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ORIGINAL RESEARCH ARTICLE

Thermogenic flux induced by lignoceric acid in peroxisomes isolated from HepG2 cells and from X-adrenoleukodystrophy and control fibroblasts

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Abstract

This work analyzes the thermogenic flux induced by the very long-chain fatty acid (VLCFA) lignoceric acid (C24:0) in isolated peroxisomes.

Specific metabolic alterations of peroxisomes are related to a variety of disorders, the most frequent one being the neurodegenerative inherited disease X-linked adrenoleukodystrophy (X-ALD). A peroxisomal transport protein is mutated in this disorder. Due to reduced catabolism and enhanced fatty acid (FA) elongation, VLCFA accumulates in plasma and in all tissues, contributing to the clinical manifestations of this disorder. During peroxisomal metabolism, heat is produced but it is considered lost. Instead, it is a form of energy that could play a role in molecular mechanisms of this pathology and other neurodegenerative disorders.

The thermogenic flux induced by lignoceric acid (C24:0) was estimated by isothermal titration calorimetry in peroxisomes isolated from HepG2 cells and from fibroblasts obtained from patients with X-ALD and healthy subjects.

Heat flux induced by lignoceric acid in HepG2 peroxisomes was exothermic, indicating normal peroxisomal metabolism. In X-ALD peroxisomes the heat flux was endothermic, indicating the requirement of heat/energy, possibly for cellular metabolism. In fibroblasts from healthy subjects, the effect was less pronounced than in HepG2, a kind of cell known to have greater FA metabolism than fibroblasts. Our hypothesis is that heat is not lost but it could act as an activator, for example on the heat-sensitive pathway related to TRVP2 receptors. To investigate this hypothesis we focused on peroxisomal metabolism, considering that impaired heat generation could contribute to the development of peroxisomal neurodegenerative disorders.

KEYWORDS

beta-oxidation, heat, isothermal calorimetry, peroxisomal disorders, very long-chain fatty acid (VLCFA), X-linked adrenoleukodystrophy

1 | INTRODUCTION

Peroxisomes are organelles with complex and differentiated enzymatic activities. The importance of peroxisomes emerged due to the growing understanding that the alterations of specific metabolic pathways, particularly those concerning lipid metabolism, were associated with various peroxisomal disorders, characterized by a variety of genotypes and clinical manifestations, most of which lead to progressive neurodegeneration (De Duve & Baudhuin, 1966; Igarashi et al., 1976; H. W. Moser, 1997).

X-linked adrenoleukodystrophy (X-ALD) is the most common peroxisomal disorder (Bezman et al., 2001; H. W. Moser, 1997). The mutated gene *ABCD1* (Xq28) encodes for the peroxisomal ABC half-transporter (ALDP; J. Mosser et al., 1993).

Because of ALDP dysfunction, the degradation of very long-chain fatty acids (VLCFA) to shorter fatty acid (FA), in a process called peroxisomal beta-oxidation, is impaired, with consequent abnormal accumulation of the VLCFA lignoceric acid (C24:0) and hexacosanoic acid (C26:0) in plasma and all tissues (Singh, Moser, Goldfischer, & Moser, 1984; Tsuji, Ohno, Miyatake, Suzuki, & Yamakawa, 1984). VLCFA accumulation is also partly due to the enhancement of their elongation from shorter activated FA (Kemp, Berger, & Aubourg, 2012; Van den Bosch, Schutgens, Wanders, & J. M. Tager, 1992).

VLCFA are considered the biochemical markers of the pathology. Their abnormal accumulation is ubiquitous, although it mainly affects the nervous system, adrenal cortex, and testis, indicating a secondary alteration in steroid hormone homeostasis (Assies, van Geel, & Barth, 1998; Petroni, Cappa, Blasevich, Solinas, & Uziel, 2004). Other metabolic processes, including oxidative processes, the development of secondary inflammatory pathways, typical of neurodegenerative, and in general long-lasting disorders, contribute to the clinical manifestation of X-ALD (Uttara, Singh, Zamboni, & Mahajan, 2009). Nevertheless, the complete molecular mechanisms leading to the progressive neurodegenerative clinical manifestations of X-ALD are not clearly understood.

During metabolic processes, peroxisomes do not produce ATP but develop heat, which is considered to be dispersed in the cell (Schulz, 1996). Heat is a form of energy that could be useful for cellular maintenance and it might have a role in the pathogenesis of peroxisomal disorders (Petroni, 2013).

To investigate heat/energy expenditure of peroxisomes, we used isothermal microcalorimetry, a sensitive tool that can provide a variety of kinetic and thermodynamic information on complex biological systems and thus can help in the description of complex metabolic steps.

For this purpose, we first tested the procedures in peroxisomes isolated from HepG2 cells, as described (Santos, Ojeda, Garrido, & Leighton, 1985; Watkins, Ferrel, Pedersen, & Hoefler, 1991), and then applied them to fibroblasts obtained from patients with X-ALD and healthy subjects (controls).

Since VLCFA peroxisomal beta-oxidation is impaired in X-ALD fibroblasts, the aim of our research was to analyze the thermogenic flux induced by the VLCFA lignoceric acid (24:0) in peroxisomes

isolated from cultured X-ALD and control fibroblasts to determine if impaired heat generation plays a role in X-ALD lipid metabolism.

2 | MATERIALS AND METHODS

2.1 | Patients and cell culture

Human fibroblasts (control and X-ALD) were used as in a previous study and according to established institutional guidelines (Petroni et al., 2004). In brief, skin biopsies from three males affected by X-ALD were obtained thanks to the kind assistance of Dr. Graziella Uziel (Istituto Neurologico Carlo Besta, Milan) and Dr. Marco Cappa (Ospedale Bambino Gesù, Rome, Italy). The age of the patients was 20 to 45 years; controls were healthy males without metabolic disease and of matching age. The diagnosis of X-ALD was based on molecular genetic testing, clinical examination, brain MRI, and plasma VLCFA.

The standards and the materials for tissue culture and analyses were from Sigma-Aldrich (St. Louis, MO). Fibroblasts were subcultivated and grown in minimum essential medium containing 10% heat-inactivated fetal bovine serum (100 U/ml), 100 µg/ml streptomycin, and L-glutamine, and they were maintained at 37°C, in 95% air and 5% CO₂. Cells were frozen according to standard procedures and kept in liquid nitrogen.

Fibroblasts, used at passages below seven, were thawed and expanded. Cells were used at confluency, which was reached within 1 week. Cells were fed every 2 to 3 days with fresh growth medium. The normal growth was constantly checked under an optical microscope. Protein/lipid measurements and cell counting were performed at confluency.

2.2 | Homogenization of HepG2 cells and fibroblasts from X-ALD patients and controls

HepG2 cells and fibroblasts from confluent 10 cm diameter dishes were harvested by trypsinization, washed with culture medium and then with phosphate buffered saline (PBS). The cell suspensions were centrifuged at 1,000g for 8 min and the pellet was resuspended in PBS and centrifuged at 1,000g for 8 min. The resulting pellet was homogenized in 250 mM sucrose, 1 mM Tris (Cl⁻), pH 7.5, and 1 mM ethylenediaminetetraacetic acid with a Dounce or Potter-Elvehjem tissue grinder, followed by centrifugation at 500g for 5 min. The subsequent steps for peroxisome isolation and evaluation of the peroxisomal marker enzyme catalase were according to Santos et al. (1985) and Watkins et al. (1991). The experiments for calorimetry and VLCFA measurement were performed with protein concentrations ranging from 35 to 400 µg/ml according to Bradford (1976).

2.3 | Lignoceric acid incubation and isothermal titration calorimetry

Lignoceric acid was dissolved in ethanol, dried under N₂ stream, dissolved in alpha-cyclodextrin in 10 mM Tris-HCl and sonicated

according to standard procedures (Watkins et al., 1991). In all the reported experiments, the final concentration of lignoceric acid was 10 μ M.

The preparations, diluted to the chosen protein concentration, were placed in the calorimetry instrument for 2 hr, after which the aliquot of lignoceric acid was added.

Details of the calorimetric method and instrument are reported elsewhere (Fessas & Schiraldi, 2017; Gardikis et al., 2017). Measurements were performed at 37°C. A 5 ml suspension containing a peroxisome preparation with protein concentration ranging from 35 to 40 μ g/ml was loaded in the measurement vessel under bland steering. After a 2-hour instrumental equilibration, lignoceric acid was added by injection to achieve the final concentration of 10 microM within the calorimetric vessel. The error on the overall thermal effect was below 6% based on three replications for each experiment.

2.4 | VLCFA measurement

At the end of the incubations, the preparations were washed with PBS, lipids were extracted according to Folch, Lees, and Sloane Stanley (1957), FA methyl esters were prepared according to Petroni, Blasevich, and Uziel (2003) and injected into a gas chromatograph (DANI GC 1000) with a Supelco Omegawax column (30 m). Further identification was carried out by liquid chromatography-mass spectrometry according to Banni et al. (2004).

2.5 | Statistical analysis

Statistical evaluation was performed by two-way analysis of variance followed by Student's *t* test.

3 | RESULT

3.1 | Isothermal titration calorimetry

We used HepG2 peroxisomes for comparison because they have very active VLCFA beta-oxidation. Figure 1 shows the isothermal titration calorimetry thermograms (heat flow vs. time) of peroxisomes isolated from HepG2 cells and from X-ALD and healthy fibroblasts. Isolated peroxisomes were incubated in independent experiments with lignoceric acid (10 μ M) at the peroxisomal protein concentration of 40 μ g/ml. When peroxisomes were incubated with lignoceric acid vehicle or only buffer without the organelles, no signal was detected by microcalorimetry.

The thermogenic curve from HepG2 peroxisomes showed that heat flux was exothermic, with a consistent amplitude, indicating normal heat production.

In contrast, the curves of peroxisomes from X-ALD fibroblasts, incubated in the same conditions, showed an opposite effect with heat adsorption (endothermic) and a delay in the onset time of heat flux.

The observed endothermic flux reveals the requirement of this form of energy by the organelles and its detection was related to the reduction of heat production in this system. Indeed, the heat flux of peroxisomes isolated from controls showed a possible balance between exothermic and endothermic activity.

The metabolic thermogenic curves of control and X-ALD peroxisomes showed less amplitude than those of HepG2 peroxisomes. This is in agreement with HepG2 cellular characteristics, in that they present elevated lipid metabolic activity, in particular concerning VLCFA beta-oxidation, in comparison with fibroblasts (Watkins et al., 1991).

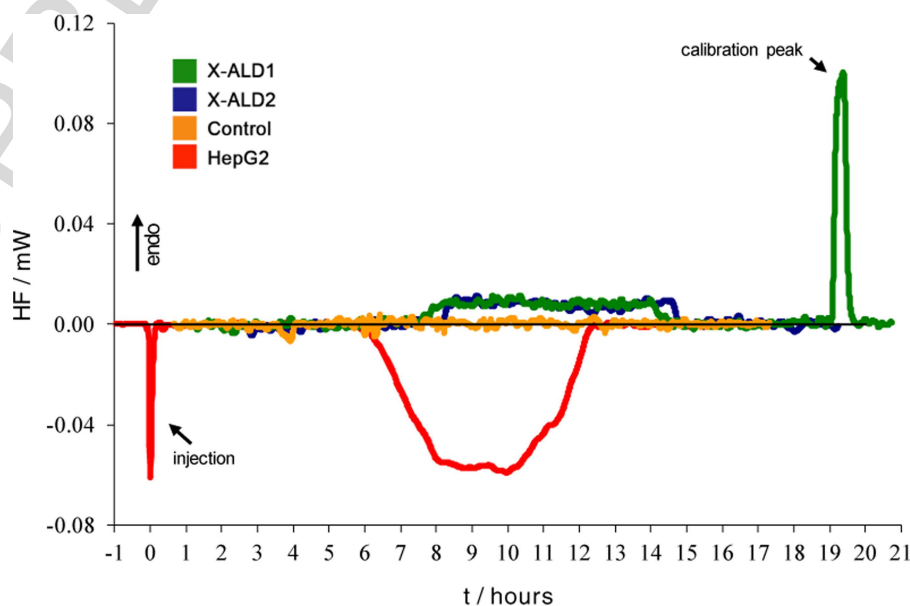


FIGURE 1 Representative ITC thermograms (heat flow vs. time) of peroxisomes isolated from HepG2 cells and from X-ALD and control fibroblasts, at a protein concentration of 40 μ g/ml. X-ALD1 and X-ALD2 represent two independent experiments. The experiments were performed more than three times for each kind of cell. ITC: isothermal titration calorimetry; X-ALD: X-linked adrenoleukodystrophy

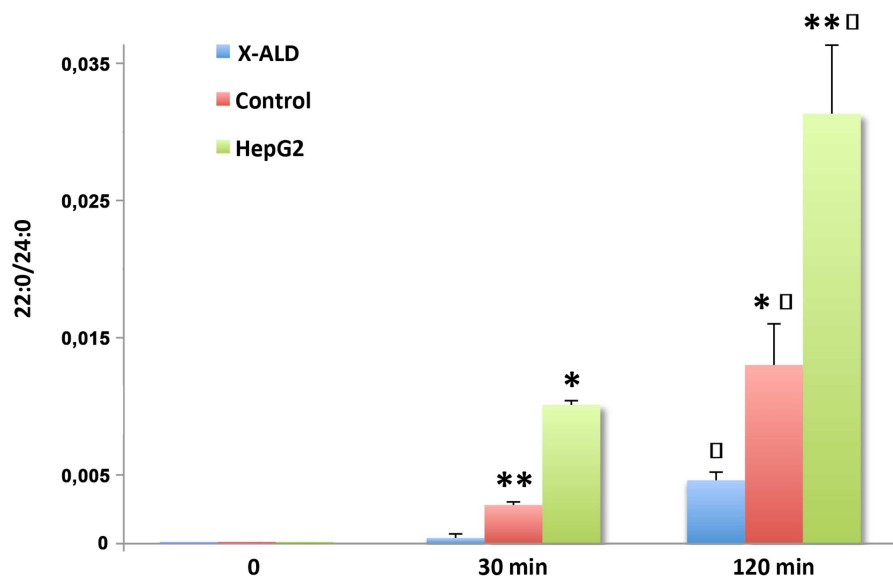


FIGURE 2 The 22:0/24:0 ratio at 30 and 120 min incubation. The columns represent the mean \pm SEM of three measurements. * $p < 0.05$, ** $p < 0.005$, vs. X-ALD of the same time; $p < 0.005$, vs. corresponding cells at 30 min; where not indicated, the difference was not significant. SEM: standard error of the mean; X-ALD: X-linked adrenoleukodystrophy

3.2 | VLCFA measurement

VLCFA beta-oxidation, expressed as 22:0/24:0, was evaluated with the same concentration of agonist, at 200 $\mu\text{g/ml}$ proteins (Figure 2). The ratio was significantly reduced in X-ALD preparations with respect to control and HepG2 organelles.

This is in agreement with the extensive literature, since VLCFA beta-oxidation is impaired but not absent in the X-ALD phenotype.

4 | DISCUSSION

The main result of our study is the endothermic effect induced by lignoceric acid in X-ALD peroxisomes in comparison to controls and HepG2 peroxisomes. This indicates a reduction of heat/energy production and the requirement of energy by X-ALD organelles. The reduction was related to VLCFA beta-oxidation impairment, as shown here and reported in the literature (Igarashi et al., 1976).

In stimulated HepG2 the effect was exothermic, indicating normal heat/energy production, and was more enhanced than in control fibroblasts. This was related to normal VLCFA beta-oxidation, which is more efficient in the HepG2 cell line than in fibroblasts (Watkins et al., 1991).

VLCFA accumulation in all tissues, in particular in the brain, has been considered a major determinant in X-ALD pathogenesis. Their accumulation is due to reduced peroxisomal beta-oxidation and the enhancement of FA elongation. Peroxisomal beta-oxidation could be a heat-dependent system. Heat could sustain this pathway or cellular metabolism in general, influencing for example the velocity or the rate of specific metabolic pathways.

Specific elongases are required for FA elongation. These pathways, enhanced in X-ALD, could require a low level of heat/energy, thus being more facilitated with respect to beta-oxidation metabolism.

We can also speculate that the observed thermodynamic modifications could play a role in the activity of the ABC peroxisomal transporters, affecting dimerization processes of the mutated ALDP and related proteins and mechanisms underlying X-ALD pathogenesis.

Heat is generated in a variety of cellular metabolic pathways, although the energy reserve is rarely considered in biological studies, even though it could modulate different parameters. The heat produced by peroxisomes is considered lost (Petroni, 2013; Schulz, 1996), whereas it could have a role in X-ALD pathogenesis.

5 | CONCLUSION

This is the first publication reporting data on a possible role of heat flux in X-ALD cell metabolism. The molecular mechanisms underlying this disorder and related neurodegenerative disorders are not clearly understood. The extensive literature has demonstrated that the serious dysfunctions characterizing this pathology are partly caused by VLCFA accumulation. However, the modification of heat generation during peroxisomal metabolism should be considered for an understanding of X-ALD and neurodegenerative disorders.

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CONFLICT OF INTERESTS

The authors declare that they have no conflict of interests.

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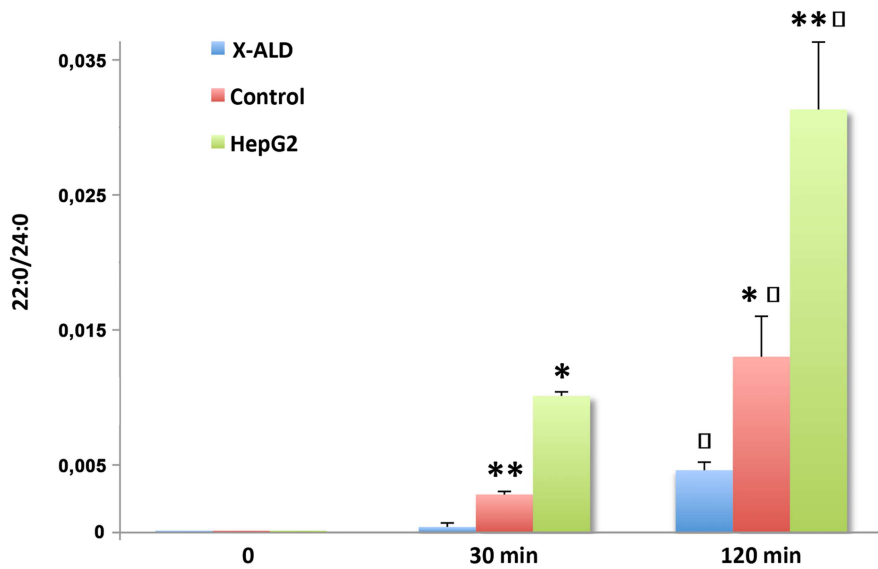
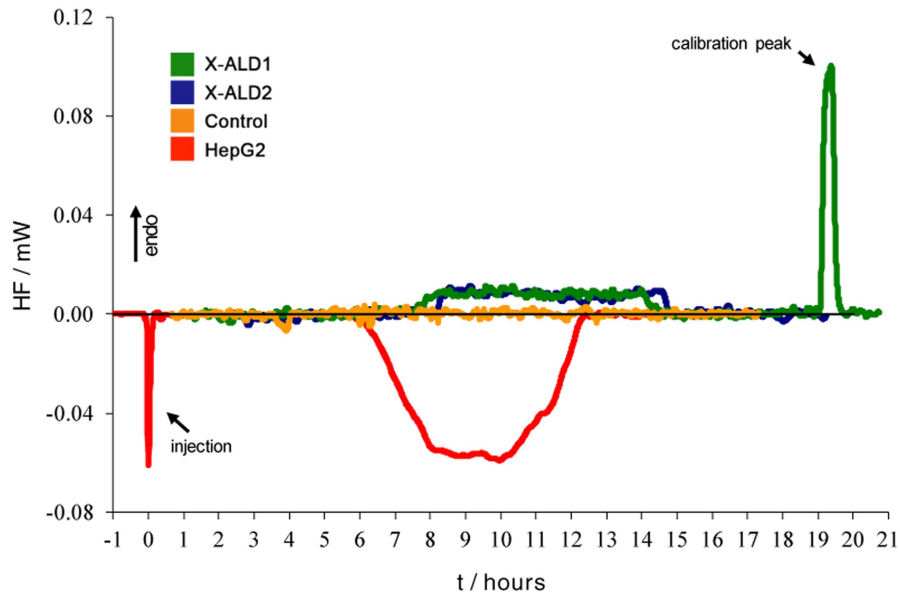
REFERENCES

- Assies, J., van Geel, B., & Barth, P. (1998). Low dehydroepiandrosterone sulphate (DHEAS) levels in X-linked adrenoleukodystrophy. *Clinical Endocrinology*, 49, 691–693.
- Banni, S., Petroni, A., Blasevich, M., Carta, G., Angioni, E., Murru, E., ... Ip, C. (2004). Detection of conjugated C16 PUFAs in rat tissues as possible partial beta-oxidation products of naturally occurring conjugated linoleic acid and its metabolites. *Biochimica Biophysica Acta*, 1682, 120–127.
- Bezman, L., Moser, A. B., Raymond, G. V., Rinaldo, P., Watkins, P. A., Smith, K. D., ... Moser, H. W. (2001). Adrenoleukodystrophy: Incidence, new mutation rate and results of extended family screening. *Neurology*, 49, 512–517.
- Bradford, M. M. (1976). A rapid and sensitive method for the quantitation of microgram quantities of protein utilizing the principle of protein-dye binding. *Analytical Chemistry*, 72, 248–254.
- De Duve, C., & Baudhuin, P. (1966). Peroxisomes (microbodies and related particles). *Physiological Review*, 46, 323–357.
- Fessas, D., & Schiraldi, A. (2017). Isothermal calorimetry and microbial growth: Beyond modeling. *Journal of Thermal Analysis and Calorimetry*, 130, 567–572.
- Folch, J., Lees, M., & Sloane Stanley, G. H. (1957). A simple method for the isolation and purification of total lipids from animal tissues. *Journal of Biological Chemistry*, 226, 131–139.
- Gardikis, K., Signorelli, M., Ferrario, C., Schiraldi, A., Fortina, M. G., Hatziantoniou, S., ... Fessas, D. (2017). Microbial biosensors to monitor the encapsulation effectiveness of doxorubicin in chimeric advanced drug delivery nano systems: A calorimetric approach. *International Journal of Pharmaceutics*, 516, 178–184.
- Igarashi, M., Schaumburg, H. H., Powers, J., Kishimoto, Y., Koloday, E., & Suzuki, K. (1976). Fatty acid abnormalities in adrenoleukodystrophy. *Journal of Neurochemistry*, 4, 851–860.
- Kemp, S., Berger, J., & Aubourg, P. (2012). X-linked adrenoleukodystrophy: Clinical, metabolic, genetic and pathophysiological aspects. *Biochimica et Biophysica Acta*, 1822, 1465–1474.
- Moser, H. W. (1997). Adrenoleukodystrophy: Phenotype, genetics, pathogenesis and therapy. *Brain*, 120, 1485–1508.
- Mosser, J., Douar, A. M., Sarde, C. O., Kioschis, P., Feil, R., Moser, H., ... Aubourg, P. (1993). Putative X-linked adrenoleukodystrophy gene shares unexpected homology with ABC transporters. *Nature*, 361, 726–730.
- Petroni, A. (2013). Peroxisomal heat generation and possible relationship with peroxisomal disorders. *Advances in Biochemistry*, 1(1), 5–6.
- Petroni, A., Blasevich, M., & Uziel, G. (2003). Effects of the testosterone metabolite dihydrotestosterone and 5alpha-androstan-3alpha,17beta-diol on very long-chain fatty acid metabolism in X-adrenoleukodystrophic fibroblasts. *Life Sciences*, 73, 1567–1575.
- Petroni, A., Cappa, M., Blasevich, M., Solinas, M., & Uziel, G. (2004). New findings on X-linked adrenoleukodystrophy: 5alpha-reductase isoform 2 relative gene expression is modified in affected fibroblasts. *Neuroscience Letters*, 367, 269–272.
- Santos, M. J., Ojeda, J. M., Garrido, J., & Leighton (1985). F. Peroxisomal organization in normal and cerebrohepatorenal (Zellweger) syndrome fibroblasts. *Proceedings of the National Academy of Science USA*, 82, 6556–6560.
- Schulz, H. (1996). Beta-oxidation in peroxisomes. In D. E. Vance, & J. Vance (Eds.), *Biochemistry of lipids, lipoproteins and membranes* (pp. 91–93). Amsterdam: Elsevier.
- Singh, I., Moser, A. B., Goldfischer, S., & Moser, H. W. (1984). Lignoceric acid is oxidized in the peroxisome: Implications for the Zellweger cerebro-hepato-renal syndrome and adrenoleukodystrophy. *Proceedings of the National Academy of Science USA*, 81, 4203–4207.
- Tsuji, S., Ohno, T., Miyatake, T., Suzuki, A., & Yamakawa, T. (1984). Fatty acid elongation activity in fibroblasts from patients with adrenoleukodystrophy (ALD). *The Journal of Biochemistry*, 96, 1241–1247.
- Uttara, B., Singh, A. V., Zamboni, P., & Mahajan, R. T. (2009). Oxidative stress and neurodegenerative diseases: A review of upstream and downstream antioxidant therapeutic options. *Current Neuropharmacology*, 7(1), 65–74.
- Van den Bosch, H., Schutgens, R. B., Wanders, R. J., & J. M. Tager, J. M. (1992). Biochemistry of peroxisomes. *Annual Review of Biochemistry*, 61, 157–197.
- Watkins, P. A., Ferrel, E. V., Jr., Pedersen, J. I., & Hoefler, G. (1991). Peroxisomal fatty acid beta-oxidation in HepG2 cells. *Archives of Biochemistry and Biophysic*, 289(2), 329–336.

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X-linked adrenoleukodystrophy (X-ALD) is the most frequent peroxisomal disorder. During peroxisomal metabolism, heat is produced but it is considered lost; instead, it is a form of energy that could play a role in molecular mechanisms. To investigate the heat/energy expenditure of peroxisomes, we used isothermal microcalorimetry, a sensitive tool that can provide a variety of kinetic and thermodynamic information.