Transiently Transfected Purine Biosynthetic Enzymes Form Stress Bodies

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Abstract

It has been hypothesized that components of enzymatic pathways might organize into intracellular assemblies to improve their catalytic efficiency or lead to coordinate regulation. Accordingly, de novo purine biosynthesis enzymes may form a purinosome in the absence of purines, and a punctate intracellular body has been identified as the purinosome. We investigated the mechanism by which human de novo purine biosynthetic enzymes might be organized into purinosomes, especially under differing cellular conditions. Irregardless of the activity of bodies formed by endogenous enzymes, we demonstrate that intracellular bodies formed by transiently transfected, fluorescently tagged human purine biosynthesis proteins are best explained as protein aggregation.

Introduction

Enzymes have previously been found to organize into intracellular assemblies that may improve their catalytic efficiency or lead to coordinate regulation. For example, the trypanosome glycolytic pathway is organized into a glycosome. Similarly, many such cellular bodies occur naturally, functioning in degradation or storage (e.g., P bodies [1] or actin bodies [2]).

While such functional bodies continue to be found, it is also the case that overexpression of proteins in cells can lead to aggregation [3,4], such as the formation of inclusion bodies. While measuring the localization of green fluorescent protein (GFP)-tagged proteins, we identified a surprisingly large number of punctate bodies that accumulated in yeast cells during nutrient starvation [5]. This led us to further speculate whether such bodies were representative of endogenous, functional assemblies, or accidental or pathological aggregates.

Among >100 proteins forming such bodies, we observed that the de novo purine biosynthetic enzyme phosphoribosyl pyrophosphate amidotransferase (encoded by the yeast gene ADE4, homolog of the human enzyme PPAT) reversibly formed intracellular bodies in the presence and absence of adenine. We initially thought that such bodies might be depots for functional enzymes, and the observation of intracellular bodies associated both with the yeast Ade4 and the human PPAT enzyme suggested the possibility of a functional purine-biosynthetic intracellular body conserved between yeast and humans. In particular, Benkovic and co-workers have identified a cellular body (composed in part of the PPAT enzyme) that they called the purinosome [6], forming in human cell culture in the absence of purines, whose assembly has been shown to be assisted by microtubules and perturbed by casein kinase II inhibitors [7,8] and which may be under GPCR control [9].

However, the possibility remained that the manipulation of the gene (via fusion to GFP) or its expression (via transfection and/or starvation) [6] had led to aggregate formation. Therefore, we sought to establish whether the observed punctate bodies [6] were formed by transiently expressing the recombinant enzymes in cultured cells. We have now characterized punctate formation in greater depth and devised explicit tests to determine whether the bodies may have arisen due to non-native protein expression, stress, or aggregation.

Results and Discussion

We studied the mechanism by which transfected genes encoding human purine biosynthetic enzymes might be organized into cellular aggregates, especially under different nutrient conditions. HeLa and HEK293 cells transiently expressing GFP- or RFP-tagged purine biosynthetic enzymes were cultured in purine-poor media. Punctate bodies were observed, resembling those seen by An et al. [6]. While variable penetrance of different purine biosynthetic enzymes was confirmed (Fig. 1A,B), we were surprised to find that this penetrance was largely independent of the presence or absence of purines in the growth medium (Fig. 1A,B).

We speculated that the formation of the punctate bodies might be due to factors other than purine starvation. Previously, the formation of yeast punctate bodies [3] was correlated with protein aggregation potentials (TANGO scores [10]), which suggested the possibility that the recombinant human proteins might be aggregating. Indeed, the fraction of cells in which individual
purine biosynthetic enzymes formed detectable punctate bodies correlated well with the predicted aggregation potentials of the enzymes (Fig. 1C).

If the proximal cause of the formation of punctate bodies was aggregation, then it seemed likely that: (a) such bodies could be induced by cellular stress, and (b) such bodies would contain proteins normally associated with aggregates, such as chaperones and ubiquitin. Indeed, we found that hydrogen peroxide can induce the formation of punctate bodies irrespective of the presence or absence of purines in the medium (Fig. 2). The punctate bodies appeared to be universally associated with the chaperone HSP70, as measured by co-transfection with a plasmid expressing fluorescently-tagged HSP70 (Fig. 3A-I). We similarly observed the chaperone HSP90 to co-localize to punctate bodies (Fig. 3J-L). We confirmed these results for endogenous HSP70, verifying using immunofluorescence that endogenous HSP70 chaperone inhibitor geldanamycin increased accumulation of the punctate bodies (Fig. 5A). However, longer term treatment with low doses of geldanamycin is known to induce HSP70 chaperone production via hormesis [11], and accordingly longer term treatment with geldanamycin not only inhibited the formation of these bodies, but also prevented their induction by stressors, such as hydrogen peroxide (Fig. 5B).

As heat shock proteins are highly multifunctional and interact with many proteins in the cell, we further tested whether punctate bodies might be aggregates by assaying for their association with ubiquitin. Using immunofluorescence, we observed the co-localization of endogenous ubiquitin to the punctate bodies (Fig. 4D-F). In contrast, control experiments employing only secondary antibody or primary antibody targeting the unrelated enzymes GAPDH or glutamine synthetase showed no such co-localization with the punctate bodies (Fig. 4G-O). The association with

Figure 1. The formation of intracellular foci by transfected purine biosynthetic enzymes correlated well with the aggregation potential of the proteins but was not strongly influenced by purine availability. (A) The frequency with which intracellular bodies appeared across the population of transfected cells varied for individual de novo purine biosynthetic enzymes, and did not depend upon the purine content of the cell growth medium, shown here for HEK293 cells cultured using the optimized medium formulations as described in the Methods. Unless otherwise specified, for HEK293 cells assays in this and later figures, 200 ng of DNA were transfected for each construct. Bars in all experiments represent the average and +/- 1 s.d. from at least 3 replicates, counting n = 769, 742, 755, 665, 1225, and 1016 cells, respectively. See Methods for abbreviations of protein names. (B) This trend was similar for HeLa cells cultured in purine-rich medium versus purine-depleted medium, formulated as in [6]. Unless otherwise specified, for HeLa cell assays in this and later figures, 1.6 μg of DNA were transfected for each construct. Bars in all experiments represent the average and +/- 1 s.d. from at least 3 replicates, here counting n = 578, 821, 662, 832, 1519, 732, 674, and 791 total cells, respectively. See Methods for abbreviations of protein names. (C) The frequency with which transfected cells exhibited bodies (shown here for values reported in [6]) correlated strongly with each protein’s predicted aggregation potential [10]. For comparison, the unrelated enzyme GAPDH has a TANGO score of 879.

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Figure 2. The cell stressor hydrogen peroxide (H₂O₂) strongly induced purine biosynthetic enzyme punctate bodies regardless of hypoxanthine (Hx) presence. Base medium is DMEM supplemented with 10% FBS. As indicated, medium was also supplemented with 1 mM H₂O₂ and/or 35 μM Hx as described in Methods. For HEK293 PPAT-EGFP cells, n = 4419, 2652, 3088, 3182 cells per bar. For HEK293 PAICS-RFP cells, n = 4419, 2652, 3088, 3182 cells per bar. For HEK293T PAICS-RFP cells, n = 4612, 3660, 4211, 3760. Bars indicate average +/- 1 s.d. across at least 3 replicates.

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ubiquitin suggested the possible involvement of the ubiquitin-proteasome proteolytic pathway; consistent with this involvement, we observed strong induction of punctate body formation by treatment with the proteasome inhibitor MG-132 (Fig. 6). The induction of punctate body formation by blocking protein degradation suggests that punctate bodies form in cases of excess protein buildup or compromised protein homeostasis, conditions which can lead to intracellular aggregation of proteins. (Interestingly, cell lines from purine salvage enzyme deficient mice are not hypersensitive to proteasome inhibitors [12].) Should the punctate bodies represent large protein aggregates, the bodies would likely be marked for degradation, consistent with their co-localization with ubiquitin. Thus, collectively, these data indicate that the punctate bodies associated strongly with known markers of aggregation, and that their formation is modulated in a manner consistent with aggregation [13,14,15,16].

If the observed bodies resulted from cellular stress and aggregation, then they might be the natural outcome of individual enzyme aggregation. If so, we would predict that other purine biosynthetic enzymes should not co-localize to the same protein puncta. Indeed, only ~1% on average of co-transfected cells showed co-localization of different purine biosynthetic enzymes to the same puncta (Fig. 7, Table S1). As aggregation was proportional to the expression levels of individual enzymes (Fig. 8) and was stimulated by stress (Fig. 2), this limited co-localization likely derived from stress-induced co-aggregation. More generally, co-transfection tended to suppress puncta formation, or individual enzymes aggregated separately (Table S1). We also attempted to co-localize endogenous purine biosynthetic enzymes with protein puncta by use of immunofluorescence with antibodies against endogenous PPAT, PAICS, and TriGART (data not shown). However, we did not find commercial antibodies satisfactory for immunofluorescence imaging, as all showed speckling in immunofluorescent experiments, irregardless of purine availability, and no co-localization with transfected recombinant proteins, arguing that the speckles are immunofluorescence artifacts.

Figure 3. Co-expressed HSP70 and HSP90 chaperones marked purine biosynthesis bodies. Intracellular bodies formed by (A) PPAT-EGFP, (D) FGAMS-EGFP and (G) PAICS-RFP co-localized with co-transfected (B,E) HSP70-RFP or (H) HSP70-GFP, shown here in HeLa cells. (C, F and I) show merged images. (J-L) Intracellular bodies were also often observed to co-localize with co-transfected HSP90, shown here with PPAT-EGFP.

Figure 4. Endogenous markers of aggregated proteins associated with intracellular foci of transfected purine biosynthetic enzymes. (A-C) Endogenous HSP70 and (D-F) ubiquitin co-localized with bodies formed in cells transfected with FGAMS-EGFP, as assayed using immunofluorescence. Immunofluorescence against endogenous glutamine synthetase (G-I) or GAPDH (J-L), which are not markers for protein aggregation, confirmed that these proteins did not co-localize with the bodies. (M-O) Additional control experiments employing only the secondary antibodies (tested for both secondary antibodies and shown here for Alexa Fluor 594-conjugated goat anti-rabbit) exhibited no positive signal and, with the experiments in panels (G-L), ruled out the possibility of non-specific antibody-mediated localization to the bodies.
We saw additional support for the association between punctate bodies and cellular stress from time-lapse microscopy experiments of transfected cells, during which we observed variation both in punctate body dynamics as well as in cell survival (Fig. 9). Microscopy analysis of individual transfected cells over the course of 1-2 hours revealed that transfected cells without punctate bodies exhibited significantly higher survival rates than cells marked by punctate bodies (Fig. 10). This difference in survival rate was largely independent of purine availability, and persisted even when the cell growth medium was exchanged from purine-depleted to purine-rich. The observation that cells marked by punctate bodies died at greater rates than transfected cells lacking such aggregates suggested that these cells likely experienced greater levels of stress, although lacking in an established mechanism and known molecular signals, it is not clear from these data if the punctate bodies were a cause or consequence of that stress.

Conclusions
These results serve as a cautionary tale for interpreting the cellular biology of transfected constructs. Our study encompassed transiently transfected cells expressing recombinant purine biosynthetic enzyme constructs, reproducing the conditions in which the original purinosome body was identified. We did not pursue the arduous task of generating stable cell lines. However, we point out that Goolijarsingh et al. [17] have tested a stably transfected TrifGART HEK293T cell line in which “the levels of GFP fusion protein produced within the cells are substantially reduced and are similar to wild-type levels” and found diffused signal throughout the cytoplasm and an absence of purinosome bodies. Although their cell growth condition of 7.7% FBS does not lead easy
direct comparison with our work or An et al.’s, it still supports our claim that transient transfection or otherwise out of context protein expression may play a role in generating punctate bodies.

In this regard, a consideration of the experiments of An et al. revealed that the purine-deficient medium did not merely lack purines, but differed in multiple other ways, as well (including the base medium, FBS concentration, and removal of nutrients up to 25 kDa), from the purine-rich medium, and the resulting cellular stress may have induced the aggregation of the recombinant proteins. A further indication that their purine-depleted medium may have been stress-inducing is their report that HTB-125 cells did not survive in their purine-depleted medium. These results also raise interesting questions regarding how growth conditions and cellular physiology impact the formation of single protein aggregates in general, especially proteins altered from their native state, for example whose levels may no longer correspond to normal, endogenous levels or which are expressed as fusion proteins. While it may be possible that purinosome bodies may shift to stress body-like protein complexes with increasing levels of protein expression or varying methods of cellular insults, we note that in general, the formation of purinosome bodies did not vary strongly amongst the different cell lines, and at least for PAICS-RFP, we observed the recombinant enzyme to be expressed within two-fold of endogenous levels (Figure S1).

While it is possible that there are indeed endogenous purinosomes—notably, experiments by An and colleagues [6]...
and Barasova and colleagues [18] both present evidence for intracellular foci composed of endogenous purine biosynthetic enzymes as detected by immunofluorescence—these additional data suggest that the formation of punctate bodies in human cell culture following transient transfection of clones expressing purine biosynthesis enzymes can possibly be explained as protein aggregation.

Materials and Methods

Abbreviations of protein names
PPAT, phosphoribosyl pyrophosphate amidotransferase; Trif-GART, the trifunctional enzyme glycaminide ribonucleotide (GAR) synthetase, GAR transformylase, and aminoimidazole ribonucleotide synthetase; FGAMS, formylglycinamidine ribonucleotide synthetase; PAICS, the bifunctional enzyme carboxyaminoimidazole ribonucleotide synthetase and succinylaminomimidazolecarboxamidase ribonucleotide synthetase; ADSL, adenylosuccinate lyase; ATIC, the bifunctional enzyme carboxyaminoimidazolecarboxamidase ribonucleotide transformylase and IMP cyclohydrolase; GFP, green fluorescent protein; RFP, red fluorescent protein; EFGP, enhanced GFP; GAPDH, glyceraldehyde 3-phosphate dehydrogenase; GLNS, glutamine synthetase.

Expression Vectors
The PPAT-EGFP and FGAMS-EGFP expression plasmids were generously provided by An and Benkovic [6]. The ADSL-EGFP expression plasmid was constructed by Gateway cloning into the FGAMS-EGFP expression plasmid, modified to introduce the Invitrogen pDEST47 Gateway cassette in place of FGAMS. All other cDNAs were obtained from the human ORFeome collection (OpenBiome) and cloned using Gateway cloning into either the pcDNA-DST47 plasmid (Invitrogen) for carboxy-terminal GFP-tagged expression clones or the pTagRFP-N plasmid (Evrogen) modified to introduce the Invitrogen pDEST47 Gateway cassette for carboxy-terminal RFP-tagged expression clones.

Cell Culture
To replicate conditions reported by An et al. to give rise to the purinosome bodies, HeLa cells obtained from the American Type Culture Collection (ATCC) were cultured at 37°C with 5% CO2 in the following medium as previously described [6]: Purine rich medium: MEM medium supplemented with 10% FBS (fetal bovine serum) and 50 μg/mL gentamycin sulfate (Invitrogen); purine-depleted medium: RPMI medium supplemented with 5% dialyzed FBS and 50 μg/mL gentamycin. FBS was dialyzed against 100-fold volumes of 0.9% NaCl at 4°C for 2-4 days using a 25 kDa molecular weight cut-off (MWCO) dialysis membrane with daily exchange of the dialysis solution.

As a test of alternate cell lines and matched purine-rich and purine-depleted media formulations, HEK293 or HEK293T cells obtained from ATCC were cultured at 37°C with 5% CO2 in the following media: (1) Optimized purine-rich medium: DMEM medium supplemented with 10% FBS and 35 μM hypoxanthine, or (2) Optimized purine-depleted medium: DMEM medium supplemented with 10% dialyzed FBS. To remove purines but retain other important factors, FBS was dialyzed against 100-fold volumes of 0.9% NaCl at 4°C for 2-4 days using a 1 kDa MWCO dialysis membrane with daily exchange of the dialysis solution [19,20].

One day prior to transfection, HeLa cells were plated in 6-well glass bottom plates in either purine-rich or purine-depleted medium lacking antibiotics. HEK293 or HEK293T cells were plated in 96-well glass bottom plates in the optimized purine-rich medium lacking antibiotics. Plasmids were transfected into cells using Lipofectamine 2000 (Invitrogen) and Opti-MEM reduced serum medium (Invitrogen) according to the manufacturer’s instructions. The transfection medium was replaced with fresh purine-rich or purine-depleted medium 5 hours after transfection. Immediately before imaging, cells were washed once with their respective growth medium. Cells were imaged in their respective growth medium to minimize stress imposed by nutrient shifts. We note that the imaging medium therefore contained phenol red, which could potentially reduce the signal-to-noise ratio, but in practice, we did not experience problems in detecting and visualizing purinosome bodies. Live cells were imaged ~20–24 hours after transfection.

For MG-132 experiments, HeLa cells were plated in 96-well glass bottom plates in DMEM medium supplemented with 10% FBS. Due to the toxic nature of MG-132, we opted to use a less toxic transfection reagent than Lipofectamine 2000, and therefore, we transfected the PPAT-EGFP-expressing plasmid into cells using Fugene HD (Roche Applied Sciences) and Opti-MEM reduced serum medium (Invitrogen) using a 2:4.5 DNA to transfection reagent ratio following the manufacturer’s protocol. The transfection medium was replaced with fresh DMEM+10% FBS about 5 hours following transfection.

Immunofluorescence
For immunofluorescence experiments, cells were fixed with 3.5% methanol-free formaldehyde freshly diluted from 16% stock (28908, Thermo Scientific) at 37°C for 15-20 minutes, blocked with 5% goat serum in PBS-T buffer for 30-60 minutes at room temperature, then incubated with primary antibody overnight at 4°C. Cells were washed with PBS buffer, then incubated with secondary antibody for 1 hour. Primary antibodies used: HSP70 (ab5439, Abcam), ubiquitin (ab7780, Abcam), GAPDH (uc-32233, Santa Cruz Biotech.), glutamine synthetase (uc-9067, Santa Cruz Biotech.). Secondary antibodies used: Alexa Fluor 594-conjugated goat anti-mouse (Invitrogen), Alexa Fluor 594-conjugated goat anti-rabbit (Invitrogen). All antibodies were used at the manufacturer’s recommended concentrations. We additionally tested the protocol using 2% methanol-free formaldehyde fixation or 2% goat serum in PBS-T block at the appropriate steps to find no appreciable differences.

Tested antibodies that proved unsuitable or inconclusive for immunofluorescence included the anti-PPAT antibodies sc-101892 (Santa Cruz Biotech.) and ab71340 (Abcam), the anti-PAICS antibody sc-16150 (Santa Cruz Biotech.), and the anti-TrifGART antibody H00002618-M01 (Novus Biologicals).

Western Blot
For western blotting, cells were grown in DMEM media containing 10% FBS and transfected as described above with 1.6 μg or 200 ng DNA (HeLa or HEK cells, respectively), in regular 96-well plates (HEK293 and HEK293T) or 6-well plates (HeLa). Cells were washed with cold PBS buffer then extracted with Laemmli sample buffer containing 5% beta-mercaptoethanol. Total protein was separated on a 4–15% Mini-PROTEAN TGX precast gel (BioRad) and blotted with PVDF membrane. Primary antibodies used: anti-PAICS (HPA035895, Sigma) and anti-GAPDH (uc-32233, Santa Cruz Biotech.). Secondary antibodies: goat anti-rabbit IgG Fab’(ab’)2-HRP (sc-2037, Santa Cruz Biotech.) and goat anti-mouse IgG Fab’(ab’)2-HRP (sc-3697, Santa Cruz Biotech.). Membranes were scanned with an ImageQuant LAS 4000 (GE Healthcare).
Fluorescent Microscopy of Cells

Live cells were imaged using a Nikon Eclipse TE2000-E inverted microscope inside a chamber maintained at 37°C and 5% CO₂. Fixed cells were imaged at room temperature without CO₂ supplement. Images were acquired using a Photometrics Cascade II 512 camera and Nikon Plan Apo 40x/0.95, 60x/0.95, 60x/1.40, or 100x/1.40 objectives. GFP detection was accomplished using a ET490/20x excitation filter (Chroma Technology), ET525/36 m emission filter (Chroma Technology) and 98/100 bs dichroic (Chroma Technology), and RFP detection was carried out using a ET555/25x excitation filter (Chroma Technology), ET605/52 m emission filter (Chroma Technology) and 89100 bs dichroic (Chroma Technology). The same excitation and emission conditions were used for each GFP variant. We did not detect notable fluorescence emission bleedthrough or photobleaching with these settings. Filter wheels (Sutter Lambda 10-3), motorized stage (Prior H117), and image acquisition were driven by NIS Elements AR Imaging software.

Cell Counts

Cell counts measuring penetration of punctate bodies reflect apparently healthy adherent cells only. Mitotic cells identified as rounded cells with a visible rod-like central structure spanning the cell’s DIC image accompanied by a corresponding decrease of fluorescent signal; see cell m in Fig. 9C and C-DIC for an example) and dead cells were not counted for measuring penetration of punctate body formation. (For the survival rate example) and dead cells were not counted for measuring fluorescent signal; see cell m in Fig. 9C and C-DIC for an example). We note that it is nonetheless possible that only a subset of cell death events—those accompanied by marked visible cellular reorganization—are counted by this assay.) Cells displaying any morphology of punctate bodies were counted, regardless of number of punctate bodies per cell or ranging sizes of those punctate bodies. We observed a spectrum of morphologies which were dynamic; for example, over the course of unperturbed growth, we observed both increases and decreases in the numbers of punctate bodies per cell. Future work may add and characterize the different morphologies and kinetics of punctate bodies.

Drug Treatments

For all drug treatments, drugs were diluted to their final concentrations in prewarmed cell growth medium then added to cells for the duration of the treatment. Hydrogen peroxide was added to cells at a concentration of 1 mM for 0.5-1.5 hours before imaging. For short-term treatment, geldanamycin was added to cells at the indicated concentrations at the time of post-transfection medium replacement, and the cells were incubated for 18 hours. For long-term treatment, geldanamycin was added to cells at the indicated concentrations 16 hours prior to transfection, for a total incubation time of 40 hours. MG-132 was added to cells at a final concentration of 20 μM for the indicated time intervals. Cells were fixed at each MG-132 time point with 3.7% formaldehyde for 15-20 minutes and washed 3 times with PBS before imaging.

Supporting Information

Figure S1 Representative expression levels of PAICS-RFP transfected constructs in comparison to the endogenous PAICS protein. HeLa (A,D), HEK293 (B,E) and HEK293T (C,F), untransfected (U) or transfected with PAICS-RFP (T) under representative growth conditions are probed with anti-PAICS antibody. (D-F) Representative variation of construct expression between independent transfections. Quantified expression level ratios of PAICS-RFP construct to endogenous PAICS (G) across all cells, adjusted with respective transfection rates (ranging from 6-32%) measured by microscopy, show comparable expression (within two-fold) of the construct and endogenous enzyme in transfected cells, under conditions (see Methods) in which PAICS-RFP foci are observed. Expression ratios are calculated as (PAICS-RFP Western blot intensity)/endogenous PAICS Western blot intensity * transfection efficiency. Bars indicate average +/- 1 s. d. across at least 3 replicates. (H) Representative HEK293 cells corresponding to the bar in (G) and showing visible foci. (I) Expression levels of PAICS-RFP are not strongly altered by hydrogen peroxide and/or hypoxanthine treatment for HEK293 (lanes 2-5) and HEK293T cells (lanes 1, 6-9). (TIF)

Table S1 Co-transfection significantly suppresses formation of intracellular punctate bodies, arguing against the bodies representing active purinosomes, as quantified for different purine biosynthetic enzyme pairs, reporting counts and percentages of successfully cotransfected HeLa cells as a function of protein localization.

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Author Contributions

Conceived and designed the experiments: AZ MT ADE EMM. Performed the experiments: AZ JS LC. Analyzed the data: AZ MT LC ADE EMM. Wrote the paper: AZ MT ADE EMM.

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