product (GlcCer) to substrates (CDH) was examined. Ratios for hydroxylated C24 isoforms (GlcCer-C24-OH:CDH C24-OH) allowed a significant separation of the

**FIGURE 3**  Univariate analysis of Niemann–Pick type C disease (NPC) biomarkers from multivariate analysis comparing unaffected comparison individuals (n = 10), confirmed NPC (n = 5), neurodevelopmental patients with known mutations (n = 30), and undiagnosed patients with unknown mutations (n = 40). Significance determined by Kruskal–Wallis analysis of variance with Dunn’s post hoc test; **p < 0.01, ***p < 0.001, ****p < 0.0001. Patients with known mutations with biomarker values in the NPC range are indicated on the graphs. (a) Known NPC biomarkers N-palmitoyl-O-phosphocholineserine (PPCS) ratioed to lyso-sphingomyelin and the bile acid 3β,5α,6β-trihydroxy-cholanoyl-glycine. (b) C24 isoforms of hexosylceramides implicated in orthogonal projections to latent structures discriminant analysis. (c) Ratio of hexosylceramide isoforms to corresponding dihexosylceramide precursor isoforms improves specificity. Abbreviation: Lyso-SM, lyso-sphingomyelin.
NPC group compared with the others (Figure 3c), with no other patient samples overlapping.

**DISCUSSION**

NPC is clinically heterogeneous, with patients often manifesting non-specific signs early in the disease course. Diagnostic delays often ensue, especially because of the condition’s rarity and limited suspicion among non-specialists, the lack of ultra-specific diagnostic markers, and because many rare neurogenetic and neurometabolic disorders have similar presentations. Our patient cohort reflected this, with recruited children manifesting a wide array of often non-specific neurodevelopmental issues and an insidious disease onset. Even with access to next-generation sequencing, difficulties in establishing a diagnosis of NPC often arise, for example when trying to decipher identified variants of unknown significance. Hence, the development of robust disease-specific markers for diagnosis, accurate disease monitoring, and assessment of efficacy of emerging novel, targeted therapies is crucial.

Until recently, filipin staining of unesterified cholesterol in cultured skin fibroblasts was used as the confirmatory test for NPC. However, this test has many limitations as it is invasive, time consuming, and requires a high level of expertise in the analysis. More recently, plasma oxysterols, PPCS with lyso-sphingomyelin, and the 3β,5α,6β-trihydroxy-cholanoyl-glycine bile acid have shown promise as NPC-specific biomarkers, however, apart from their occasional non-specificity and lack of data about efficacy in accurately monitoring disease progression or response to therapeutic interventions, their levels have not been sufficiently examined in populations of similarly affected patients without NPC with other neurological disorders.

To address the continuous need for improved biomarkers in NPC, we developed a panel of markers that included both established markers and others with levels predicted to be affected by disease pathophysiology. Our final panel included measurements of PPCS, lyso-sphingomyelin, 3β,5α,6β-trihydroxy-cholanoyl-glycine, GlcCer C24-OH, and CDH C24-OH. Other studies have assumed similar approaches of combining biomarker analysis for NPC diagnosis, but using both DBS and serum. We selected performing the assay only on DBS, which has advantages of simplified sample collection, transport, and preservation. Another advantage of using DBS was that most compounds of interest were quickly, easily, and reliably extractable, hence our assay would be associated with low costs and rapid turnaround times. Moreover, this approach also removed the need for multiple laboratories to test single analytes. Importantly, multivariate analysis of the biomarker panel in the multiplex assay significantly increased the specificity in discriminating NPC from non-NPC cases.

Furthermore, we examined whether any other compounds were altered in NPC. Multivariate analysis revealed that, as well as the known biomarkers, an elevated ratio of hydroxylated C24 species of GlcCer to CDH (GlcCer-C24-OH:CDH C24-OH) gave good specificity for NPC. GlcCer has previously been observed to be increased in NPC but its role in the disease is not well studied. It is hydrolysed to ceramide by glucocerebrosidase (GBA) for which there are two forms: GBA1 (lysosomal, usually implicated in Gaucher disease) and GBA2 (membrane-associated). GBA activity is influenced by other lipids such as cholesterol and sphingomyelin, which are affected in NPC, while GBA2 has been shown to regulate endo-lysosomal function in NPC. Hydroxylated isoforms of glycosphingolipids are particularly enriched in biological membranes, and are therefore more likely to be substrates of GBA2 than GBA1. More studies are warranted to determine this new biomarker’s clinical utility and to identify links to NPC pathophysiology.

Univariate analysis of other identified biomarkers distinguished patients with NPC, but with occasional overlap with other patient groups, which suggests that using just one marker can be of limited use. We observed that PPCS/lyso-sphingomyelin was increased in three patients without NPC who had gene defects in SCN1A, ABA-T, and KMT2B (Figure 3a). However, all three of these patients had normal levels of 3β,5α,6β-trihydroxy-cholanoyl-glycine, which eliminated suspicions of NPC (Figure 3a). Moreover, only one patient without NPC (chromosome 7 paternal uniparental disomy presenting with cholestasis) had 3β,5α,6β-trihydroxy-cholanoyl-glycine levels in the NPC range; conversely, PPCS/lyso-sphingomyelin levels were normal; therefore, the use of more than one biomarker in a multiplex assay can greatly reduce false-positive results and improve the specificity of NPC diagnosis.

There were limitations in our work. Our cohort was fairly large, but only five patients with NPC were included and analysed. Our population norm comparison cohort was not age-matched, hence more studies are warranted to ensure that paediatric metabolite values from typically developing children do not overlap with NPC. Patients with Niemann–Pick disease types A and B, as well as NPC1 carriers, should also be included in future studies, as they can exhibit biomarker (e.g. PPCS) values that overlap with those of patients with NPC. Moreover, oxysterols were not included in the analysis owing to expected artefacts in extraction from DBS samples. Nevertheless, our results clearly separate patients with NPC from age-matched patients with similar clinical presentations and provide a paradigm for similar approaches in other monogenic conditions.

In summary, we describe an approach to aiding diagnosis in NPC by designing a biochemically relevant liquid chromatography with tandem mass spectrometry-based biomarker panel, and subsequent multivariate analysis. This tool accurately distinguished patients with NPC within a clinically and genetically heterogeneous cohort of age-matched children presenting with symptoms and signs of neurometabolic disease. This selection of biomarkers could be applicable in the interpretation of NPC1/2 genetic variants, thus expediting diagnosis and subsequent implementation of appropriate treatments. Overall, our results provide proof-of-concept that such methodologies can be useful in
many other neurological and metabolic disorders that, like NPC, are not linked to specific enzyme deficiencies and lack reliable functional assays or single clinical, radiological, or biochemical diagnostic markers.

ACKNOWLEDGMENTS

Members of the clinical cohort recruitment and characterization group are as follows: Matthew P. Wilson, Youssef Khalil, Emma Footitt, Spyros Batziros, James Davison, Lucinda J. Carr, Marios Kaliakatsos, Robert Robinson, Hywel Williams, and Peter T. Clayton.

All research at Great Ormond Street Hospital NHS Foundation Trust and UCL Great Ormond Street Institute of Child Health was made possible by the NIHR Great Ormond Street Hospital Biomedical Research Centre. The views expressed are those of the authors and not necessarily those of the NHS, the NIHR, or the Department of Health.

DATA AVAILABILITY STATEMENT

Most data that supports the findings of this study are available in the supplementary material of this article. Other data that support the findings of this study are available on request from the corresponding author, but are currently not publicly available due to privacy or ethical restrictions.

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**SUPPORTING INFORMATION**

The following additional material may be found online:

- **Appendix S1:** Sample preparation protocol.
- **Appendix S2:** Multiplex compound inclusion criteria.

**Figure S1:** Glycosphingolipid degradation pathway.

**Figure S2:** Orthogonal projections to latent structures discriminant analysis biplot of Figure 2 data.

**Table S1:** Internal standard compound stock solutions.

**Table S2:** Table of mass spectral analysis parameters.

**Table S3:** Gradient parameters 1.

**Table S4:** Gradient parameters 2.

**Table S5:** Mass transitions and conditions for all compounds and internal standards.

**Table S6:** Orthogonal projections to latent structures discriminant analysis parameters.

**Table S7:** Genetic diagnoses established in our patient cohort.

**Table S8:** Individual biomarker statistics.

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How to cite this article: Papandreou A, Doykov I, Spiewak J, Komarov N, Habermann S, Kurian MA, Clinical cohort recruitment and characterization group. Niemann–Pick type C disease as proof-of-concept for intelligent biomarker panel selection in neurometabolic disorders. Dev Med Child Neurol. 2022;00:1–8. [https://doi.org/10.1111/dmcn.15334](https://doi.org/10.1111/dmcn.15334)