High cystine in platelets from patients with nephropathic cystinosis: a chemical, ultrastructural, and functional evaluation

L Olcay, E Erdemli, M Kesimer, Y Büyükasık, H Okur, H S Kalkanoğlu, T Coşkun, Ç Altay

Aim: To investigate the morphology and function of platelets in nephropathic cystinosis (NC).

Methods: Seven patients (mean age, 6.5 years; SD, 20 months) with NC were investigated. Their platelets were examined by transmission electron microscopy (TEM) and the characteristics of the dense granules (DGs) were determined by mepacrine labelling and the uranaffin reaction. Bleeding time, turbidometric aggregation, and luminescence aggregation were studied and intraplatelet cystine was measured.

Results: Increased intraplatelet cystine, primary and secondary aggregation defects, and the absence of ATP release were demonstrated. TEM revealed DGs of various shapes and sizes and lamellary or amorphous cytoplasmic inclusions. Viscous material had been released into the vacuolar spaces and enlarged open canalicular system. Mepacrine labelling revealed that the numbers of DGs/platelet were comparable between the patients and the controls (mean, 2.9 (SD, 0.22) v 3.32 (0.18); p = 0.34). The uranaffin reaction revealed that the numbers of type 1, 3, and 4 DGs were comparable between the patients and the controls, but that there were fewer type 2 DGs in the patients (mean, 8.5 (SD, 1.95) v 17.22 (1.58); p = 0.01). TEM for platelet aggregation revealed a lack of induction and/or defective execution and/or delayed transmission. The patients’ intraplatelet cystine concentrations were higher than the controls (mean, 1.56 (SD, 0.84) v 0.08 (0.01) nmol/mg protein; p = 0.009).

Conclusions: This is the first report to demonstrate raised intraplatelet cystine, abnormal platelet ultrastructural findings, and defective aggregation in NC.

Cystinosis is a disorder with autosomal recessive transmission caused by defective carrier mediated transport of the amino acid cystine across the lysosomal membrane. In the “nephropathic” form of the disease, children develop renal Fanconi syndrome, usually at around 6–12 months of age; and then hypothyroidism, photophobia, chronic renal failure, myopathy, difficulty in swallowing, retinal blindness, diabetes mellitus, male hypogonadism, pulmonary dysfunction, and central nervous system dysfunction.

Transmission electron microscopy (TEM)

Venous blood drawn from the patients was mixed with acid citrate dextrose anticoagulant in a 6 to 1 ratio and platelet rich plasma (PRP) was prepared by centrifugation at 150 × g for 20 minutes at room temperature. After fixation in 3% gluteraldehyde in phosphate buffer, platelets were washed with phosphate buffer. Postfixation with 1% osmium tetroxide was performed and the specimens were dehydrated in a graded series of alcohol and embedded in araldite. Ultrathin serial sections were cut on a Leica Ultracut R ultramicrotome (Leica, Vienna, Austria), stained with uranyl acetate and lead citrate, and examined by a Leo 906 E TEM.

Abbreviations: CRF, chronic renal failure; DG, dense granule; NC, nephropathic cystinosis; PPP, platelet poor plasma; PRP, platelet rich plasma; δ-SPD, delta storage pool disease; TEM, transmission electron microscopy.
Mepacrine labelling of the dense granules

Blood was collected in a plastic tube containing ACD solution. PRP was prepared by centrifugation at 100 g for 10 minutes at 14°C. Mepacrine (1 μM; Sigma, St Louis, Missouri, USA) was added to 1 ml of PRP and the solution was incubated for 20 minutes at 37°C.2,4 Samples were then resuspended in buffer (130 mmol NaCl, 5 mmol KCl, 1 mmol NaH₂PO₄, 24 mmol NaHCO₃, 2 mmol EDTA Na₂, 10 mmol resuspended in buffer (130 mmol NaCl, 5 mmol KCl, 1 mmol NaH₂PO₄, 24 mmol NaHCO₃, 2 mmol EDTA Na₂, 10 mmol NaHCO₃) in 0.2 mol/litre cacodylate buffer for 90 minutes at 700 g. After the third wash, the platelets were combined with 2% osmic acid in distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed twice for five minutes each with distilled water, centrifuged to form pellets, and combined samples were washed two...
remaining platelets was aspirated and discarded and the last layer, which was just above the red blood cells and contained leucocytes, was transferred into another 10 ml centrifuge tube. The platelets were washed and centrifuged twice at 1750 \( g \) for 10 minutes, and the final pellet was resuspended in 0.2 ml distilled water. The leucocyte layer was centrifuged at 1000 \( g \) for five minutes and the supernatant was discarded. The pellet was washed repeatedly with distilled water to haemolyse the red blood cells until a clear leucocyte solution was obtained. The clear leucocyte pellet was resuspended with 0.2 ml distilled water. The suspensions of platelets and leucocytes were sonicated on ice three times for 10 seconds, with cooling intervals of 10 seconds. The suspensions were centrifuged at 15000 \( g \) for 10 minutes. Protein concentrations in the supernatants were determined with microprotein-PR™ kit (Sigma) on a Hitachi F800 automatic. Next, the supernatants were deproteinised by the addition of SERAPREP™ (Pickering Laboratories, Mountain View, California, USA) at a ratio of 1 to 1, an internal standard was added (sarcosine, 10 \( \mu \)mol/litre), and they were centrifuged for 10 minutes at 15000 \( g \). Finally the cystine concentration of the supernatants was measured by ion exchange high performance liquid chromatography with post column derivatisation by ninhydrin (Series 1100 (Agilent Technologies, Palo Alto, California, USA) combined with PCX 3100 post-column reaction module (Pickering Laboratories).

Statistics
The Mann Whitney U test was performed for comparison of the mean granule number/platelet, the cystine concentrations within the platelets and leucocytes, and the different types of uranaffin granules in the patients compared with those of the control group.

RESULTS
Platelet and dense granule morphology evaluated by TEM
Dense granules varied in size and shape, with most being round and some being oval (control, fig 1A; patients fig 1B, C). Some platelets showed features of immaturity, such as those from patient 4, which contained Golgi complex (fig 1C). The platelets of all patients contained a bundle of microtubules around the periphery, which gave them their lenticular shape (fig 1B, C). There were many vacuolar spaces, which were composed of viscous, electron dense material (fig 2A). All of the patients' platelets contained heterogeneous cytoplasmic inclusions of various sizes, which were lamellous or amorphous in nature and were not encircled by a unit membrane (figs 1B, C, 2B). Myelin figures were seen in some of these inclusions (figs 1B, 2B). Interestingly, the same kind of cytoplasmic inclusion was also detected in the cytoplasm of a lymphocyte from patient 3 (fig 2C). Centrifugation caused spontaneous platelet release (figs 1C, 2B) and the cytoplasmic inclusions were seen to be intact in those platelets during spontaneous release. The released material seen nearby the platelets had viscous, amorphous, and lamellary characteristics (fig 2B). The uranaffin reaction on thin section electron microscopy was used to classify the dense granules into types 1–4. Mean (SD) values for the patients versus the controls were: type 1 granules in 40.31% (10.13%) versus 31.2% (4.11%) (p = 0.67); type 2 in 8.51% (1.95%) versus 17.22% (1.58%) (p = 0.01); type 3 in 31.42% (8.45%) versus 37.98% (2.21%) (p = 0.91); and type 4 in 19.73% (3.31%) versus 13.6% (1.86%) (p = 0.11) of all DGs (fig 2D).

DGs evaluated by fluorescence microscopy
The mean (SD) number of fluorescent granules/platelet was 2.9 (0.2) in the patients and 3.3 (0.2) in the controls (p = 0.34).

Aggregation tests
The bleeding time was prolonged in four of the seven patients. Primary or secondary aggregation defects were seen with 2\( \mu \)M ADP in five of the seven patients. By increasing the ADP dose to 10\( \mu \)M in four of these five patients, normal
aggregation was attained in one, and minimal improvement was seen in the other three. Collagen (2 µg/ml) and epinephrine (10 µM) induced platelet aggregation were found to be defective in one of the seven and three of the five patients studied, respectively. The absence of ATP release was confirmed by lumiaggregometry in two of three patients who had a platelet aggregation defect (table 1); minimal improvement was seen after increasing the ATP dose to 10µM.

**TEM evaluation of platelet aggregation**

*Second minute*

Although the platelets of patients 2, 3, 4, and 5 fitted together tightly and had abundant pseudopods, as in normal platelets, those of patient 1 were distant from each other and only a few displayed pseudopods, which were blunted. Whereas normal platelets should have released almost all of their granules by this time, degranulation was seen in almost none of the platelets from patients 1, 3, and 5 and only in a variable number of platelets from patients 2 and 4. Granules were still dispersed throughout the platelet and execution (central clustering) was present in a few (patient 1) to a moderate number (patients 3 and 5) of platelets (fig 3).

*Eighth minute*

Whereas the platelets from patients 2 and 4 still fitted together tightly, as is normal, the interplatelet distance increased in patients 3 and 5 compared with the second minute. The platelets of patient 1 were still remote from each other. Although some (patients 2, 3, 4, and 5) still had pseudopods, the platelets from patient 1 had none. In patients 2 and 4, although degranulation took place in most of the platelets, a few still showed central clustering of granules. Platelets from patients 1, 3, and 5 showed different stages of transformation, some having completely or incompletely degranulated, yielding an enlarged open canalicular system, some still displaying central clustering of granules, and some having granules dispersed throughout (fig 3).

*14th minute*

Although the platelets still fitted together tightly and displayed pseudopods in patients 2 and 4, as is normal, the interplatelet distance increased even more than at the eighth minute in patients 1, 3, and 5. In patients 2 and 4, central clustering of granules was not seen, and most of the platelets were degranulated and displayed an enlarged open canalicular system. In a few, granules persisted throughout the platelet, implying partial degranulation. Some platelets from patients 3 and 5 still displayed pseudopods and were still in different stages of transformation, with most showing granules dispersed throughout the platelet and some having partially or completely degranulated, yielding an enlarged open canalicular system to a varying extent. Pseudopods were lacking and granules were dispersed throughout most of the platelets in patient 1.
Evaluation of intraplatelet cystine concentrations

The mean (SD) intraplatelet cystine concentration was higher in the patients than in the control group (1.56 (0.84) v 0.60 (0.08) mmol/mg protein; p = 0.008). The mean (SD) cystine concentration within the leucocytes was also higher in the patients than in the controls (8.64 (3.81) v 0.11 (0.01) mmol/mg protein; p = 0.008). The mean (SD) protein content of the platelets (164.4 (81.8) v 407.4 (72.2) mg/dl; p = 0.08) and the leucocytes (126.2 (42.7) v 165.4 (24.4) mg/dl; p = 0.29) was similar in the patients and the controls.

Discussion

In cystinosis, free cystine classically accumulates in the lysosomes, cytoplasm, nucleus, mitochondria, and the intracellular spaces of bone marrow in the form of characteristic crystals, granular, fine, dark material, or with no visible form in various cells.

We showed that in NC cystine accumulates in platelets in neither the crystalline nor the darkened material form.

To our knowledge, intraplatelet accumulation of cystine has not been demonstrated before. In addition, intraplatelet cystine concentrations seen here may be lower than the pretreatment values because intraleucocyte values are reduced by treatment with cysteamine. Active pinocytosis and phagocytosis have been suggested to provide the mechanism for cystine accumulation, and this has been demonstrated in polymorphonuclear leucocytes and monocytes, but not in lymphocytes and erythrocytes. The relatively lower concentrations of cystine seen in platelets compared with leucocytes in our patients may reflect the fact that endocytosis and pinocytosis (which take place to supply some of the granule contents) are less active in megakaryocytes and platelets than in leucocytes.

It seems reasonable to assume that the lamellary and amorphous cytoplasmic inclusions in our patients are cystine accumulations. The existence of the same types of inclusions in the lymphocytes of patient 3 and their presence in all of the patients studied is striking. However, we did not investigate the presence of sulfur in these inclusions or in other parts of the platelets using electron probes, although we plan to do so in the near future.

It has been reported that vacuoles and vesicles are parts of the open canalicular system that have become dilated and are not characteristic for a particular disorder. DGs in different stages of transformation and spontaneous platelet release caused by cytocentrifugation are seen normally. It is striking that the microtubule system is intact, giving the platelets their lens-like shape. Some of the platelets displayed features of immaturity, such as the presence of the Golgi complex seen in patient 4, as seen in preleukaemia.

The contrast between the morphological changes and the degree of cystine accumulation in our patients’ platelets suggests that mechanisms other than cystine accumulation (which have not been identified yet) may be involved in the morphological and functional abnormalities of the platelets, as in the proximal tubuli.

Among the seven patients, five had abnormal aggregation tests, four had a prolonged bleeding time (patients 1, 3, 5, and 6) and only two (patients 1 and 6) had a bleeding tendency. These findings are similar to previous reports in that bleeding symptoms and/or abnormal laboratory tests (bleeding time and/or abnormal aggregation tests) are not present in all patients with platelet release defects, so that electron microscopic analysis is the most reliable tool for the diagnosis of platelet defects. Although CRF cannot be excluded as a cause of platelet dysfunction in patients 1 and...
adherent, non-steroidal anti-inflammatory drugs, or antihista-
mines for 10 days before the tests.

Because exogenous ADP is the main stimulator of DGs and to a less extent z granules, the lack of mainly secondary aggregation with ADP stimulation in our patients and the absence of grey platelets in light microscopic examination led us to distinguish between the two release defects involving DGs. The first is delta storage pool disease (d-SPD), in which the numbers of DGs are reduced but central clustering of the granules after stimulation take place normally, but temporarily, and the other is an aspirin-like defect in which granule numbers are normal but central clustering does not occur. This second defect may be hereditary or may be caused by the ingestion of aspirin and other drugs (anti-inflammatory agents, phenothiazines, antihistamines, tricyclic antidepressants), and the execution defect may be a component of aggregation defects in leukaemia and refractory anaemia.

"The aggregation pathology seen in our patients is similar to that seen in leukaemia and refractory anaemia, rather than an aspirin-like platelet defect, because it does not just involve the execution process but also induction and transmission."

We first labelled the platelets with mepacrine, a fluorescent drug that selectively accumulates in platelet DGs, binding to serotonin or 5-hydroxytryptamine, and ATP3-5 to determine the number of DGs precisely, and demonstrated that the numbers of DGs were similar in patients and controls. Ultrastructural evaluation of DGs by the uranaffin reaction in which staining of DG membranes produces different staining patterns in normal and pathological conditions showed that degranulation took place in most platelets later than normal and empty (type 4) granules predominate. After confirming that the DGs of our patients had neither a quantitative nor morphological defect suggestive of d-SPD, we examined the ultrastructural changes that take place during aggregation, to determine whether a functional abnormality existed.

The ultrastructural platelet analysis of our patients revealed different degrees of abnormality at different stages of aggregation. For example, platelets from patients 2 and 4 showed that degranulation took place in most platelets later than in the controls, but a few incompletely degranulated platelets were left after 14 minutes. However, platelets from patients 1, 3, and 5 showed different stages of transformation, a lack or delay of central clustering of granules, and a moderate to high number of degranulated platelets. Adherence established at the second minute of aggregation was found to disappear at the eighth minute, implying intact primary but defective secondary aggregation in patients 3 and 5. The lack of evident shape change and few and blunt pseudopods, in addition to lack of platelet adherence, in patient 1 at the second minute of aggregation is consistent with a primary aggregation defect.

The aggregation pathology seen in our patients is similar to that seen in leukaemia and refractory anaemia, rather than an aspirin-like platelet defect, because it does not just involve the execution process (patients 1, 3, and 5) but also induction (patient 1) and transmission (patients 2, 4 mainly and 1, 3, and 5).

However, the aggregating effect of CRF on both adhesion and execution in patient 1 should also be taken into account. The wide spectrum in the severity of defective aggregation may result from different mutations, which have yet to be defined, in the affected children.

Secretion depends upon active ATP production, and abnormal platelet aggregation has been reported in disorders with abnormal glucose metabolism. The ADP/ATP content of DGs was reported to be both normal and low in execution defects, including aspirin-like platelet defects. Subnormal degradation of metabolic ATP into its metabolites inosine monophosphate and hypoxanthine was reported in aggregation defects (including all phases of induction, transmission, and execution) in leukaemia, unlike d-SPD. We did not have the opportunity to measure ATP or its metabolites. However, the predominance of full DGs (type 1) in our patients, unlike that seen in d-SPD, and the low numbers of semi-empty DGs (type 2), seem to exclude insufficiency of adenine nucleotide content and impaired ATP transport across the granule membrane in platelets from our patients. In our patients' platelets, intracellular cystine accumulation may reduce ATP production by impairing oxidative phosphorylation, and may also reduce both intracellular phosphate and calcium concentrations, just as it does in the kidney cells of patients with NC. If this were the case, adequate messenger molecules would not be generated after interaction of agonists with their receptors, so that contractile mechanisms, centralisation of granules, and release would be partially or completely blocked.

The dilated open canalicular system seen in disorders with execution defects may reflect ineffective reception and transmission of release inducing stimuli and thereby may impair the release mechanism more. Inadequate evacuation may be aggravated by the residual material, which may increase after each secretion attempt.

To our knowledge, this is the first report to demonstrate raised intraplatelet cystine, abnormal platelet ultrastructure, and an aggregation defect in NC. We suggest that platelets should be added to the list of cells in which cystine accumulates and compromises cellular function, and that cystinosis should be added to the disorders that give rise to aggregation defects involving mainly the transmission and execution phases. We think that patients with NC, even if they have a normal bleeding screening assessment, might be at increased risk for bleeding during surgical intervention or traumatic accidents, particularly if they have CRF.

ACKNOWLEDGEMENTS

We are grateful to B Solmar for his valuable technical assistance in the preparation of the ultrastructural samples, to Dr N Erarslan for her kind help in performing some aggregation tests, to Professor G Yildirim for her helpful advice, to Drs A Can, A Olcay, and G Ersoy for their technical assistance in the preparation of the manuscript, and to
our diligent nurse S Kirogil for drawing blood samples. This study was partially supported financially by the METVAK and DOKAV foundations.

Authors' affiliations

L Olcay, Section of Paediatric Haematology, Ankara Oncology Hospital, Demetevler, 06200, Ankara, Turkey

E Erdemli, Department of Histology-Embryology, Faculty of Medicine, Ankara University, 06100, Ankara, Turkey

M Kesimer, H S Kalkanoglu, T Coskun, Section of Nutrition and Metabolism, Department of Paediatrics, Faculty of Medicine, Hacettepe University, 06100, Ankara, Turkey

Y Büyükakş, Section of Haematology, Department of Internal Medicine, Faculty of Medicine, Hacettepe University

H Okur, C Altay, Section of Paediatric Haematology, Department of Paediatrics, Faculty of Medicine, Hacettepe University

REFERENCES


High cystine in platelets from patients with nephropathic cystinosis: a chemical, ultrastructural, and functional evaluation

L Olcay, E Erdemli, M Kesimer, Y Büyükasik, H Okur, H S Kalkanoglu, T Coskun and Ç Altay

J Clin Pathol 2005 58: 939-945
doi: 10.1136/jcp.2005.027177

Updated information and services can be found at:
http://jcp.bmj.com/content/58/9/939

These include:
References
This article cites 23 articles, 5 of which you can access for free at:
http://jcp.bmj.com/content/58/9/939#BIBL

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/